

# Alberta Bone Marrow and Blood Cell Transplant Program: Standard Practice Manual



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# Indications



# Acute Myeloid Leukemia (AML)

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## Summary

- Disease risk stratification will be based on the cytogenetic and molecular features of the tumour cells, response to first induction, presence of secondary or therapy related disease, white blood cell (WBC) at diagnosis and measurable residual disease when available.
- Patients with favourable cytogenetics and no unfavourable molecular changes show good response to chemo-immunotherapy and in the majority of cases will enter a second remission if relapse occurs. Patients with t(8;21) or inv(16)/t(16;16) without evidence of MRD should undergo allogeneic stem cell transplant in CR2.
- Patients with a normal karyotype who are FLT3 ITD negative and either NPM1 mutation positive or bZIP in-frame CEBP $\alpha$  biallelic mutation positive are expected to have a favourable outcome to chemo-immunotherapy and should be offered an allogeneic stem cell transplant in CR2.
- Patients in the intermediate genetic risk group may be offered a transplant from a matched sibling or a matched unrelated donor in CR1. This includes patients with a normal karyotype as well as non-informative cytogenetic changes. Patients with t(8;21) or inv(16) / t(16;16) and a KIT mutation appear to fall into this risk group. Patients with FLT3 ITD at low allelic ratio and with FLT3 TKD mutation also appear to fall into this risk group.
- Patients with high-risk features will likely not be salvageable at relapse and should be offered transplant in first complete remission. This includes high-risk cytogenetics, those with a normal karyotype who are FLT3 ITD positive, various molecular findings on NGS, those requiring more than one chemotherapy cycle to achieve a complete remission, as well as those with secondary or therapy related disease or measurable residual disease after two cycles of chemotherapy.
- Regarding TP53 mutations, it remains controversial whether HCT should be offered. Many factors need to be considered (Table 4).
- Patients who relapse after conventional chemotherapy should undergo stem cell transplantation in CR2.
- It is preferable for patients to be in complete remission (defined as fewer than 5% blasts and no active extra-medullary disease) at the time of transplantation. Patients with untreated or refractory CNS disease or with circulating blasts are not eligible for transplantation.
- Patients should receive at least one cycle of post-remission therapy prior to transplantation if transplantation cannot occur within 4 weeks of the complete remission being achieved.
- Regarding post transplant maintenance therapy, at present we do not routinely use it.

## Background

Risk stratification in AML has traditionally relied on patient and disease characteristics at diagnosis (chiefly: age, cytogenetics, molecular findings, white blood cell count at diagnosis and the presence of an antecedent hematological disorder or therapy related disease) and on the response to induction

chemotherapy. While patients in favourable risk categories may enjoy long-term disease free survival, AML may be virtually incurable with conventional treatment in patients with high-risk features and those with poor response to chemotherapy. Risk-adapted therapy attempts to avoid exposing favourable-risk patients to the morbidity and mortality risks of stem cell transplant while directing high-risk patients to up-front transplant in order to minimize relapse risk early in the course of therapy. Measurable residual disease (MRD) after induction and/or consolidation chemotherapy is also becoming more reliably prognostic by flow cytometry as well as by quantitative PCR for certain specific genes.

## Prognosis

### Combined Cytogenetic and Molecular Risk Groups

In addition to basic cytogenetic analysis, molecular markers refine prognostic groups.

Patients with normal cytogenetics make up the largest group of patients with AML, yet they show significant variability in outcomes with standard treatment. The likely explanation for this finding is the influence of molecular abnormalities that go undetected by standard cytogenetics. Among these abnormalities, mutations of *NPM-1* and bZIP in-frame mutated *CEBPA* are associated with significantly better overall survival (OS) compared to patients with the wild-type loci. Internal tandem duplications (ITD) to *FLT-3* confer inferior OS on patients who harbor these mutations. Next-generation sequencing is now done routinely in transplant eligible patients and allows for the detection of many other known mutations of potential clinical significance.

AML with myelodysplasia-related gene mutations is categorized in the adverse-risk group. These mutations, typically associated with AML following an antecedent hematologic disease, are also prevalent in de novo AML and indicate adverse risk even in the absence of myelodysplasia-related cytogenetic abnormalities.<sup>6,26,42,44,45</sup> These include mutations in the *RUNX1*, *ASXL1*, *BCOR*, *EZH2*, *SF3B1*, *SRSF2*, *STAG2*, *U2AF1*, or *ZRSR2* genes. Additional disease-defining recurring cytogenetic abnormalities are included in the adverse-risk group, including t(3q26.2;v) involving the *MECOM* gene,<sup>31,73</sup> or t(8;16)(p11.2;p13.3) associated with *KAT6A::CREBBP* gene fusion.<sup>46</sup>

The presence of a pathogenic *TP53* mutation (at a variant allele fraction of at least 10%, with or without loss of the wild-type *TP53* allele) defines the new entity AML with mutated *TP53*. This subtype of AML is associated with a very poor prognosis and the utility of a stem cell transplant may be debated.

Table 1 outlines the risk groups according to the most recent European LeukemiaNet (ELN) classification with Table 2 showing prognosis according to the defined risk groups.

**Table 1.** ELN risk classification 2022

| Risk category† | Genetic abnormality                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       |
|----------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Favourable     | <ul style="list-style-type: none"> <li>t(8;21)(q22;q22.1)/<i>RUNX1::RUNX1T1</i></li> <li>inv(16)(p13.1;q22) or t(16;16)(p13.1;q22)/ <i>CBFB::MYH11</i></li> <li>Mutated <i>NPM1</i>†,§ without <i>FLT3</i>-ITD</li> <li>bZIP in-frame mutated <i>CEBPA</i></li> </ul>                                                                                                                                                                                                                                                                                                                                                                                     |
| Intermediate   | <ul style="list-style-type: none"> <li>Mutated <i>NPM1</i>†,§ with <i>FLT3</i>-ITD</li> <li>Wild-type <i>NPM1</i> with <i>FLT3</i>-ITD (without adverse-risk genetic lesions)</li> <li>t(9;11)(p21.3;q23.3)/<i>MLL3::KMT2A</i>,</li> <li>Cytogenetic and/or molecular abnormalities not classified as favourable or adverse</li> </ul>                                                                                                                                                                                                                                                                                                                    |
| Adverse        | <ul style="list-style-type: none"> <li>t(6;9)(p23.3;q34.1)/<i>DEK::NUP214</i></li> <li>t(v;11q23.3)/<i>KMT2A</i>-rearranged</li> <li>t(9;22)(q34.1;q11.2)/<i>BCR::ABL1</i></li> <li>t(8;16)(p11.2;p13.3)/<i>KAT6A::CREBBP</i></li> <li>inv(3)(q21.3q26.2) or t(3;3)(q21.3;q26.2)/ <i>GATA2</i>, <i>MECOM(EVI1)</i></li> <li>t(3q26.2;v)/<i>MECOM(EVI1)</i>-rearranged</li> <li>–5 or del(5q); –7; –17/abn(17p)</li> <li>Complex karyotype, monosomal karyotype</li> <li>Mutated <i>ASXL1</i>, <i>BCOR</i>, <i>EZH2</i>, <i>RUNX1</i>, <i>SF3B1</i>, <i>SRSF2</i>, <i>STAG2</i>, <i>U2AF1</i>, and/or <i>ZRSR2</i></li> <li>Mutated <i>TP53</i></li> </ul> |

**Table 2.** Prognosis by 2022 European Leukemia Net Risk group in patients receiving intensive induction in the multicentre Beat AML Cohort

| Risk category† | N   | CR/CRi (%) | OS (%) |
|----------------|-----|------------|--------|
| Favourable     | 117 | 86         | ~65    |
| Intermediate   | 92  | 59         | ~35    |
| Adverse        | 136 | 49         | ~20    |

N=number of patients, CR/CRi=complete remission/complete remission with incomplete count recovery, OS=overall survival.

Conventional induction chemotherapy for patients with non-promyelocytic AML consists of combination chemotherapy on a backbone of anthracycline and cytarabine. In patients with a *FLT3* mutation a *FLT3* inhibitor is added. Patients with good risk cytogenetics and *NPM1* mutations as well as certain patients with intermediate risk cytogenetics are offered gemtuzumab ozogamicin along with their chemotherapy. In the setting of therapy-related or antecedent hematologic malignancies patients CPX-351 is substituted. Patients with acute promyelocytic leukemia are offered induction with Arsenic trioxide and ATRA. Azacytidine + venetoclax is an emerging option which could in the future replace anthracycline + cytarabine based chemotherapy.

### Minimal Residual Disease

Despite the above clinical and genetic risk factors present at diagnosis the outcome of individual patients is still highly variable indicating other factors are at play. The detection of measurable residual disease at various time points during therapy likely reflects these yet unexplained factors. Several studies have indicated that undetectable or low MRD values at any time point distinguish patients with more favourable outcomes in terms of relapse-free survival (RFS) and OS than those

with higher values including pre-transplantation. Post two cycles of intensive chemotherapy may be the most informative. The loss of previous MRD negativity is also a potential indication for intervention with transplantation. Molecular MRD testing is routinely done for. How to use this information is currently being investigated with active intervention clinical trials. The main subtypes for which PCR-based detection is routinely available include PML-RARA, RUNX1-RUNX1T1 associated with t(8;21), CBFB-MYH11 associated with inv(16) and variants, and NPM1-mutated AML. FLT 3 ITD testing is currently being investigated in Canada.

Table 3 adds measurable residual disease after 2 cycles of chemotherapy (e.g., 1 induction and 1 consolidation) and other prognostic factors to the cytogenetic and molecular risk stratification to further help with decision on allogeneic stem cell transplantation in first complete remission.

**Table 3.** Cytogenetic and molecular risk stratification including minimal residual disease and other Prognostic factors. (From Cornelissen et al: Blood 2016)<sup>59</sup>

| AML risk group‡ | AML risk assessment criteria at diagnosis                                                                                                               | MRD after cycle 2    | Risk of relapse following consolidation approach |               | Prognostic scores for RM that indicate alloH SCT as preferred consolidation |              |              |
|-----------------|---------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------|--------------------------------------------------|---------------|-----------------------------------------------------------------------------|--------------|--------------|
|                 |                                                                                                                                                         |                      | Chemotherapy or autoH SCT (%)                    | AlloH SCT (%) | EBMT score                                                                  | HCT-CI score | NRM risk (%) |
| Good            | -t(8;21) or AML1-ETO, WBC < 20<br>-inv16/t(16;16) or CBFB-MYH11<br>-CEBPA-biallelic mutant-positive<br>-t(8;21) or AML1-ETO plus WBC > 20 or mutant KIT | Positive or negative | 35-40                                            | 15-20         | NA(≤1)                                                                      | NA (<1)      | 10-15        |
| Intermediate    | -CN-X-Y, WBC < 100, CRe<br>-t(8;21) or AML1-ETO plus WBC > 20 or mutant KIT                                                                             | Negative             | 50-55                                            | 20-25         | ≤2                                                                          | ≤2           | <20-25       |
| Poor            | -CN -X-Y, WBC < 100, CRe                                                                                                                                | Positive             | 70-80                                            | 30-40         | ≤3-4                                                                        | ≤3-4         | <30          |
|                 | -t(8;21) or AML1-ETO, WBC > 20 and/or mutant KIT                                                                                                        | Positive             |                                                  |               |                                                                             |              |              |
|                 | -CN-X-Y, WBC < 100, n CRe                                                                                                                               | Negative             |                                                  |               |                                                                             |              |              |
|                 | -CN-X-Y, WBC > 100                                                                                                                                      | Negative             |                                                  |               |                                                                             |              |              |
|                 | -CA, but non-CBF, MK-negative, no abn3q26                                                                                                               |                      |                                                  |               |                                                                             |              |              |
| Very Poor       | -CN -X -Y, WBC > 100                                                                                                                                    | Positive             | >90                                              | 40-50         | ≤5                                                                          | ≤5           | <40          |
|                 | -CA, but non-CBF, MK-negative, no abn3q26, EV1-negative                                                                                                 | Positive             |                                                  |               |                                                                             |              |              |
|                 | -MK-positive                                                                                                                                            | Positive or negative |                                                  |               |                                                                             |              |              |
|                 | -abn3q26<br>-Non-CBF, EVI1-positive<br>-Non-CBF with mutant p53, or                                                                                     |                      |                                                  |               |                                                                             |              |              |

|  |                                                                                                     |  |  |  |  |  |  |
|--|-----------------------------------------------------------------------------------------------------|--|--|--|--|--|--|
|  | -mutant RUNX1, or mutant ASXL1<br>-or biallelic FLT3-ITD with<br>-FLT3-ITD:FLT3 WT ratio of<br>>0.6 |  |  |  |  |  |  |
|--|-----------------------------------------------------------------------------------------------------|--|--|--|--|--|--|

Abbreviations: CA = cytogenetic abnormalities; CBF = core binding factor; CN = cytogenetically normal; CRe = early complete remission; EBMT = European Group for Blood and Marrow Transplantation; HCT-CI= hematopoietic cell transplantation comorbidity index; ITD = internal tandem duplication; MK = monosomal karyotype; NA = not applicable; NRM = non-relapse mortality; -X -Y = deleted X or Y chromosome.

## Treatment

If CR has been achieved further therapy is necessary for potential cure. The nature of consolidation therapy must be individualized for each patient based on a risk analysis of the risk of relapse of the AML versus the risk of the proposed consolidation therapy. This will depend on prognostic features of the leukemia, response to therapy, performance status and type of hematopoietic stem cell donor available. High dose Ara-c (HiDAC) is the mainstay of consolidation chemotherapy as there has been shown to be a dose intensity effect to cytarabine suggesting that HiDAC is necessary in induction or consolidation. Additional agents such as FLT3 inhibitors or gemtuzumab ozogamicin are added as indicated and CPX-351 can replace HiDAC in the appropriate patients. Generally, at least one cycle is administered in all patients if only to allow for planning of an allogeneic stem cell transplant although the absolute need for this is controversial. Autologous stem cell transplantation shows some superiority in event-free survival over chemotherapy alone for consolidation, however, is not routinely recommended unless a donor is not available.

- **Favourable risk patients:** In patients with AML with t(8;21), inv 16, isolated NPM1 mutation or bZIP in-frame mutated CEBPA data suggests that provided there are no additional risk factors multiple cycles of HiDAC provide higher overall survival than lower doses of cytarabine or stem cell transplant. MRD assessment is paramount to making decisions around the use of allo-HSCT in CR1. In those not proceeding to allo-HSCT in CR1, Our recommendation is 2-4 cycles of HiDAC post induction chemotherapy accompanied by gemtuzumab ozogamicin for the first two cycles.
- **Intermediate risk patients:** HiDAC has been shown to be preferable to lower dose cytarabine in this cytogenetic group as well but its superiority over stem cell transplantation has not been established. It is generally recognized that an allogeneic stem cell transplant provides a decreased relapse rate at a cost of increased treatment related mortality when compared to consolidation chemotherapy or autologous transplantation. The transplant related mortality gap between match related and unrelated donors has been shown to be significantly reduced in recent years. A suitable hematopoietic stem cell donor should be sought in all cases to aid in decision-making about HSCT. If a matched sibling donor is found, a related myeloablative stem cell transplant should proceed as soon as possible, ideally after one cycle of HiDAC. If there are no suitable family donors, the patient should proceed through 3-4 cycles of HiDAC +/- GO or midostaurin consolidation while a matched-unrelated donor is obtained. If no matched donor is

available, the decision to proceed with a haploidentical or mismatched unrelated donor allo-HSCT should be individualized based on AML characteristics, patient comorbidities and patient preference.

- **High risk patients:** All efforts should be undertaken to find a suitable donor for eligible high-risk patients. During that time the patient should receive ongoing cycles of HiDAC chemotherapy up to a total of 4 cycles or CPX-351 up to 2 cycles. The patient should proceed to allogeneic stem cell transplantation as soon as a donor is identified.
- **TP53:** A patient with TP53-mutated AML should be considered for Allo HSCT only if fit and in remission, but the decision should be individualized based on molecular and cytogenetic risk factors as well as response to therapy as outcomes remain poor and careful risk stratification is essential. Figure 1, adapted from Baranwal et al reviews outcomes of patients with myeloid neoplasms according to various diagnostic and post treatment scenarios. CK = complex karyotype.

**Figure 1:**

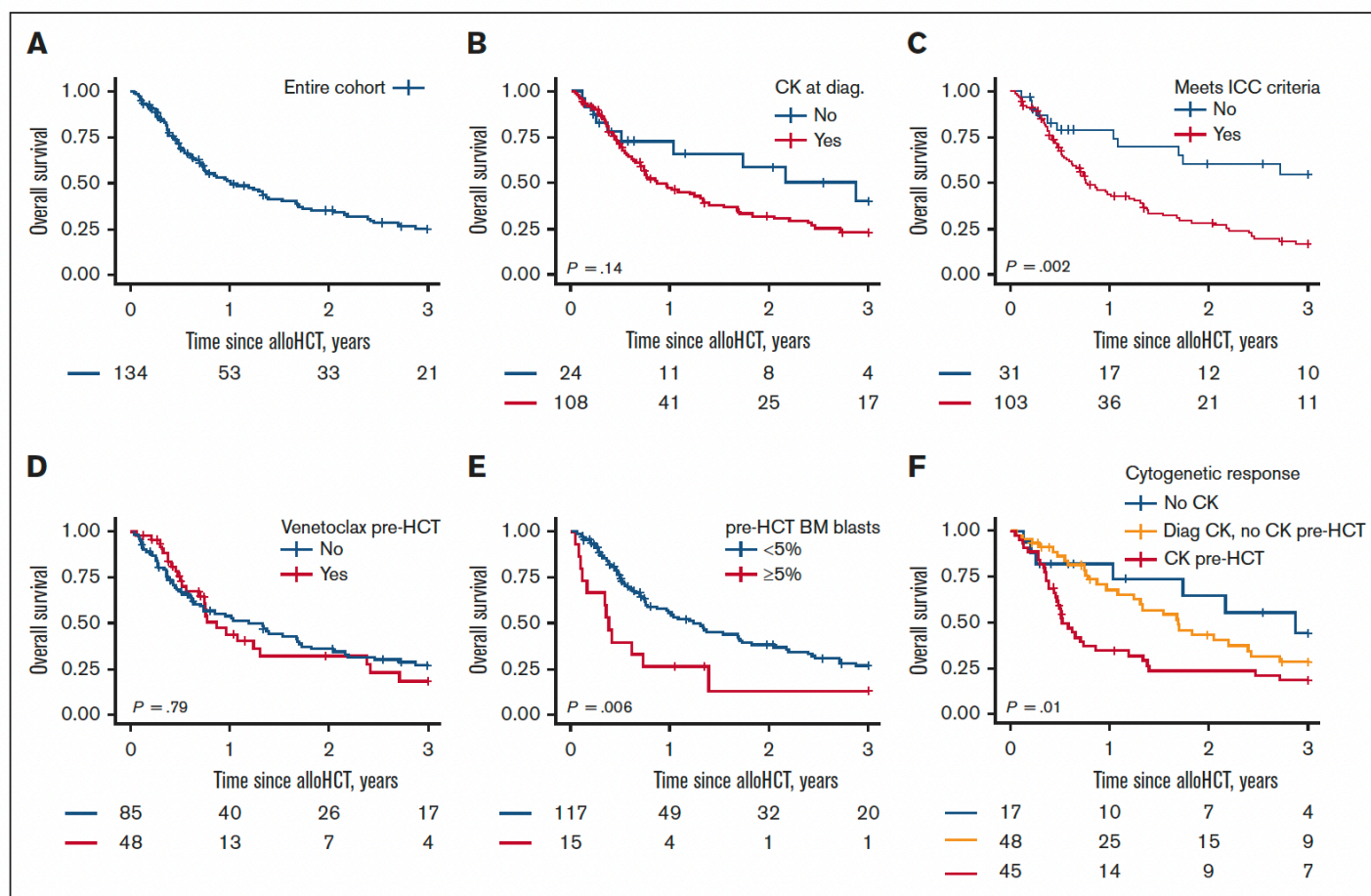




Table 4, adapted from Nawas et al, provides a good framework for decision making in these patients. There is no conclusive evidence regarding the intensity of conditioning in these patients.

**Table 4.** Summary of proposed recommendations regarding pursuit of allo-HCT in patients with TP53-mutated MDS/AML according to disease and patient characteristics

| Disease characteristics                                    | Patient characteristics  | Offer allo-HCT?                | Comments                                                                                                                                                                                                                                                                                                                                                                                                                                    |
|------------------------------------------------------------|--------------------------|--------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Monoallelic <i>TP53</i> mutation without complex karyotype | Fit for allo-HCT         | Yes, if both criteria were met | Standard fitness criteria applied.                                                                                                                                                                                                                                                                                                                                                                                                          |
| Biallelic <i>TP53</i> loss ± complex karyotype             | KPS ≥ 90 and HCT-CI < 4* | Consider                       | *Score intended as provisional guidance rather than absolute threshold. Consider the use of alternative nonrelapse mortality scoring systems. <sup>70,71</sup><br>Donor options may modify risk assessment (eg, we are less likely to pursue allo-HCT in an older patient with these characteristics if only alternative donor is available).<br>Must clearly discuss risks, benefits, and expectations regarding posttransplant prognosis. |
| Biallelic <i>TP53</i> loss ± complex karyotype             | KPS < 90<br>HCT-CI ≥ 4*  | No                             | Consider referral for second opinion regarding allo-HCT.                                                                                                                                                                                                                                                                                                                                                                                    |

## Maintenance:

The use of various post transplant therapies in the maintenance setting is being investigated in the FLT3 positive setting. Studies are also ongoing looking at the use of HMA's alone or in combination in high risk scenarios. No such intervention is currently recommended.

**CR2:** Patients with any initial risk profile who relapse after standard treatment not including stem cell transplantation should undergo transplant should they achieve a second complete remission.

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# Acute Lymphoblastic Leukemia (ALL)

Presented by: Lynn Savoie

## Summary

- The search for a donor should be undertaken for all patients, including those with standard risk disease until it has been proven that they can tolerate the intensification portion of the chemotherapy protocol.
- Transplantation in first complete remission will be offered to patients who meet other eligibility criteria and who have any one of the following:
  - Complex (>5 abnormalities), low hypodiploid (30-39 chromosomes) or near triploid (66-79 chromosomes) karyotypes.
  - t(8;14), KMT2A gene rearrangements or IKZF1 mutations.
  - Philadelphia-like disease.
  - Failure to enter a morphologic remission within 28 days of starting induction chemotherapy.
  - Patients who are MRD +ve at  $>10^{-3}$  (or  $>0.1\%$ ) after induction therapy, or  $>10^{-4}$  ( $<0.01\%$ ) at 16-18 weeks
  - Ph +ve patients with persistent molecular positivity
  - Intolerance of post-induction chemotherapy such that less than 80% of scheduled chemotherapy is likely to be delivered.
- Patients without documented CNS disease should receive at least four doses of intrathecal chemotherapy for CNS prophylaxis prior to transplant.
- Patients should be in remission (defined as fewer than 5% blasts in a normocellular bone marrow and no active extramedullary disease or circulating blasts) at the time of transplantation.
- BCR-ABL will be monitored post-transplant and TKI therapy re-instituted upon any evidence of molecular positivity. TKI prophylaxis should be considered for those with high-risk disease at the time of transplant (see below for details).
- Stem cell transplantation should be offered to all transplant-eligible patients with recurrent ALL, a suitable donor and meeting general eligibility criteria (including remission status) for transplantation.
- CAR-T therapy with Tisagenlecleucel or Brexucabtagene autoleucel may be offered in the setting of relapsed or refractory disease.

## Background

Acute Lymphoblastic Leukemia (ALL) is a highly aggressive hematological-malignancy resulting from the proliferation and expansion of lymphoid blasts in the blood, bone marrow and other organs<sup>1</sup>. ALL occurs with a bimodal distribution with an early peak in children 4-5 years old followed by a second peak at ~ 50 years of age<sup>2</sup> with the worldwide incidence being ~ 1- 4.75/100,000 individuals with a male:female prevalence of roughly 1:3:1<sup>1</sup>. It is the most common childhood acute leukemia

accounting for ~ 80% of the pediatric leukemias but contributing to only 20% of adult leukemias. Although significant progress has been made in treating adult ALL the overall survival amongst adults 18 to 60 years old is only 35% in contrast to childhood ALL in which overall survival at five years is more than 80%<sup>1</sup>.

Over the past two decades the treatment of adult ALL has changed significantly with the introduction of pediatric protocols for the treatment of adolescents and young adults, the addition of tyrosine kinase (TKI) inhibitors for the treatment of Philadelphia positive/BCR-ABL positive ALL, a reevaluation of the role of allogeneic stem cell transplantation for standard risk ALL patients, the incorporation of minimal residual disease into risk assessments and most recently the introduction of novel agents such as blinatumomab, inotuzumab and chimeric antigen receptor- T cell therapy for relapsed/refractory ALL.

With current treatment regimens, the cure rate among children with ALL is approximately 80-90%<sup>2</sup>. The long-term prognosis for adults with ALL treated with conventional chemotherapy regimens, however, remain poor, with cure rates of only 30 to 40%<sup>3-10</sup>. This reflects the greater tendency for older individuals to have adverse chromosomal markers (notably t (9; 22)) and other unfavorable prognostic indicators (high white blood cell (WBC) count, longer time to complete response and measurable disease negativity). Multidrug chemotherapy regimens have been the standard approach to treatment of adults with ALL. Such regimens generally consist of 4- or 5-drug induction protocols followed by intensive re-induction, consolidation or intensification to address residual disease. These regimens also feature CNS prophylaxis in the form of whole brain radiotherapy or intrathecal chemotherapy and prolonged antimetabolite-based maintenance, as has been used successfully in management of pediatric cases.

Subsequently, a growing body of data has shown that, at least for late adolescents and young adults (defined variably up to 40 years of age), treatment with pediatric-based protocols produces superior outcomes to the regimens standardly used in adults<sup>11-15</sup>. Canadian data has shown that a pediatric approach can safely be extended to adults up to the age of 60 with only minor modifications<sup>16</sup>. This protocol is heavily dependent on L-Asparaginase in intensification and has been shown to have the best outcomes if 80% of L-Asparaginase doses can be delivered; this has been shown to be possible in 80% of patients. Meaningful comparisons of this strategy with early transplantation have yet to be published.

Finally, many novel therapies have come along in the management of ALL and have affected the need for or the outcome of stem cell transplantation. These include Blinatumomab<sup>17</sup>, Inotuzumab<sup>18</sup>, the addition of Rituximab to chemotherapy<sup>19</sup> and CAR-T cell therapy<sup>20</sup>.

CNS prophylaxis in the form of, intrathecal chemotherapy and/or high dose systemic chemotherapy has been shown to be necessary throughout chemotherapy and prior to stem cell transplantation.

## Risk Stratification in ALL

Over the past twenty years there has been continued debate regarding the risk stratification of patients with ALL with different groups using the above clinical, immunophenotype, cytogenetics and molecular tests to variably group patients into those that are standard risk, high risk or very high risk of having a leukemia relapse. As discussed, many of these discrepancies may be related to the type of treatment regimens used (adult vs pediatric-based), and whether MRD analysis is taken into consideration in risk stratification.

Recent studies have found that, using pediatric-based protocols, most cytogenetic abnormalities are not independent predictors except for certain abnormalities, such as KMT2A-based abnormalities, t(9;22) and possibly hypodiploidy<sup>21-22</sup>. Given the poor prognosis of patients with Ph+ve ALL, the initial risk stratification for all patients should be based on the presence or absence of t(9;22)/BCR-ABL1. Amongst Ph-ve patients the NCCN considers those with hypodiploidy (<44 chromosomes), t(v;11q23)/KMT2A rearrangements or complex karyotype (≥5 chromosomal abnormalities) as high risk<sup>23</sup>.

Although prior studies have shown that a WBC > 30 for B-cell ALL and >100 for T-cell ALL were important risk factors<sup>16</sup>, MRD has since been demonstrated to be a more important predictor on multivariate analysis. However, age remains an important predictor of outcome in nearly all studies, with patients > age 30-35 having worse outcomes than so-called AYA (adolescent and young adult) patients<sup>16</sup>.

Based on the pediatric literature, a number of studies over the past 15 years have explored the role of MRD in adults with ALL. Most of these have suggested that MRD may be the single most important factor predicting clinical outcomes.<sup>24</sup>

A study by the GRAALL group, using a pediatric inspired protocol for adults with B-ALL, evaluated the role of allogeneic HSCT according to MRD response with induction therapy.<sup>25</sup> Patients who were MRD positive, defined as a level > 10<sup>-3</sup> following induction therapy, had a significantly higher relapse rate and inferior OS as compared to those who achieved a level < 10<sup>-3</sup>. Furthermore, those with MRD levels > 10<sup>-3</sup> who underwent subsequent HSCT in CR1 had a significantly superior RFS and OS as compared to those who were not transplanted. In contrast, those with MRD levels < 10<sup>-3</sup> following induction did not benefit from HSCT. These effects were seen for both B-ALL and T-ALL.

The Edmonton group subsequently analyzed outcomes following DFCI induction therapy. Between 2013-2019 patients with BCR-ABL negative ALL underwent induction therapy with this protocol, and MRD post-induction was assessed by multiparameter flow cytometry, with a sensitivity of 0.1%. Of 46 patients who achieve CR, 26 (57%) were MRD negative and 43% were MRD positive. The cumulative incidence of relapse was 45% in patients who were MRD positive at a level of >0.1%, as



compared with 12% in MRD negative patients ( $p=0.05$ ) (unpublished data). These results essentially mirror those reported by the GRAALL group as described above.

These data further support the conclusion that patients who fail to achieve a 3 log reduction in MRD levels with intensive induction therapy represent a high-risk group for relapse, and that these patients should be considered for HSCT in CR1. In contrast, those who achieve a  $>3$  log reduction with induction can be successfully managed by chemotherapy alone with a low relapse risk and favorable prognosis.

Recent studies in adults have demonstrated the impact of molecular status on relapse risk in Ph+ ALL. The MD Anderson Cancer Center<sup>26</sup> found that molecular positivity at a variety of time points was associated with a significantly higher cumulative incidence of relapse. The French GRAAPH study<sup>27</sup> found that the benefit of allo-HSCT in CR1 was restricted to patients who were molecularly positive post-induction chemotherapy. On subgroup analysis, patients who had achieved MRD negativity (defined as a  $>4$  log reduction in BCR-ABL transcripts by PCR) by the end of the second induction cycle had a similar RFS with or without allo-HSCT. In contrast patients who were MRD positive at that timepoint had a significantly better RFS with allogeneic HSCT, compared to those who were not transplanted. Therefore, MRD assessment with this regimen could be used to help identify a high-risk cohort for which HSCT was clearly indicated.

In contrast, a study using nilotinib plus chemotherapy<sup>28</sup> found that overall survival was superior in patients undergoing HSCT regardless of MRD status; however, 60% of patients achieving molecular negativity remained free of relapse at 30 months without a transplant. Taken together, these data suggest that some patients who achieve MRD negativity early on may remain relapse-free with continuing chemotherapy + TKI, potentially sparing them the toxicity associated with transplant.

## Hematopoietic Stem Cell Transplant (HSCT)

### Transplantation in First Complete Remission

It remains unclear whether adult patients treated with pediatric protocols gain a benefit from SCT. In a study of 156 BCR-ABL –ve patients treated with the DFCI protocol the 5 year OS amongst patients receiving an allogeneic SCT was 44% while for those not undergoing a SCT the survival was 74% with the difference possibly related to transplant related mortality. Seftel and colleagues compared 422 Ph-ve ALL patients aged 18-50 years with 108 patients receiving DFCI chemotherapy<sup>29</sup>. Expectedly, treatment related mortality was higher in those receiving a SCT (37% vs. 6%). At 4 years, the incidence of relapse was similar for those receiving SCT and chemotherapy (24% vs. 23%), however, both DFS and OS were improved for those receiving chemotherapy alone (40% vs. 71% for DFS and 45% vs. 73% for OS). Dhedin and colleagues further assessed the role of allogeneic stem cell transplantation in Ph-ve ALL adult patients with at least 1 conventional high-risk factor treated with the pediatric inspired GRAALL 2003 and 2005 protocols<sup>30</sup>. In all, 522 patients age

15 to 55 years old were candidates for SCT in first complete remission. Among these, 282 (54%) received a transplant in first complete remission while 240 (46%) did not. As with previous studies, the lower CIR observed in the SCT was counterbalanced by a higher NRM. When analyzing SCT in CR1 as a time-dependent event RFS and OS were not significantly improved in the SCT cohort. No significant effect of SCT on RFS was noted in patients younger or older than age 45 or on any prespecified baseline risk factor. RFS and OS were significantly longer, however, in patients who presented with morphologic poor early BM blast clearance or in late CR patients. Furthermore, SCT was associated with longer RFS in patients with postinduction minimal residual disease (MRD)  $>10^{-3}$  but not in good MRD responders.

For pediatric-based protocols, asparaginase is a critical agent in achieving the high cure rates reported. There is evidence in both pediatric and adult studies with the DFCl protocol that the inability to deliver the intended asparaginase dosing during intensification (defined as  $\geq 80\%$ ) is associated with a higher relapse rate<sup>16,31</sup>. Therefore, the inability to deliver effective asparaginase dosing (e.g. due to pancreatitis) places the patients in a higher risk category and warrants consideration of allogeneic HSCT.

Transplantation in first complete remission will be offered to patients who meet other eligibility criteria and who have any one of the following:

- Complex ( $>5$  abnormalities), low hypodiploid (30-39 chromosomes) or near triploid (66-79 chromosomes) karyotypes.
- t(8;14), KMT2A gene rearrangements or IKZF1 mutations.
- Philadelphia-like disease.
- Failure to enter a morphologic remission within 28 days of starting induction chemotherapy.
- Patients who are MRD +ve at  $>10^{-3}$  (or  $>0.1\%$ ) after induction therapy, or  $>10^{-4}$  ( $<0.01\%$ ) at 16-18 weeks
- Ph +ve patients with persistent molecular positivity
- Intolerance of post-induction chemotherapy such that less than 80% of scheduled chemotherapy is likely to be delivered.

### **Philadelphia-positive Acute Lymphoblastic Leukemia**

BCR/ABL monitoring should be done every 3 months for the first year post transplant then with every visit. If there is re-appearance of a TKI transcript treatment with a TKI should be reinitiated. The choice of TKI would depend on responses pre-transplant and comorbidities. There is no data to suggest duration of TKI therapy in this setting but would be at least for many years after return to negativity or possibly indefinitely.

TKI maintenance may have a potential role in reducing the risk of relapse following HSCT, however pre-emptive therapy has also led to favorable long term outcomes<sup>32-33</sup>. A meta-analysis of the use of TKI's post transplant showed that use of TKIs (all generations) after allo-HSCT for patients in CR1

improved OS when given as a prophylactic or pre-emptive regimen. Limited data suggest that second-generation TKIs (i.e. dasatinib) have a better OS, especially in patients with MRD-positive status<sup>34</sup>. Consideration should be given to the use of TKI's post transplantation for those with high risk disease such as BCR/ABL MRD positivity or in second complete remission at the time of transplantation.

### **Transplantation beyond First Complete Remission**

The outcome for patients with ALL who fail to achieve a remission or who relapse remains poor, and such patients are generally offered alloHSCT from a matched or mismatched sibling, a volunteer unrelated donor or with umbilical cord blood stem cells. Long-term prognosis depends on time from remission to relapse, with shorter remissions being associated with worse prognosis. Allogeneic sibling HSCT in second CR results in 15-35% LFS, while for patients with refractory relapse, LFS between 8 – 33% have been reported. It is generally recommended that patients complete a course of CNS prophylaxis between relapse and transplantation.

### **CAR-T Therapy**

Tisagenlecleucel is indicated in Canada for the treatment of relapsed or refractory acute lymphoblastic leukemia in patients under the age of 26. This indication is based partly on the results of a single-cohort, multi-center phase II trial (ELIANA) conducted by Maude et al<sup>35</sup>. In this study, children and young adults with relapsed or refractory CD19+ B-cell ALL received a single infusion of tisagenlecleucel after lymphodepleting chemotherapy. The primary end point, the overall remission rate at 3 months, was 82%, with all responding patients negative for MRD by flow cytometry. Event-free and overall survival were 50% and 76% at one year. Cytokine release syndrome occurred in 77% of patients, with tocilizumab required in 48%. Neurological events occurred in 40%. There were significant improvements in patient-reported outcomes (PedsQL and EQ-5D) three months after treatment<sup>36</sup>. Real world evidence in 255 patients with relapsed or refractory ALL further supports the use of tisagenlecleucel in this context. In this report twelve-month EFS and OS were 52.4% and 77.2%<sup>37</sup>.

Brexucabtagene autoleucel is indicated in Canada for use in adults with relapsed or refractory ALL based on the ZUMA-3 phase 2 study. This showed that in patients aged 28-52 with a median followup of 16.4 months, 39 patients had complete remission or complete remission with incomplete haematological recovery, with 31 (56%) patients reaching complete remission. Median duration of remission was 12.8 months, median relapse-free survival was 11.6 months and median overall survival was 18.2 months. Among responders, the median overall survival was not reached, and 38 (97%) patients had MRD negativity<sup>38</sup>.

At this time, in the relapsed/refractory setting, the optimal sequencing of transplantation versus CAR-T has not been established.



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# Myelodysplastic Syndromes (MDS)

Presented by: Michelle Geddes

## Summary

- All patients should have risk stratification including calculation of the Revised International Prognostic Scoring System (R-IPSS, <https://www.mds-foundation.org/ipss-r-calculator/>) and molecular prognostication with IPSS-M, (<https://mds-risk-model.com/>) at diagnosis.
- Patients with high (>4.5 to 6 points) or very high (>6 points) R-IPSS should be offered HCT if they are transplant-eligible, or they have moderate-high, high or very high-risk M-IPSS score at diagnosis.
- Patients with intermediate (>3 to 4.5 points) R-IPSS with symptomatic cytopenias or evidence of disease progression, who are transplant-eligible, can be considered for allogeneic HCT; with consideration of patient values and discussion around risks of transplant compared to the underlying disease.
- Sibling typing should be initiated at the earliest opportunity for all transplant-eligible patients.
- Disease control as a bridge to transplant with induction chemotherapy (consider if blasts >10% and no TP53 mutation and/or adverse cytogenetic profile) or hypomethylating agents such as azacitidine should be considered for patients with higher risk disease or elevated blast counts at presentation, unless transplant can be done rapidly (1-2 months). The optimal therapy in this setting is not clear.
- Minimize iron overload pretransplant with iron chelation therapy to minimize the adverse effects of iron overload on treatment-related mortality.
- Myeloablative conditioning is preferred over non-myeloablative conditioning in patients who are fit to improve patient outcomes. Our standard conditioning is myeloablative busulfan + fludarabine + 4Gy TBI (see Conditioning chapter). Treosulfan-based reduced intensity conditioning can be considered in patients with multiple comorbidities, especially liver disease.
- Patients under the age of 40 or with an appropriate family history should be screened for congenital causes of MDS (i.e. Fanconi, dyskeratosis congenita). Appropriate consideration of hereditary syndromes should be made during donor search and planning of conditioning; this testing should be initiated early i.e. before transplant referral in patients with NGS or family history suggestive of germline mutations.
- In very high-risk patients, i.e., complex karyotype and p53 mutation, alternatives to transplant should be considered, and frank discussion with patients about poor outcomes with transplant should be initiated.

## Background

Myelodysplastic syndromes are a heterogeneous group of related clonal stem cell disorders featuring dysplastic changes in one or more bone marrow cell lines, ineffective hematopoiesis, bone marrow failure, and often clonal evolution and/or transformation to acute leukemia. It is a disorder of the

elderly, with a median age of 65-70 years at diagnosis. Allogeneic stem cell transplantation remains the only curative option; however, the majority of patients are not eligible for transplantation due to age and/or comorbidity. For those who are eligible, the variable natural history of the disease and relative toxicity of transplant are important factors in the decision between supportive care, hypomethylating agents, lenalidomide, medical therapy including growth factors and allogeneic transplantation, and clinical trials.

## Etiology

A history and physical exam should investigate for potential etiology of MDS:

- Ionizing radiation
- Cytotoxic agents (i.e., alkylating agents, topoisomerase inhibitors)
- Occupational or environmental carcinogens (i.e., viruses, benzenes, heavy metals)
- Inherited disorders (i.e., Fanconi anemia) especially in consideration of related donors
- Antecedent hematologic disorders (i.e. paroxysmal nocturnal hemoglobinuria, aplastic anemia)
- Rule out B12 and folate deficiency

Cytogenetic abnormalities are found in 40-70% of *de novo* MDS, and 95% of therapy-related MDS.

**Table 1:** WHO 2022 Classification of MDS (from Khoury JD, et al. Leukemia. 2022. PMID: 35732831)

|                                                                                 | Blasts                             | Cytogenetics                                                                        | Mutations                                                                                               |
|---------------------------------------------------------------------------------|------------------------------------|-------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------|
| <b>MDS with defining genetic abnormalities</b>                                  |                                    |                                                                                     |                                                                                                         |
| MDS with low blasts and isolated 5q deletion (MDS-5q)                           | <5%BM and <2% PB                   | 5q deletion alone, or with 1 other abnormality other than monosomy 7 or 7q deletion |                                                                                                         |
| MDS with low blasts and <i>SF3B1</i> mutation <sup>a</sup> (MDS- <i>SF3B1</i> ) |                                    | Absence of 5q deletion, monosomy 7, or complex karyotype                            | <i>SF3B1</i>                                                                                            |
| MDS with biallelic <i>TP53</i> inactivation (MDS-bi <i>TP53</i> )               | <20% BM and PB                     | Usually complex                                                                     | Two or more <i>TP53</i> mutations, or 1 mutation with evidence of <i>TP53</i> copy number loss or cnLOH |
| <b>MDS, morphologically defined</b>                                             |                                    |                                                                                     |                                                                                                         |
| MDS with low blasts (MDS-LB)                                                    | <5% BM and <2% PB                  |                                                                                     |                                                                                                         |
| MDS, hypoplastic <sup>b</sup> (MDS-h)                                           |                                    |                                                                                     |                                                                                                         |
| MDS with increased blasts (MDS-IB)                                              |                                    |                                                                                     |                                                                                                         |
| MDS-IB1                                                                         | 5-9% BM or 2-4% PB                 |                                                                                     |                                                                                                         |
| MDS-IB2                                                                         | 10-19% BM or 5-19% PB or Auer rods |                                                                                     |                                                                                                         |
| MDS with fibrosis (MDS-f)                                                       | 5-19% BM; 2-19% PB                 |                                                                                     |                                                                                                         |

<sup>a</sup>Detection of ≥15% ring sideroblasts may substitute for *SF3B1* mutation. Acceptable related terminology: MDS with low blasts and ring sideroblasts.

<sup>b</sup>By definition, ≤25% bone marrow cellularity, age adjusted; BM bone marrow, PB peripheral blood, cnLOH copy neutral loss of heterozygosity

**Table 2:** International Consensus Classification of MDS and MDS/AML (from Blood (2022) 140 (11): 1200–1228).

| Myelodysplastic syndromes (MDS) and myelodysplastic syndrome/acute myeloid leukemia (MDS/AML) |                           |            |                        |                               |                                                                 |                                                                |
|-----------------------------------------------------------------------------------------------|---------------------------|------------|------------------------|-------------------------------|-----------------------------------------------------------------|----------------------------------------------------------------|
|                                                                                               | Dysplastic lineages       | Cytopenias | Cytoses*               | BM and PM Blasts              | Cytogenetics <sup>†</sup>                                       | Mutations                                                      |
| MDS with mutated <i>SF3B1</i> (MDS- <i>SF3B1</i> )                                            | Typically ≥1 <sup>‡</sup> | ≥1         | 0                      | <5% BM<br><2% PB              | Any, except isolated del(5q), -7/del(7q), abn3q26.2, or complex | SF3B1 (≥10% VAF), without multi-hit TP53, or RUNX1             |
| MDS with del(5q) [MDS-del(5q)]                                                                | Typically ≥1 <sup>‡</sup> | ≥1         | Thrombocytosis allowed | <5% BM<br><2% PB <sup>§</sup> | del(5q), with up to 1 additional, except -7/del(7q)             | Any, except multi-hit TP53                                     |
| MDS, NOS without dysplasia                                                                    | 0                         | ≥1         | 0                      | <5% BM<br><2% PB <sup>§</sup> | -7/del(7q) or complex                                           | Any, except multi-hit TP53 or SF3B1 (≥10% VAF)                 |
| MDS, NOS with single lineage dysplasia                                                        | 1                         | ≥1         | 0                      | <5% BM<br><2% PB <sup>§</sup> | Any, except not meeting criteria for MDS-del(5q)                | Any, except multi-hit TP53; not meeting criteria for MDS-SF3B1 |
| MDS, NOS with multilineage dysplasia                                                          | ≥2                        | ≥1         | 0                      | <5% BM<br><2% PB <sup>§</sup> | Any, except not meeting criteria for MDS-del(5q)                | Any, except multi-hit TP53; not meeting criteria for MDS-SF3B1 |
| MDS with excess blasts (MDS-EB)                                                               | Typically ≥1 <sup>‡</sup> | ≥1         | 0                      | 5-9% BM, 2-9% PB <sup>§</sup> | Any                                                             | Any, except multi-hit TP53                                     |
| MDS/AML                                                                                       | Typically ≥1 <sup>‡</sup> | ≥1         | 0                      | 10-19% BM or PB <sup>  </sup> | Any, except AML-defining <sup>¶</sup>                           | Any, except NPM1, bZIP CEBPA or TP53                           |

\*Cytoses: Sustained white blood count ≥13 x 10<sup>9</sup>/L, monocytosis (≥0.5 x 10<sup>9</sup>/L and ≥10% of leukocytes) or platelets ≥450 x 10<sup>9</sup>/L; thrombocytosis is allowed in MDS-del(5q) or in any MDS case with inv(3) or t(3;3) cytogenetic abnormality.

†BCR::ABL1 rearrangement or any of the rearrangements associated with myeloid/lymphoid neoplasms with eosinophilia and tyrosine kinase gene fusions exclude a diagnosis of MDS, even in the context of cytopenia.

‡Although dysplasia is typically present in these entities, it is not required.

§Although 2% PB blasts mandates classification of an MDS case as MDS-EB, the presence of 1% PB blasts confirmed on 2 separate occasions also qualifies for MDS-EB.

|| For pediatric patients (<18y), the blast thresholds for MDS-EB are 5% to 19% in BM and 2% to 19% in PB, and the entity MDS/AML does not apply.

¶AML-defining cytogenetics are listed in the AML section.

**Table 3.** Revised IPSS (R-IPSS) for MDS<sup>2</sup>

| Prognostic Variable                  | Score     |           |          |     |              |      |           |
|--------------------------------------|-----------|-----------|----------|-----|--------------|------|-----------|
|                                      | 0         | 0.5       | 1        | 1.5 | 2            | 3    | 4         |
| Cytogenetics*                        | Very good |           | Good     |     | Intermediate | Poor | Very poor |
| Bone marrow blast (percent)          | ≤2        |           | >2 to <5 |     | 5 to 10      | >10  |           |
| Hemoglobin (g/dL)                    | ≥10       |           | 8 to <10 | <8  |              |      |           |
| Platelets (cells/μL)                 | ≥100      | 50 to 100 | <50      |     |              |      |           |
| Absolute neutrophil count (cells/μL) | ≥0.8      | <0.8      |          |     |              |      |           |

\*Cytogenetic definitions: Very good: -Y, del(11q)

Good: Normal, del(5q), del(12p), del(20q), double including del(5q)

Poor: -7, inv(3)/t(3q)/del(3q), double including -7/del(7q), complex: 3 abnormalities

Very poor: Complex: >3 abnormalities.

**Table 4.** Survival based on total score from the R-IPSS

| Risk Group   | IPSS-R score | Median overall survival (years) | Median time to 25% AML evolution (years) |
|--------------|--------------|---------------------------------|------------------------------------------|
| Very low     | ≤1.5         | 8.8                             | >14.5                                    |
| Low          | >1.5 to 3    | 5.3                             | 10.8                                     |
| Intermediate | >3 to 4.5    | 3                               | 3.2                                      |
| High         | >4.5 to 6    | 1.6                             | 1.4                                      |
| Very high    | >6           | 0.8                             | 0.7                                      |

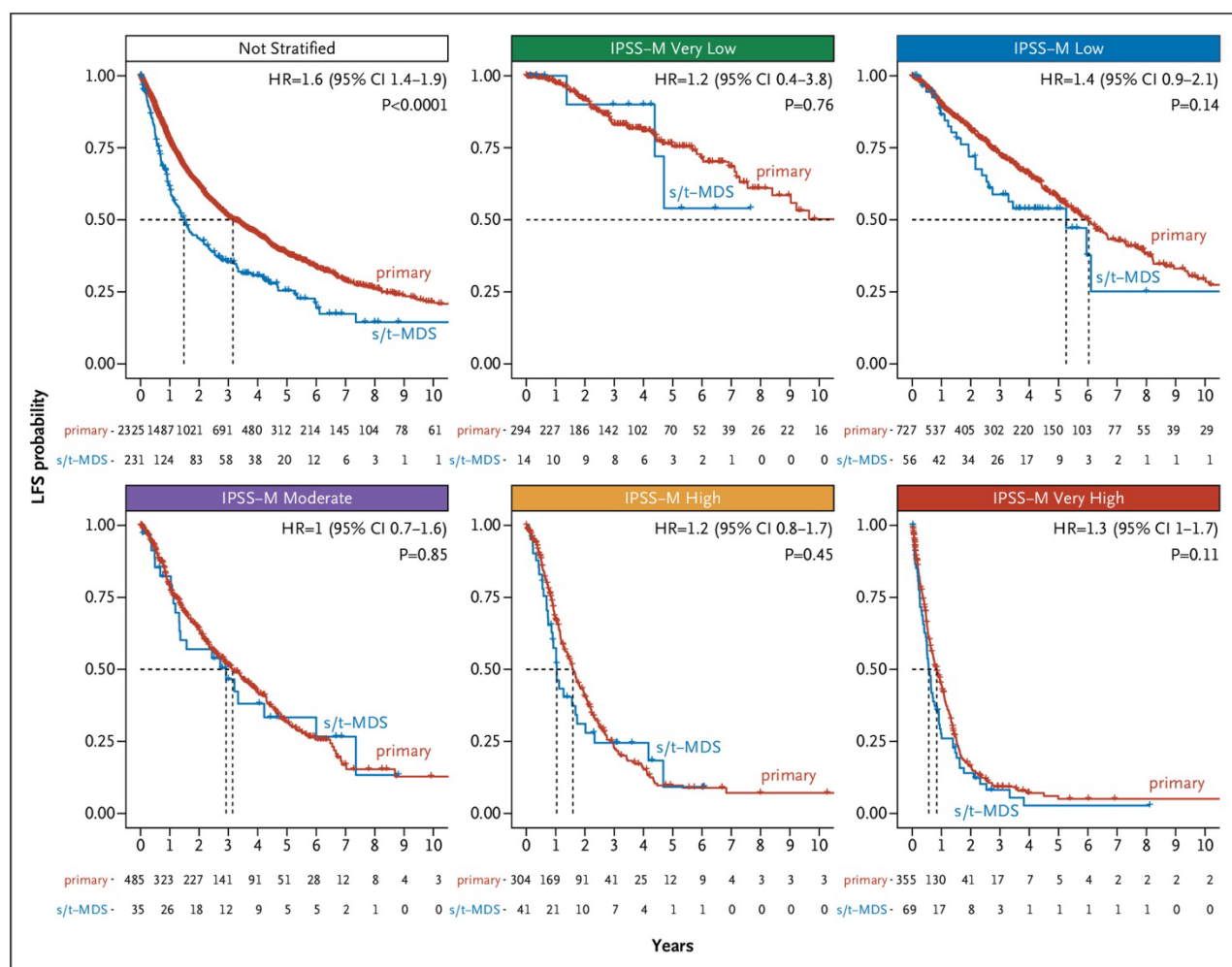
Automatic IPSS-R Calculator can be accessed at: <https://www.mds-foundation.org/ipss-r-calculator/>

For these and other online calculators also see <http://www.mdsclearpath.org>

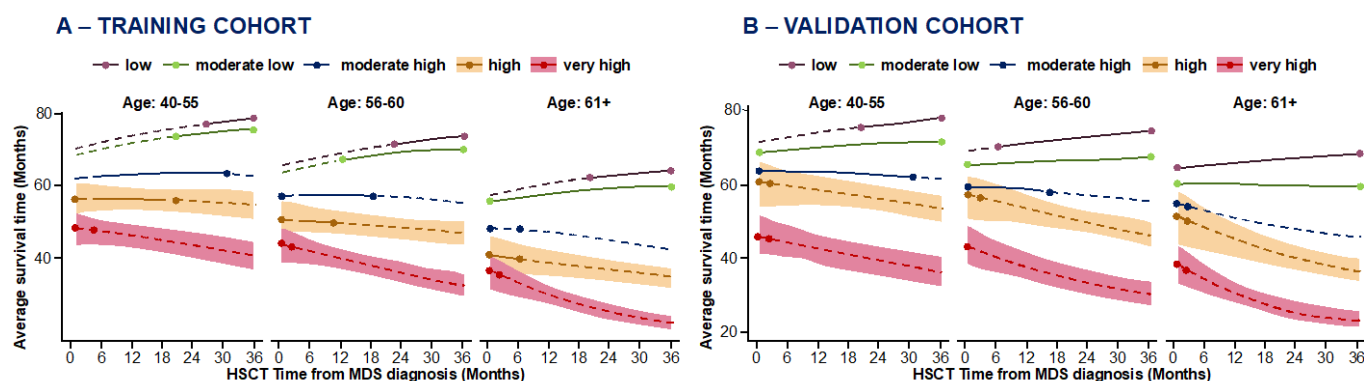
Incorporation of molecular data into the Revised IPSS score has been found to improve prognostication in MDS. Independent significant prognostic factors for survival include age, R-IPSS, EZH2, SF3B1 and TP53<sup>3</sup>. A linear predictive model was built which resulted in four prognostic groups (low, intermediate-1, intermediate-2 and high) with a median overall survival of 37.4, 23.2, 19.9 and 12.2 months, respectively, (P<0.001).

Recently a newer prognostic scoring system, the Molecular International Prognostic Scoring System (M-IPSS) incorporating clinical, cytogenetic and molecular mutations has shown improved predictive value of outcomes in patients with MDS<sup>4</sup>. This divides patients into very low, low, moderate, high and very high categories using a website calculation, and has been prospectively validated and applies to the secondary and therapy-related MDS populations.

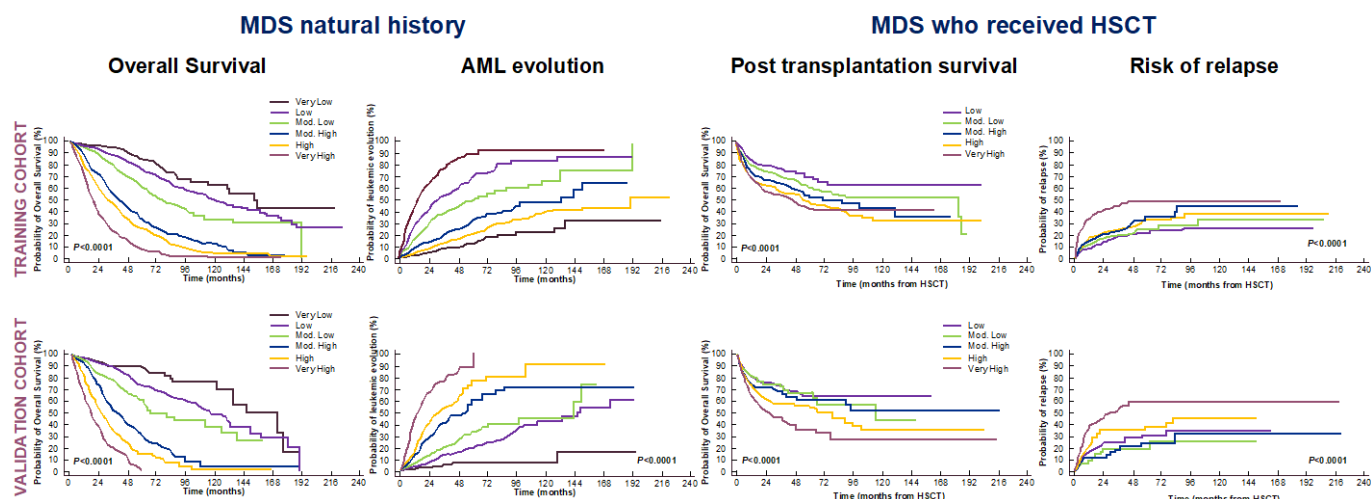




An abstract at ASH 2023 used a decision support system to predict optimal timing of transplant based on the M-IPSS with a training cohort of 4627 patients and validation cohort of 2491 patients<sup>5</sup>. Results were used to analyze the average survival time over an 8-year time period when patients with variable M-IPSS prognostic scores underwent HCT. Patients with either low- or moderate-low risk benefited from a delayed transplantation policy, patients in moderate-high, high- and very-high risk categories had improved survival with immediate transplantation.







## Intensity of Treatment in Allogeneic Stem Cell Transplantation for MDS

Reduced intensity conditioning therapy is known to have a higher relapse rate in MDS but lower treatment-related mortality. In a retrospective study of 836 patients with MDS transplanted with an HLA matched sibling, the 3-year relapse rate was higher in patients given reduced intensity conditioning (HR 1.6,  $p=0.001$ ) but a corresponding decrease in 3-year non-relapse mortality (NRM) resulted in similar progression-free survival (PFS, 33% vs. 39%) and overall survival (OS) rates (41% vs. 45%)<sup>6</sup>. The role of treatment intensity was evaluated in a randomized multicenter phase III clinical trial comparing reduced intensity conditioning (RIC) including FluBu2 (2 days busulfan) to myeloablative conditioning regimens including our current conditioning fludarabine and busulfan without TBI<sup>7</sup>. The study was stopped early due to increased relapse rates with RIC (48% vs 13.5%) with a nonsignificant reduction in OS at 18 months (68% vs 78% for RIC and MAC, respectively) and a lower relapse-free survival with RIC (47% vs 68%). Notably, TRM was lower at 4.4% with RIC vs 15.8% with MAC. Outcomes with FLUBUP/TBI remain to be determined but compare favorably with these series. Local outcomes comparing the use of TBI in patients with MDS from 1999-2010 suggest improved 2-year DFS in patients given FLUBUP/TBI compared to FLUBUP alone (2-year DFS 67% vs. 41%) although with small numbers the difference is not statistically significant. The decision has been made to incorporate TBI in the transplant regimen of patients with MDS.

Myeloablative conditioning is preferred over non-myeloablative conditioning in patients who are fit to improve patient outcomes<sup>8</sup>. Our standard conditioning is myeloablative busulfan + fludarabine + 4Gy TBI (see Conditioning chapter). Regarding reduced intensity conditioning, a large systemic review presented in a recent abstract evaluated transplant outcomes in patients with busulfan (N=4706) vs treosulfan (N=1979) based conditioning protocols for patients with MDS and AML, in patients with a median age of 55-61y<sup>9</sup>. The treosulfan and busulfan doses administered ranged from 10-42 g/m<sup>2</sup> and 0.8-12.8 mg/kg, respectively. Treosulfan was associated with better OS (RR 1.39, 95% CI 1.15-1.68,  $p$ -value 0.0007), DFS (RR 1.39, 95% CI 1.10-1.75,  $p$ -value 0.006,) and NRM (RR 0.95, 95% CI 0.90-1.00,  $p$ -value 0.05) with no difference in relapse (RR 0.96, 95% CI 0.89-1.04,  $p$ -value 0.33), aGvHD (RR 1.00, 95% CI 0.65-1.54,  $p$ -value 1.00, I<sup>2</sup> 94%) and cGvHD (RR 1.00, 95% CI 0.95-1.05,  $p$ -value

0.92). Other studies have also shown better outcome with Flu/Treo at 10-14g/m<sup>2</sup> in patients with MDS with better NRM, EFS and OS (72.8% in FluTreo at 1 year compared to 49.1% in Flubu,  $p=0.018$ )<sup>10</sup>, and similar outcomes between these regimens in a retrospective single centre analysis of patients older than 65<sup>11</sup>. Improved overall survival, RFS and GRFS were seen with treo-based regimens in Toronto in matched propensity analysis<sup>12</sup>. However, most of the above comparisons of treosulfan against busulfan did not use PK-adjusted busulfan dosing. In conclusion, treosulfan-based regimens are a promising option especially in older patients or patients with significant comorbidities undergoing HCT for MDS. Treosulfan-based reduced intensity conditioning can be considered in patients with multiple comorbidities especially liver disease<sup>12</sup>.

### Outcomes with Allogeneic Transplantation

An EBMT review of 1333 patients age >50 with high risk MDS or secondary AML who received allogeneic sibling (61%) or unrelated donor (39%) hematopoietic stem cell transplant with a myeloablative (38%) or reduced intensity conditioning (62%) regimen<sup>13</sup>. Of the 1333 patients, 449 (34%) were >60y of age. *Four-year OS was 31%*. Factors associated with higher risk of relapse include use of RIC (HR 1.44,  $P<0.01$ ) and advanced disease stage at transplantation (HR, 1.51;  $P<0.01$ ). Factors associated with increased non-relapse mortality include advanced disease stage at transplantation (HR, 1.43; 95% CI, 1.13 to 1.79;  $P = 0.01$ ), use of an unrelated donor ( $P = 0.03$ ), and RIC (HR, 0.79; 95% CI, 0.65 to 0.97;  $P = 0.03$ ). *The major factor associated with 4-year OS was disease stage at transplantation* (HR, 1.55;  $P<0.01$ ) and challenges remain in with both higher relapse rates posttransplant and higher treatment-related mortality with MDS compared to de novo AML.

A single-centre study at the University of Wisconsin describes the importance of pre-transplant disease burden (as reflected by the proportion of bone marrow blasts at transplant). Patients entering transplant with < 5% blasts had a lower probability of relapse at 1-year than those entering transplant with 5-20% blasts (18% (8-28%) vs. 35% (16-54%),  $p=0.07$ )<sup>14</sup>. The use of chemotherapy to achieve fewer than 5% blasts did not adversely affect the outcome of transplant in this cohort. The use of myeloablative conditioning was unable to overcome the adverse effect of high disease burden (>5% blasts): relapse rates were similar for patients with >5% blasts, regardless of whether myeloablative or non-myeloablative conditioning was used (28% (CI 8-48%) vs. 50% (CI 18-82%),  $p=0.33$ ).

As NGS studies become available and more information is available about disease prognostication with transplant. A Japanese study of 797 patients with MDS showed that in patients with cytogenetics and NGS testing, cox regression analysis showed approximately 70% of the hazard ratio of transplant was related to clinical factors i.e. performance status, comorbidities, transfusion history and 30% contributed by adverse genetic risk<sup>15</sup>. An especially high-risk category of patients with both mutation TP53 and complex karyotype did very poorly with transplant with a median survival of 4.8 months; 38% died before day 100 and >80% within 2 years of transplant, largely due to early relapse in 60% of patients. A retrospective review of 1514 patients with MDS undergoing stem cell transplantation

showed shorter survival and time to relapse in patients with TP53 mutations, and shorter survival in patients with JAK2 and RAS mutations<sup>16</sup>.

*Comparison of HCT vs non-HCT.* A landmark decision analysis by the CIBMTR compared outcomes in newly diagnosed MDS between three treatment strategies: transplantation at diagnosis, transplantation at leukemic progression, and transplantation at an interval from diagnosis but before leukemic progression<sup>17</sup>. Low and intermediate-1 IPSS groups maximized survival with delayed transplantation, especially in patients <40y old, and outcomes were better with transplantation prior to leukemic transformation. Patients in Int-2 and high risk IPSS groups maximized survival with transplantation at diagnosis. An updated cohort study with Markov decision analysis in 2013 using older patients (age 60-70y) stratified by IPSS and reduced intensity conditioning transplant vs nontransplant strategies (basic supportive care, ESAs if anemia, hypomethylating agents for Int-2 and high risk disease) showed improved life expectancy with RIC transplant for int-2 and high risk MDS, and longer life expectancy with non-RIC treatments for low and int-1 disease<sup>18</sup>.

*Bridge to HCT.* The use of hypomethylating agents provide further options for care and potentially for bridge to transplantation and cytoreduction. Several case series using azacitidine as bridge to transplantation shows this treatment is feasible; effect on transplant outcomes is being determined<sup>19-21</sup>. There is no randomized controlled data supporting outcomes for bridging vs direct to transplant approach; two meta-analyses support going direct to transplant for very high or high risk MDS without high blast count i.e. >10% if patients can get to transplant in 1-2 months, however this time frame is not often feasible<sup>22,23</sup>. An EBMT retrospective review of 209 patients with higher risk MDS showed that outcomes at 3 years were not significantly different between patients treated with hypomethylating agents or chemotherapy prior to HCT with respect to OS (42% versus 35%), RFS (29% versus 31%), cumulative incidence of relapse (45% versus 40%), and NRM (26% versus 28%), despite younger age and a higher proportion of patients with primary refractory disease in the hypomethylating group arm<sup>24</sup>. In patients with very high blast counts >10% and a planned rapid progression to transplant, chemotherapy can provide a faster response and is more likely to result in a CR to help bridge to transplant but has more toxicities. For patients with monosomy 7, del 7q, ≥3 chromosomal abnormalities, or mutation of *TP53*, hypomethylating agents are recommended over intensive chemotherapy or supportive care due to poor response rates to intensive chemotherapy in MDS patients with these features. For patients with high-risk disease, treatment is recommended as a bridge to curative therapy during transplant workup.

*Pre-HCT (pre-conditioning) chemotherapy.* Retrospective studies suggest that patients with excess blasts have better HCT outcomes if they achieve blast count <5% with chemotherapy pre-HCT than if they proceed directly into HCT. However, it is unclear whether this is due to selection bias (patients who died during pre-HCT chemotherapy discounted). Time to transplant may be a better predictor of HCT outcomes than achieving complete remission; priority should be given to early transplant over achieving disease response<sup>25</sup>.

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# Chronic Myelogenous Leukemia

Presented by: Lynn Savoie

## Summary

### Chronic Phase

#### First line Therapy:

- First line therapy is with a tyrosine kinase inhibitor (TKI) (imatinib, dasatinib or nilotinib)
- Molecular monitoring with quantitative PCR (polymerase chain reaction) every 3 months
  - Cytogenetics and mutation analysis as per the chronic myeloid leukemia (CML) treatment guidelines
  - Assess milestones as per provincial CML treatment guidelines

#### Subsequent Therapy:

- Adjust TKI therapy as per CML treatment guidelines for patients showing resistance or intolerance to first-line therapy, options include imatinib, dasatinib, nilotinib, bosutinib, ponatinib or asciminib
- In patients on a second generation TKI showing warning signs, who experience a suboptimal response or failure and are otherwise transplant eligible, perform human leukocyte antigen (HLA) typing of patient and siblings followed by a search for a volunteer unrelated donor (VUD) if no suitable family member is identified
- Transplantation should be undertaken for eligible patients who fail to meet the provincial treatment guideline milestones for response to two second generation TKIs, ponatinb and/or asciminib
- Transplantation should be undertaken in eligible patients who are unable to tolerate tyrosine kinase inhibitors such that compliance becomes an issue
- Patients found to carry the T315I mutation should receive ponatinib or asciminib during the donor search and workup periods

### Blast Phase

- HLA type patients and siblings and proceed with VUD or alternate donor search if no family match identified
- Attempt to induce CP2 prior to allogeneic stem cell transplantation with chemotherapy and TKIs
- Transplantation is contraindicated in blast phase

### Monitoring for Relapsed/Refractory CML Post Transplantation

- Quantitative peripheral blood PCR for bcr/abl transcript every 3 months for 2 years then every 6 months to 5 years and then yearly to coincide with scheduled follow up appointments



## **Treatment of Relapsed Disease Posttransplant**

### **Molecular Relapse or Relapse in Chronic Phase:**

- Minimize immunosuppression
- Initiate therapy with at least a second generation TKI, preferably one not used pre-transplant, if a positive nested PCR is detected at  $\geq 6$  months post-transplant or if the BCR/ABL ratio is rising prior to that
- If no response to TKI a mutation analysis should be sent prior to proceeding to escalating doses of TKI

### **Blast Phase Relapse:**

- Minimize immunosuppression
- Perform mutation analysis
- Re-induce chronic phase prior to a second transplant in eligible patient– overall prognosis poor; palliation is a reasonable choice

## **Background**

Chronic myelogenous leukemia makes up 14% of new leukemias, with a median age of 67 years. It is associated with the Philadelphia chromosome t(9;22) and p190, p210 or p230 bcr/abl fusion proteins. The Philadelphia chromosome is found in multiple cell lineages including granulocyte, erythroid, megakaryocyte, and B lymphocyte lineages. Progression of disease is often associated with cytogenetic evolution with common additional abnormalities including +Ph, +8, i(17q) and +19.

### **Natural History of CML**

The natural history of CML involves a chronic phase, and blast phase. Prior to tyrosine kinase inhibitor (TKI) therapy, without stem cell transplantation progression to blast phase occurred on average 3-5 years after diagnosis. In the pre-imatinib era, with sudden onset of blast crisis pre-imatinib in 0.4% of patients in the first year, 1.8% in the second year, and 2.6% in the third year<sup>1</sup>. In the tyrosine kinase era life expectancy approximates normal<sup>2</sup>.

### **Accelerated Phase: World Health Organization (WHO) Classification<sup>3</sup>**

- Blasts 10-19% in peripheral blood or bone marrow
- Peripheral blood basophils  $\geq 20\%$
- Persistent platelets  $< 100/\text{nl}$  unrelated to therapy or  $> 1000/\text{nl}$  unresponsive to therapy
- Increasing spleen size and/or white blood cell count unresponsive to therapy
- Clonal cytogenetic evolution

### **Blast Phase: WHO Classification<sup>3</sup>**

- Blasts  $\geq 20\%$  in peripheral blood or bone marrow
- Extramedullary blast proliferation
- Large foci or clusters of blasts in bone marrow biopsy

## **Treatment**

### **Front Line Treatment**

Treatment with a TKI as first-line treatment for all newly diagnosed CP-CML patients is recommended. Currently in Alberta therapy is begun with imatinib, dasatinib or nilotinib. The choice of TKI should be guided by an individual patient's comorbidities. Patients having achieved their therapeutic milestones with and tolerant of a TKI should continue on it.

Peripheral blood Q-RT-PCR should be performed every 3 months. If a molecular response greater than 4.7-log reduction (MMR) is reached and stable for 2 years, the frequency of Q-RT-PCR may be decreased to every 4-6 months. The recommended definition of first-line optimal treatment response to tyrosine kinase inhibitors (TKIs) in accordance with European Leukemia Net guidelines<sup>4</sup>, are defined as:

- BCR-ABL1  $\leq 10\%$  (at least a 1-log reduction)
- BCR-ABL1  $< 1\%$  (2-log reduction) and or Ph+ 0 at 6 months
- BCR-ABL1  $\leq 0.1\%$  ( $\geq 3$ -log reduction) at 12 months, and thereafter

A second-generation TKI (nilotinib, dasatinib or bosutinib) is recommended for patients with imatinib resistance/ intolerance, or who fail to achieve any of the treatment milestones while on imatinib. The choice of a second-generation TKI should be guided by an individual patient's comorbidities. The presence of specific mutations will override other considerations when determining the optimal agent to employ. Asciminib, an allosteric inhibitor of the myristoyl site of the BCR-ABL protein, is available in the third line for patients who have failed or are resistant to at least 2 prior traditional TKI's. The third-generation drug, Ponatinib, is also available for patients in whom other TKI therapy is not appropriate, including CML that is T315I mutation positive or when there is resistance or intolerance to all other TKI therapy.

Human leukocyte antigen (HLA) typing of the patient and siblings is recommended when a patient presents in AP or BC or when there is suboptimal response, loss of a previously obtained response or significant intolerance.

### **Syngeneic Transplantation for CML**

Although not commonly used, syngeneic transplantation provides evidence that graft-versus-leukemia effect is useful but not necessary for the cure of CML with high dose chemotherapy. A 1982 series of 22 patients, including 12 in chronic phase, resulted in 7 of 12 patients alive at 20-26 years<sup>5</sup>.

Syngeneic transplants remain a viable option for a small number of patients, especially without other

donor options. Registry analysis shows a much higher relapse rate of 40% compared to 7% in allogeneic transplantation thought secondary to lack of graft versus leukemia effect. Supporting the importance of this effect is the higher relapse rate in T-cell depleted transplants and effectiveness of donor lymphocyte infusion (DLI). However, toxicities due to GVHD in syngeneic transplants are minimal.

### **Allogeneic Transplantation for CML**

Allogeneic transplantation is a potentially curative modality for CML associated with increased toxicity up front compared to non-transplant therapy. An IBMTR (International Bone Marrow Transplant Registry) comparison of allogeneic stem cell transplantation with German CML Study Group trials using hydroxyurea or interferon showed that in the first 18 months the relative risk of death with transplant was 5.9, with similar mortality between the two groups between 18 and 56 months, and lower overall mortality with transplant after 56 months<sup>6</sup>. Seven-year survival was higher in the transplant group (58% versus 32%). Registry data reveal a 5-year survival post-transplant of 50 to 70% for matched related donor transplants and 40 to 60% for unrelated donors<sup>7</sup>. Advanced disease is associated with poor outcomes in allogeneic matched sibling transplantation; survival at 3 years with BuCy2 was 58% in chronic phase versus 41% in accelerated phase and 25% in blast phase, with relapse in 3%, 12%, and 27% of patients in each group<sup>7</sup>.

The importance of obtaining a second chronic phase in patients in blast crisis pretransplant was seen in a small trial randomizing 10 patients to upfront allogeneic transplantation and 10 patients to induction chemotherapy followed by allotransplant<sup>8</sup>. All 10 patients transplanted in blast crisis died; 8 of 10 given induction chemotherapy achieved a second chronic phase, 7 patients were transplanted, and all of the 6 patients in the second chronic phase at the time of transplant achieved molecular remission. Median OS in this group was 23 months versus 6 months in those transplanted up front.

Data using the FLUBUP (fludarabine + busulfan) protocol in the first 21 CML patients in Calgary show a projected 3-year OS of 86% with FLUBUP/ATG (anti-thymocyte globulin), compared to a 3-year OS of 76% with the BuCy (busulfan + cyclophosphamide) protocol (p-value not significant). Transplant-related mortality at 3 years was 0% compared to 24% with BuCy (p=0.03). Further data is being accrued.

### **Allotransplants in the Post TKI Era**

There is no evidence that transplant outcomes are worse in patients who have received prior tyrosine kinase inhibitors. An IBMTR analysis of 409 patients transplanted with prior imatinib exposure (9% imatinib intolerance, 37% imatinib failures, remainder planned transplants up front) and 900 patients without imatinib exposure revealed that in patients transplanted in first chronic phase, prior imatinib was associated with better overall survival, and no difference in transplant-related mortality, relapse, or leukemia-free survival.<sup>9</sup> This was confirmed in a matched pairs analysis. In patients with advanced CML, there was no difference between groups in transplant-related mortality, relapse, leukemia-free survival, and overall survival. No difference was seen in rates of acute GVHD. A single institution

study of 12 patients receiving a second generation TKI after imatinib failure showed no negative impact on transplant engraftment, relapse rate of transplant-related toxicity when compared to historical controls<sup>10</sup>.

A report by the Swedish CML registry<sup>11</sup> reviewed 118 patients transplanted between 2002 and 2017. 47.4% received an allo-HSCT in first CP. TKI resistance was the most common transplant indication (62.5%). For patients diagnosed with CML in CP at <65 years of age, the cumulative probability of undergoing allo-HSCT within 5 years was 9.7%. Overall 5-year survival was 96.2%, 70.1% and 36.9% when transplanted in first CP, second or later CP, and in accelerated phase or blast crisis, respectively. Non-relapse mortality for patients transplanted in CP was 11.6%.

Timing of Transplantation

Multiple studies showed better outcomes in the pre-imatinib era if patients are transplanted in the first year after diagnosis. For example, in one study, patients transplanted within one year of diagnosis in chronic phase had a survival of 70% compared with 40% when transplanted beyond one year<sup>12</sup>. In the TKI era, early transplantation is no longer undertaken in patients meeting their milestones. In patients not meeting their milestones a study has shown that the degree of response immediately before transplantation was directly associated with worse outcomes for patients with only a hematological response compared with patients with a cytogenetic or molecular response advocating for the best response possible prior to transplantation<sup>12</sup>.

Blood versus Marrow Stem Cell Source

Less relapse is seen in patients treated with peripheral blood stem cells (PCR positivity 44% with bone marrow versus 7% with peripheral blood at 4 years, p<0.009) but more chronic GVHD with peripheral blood<sup>13</sup>. Overall survival has been higher in peripheral blood transplants than bone marrow stem cell sources. *In vivo* T cell depletion with ATG decreases GVHD. The impact that ATG makes on altering relapse and GVHD outcomes between peripheral blood and bone marrow with the FLUBUP protocol is not fully understood.

Prognostication Pre-Allotransplant – EBMT Transplant Risk Score<sup>14</sup>

Table 2. European Group for Blood and Marrow Transplantation risk factor assessment

| EBMT Risk Factor Assessment <sup>4</sup> |                    |    |                |    |                          |    |    |     |
|------------------------------------------|--------------------|----|----------------|----|--------------------------|----|----|-----|
| Points                                   | 0                  |    | 1              |    | 2                        |    |    |     |
| Risk Factors                             |                    |    |                |    |                          |    |    |     |
| Age                                      | <20 years          |    | 20-40 years    |    | >40 years                |    |    |     |
| Stage                                    | 1 <sup>st</sup> CP |    | AP             |    | BP or 2 <sup>nd</sup> CP |    |    |     |
| Donor                                    | HLA sib            |    | MUD            |    |                          |    |    |     |
| Sex Match                                | All others         |    | Female to Male |    |                          |    |    |     |
| Time to Therapy                          | <12 months         |    | >12 months     |    |                          |    |    |     |
| Points                                   | 0                  | 1  | 2              | 3  | 4                        | 5  | 6  | 7   |
| TRM                                      | 20                 | 23 | 31             | 46 | 51                       | 71 | 73 | N/A |
| OS                                       | 72                 | 70 | 62             | 48 | 40                       | 18 | 22 | N/A |

Abbreviations: AP = accelerated phase; BP = blast phase; CP = chronic phase; EBMT = European Group for Blood and Marrow Transplantation; TRM = transplant-related mortality; MUD = matched unrelated donor

## **Molecular Monitoring Post HCT for CML**

A retrospective review of 346 patients followed with PCR every 3 months post-transplant found that while in the first 3 months post-transplant, PCR positivity did not correlate with worse outcome. At 6 months or later, it was highly correlated with relapse (42% PCR+ relapse versus 3% PCR-,  $p<0.0001$ ; 4-year OS 74% versus 93%,  $p=0.002$ )<sup>15</sup>. Between 6 and 12 months, the PCR+ patients had a relative risk of relapse of 26.0. However, at greater than 36 months, the short-term risk of relapse was much less; 15/59 were qualitative PCR+ but only 1 patient relapsed.

Quantitative PCR can be helpful in predicting relapse risk; at 3 to 5 months post-transplant, increasing PCR positivity is associated with increased risk of relapse. Relapse risk is 17% if PCR-, 43% if low level PCR+ ( $<0.02\%$ ), and 86% if PCR+ is  $>0.02\%$ )<sup>16</sup>. In a study of 379 patients alive at 18 months, 90 had at least 1 positive test at 18 months, but only 14% relapsed (median 40,000 copies/ug) compared to 1% of PCR- patients relapsing (69 had only 1 test positive with mean 24 copies/ug)<sup>19</sup>. In a study of 98 patients, 69 had undetectable, decreasing, or low  $<50$ copies/ug PCR titers and only one relapsed. There was a 72% relapse rate in patients with persistent or high ( $>50$  copies/ug) titers ( $p<0.00001$ )<sup>17</sup>. The correlation between blood and marrow PCR positivity is approximately 90%.

Based on this quantitative peripheral blood PCR for bcr/abl transcript should be done every 3 months for 2 years then every 6 months to 5 years and then yearly to coincide with scheduled follow up appointments.

## **Treatment of Relapsed Disease**

### **Treatment of Relapsed Disease Post-AlloHCT:**

Imatinib is one therapy with moderate effectiveness in advanced relapsed disease post allogeneic HCT; in a review of 28 (5 chronic phase, 15 accelerated phase, 8 blast phase, 13 with previous DLI) imatinib-naïve patients who relapsed post-allotransplant, overall response to imatinib was 22/28, CCR 9/28 (35%), complete molecular response (CMR) 4/28<sup>18</sup>. All chronic phase patients attained CHR compared to 83% of the accelerated phase patients and 43% of the blast phase patients; one year overall survival was 74%. Five patients reactivated GVHD; three had grade III disease.

The second and third generation TKI's and asciminib have not been systemically studied post transplantation but are presumed to be equally if not more efficacious than imatinib. One study published in abstract form only supports this<sup>20</sup>.

DLI is also effective and can induce a complete molecular response in about 70% of patients. These can be durable, with a probability of 80-90% DFS at three years and improvement of OS from 53% without DLI to 95% with DLI at three years ( $p=0.0001$ )<sup>7,16</sup>. There is an approximately 40% chance of GVHD greater than or equal to grade 2 and 30% chance of myelosuppression post-DLI. Responses are not generally durable in second chronic phase disease. The role of imatinib plus DLI is being investigated and a small number of patients have been reported with encouraging results<sup>21</sup>.

A CIBMTR study retrospectively reviewed the outcomes of TKI vs. DLI vs. DLI + TKI in the setting of post-transplant relapse in the TKI era<sup>22</sup>. They found that patients who received a DLI alone had inferior survival compared with those who received a TKI with a DLI. Those who received a TKI alone had similar survival compared with those who received a TKI with a DLI supporting the use of a TKI alone following relapse. Therefore, initiation of TKI therapy with a second generation TKI, preferably one not used pre-transplant should be initiated if a positive PCR is detected 6 months post-transplant or if the BCR/ABL ratio is rising prior to that. If there is no response to TKI alone a mutation analysis should be sent. If there is no mutation proceed to escalating doses of TKI as per the relapse guideline section.

There is very little current data for more advanced phase relapses post-transplant. A mutation analysis should be sent. The next step is the minimization of immunosuppression as well as TKI therapy with- or without DLI should be considered in the accelerated phase potentially as a bridge to second transplant if the patient remains eligible. In the blast phase induction type chemotherapy in conjunction with TKI therapy should be undertaken prior to a second transplant if eligible. In this situation the overall prognosis is poor and palliation is a reasonable choice.



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# BCR-ABL-Negative Myeloproliferative Neoplasms

Presented by: Michelle Geddes

## Summary

- Transplant eligible patients with myelofibrosis (MF) with *intermediate-2 or high-risk* disease according to the Dynamic International Prognostic Scoring System-plus (DIPPS-plus) criteria should be considered for allogeneic stem cell transplantation. This applies to both primary and post-PV/ET MF.
- Medically fit patients with *intermediate-1 risk* can be considered for transplant especially if age <65, with refractory, transfusion-dependent anemia, peripheral blood blasts >2%, or adverse cytogenetics, and should have a donor search performed.<sup>1</sup>
- All patients being considered for allotransplant should have mutation analysis by NGS and cytogenetics performed, if possible, to inform decision-making.
- Patients in blast phase (>20% bone marrow blasts) should be given induction chemotherapy prior to proceeding with stem cell transplantation.
- There is no convincing data to support the requirement for splenectomy or splenic radiation before transplantation. We do not recommend routine splenectomy or splenic irradiation pre-transplant.
- Our standard conditioning is myeloablative busulfan + fludarabine + 4Gy TBI (see Conditioning chapter).
- The use of JAK inhibitors pre-transplant is associated with improvement in constitutional symptoms and performance status, and decrease in spleen size, and can help improve clinical status prior to transplant. JAK inhibitors should be discontinued at the start of the conditioning for HSCT.

## Background

Myeloproliferative neoplasms (MPNs) originate from acquired mutations that target the hematopoietic stem cell and induce dysregulation of kinase signaling, clonal myeloproliferation, and abnormal cytokine expression. The JAK2 V617F mutation is most frequent. Other mutations include CALR, MPL, and other mutations including some with adverse prognostic implications such as ASXL1 (38%), EZH2 (7%), IDH1/2 (45%), SF3B1 (14%) and U2AF1Q157 (8%) mutations<sup>2,3,4</sup>. Patients with triple negative disease status for JAK2, CALR and MPL are recognized to have adverse prognosis.

The 2016 WHO MPN classification is used to diagnose MPNs into categories including polycythemia vera (PV), essential thrombocythemia (ET), primary myelofibrosis (PMF), chronic neutrophilic leukemia, atypical CML, myeloid/lymphoid neoplasms associated with eosinophilia and rearrangements of PDGFRA, PDGFRB, or FGFR1 or with PCM1-JAK2, and overlap syndromes including chronic myelomonocytic leukemia and myelodysplastic/myeloproliferative neoplasm with ring sideroblasts and thrombocytosis.<sup>5</sup>

Hematopoietic cell transplantation is generally considered for patients with myelofibrosis (idiopathic or post PV or ET), and overlap syndromes with poor prognosis.

## Myelofibrosis

Myelofibrosis refers to the MPN classified by the WHO system as primary myelofibrosis or the phenotypically similar condition that develops in the setting of either polycythemia vera (post-PV MF) or essential thrombocythemia (post-ET MF). It is the least common of the three MPNs, with annual incidence of 0.2-1.5 cases/100 000, and carries the worst prognosis, with a median survival of around 6 years.<sup>6</sup> Median age at diagnosis is 65; MF is uncommon in young patients (~20% age <55). It is characterized by marrow fibrosis, myeloid proliferation and abnormal megakaryocyte morphology/clustering, splenomegaly, leukoerythroblastosis, and extramedullary hematopoiesis. Ultimately, this disease results in one of two outcomes: leukemic transformation or bone marrow failure. Currently, allogeneic stem cell transplantation is the only curative option, as all other available treatments are considered palliative.

### Prognostic factors in myelofibrosis:

DIPPS can be used at any time in the disease course, and includes factors such as age >65 years, hemoglobin level <100 g/L, leukocyte count >25 x10<sup>9</sup>/L, circulating blasts ≥1%, and presence of constitutional symptoms; this was upgraded to DIPSS-plus to incorporate three additional independent risk factors, including red cell transfusion need, platelet count <100 x10<sup>9</sup>/L, and unfavorable karyotype (includes complex karyotype, or 1-2 abnormalities that include +8, -7/7q-, i(17q), inv(3), -5/5q-, 12p-, or 11q23 rearrangement).<sup>7,8</sup> The eight DIPSS-plus risk factors are used to define low, intermediate-1, intermediate-2, and high risk groups, as described in the table below. A link to a DIPSS calculator is [DIPSS Prognosis in Myelofibrosis | QxMD](#).

### Dynamic International Prognostic Scoring System - plus (DIPSS):

| Risk Factors                                                                                                                                                                                                                                                                                                     |                        |                   |
|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------|-------------------|
| Age > 65<br>Hemoglobin < 100 gm/L<br>Constitutional symptoms<br>Leukocytes > 25 x 10 <sup>9</sup> /L<br>RBC transfusion requirement<br>Platelets < 100 x 10 <sup>9</sup> /L<br>Unfavourable karyotype (complex or including -5/5q-, -7/7q-, +8, abnormal 11q23, inv(3), 12p-, i(17q))<br>Circulating blasts > 1% |                        |                   |
| Prognostic Group                                                                                                                                                                                                                                                                                                 | Number of Risk Factors | Median OS (years) |
| Low                                                                                                                                                                                                                                                                                                              | 0                      | 15.4              |
| Intermediate-1                                                                                                                                                                                                                                                                                                   | 1                      | 6.5               |
| Intermediate-2                                                                                                                                                                                                                                                                                                   | 2-3                    | 2.9               |
| High                                                                                                                                                                                                                                                                                                             | ≥ 4                    | 1.3               |

## MIPSS70 and MIPSS70-plus include molecular data for prognosis:<sup>9,10</sup>

Recently, integrated clinical, genetic and molecular models with (MIPSS70-plus v.2) or without (MIPSS-70) cytogenetics have been developed that better risk stratify patients who are transplant eligible. A link of a D-IPSS calculator is [MIPSS70 score | MIPSS70-plus version 2.0 score](#).

| MIPSS-70                            | MIPSS-70 Plus v.2:                             |
|-------------------------------------|------------------------------------------------|
| Hemoglobin <100 g/L (1)             | Severe anemia <800 g/L                         |
| WBC >25 x 10 <sup>9</sup> /L (2)    | Moderate anemia (Hb 80-100 g/L)                |
| Platelets <100 g/L(2)               | Leukocytosis >25 x 10 <sup>9</sup> /L          |
| Circulating blasts ≥ 2% (1)         | Thrombocytopenia (<100 x 10 <sup>9</sup> /L)   |
| Bone marrow fibrosis ≥ 2 (1)        | Circulating blasts ≥ 2%                        |
| Constitutional symptoms (1)         |                                                |
| Absence of CALR type 1 mutation (1) | Bone marrow fibrosis grade ≥2                  |
| Presence of HMR mutation* (1)       | Constitutional symptoms                        |
| Presence of ≥ HMR mutations* (2)    | Absence of CALR type 1/like mutation (2)       |
|                                     | Presence of HMR mutation <sup>1</sup> (1)      |
|                                     | Presence of ≥ 2 HMR mutations <sup>2</sup> (2) |
|                                     | Unfavorable karyotype <sup>3</sup> (3)         |
|                                     | Very high risk karyotype                       |
| MIPSS-70 Survival:                  | MIPSS-70 Plus Survival:                        |
| Low (0-1): 27.7 years               | Low (0-2): 20.0 years                          |
| Intermediate (2-4): 7.1 years       | Intermediate (3): 6.3 years                    |
| High (≥ 5): 2.3 years               | High (4-6): 3.9 years                          |
|                                     | Very high (≥7): 1.7 years                      |

<sup>1</sup>HMR: High molecular risk mutations: ASXL1, EZH2, SRSF2, IDH1/2, U2AF1.

<sup>2</sup> Unfavorable karyotype: any abnormal karyotype other than normal karyotype or sole abnormalities of 20 q-, 13q-, +9, chromosome 1 translocation/duplication, -Y, or sex chromosomes other than -Y.

<sup>3</sup>single/multiple abnormalities of -7, i(17q), inv(3)/3q21, 12p-/12p11.2, 11q-/11q23, or other autosomal trisomies not including +8, +9 (eg +21, +19); "Favorable": normal karyotype or sole abnormalities of 13q-, +9, 20q-, chromosome 1 translocation/duplication or sex chromosome abnormality including -Y; "Unfavorable": all other abnormalities

A personalized MPN prediction model was developed in an analysis of a total of 2035 patients and then validated with an external cohort. This calculator is found at <https://cancer.sanger.ac.uk/mpn-multistage/> and can be used to try to individualize prognosis and inform decisions about potential transplantation.<sup>11</sup>

## Genetics-based Prognostic Scoring System (GPSS):

Patient prognosis based on the MIPSS can be augmented by the genetics-inspired prognostic scoring system (GPSS), which incorporates high-risk karyotypes: 5q-, +8, inv(3), i(17q), -7/7q-, 11q or 12p

abnormalities, autosomal trisomies (except +9), monosomal and complex non-monosomal karyotypes. High risk GPSS was also associated with higher blast transformation rate (HR 7.4, 95% CI 2.1-26.3). In patients who had ambiguous prognosis based on differing scores by GPSS and by DIPSS-plus, those found to be higher risk on the GPSS were associated with poorer survival outcomes.<sup>12</sup>

### Secondary Myelofibrosis:

None of the above prognostic tools were developed in patients who have secondary myelofibrosis after progression from polycythemia vera or essential thrombocythosis. In practice, the tools above are used but in a non-validated setting. There has been a model for prognosis developed in patients post-PV and post-ET: Myelofibrosis Secondary to PV and ET-Prognostic Model (MYSEC-PM), based on multivariate analysis of 685 patients with a median survival of 9.3 years. Secondary MF patients were divided into four risk categories based on: hemoglobin, circulating blasts, CALR status, platelet count and constitutional symptoms. Median survival according to risk group was: low (survival not reached), intermediate-1 (9.3 years), intermediate-2 (4.4 years), and high risk (2 years).<sup>13</sup> A calculator is available for the MYSEC-PM: <http://www.mysec-pm.eu/>

**Table 5: Myelofibrosis Secondary to PV and ET-Prognostic Model (MYSEC-PM).**

| Secondary MF: MYSEC-PM       |
|------------------------------|
| <b>Risk factor (Points):</b> |
| Hemoglobin <110 (2)          |
| Circulating blasts ≥ 3% (2)  |
| CALR UNMUTATED (2)           |
| Platelets <150 (1)           |
| Constitutional symptoms (1)  |
| <b>Median Survival:</b>      |
| Low risk (points): NR        |
| Int-1 (points): 9.3 yrs.     |
| Int-2 (points): 4.4 yrs.     |
| High (points): 2 years       |

### Transplantation outcomes in myelofibrosis:

Allogeneic HCT is currently the only treatment option in myelofibrosis that is capable of inducing complete hematologic, cytogenetic, and molecular remissions. However, there are associated risks of treatment-related mortality, graft failure, and disease relapse.

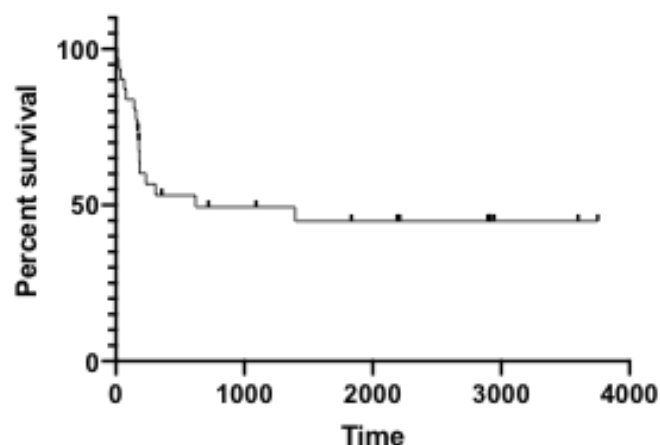
*A multicentre analysis of 100 consecutive transplants for myelofibrosis in patients treated with JAK inhibitors found that 2-year overall survival (OS) was 61%, but 91% for those who experienced clinical improvement pretransplant on JAK inhibitors, and 32% for those with leukemic transformation on JAK inhibitor therapy.*<sup>14</sup> Age of the patients was 32-72 years (median 59). Response to JAK

inhibitor ( $p=0.03$ ), DIPSS score before starting JAK inhibitor ( $p=0.003$ ), and donor type ( $p=0.006$ ) were independent predictors of survival. Intensity of therapy is important in this disease and comparison of nonmyeloablative vs reduced intensity conditioning protocols showed higher levels of graft failure and poorer outcomes with nonmyeloablative regimens.<sup>15</sup> A retrospective analysis of 217 patients given reduced intensity conditioning regimens including Bu 3.2mg/kg vs 6.4mg/kg with fludarabine 30 g/m<sup>2</sup> daily for 4 days showed no difference in outcomes between the two regimens.<sup>16</sup> Age and comorbidities affect outcomes and need to be considered as with transplants for other indications.

**Comparison of HCT vs non-HCT.** There are no randomized trials to compare outcomes in patients treated with JAK1/2 inhibitors vs transplantation. However, a retrospective review of 443 patients under the age of 65, not treated with JAK inhibitors, who received transplant or nontransplant (censored at time of transplant) therapy showed a survival benefit to transplant with int-2 or high risk disease, and this is used by most guidelines as indication for transplant.<sup>17,18</sup> The RR of mortality in patients receiving allogeneic SCT vs conventional therapies was 5.6 (95% CI, 1.7-19;  $P = .0051$ ) for low-risk DIPSS, 1.6 (95% CI, 0.79-3.2;  $P = .19$ ) for int-1 risk, 0.55 (95% CI, 0.36-0.83;  $P = .005$ ) for int-2 risk, and 0.37 (95% CI, 0.21-0.66;  $P = .0007$ ) for high-risk DIPSS patients. Comparison of 5-year OS between the transplant and nontransplant cohorts showed 69% and 95% for low-risk, 52% and 77% for int-1, 50% and 41% for int-2, and 32% and 11% for high-risk patients.

Analysis of retrospective data does not provide clear support for splenectomy prior to transplantation to improve engraftment or outcomes.<sup>19</sup>

**Survival in patients transplanted for myelofibrosis 2009-2018**



Kaplan Meier curve for outcomes in 31 patients transplanted for myelofibrosis in Calgary 2009-2018. Four of the 31 patients relapsed in the follow up period.

### **Polycythemia Vera and Essential Thrombocythemia**

Hematologic transformations towards myelofibrosis and/or acute leukemia, although uncommon, represent a major cause of death in these disorders. In the case of ET, risk of myelofibrotic



transformation increases with disease duration, affecting 3-10% in the first decade after diagnosis and 6-20% in the second decade. Progression to acute leukemia occurs in a small minority of patients, with incidences of 1-2.5% in the first decade after diagnosis, and 5-8% in the second decade, and continuing to increase thereafter. Similar patterns are seen with PV, with leukemic transformation reported as high as 20%. The use of cytoreductive therapy, including alkylating agents, is known to increase the rate of leukemic transformation, and thus the true rate of transformation is unknown. Very little literature exists of transplantation for these diseases, usually in the form of case reports. Prognosis with DIPSS-plus score is not validated in this population although it is commonly used. The problems and complications associated with myelofibrotic transformation of either ET or PV are similar to de novo PMF, thus therapy of post-ET MF or post-PV MF should be approached in the same manner.

### **Use of JAK2 Inhibitors Prior to HSCT for Myelofibrosis**

The JAK2V617F activating kinase mutation is seen in the many patients with BCR-ABL1 negative myeloproliferative patients, and Ruxolitinib, an oral JAK1/JAK2 inhibitor, is approved for the treatment of patients with symptomatic myelofibrosis, based on the data from two randomized phase 3 studies. Treatment is effective in patients without this specific mutation as other mutations in this pathway also cause symptoms. COMFORT-I and COMFORT-II compared ruxolitinib with placebo and best-available therapy (BAT), respectively, and found significant reductions in splenomegaly and improvement in constitutional symptoms.<sup>20,21</sup> Increased caloric intake and enhanced performance status as a result of improved constitutional symptoms and reduced splenomegaly could contribute to improved survival estimates for patients treated with ruxolitinib (71% vs. 54%, HR 0.48).<sup>22,23</sup> Longer follow-up will be needed.

It has been postulated that the anti-JAK2 mediated reduction in both cytokines and splenomegaly, as well as improvement in performance status, might improve outcome after allogeneic HSCT in patients with myelofibrosis. Some patients improve performance status and become transplant eligible. The down-regulation of inflammatory cytokines might have a beneficial impact on graft failure and has been seen to provide benefit in acute GVHD. The largest retrospective study examining transplant outcomes post ruxolitinib is outlined above and shows that pretransplant ruxolitinib therapy is feasible and patients responding to ruxolitinib have overall better transplant outcomes.<sup>14</sup>

There were concerns that abrupt discontinuation of ruxolitinib in advance of transplant may result in cytokine storm reaction and severe inflammatory response. Preliminary reports from the JAK ALLO trial<sup>24</sup> of ruxolitinib prior to HSCT included ten patients who discontinued ruxolitinib, 7 of whom developed life-threatening events (including cardiogenic shock, tumor lysis syndrome, severe GVHD), with two deaths within 3 weeks of drug withdrawal. This pattern has not been seen in subsequent studies; the retrospective series of 100 patients above showed two with significant adverse events after they stopped drug more than 6 days pretransplant.<sup>14</sup> For this reason it is recommended to continue JAK1/2 inhibitors until the day before conditioning.<sup>18</sup>

The average time to treatment failure with JAK 1/2 inhibitors in myelofibrosis is between two and three years. Ideally, patients should be referred for consideration of HSCT before they lose their response to these agents in order for them to undergo transplantation during a time of relatively good health. Several factors have been associated with a short (less than one year) time to treatment failure. These factors include “triple negative” myelofibrosis (negative for JAK2, MPL and CALR mutations) and ASXL-1 and EZH2 mutations, a high DIPSS-Plus score and those requiring transfusions at the time JAK 1/2 inhibitors are started. Patients with any of these risk factors should be referred at the time JAK1/2 inhibitors are started so that they can proceed to HSCT within one year or sooner. Patients with mutated CALR, 0-2 subclonal mutations without ASXL-1 or EZH2 mutations and those with mismatched donors should be followed closely and transplanted at the first sign of progression.<sup>25</sup>

Fedratinib, another JAK2 inhibitor selective for JAK2 relative to other kinases has now been approved by Health Canada and funding recommendations made by CADTH for treatment of splenomegaly and/or disease-related symptoms in adult patients with intermediate-2 or high-risk primary myelofibrosis, post-polycythemia vera myelofibrosis, or post-essential thrombocythemia myelofibrosis, including patients who have been previously exposed to ruxolitinib. It is recommended for patients who are intolerant of ruxolitinib and not for patients with progressive symptoms after ruxolitinib therapy. Approval was made on the basis of the JAKARTA phase 3 randomized MF study for intermediate 2 and high risk disease with placebo and JAKARTA-2 in patients previously treated with ruxolitinib and who were refractory or intolerant. Reduction in spleen volume by 35% was seen in 30-40% of patients and there was improvement in Total Symptoms Score by ≥50% in one third of patients. Thiamine levels are required and supplementation is necessary, and monitoring for symptoms of encephalopathy is required. Other side effects include cytopenias and GI intolerance that should be proactively managed. Current data supports fedratinib’s use post ruxolitinib failure if funding is available. For patients planned for allogeneic stem cell transplant, transplant should occur earlier in the course of disease prior to multidrug resistance and before second line therapy is required for refractory disease. If fedratinib is used frontline prior to stem cell transplant, there is no evidence around management of medication prior to BMT and we would discontinue medication as per ruxolitinib protocol.

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# Chronic Lymphocytic Leukemia (CLL)

Presented by: Robert Puckrin and Mona Shafey

## Summary

- Factors favoring allogeneic stem transplantation over conventional therapy include higher disease risk (high risk cytogenetics, MRD positive, short duration of response, Richter transformation) *and* low transplant risk (younger age, lack of co-morbidities, well-matched donor)
- Allogeneic stem cell transplantation may be offered to CLL patients with:
  - Relapse after at least 1 prior novel agent (BTKi and/or BCL2 inhibitor), preferably responding to a second agent as salvage treatment
  - Richter transformation responding to treatment, particularly if high-risk features such as previously treated CLL, TP53 mutated or clonally related Richter transformation, relapsed disease, or failure to achieve complete response to R-CHOP
- Autologous stem cell transplantation is not indicated to treat CLL but may be considered in selected patients with chemosensitive Richter transformation

## Background

Chronic lymphocytic leukemia (CLL) represents one of the most common lymphoid malignancies of adults. With a median age at diagnosis of 70 years, many patients with this disease will die of other causes. For young patients however, this diagnosis represents a serious threat to life and aggressive management with high-dose therapy and blood stem cell transplantation (SCT) is a reasonable treatment option. This is particularly the case for patients whose CLL is associated with deletion chromosome 17p13.1 [del(17p)], which is observed in 5% of untreated CLL cases but in up to 30% of relapsed and refractory cases. CLL with del(17p) usually require therapy within 1 year of diagnosis and is now being treated with frontline BTK inhibitors such as ibrutinib. Even novel agents such as ibrutinib do not control del(17p)-associated CLL for long durations of time. A recent study by O'Brien and colleagues involving 144 patients with relapsed del(17p) CLL reported 2-year progression-free survival (PFS) rates of approximately 60% (mPFS of 30mo) and 24-month OS of 75%<sup>1</sup>.

For a review of the diagnosis, staging, prognosis, assessments of patient fitness and response, and current treatment recommendations of the Alberta Provincial Hematology Tumour Team, please refer to the [CLL Clinical Practice Guideline \(LYHE-007\)](#).

## Stem Cell Transplantation in CLL

Data from the Center for International Blood and Marrow Transplant Research (CIBMTR) suggests that CLL is an infrequent indication for transplant. The majority of transplants reported were allogeneic, many of which were carried out after reduced intensity or non-myeloablative conditioning.

## Allogeneic Stem Cell Transplantation in CLL

Long-term survival after allogeneic SCT for CLL of the largest series of patients (n=2589), who underwent transplant between 2000 and 2010, has been reported from the European Group for Blood and Marrow Transplantation (EBMT)<sup>2</sup>. In this series, long-term disease control was established for patients, but with longer follow-up both event-free and overall survival decreased over time (62% at 2 years vs. 35% at 10 years for OS; 49% at 2 years vs. 28% at 10 years for EFS). The incidence of relapse was 21% at 2 years vs. 32% at 10 years, and non-relapse mortality 30% at 2 years vs. 40% at 10 years. The presence or absence of del(17p)/TP53 mutation has not been shown to impact outcome of alloSCT<sup>3,4</sup>. Risk factors for relapse included active disease at time of transplant, T-cell depletion with alemtuzumab, prior autologous SCT, and use of mismatched donor, while absence of MRD at 12 months was highly prognostic for reduced relapse risk<sup>2,3,4</sup>.

In regards to donor selection, the EBMT analyzed 368 chronic lymphocytic leukemia patients who underwent allogeneic hematopoietic stem cell transplantation between 1995 and 2007<sup>5</sup>. There were 198 HLA-identical siblings; among unrelated transplants, 31 were well matched in high resolution ('well matched' unrelated donor, WMUD), and 139 were mismatched (MM), including 30 matched in low resolution; 266 patients (72%) received reduced-intensity conditioning and 102 (28%) received standard. There was no difference in OS at 5 years between HLA-identical siblings (55% (48-64)) and WMUD (59% (41-84)), p=0.82. In contrast, OS was significantly worse for MM (37% (29-48) p=0.005) due to a significant excess of transplant-related mortality. HLA matching had no significant impact on relapse (siblings: 24% (21-27); WMUD: 35% (26-44), p=0.11 and MM: 21% (18-24), p=0.81); alemtuzumab T-cell depletion and stem cell source (peripheral blood) were associated with an increased risk. As the toxicity of haploidentical transplants have been greatly decreased with the use of post-transplant cyclophosphamide, donor availability for transplantation has increased. The EBMT has reported the outcome of 117 CLL patients who had received an allogeneic SCT with a haploidentical donor, with results appearing almost identical to those with HLA-matched donors; 5-yr OS 38%, PFS 31%, CI of relapse 26%, and NRM 44%<sup>6</sup>.

Myeloablative conditioning (MAC) is associated with improved overall survival in acute myeloid leukemia (AML), where the increased risks of NRM and acute and chronic GVHD associated with MAC were offset by the significant reduction in relapse<sup>7</sup>. It is unclear if these results can be extrapolated to patients with CLL, given that CLL is an indolent disease with high susceptibility to GVL which may not necessarily require intensive chemotherapy; patients with CLL may be at higher risk of NRM due to older age and underlying immune dysregulation; allotransplant recipients with CLL are heavily pre-treated which may confer greater resistance to MAC; and there may be more options to treat relapse after allotransplant in CLL, such as donor lymphocyte infusion and emerging therapies (e.g. non-covalent BTK inhibitors, CAR-T cells). Retrospective and transplant registry studies conclude that reduced intensity conditioning (RIC) is associated with lower risks of NRM, less



acute and chronic GVHD, and similar or improved OS compared to MAC in CLL<sup>8, 9, 10, 11, 12, 13</sup>. Indeed, intensive conditioning is not necessarily indicated since CLL has among the lowest relapse risk after non-myeloablative (NMA) conditioning of any hematologic malignancy<sup>14</sup>. An EBMT registry study also concluded that selected patients receiving NMA conditioning have similar relapse incidence, relapse-free survival (RFS), and overall survival as recipients of RIC in CLL<sup>15</sup>. As a result, the majority of allotransplants (75-80%) performed for CLL in North America and Europe employ RIC or NMA conditioning, and ASTCT guidelines recommend RIC or NMA conditioning for all patients undergoing allotransplant for CLL<sup>2, 16, 17</sup>. MAC with FluBu4 was utilized for most allotransplant recipients with CLL in Alberta until 2022, at which time a review of local outcomes revealed high rates of acute and chronic GVHD and low mod/severe chronic GVHD and RFS (cGRFS) despite the use of ATG (Table 3). As a result, in February 2023 RIC was adopted as the preferring conditioning regimen for patients with CLL to reduce their risks of acute and chronic GVHD as well as other acute toxicities and NRM.

**Table 1. Summary of transplant characteristics and survival in prospective studies of RIC HSCT in CLL (modified from Gribben 2018<sup>18</sup>).**

|                          | Fred Hutchinson Cancer Center <sup>8</sup> | German CLL Study Group <sup>9, 10</sup> | MD Anderson Cancer Center <sup>11</sup> | Dana-Farber Cancer Institute <sup>12</sup> |
|--------------------------|--------------------------------------------|-----------------------------------------|-----------------------------------------|--------------------------------------------|
| Number of patients       | 82                                         | 90                                      | 86                                      | 76                                         |
| Conditioning regimen     | Flu/low-dose TBI                           | Flu/Cy ± ATG                            | Flu/Cy ± R                              | Flu/Bu                                     |
| Donors, % sibling/% MUR  | 63/37                                      | 41/59                                   | 50/50                                   | 37/63                                      |
| Median follow-up, months | 60                                         | 72                                      | 37                                      | 61                                         |
| Median PFS, %            | 39 (at 5 y)                                | 38 (at 6 y)                             | 36 (at 6 y)                             | 43 (at 6 y)                                |
| Median OS, %             | 50 (at 5 y)                                | 58 (at 6 y)                             | 51 (at 6 y)                             | 63 (at 6 y)                                |

Abbreviations: ATG = antithymocyte globulin; BU = busulfan; CLL = chronic lymphocytic leukemia; Cy = cyclophosphamide; Flu = fludarabine; HSCT = hematopoietic stem cell transplantation; MUR = matched unrelated donor; OS = overall survival; PFS = progression-free survival; R = rituximab; RIC = reduced-intensity conditioning; TBI = total body irradiation; y = years.

**Table 2. Summary of recent studies using RIC HSCT in CLL**

| Outcome          | Dana Farber retrospective <sup>19</sup> | Multicenter retrospective <sup>12</sup> | France phase II trial <sup>13</sup> |
|------------------|-----------------------------------------|-----------------------------------------|-------------------------------------|
| No. patients     | 30                                      | 65                                      | 42                                  |
| Conditioning     | FluBu1 or FluBu2                        | Various RIC (95%)                       | FluBu2                              |
| GVHD prophylaxis | CNI + MTX +/- Sir                       | CNI + MTX (84%)                         | ATG+CNI+MTX                         |
| Overall survival | 3-yr 87%                                | 2-yr 81%                                | 3-yr 87%                            |
| PFS              | 3-yr 72%                                | 2-yr 63%                                | 3-yr 63%                            |
| Relapse          | 3-yr 21%                                | 2-yr 27%                                | 3-yr 30%                            |
| NRM              | 3-yr 7%                                 | 2-yr 13%                                | 3-yr 10%                            |
| Mod-severe cGVHD | NR                                      | 27%                                     | 23%                                 |

The EBMT studied 44 patients with 17p- CLL who received allogeneic hematopoietic SCT between March 1995 and July 2006 from a matched sibling (n = 24) or an alternative donor (n = 20)<sup>20</sup>. Patients had received a median of 3 lines of chemotherapy before SCT, and at the time of transplantation, 53% of patients were in remission. RIC was applied in 89% of patients. Acute or extensive GVHD occurred in 43% and 53% of patients, respectively. Nineteen patients were alive at the last follow-up (median observation time 39 months), and no late relapse occurred in 9 patients with a follow-up longer than 4 years. The 3-year OS and PFS rates were 44% and 37%, respectively.

It is clear that with the approval of novel agents including BTK inhibitors (e.g. ibrutinib, acalabrutinib), PI3 kinase inhibitors (e.g. idelalisib), and the BCL2 inhibitor venetoclax, the number of transplants being performed for CLL continues to decline, particularly for those patients without high risk cytogenetics. In a study of 65 patients with prior novel agent exposure who underwent alloSCT, the 24 month PFS, OS, NRM, and relapse incidence was 63%, 81%, 13%, and 27%, respectively<sup>12</sup>. Poor-risk disease characteristics, prior NA exposure, complete vs. partial remission status, and transplant characteristics were not independently associated with PFS; 1 vs.  $\geq$  novel agents, or ibrutinib vs. venetoclax as the line of therapy immediately pre-alloHSCT had no impact on PFS or OS; only HCT-CI independently predicted for PFS. It remains to be seen whether combinations of these agents will alter the natural history of the disease, or whether they are just delaying the use of allogeneic SCT until later in the disease course. Allogeneic SCT will continue to have a role for patients who fail or are intolerant to these therapies.

Overall, allogeneic stem cell transplantation (HSCT) should be considered for fit patients who are younger than 65 years of age and have CLL that did not respond or have progressed after treatment with at least one novel agent (either BTKi and/or venetoclax). This is especially important in patients with higher disease risk, including complex karyotype, high risk cytogenetics (del17p/TP53), short duration of response, and MRD positive (on venetoclax).

### **Richter Transformation:**

Richter transformation (RT) of CLL into an aggressive DLBCL occurs in 1-5% of patients with CLL. RT is associated with a dismal prognosis with PFS <25% with CHOP-based chemoimmunotherapy and median survival 6-12 months<sup>21, 22, 23, 24</sup>. Given these poor outcomes, consolidation with allogeneic HCT should be considered for eligible patients with responding disease who have relapsed RT or other high-risk features, such as previous therapy for CLL, failure to achieve a complete response to R-CHOP, TP53 aberrations, or clonally related RT<sup>24, 25, 26, 27</sup>. Although <10% of all patients with RT will ever undergo allogeneic HCT, this may represent a curative therapy for selected cases with 3-year PFS 43%, OS 52%, relapse incidence 30%, and NRM 27% in a CIBMTR study of 118 allogeneic HCT recipients<sup>28</sup>. Outcomes were best for patients in complete response (3-year PFS 66%) or partial response (3-year PFS 43%) at the time of HCT compared to those with resistant disease (3-year PFS 5%). The majority of patients with RT receive reduced intensity conditioning, which has been

associated with similar to improved outcomes in this setting compared to myeloablative conditioning<sup>28, 29, 30, 31</sup>.

Importantly, patients with RT arising from treatment naïve CLL appear to have comparatively better outcomes with R-CHOP and may not necessarily require allogeneic HCT<sup>23, 24</sup>. In an Alberta study of 99 patients with RT, those with treatment naïve CLL had higher response rates to first line chemoimmunotherapy (71% versus 40%) and superior 2-year OS (51% versus 28%) compared to those with previously treated CLL. Nevertheless, 2-year OS remained suboptimal at 53% for the 13 patients ≤70 years old with RT and treatment naïve CLL, which suggests there may be a role to consider consolidation with autologous HCT for these patients, similarly as other high risk aggressive lymphomas. Although data is lacking on the outcomes of autologous HCT as part of first-line therapy for RT, a CIBMTR study of 53 patients undergoing autologous HCT for predominantly relapsed RT reported 3-year PFS 48%, OS 52%, and relapse incidence 37%<sup>28</sup>. In an EBMT study of 34 patients who underwent autologous HCT, only 11 of 17 relapses were related to RT (the remainder were due to CLL), suggesting autologous HCT may eradicate the RT component in many patients even though the underlying CLL may persist<sup>29</sup>. It should be noted that even if allogeneic HCT may not be required as a part of primary therapy for patients with lower-risk RT, a referral for transplant consultation and HLA typing is suggested at diagnosis in all patients who are eligible for allogeneic HCT by age and/or comorbidities, given the significant risk of relapse/refractory disease with RT.

Less commonly, patients with CLL may develop a Hodgkin lymphoma variant of RT which is often clonally unrelated to the CLL. Available evidence suggests that Hodgkin-variant RT has similar outcomes with standard chemotherapy as de novo Hodgkin lymphoma in this age group<sup>32, 33</sup>. As such, there is not an established role for consolidation with HCT in these cases<sup>27</sup>.

## **Autologous Stem Cell Transplantation in CLL**

Case series from a number of institutions report high overall survival (4-year OS 65-94%) with low TRM (4-10%) of autologous stem cell transplantation (ASCT) for CLL; however, to date, no randomized study has demonstrated an OS advantage for the use of ASCT in CLL. Despite a strong PFS advantage in the published studies of ASCT, ASCT is now rarely used for CLL<sup>17-19</sup>. This is because FCR (fludarabine, cyclophosphamide, and rituximab) is now used as front-line therapy for most young CLL patients, as it has been shown in a randomized, controlled trial, to provide an OS advantage. Published studies of ASCT in CLL predate the introduction of FCR chemotherapy so the role that ASCT could play in the era of FCR is unclear. Additionally, the use of 6 cycles of a fludarabine-containing regimen significantly impairs the subsequent ability to mobilize and collect autologous blood stem cells. With the emergence of novel agents for relapsed CLL, the role of ASCT is even more unclear. At present, there are no definite indications for ASCT for CLL.

**Figure 1A: Progression-free survival and overall survival of patients with CLL undergoing allotransplant in Calgary between 2000-2021**

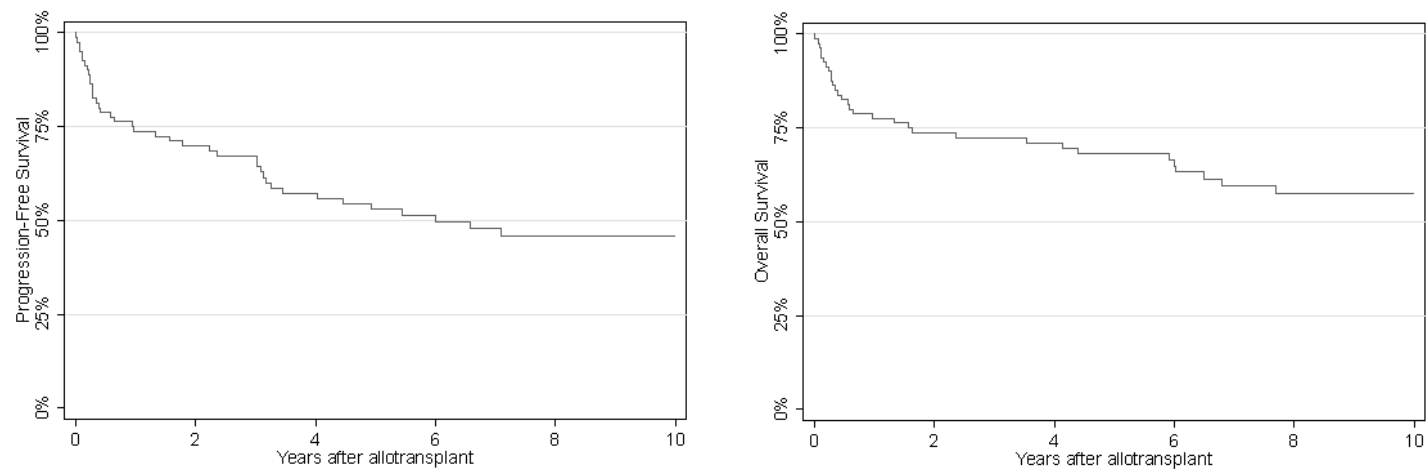


**Figure 1B: Progression-free survival and overall survival of patients with DLBCL-type Richter transformation of CLL undergoing allotransplant in Calgary between 2000-2021**

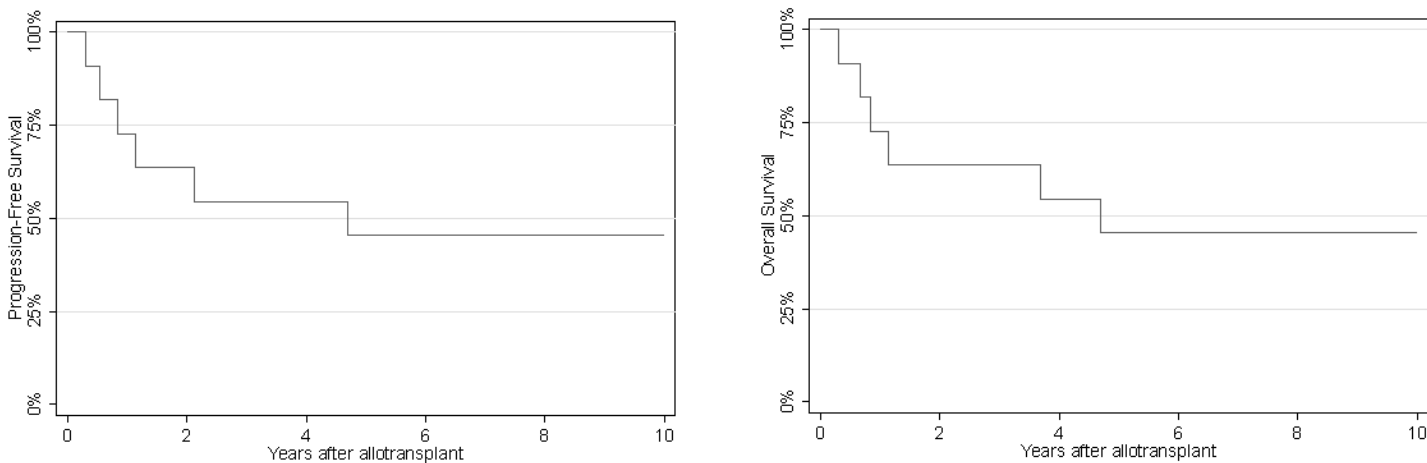


**Table 3. Outcomes of patients undergoing allogeneic HCT with FluBu4 +/- TBI MAC for CLL or DLBCL-type Richter transformation in Calgary between 2000-2021**

| <b>Outcome (at 5 years)</b>     | <b>CLL (n=79)</b> | <b>Richter (n=11)</b> |
|---------------------------------|-------------------|-----------------------|
| Overall survival                | 68%               | 45%                   |
| Progression-free survival       | 53%               | 45%                   |
| Relapse                         | 28%               | 18%                   |
| Non-relapse mortality           | 20%               | 36%                   |
| Grade II-IV acute GVHD          | 43%               | 27%                   |
| Grade III-IV acute GVHD         | 18%               | 9%                    |
| Moderate-to-severe chronic GVHD | 41%               | 45%                   |
| Chronic GVHD and RFS (cGRFS)    | 24%               | 18%                   |

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# Hematopoietic Cell Transplantation/Cellular Therapy for Lymphoma

Presented by: Robert Puckrin

## Summary

### CAR-T Cell Therapy

- Axicabtagene autoleucel is expected to soon be funded as second-line therapy for diffuse large B-cell lymphoma (DLBCL) which is refractory or has relapsed within 12 months of the end of first-line therapy
- Axicabtagene autoleucel, tisagenlecleucel, and soon lisocabtagene maraleucel are funded for relapsed or refractory DLBCL after  $\geq 2$  lines of systemic therapy
  - Eligibility includes most subtypes of DLBCL, including transformed indolent lymphoma and follicular large B-cell lymphoma (previously known as follicular lymphoma grade 3B)
  - Patients with Richter transformation arising from previously-treated chronic lymphocytic leukemia/small lymphocytic lymphoma (CLL/SLL) are not currently eligible
  - Patients with rare subtypes of DLBCL which typically lack B-cell antigens should have demonstration of CD19 expression to be eligible for CD19-directed CAR-T
- Brexucabtagene autoleucel is funded for relapsed or refractory mantle cell lymphoma after  $\geq 2$  lines of systemic therapy including a BTK inhibitor
- Axicabtagene autoleucel is expected to be funded for relapsed or refractory follicular lymphoma after  $\geq 2$  lines of systemic therapy
- Tisagenlecleucel is expected to be funded for follicular lymphoma which is refractory to a second or later line of systemic therapy, relapsed within 6 months after completion of a second or later line of systemic therapy, relapsed during anti-CD20 antibody maintenance (following at least 2 lines of therapies as above) or within 6 months after maintenance completion, or relapsed after autologous HSCT.
- Lymphodepletion is typically with fludarabine and cyclophosphamide.
- Patients must meet the following criteria in addition to the above disease criteria:
  - Clinically stable and expected to remain so until the planned CAR-T cell infusion date with adequate organ function and performance status (ECOG 0-2)
  - No prior treatment with a CAR-T cell product targeting the same antigen
  - Patients with previous or active secondary CNS involvement are eligible for CAR-T but patients with active primary CNS lymphoma are not currently eligible

### Autologous HCT (ASCT) and Allogeneic HCT- Eligibility and Conditioning

- Patients are generally considered eligible for autologous stem cell transplant (ASCT) or allogeneic HCT if they are approximately  $\leq 75$  years old and have controlled disease status (e.g. complete or partial response), a good performance status (e.g. ECOG 0-2), no serious uncontrolled infections, adequate organ function, and  $> 2 \times 10^6$  CD34+ cells/kg (ASCT) or an available stem cell donor (allogeneic HCT)

- Conditioning for ASCT:

|                              |                                                                                          |
|------------------------------|------------------------------------------------------------------------------------------|
| Aggressive NHL (DLBCL, PTCL) | (R) + Bu(13500uM.min) + Mel(140mg/m <sup>2</sup> )                                       |
| Indolent NHL (FL, MZL, LPL)  | (R) + Mel(180mg/m <sup>2</sup> ) + TBI(5Gy x1)                                           |
| Mantle cell lymphoma         | (R) + Mel(180mg/m <sup>2</sup> ) + TBI(5Gy x1)                                           |
| Hodgkin lymphoma             | Gemcitabine 1500 mg/m <sup>2</sup> +<br>Melphalan 200 mg/m <sup>2</sup>                  |
| Primary CNS lymphoma         | (R) + Thiotepa(600mg/m <sup>2</sup> ) +<br>Bu(13500 uM.min)                              |
| Secondary CNS lymphoma       | (R) + Thiotepa(500mg/m <sup>2</sup> ) + Bu(13500uM.min) +<br>Mel (100mg/m <sup>2</sup> ) |

- Reduced intensity conditioning allogeneic HCT is recommended for most patients who have heavily pretreated lymphoma (e.g. relapse after ASCT or multiply-relapsed disease), intrinsically chemoresistant lymphomas (e.g. TP53-mutated mantle cell lymphoma or cutaneous T-cell lymphomas), or indolent lymphomas with high susceptibility to the graft-versus-lymphoma effect (e.g. SLL/CLL or follicular lymphoma).

## Indications for HCT/CT

### Diffuse Large B-Cell Lymphoma:

- Consolidative ASCT may be considered for high-risk patients with DLBCL responding to first-line therapy, such as those with (1) high IPI score 4-5 and partial metabolic response on interim PET or (2) high-grade B-cell lymphoma with MYC and BCL2 rearrangements with IPI score 2-5 (1-5). This practice will require re-evaluation once second-line CAR-T cell therapy and novel first-line therapies are available.
- Once funded, second-line CAR-T cell therapy is recommended for eligible patients with DLBCL refractory to or relapsing within 12 months of the end of first-line chemoimmunotherapy (6-9)
- ASCT is recommended for eligible patients with chemosensitive relapse of DLBCL occurring >12 months after completion of first-line chemoimmunotherapy (10). Examples of appropriate salvage regimens before ASCT include R-DICEP, R-GDP, R-DHAP, or R-ICE (11-15).
- Third-line CAR-T cell therapy is recommended for patients with relapsed/refractory DLBCL after ≥2 lines of therapy who have not previously received CAR-T cell therapy (16-19).
- Allogeneic HCT is rarely performed for DLBCL but may be considered for fit, motivated patients who relapse after CAR-T cell therapy and achieve an adequate response to pre-transplant therapy (20, 21).

### Central Nervous System (CNS) Lymphoma:

- Thiotepa/busulfan-based ASCT is recommended for eligible patients with primary CNS lymphoma responding to high-dose methotrexate and cytarabine based induction (see Lymphoma guideline) (22-27)

- Thiotepe/busulfan/melphalan-based ASCT is recommended for eligible patients with chemosensitive secondary CNS lymphoma (SCNSL), with favorable outcomes observed among those with SCNSL at diagnosis or isolated CNS relapse of systemic lymphoma (see Lymphoma guideline) (28, 29)
- The prognosis of patients with early concurrent CNS and systemic relapse is poor and the optimal treatment is unclear, so these patients should be considered for ASCT or second-line CAR-T cell therapy once funded on a case-by-case basis (28-30).
- Patients with SCNSL undergoing CAR-T cell therapy should receive appropriate CNS-directed bridging therapy before infusion since outcomes appear poor if there is active CNS disease at the time of infusion (30).
- Allogeneic HCT is not well established for CNS lymphoma (31).

### **Transformed Lymphoma:**

- ASCT is recommended as consolidation therapy of R-CHOP for patients with chemosensitive transformed DLBCL if they have previously received chemotherapy for indolent B-cell lymphoma (32, 33)
- Patients who develop transformed DLBCL within 12 months of receiving R-CHOP or R-CEOP for indolent lymphoma may be considered for second-line CAR-T cell therapy once funded (6-9). Otherwise, CAR-T cell therapy is recommended for patients with relapsed/refractory transformed lymphoma after  $\geq 2$  lines of therapy (e.g. 1 for indolent lymphoma and 1 for DLBCL) (16, 18, 19)
- Allogeneic HCT is rarely performed for transformed lymphoma but may be considered for fit, motivated patients who relapse after CAR-T cell therapy and achieve an adequate response to pre-transplant therapy (20, 21, 33).

### **Burkitt Lymphoma:**

- Relapsed Burkitt lymphoma has a poor prognosis but ASCT or rarely allogeneic HCT may be considered for patients with chemosensitive relapses (34)

### **Mantle Cell Lymphoma:**

- Consolidative ASCT is recommended for eligible patients responding to first-line cytarabine-containing chemoimmunotherapy (35-40)
- Maintenance rituximab is recommended every 2 months for 3 years after ASCT (41, 42)
- CAR-T cell therapy is recommended for eligible patients with relapsed MCL after  $\geq 2$  lines of therapy including a BTK inhibitor (43, 44)
- Allogeneic HCT may be considered for fit, motivated patients who relapse after CAR-T cell therapy and achieve an adequate response to pre-transplant therapy (45)
- Patients with TP53-mutated mantle cell lymphoma have poor outcomes with chemotherapy and do not benefit from ASCT (46). These patients should be prioritized for first-line allogeneic HCT or third-line CAR-T cell therapy on a case-by-case basis.

### **Follicular Lymphoma:**

- Given the poor prognosis with standard chemotherapy, ASCT is recommended for eligible patients with chemosensitive relapse of follicular lymphoma occurring within 24 months of first-line treatment (POD24) (47)
- ASCT achieves durable remissions and is also a recommended option for fit patients with first or second relapse of follicular lymphoma arising >24 months after first-line treatment (48-50)
- Once funded, CAR-T cell therapy is a recommended option for eligible patients with relapsed/refractory follicular lymphoma after  $\geq 2$  lines of systemic therapy (51, 52)
- Allogeneic HCT is rarely performed for follicular lymphoma but may be considered for fit, motivated patients who have relapsed after, or are unable to receive, ASCT and/or CAR-T cell therapy and who achieve an adequate response to pre-transplant therapy (53).

### **Other Indolent B-Cell Lymphomas:**

- ASCT may be considered for selected patients with first or second chemosensitive relapse of marginal zone lymphoma (MZL), lymphoplasmacytic lymphoma (LPL), or nodular lymphocyte-predominant B-cell lymphoma (NLPBL), particularly for those with early relapses, aggressive clinical behavior, and in patients who prioritize the possibility of long-term disease control or who lack other treatment options (54-56)
- Allogeneic HCT is seldomly performed for rare indolent B-cell lymphomas but may be considered for fit, motivated patients who lack other therapeutic options and have relapsed after, or are unable to receive, ASCT and achieve an adequate response to pre-transplant therapy (57)

### **Hodgkin Lymphoma (HL):**

- ASCT is recommended for eligible patients with relapsed/refractory HL who respond to second-line chemotherapy or immune checkpoint inhibitors (ICI) (58-62)
- Maintenance brentuximab vedotin (BV) every 3 weeks for 16 cycles starting 4-6 weeks after ASCT is funded for patients with primary refractory HL, relapsed HL <12 months from the end of frontline therapy, or extranodal disease at relapse. The benefits must be weighed against the risks given the lack of survival benefit, increased toxicity, and potential overtreatment of patients already cured by ASCT. The benefit of maintenance BV may be more pronounced in patients with 2-3 risk factors or with a positive PET before ASCT (63, 64).
- Allogeneic HCT is recommended for fit, motivated patients with HL who have exhausted other treatment options (65). Eligible patients should be referred for discussion of allogeneic HCT when starting their second novel agent (i.e. ICI or BV). For patients with highly refractory disease or short duration of remissions, consider using the second novel agent to achieve a response as a bridge to allogeneic HCT. Other patients who achieve a complete response to the second novel agent may reasonably defer allotransplant to the next relapse provided that effective bridging therapy is expected to be available. A washout period of the ICI for 6-12 weeks before allogeneic HCT is recommended to reduce the risk of GVHD (65, 66).

**Peripheral T-Cell Lymphomas (PTCL) and Natural Killer-Cell (NK) Lymphomas:**

- Given the poor prognosis with standard chemotherapy, consolidative ASCT is recommended for eligible patients responding to first-line therapy with advanced stage or high IPI score PTCL NOS, angioimmunoblastic T-cell lymphoma, ALK-negative anaplastic large cell lymphoma (ALCL), advanced-stage NK/T-cell lymphoma, or enteropathy-associated T-cell lymphoma (67-73). Although ALK-positive ALCL usually has a favorable prognosis with BV-CHP, consolidative ASCT may be considered for selected cases with a high IPI score given their poorer outcomes with standard chemotherapy and the uncertain benefit of brentuximab vedotin in this high-risk subgroup (72-75).
- First-line allogeneic HCT is recommended for eligible patients with certain poor prognosis lymphomas responding to first-line therapy, such as hepatosplenic T-cell lymphoma, acute or lymphoma-type adult T-cell leukemia/lymphoma, aggressive NK cell leukemia, and selected cases of advanced-stage NK/T-cell lymphoma (71, 73)
- ASCT is recommended for patients with relapsed PTCL who have not previously received ASCT and who demonstrate a good response to second-line chemotherapy (e.g. DICEP or GDP) (76, 77)
- Allogeneic HCT is also recommended for patients with PTCL who relapse after ASCT or who are unable to receive ASCT due to chemorefractory disease, provided that an adequate response to pre-transplant therapy is achieved (78)

**Cutaneous T-Cell Lymphomas (CTCL):**

- ASCT is not routinely recommended for patients with mycoses fungoides or Sezary syndrome (MF/SS) but allogeneic HCT may be considered for selected high-risk cases (79, 80)



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# Myeloma and Amyloidosis

Presented by: The Myeloma Group

## Summary

(Also see AHS Myeloma Guidelines)

1. For **symptomatic multiple myeloma**, an autologous stem cell transplant (ASCT) should be offered to patients who are ≤65 years old, without significant co-morbidities, and have achieved at least partial response after induction chemotherapy.
  - For individuals between ages 65 and 70 years, ASCT will be considered if the IMWG Frailty index is <1. (<http://www.myelomafrailtyscorecalculator.net/>)
    - i. This is particularly pertinent where upfront use of Daratumumab/Lenalidomide and dexamethasone (DRd) may provide better outcomes than ASCT.
  - We do NOT recommend ASCT for individuals ≥ 70 years.
  - We recommend an early referral (after 2 cycles of induction chemotherapy) to the transplant team for ASCT consideration
2. Collecting enough stem cells for 2 transplants (for salvage, tandem or boost) is recommended but will be discussed on a case-by-case basis
  - Adequate stem cells will be collected for 2 ASCT in patients <60 years and adequate stem cells will be collected for 1 ASCT in patients >60 years.
  - Other considerations include 1) patient factors, 2) disease factors, 3) evolving myeloma therapies, and 3) local resource factors
3. The preferred conditioning chemotherapy for ASCT is Melphalan 200mg/m<sup>2</sup>
  - The use of Busulfan with melphalan conditioning is reasonable
  - Dose reductions of melphalan is reasonable in those with renal dysfunction and/or frailty.
4. The preference is to offer an upfront ASCT post-induction as opposed to an ASCT on relapse.
5. Offering a second ASCT on relapse is reasonable if the disease has been in control for at least 2 years (no maintenance) and at least 4 years (with maintenance).
6. Tandem ASCT is not recommended. However, for patients with high-risk disease that may benefit from this approach, tandem ASCT will be considered on a case-by-case basis.
7. The use of consolidation therapy post-ASCT is recommended in patients with high-risk disease and in patients who achieved < very good partial response (VGPR) after ASCT.
8. The use of maintenance therapy with lenalidomide is recommended post-ASCT until disease progression (in contrast to fixed duration). The use of combination lenalidomide with bortezomib or carfilzomib, but not ixazomib as maintenance is reasonable in patients with high-risk disease.
9. Overall, the use of daratumumab with induction, consolidation and maintenance therapy is strongly recommended in those with access to funding. It would be ideal to have Minimal



Residual Disease (MRD) testing to guide potential discontinuation of therapy after 2 years (PERSEUS study), where up to 65% of patients may achieve this outcome.

10. Allogeneic transplant is not indicated.

11. For **systemic AL amyloidosis**, ASCT may be offered to patients with the following eligibility criteria:

- Age <65 years,
- Performance status (Eastern Cooperative Oncology Group) 0 to 2,
- NT-proBNP is <5000 ng/l and/or cardiac troponin T is <0.06 ng/ml,
- Estimated glomerular filtration rate >30 ml/min per 1.73 m<sup>2</sup> unless on dialysis,
- New York Heart Association class <III, cardiac ejection fraction >45%, systolic blood pressure >90 mm Hg (standing), and
- Lung CO diffusion capacity >50%.

12. Induction chemotherapy for patients with AL amyloidosis should be recommended for patients with >10% monoclonal plasma cells in the bone marrow at presentation.

13. A salvage ASCT for AL amyloidosis will be considered on a case-by-case basis.

# Symptomatic Myeloma

## Background

Multiple myeloma is a chemotherapy-responsive tumor that demonstrates significant dose-response effects. The availability of increased and superior therapeutic options has led to improvements in depth and duration of response. Ultimately, this translates to better Progression-Free Survival (PFS), Overall Survival (OS) and Health related Quality of Life (HRQOL). A more thorough discussion on myeloma care can be found within the AHS Myeloma Guidelines.

Hematopoietic stem cell transplantation involves the use of high-dose chemotherapy followed by an infusion of either autologous, allogeneic, or syngeneic stem cells. Data from the CIBMTR, suggests improvements in myeloma outcomes with autologous stem cell transplantation (ASCT)<sup>1</sup>. In historical randomized controlled trials (RCT), the use of high-dose chemotherapy with ASCT to standard induction protocols demonstrates improved PFS, OS and HRQOL (Table 1).

**Table 1.** Review of Historic RCTs Comparing Standard Therapy (SDT) to High-Dose Chemotherapy (HDT) with ASCT.

| Study                                                  | N   | Age   | SDT versus HDT (p-value) |                       |                       |
|--------------------------------------------------------|-----|-------|--------------------------|-----------------------|-----------------------|
|                                                        |     |       | CR/nCR (%)               | Median EFS/PFS(mths)  | Median OS (mths)      |
| Attal <i>et al.</i> 1996 IFM90 <sup>2</sup>            | 200 | ≤ 65  | 5 vs. 22 (p<.001)        | 18 vs. 28 (p=.01)     | 44 vs. 57 (p=.03)     |
| Ferland <i>et al.</i> 1998 MAG-95 <sup>3</sup>         | 190 | 55-65 | 20 vs. 36 (p=NR)         | 18.7 vs. 25.3 (p=.07) | 47.6 vs. 47.8 (p=.91) |
| Child <i>et al.</i> 2003 MRC VII <sup>4</sup>          | 407 | ≤ 65  | 8 vs. 44 (p<.001)        | 19 vs. 31 (p=.001)    | 42 vs. 54 (p<.001)    |
| Palumbo <i>et al.</i> 2004 M97G <sup>5</sup>           | 194 | 50-70 | 25 vs. 6 (p=0.002)       | 16 vs. 37 (p<0.001)   | 62 vs. 77 (p<0.001)   |
| Blade <i>et al.</i> 2005 PETHEMA <sup>6</sup>          | 216 | ≤ 65  | 11 vs. 30 (p=.002)       | 33 vs. 42 (p=NS)      | 61 vs. 66 (p=NS)      |
| Barlogie <i>et al.</i> 2006 US Intergroup <sup>7</sup> | 516 | ≤ 70  | 15 vs. 17 (p=NS)         | 21 vs. 25 (p=.05)     | 53 vs. 58 (p=NS)      |

## Rationale For Use of Autologous Stem Cell Transplantation

Autologous stem cell transplantation (ASCT) represents a significant advancement in care for patients with myeloma, where the chemotherapeutic options were historically limited. Multiple randomized controlled trials (RCT) have demonstrated the superiority of ASCT over standard care/conventional cytotoxic chemotherapy – improved depth of response, PFS/EFS and OS (Table 1). A meta-analysis of these historic studies supports the use of ASCT with improvements in PFS, but not OS<sup>8</sup>. Additionally, one RCT demonstrates better HRQOL as evaluated by a composite endpoint of a longer period without symptoms, treatment, and treatment toxicity (TwisTT)<sup>9</sup>.

With the availability of newer chemotherapeutics such as proteasome inhibitors and immunomodulatory agents, they have been incorporated into myeloma care<sup>10</sup>. Such combinations have led to deeper and more durable responses either in induction, consolidation, and maintenance therapy<sup>11,12</sup>. Given these improved outcomes, many have challenged whether ASCT still has a role in myeloma care. Table 2 summarizes more recent Phase III RCTs performed in the current era of chemotherapeutics for induction. These studies have been variably subject to pooled analyses and meta-analyses demonstrating ACST's value<sup>13,14</sup>. More recently, the Determination study led by Richardson *et al.* compared 8 cycles of lenalidomide, bortezomib and dexamethasone (RVD) to 3

cycles of RVD, ASCT with 2 cycles of RVD consolidation. Subsequently, both groups received lenalidomide maintenance. With a median follow-up of 76 months, they demonstrate a median PFS that favors the ASCT group (46.2 vs. 67.5 months) affirming the continued role of ASCT.

**Table 2:** Review of RCTs Comparing Standard Therapy (SDT) to High-Dose Chemotherapy (HDT) with ASCT in the era of novel agent therapy.

| Study                                                     | N    | Age | SDT versus HDT (p-value) |                            |                        |
|-----------------------------------------------------------|------|-----|--------------------------|----------------------------|------------------------|
|                                                           |      |     | VGPR/CR (%)              | Median PFS (mths)/3 yr PFS | 3-,4- or 5 year OS (%) |
| Palumbo <i>et al.</i> 2014 <sup>15</sup>                  | 402  | <65 | 15.7 vs. 20              | 22.4 vs. 43 (p<0.001)      | 65.3 vs. 81.6 (p=0.02) |
| Gay <i>et al.</i> 2015 <sup>16</sup>                      | 389  | ≤65 | ?20s vs. 30s             | 28.6 vs. 43.3 (p<0.001)    | 84 vs. 87              |
| Cavo <i>et al.</i> 2017 <sup>17</sup>                     | 1503 | ≤65 | 75 vs. 84                | 57% vs. 64% (p=0.002)      | NR                     |
| Attal <i>et al.</i> 2017 IFM/DFCI2009 <sup>18</sup>       | 700  | ≤65 | 48 vs. 59 (p=0.03)       | 36 vs. 50 (p<0.001)        | 82 vs. 81 (p=0.87)     |
| Richardson <i>et al.</i> 2022 DETERMINATION <sup>19</sup> | 722  | ≤65 | 42 vs.46.8               | 46.2 vs.67.5               | 79.2 vs. 80.7 (p=1.0)  |

Collectively, these studies suggest that ASCT consistently improves responses, PFS and OS. ASCT continues to be a key intervention in the current era of chemotherapeutics.

### Timing of ASCT

The optimal timing of ASCT has been debated – should it be offered after successful induction therapy or on relapse? The only Phase III RCT was performed prior to the availability of novel agent, where it demonstrates similar survival, but patients undergoing early ASCT had superior HRQOL as measured by TwisTT<sup>9</sup>. In the current therapeutic era, there are several single institution observational studies compared early vs. delayed ASCT<sup>20-23</sup> and systematic reviews<sup>14</sup> suggesting a superior depth of response and PFS but similar OS with early ASCT. The IFM-DFCI RCT by Attal *et al.* compared 3 cycles of RVD induction with ASCT vs. 8 x RVD. In the 8 x RVD group, 79% of symptomatic patients received a salvage (delayed) ASCT<sup>18</sup>. This study may allow an indirect comparison of early vs. delayed ASCT. Both PFS and HRQOL was favored in the patients receiving early ASCT in the IFM-DFCI study<sup>18</sup>. More recently, patients participating in phase III EMN02/HO95 study who were randomized to upfront ASCT were compared (not randomized) to those who received ASCT at the time of progression after primary randomization to VMP<sup>24</sup>. After a median follow-up of 85 months in the upfront auto-HSCT group with a median follow-up of 51 months in the delayed auto-HSCT group, the median PFS2 rate was 55% vs 32% (HR, 0.52; 95% CI, 0.40–0.66; p < 0.0001) respectively. Moreover, median OS was not reached in the upfront auto-HSCT group vs 81 months in the delayed auto-HSCT group, with OS rates of 69% vs 58% (HR, 0.68; 95% CI, 0.51–0.93; p = 0.0164).

**Table 3:** Review of RCTs Comparing Upfront or Delayed ASCT

| Study                                               | N   | Age (yrs) | Upfront versus Delayed (p-value) |                       |                          |
|-----------------------------------------------------|-----|-----------|----------------------------------|-----------------------|--------------------------|
|                                                     |     |           | Response                         | Median EFS/PFS (mths) | Median OS (mths)         |
| Ferland <i>et al.</i> 1998 <sup>9</sup>             | 202 | ≤56       | NR                               | 39 vs. 13 p=S         | 64.6 vs. 64 p=0.92       |
| Attal <i>et al.</i> 2017 IFM/DFCI2009 <sup>18</sup> | 700 | ≤65       | 59% vs. 48% (CR)                 | 50 vs. 36 p<0.001     | 81% vs. 82% at 4yrs p=NS |

Taken together, it is ideal that ASCT be considered upfront in patients who are eligible. However, it is reasonable to delay ASCT due to personal and/or psychosocial reasons given similar OS. It must be recognized that by deferring an ASCT may mean that in the future, an ASCT may not be possible or warranted. This risk has been estimated to be around 10% from a single institution study<sup>25</sup>. Moreover, it may be preferable to strongly consider an upfront ASCT in individuals who are deemed high risk by disease presentation, cytogenetics, or gene expression profiling.

## **Eligibility for ASCT**

### **Chronologic Age:**

Given that most RCTs evaluate patients who are ≤65 years for ASCT, this constitutes the highest level of evidence for practice. However, there have been numerous observational studies that suggest that ASCT is feasible in patients > 65 years with careful selection<sup>26-30</sup>. In addition, there have been movements to consider physiologic age as opposed to chronologic age<sup>31</sup>. As such, it is reasonable to consider ASCT in patients >65 with careful attention to comorbidities and assessments of frailty. Assessments of frailty<sup>32-34</sup> are nuanced and in the myeloma setting, the IMWG Frailty score<sup>35</sup> is often used: <http://www.myelomafrailtyscorecalculator.net/>

As such, it is reasonable to consider the IMWG Frailty score as an anchor to help determined ASCT eligibility<sup>36</sup>. A more in-depth assessment of frailty may be required, and a review is available in the AHS BMT guidelines chapter: Patient Eligibility.

However, given the advancements in myeloma therapeutics, the relative value of ASCT in older patients will likely decrease. For instance, the use of Daratumumab/Lenalidomide and dexamethasone (DRd) can provide a 5-year PFS rate of 52.5%<sup>37</sup>.

### **Patient Variables:**

See AHS BMT guidelines section: Patient Eligibility

### **Depth of Response:**

In general depth of response pre-ASCT correlates with post-ASCT outcomes<sup>38</sup>. However, a historic registry study at IBMTR suggests deepening responses beyond partial response (PR) did not translate to better OS. More recent data from the CIBMTR suggest that patients achieving less than a PR to initial induction therapy, including with novel agent combinations, additional pre-ASCT salvage chemotherapy improved the depth of response and pre-ASCT disease status but was not associated with survival benefit<sup>39</sup>. Additionally, patients with refractory myeloma (<PR) might still derived benefit from ASCT. Notably, current induction therapies (e.g., CYBORD or RVD) do not generally include cytotoxic therapy – meaning that a less than responsive disease might still derived benefit from ASCT where high dose melphalan is used for conditioning<sup>40</sup>.

In the “novel therapy” era, the MRC XI<sup>41</sup> evaluated whether there is value in deepening responses prior to ASCT. They randomized patients who only achieved a MR or PR after immunomodulatory

based triplet therapy to receive either no additional therapy or additional therapy with bortezomib, cyclophosphamide and dexamethasone (CYBORD). The additional therapy improved the pre-ASCT responses (PR improved to VGPR in 41% of evaluable cases) and translated to improved PFS (55 months vs. 30 months,  $p=0.0003$ ), but no differences in OS.

## Stem Cell Mobilization

See AHS BMT guidelines section: Donor Management, Mobilization

## Should We Be Collecting Enough Stem Cells for 2 ASCTs:

Table 4 illustrates ASCTs performed in Calgary from 2010, including the proportion of 2<sup>nd</sup> and 3<sup>rd</sup> ASCT (planned tandem, delayed ASCT or stem cell boost).

Considerations for collection amount include: 1) patient factors such as age, co-morbidities, 2) disease factors such as high-risk disease, 3) evolving myeloma therapies, and 3) local resource factors. Taken together, collecting for 1 or 2 ASCT will be discussed on a case-by-case basis.

**Table 4:** ASCTs performed in Calgary from 2010

|                        | 2010 | 2011 | 2012 | 2013 | 2014 | 2015 | 2016 | 2017 | 2018 | 2019 | 2020 | 2021 |
|------------------------|------|------|------|------|------|------|------|------|------|------|------|------|
| 1 <sup>st</sup> ASCT   | 28   | 29   | 24   | 27   | 40   | 41   | 40   | 25   | 43   | 32   | 38   | 38   |
| ≤65 years <sup>1</sup> | 28   | 28   | 24   | 25   | 38   | 32   | 33   | 23   | 40   | 31   | 32   | 31   |
| >65 years <sup>1</sup> | 0    | 1    | 0    | 2    | 2    | 9    | 7    | 2    | 3    | 1    | 6    | 7    |
| 2 <sup>nd</sup> ASCT   | 3    | 0    | 1    | 1    | 5    | 6    | 6    | 5    | 7    | 3    | 2    | 5    |
| ≤65 years <sup>2</sup> | 3    | 0    | 1    | 1    | 4    | 5    | 5    | 3    | 4    | 2    | 1    | 4    |
| >65 years <sup>2</sup> | 0    | 0    | 0    | 0    | 1    | 1    | 1    | 2    | 3    | 1    | 1    | 1    |
| 3 <sup>rd</sup> ASCT   | 0    | 0    | 0    | 0    | 2    | 0    | 0    | 1    | 0    | 0    | 1    | 0    |
| ≤65 years <sup>3</sup> | 0    | 0    | 0    | 0    | 2    | 0    | 0    | 1    | 0    | 0    | 0    | 0    |
| >65 years <sup>3</sup> | 0    | 0    | 0    | 0    | 0    | 0    | 0    | 0    | 0    | 0    | 1    | 0    |
| Total ASCT             | 31   | 29   | 25   | 28   | 47   | 47   | 46   | 31   | 50   | 35   | 41   | 43   |

2<sup>nd</sup>/3<sup>rd</sup> ASCT may include tandem ASCT or stem cell boosts

## Conditioning Regimen

The current standard for conditioning is Melphalan 200mg/m<sup>2</sup> following a RCT demonstrating that Melphalan 200mg/m<sup>2</sup> (Mel200) was superior to Melphalan 140mg/m<sup>2</sup> (Mel140) with 8 Gy Total Body Irradiation<sup>42</sup>. A large database study from EBMT suggests that Mel200 may be preferable to Mel140 to patients who do not achieve a ≥PR post-induction chemotherapy. In contrast, patients appear to do just as well with Mel140 in those with ≥VGPR post-induction chemotherapy<sup>43</sup>.

More recently, Roussel *et al.* compared Mel200 with Mel200+Bortezomib. In this RCT carried out by the IFM, the combination of Mel200+Bortezomib did not result in superior depth of response, PFS or OS<sup>44</sup>. Additionally, Bashir *et al.* evaluated the combination of Mel140+Busulfan 130mg/m<sup>2</sup> vs. Mel200 in a RCT with the combination therapy associated with an improved PFS (65 months vs. 44 months)<sup>45</sup>. Interestingly, this trial showed an improvement in PFS with the Bu-Mel conditioning regimen in patients with newly diagnosed myeloma with HR cytogenetics, where the median PFS was 44.7 months and 25.7 months in the Bu-Mel and Mel arms, respectively ( $P = .044$ )<sup>46</sup>. A meta-analysis

of 10 observation studies with a total of 2855 patients examined the use of Bu-Mel conditioning compared with Mel200 alone. Bu-Mel conditioning correlates with longer PFS (HR 0.77; p=0.0002) but similar OS (HR 1.08; p=0.35)<sup>47</sup>.

The dose of melphalan should be reduced to 140mg/m<sup>2</sup> in individuals with renal dysfunction or on dialysis<sup>48-50</sup>. Further, consideration could be given to a 140mg/m<sup>2</sup> melphalan dosing in those who >65 years with deemed frailty<sup>50-52</sup>.

**Table 5:** Review of RCTs Comparing Conditioning Regimens in ASCT

| Study                                                | N   | Age (yrs)<br>Median | Mel200 versus "other" (p-value) |                                          |                                  |
|------------------------------------------------------|-----|---------------------|---------------------------------|------------------------------------------|----------------------------------|
|                                                      |     |                     | CR (%)                          | Median EFS/PFS                           | OS (%)                           |
| Moreau <i>et al.</i> 2002 IFM 9502 <sup>42</sup>     | 282 | <65                 | NA                              | EFS of 20.5 mths vs. 21 mths (p=0.6)     | 65.8 vs. 45.5 (at 45mths) p=0.05 |
| Roussel <i>et al.</i> 2022 IFM 2014-02 <sup>44</sup> | 300 | 58                  | 22.1 vs. 20.5                   | PFS of 34 mnths vs. 29.6 mnths (p=0.244) | 89.5 vs. 89.5 (at 3 years)       |
| Bashir <i>et al.</i> 2019 <sup>53</sup>              | 202 | 58-59               | 26 vs. 33                       | PFS of 43.5 mths vs. 64.7 mths (p=0.022) | NS at (p=0.75)                   |

## Role of Tandem ASCT

### Tandem ASCT with Historic Induction Chemotherapy:

A tandem ASCT can be defined as a pre-planned second ASCT within 6 months of the first ASCT, where the goal is avail of additional high dose melphalan to achieve a deeper hematological response.

There has been several RCTs comparing tandem with single ASCTs prior modern induction chemotherapy<sup>52,54-56</sup>. These older studies demonstrate that tandem RCTs can improve either EFS or PFS with only one demonstrating OS advantage limited to the subgroup of patients not achieving VGPR after the 1<sup>st</sup> ASCT<sup>54</sup>. Two meta-analyses would confirm these observations<sup>57,58</sup>.

**Table 6:** Review of historic RCTs Comparing Tandem ASCT with Single ASCT

| Study                                            | N   | Age (yrs) | Tandem versus single (p-value) |                            |                                 |
|--------------------------------------------------|-----|-----------|--------------------------------|----------------------------|---------------------------------|
|                                                  |     |           | Responses (%)                  | Median EFS/PFS (mths)      | Median OS (mths)                |
| Attal <i>et al.</i> 2003 <sup>54</sup>           | 399 | <60       | 50 vs. 42 (>VGPR)              | 20% vs. 10% at 7yrs p=0.03 | 42% vs. 21% (at 7 years) p=0.01 |
| Cavo <i>et al.</i> 2007 Bologna 96 <sup>55</sup> | 321 | ≤60       | 47 vs. 33 (≥nCR)               | 25 vs. 32 p=0.19           | NR                              |
| Sonnevald <i>et al.</i> 2007 <sup>52</sup>       | 303 | ≤65       | 32 vs. 13 (CR)                 | 17% vs. 9% at 6yrs p=0.006 | 39% vs. 36% at 6yrs p=0.51      |
| Mai <i>et al.</i> 2016 GMMG-HD2 <sup>56</sup>    | 358 | ≤66       | 19.4 vs. 16 (CR)               | 33.1 vs. 26.2 p=NS         | 75.3 vs. 73.0 p=NS              |

### Tandem ASCT with Modern Induction Chemotherapy:

More recently, a post-hoc analyses using pooled data from 3 independent Phase III RCTs was performed to elucidate the potential value of tandem ASCT<sup>59</sup>. All 3 RCTs utilized bortezomib-based



induction regimen. There was a significant improvement in the median PFS and 5-year OS in favor of tandem ASCT. This benefit was more apparent in patients with high-risk cytogenetics such as t(4;14) and/or deletion 17p who had not achieved a CR after induction therapy (70% vs. 17%). Additionally, these results were further affirmed with their 10 year follow-up data where patients with ISS stage II+III, high-risk cytogenetics and failure to achieve CR benefitted from a tandem ASCT approach<sup>60</sup>.

These findings were also echoed by the preliminary findings in the EMN02/HO95 MM study<sup>17</sup>, where there was a significant improvement in the 3-year PFS (73% vs. 64%) and 3-year OS (89% vs. 82%) in favor of tandem ASCT compared to single ASCT. The superiority of a tandem ASCT was again evident in patients with high-risk cytogenetics in both 3-year PFS and OS. In this study, most patients received bortezomib based induction (i.e., no lenalidomide).

In contrast, the BMT-CTN 0702 STAMINA study<sup>61</sup> did not demonstrate any difference between patients (after initial ASCT who either received a 1) tandem ASCT followed by maintenance lenalidomide, 2) 4 cycles of consolidation (lenalidomide, bortezomib and dexamethasone (RVD)) followed by maintenance lenalidomide, or 3) maintenance lenalidomide alone. Of note, most patients in this study received RVD induction therapy prior to initial ASCT.

**Table 7:** Review of recent RCTs Comparing Tandem ASCT with Single ASCT

| Study                                               | N    | Age    | Tandem versus single/other    |               |               |
|-----------------------------------------------------|------|--------|-------------------------------|---------------|---------------|
|                                                     |      |        | Randomized Gps                | Median PFS    | Median OS     |
| Stadtmauer <i>et al.</i> 2016 STAMINA <sup>61</sup> | 758  | <71yrs | Mel200 x1 - R maint           | 52 mths       | 83 mths       |
|                                                     |      |        | Mel200 x2 - R maint           | 57 mths       | 86 mths       |
|                                                     |      |        | Mel200 x1- RVD conso- R maint | 57 mths       | 82 mths       |
| Cavo <i>et al.</i> 2017 EMN02/HO95 <sup>17</sup>    | 1503 | ≤65yrs | Mel200 x1 - +/- RVD - R maint | 64% (3yr PFS) | 82% (3yr PFS) |
|                                                     |      |        | Mel200 x2 - +/- RVD - R maint | 73% (3yr PFS) | 89% (3yr PFS) |

### Consolidation Post-ASCT

The goal of consolidation therapy post-ASCT is to improve and augment responses – to suppress residual disease. There have several RCTs suggesting that consolidation can deepen responses and prolong PFS. However, its effect on OS is less clear.

Indeed, the above mentioned BMT-CTN 0702 STAMINA study<sup>61</sup> did not demonstrate benefit of consolidation therapy. In contrast, preliminary data from the EMN02/HO95 MM study<sup>62</sup> suggest a PFS benefit with 2 cycles of lenalidomide, bortezomib and dexamethasone (VRD) consolidation (p=0.013) without OS benefit (p>0.05).

Both the CASSIOPEIA and PERSEUS seek to understand the value of Daratumumab in patients eligible for ASCT. The PERSEUS randomly assigned 709 transplantation-eligible patients with newly diagnosed multiple myeloma to receive either subcutaneous daratumumab combined with VRd induction and consolidation therapy and with lenalidomide maintenance therapy (D-VRd group) or VRd induction and consolidation therapy and lenalidomide maintenance therapy alone (VRd group). Similarly, the CASSIOPEIA evaluated daratumumab in combination with bortezomib, thalidomide,

and dexamethasone (D-VTd) as induction therapy before and consolidation therapy after autologous stem-cell transplantation (ASCT), followed by a second randomization to daratumumab maintenance therapy (up to 2 years) or observation alone. This means that the independent value of consolidation post-ASCT in these 2 studies cannot be assessed – see Section Maintenance with anti-CD38 monoclonal antibody for more details.

**Table 8:** Review of recent RCTs Comparing Consolidation Therapy with No Consolidation Therapy

| Study                                                    | N   | Age    | Consolidation versus none     |                   |                  |
|----------------------------------------------------------|-----|--------|-------------------------------|-------------------|------------------|
|                                                          |     |        | Randomized Gps                | Median PFS (mths) | Median OS (mths) |
| Stadtmauer <i>et al.</i> 2019<br>STAMINA <sup>63</sup>   | 758 | <71yrs | Mel200 x1 - R maint           | 52                | 83               |
|                                                          |     |        | Mel200 x2 - R maint           | 57                | 86               |
|                                                          |     |        | Mel200 x1- RVD conso- R maint | 57                | 82               |
| Sonneveld <i>et al.</i> 2020<br>EMN02/HO95 <sup>62</sup> | 903 | ≤65yrs | RVD consolidation x 2         | 59                | 75% at 6yrs      |
|                                                          |     |        | No consolidation              | 43                | 69% at 6yrs      |

### Maintenance Post-ASCT

A typical maintenance therapy is low dose with limited toxicity administered over a prolonged period to deepen responses and/or maintain responses. Thalidomide was historically for maintenance therapy post-ASCT with improvements in depth of responses, PFS and possibly OS. This came at a cost of clinically significant peripheral neuropathy and therapy related fatigue. Interestingly, thalidomide maintenance therapy has been associated with worse outcomes in patients with high-risk cytogenetics.

### Maintenance with Lenalidomide Monotherapy

Maintenance with lenalidomide has been considered standard of care following the publications by CALGB<sup>64</sup> and IFM<sup>65</sup>. Both studies demonstrate improvements in PFS but only the CALGB study demonstrates improvements with OS on lenalidomide maintenance. Both the GIMEMA<sup>15</sup> and Myeloma IX<sup>41</sup> study would confirm the benefit of maintenance lenalidomide on PFS. The results of these studies have been subject to systematic reviews/meta-analyses which confirms the efficacy of maintenance lenalidomide<sup>66-70</sup>.

The duration of maintenance therapy with lenalidomide has been subject to some discussion. In the IFM/DFCI2009 study<sup>18</sup> by Attal *et al.*, maintenance therapy with lenalidomide post-ASCT was for 1 year. In contrast, the Determination study<sup>19</sup> by Richardson *et al.* employed maintenance therapy with lenalidomide until progression. An indirect comparison suggests a median PFS of 35 months and 46.2 months respectively. This would support a preference for lenalidomide maintenance until progression.

**Table 9:** Review of recent RCTs Comparing Maintenance Therapy with Lenalidomide with No Maintenance Therapy

| Study                                           | N    | Age | Lenalidomide Maintenance versus none (p-value) |                         |                                   |
|-------------------------------------------------|------|-----|------------------------------------------------|-------------------------|-----------------------------------|
|                                                 |      |     | >VGPR                                          | Median PFS (mths)       | Median OS (mths)                  |
| Attal <i>et al.</i> 2012 IFM <sup>65</sup>      | 614  | <65 | 84% vs. 76% (p=0.009)                          | 41 vs. 23 (p<0.001)     | 73% vs. 75% at 4yrs (p=NS)        |
| McCarthy 2012 CALGB <sup>64</sup>               | 460  | <71 | NR                                             | 46 vs. 27 (P<0.001)     | 88% vs. 80% at 3yrs (p=0.03)      |
| Palumbo <i>et al.</i> 2014 GIMEMA <sup>15</sup> | 273  | <65 | NR                                             | 41.9 vs. 21.6 (p<0.001) | 88% vs. 79.2% at 3yrs (p=NS)      |
| Jackson <i>et al.</i> Myeloma IX <sup>41</sup>  | 2568 | NR  | NR                                             | 57 vs. 30 (p<0.0001)    | 87.5% vs. 80.2% at 3yrs (p=0.014) |

### Maintenance with Proteasome Inhibition Monotherapy:

The effect of maintenance bortezomib post-ASCT has also been evaluated. The HOVON-65/GMMG-HD4 study suggest that patients with del 17p might benefit from a proteasome inhibitor maintenance<sup>71,72</sup>. More recently, the TOURMALINE MM3 study<sup>73</sup> evaluated ixazomib maintenance post-ASCT compared to placebo. After a median follow-up of 31 months with 54% of PFS events, there was a 28% reduction in the risk of progression/death, corresponding to a 39% improvement in PFS with ixazomib vs placebo (median 26.5 vs 21.3 months; hazard ratio [HR] 0.72; 95% CI: 0.582, 0.890; p=0.002). In a landmark analysis from ASCT, PFS was 30.7 vs 24.9 months (HR 0.684; 95% CI: 0.551, 0.848; p<0.001).

**Table 10:** Review of recent RCTs Comparing Maintenance Therapy with Proteasome Inhibition (PI) with No Maintenance Therapy

| Study                                                             | N   | Age (yrs) | Proteasome Inhibition versus other (p-value) |                         |                     |
|-------------------------------------------------------------------|-----|-----------|----------------------------------------------|-------------------------|---------------------|
|                                                                   |     |           | Response (%)                                 | Median PFS (mths)       | OS                  |
| Goldschmidt <i>et al.</i> 2018 HOVON-65/GMMG-HD4 <sup>71,72</sup> | 827 | 57        | 36 vs. 24 CR                                 | 35 vs. 28 (p=0.002)     | 61% vs. 55% at 5yrs |
| Dimopoulos <i>et al.</i> 2019 TOURMALINE MM3 <sup>73</sup>        | 656 | 57        | 12 vs. 7 MRD -ve                             | 26.5 vs. 21.3 (p=0.002) | NR                  |

### Dual Maintenance with Proteasome Inhibition and Lenalidomide:

Maintenance therapy with ixazomib, lenalidomide and dexamethasone (IRd) was compared to lenalidomide with dexamethasone in the GEM2014MAIN trial led by Rosinol *et al*<sup>74</sup>. With a median follow-up of 56 months, they demonstrate there was no difference in PFS between the two maintenance arms (median not reached, PFS at 5 years: 62% vs. 63% with IRd and Rd, respectively, p=0.785).

Similarly, concurrent carfilzomib with lenalidomide was also compared with lenalidomide as part of a second randomization in the FORTE study<sup>75</sup>. Here, the 3-year PFS from the second randomization was 75% in patients treated with carfilzomib + lenalidomide (95% CI, 68–82, median, not reached [NR]; 95% CI, NR–NR) versus 65% with lenalidomide alone (95% CI, 58–72, median, NR; 95% CI, NR–NR) (hazard ratio, 0.64; 95% CI ,0.44–0.94; p = 0.023).

**Table 11: Review of recent RCTs Comparing Dual Maintenance Therapy with PI and Lenalidomide with Lenalidomide alone Therapy**

| Study                                                | N   | Age (yrs.) | Proteasome Inhibition and Lenalidomide versus other (p-value) |                                  |                        |
|------------------------------------------------------|-----|------------|---------------------------------------------------------------|----------------------------------|------------------------|
|                                                      |     |            | Response (%)                                                  | Median PFS (mths)                | OS                     |
| Gay <i>et al.</i> 2021 FORTE <sup>75</sup>           | 356 | 56-57      | 68 vs. 65 sCR                                                 | 75% vs. 65% at 3 years (p=0.023) | 94% vs. 90% at 3 years |
| Rosinol <i>et al.</i> 2021 GEM2014MAIN <sup>74</sup> | 332 | 58-59      | 73.7 vs. 65.2 sCR/CR                                          | 62% vs. 63% at 5 years           | Not available          |

### Maintenance with Anti-CD38 Monoclonal Antibody:

Again, CASSIOPEIA was a two-part, randomized phase 3 trial that evaluated daratumumab in combination with bortezomib, thalidomide, and dexamethasone (D-VTd) as induction therapy before and consolidation therapy after autologous stem-cell transplantation (ASCT), followed by a second randomization to daratumumab maintenance therapy (up to 2 years) or observation alone, in transplant-eligible patients with newly diagnosed multiple myeloma. With a follow-up of 80.1 months, progression-free survival from second randomization was significantly longer in the daratumumab maintenance group than the observation-alone group (see Table 12).

Likewise, the PERSEUS randomly assigned 709 transplantation-eligible patients with newly diagnosed multiple myeloma to receive either subcutaneous daratumumab combined with VRd induction and consolidation therapy and with lenalidomide maintenance therapy (D-VRd group) or VRd induction and consolidation therapy and lenalidomide maintenance therapy alone (VRd group). Daratumumab was stopped at 2 years if complete response or better and had sustained minimal residual disease (MRD)–negative status (the absence of malignant cells at a sensitivity threshold of  $10^{-5}$  or lower) for at least 12 months. PERSEUS trial did NOT have a second randomization to maintenance therapy – meaning the independent value of maintenance daratumumab is less clear.

It is ideal to have Minimal Residual Disease (MRD) testing to guide potential discontinuation of therapy after 2 years, a management strategy validated by the PERSEUS study, where up to 65% patients may achieve this outcome.

**Table 12: Review of recent RCTs Comparing anti-CD38 antibody with No maintenance/ Lenalidomide alone Therapy**

| Study                                | N   | Age (yrs.) | Anti-CD38 versus other (p-value) |                                   |                       |
|--------------------------------------|-----|------------|----------------------------------|-----------------------------------|-----------------------|
|                                      |     |            | Response (%)                     | Median PFS (mths)                 | OS                    |
| Sonneveld <i>et al.</i> 2023 PERSEUS | 709 | 60         | 65.1 vs. 32.2 (MRD $10^{-6}$ )   | 84.3% vs 67.7% at 4 years         |                       |
| Moreau <i>et al.</i> 2024 CASSIOPEIA | 886 | 58         | 58.1 vs. 48.9 (MRD $10^{-6}$ )*  | Not reached vs. 72.1 (p=0.0481) * | 85% vs.84% at 80 mths |
|                                      |     |            | 43.7 vs 26.5 (MRD $10^{-6}$ )^A  | Not reached vs. 32.7 (p<0.0001)^A |                       |

\*received D-VTD induction; ^received VTD induction

## Role of Second ASCT for Salvage Therapy

Given that most if not all patients will relapse, a second ASCT can be considered as salvage treatment. There has been 1 RCT that evaluates the use of 2<sup>nd</sup> salvage ASCT vs. conventional care, demonstrating that a 2<sup>nd</sup> salvage improves PFS but not OS<sup>76</sup>. In contrast, there have been numerous observational studies that support the use of a 2<sup>nd</sup> salvage ASCT<sup>77-83</sup>. Given that the duration of response with a 2<sup>nd</sup> ASCT will be shorter than the 1<sup>st</sup> ASCT, an arbitrary cutoff of at least 2 years from the 1<sup>st</sup> ASCT before 2<sup>nd</sup> ASCT should be considered. However, the routine use of consolidation and maintenance may change this duration of response “cutoff”.

**Table 13:** Review of RCTs of 2<sup>nd</sup> salvage ASCT.

| Study                                                        | N   | Age | Salvage 2 <sup>nd</sup> ASCT vs. conventional (p-value) |                     |                        |
|--------------------------------------------------------------|-----|-----|---------------------------------------------------------|---------------------|------------------------|
|                                                              |     |     | Overall Response (%)                                    | Median PFS (mths)   | 3-Year OS (%)          |
| Cook <i>et al.</i> 2014 NCRI myeloma X Relapse <sup>76</sup> | 297 | 61  | 83 vs. 75                                               | 19 vs. 11 (p<0.001) | 80.3 vs. 62.9 (p=0.19) |

## Role of Allogeneic Transplant

Evidence for a graft-versus-myeloma effect has been weak. Allogeneic transplants (myeloablative) are associated with significant treatment related toxicity with unclear long-term benefits. Given these toxicities, reduced intensity conditioning (RIC) allogeneic transplants have been advocated to mitigate concerns surrounding transplant related mortality. There have been several RCTs and quasi-RCTs that have evaluated tandem ASCT vs. ASCT followed by RIC allogeneic transplants<sup>84-89</sup>. In general, there is a lack of meaningful benefit with an ASCT-RIC allogeneic transplant approach to myeloma care<sup>90,91</sup>. Allogeneic transplant in relapsed disease is poorly tolerated with marginal effectiveness over other available therapies<sup>90,91</sup>.

**Table 14:** Review of RCT and quasi-RCTs comparing studies Tandem ASCT vs. ASCT-RIC Allo

| Study                                                         | N   | Mean/Median Age (yrs) | Tandem ASCT vs. ASCT-RIC Allo (p-value) |                                 |                                |
|---------------------------------------------------------------|-----|-----------------------|-----------------------------------------|---------------------------------|--------------------------------|
|                                                               |     |                       | CR (%)                                  | Median PFS/EFS                  | Median OS                      |
| Moreau <i>et al.</i> 2008 IFM99-03 and IFM99-04 <sup>87</sup> | 284 | 58 vs. 54 (p=0.006)   | 38 vs. 62                               | 22mths vs. 19mths EFS (p=0.58)  | 48mths vs. 34mths (p=0.07)     |
| Rosinol <i>et al.</i> 2008 PETHEMA <sup>88</sup>              | 752 | 55 vs. 52             | 11 vs. 40 (p=0.001)                     | 31mths vs. NR PFS (p=0.08)      | 34mths vs. 58mths (p=0.9)      |
| Krishnan <i>et al.</i> 2011 BMT-CTN0102 <sup>85</sup>         | 710 | <70                   | 45 vs. 58                               | 46% vs. 43% at 3 yrs (p=0.671)  | 80% vs. 77% at 3 yrs (p=0.191) |
| Giaconne <i>et al.</i> 2011 <sup>86</sup>                     | 162 | 55                    | 26 vs. 55                               | 2.4yrs vs. 2.8yrs EFS (p=0.005) | 4.25yrs vs. NR (p=0.001)       |
| Gahrton <i>et al.</i> 2013 EBMT-NMAM2000 <sup>84</sup>        | 357 | ≤69                   | 41 vs. 50                               | 12% vs. 22% at 8 yrs (p=0.027)  | 36% vs. 49% at 8 yrs (p=0.154) |
| Knop <i>et al.</i> 2014 <sup>89</sup>                         | 199 | 53                    | 31 vs. 59                               | 23mths vs. 35mths PFS (p=0.005) | 72mths vs. 70mths (p=NS)       |

# Systemic AL Amyloidosis

## Background

Systemic immunoglobulin light chain amyloidosis is a protein misfolding disease caused by the conversion of immunoglobulin light chains from their soluble functional states into highly organized amyloid fibrillar aggregates that lead to organ dysfunction<sup>92</sup>. Light-chain (AL) amyloidosis is the most common form of systemic amyloidosis, accounting for 70% of patients with amyloidosis<sup>93</sup>. AL amyloidosis (historically referred to as primary amyloidosis) is an uncommon disorder and its exact incidence is unknown. However, in the USA the incidence ranges from 9–14 cases per million person years<sup>94</sup>. AL amyloidosis is a disease of the elderly with a median age at diagnosis of 63 years<sup>95</sup>. There is a male predominance, with men accounting for 55% of cases<sup>96</sup>. AL amyloidosis occurs in all races and geographic locations, but data are limited regarding the incidence of AL amyloidosis across different ethnic groups.

Patients with a new diagnosis of AL amyloidosis should be referred to a center with expertise in the treatment of this entity, especially if considering for autologous stem cell transplantation (ASCT).

## Autologous Stem Cell Transplantation (ASCT):

Most AL amyloidosis patients are not eligible for ASCT due to the presence of significant comorbidities. Early studies reported high mortality rates during ASCT for AL patients, however, due to improved supportive care and careful patient selection, the mortality in ASCT has decreased significantly<sup>97</sup>.

A recent report by the Mayo Clinic showed an early mortality rate (before day 100) of only 1.1% when Mayo stage III patients are excluded from transplant<sup>98</sup>. In addition, a recent long-term report on 20 years of experience with ASCT for AL amyloidosis at the Mayo Clinic Rochester highlighted the benefits of supportive care and patient selection in the setting of ASCT for AL Amyloidosis<sup>99</sup>. In brief, 672 consecutive patients receiving ASCT for AL amyloidosis were divided into three cohorts based on date of transplantation (cohort 1, 1996-2002 [n = 124]; cohort 2, 2003-2009 [n = 302]; and cohort 3, 2010-2016 [n = 246]). The median age for the entire cohort was 59 years, with patients in cohort 3 being slightly older than those in the other two cohorts (60 vs 58 vs 54 years for cohorts 3, 2, and 1, respectively;  $p < 0.001$ ). More patients received pre-transplantation therapy in cohort 3 compared with earlier time periods (49% vs 38% vs 42% for cohorts 3, 2, and 1, respectively;  $p = 0.02$ ). Hematologic response was higher in cohort 3 (84% vs 79% vs 69% for cohorts 3, 2, and 1, respectively;  $p = 0.002$ ). Median overall survival for the entire cohort was 122 months and improved over time (not reached vs 120 months vs 75 months for cohorts 3, 2, and 1, respectively;  $p < 0.001$ ). Treatment-related mortality declined over time (2.4% vs 8.6% vs 14.5% for cohorts 3, 2, and 1, respectively;  $p < 0.001$ ).

The improved survival and markedly reduced treatment-related mortality in eligible patients indicate that this will remain an important first-line option even in the era of treatment approaches.



## Eligibility Criteria

ASCT was reported in 1996 as a form of treatment for AL amyloidosis<sup>100</sup>. High-dose dexamethasone was introduced later in 1997<sup>101</sup>. Since then, multiple advances in the treatment (novel agents) and supportive care have been developed. The first randomized clinical trial on AL amyloidosis led by the MAG and IFM group reported that the outcome of treatment of AL amyloidosis with high-dose melphalan plus ASCT was not superior to the outcome with standard-dose melphalan plus dexamethasone<sup>102</sup>. However, no cardiac biomarker selection was made on those patients and 29 centers were included for the study.

Requirements for safe ASCT currently include<sup>97,98,103-105</sup>:

1. Age <65 years,
2. Performance status (Eastern Cooperative Oncology Group) 0 to 2,
3. NT-proBNP is <5000 ng/l and/or cardiac troponin T is <0.06 ng/ml,
4. Estimated glomerular filtration rate >30 ml/min per 1.73 m<sup>2</sup> unless on dialysis,
5. New York Heart Association class <III, cardiac ejection fraction >45%, systolic blood pressure >90 mm Hg (standing), and
6. Lung CO diffusion capacity >50%.

Non-transplant candidates can be offered melphalan-dexamethasone or cyclophosphamide-bortezomib-dexamethasone. Daratumumab appears to be highly active in AL amyloidosis. Currently, a clinical trial incorporating daratumumab to CyBorD is ongoing. Antibodies designed to dissolve existing amyloid deposits are under study. So far, only one amyloid removal antibody trial is ongoing (CAEL 101).

## Induction and Stem Cell Collection

One of the first issues to consider for AL patients eligible for ASCT is the stem cell collection process<sup>106</sup>. Patients with AL commonly suffer from kidney and heart involvement and during the collection process often tend to accumulate fluids during filgrastim (granulocyte colony stimulating factor) mobilization<sup>107</sup> and thus, fluid balance should be meticulously followed and maintained.

The second issue in transplantation of AL amyloidosis patients is whether an induction before SCT improves outcomes. A single-center, prospective randomized trial reported on the role of induction (two cycles of bortezomib and dexamethasone) versus no induction in 56 AL amyloidosis patients. Overall hematologic (ORR) and organ response rates (OR) in the whole cohort after ASCT were 77% and 58%. The ORR and OR in the bortezomib pretreated group were 92% and 75% vs. 69% and 54% in the group that received no pretreatment. The median time to maximum hematologic response after ASCT was reduced in the group that received bortezomib induction (3 vs. 14 months). Overall cardiac response rate was 60%; 100% in patients pretreated with bortezomib and 43% in those without induction treatment. With a median follow-up of 2.9 years, the 3-year progression-free and overall survival rates in the entire cohort were 66% and 73% and in those with cardiac involvement, 73% and 80%<sup>108</sup>.

In a study from the MDACC the type of induction therapy and its impact on the outcome of autologous hematopoietic stem cell transplantation in AL was evaluated in 128 patients. The patients were

divided into 3 groups: no induction (20 patients), conventional chemotherapy-based induction (melphalan and steroids; 25 patients), and IMiD/proteasome inhibitor (PI)-based induction (83 patients). Overall, the hematological response on day 100 was highest in the IMiD/PI group, and organ response at 1 year was highest in the conventional chemotherapy-based induction. The 2-year PFS rates were 67, 56, and 73% in the no induction, CC, and IMiD/PI groups, respectively, and OS rates at 2 years were 73, 76, and 87%, respectively<sup>109</sup>.

Among 415 AL patients, 35% had induction prior to ASCT at the Mayo Clinic<sup>110</sup>. Post-ASCT hematologic CR plus VGPR rates were significantly higher in those with baseline BMPC  $\leq 10\%$  compared to BMPC  $>10\%$  (58% vs 40%,  $p = 0.0013$ ). Significant risk factors for lack of attainment of CR included attenuated dose melphalan conditioning, baseline BMPC  $> 10\%$ , no induction, and male gender. The 5-year OS for the entire group was 65%. Mayo Clinic patients eligible for ASCT that have bone marrow plasma cells lower than 10% are sent directly to ASCT.

We recommend induction therapy for those with  $>10\%$  BMPC's as outcomes appear to be better.

### **Conditioning**

In immunoglobulin light-chain (AL) amyloidosis, the depth of hematologic response to treatment is associated with improved survival and organ responses. A recent clinical trial using bortezomib in induction and in conditioning with melphalan before ASCT for AL amyloidosis was reported by the Boston University (BU) group<sup>111</sup>. The long-term results of this clinical trial with a median follow-up of 77 months in 35 patients enrolled showed a hematologic complete response and very good partial response (VGPR) of 100% (27 of 27) of the evaluable patients at 6 months post-ASCT. Four patients (15%) had hematologic relapse at a median of 42 months, and 1 patient (3.7%) had organ progression despite maintaining a VGPR at 37 months. The median overall survival and progression-free survival have not yet been reached at the time of the report. Renal and cardiac responses occurred in 65% and 88%, respectively, at 5 years post-ASCT. The median time to renal and cardiac response was 12 months and 6 months, respectively.

In conclusion, incorporating bortezomib into induction and conditioning yielded durable hematologic responses of AL amyloidosis, with corresponding organ responses and prolonged survival. At our center bortezomib and melphalan as well as melphalan are the recommended conditioning regimens for transplantation in AL amyloidosis. It should be noted that recent data from the IFM in MM showed that Bor-HDM did not improve clinical outcomes or degree of response<sup>112</sup> and thus this conditioning regimen has been discouraged in MM, no data is available in this regard for AL amyloidosis.

### **Consolidation**

It has been reported that bortezomib in combination with dexamethasone (BD) followed by ASCT can significantly improve both the hematological and organ response rates of AL amyloidosis patients compared to ASCT alone.

An initial phase II trial using bortezomib and dexamethasone (BD) as consolidation was reported by Landau *et al.*<sup>112</sup>. Forty untreated patients with renal (70%), cardiac (65%), liver/gastrointestinal (15%) or nervous system (13%) AL were assigned MEL 100, 140 or 200 mg/m<sup>2</sup> based on age, renal function, and cardiac involvement. Hematological response was assessed at 3 months post stem cell transplant (SCT); patients with less than complete hematological response (CR) received BD consolidation. Four patients with advanced cardiac AL died within 100 days of SCT (10% treatment-related mortality). Survival at 12- and 24-months post treatment start was 88 and 82% overall and was 81 and 72% in patients with cardiac AL. At 3 months post SCT, 45% had  $\geq$  partial response (PR) including 27% CR. Twenty-three patients received consolidation and in 86% response improved; all patients responded in one cycle. At 12 and 24 months, 79 and 60% had  $\geq$  PR, 58 and 40% CR. Organ responses occurred in 55 and 70% at 12 and 24 months. Eight patients relapsed/progressed. One patient with serologic progression had organ impairment at time of progression.

Based on this study, a small non-randomized trial has been recently conducted to evaluate the efficacy and safety of bortezomib in combination with ASCT in the induction, conditioning, and consolidation of patients with newly diagnosed AL amyloidosis<sup>113</sup>. The overall response (OR) rate after induction therapy of two cycles of BD was 57.1% and the CR rate was 28.5%. Eight (8/20, 40%) patients achieved hematologic CR after ASCT and 10 (10/20, 50%) after consolidation therapy. According to intention-to-treat (ITT) analysis, the CR rate was 47.6% (10/21) at 12 months after ASCT. The hematologic very good partial response (VGPR) rate reached 40% (8/20) after ASCT and 30% (6/20) after consolidation therapy. The OR rate was 90% (18/20) at 12 months after ASCT in evaluable patients and 85.7% (18/21) according to ITT analysis. The hematologic response rate of CR and VGPR increased from 52.4% to 80% after ASCT treatment. The OR rate was similar after ASCT, and the CR rate increased from 40% to 50% after consolidation therapy.

Based the paucity of data, consolidation is not recommended as the standard treatment for patients with AL amyloidosis undergoing ASCT at our center.

### Deferred ASCT

Modern chemotherapy agents can induce hematologic and organ responses in AL, including those with cardiac involvement, but durability of response remains uncertain<sup>114,115</sup>. No study has demonstrated the prolonged progression-free survival (PFS) patients treated with non-transplant regimes akin to that achieved with ASCT.

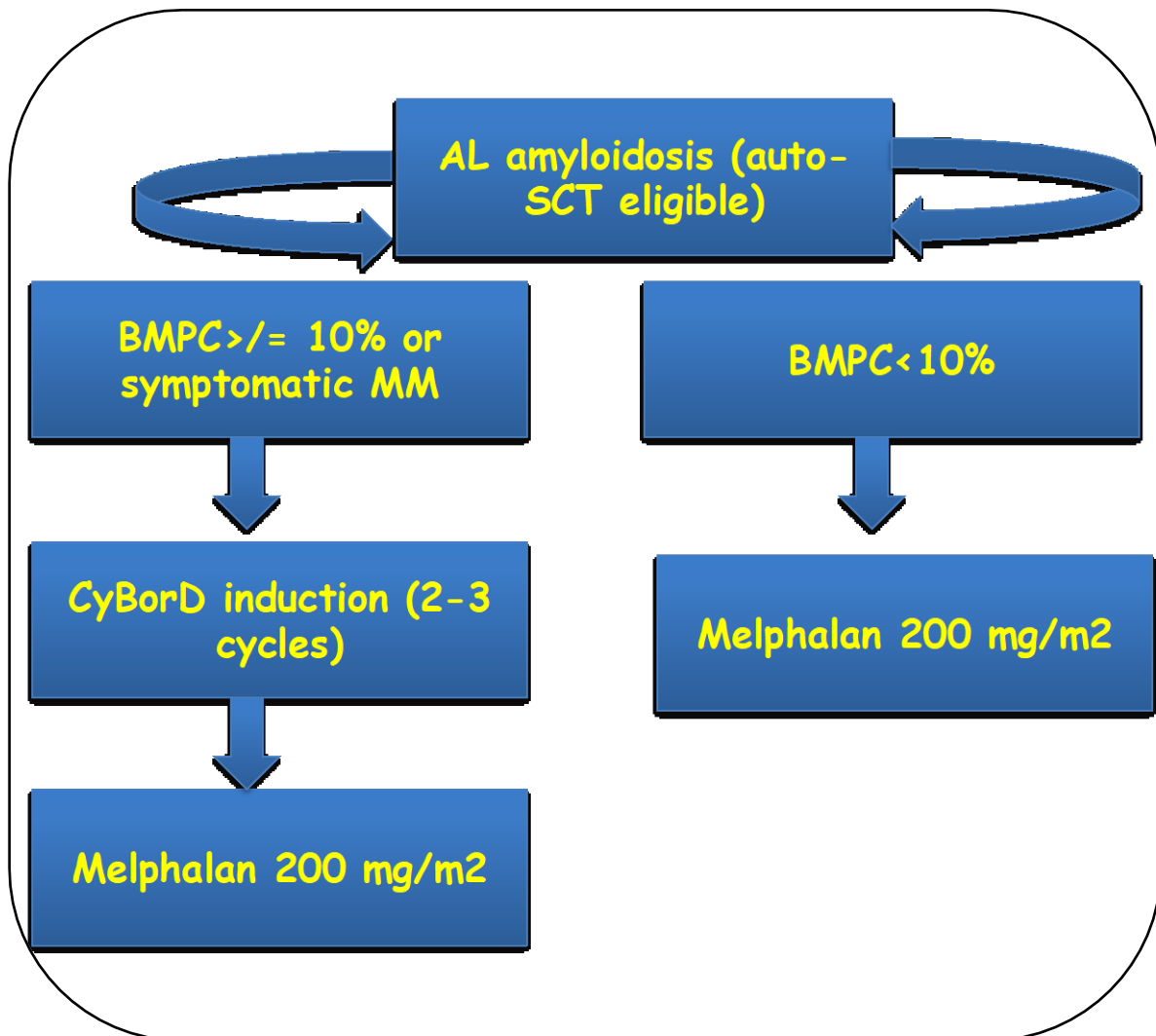
It is now apparent that a proportion of patients with significant cardiac involvement will substantially improve after achieving a good response to chemotherapy. While studies have examined the role of bortezomib-based induction chemotherapy immediately prior to ASCT, no studies to date have specifically focused on the role of deferred ASCT in transplant-ineligible patients.

A recent study by Manwani *et al.*<sup>116</sup> reported on 22 AL patients who underwent deferred ASCT. All patients were transplant-ineligible at presentation, predominantly due to advanced cardiac involvement. All received bortezomib-based therapy, with 100% hematologic response (86% complete response (CR)/very good partial response (VGPR)), enabling reversal of ASCT exclusion criteria. Patients underwent deferred ASCT for hematologic progression (45%) or consolidation

(55%). There was no transplant-related mortality. Hematologic responses post-ASCT: CR 50%, VGPR 27%, PR 18%, non-response 5%. In all, 85.7% achieved cardiac responses. Median overall survival (OS) was not reached, and median progression-free survival (PFS) was 54 months. This selected cohort achieved excellent hematologic responses, organ responses, PFS and OS with deferred ASCT.

If larger studies confirm these findings, this may widen the applicability of ASCT in AL Amyloidosis.

**Figure 1:** Recommended algorithm for ASCT in AL amyloidosis at Tom Baker Cancer Center.



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# Hematopoietic Cell Transplantation for Severe Aplastic Anemia

Presented by: Lynn Savoie

## Summary

- All patients with severe aplastic anemia should have HLA typing and a search for a related donor carried out at diagnosis.
- Patients less than 40 years old with a matched sibling donor should proceed directly to stem cell transplantation provided no contraindication to transplant exists.
- Patients greater than 40 years old and patients less than 40 years old without a matched sibling donor should receive immunosuppressive therapy with cyclosporine and equine anti-thymocyte globulin. They should proceed to stem cell transplantation from a matched sibling, matched unrelated donor, or a haploidentical donor if there is no clinically significant response after 6 months or if relapse.
- Expert opinion is divided on whether platelet transfusion-dependent patients should receive immunosuppressive therapy given the propensity of this treatment to increase platelet requirements and induce platelet refractoriness. These patients should be considered for early HCT if an appropriate donor can be identified in a suitable timeframe.
- A search for a MUD or a haploidentical donor should be initiated on patients without a matched sibling who show no response to immunosuppressive therapy after 3 months to allow a transplant to take place at 6 months.
- Conditioning
  - For matched sib HCT, fludarabine, cyclophosphamide and rabbit ATG. Additional GVHD prophylaxis will consist of methotrexate (day 1, 3, 6, 11), and cyclosporine for at least 6 months.
  - For matched unrelated donor and haploidentical HCT, rabbit ATG (day -9 to -7), fludarabine, low dose cyclophosphamide, and 2 Gy TBI (4 Gy if no previous immunosuppressive therapy). GVHD prophylaxis will consist of post-transplant cyclophosphamide (day 3 and 4), mycophenolate mofetil until day 35 and tacrolimus for at least 6 months.
  - See chapters on Conditioning and GVHD prophylaxis for details.
- Bone marrow will be the preferred source of stem cells in aplastic anemia. If considering an unrelated donor, choose one from a donor center that has a history of collecting consistently  $>2.5 \times 10^8$  NC/kg.
- Patients with recurrence of SAA after stem HCT may be considered for repeat transplantation or immunosuppressive therapy.

## Background<sup>1-4</sup>

Severe aplastic anemia (SAA) is an uncommon condition with an annual incidence rate of approximately 2 per million. While the majority of cases seen clinically are idiopathic, acquired SAA



has been described in relation to medications (chloramphenicol, gold salts, anticonvulsants), infection (e.g. non-A, B, C hepatitis or HIV), immune diseases (thymoma, eosinophilic fasciitis, graft-versus-host disease) and paroxysmal nocturnal hemoglobinuria (PNH). In children and young adults, hereditary conditions such as Fanconi anemia, dyskeratosis congenita and Schwachman- Diamond syndrome are important considerations and are frequently associated with non-hematological abnormalities. The manifestations of SAA occur as a result of damage to the hematopoietic stem cell compartment, making stem cell transplantation a natural treatment choice in this disease. For the purposes of these guidelines, SAA will be defined as follows:

- Bone marrow cellularity < 25% on an adequate biopsy and any two of the following:
- ANC (absolute neutrophil count) <  $0.5 \times 10^9 / L$
- Platelets <  $20 \times 10^9/L$
- Reticulocyte index < 1.0

## Results with Standard Treatment<sup>5-17</sup>

Immunosuppressive treatment with the combination of antithymocyte globulin (ATG) and cyclosporine has become standard treatment in SAA. Recent trials outlined in the table below show response rates of 65 to 75% and survival rates of 75 to 80% using this approach. Responses are generally delayed, with 85% of responses occurring in the first 3 months after treatment. As a general rule, response rates at 3, 6 and 12 months are 67%, 71% and 78%, respectively.

**Table 1.** Results of recent trials of standard treatment for severe aplastic anemia

| Study   | N   | Ages    | Response  | OS          | Relapse      |
|---------|-----|---------|-----------|-------------|--------------|
| NIH     | 122 | 35      | 61%       | 55% (7 y)   | 35% (5 year) |
| EBMT    | 182 | 25      | 83-85%    | NA          | NA           |
| Germany | 51  | 43      | 70% (6 m) | 64% (3.5 y) | 11%          |
| EBMT    | 46  | 29      | 74% (6 m) | 93% (4 y)   | NA           |
| Korea   | 83  | 14 – 40 | 47% (6 m) | 69% (6 y)   | 7.1%         |

Complications of immunosuppressive treatment include serum sickness due to heterologous protein in ATG, renal dysfunction and infectious illnesses. Over the longer term, patients are at risk of developing secondary myelodysplasia or AML: clonal disorders occur in 10 to 20% of SAA patients treated in this way. Relapses are not uncommon and may coincide with discontinuation of cyclosporine. Patients who fail a first course of immunosuppressive therapy (IST) for SAA may respond to retreatment with a similar regimen. Response rates in this situation are 43 to 77%. Response to IST is poorly defined, but at a minimum should include freedom from transfusions and neutropenic infections. Many patients will continue with abnormal blood counts indefinitely following successful IST.

In a randomized phase III study, the addition of Eltrombopag to standard immunosuppressive therapy increased the rapidity of response and the number of complete and overall responses without increased toxicity, as reported by Peffault de Latour in 2022. The percentage of patients who had a complete response at 3 months increased to 22% from 10% and the percentage of overall response at 6 months increased to 68% from 41%. The median time to first response decreased to 3.0 months from 8.8 months. EFS increased to 46% from 34% but this is an early timepoint for this assessment as many events happen late.

## Bone Marrow Transplantation in SAA<sup>17-26</sup>

Matched sibling bone marrow transplantation is the treatment of choice for young patients with a suitable donor, as these patients enjoy excellent long-term survival with few relapses. Outcome of transplantation in this group of patients is limited by graft rejection (reported in 3 to 23% of recipients) and GVHD but overall survival is reported to be 63 to 93% in single institution reports. The CIBMTR reported results on 1699 recipients of allogeneic transplantation for this disease, with 5 year survival rates of 75%, 68% and 35% for patients aged <20, 20-40 and >40, respectively.

Age at transplant has emerged as the major determinant of outcome and is used in most clinical algorithms to direct patients to the most appropriate treatment. Few quality of life studies have been carried out in this field; one such report found similar survival, event-free survival and quality-adjusted time without symptoms and toxicity (Q-TWiST) for bone marrow transplantation (BMT) and immunosuppression (IS), with BMT-treated patients enjoying longer periods free of symptoms and IS-treated patients requiring closer medical care, transfusion support and medications.<sup>24</sup>

The existing literature fails to distinguish outcomes for patients who undergo SCT as up-front treatment from those in whom it is used as second-line or salvage therapy. Small reports suggest that the outcome of SCT after failure of immunosuppressive therapy may approach that of first-line therapy,<sup>6</sup> while others have found a higher rate of graft rejection when transplant is undertaken in these circumstances.<sup>25</sup> The table below summarizes selected reports of outcome of BMT in SAA.

**Table 2.** Outcomes of BMT in severe aplastic anemia

| Study | Regimen | N    | Age | Engraftment | OS (time) | GVHD % (a/c) |
|-------|---------|------|-----|-------------|-----------|--------------|
| FHCRC | Cy-ATG  | 94   | 26  | 96%         | 88% (6y)  | 29/32        |
| GITMO | CyA-Mtx | 37   | 20  | 97%         | 94% (5y)  | 30/44        |
| EBMT  | BMT     | 1567 | NR  | NR          | 73% (10y) | NR           |
|       | IS      | 912  | NR  | NR          | 68% (10y) |              |
| IBMTR | Various | 471  | 20  | 84%         | 66%       | 19/32        |

Abbreviations: Cy-ATG = cyclosporine + antithymocyte globulin; CyA-Mtx = cyclosporine + methotrexate; GVHD = graft-versus-host-disease.

Early application of HCT to patients with IST-refractory SAA is essential. Our local results are in

keeping with those of other groups, which have shown that patients who receive a transplant for SAA more than two years after diagnosis have poor outcomes as shown below. It is essential that patients be taken to transplant as soon as possible (provided there are no contraindications) once patients are identified as being IST-refractory.

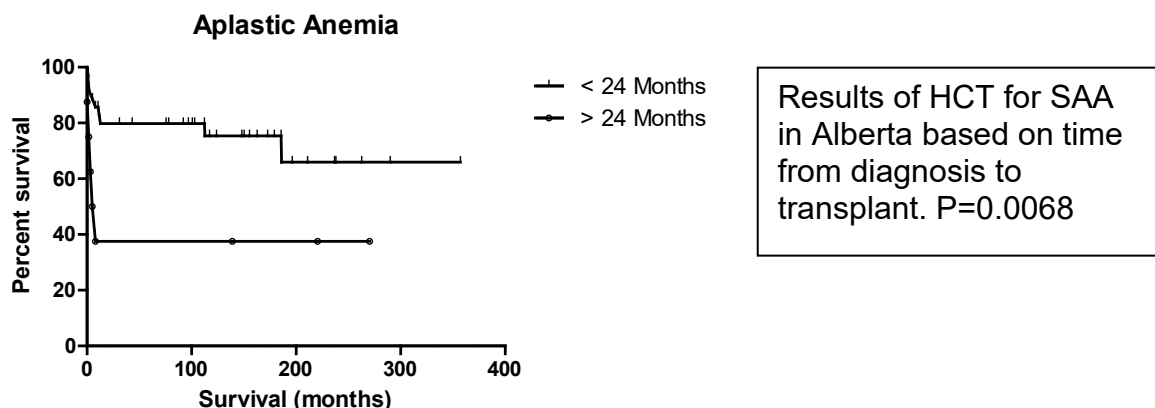


**Figure 1.** Percent survival over time for patients with aplastic anemia

Experience has been developed in the area of haploidentical HCT for SAA. Previous experience with haploidentical donors for transplantation in other contexts has demonstrated a high rate of graft failure, infection and treatment-related mortality. In aplastic anemia experience is limited but the results appear promising. Two publications have described the outcome of SAA patients who have received non-myeloablative HCT followed by G-CSF (granulocyte colony stimulating factor) mobilized peripheral blood stem cell (PBSC) grafts.<sup>31,32</sup> GVHD prophylaxis was with post-transplant cyclophosphamide, tacrolimus, and mycophenolate mofetil. Informal meta-analysis of these two reports indicates that engraftment occurs in approximately 90% of cases, and that overall survival at 1-2 years is 70-80%.<sup>33</sup> Further improvement appears to have been achieved by including rabbit antithymocyte globulin into the conditioning. The Johns Hopkins (Baltimore) group reported on 20 patients with relapsed/refractory SAA, who received ATG, fludarabine, low dose cyclophosphamide, 200 cGy TBI, marrow graft, and posttransplant cyclophosphamide, MMF and tacrolimus for haploidentical transplants<sup>35</sup>. Overall survival with a median follow-up of 32 months was 100%, and no graft failure or moderate to severe cGVHD occurred. They also report on 17 treatment naïve patients treated with the same protocol with similar results, the exception being an increase in graft failure that was resolved by increasing the TBI to 400cGy. A BMT CTN study using this same protocol<sup>36</sup> in 31 patients for haploidentical transplants in relapsed/refractory patients produced a 1-year OS of 81%, with deaths due to disease and unsuccessful bone marrow transplant (mostly graft failure?). A similar protocol has been used in 42 published patients with hemoglobinopathies (no previous immunosuppression other than hydroxyurea). In this setting, it was found that a higher dose of TBI may be associated with a decreased incidence of graft failure (6/14 haploidentical HCT recipients

developed GF after 200 cGy,<sup>37</sup> 1/8 after 300 cGy,<sup>38</sup> and 1/17 after 400 cGy TBI<sup>39</sup>). Given the encouraging results, we will offer haploidentical HCT to patients with relapsed/refractory SAA, using the Baltimore protocol with 200 cGy TBI. Despite we do not yet plan to routinely offer haploidentical HCT as primary treatment for SAA (without previous immunosuppressive therapy), if such a need arose, it would be prudent to use 400 instead of 200 cGy TBI.

## Transplant Details

In transplantation for malignant disease, the presence of graft-versus-host disease is associated with improved disease control and translates into superior disease-free survival. In aplastic anemia, graft-versus-host disease is deleterious to survival and has significant impact on patients' quality of life. Given the association between transplantation of stem cells from G-CSF mobilized peripheral blood and chronic GVHD (cGVHD), we will use bone marrow as the primary source of stem cells for transplantation in SAA.<sup>30</sup> Cyclosporine and short-course methotrexate will be used for GVHD prophylaxis given the results of randomized studies showing greater overall survival among patients treated in this way.

The conditioning regimen for patients undergoing stem cell transplantation for severe aplastic anemia from matched siblings has consisted of cyclophosphamide and ATG. With this approach it has been difficult to reduce the graft rejection rate below 10%, with consequent high transplant-related mortality (TRM) especially among older patients or those receiving transplants from mismatched or unrelated donors. The addition of fludarabine to Cy-ATG (FCA) has probably improved engraftment rates, and some series report engraftment rates of as much as 100% (see table below).<sup>27</sup> Retrospective comparisons of FCA with Cy-ATG show a trend to reduced rates of engraftment failure among those treated with FCA (0% vs. 11%,  $p=0.09$ ).<sup>28</sup> We use fludarabine 30 mg/m<sup>2</sup> daily x 4 days (days -5, -4, -3, -2), cyclophosphamide 60 mg/kg daily x 2 days (-3 and -2) and thymoglobulin 4.5 mg/kg (0.5 mg/kg on day -2 followed by 2 mg/kg on day -1 and day 0) for patients receiving transplants from HLA-matched related donors. Additional GVHD prophylaxis is with MTX (d 1,3,6,11) and CSA from day -1 to 6 months postHCT.

For matched unrelated donor and haploidentical transplants, we use the Baltimore protocol.<sup>34,39</sup> Conditioning consists of Thymoglobulin (0.5 mg/kg on day -9, 2 mg/kg on day -8, and 2 mg/kg on day -9), fludarabine (30 mg/m<sup>2</sup>/day on days -6 to -2), cyclophosphamide (14.5 mg/kg/day on days -6 and -5), and TBI 2 Gy on day -1 (4 Gy in a single fraction on day -1 if no previous immunosuppressive therapy). Marrow graft will be infused on day 0. Patients receive cyclophosphamide 50 mg/kg/day on days +3 and +4. Additional GVHD prophylaxis is with MMF 15 mg/kg (max 1 g tid), day 5 to 35, and Tacrolimus, day 5 to 6 mo postHCT.

Evidence has repeatedly shown that cell dose is important in order to avoid graft failure in patients with SAA, including in the haploidentical setting using ATG, fludarabine, low dose cyclophosphamide, low dose TBI, and additional GVHD prophylaxis with PTCy, MMF, and tacrolimus<sup>36</sup>. If using an

unrelated donor we will prioritize the use of donor centers with histories of collecting consistently  $>2.5 \times 10^8$  NC/kg.

**Table 3.**

| Study               | N  | Conditioning        | Product               | Graft Failure | aGVHD II-IV      | cGVHD | TRM             | OS (%)       |
|---------------------|----|---------------------|-----------------------|---------------|------------------|-------|-----------------|--------------|
| Bacigalupo EBMT MUD | 38 | Flu/CY/rATG         | BM=36; PBSC=2         | 18%           | 11%              | 27%   |                 | 72 (2 years) |
| Kang                | 5  | Flu/CY/rATG         | BM                    | 0             | 0 (1/5, grade I) | 0     |                 | 80           |
| Gupta               | 7  | Flu/CY/alemtuzumab  | BM                    | 0             | 3/7              | 1/6   | 2/7             |              |
| Chan                | 5  | Flu/CY/ATG          |                       | 0             | 80%              | 80%   | 0               |              |
| Urban               | 3  | Flu + other         | PBSC/CD34+ cells      | 0             |                  |       |                 |              |
| Vassiliou           | 8  | Alemtuzumab/CY/TBI  | MUD=7; haplo sib=1    | 0             | 25% (grade II)   | 0     | 0               | 100          |
| <b>MRD</b>          |    |                     |                       |               |                  |       |                 |              |
| George              | 35 | Flu/CY $\pm$ ATG    | G stim; BM=7; PBSC=28 | 2.8%          | 29% (I-IV)       | 32%   | 17.1% (day 100) | 82           |
| Resnick             | 13 | Flu/CY/ATG          | BM=4; PBSC=9          | 0             | 8.3%             | 12.5% |                 | 84 (5 years) |
| Koh                 | 8  | Flu/TBI             | PBSC; MRD=7; MUD=1    | 0             | 37.5%            | 60%   | 25%             | 75           |
| Rzepecki            | 5  | Flu/alemtuzumab/Mel | BM=2; PBSC=2          | 0             | 0                | 0     | 0               | 100          |
| Srinivasan          | 26 | Flu/CY/ATG          | PBSC; MRD=22; MMRD=4  | 0             | 65%              | 56%   | 3.8%            | 92           |
| Gupta               | 33 | CY/alemtuzumab      | BM=32; PBSC=1         | 24%           | 14%              | 4%    | 6/33            | 81 (5 years) |
| Gomez-Almaguer      | 23 | Bu/CY/Flu           | PBSC=23               | 26%           | 17.3%            | 26%   | 2/23            | 91           |

Abbreviations: aGVHD = acute GVHD; Bu = busulfan; cGVHD = chronic GVHD; CY = cyclophosphamide; Flu = fludarabine; MMRD = mismatched related donor; MRD = matched related donor; MUD = matched unrelated donor; PBSC = peripheral blood stem cells; TRM = treatment/transplant-related mortality.

**Table 4.** Results of haploidentical transplants in SAA (from Bacigalupo, Hematology 2018)<sup>33</sup>

| Ref.  | No. of Patients | Age, y | Conditioning | GVHD Proph               | SC source    | Engraftment | GVHD 2-4 | Alive at 1 y |
|-------|-----------------|--------|--------------|--------------------------|--------------|-------------|----------|--------------|
| 36    | 26              | 30     | RIC          | ATG CsA                  | BM           | 92%         | 10%      | 84%          |
| 37    | 21              | 14     | NMA          | CD3 dep                  | PB           | 96%         | 30%      | 94%          |
| 38    | 8               | 30     | NMA          | PTCY, FN, MMF            | GPregimen; B |             |          | 75%          |
| 39    | 17              | 19     | NMA          | ATG, Basilix, CsA        | GBM + GPB    | 90%         | 25%      | 65%          |
| 40    | 26              | 30     | NMA          | ATG, CsA, MTX, MMF       | GBM + GPB    | 92%         | 12%      | 84%          |
| 41    | 77              | 8      | NMA          | ATG, CsA, MTX, MMF + MSC | GBM + GPB    | 92%         | 12%      | 93%          |
| 42    | 13              | 30     | RIC          | PTCY, FK, MMF            | BM           | 100%        | 10%      | 100%         |
| 43    | 89              | 25     | RIC          | ATG, CsA, MTX, MMF       | GBM + GPB    | 97%         | 30%      | 86%          |
| Total | 277             | 27     |              |                          |              | 92%         | 12%      | 85%          |

BM, bone marrow; FK, tacrolimus; GVHD Proph, GVHD prophylaxis; MMF, mycophenolate; NMA, nonmyeloablative regimen; PB, peripheral blood; PTCY, high-dose posttransplant cyclophosphamide; RIC, reduced intensity conditioning regimen; SC source, stem cell source.



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# Hemoglobinopathies

Presented by: Kareem Jamani

## Summary

### Sickle Cell Disease

- Referrals for allo-HCT for SCD (typically sickle cell anemia and sickle cell  $\beta^0$  thalassemia) will be accepted from the Northern and Southern Alberta Rare Blood Disorders programs.
- Requirements for allo-HCT include:
  - An HLA-matched sibling or a haploidentical relative without SCD (sickle cell trait is acceptable).
  - Demonstrated compliance with medications and follow-up.
  - KPS >70, GFR >30 mL/minute, LVEF >40% and DLCO >50% predicted.
  - No evidence of cirrhosis or active hepatitis.
  - RBC allo-antibodies directed towards donor RBC antigens (including major ABO incompatibility) can lead to prolonged transfusion requirement post-HCT but do not appear to be associated with graft failure. The decision to proceed with HCT in this setting should be individualized.
- Indications for allo-HCT include any one of the following:
  - SCD-related end-organ complication (previous cerebrovascular event, sickle nephropathy, hepatopathy, or pulmonary artery hypertension by right heart catheterization or echocardiogram (TRV >2.5 m/s).
  - Reversible SCD-related complication not ameliorated by hydroxyurea (>2 vaso-occlusive crises/year requiring medical attention, >1 lifetime episode of acute chest syndrome, >1 episode of priapism/year requiring medical attention, proliferative retinopathy with visual impairment, >1 joint with avascular necrosis).
  - Red blood cell alloimmunization complicating chronic transfusion therapy.
  - Patients with combinations of clinical characteristics such as elevated WBC, elevated LDH, history of sepsis, age >35 and chronic transfusion who are at moderate-high risk of short-term mortality.
- Matched sibling donor HCT is performed according to the NIH protocol:
  - Conditioning is non-myeloablative and includes alemtuzumab (0.03 mg/kg D-7, 0.1 mg/kg D-6, 0.3 mg/kg D-5, -4, and -3) followed by TBI 3 Gy in a single fraction on D-2.
  - Grafts will be G-CSF mobilized PBSCs with a target of  $10 \times 10^6$  CD34+ cells/kg recipient weight.
  - GVHD prophylaxis is in the form of sirolimus starting on D-1 with a trough serum level of 5-15 ng/mL. Sirolimus should be maintained for at least 1 year and should be tapered thereafter only when donor T-cell chimerism is >50% in the absence of GVHD.
  - In the setting of sirolimus toxicity, alternate immunosuppression with mycophenolate should be considered as posterior reversible encephalopathy syndrome has been reported with calcineurin inhibitor use in this setting.

- Myeloid and T-cell chimerism should be measured at days 90, 180 and 365 post-HCT and yearly thereafter (however, if sirolimus is continued beyond 1 year, chimerism may be monitored more frequently, i.e., q. 3-6 months). RBC chimerism can also be monitored at these time points via Hb electrophoresis/HPLC.
- Haploidentical HCT is performed according to the Baltimore protocol:
  - Conditioning is non-myeloablative and includes Thymoglobulin (0.5 mg/kg on day -9, 2 mg/kg on day -8, 2 mg/kg on day -7), Thiotepa (10 mg/kg on day -7), Fludarabine (30 mg/m<sup>2</sup> daily from day -6 to -2), Cyclophosphamide (14.5 mg/kg daily on day -6 and -5), and TBI (2 Gy in a single fraction on day -1).
  - Bone marrow graft.
  - GVHD prophylaxis consists of posttransplant cyclophosphamide (50 mg/kg daily on day +3 and +4), mycophenolate mofetil from day +5 to +35 (15 mg/kg/d tid, max 1 g tid), and sirolimus from day +5 to (target 5 to 15 ng/dL). Sirolimus should be maintained for at least 1 year and should be tapered thereafter only when donor T-cell chimerism is >50% in the absence of GVHD.
- Supportive care measures will be provided as outlined in the ABMTP standard practice guidelines, with the following modifications:
  - Patients should undergo exchange transfusion with a goal HbS <30% and Hb 90-100 g/L on D-10. Extended phenotype-matched RBC units (ABO, Rh D, C/c, E/e & Kell) should be used for exchange transfusion (the need for, on average, 7 units should be communicated to transfusion medicine in advance).
  - The transfusion target for Hb and platelets post-HCT should be 90-100 and 50, respectively.
  - If RBC allo-antibodies are identified it should be ensured that enough antigen negative units will be available for transfusion post-HCT (on average 6 units).
  - Hydroxyurea should be discontinued on 1 day before starting ATG or alemtuzumab.
  - G-CSF should be avoided altogether given the adverse outcomes associated with this medication in SCD.
  - Penicillin V prophylaxis should be provided until completion of pneumococcal vaccination, i.e., 2 years posttransplant (in addition to trimethoprim-sulfamethoxazole until 3 mo after discontinuation of immunosuppression).
- Patients should be counseled regarding possible late toxicities of allo-HCT including infertility (and preservation options) and therapy-related myeloid neoplasms.

## Thalassemia

- At this time, allo-HCT for adults with thalassemia should not be offered outside of a clinical trial.

# Allo-HCT for Sickle Cell Disease

## Background

Sickle cell disease (SCD) is a severe monogenic autosomal recessive multisystem disease characterized by “sickled” erythrocytes. While SCD is an overarching term referring to all genotypes that cause this clinical syndrome, sickle cell anemia (SCA) refers to the most common form of the disease (70% of cases) resulting from homozygosity for the sickle cell allele (the majority of remaining cases result from hemoglobin SC and sickle cell/ $\beta$ -thalassemia)<sup>1</sup>. Sickled hemoglobin (Hb S) results from a point mutation in the  $\beta$ -globin gene in which a single nucleotide of glutamic acid is replaced with valine. The consequence is a hydrophobic patch on the  $\beta$ -globin molecule, which allows binding of  $\beta$ -globin chains of two hemoglobin molecules when deoxygenated and thus polymerization of hemoglobin molecules<sup>1</sup>. Ultimately, the result is a distortion in the shape of the erythrocyte and a significant loss of its flexibility.

The underlying pathophysiology of SCD is complex. At the most basic level, sickled erythrocytes contribute to both chronic hemolysis and vaso-occlusion with resultant tissue hypoxia. Recent work has produced additional insights into SCD pathophysiology including the role of vasculopathy and endothelial cell dysfunction, dysregulated inflammatory responses and innate immunity, oxidant stress and iron dysregulation, and sensitization of the nervous system to pain stimuli<sup>2</sup>. The resultant clinical manifestations of SCD are summarized in Table 1.

**Table 1.** Clinical manifestations of sickle cell disease

| SCD Pathology or Outcome | Clinical Manifestation                    |
|--------------------------|-------------------------------------------|
| Chronic hemolysis        | Pulmonary hypertension                    |
|                          | Gallstones                                |
|                          | Fatigue                                   |
| Vaso-occlusive events    | Acute pain                                |
|                          | Chronic pain                              |
|                          | Acute chest syndrome                      |
|                          | Osteonecrosis                             |
|                          | Priapism                                  |
| Vasculopathy             | Retinopathy                               |
|                          | Stroke/Moyamoya and neurologic impairment |
|                          | Nephropathy                               |
|                          | Hepatopathy                               |
|                          | Asplenia and infection                    |
|                          | Hypercoagulability                        |
| Chronic Transfusion      | Iron overload                             |
|                          | RBC allo-immunization                     |
| Poor Quality of Life     | Poor educational outcomes                 |
|                          | Lack of employment                        |
|                          | Mental illness                            |
|                          | Stigma                                    |



Advances in SCD care; notably newborn screening, penicillin prophylaxis, vaccination, transcranial Doppler monitoring with pre-emptive transfusion therapy for primary stroke prevention and hydroxyurea therapy<sup>3</sup>; have led to significant improvements in survival in children with SCD<sup>4</sup>. Hydroxyurea, the only approved disease-modifying pharmacotherapy for SCD, has been shown to reduce the incidence of vaso-occlusive pain crises, acute chest syndrome and red cell transfusion as well as improve survival in SCD<sup>3</sup>. Yet, over the last 30 years, there has been no improvement in the survival of adults with SCD. In a large American longitudinal study, mortality in adults with SCD appeared to increase by 1% in each year studied from 1979 to 2005 and the median age at death in 2005 was 42 and 38 years for females and males, respectively<sup>5</sup>. In another recent American prospective observational cohort, those with SCA had a median survival of 58 years<sup>6</sup>. In recent years, the most common cause of death in SCD is chronic cardiopulmonary disease, including chronic lung disease, pulmonary hypertension, congestive heart failure, myocardial ischemia and venous thromboembolic disease<sup>7, 8</sup>. There is no convincing evidence to suggest that hydroxyurea alters the incidence or course of chronic SCD-related cardiopulmonary disease<sup>3, 9</sup>. Thus, in adults, despite hydroxyurea and improvements in supportive care, SCD continues to reduce life expectancy.

### **Allo-HCT for SCD**

The recognition that those with SCD continue to suffer poor outcomes has led to growing interest in the development of disease-modifying and potentially curative therapy, including allogeneic hematopoietic cell transplantation (allo-HCT). In 1996, Walters et al demonstrated that allo-HCT from HLA-matched siblings with myeloablative conditioning (Bu/Cy/ATG) was feasible in children and resulted in sustained engraftment, elimination of vaso-occlusive episodes and stability in SCD-related end-organ damage present pre-transplant<sup>10</sup>. In children, experience with allo-HCT has rapidly expanded since that time; outcomes with a variety of conditioning strategies are excellent with CIBMTR (Center for International Blood and Marrow Transplant Research) and EBMT (European Group for Blood and Marrow Transplantation) registries reporting >90% 1 year survival and low rates of graft-versus-host-disease (GVHD) for those receiving HLA-matched sibling HCT<sup>11</sup>.

In adults, there are now several published reports of allo-HCT for SCD. Encouraging results, particularly with non-myeloablative approaches have been reported (summarized in table 2). In the earliest attempt at myeloablative conditioning, the Chicago group reported on 2 patients receiving HLA-matched sibling peripheral blood stem cells (PBSC) after conditioning with Flu/Mel/ATG. Both patients engrafted and neither had SCD-related complications post-HCT, however, both died before 1 year from GVHD/infection<sup>12</sup>. A French group reported on 15 patients receiving HLA-matched sibling bone marrow after conditioning with Bu/Cy/ATG. All patients engrafted and one patient experienced early mortality due to cerebral hemorrhage in the setting of severe cerebral vasculopathy. At a median follow-up of 3.4 years: DFS was 93%, half of patients developed steroid-responsive grade 2-3 aGVHD, 2 patients developed moderate cGVHD, donor chimerism was sustained with all patients off immunosuppression, and all patients enjoy normal quality of life per the authors<sup>13</sup>. More recently, a multi-centre prospective American pilot study reported on 22 patients receiving HLA-matched sibling



(17) or unrelated bone marrow (5) after conditioning with Flu/Bu/ATG. All patients engrafted and two patients experienced early mortality (intracranial hemorrhage and GVHD). One year OS and EFS were 91% and 86%. Four patients developed grades 2-3 acute GVHD, 3 developed moderate-severe chronic GVHD and one developed secondary graft failure and is alive after a second transplant. Significant improvements in health-related quality of life and pain were observed<sup>14</sup>.

**Table 2.** Studies of allo-HCT for Sickle Cell Disease

| Ref                      | N  | Donors/<br>Graft                    | Conditioning<br>and GVHD<br>prophylaxis               | Engraftment                     | GVHD                                                          | TRM  | SCD-Specific Outcome                                                                                      |
|--------------------------|----|-------------------------------------|-------------------------------------------------------|---------------------------------|---------------------------------------------------------------|------|-----------------------------------------------------------------------------------------------------------|
| <b>Myeloablative</b>     |    |                                     |                                                       |                                 |                                                               |      |                                                                                                           |
| 12                       | 2  | MSD /<br>PBSC                       | Flu/Mel/ATG<br>MTX/Tac                                | 2/2                             | 1 acute/1<br>chronic                                          | 2/2  | No acute SCD complications                                                                                |
| 13                       | 15 | MSD /<br>BM                         | Bu/Cy/ATG<br>MTX/Csa                                  | 15/15                           | Acute: 7 grade II<br>1 grade III<br>Chronic: 2 mod-<br>severe | 1/15 | 14/15 “normal” QoL & no<br>immune suppression                                                             |
| 14                       | 22 | MSD (17)<br>or MUD<br>(5) / BM      | Flu/Bu/ATG<br>MTX/Csa                                 | 22/22 (1 late<br>graft failure) | Acute: 4 grade<br>2-3<br>Mod-Severe<br>Chronic: 3             | 2/22 | No SCD recurrence post<br>HCT. ↑ HR-QoL and ↓ pain.                                                       |
| <b>Non-myeloablative</b> |    |                                     |                                                       |                                 |                                                               |      |                                                                                                           |
| 15                       | 30 | MSD /<br>PBSC                       | Alem/TBI<br>Sirolimus                                 | 26/30                           | None                                                          | 0/30 | ↓TRV ↓Hospitalization ↓Narcs<br>No recurrent neurologic<br>events<br>15/26 off sirolimus @ med<br>2.1 yrs |
| 16                       | 13 | MSD /<br>PBSC                       | Alem/TBI<br>Sirolimus                                 | 12/13                           | None                                                          | 0/13 | ↑QoL ↓BNP ↑FEV1&FVC<br>4/12 off sirolimus at med f-up<br>22 mos                                           |
| 20                       | 17 | Haplo<br>(14) or<br>MSD (3) /<br>BM | ATG/Flu/Cy/<br>TBI200/PTCy/<br>MMF/Tacro or<br>Sirol  | 11/17                           | No gr 2-4<br>aGVHD,<br>No mod-sev<br>cGVHD                    | 0/17 | 10/17 disease-free<br>(transfusion-independent, off<br>narcs)                                             |
| 21                       | 8  | Haplo /<br>PBSC                     | ATG/Flu/Cy/<br>TBI300/PTCy/<br>MMF/Sirol              | 7/8                             | 2 gr 2-4 aGVHD,<br>1 mod-sev<br>cGVHD                         | 1/8  | 6/8 disease-free                                                                                          |
| 27                       | 17 | Haplo /<br>BM                       | ATG/Flu/Cy/<br>TBI400/PTCy/<br>MMF/Sirol              | 16/17                           | 5 gr 2-4 aGVHD,<br>1 mod-sev<br>cGVHD                         | 0/17 | 16/17 disease-free                                                                                        |
| 28                       | 42 | Haplo/<br>BM                        | ATG/Flu/Thiote<br>pa/Cy/TBI200/<br>PTCy/MMF/Sir<br>ol | 39/42                           | 26% gr 2-4<br>aGVHD, 7%<br>mod-sev<br>cGVHD                   | 2/42 | Not yet reported                                                                                          |
| 29                       | 38 | Haplo/<br>BM                        | ATG/Flu/Thiote<br>pa/Cy/TBI200/<br>PTCy/MMF/Sir<br>ol | 38/38                           | 8% gr 3-4<br>aGVHD, 3%<br>mod-sev<br>cGVHD                    | 3/38 | Not yet reported                                                                                          |

Abbreviations: Alem = alemtuzumab; ATG = anti-thymocyte globulin; ATLg = anti-Jurkat T cell globulin; BM = bone marrow; Csa = cyclosporine; Flu = fludarabine; Mel = melphalan; MMF = mycophenolate mofetil; MSD = matched sibling donor; MTX = methotrexate; MUD = matched unrelated donor; PTCy = post-transplantation cyclophosphamide; Tac = tacrolimus; TBI = total body irradiation; TRM = treatment-related mortality.

The most extensively reported experience in adults, and the approach to be used in the Alberta Bone Marrow Transplant Program (ABMTP), is with non-myeloablative conditioning from matched sibling or haploidentical donors. This approach aims to produce mixed chimerism to alleviate the SCD phenotype while maintaining low non-relapse mortality (NRM). The group at the NIH has reported results of a phase 1/2 trial involving 30 patients given alemtuzumab and low dose TBI conditioning followed by infusion of sibling HLA-matched PBSCs and sirolimus for GVHD/graft failure prophylaxis<sup>15</sup>. Patients were followed for a median of 3.4 years. All patients initially engrafted but 4 subsequently experienced graft failure with recurrence of SCD and one of these patients died from intracranial hemorrhage. In patients who had sustained engraftment, mean donor T-cell and myeloid chimerism were 48% and 86%, respectively. Chimerism was monitored frequently and withdrawal of sirolimus was considered at 1 year or more post-HCT if T-cell chimerism was >50% donor. Fifteen patients were able to discontinue immunosuppression at a median of 2.1 years and the remainder continue due to inadequate T-cell chimerism. NRM and GVHD were not observed. In those with sustained engraftment, specific SCD outcomes included reduction in tricuspid regurgitant velocity (TRV), no recurrent neurologic events, reduction in hospitalization rate and reduction in narcotic use. These findings have recently been replicated by the Chicago group in 13 patients<sup>16</sup>. At a median follow-up of 22 months; 1 patient experienced secondary graft failure (non-compliant with sirolimus) and the rest had stable mixed chimerism, 4 were able to discontinue sirolimus, quality of life scores improved at 1-year post-HCT and no TRM or GVHD were observed. There was significant improvement in cardiopulmonary parameters at 1 year. Of note, 2 patients were transplanted across major ABO incompatibility without engraftment concerns.

### Late Outcomes

Beyond improvements in quality of life and pain after allo-HCT as noted above, emerging data appears to confirm the protective effect of allo-HCT on organ function. Specifically (as well-reviewed by Hulbert *et al.*)<sup>17</sup>, allo-HCT appears to be associated with: a reduced incidence of stroke, stable or improved neurocognitive outcomes, a reduction in glomerular hyperfiltration, stability or improvement in restrictive or obstructive lung disease, improvements in diastolic filling/cardiac size and a reduction in tricuspid regurgitant velocity.

Regarding toxicities: acute kidney injury and hypertension appear relatively common after allo-HCT, warranting careful attention to nephrotoxins in the peri-HCT period and careful monitoring of blood pressure post-transplant<sup>17</sup>. An additional concern is the development of therapy-related myeloid neoplasms (TRMN) after non-myeloablative conditioned allo-HCT for sickle cell disease: Investigators at the NIH recently reported that 5/120 recipients (4.2%) developed a TRMN, all in the setting of residual host hematopoiesis (4 graft failure and 1 loss of myeloid chimerism). The investigators hypothesize that the TRMNs after non-myeloablative HCT are a result of selective pressure, induced by graft failure, placed on autologous preleukemic myeloid clones (which are especially noted to be present in older adults with more severe sickle cell disease)<sup>18</sup>.

## Use of Alternative Donors

Most SCD patients will not have a suitable matched sibling donor available, thus, there is significant interest in the use of alternative donors. The use of MUDs has predominantly been described in children with the largest series (29 patients) reporting a 28% treatment-related mortality after reduced-intensity conditioning with alemtuzumab, melphalan, and fludarabine, predominantly due to GVHD<sup>19</sup>. Haploidentical allo-HCT with post-transplant cyclophosphamide is promising. Initially, it was hampered by high rates of graft failure<sup>20, 21</sup>. However, with the newest version of the Baltimore protocol (using pretransplant ATG and posttransplant cyclophosphamide), of 17 patients who underwent haploidentical HCT, all 17 survived, only one developed moderate-severe cGVHD, only one developed graft failure, and 13/16 engrafted patients demonstrated full donor chimerism<sup>27</sup>. A modification of the Baltimore protocol (addition of a single dose of thiotepa and reduction in the TBI dose to 200 cGy) was subsequently studied in two prospective clinical trials (the BMT CTN 1507 and the Vanderbilt International Learning Collaborative) enrolling a total of 80 patients.<sup>28, 29</sup> Combined, 77/80 patients engrafted, the rates of grade 3-4 aGVHD and moderate-severe cGVHD were <10% each and TRM was reported in 5/80. Importantly, all engrafted patients demonstrated 100% donor chimerism at the 1-year mark. Full donor chimerism is particularly important to reduce the risk of TRMN as described above. Thus, we will use the modification of the Baltimore protocol (thiotepa and 2 Gy TBI), as outlined in the Summary, above. Novel approaches involving ex-vivo T-cell depletion, such as  $\alpha/\beta$  T-cell depletion, have shown promise but are in their infancy<sup>22</sup>. The use of umbilical cord grafts has not been described in adult SCD patients.

## Patient Selection

SCD results in phenotypic diversity. Recent efforts have focused on identifying specific clinical features that are associated with risk of mortality with standard SCD care. In a recent review of observational SCD studies: elevated TRV, leukocytosis and chronic transfusion were associated with 10% 2-year mortality, while elevated NT-proBNP, history of sepsis, elevated LDH (lactate dehydrogenase) and age >35 were associated with 5-9% 2-year mortality. Having a combination of two of these features led to 7-24% 2-year mortality<sup>23</sup>. Other end organ complications like sickle hepatopathy, sickle nephropathy, cerebrovascular events and acute chest syndrome are also associated with mortality<sup>24</sup>. In addition, recurrent vaso-occlusive crises, sickle retinopathy and osteonecrosis lead to significant morbidity. Given the low NRM, patients with over 5% 2-year mortality are likely to benefit from matched sibling HCT. In contrast, only patients with higher (>10%) estimated 2-year mortality are likely to benefit from higher risk grafts (MUD, haploidentical and umbilical cord)<sup>23</sup>. Specific indications for allo-HCT in the SCD in the two non-myeloablative trials described above include: end-organ complication (previous cerebrovascular event, sickle nephropathy or hepatopathy, TRV >2.5 m/s), a reversible complication not ameliorated by hydroxyurea (>2 vaso-occlusive crises/year requiring medical attention, >1 lifetime episode of acute chest syndrome, >1 episode of priapism/year requiring medical attention, proliferative retinopathy with visual impairment or >1 joint with avascular necrosis) or red blood cell alloimmunization during chronic transfusion therapy<sup>15, 20</sup>. RBC allo-antibodies directed towards donor RBC antigens (including major ABO incompatibility) can

lead to prolonged transfusion requirement post-HCT but do not appear to be associated with graft failure. The decision to proceed with HCT in this setting should be individualized. Given the risk of secondary graft failure and infectious or toxic complications of allo-HCT, demonstrated compliance with medications and follow-up is crucial. Candidates for allo-HCT should be referred by an SCD expert after a comprehensive assessment of SCD status. Most patients who meet the above inclusion criteria will have an elevated HCT-CI (hematopoietic cell transplantation comorbidity index), making non-myeloablative conditioning an attractive option. Minimal functional status and organ function criteria, however, in the above trials has included: KPS >70, GFR >30 mL/minute, LVEF >40% and DLCO (diffuse capacity of lung for carbon monoxide) >50% predicted. Active hepatitis and a diagnosis of cirrhosis are exclusion criteria.

### **SCD-Specific Supportive Care for Allo-HCT**

Because of the unique physiological circumstances in SCD and the potentially toxic nature of allo-HCT, additional supportive care measures will apply to these patients in addition to standard allo-HCT care.

1. There is a risk of gonadal failure after low dose TBI. Patients should be counseled about fertility preservation options. Testicular shielding will be used during TBI treatment. Our center does not have the capacity to provide ovarian shielding.
2. Medication management: hydroxyurea should be discontinued the day before conditioning begins and G-CSF should be avoided given its association with severe SCD-related acute complications (vaso-occlusive events, acute chest syndrome, multi-organ failure and death)<sup>25</sup>.
3. Transfusion medicine: As per standard allo-HCT practice, transfused blood products should be irradiated. The target hemoglobin (Hb) in the peri-transplant period is 90-100 g/L. The need for extended phenotype-matched RBC units (ABO, Rh D, C, E & Kell) should be communicated to transfusion medicine. A median of 6 (range 0-15) units of RBCs transfused has been reported with the NIH non-myeloablative protocol. An RBC antibody screen should be performed during pre-HCT workup and if RBC allo-antibodies are identified, it should be ensured that enough antigen negative units will be available for transfusion post HCT. Given the physiologic stress (fever, infection, volume depletion etc.) likely to be encountered post-HCT and the associated risk of an SCD-related acute event, patients should undergo exchange transfusion with a goal HbS <30% and Hb 90-100 g/L (using the above noted RBC unit attributes) prior to beginning conditioning. Given the risk of CNS bleeding in the setting of vasculopathy and thrombocytopenia, the transfusion target for platelets post-HCT should be 50. A median of 4 platelet units (range 0-19) were required to achieve this target with the NIH protocol.
4. Additional supportive care measures should include careful attention to hydration status, encouraging mobilization and out of hospital passes when appropriate, pharmacologic venous thromboembolism prophylaxis if the patient remains on the inpatient unit and platelets are >50 and use of incentive spirometry when on the inpatient unit.
5. Infectious prophylaxis, including CMV monitoring and pre-emptive therapy, should be per current ABMTP practice guidelines, with the following modifications:

- a. Penicillin V prophylaxis should be provided until completion of pneumococcal vaccination, i.e., 2-years posttransplant (in addition to trimethoprim-sulfamethoxazole until 3 mo after discontinuation of immunosuppression).
- b. While EBV viremia is expected to be uncommon, the approach should be individualized given the risk of secondary graft failure or GVHD with tapering immunosuppression, i.e., use of rituximab only (without immunosuppression taper) should be considered.

## Allo-HCT for Thalassemia

There is very limited experience with allo-HCT for adults with  $\beta$ -thalassemia major. Myeloablative approaches have resulted in high non-relapse mortality and outcomes are primarily determined by hepatic iron overload status<sup>26</sup>. There are no significant reports of reduced intensity or non-myeloablative approaches in this patient population. At this time, allo-HCT for adults with thalassemia should not be routinely offered outside of a clinical trial.

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# Multiple Sclerosis

Presented by: Jodie Burton and Jan Storek

## Summary

- Eligibility for autologous hematopoietic stem cell transplantation (autoHCT) includes poorly controlled relapsing-remitting multiple sclerosis (RRMS) or apparent pseudo-progression in highly select group of patients
- Relapsing-remitting patients will be eligible if they have failed a second disease modifying therapy (DMT), or are intolerant of multiple DMTs. In special cases, RRMS patients might be eligible having failed only one DMT (e.g., high risk of PML).
- “Pseudo-progressive” patients will be eligible if they meet stringent criteria and consensus agreement by an MS neurologist experienced with the use of AHSCT in MS and a transplant physician
- For transplant technique, we use mobilization with cyclophosphamide + GCSF + dexamethasone, no CD34 enrichment, and conditioning with cyclophosphamide (200 mg/m<sup>2</sup>) + Thymoglobulin (4.5 mg/m<sup>2</sup>), and more intense infection prophylaxis than for patients with malignancies.

## Background

Multiple Sclerosis (MS) is the most common neurodegenerative disease of non-elderly adults in North America, with a prevalence of roughly 1/385 in Alberta, Canada<sup>1,2</sup>. It is characterized by central nervous system (CNS) demyelination and axonal loss/degeneration. Most patients present with the relapsing-remitting (RRMS) form of the disease, characterized by episodes of CNS dysfunction that typically last weeks with fair to good recovery<sup>3</sup>. The average patient is female, age 32, and while there is a small impact on life expectancy, it is typically in single digit years, thus patients will incur disability over decades and all the direct and indirect costs that entails<sup>3</sup>.

### First-Line Multiple Sclerosis Disease Modifying Treatment

Since the mid 1990s, parenteral agents, interferon beta (Avonex®, Rebif®, Betaseron®) and glatiramer acetate (Copaxone®), to reduce relapse frequency in RRMS have been available<sup>4-7</sup>. While mildly to moderately effective, these agents reduce relapse rates by roughly 30%, and 30% or more of patients on these agents are considered treatment failures<sup>4-7</sup>. An additional subset of patients fails to tolerate these agents due to common adverse events of flu-like symptoms, leucopenia, transaminitis and a variety of skin manifestations<sup>4-7</sup>. In 2013, dimethyl fumarate (Tecfidera®), an oral agent taken twice daily, was approved for RRMS, soon thereafter joining the approved first-line agents in Alberta. This agent has demonstrated roughly a 50% reduction relapse rate versus placebo and ~ 34% versus Copaxone®<sup>8,9</sup>. Tecfidera® is associated with a small risk of lymphopenia, typically

manifesting in the first 6-months of use, which typically persists, and if grade 3 or higher, requires discontinuation to avoid immunosuppressive complications. There have also been a small number of cases of PML, most of which have been linked to ongoing lymphopenia and ongoing use of Tecfidera®.<sup>10</sup> Teriflunomide (Aubagio®), a once daily oral agent approved in 2013, has also been added to the first-line arsenal. In the pivotal trials, Aubagio® showed a 31-36% relative reduction in relapse activity, with adverse events that include hair thinning/loss and the risk of teratogenicity (based mostly on animal data)<sup>11</sup>. A proportion of patients, approximately 4-14%, have what is considered to be aggressive multiple sclerosis, defined as reaching a high degree of disability within 5 years of disease onset or age 40, or transitioning to progressive MS within only 3 years of disease onset<sup>12</sup>.

Ocrelizumab/Ocrevus®, approved in Canada in 2018 for both RRMS and PPMS, is a humanized anti-CD20 monoclonal antibody given by infusion every 6 months (similar to rituximab which is not approved as a DMT in Canada). In pivotal trials in RRMS, relapse rates were reduced by 46-47% vs Rebif, with a relatively tolerable adverse event profile<sup>13</sup>. The apparent small increased risk of breast cancer associated with Ocrelizumab has since been disproven<sup>14</sup>. Extension and post-marketing trials of Ocrelizumab have shown it to be consistently highly efficacious in RRMS<sup>15</sup>, but associated with hypogammaglobulinemia, albeit rarely symptomatic<sup>16</sup>. This agent was approved by Alberta Blue Cross as a first-line agent in April 2019.

Ofatumumab/Kesimpta®, approved in Canada in 2021 for RRMS, is a fully humanized monoclonal antibody targeting a different component of the CD20 receptor, and is given subcutaneously every month. In pivotal trials, relapse rates in patients receiving ofatumumab versus teriflunomide were 0.11 vs 0.22 and 0.10 vs 0.25<sup>17</sup>. A similar risk profile to Ocrelizumab with respect to typically asymptomatic hypogammaglobulinemia was seen<sup>17</sup>. This agent was approved by Alberta Blue Cross as a first-line agent in May 2022.

Of note, a sizable proportion of RRMS patients, particularly those who are newly diagnosed, start with ocrelizumab or ofatumumab.

## **Second Line-Escalation Disease Modifying Treatment**

In truth, escalation agents (typically classic immunosuppressants such as azathioprine and cyclophosphamide) have been used for decades, but those with randomized control trial evidence have only been available since 2000. Mitoxantrone (Novantrone®) was approved for use in worsening RRMS and secondary progressive MS in 2000, although its use has decreased considerably in the wake of relatively high rates of serious adverse events including cardiac dysfunction, leukemia and bone marrow damage<sup>18</sup>. In 2006, Natalizumab (Tysabri®) was approved for use in RRMS in the context of marked failure on conventional agents<sup>19,20</sup>. Although highly effective, it has become evident that the risk of progressive multifocal leukoencephalopathy (PML) from JC virus entry into the CNS is as high as 1/30 patients based on risk factor stratification<sup>21,22</sup>. In

Alberta, all currently approved therapies not categorized as first-line therapy are considered second-line (i.e. there are no therapies solely categorized by Alberta and Blur Cross as third-line). These agents include:

#### **Fingolimod/Gilenya®:**

Gilenya® was the first oral agent in RRMS was approved (for RRMS) in Canada. This agent has a novel mechanism of action characterized by activation of lymphocyte S1P1 via high-affinity receptor binding that subsequently induces S1P1 down-regulation, preventing lymphocyte egress from lymphoid tissues and thus reducing autoaggressive lymphocyte infiltration into the central nervous system (CNS)<sup>23,24</sup>.

In pivotal trials, there was a 54% relative reduction in relapses versus placebo (52% versus Avonex®), as well as significant reductions in MRI lesion load, and markers of disability progression<sup>23,24</sup>. It is also associated with rare cardiac, respiratory adverse events as well as viral infectious (namely varicella zoster virus reactivation, i.e. shingles) and leads to an expected apparent lymphopenia due to its mechanism of action<sup>23,24</sup>. It, like all agents mentioned below is considered a second-line/escalation agent in Canada<sup>25</sup>. Since its approval, there have been upwards of 15 cases of PML associated with Gilenya® use, with a cited risk of 3.12 per 100,000<sup>26</sup>. The only risk factor identified thus far is duration of use.

#### **Alemtuzumab/Lemtrada®:**

As well, Alemtuzumab, a very potent intravenous escalation agent with compelling results was approved in Canada in December 2013<sup>27</sup>. It is currently covered in the province of Alberta as a second-line treatment. Use of Alemtuzumab requires long-term monitoring of a minimum of four to five years of monthly blood and urine testing for potentially significant side effects (thyroid dysfunction, idiopathic thrombocytopenia purpura and Goodpasture syndrome)<sup>27</sup>. More recently, additional risks including Acute acalculous cholecystitis and stroke during infusions have been reported<sup>28,29</sup>.

#### **Cladribine/Mavenclad®:**

Cladribine, approved for use in RRMS in Canada in 2018, is a purine nucleoside analogue that selectively depletes peripheral lymphocytes without a major impact on cells of the innate immune system. It is given in oral form given as a weight-based dose in two relatively short courses over two annual cycles. Oral cladribine results in the peripheral depletion of lymphocytes that is gradual, occurring over several weeks, and is not associated with a cell lysis syndrome, has a greater impact on B cells than T cells, and is followed by gradual reconstitution of the peripheral lymphocyte counts over several months<sup>30</sup>. In pivotal trials, cladribine patients had a relative relapse reduction of 57% compared to placebo. Beyond typical mild adverse events, there is a risk of lymphopenia with cladribine, which may lead to a delay or cancellation of the second cycle of treatment if persistent<sup>31</sup>. Recent studies have demonstrated that the duration of “no evidence of disease activity” with Mavenclad® is relatively short, although not synonymous with needing further treatment<sup>32</sup>.

## The History of Transplantation Therapy in MS

Multiple randomized studies have been initiated comparing autologous transplantation to conventional therapy in MS or other autoimmune diseases. Over the history of these trials, both efficacy and toxicity has improved, due in part to improved patient selection restricting enrollment to less advanced patients. Transplant-related mortality for MS in Europe dropped from 7.3% in 1995-2000 to 1.3% in 2001-2007<sup>33</sup>. Trial regimens include the use of agents such as busulfan or BEAM. According to the European Bone Marrow Transplant Registry (EBMTR) and the Center for International Blood and Marrow Transplant Research (CIBMTR), more than 250 patients have received autologous stem cell transplants for the treatment of refractory MS. Current trials for the most part employ a non-ablative hematopoietic stem cell transplant regimen, and enrolment criteria of these modern trials have focused on younger patients who have yet to reach advanced disability, and have not required failure of multiple agents. These choices are likely contributory to the reduced morbidity, mortality and toxicity in present trials. Atkins et al recently published the results and pearls learned from over 600 cases of transplant in MS in the literature supporting these lesions.<sup>34</sup> And in 2016, Atkins et al published the results of their landmark autoHCT trial using busulfan, revealing that no patient has had any evidence of inflammatory disease activity (relapse, gadolinium (gd) enhancing lesions) since transplant<sup>35</sup>. Unfortunately, no trial have not reliably shown a halting of or reversal of disability from neurodegeneration, hence conventional progressive patients are likely to incur all the toxicity and none of the benefit of such treatment. The role of mesenchymal stem cells in transplant is still under study.

## MS Treatment

### First-Line Management of Relapsing-Remitting Multiple Sclerosis

- Interferon beta-1 alpha (Rebif®, Avonex®, Betaseron®, Extavia®)
- Glatiramer acetate (Copaxone®)
- Dimethyl Fumarate (Tecfidera®)
- Teriflunomide (Aubagio®)
- Ocrelizumab (Ocrevus®)
- Ofatumumab (Kesimpta®)

### First-Line Management of Aggressive Inflammatory Pseudo-progression in Multiple Sclerosis

- definition of aggressive inflammatory pseudo-progression:
  - very large EDSS change/major changes on neurological exam in motor/brainstem/cerebellar categories. Typically patients move from fully ambulatory to significant limitation in ambulation in < 12 months with coincident gadolinium activity on MRI and objective exam improvement after trial of high dose steroids and ≤ 45 years of age
- no approved therapy, no consensus
- typically used agents include Mitoxantrone (Novantrone®), Cyclophosphamide (Cytosan®)

## Definition of Failure of First-Line Agents for Escalation Therapy

- relapse activity unchanged or worsened despite first-line agent
- a combination of mild-moderate relapse activity and new MRI (new T2/FLAIR and/or gadolinium (gd) enhancing lesions) activity with first-line agent
- rapid progression in absence of distinct relapse events as described above

## Current Escalation Management of Relapsing-Remitting Multiple Sclerosis in Treatment Failure

In patients with evidence of failure, conventionally a switch to a **second-line** option includes:

- Fingolimod (Gilenya®)\*
- Dimethyl Fumarate (Tecfidera®)\*
- Natalizumab for a finite period of time (Tysabri®)\*
- Alemtuzumab (Lemtrada®)\*
- Ocrelizumab (Ocrevus®)\*\*
- Ofatumumab (Kesimpta®)
- Cladribine (Mavenclad®)\*

*\*only approved and covered for use in relapsing patients<sup>36</sup>*

*\*\*approved for both relapsing remitting MS and primary progressive MS in a special cohort*

Escalation treatment options in MS depend on the nature and severity of failure on first-line agents and associated comorbidities and pregnancy planning and other issues.

## Risk factors for poor outcomes on first-line agents include:

- Incomplete recovery from relapses
- High relapse frequency in first 2-5 years from onset, short interval between initial relapses
- Reaching high EDSS in the first five years of disease (EDSS >3)
- Ongoing accumulation of T2/gd lesions, brain atrophy and other measures of neurodegeneration

## Definitions of treatment failure in MS (modified from CanTOR guidelines 2020<sup>37</sup>):

### Mild Failure

- Relapse rate may be better than prior to DMT, but still active (annualized relapse rate or annualized relapse rate (ARR) ~ 0.5-1) and coupled with mild activity on MRI (new T2/gd lesions)
- Near complete recovery from relapses

### Moderate Failure

- Relapse rate unchanged from previous or worsening

- Incomplete relapse recovery with fixed FSS changes > 1 in motor/cerebellar/brainstem/sphincter/sensory domains, but EDSS still < 6.0

**OR**

- Milder relapse breakthrough but coupled with active MRI (T2/gd lesions)

#### Severe Failure

- Highly active relapse rate (ARR =>2)
- Marked residual disability from relapses, at least 0.5 point change in EDSS if 5.5 or => 2 point if EDSS ≤4.0
- Above coupled with active MRI (new T2/Gd lesions)

**OR**

- Rapid and severe progression in apparent absence of relapses in relatively young patient coupled with active MRI (gd lesions), but exam improved with trial of high dose steroids (suggesting inflammatory-based progression)

**Note that transition to classic progressive disease is not currently considered “treatment failure”. This may change in the coming years.**

## Selection Criteria For Autologous Hematopoietic Stem Cell Transplant In MS:

### Inclusion Criteria

- MS by current McDonald criteria
- Age ≤ 45
- EDSS ≤ 6.0 based on observed ambulation assessment
- If EDSS = 6.0, it cannot be for a period > 12 months
- *Failure to respond to standard MS DMT or pseudoprogession (defined below).*
- Patients must be confirmed eligible after consultation with an MS neurologist with knowledge on AHSCT and escalation therapy
- All patients require approval of an MS neurologist with knowledge on AHSCT and escalation therapy and transplant hematologist. In the event of disagreement, an additional opinion will be sought
- Patients meet “failure” as per options 1 or 2 listed directly below
- *Failure to respond to standard MS DMT is defined as:*  
While adherent to a second-line DMT\*:
  - One severe relapse or ≥ 2 moderate relapses in past 12 months regardless of MRI activity

**OR**

  - While adherent to a second-line DMT:  
One or more moderate/severe relapses in past 12 months **AND**



- MRI evidence of new inflammatory disease within the same 12 month time period (characterized by  $\geq 1$  gadolinium enhancing lesions and/ or  $> 2$  new T2 lesions).

\*Patients may be eligible at the discretion of the neurology and transplant team members if they have demonstrated evidence of severe indicators of failure on a first-line DMT (excluding interferons and glatiramer acetate) characterized by such features as high ARR, poor relapse-recovery and rapidly advancing EDSS secondary to relapses

*Special Circumstances for Users of Natalizumab<sup>\*\*\*</sup>, Ocrelizumab, Ofatumumab, or Alemtuzumab*

- While fully adherent to a minimum of 12 months on Natalizumab or Ocrelizumab, or after two annual cycles of Alemtuzumab:
    - One moderate relapse AND MRI evidence of new inflammatory disease within the same 12 month time period in the form of any new gadolinium enhancing lesions or  $> 2$  new T2 lesions
- OR**
- $\geq 2$  mild/moderate relapses over a 12 month period regardless of MRI activity
  - If the patient has to stop Natalizumab or Alemtuzumab for adverse event-related reasons, the pre-treatment disease activity profile will be used to determine eligibility

<sup>\*\*\*</sup> Natalizumab before HCT poses the following problems: 1. MS flare due to immune reconstitution could occur if the interval between natalizumab discontinuation and stem cell mobilization was too long. This may not apply to patients whose MS is not controlled by the natalizumab. 2. JCV could cause PML if the interval between natalizumab discontinuation and stem cell mobilization was too short. This may be mitigated by checking JCV by PCR in CSF and proceeding with stem cell mobilization only if negative. 3. Natalizumab increases the number of CD34 cells in blood and marrow and changes the expression of some antigens on the CD34 cells in vivo and changes their function in vitro (eg, chemokine-induced migration).<sup>45</sup> It is unknown whether this persists after natalizumab discontinuation and, if yes, whether it has any impact on stem cell mobilization and HCT outcome. Until more knowledge has been obtained, our standard practice is to aim for the interval between the last dose of natalizumab and the infusion of cyclophosphamide for stem cell mobilization of 2-3 months, if JCV PCR in CSF is negative. If the interval needs to be longer because of JCV PCR positivity or for other reasons, bridging immunosuppressive therapy can be considered (eg, cyclophosphamide 750 mg/m<sup>2</sup> iv monthly or rituximab 1 g iv once or twice 2 weeks apart).

- *Progression due to very active inflammatory disease (pseudoprogression):*
  - Rapid decline ( $< 12$  months) in EDSS (2 or more EDSS points within 12 months if EDSS  $< 5.0$  or 1 or more EDSS points if EDSS  $\geq 5.0$ ) with a cerebellar, brainstem, or pyramidal functional score of at least 3 points and impaired ambulation **AND**
  - MRI demonstrating two or more gadolinium enhancing lesions **AND**
  - Objective improvement in neurological exam with improvement in EDSS after trial of high dose steroids (as objectively determined by an MS neurologist)

## Exclusion Criteria

- DMT failure in context of poor compliance/adherence (confirmation of dispensing by pharmacy is required)
- >2 courses of cladribine is a relative contraindication (concern of poor stem cell mobilization)
- Indwelling urinary catheter during the peri-transplant period (patients could make arrangements for intermittent catheterization during the high-risk period)
- Pregnancy, inability or unwillingness to use appropriate contraception
- Inability to provide informed consent for treatment
- Previous malignancy with the exception of non-melanoma skin cancer or carcinoma in situ.
- Active infection or significant organ dysfunction.
- In patients at risk, CD4 T cell count <100/microliter (HIV infection per se is not an exclusion).
- History of congenital immune deficiency
- Myelodysplasia/leukemia (marrow aspiration is required on all patients with CBC abnormality that could be due to myelodysplasia/leukemia and on all patients with history of myelotoxic drugs).
- Absence of support/caregiver during 4 months peri-transplant
- Inability to reside within the city of Calgary in the 30 days prior to and 100 days following transplant
- Natalizumab or another anti-lymphocyte antibody should ideally be discontinued 2 months before stem cell mobilization chemotherapy.

## General Note Regarding Selection Criteria

Patients most likely to benefit from autoHCT include those of relatively younger age, with relatively short disease duration, a relapsing form of MS (although cases of disease inactivity/stabilization after autoHCT in patients with progressive MS have been described, this appears rare and thus not proposed here), accumulating disability but still ambulatory, and ongoing disease activity despite DMT.

## HCT Details

**Stem cell mobilization** is achieved with cyclophosphamide, filgrastim and dexamethasone. Cyclophosphamide, 2500 mg/m<sup>2</sup> IV over 1 h, is given in BMT clinic. Antiemetics and hydration are given per our standard practice; Mesna, 2500 mg/m<sup>2</sup> IV, should be given in two to three divided doses, the first one concurrently with cyclophosphamide and the second (third) one 4h (8h) later. Filgrastim is started on day 7 and continued until apheresis per our standard practice (see chapter “Donor Management, Including Mobilization”). Dexamethasone, 2 mg QID PO on the days of filgrastim administration, is used to enhance stem cell mobilization and to prevent filgrastim-induced flare of MS activity.

**Apheresis:** The target CD34 cell yield is  $5 \times 10^6/\text{kg}$  ( $\sim 5 \times 10^6/\text{kg}$  after CD34 cell enrichment). The minimum CD34 cell yield is  $2 \times 10^6/\text{kg}$  ( $\sim 3 \times 10^6/\text{kg}$  after CD34 cell enrichment).

**Graft processing:** Both unmanipulated and CD34 cell-enriched grafts have been used. It is currently not known whether CD34 cell enrichment is necessary, but probably not (Table 1). EBMT currently recommends no CD34 selection, except in the context of a clinical trial<sup>50</sup>, so we will follow that recommendation.

**Conditioning:** Several different regimens have been used (Table 1). Between 2014 and 2023 we used busulfan + cyclophosphamide + ATG (Ottawa regimen) (Appendix D). In 2024, we transitioned to cyclophosphamide + ATG. The reasons were:

- Multiple studies published in 2019-2024 showed similar efficacy to the Ottawa regimen (Table 1)
- The Ottawa regimen may be associated with a risk of potentially fatal liver veno-occlusive disease / sinusoidal obstruction syndrome (Atkins et al<sup>35</sup> and our experience in one patient (Appendix B)).

To avoid an unnecessarily large number of conditioning regimens that differ in minor details, we use the same regimen for MS as well as for systemic sclerosis (see chapter on Systemic Sclerosis, also Table 2 pasted here). Most of the centers that reported outcomes of the cyclophosphamide + ATG regimen used a slightly higher dose of ATG than our systemic sclerosis regimen (6 vs 4.5 mg/kg). However, this is probably negligible.

**Table 1.** Results of recent studies with ≥20 HSCT patients, at least 50% of whom had RRMS, and with NEDA\*\* reported.

|                             | Burt<br>(US)<br>2009 <sup>38</sup> | Atkins<br>(Canada)<br>2016 <sup>35</sup> | Nash<br>(US)<br>2017 <sup>46</sup> | Moore<br>(Austr)<br>2018 <sup>47</sup> | Burt<br>(US)<br>2019 <sup>39</sup> | Jespersen<br>(Denmark)<br>2023 <sup>49</sup> | Silfverberg<br>(Sweden)<br>2024 <sup>48</sup> | Vaisvilas<br>(Lithuania)<br>2024 <sup>51</sup> | Kvistad<br>(Norway)<br>2024 <sup>52</sup> | Braun<br>(Germany)<br>2024 <sup>53</sup> |
|-----------------------------|------------------------------------|------------------------------------------|------------------------------------|----------------------------------------|------------------------------------|----------------------------------------------|-----------------------------------------------|------------------------------------------------|-------------------------------------------|------------------------------------------|
| No. of patients             | 21                                 | 24                                       | 24                                 | 35                                     | 52                                 | 32                                           | 174                                           | 31                                             | 29                                        | 20                                       |
| % RRMS                      | 100%                               | 50%                                      | 100%                               | 57%                                    | 100%                               | 100%                                         | 100%                                          | 100%                                           | 100%                                      | 76%                                      |
| Age (median)                | 33 y                               | 34 y                                     | 36 y                               | 37 y                                   | 35.6 y                             | ~40 y                                        | 31 y                                          | 38 y                                           | 31 y (mean)                               | 35 y (mean)                              |
| EDSS (median)               | 3.1                                | 3.0 – 6.0                                | 6.0                                | 6.0                                    | 3.4                                | ~4.0                                         | 3.5                                           | 6.0                                            | 2.9 (mean)                                | 5.0 (mean)                               |
| Duration of MS (y, med)     | 5 y                                | 6.5 y                                    | 10 y                               | 7 y                                    | 5 y                                | 4 y                                          | 3 y                                           | 8 y                                            | 5 y (mean)                                | 6 y (mean)                               |
| Mobilization                | Cy + GCSF                          | Cy + GCSF                                | GCSF + Pred                        | Cy + GCSF                              | Cy + GCSF                          | Cy + GCSF                                    | Cy + GCSF                                     | Cy + GCSF                                      | Cy + GCSF                                 | Cy + GCSF                                |
| CD34 selection              | No                                 | Yes                                      | Yes                                | No                                     | No                                 | No                                           | No                                            | No?                                            | No                                        | No                                       |
| Conditioning                | Cy 200 + Alem 20 mg                | Bu + Cy 200 + rATG 5 mg/kg               | BEAM + rATG 5 mg/kg                | BEAM + hATG 40mg/kg                    | Cy 200 + rATG 6 mg/kg              | Most Cy 200 + rATG 6 mg/kg                   | Most Cy 200 + rATG 6 mg/kg                    | Cy 200 + rATG 6.5 mg/kg                        | Cy 200 + rATG 6 mg/kg                     | BEAM + rATG 3.75mg/kg                    |
| Follow up (y)               | 3                                  | 7                                        | 5                                  | 3                                      | 5                                  | 3-4                                          | 5.5                                           | 4                                              | 6                                         | 9                                        |
| TRM (by 1 y)                | 0%                                 | 4% (1 pt)                                | 0%                                 | 0%                                     | 0%                                 | 0%                                           | 0%                                            | 0%                                             | 0%                                        | 0%                                       |
| EDSS trend                  | Impr                               | Stabiliz                                 | Impr                               | Impr                                   | Impr                               | Stabiliz?                                    | Impr                                          | Impr                                           | Stabiliz                                  | Stabiliz                                 |
| % pts with clinical relapse | 24%                                | 0%                                       | 13%                                | ?                                      | 2% y1<br>8% y1-2<br>15% y1-5       | 23%                                          | 7%                                            | 29% at 2 y                                     | 17%                                       | 0% at 5 y                                |
| % pts with MRI progr        | 14%                                | 0%                                       | 9%                                 | ?                                      | ?                                  | 7%                                           | 9%                                            | 10% at 2 y                                     | 21%                                       | 0% at 5 y                                |
| Progression-free survival*  | 77%                                | 70%                                      | 91%                                | ?                                      | 94%                                | ?                                            | 97%? (5 pts progressed)                       | ?                                              | 90%                                       | ~75% at 5y                               |
| Survival with NEDA**        | 62%                                | 70%                                      | 69%                                | 60%                                    | 78.5%                              | 69%                                          | 73% at 5 y,<br>65% at 10 y                    | 79% at 2 y,<br>54% at 5 7                      | 69%                                       | 75% at 5 y,<br>55% at 10 y               |

**Abbreviations:** RRMS = relapsing remitting multiple sclerosis, rATG = rabbit ATG (Thymoglobulin), hATG = horse ATG, Alem = alemtuzumab, TBI = total body irradiation, Cy = cyclophosphamide, Bu = busulfan, Pred = prednisone, BM = busulfan + melphalan, BEAM = BCNU + etoposide + AraC + melphalan, TRM = transplant related mortality.

\* Survival free of EDSS progression

\*\* No Evidence of Disease Activity (NEDA), i.e., no EDSS progression, no clinical relapse, and no MRI activity

\*\*\* Prednisone 1 mg/kg/d x 10 days beginning one day before starting GCSF

**Table 2.** Transplant Conditioning/Infusion Regimen used in Calgary (since 2024):

| Day                 | -6       | -5       | -4       | -3        | -2        | -1        | 0    |
|---------------------|----------|----------|----------|-----------|-----------|-----------|------|
| Cy*                 | 50 mg/kg | 50 mg/kg | 50 mg/kg | 50 mg/kg  |           |           |      |
| Rabbit ATG**        |          |          |          | 0.5 mg/kg | 2.0 mg/kg | 2.0 mg/kg |      |
| Methyl-prednisolone |          |          |          | 40 mg bid | 40 mg bid | 40 mg bid | X*** |
| Stem cell infusion  |          |          |          |           |           |           | X    |

\* Cyclophosphamide (50 mg/kg ideal body weight in 250 mL D<sub>5</sub>W infused over 2 h) with Mesna (50 mg/kg ideal body weight in 1 L NS over 24 h starting with each Cy dose), hydration (NS at 75 mL/h starting the night before the first Cy dose and continuing till 24 h post the last Cy dose) and antiemetics (ondansetron + dexamethasone [except on day -3, when methylprednisolone is given as ATG premedication] + aprepitant + prn dimenhydrinate + prn metoclopramide)

\*\* Thymoglobulin (0.5-2.0 mg/kg in as low volume of NS as possible [0.5 mg/mL] infused over ≥4 h) with premedication (Methylprednisolone 1 mg/kg before each infusion + acetaminophen + diphenhydramine + meperidine prn)

\*\*\* Prednisone 0.25 mg/kg/d or equivalent methylprednisolone dose is given from day 0 to day 21, then tapered by day 37. This is to minimize the likelihood of fever (due to neutropenia, engraftment syndrome, or ATG-induced serum sickness) and its negative effect on neurological status.

**Infection prophylaxis** posttransplant is more stringent than after autologous transplantation for hematologic malignancies. Anti-bacterial and fungal prophylaxis early posttransplant is given to avoid neutropenic fever, which could result in the worsening of neurological status. CMV and EBV monitoring and preemptive therapy is given because of severe lymphopenia produced by CD34 enrichment of the graft and by ATG. Specific measures:

- Valacyclovir 500 mg qd until VZV vaccination per our Standard Practice (see chapters “CMV/HSV/VZV/HHV6” and “Vaccination”)
- CMV and EBV PCR weekly from ~day 7 until 3 months posttransplant, and preemptive valganciclovir or rituximab per our Standard Practice (see chapters “CMV/HSV/VZV/HHV6” and “EBV/PTLD”)
- Levofloxacin 500 mg qd po or iv during neutropenia
- Fluconazole 400 mg qd po or iv from day 0 until 1 month posttransplant
- Pneumocystis/pneumococcal prophylaxis ideally with trimethoprim-sulfamethoxazole (80/400 mg qd po) from neutrophil engraftment until 12-24 months posttransplant per our Standard Practice (see chapter “Bacterial and Pneumocystis Prophylaxis”)
- Vaccinations per our Standard Practice (see chapter “Vaccination”)

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## Appendix A: Schedule of Tests and Evaluations

|                                                               | Baseline/<br>Eligibility | Transplant Regimen    |                       | Post-Transplant<br>Haematology<br>Monitoring |   |   |    |    | Post-Transplant Neurological Monitoring |    |    |                 |     |                  |     |     |     |     |
|---------------------------------------------------------------|--------------------------|-----------------------|-----------------------|----------------------------------------------|---|---|----|----|-----------------------------------------|----|----|-----------------|-----|------------------|-----|-----|-----|-----|
|                                                               |                          | Mobilization<br>start | Conditioning<br>start | 4                                            | 6 | 8 | 10 | 12 | 26                                      | 52 | 78 | 104             | 130 | 156              | 182 | 208 | 234 | 260 |
| Week                                                          | ~ -12                    | ~ -6                  | ~ -1                  |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| Medical History                                               | X                        |                       |                       | X                                            | X | X | X  | X  | X                                       | X  | X  | X               | X   | X                | X   | X   | X   | X   |
| Physical Exam                                                 | X                        |                       |                       | X                                            | X | X | X  | X  | X                                       | X  | X  | X               | X   | X                | X   | X   | X   | X   |
| EDSS Exam*                                                    | X                        |                       |                       |                                              |   |   |    |    | X                                       | X  | X  | X               | X   | X                | X   | X   | X   | X   |
| CBC                                                           | X                        |                       |                       | X                                            | X | X | X  | X  | X                                       | X  | X  | X               | X   | X                | X   | X   | X   | X   |
| Chemistry panel                                               | X                        |                       |                       | X                                            | X | X | X  | X  | X                                       | X  | X  | X               | X   | X                | X   | X   | X   | X   |
| PT/PTT                                                        | X                        |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| Pregnancy test                                                | X                        |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| PFTs                                                          | X                        |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| MUGA or<br>Echocardiogram                                     | X                        |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| CXR, EKG                                                      | X                        |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| Urinalysis                                                    | X                        |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| TSH                                                           | X                        |                       |                       |                                              |   |   |    |    |                                         | X  |    | X               |     | X                |     | X   |     | X   |
| Ig levels for tetanus,<br>hepatitis B, measles<br>and rubella |                          |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     | X                |     |     |     |     |
| Vaccinations                                                  | X                        |                       |                       |                                              |   |   |    |    | X <sup>#</sup>                          |    |    | X <sup>##</sup> |     | X <sup>###</sup> |     |     |     |     |
| HIV1 and HIV2                                                 | X                        |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| HSV/VZV/CMV/EBV****                                           | X                        |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| Hepatitis A/B/C<br>serology                                   | X                        |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| Dental Consult                                                | X                        |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| MRI brain +/- spinal<br>cord #####                            | X                        |                       |                       |                                              |   |   |    |    |                                         | X  |    | X               |     | X                |     | X   |     | X   |
| Fertility consult                                             | X <sup>**</sup>          |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |
| Bone marrow biopsy                                            | X <sup>***</sup>         |                       |                       |                                              |   |   |    |    |                                         |    |    |                 |     |                  |     |     |     |     |

\* EDSS = Extended disability status scale (0-10), done by a neurologist collaborating with the Calgary-based Alberta Blood and Marrow Transplant Program. During late posttransplant period, the EDSS may be estimated during a virtual visit or obtained from referring neurologist.

\*\*Male patients will be offered sperm banking, female patients will be offered fertility clinic consult.

\*\*\*Only if blood cell counts are abnormal.

\*\*\*\* Pretransplant, HSV, VZV, CMV and EBV IgG should be done once. Posttransplant, CMV and EBV PCR should be done weekly until 12 weeks.

# Referral to Public Health for non-live vaccines.

## Referral to Public Health for live vaccines.

### Referral to Public Health for boosters if specific Ig levels for vaccine-preventable diseases are low.

##### Arranged by Neurology

## Appendix B: Calgary Experience as of January 2024

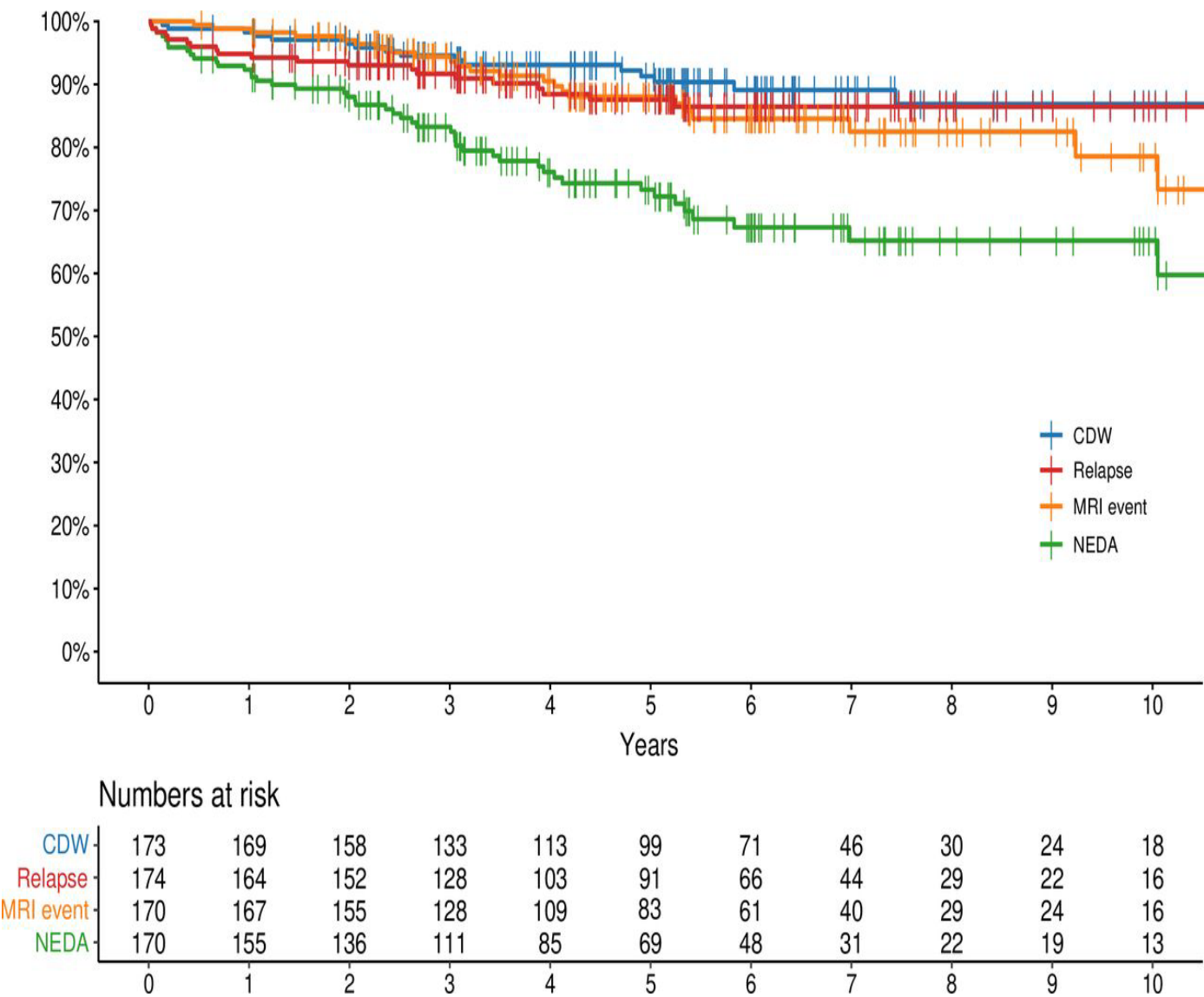
| UPN  | Year of auto HCT | EDSS pre HCT** | Mobilization | CD34 selection | Conditioning | Alive? | MS activity since HCT?* | Comment                              |
|------|------------------|----------------|--------------|----------------|--------------|--------|-------------------------|--------------------------------------|
| 986  | 2009             |                | Cy+GCSF      | No             | Cy+Thymo     | Y      | Y since ~2011           | Was on the verge of RRMS→SPMS at HCT |
| 1052 | 2010             |                | Cy+GCSF      | No             | Cy+Thymo     | Y      | Y since 2016            |                                      |
| 1355 | 2014             |                | Cy+GCSF      | Yes            | Bu+Cy+Thymo  | Y      | Y since 2017            | Was on the verge of RRMS→SPMS at HCT |
| 1604 | 2016             | 2.5            | Cy+GCSF      | Yes            | Bu+Cy+Thymo  | Y      | N                       |                                      |
| 1616 | 2017             | 4.5            | Cy+GCSF      | Yes            | Bu+Cy+Thymo  | Y      | N                       |                                      |
| 1842 | 2019             | 3.5            | Cy+GCSF      | Yes            | Bu+Cy+Thymo  | Y      | N                       |                                      |
| 1913 | 2020             | 2.5            | Cy+GCSF      | Yes            | Bu+Cy+Thymo  | Y      | N                       |                                      |
| 2166 | 2022             | 2.5            | Cy+GCSF      | Yes            | Bu+Cy+Thymo  | Y      | N                       |                                      |
| 2237 | 2023             | 2.5            | Cy+GCSF      | Yes            | Bu+Cy+Thymo  | Y      | N                       |                                      |
| 2300 | 2023             | 1.5            | Cy+GCSF      | Yes            | Bu+Cy+Thymo  | Y      | N                       |                                      |
| 2317 | 2023             | 3.0            | Cy+GCSF      | Yes            | Bu+Cy+Thymo  | Y      | N                       | Moderate VOD/SOS***                  |

\* Relapse, progression, or new or enhancing MRI lesions

\*\* EDSS (expanded disability status scale, 0-10, a higher score denotes greater disability)

\*\*\* Liver veno-occlusive disease/sinusoidal obstruction syndrome

Appendix C: Results of the largest retrospective study



Kaplan-Meier curves for the primary endpoint of no evidence of disease activity (NEDA), and for the secondary endpoints of freedom from MRI events, freedom from clinical relapse, and freedom from confirmed disability worsening (CDW), in the Swedish registry study (Silfverberg 2024).<sup>48</sup> This study included only patients with RRMS. There was no transplant-related mortality, but one patient died at >6 years of suicide related to substance abuse.

## Appendix D: Ottawa conditioning, used in Calgary 2014-2023, then retired.

| Day                                          | -10 | -9 | -8 | -7 | -6 | -5 | -4 | -3  | -2  | -1  | 0 | +7 |
|----------------------------------------------|-----|----|----|----|----|----|----|-----|-----|-----|---|----|
| Busulfan* ~2.4 mg/kg/day IV                  | X   | X  | X  | X  |    |    |    |     |     |     |   |    |
| Lorazepam 1 mg QID PO (seizure prophylaxis)  | X   | X  | X  | X  | X  |    |    |     |     |     |   |    |
| Hydration**                                  |     |    |    |    | X  | X  | X  | X   | X   | X   |   |    |
| Cyclophosphamide**<br>50 mg/kg/day IV        |     |    |    |    |    | X  | X  | X   | X   |     |   |    |
| MESNA continuous infusion<br>50 mg/kg/day IV |     |    |    |    |    | X  | X  | X   | X   |     |   |    |
| ATG*** (Thymoglobuline)<br>(mg/kg/day)       |     |    |    |    |    |    |    | 0.5 | 2.0 | 2.0 |   |    |
| Methyl-prednisolone****                      |     |    |    |    |    |    |    | X   | X   | X   | X | X  |
| Stem cell infusion                           |     |    |    |    |    |    |    |     |     |     | X |    |
| GCSF ~0.5 ug/kg/d from d7 till<br>ANC>1/nl   |     |    |    |    |    |    |    |     |     |     |   | X  |

\* **Busulfan dosing is PK-adjusted.** First dose is 2.4 mg/kg IV at a constant rate of 80 mg/hr (160 ml/hr for busulfan at 0.5 mg/ml concentration). Blood (4 ml green top (heparinized) tube) for busulfan PK is collected at the end of the infusion and at 1, 3, 5 and 7 h after the end of the infusion. Subsequent doses are adjusted to target overall busulfan AUC of <16000 µmol.min/L over four days. The last dose of busulfan should be given in the morning of day -7 to ensure >>24 h interval between busulfan and cyclophosphamide infusions.

\*\* Cyclophosphamide 50 mg/kg/day is given IV over 1 hour in 500 cc of normal saline. If actual weight is < ideal weight, cyclophosphamide is given based on actual weight. If actual weight is > ideal weight, cyclophosphamide is given as adjusted weight. Adjusted weight = ideal weight + 0.25 x (actual weight minus ideal weight). Anti-emetics, as pre-medications for Cyclophosphamide, should be given per institutional policy and medical judgement. Hydration with Normal Saline, approximately 2 liters/m<sup>2</sup>/day, should be started on day -6, and at least 6 hours before cyclophosphamide and continued until 24 hours after the last cyclophosphamide dose.

\*\*\* ATG (Thymoglobulin) 0.5 mg/kg is given IV on day -3 and 2.0 mg/kg IV on days -2 and -1 (no dose adjustment), over 4-6 hours each day. Pre-medicate with methylprednisolone 1.0 gram IV, acetaminophen 650 mg po and diphenhydramine 25 mg IV or PO 30 minutes before infusion. An in-line 0.22 µm filter should be used for ATG administration.

\*\*\*\* Methylprednisolone or prednisone is given to minimize the likelihood of fever (due to ATG, neutropenia, or engraftment syndrome) and its negative effect on neurological status, according to the following schedule:

Day -3 to -1, 1 g IV as premedication for ATG

Day 0 to 3, 0.5 mg/kg/d,

Day 4 to 7, 0.4 mg/kg/d,

Day 8 to 11, 0.3 mg/kg/d,

Day 12 to 15, 0.2 mg/kg/d,

Day 16 to 19, 0.1 mg/kg/d, then discontinue



# Autologous Hematopoietic Cell Transplant in Systemic Sclerosis (SSc): Alberta Protocol

Presented by: Jan Storek / Caylib Durand / Mo Osman

## Summary

- Autologous HCT (HCT) for SSc is indicated if:
  - Age <65, ideally younger
  - <5 years from the first non-Raynaud symptom, ideally <1 year
  - Severe skin involvement (mRSS >20), or
  - Moderate skin involvement (mRSS 15-20) with mild to moderate Interstitial lung disease (FVC/DLCO 40-80%, ideally 60-80%. No hypoxia)
  - No moderate/severe PAH (mean PAP <30 mmHg by right heart catheterization)
  - No or minimal heart involvement (no significant cardiac fibrosis, no heart failure, myocarditis/pericarditis must be in remission).
  - If GAVE, needs to be successfully treated before HCT
- Allogeneic HCT should be considered only in patients with concurrent hematologic disease or under a clinical trial.
- CAR T cells only under a clinical trial.

## Autologous HCT Indications/Contraindications by SSc Manifestation

- **Skin involvement**
  - Thickening (as defined as mRSS. See Appendix 1 Figure 2)
    - *Localized* cutaneous systemic sclerosis (morphea)
      - Not an indication for HCT due to good prognosis with standard therapies.
    - *Limited* cutaneous systemic sclerosis (hands/distal forearms (not extending beyond elbows and no truncal involvement)/face) / CREST syndrome (calcinosis of skin, Raynaud's phenomenon, esophageal dysmotility, sclerodactyly, telangiectasia)
      - Associated with anti-centromere antibody (ACA) (60%)
      - Not an indication for HCT at present due to better prognosis compared to diffuse cutaneous systemic sclerosis without HCT, and minimal data on HCT. However, may consider HCT if interstitial lung disease.
    - *Diffuse* cutaneous systemic sclerosis (dcSSc, involves also proximal skin) – clearcut indication for HCT if otherwise meets criteria
      - Associated with Scl-70 antibody (60%)
      - Indicated for HCT if moderate to severe (mRSS >20) or if associated with lung disease
  - Other skin manifestations (alone not an indication for HCT)

- Edema (early)
- Contractures (late)
- Pruritus
- Hyper/hypopigmentation (“salt-and-pepper”)
- Loss of appendicular hair
- Ulcers
- Calcinosis

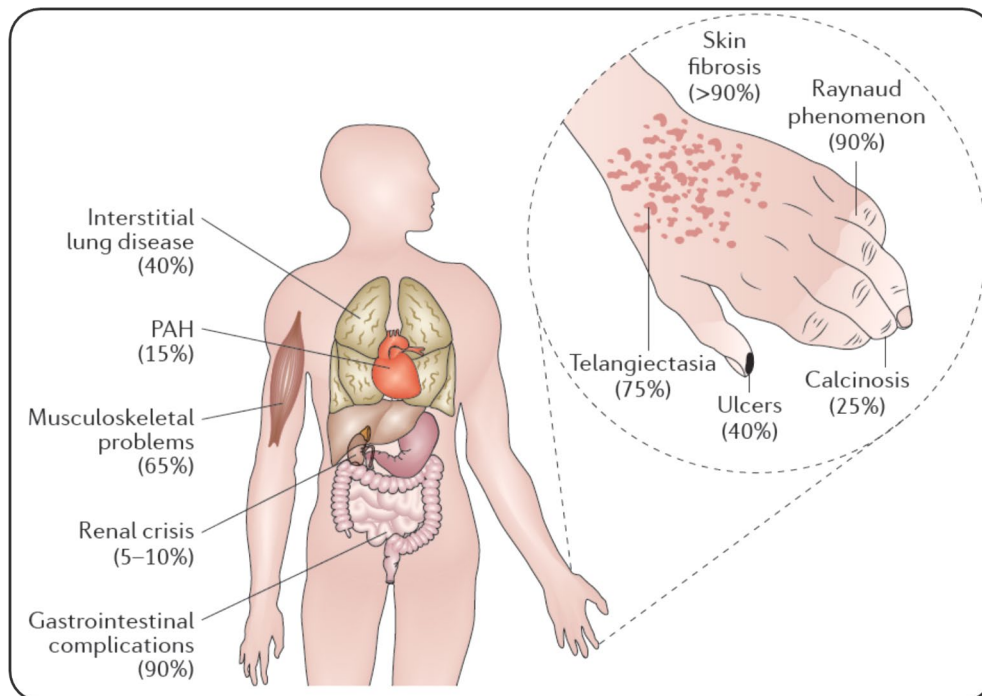


Figure 1. Manifestations of SSc (from A.Yannick et al: Nat Rev Dis Prim 2015)

## • Lung involvement

- Interstitial lung disease / fibrosing alveolitis
  - Indicated for HCT, particularly if rapidly progressing, but FVC and DLCO must be >40% predicted, ideally >60% predicted
- Pulmonary artery hypertension is a contraindication to HCT
  - Absolute: RVSP >55 mmHg by echo or resting mean PAP >30 mmHg by RHC
  - Relative: RVSP 40-55 mmHg by echo or resting mean PAP 25-30 mmHg by RHC, or significant increase in PAP with fluid/exercise
- Hypoxia requiring O<sub>2</sub> is a contraindication to HCT
- Lung cancer (5-fold higher incidence compared to general population)
  - Contraindication to HCT
- Smoking
  - Current smoking is a contraindication to HCT

- **Renal involvement**

- Renal crisis is a contraindication to HCT
  - Renal crisis is characterized by hypertension, decreased GFR, non-heavy proteinuria, thrombotic microangiopathy
- Renal failure due to any cause is a contraindication to HCT
- Hypertension must be controlled before HCT considered, especially if due to renal crisis

- **Heart involvement**

- Myocarditis and or cardiac fibrosis; myocardial ischemia; pericarditis/effusion
- Absolute contraindications to HCT<sup>1,2</sup>:
  - LVEF <40%
  - D-sign or septal bounce on echo or MRI (sign of RV overload/failure or constrictive pericarditis)
  - Non-revascularized severe coronary artery disease
  - Uncontrolled severe arrhythmia
  - Pericardial Tamponade
  - Constrictive pericarditis
- Relative contraindications to HCT:
  - LVEF 40-50%
  - Tricuspid annular plane systolic excursion (TAPSE) <18 mm on echo
  - Any sign of heart involvement with systemic sclerosis on MRI
  - No increase in cardiac output on RHC with exercise/fluid
- The above cardiac contraindications may be less important with non-cardiotoxic conditioning<sup>3,4</sup>. However, long-term efficacy of AHCT using non-cardiotoxic conditioning is uncertain.

- **Gastrointestinal involvement**

- Not a contraindication to HCT, unless severe or unless untreated Gastric Antral Venous Ectasia (GAVE)
- Manifestations:
  - Esophagus: Esophageal dysmotility and incompetence of the LES leading to chronic esophagitis, stricture, Barrett's esophagus, pulmonary micro aspiration.
  - Stomach: GAVE ("watermelon stomach") that can lead to gastric ulcers and or anemia.
    - GAVE needs to be successfully treated (e.g., with Argon Plasma Coagulation) before HCT.
  - Colon: Diarrhea or constipation, bacterial overgrowth with malabsorption.
  - Anorectum: Fecal incontinence due to sphincter muscle damage, rectal prolapse.

- **Involvement of other organs (usually has no impact on whether AHCT is indicated):**

- Systemic
  - Fatigue associated with pain similar to that described in patients with idiopathic fibromyalgia

- Weakness, may or may not be associated with elevated CK. Important to assess for myositis overlap with systemic sclerosis (may be associated with worse outcomes)
- Pain that can be directly or indirectly associated with active systemic sclerosis.
- Vascular
  - Raynaud's phenomenon and associated digital ulcers. May also be assessed by nail fold capillaroscopy or nail fold video capillaroscopy.
  - Telangiectasia.
- MSK/Joints
  - Myalgias and arthralgias with AM stiffness, due to inflammatory arthritis in addition to sclerodactyly and skin inflammation. Both can lead to fibrosis around tendons/periarticular soft tissue leading to tendon friction rubs, joint flexion contractures and decreased range of motion, which is more commonly associated with diffuse cutaneous SSc.
  - Inflammatory arthritis can be associated with systemic sclerosis directly but must also have workup for overlap with rheumatoid arthritis (CCP, RF and X rays).
- Neuromuscular
  - Myositis
  - Peripheral neuropathy, including autonomic. Less evidence but might also be associated with medications for the treatment of systemic sclerosis.
  - CNS disease is rare
- Genital
  - Erectile dysfunction, sexual dysfunction is common in both sexes.
  - Dyspareunia due to vaginal dryness/narrow introitus.

## Pathogenesis

- Poorly known but thought to involve 3 main pathways: Inflammation, fibrosis and vasculopathy (Figure 2)
- Immune dysregulation (T and B lymphocytes), endothelial cell and fibroblast abnormalities
- Autoantibodies:
  - Antibodies may contribute to the pathogenesis of systemic sclerosis:
    - Anti-Scl-70 (anti-topoisomerase on fibroblast surface)<sup>5</sup>?
    - Anti-PDGFR with profibrotic activity<sup>6</sup>?
  - Whether autoantibodies persist after AHCT is controversial<sup>7,8</sup>

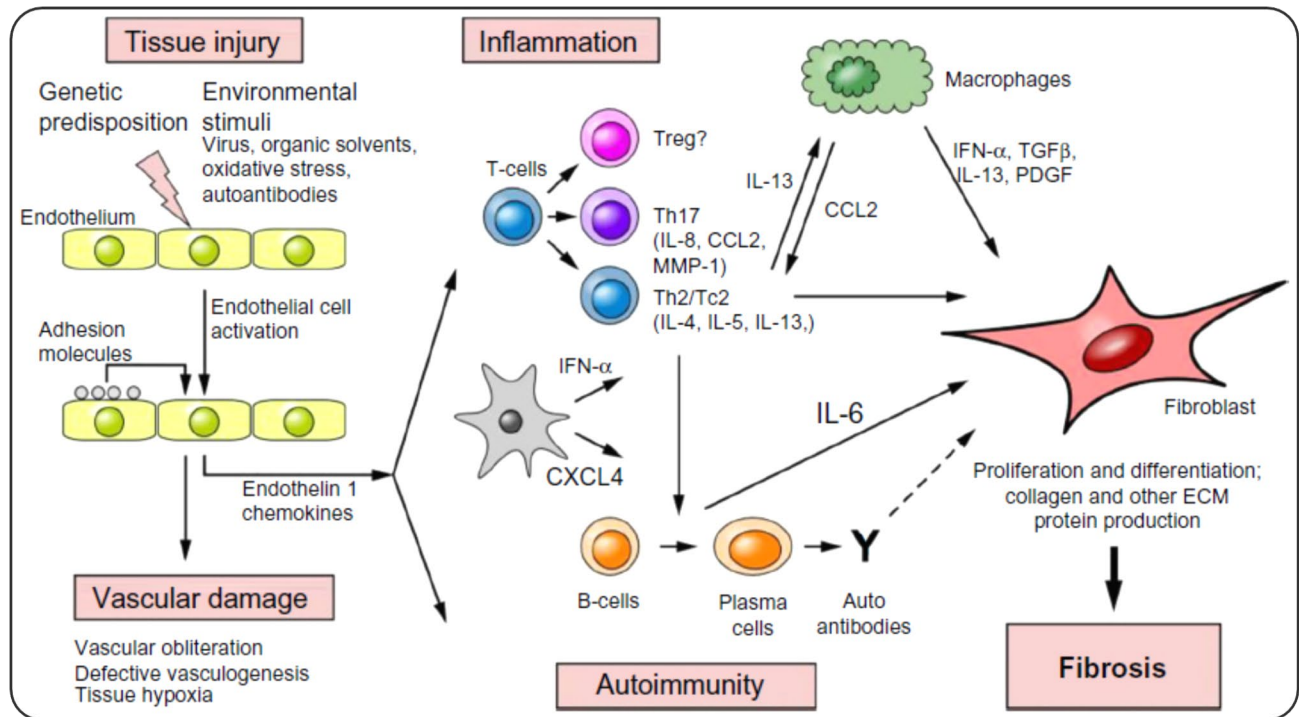


Figure 2: Proposed Pathophysiology of Systemic Sclerosis (from Fuschiotti P: Immunotargets Ther 2016).

## Incidence of SSc

- 0.6 to 122/million/year; Median 12/million/year in North American studies<sup>9</sup>
- Trend toward increasing incidence
- Females > Males, particularly ≥1-para/gravida females
- Peak age 50-60 y

## Prognosis without HCT

- Survival ~80% at 2 years, ~60% at 5 years, ~40% at 10 years per Altman et al<sup>10</sup>; consistent with more recent studies<sup>11,12</sup>
- Survival is particularly low with:
  - Diffuse cutaneous systemic sclerosis<sup>13,14</sup>
  - Heart, lung (ILD or PAH), or renal involvement (SRC)<sup>10</sup>
  - For diffuse cutaneous systemic sclerosis without or with only mild internal organ involvement, rapid Skin Thickness Progression Rate (STPR)<sup>15</sup>
    - Onset of skin thickening defined as the first time the patient's fingers became swollen and never again returned to normal
    - STPR = mRSS / time interval between the onset of skin thickening and the documented mRSS in years

- STPR >45 associated with survival of 76% at 2 y (compared to ~86% with STPR ≤45 (p=.002)
- Not validated for patients with >2 y interval between the onset of skin thickening and the documented mRSS

## SSc Treatments Other than HCT

- Systemic immunosuppressive/antifibrotic/anti-cytokine agents
  - All studies retrospective or non-randomized prospective (thus dubious efficacy), except for cyclophosphamide, which was shown to have limited efficacy in randomized studies<sup>16</sup>, and for MMF, which has efficacy similar to cyclophosphamide<sup>17</sup>.
  - Cyclophosphamide
    - In a randomized study of oral Cy vs placebo for 1 y, the Cy group had a smaller decline of FVC (1% vs 2.6% predicted, p<.03)<sup>18</sup>. There was no difference at 2 years<sup>19</sup>.
  - MMF (mycophenolate mofetil)
    - In a randomized study of MMF vs Cy, MMF was as efficacious as Cy, but Cy was more toxic<sup>17</sup>.
  - Methotrexate
    - Corticosteroid (caveat: at high dose may induce renal crisis)
  - Rituximab (RTX)
    - Several case studies and series demonstrate efficacy of RTX in systemic sclerosis patients. Recently some RCT data particularly in patients with SSc-associated ILD (see RECITAL and EVER-ILD). Particularly for combination therapy.
  - IL-6 inhibitors (Tocilizumab/Actemra)
    - Minimal efficacy for ILD but shows mild improvement in skin fibrosis and particularly useful for overlapping inflammatory arthritis.
  - Antifibrotics (Nintedanib/OFEV)
    - SENSIS trial shows stabilization of FVC and has been approved for SARD-ILD in combination with immunosuppression.
  - Efficacy of established non-AHCT therapies is typically short-term, and long-term efficacy of newer non-AHCT therapies (combination therapies like rituximab+tocilizumab) is unknown.
- Organ/Symptom-based therapies
  - Pruritus – antihistamines, topical steroids, methotrexate.
  - Raynaud / digital ulcers – Ca channel blocker, avoiding cold environment, heated gloves, Raynaud's gel, nitro patches, PDE4 inhibitors like sildenafil.
  - Contractures – physiotherapy, occupational therapy, hot wax baths.



- Renal crisis – ACE inhibitor, good blood pressure control, avoiding systemic steroids especially if RNA Pol3 autoantibodies are present.
  - These patients should never be given any type of steroid because of significant risk of scleroderma renal crisis. However, in case of HCT, we have been giving peri-ATG steroid, recommend with and ACE inhibitor.
- Esophageal dysmotility – proton pump inhibitor, antihistamines like famotidine in combination, gastric coating agents. Medications may need to be compounded because of poor bioavailability and absorption by GI complication of systemic sclerosis.
- Malabsorption/diarrhea due to bacterial overgrowth – antibiotics, dietary fiber.
- ILD – Oxygen, antifibrotics, immunosuppression, lung transplant.
- PAH – oxygen, diuretic, PAP lowering agents (macitentan, riociguat), lung transplantation.
- Arthritis – NSAID, hydroxychloroquine, methotrexate, leflunomide, sulfasalazine. Note that MMF does not work for controlling inflammatory arthritis.
- CHF – ACE inhibitor, implantable cardioverter-defibrillator

## Autologous HCT (HCT)

Multiple non-randomized and 3 randomized studies of HCT for SSc are published (Appendix 1). From these studies it can be surmised that AHCT is superior compared to pre-2015 conventional therapy (e.g., oral or monthly IV cyclophosphamide) for the following indications:

- Indications
  - SSc involving skin + lungs, if FVC or DLCO 40-80% predicted or rapid decline of FVC (>10% over 12-18 months), particularly if patient never smoked, or
  - Systemic sclerosis without lung involvement, if mRSS >20 with high ESR/CRP or rapid skin thickness progression rate
  - Disease duration <5 years, ideally <1 year
    - Insufficient data for patients with longer disease duration<sup>4</sup>. With other autoimmune disease, duration appears to matter<sup>20-23</sup>.
    - Pretreatment with systemic immunosuppressive drugs may not be a contraindication<sup>24</sup>
  - No pulmonary or cardiac contraindication as outlined above (under SSc Manifestations)
  - If GAVE, needs to be successfully treated before HCT
  - Age ≤65 years
    - Progression-free survival is worse with older age<sup>25,26</sup>.
- Benefits
  - Survival benefit
  - Skin improvement (over years; greater improvement proximally than distally)
  - Lung stabilization or slight improvement

- QOL improvement
- Risks
  - Early transplant-related mortality (TRM) (first 5 years) 5-10%
    - Organ failure, particularly heart and lung
    - Infections
  - Late toxicity
    - Myelodysplastic syndrome/Acute myeloid leukemia
    - Solid cancer (increased incidence with SSc alone)
    - Second autoimmune disease (e.g., thyroiditis, immune cytopenia)
- HCT Protocol
  - Optimal drugs/doses for stem cell mobilization and transplant conditioning are unknown. Intermediate intensity conditioning may be optimal<sup>27</sup>, but fludarabine-based low intensity conditioning appears also efficacious<sup>4</sup>. Role of CD34 selection is uncertain<sup>25,28</sup>.
  - Two protocols have been widely used. ASTIS protocol<sup>29</sup>, with conditioning based on cyclophosphamide 200 mg/kg, is typically used in Europe. SCOT protocol<sup>30</sup>, with conditioning based on cyclophosphamide 120 mg/kg + TBI 8 Gy, is typically used in the USA.
  - In Calgary, between 2016 and 2019, we used the ASTIS protocol, except we did not use graft CD34 cell enrichment due to its cost and controversial benefit. In 2020, we switched to the SCOT protocol as it appeared to be associated with lower incidence of SSc relapse/progression, and possibly lower transplant-related mortality and possibly lower incidence of a new autoimmune disease than the ASTIS protocol (ref<sup>30</sup> for SCOT, refs<sup>29,31,32</sup> for ASTIS). In 2021, a CIBMTR retrospective analysis was completed, which did not show a difference in overall survival (OS) or progression-free survival (PFS) between the ASTIS and SCOT-like protocol (Georges G et al: ACR 2021 Meeting, Abstract No. 1364). In summer 2022, a staff shortage substantially impacted our Radiation Physics department. Given that the TBI with lung and kidney shielding, used in the SCOT study, required a lot of staff resources and given the lack of evidence for the potential superiority of the SCOT over the ASTIS protocol, in October 2022 we switched back to the ASTIS protocol, including CD34 enrichment. The rationale for the CD34 enrichment was that by then there had been 4 studies, 3 of which showed at least a trend toward better outcomes with CD34 enrichment – Oliveira's retrospective EBMT study showing no benefit<sup>28</sup>, Georges' retrospective CIBMTR study showing a trend toward improved PFS (Georges G et al: ACR 2021 Meeting, Abstract No. 1364), Ayano's small retrospective study from Japan showing a significantly improved PFS<sup>33</sup>. and Henes' prospective non-interventional EBMT study showing a better SSc response but insignificant impact on PFS<sup>25</sup>. It had also become clear by 2022 that CD34 enrichment together with ATG is likely associated with a high incidence of new autoimmune diseases after HCT and the need for ATG had been questioned

(reviewed in Levin et al<sup>34</sup>). Therefore, the protocol we started to use in October 2022 was a modified ASTIS protocol with a reduced dose of Thymoglobulin (4.5 instead of 7.5 mg/kg).

- A low intensity protocol using fludarabine, low dose cyclophosphamide, rabbit ATG, and rituximab conditioning has been evaluated in a pilot study of 28 patients<sup>4</sup>. Transplant-related mortality at one year was only 4% despite patients were included who would be conventionally considered transplant-ineligible (SSc duration >5 y, DLCO as low as 30%, O<sub>2</sub>-dependent, mPAP >30 mmHg after fluid challenge, interventricular septal flattening/bounce, late Gd enhancement on CMR, pericardial effusion). Short-term efficacy (1 year) appeared to be at least as good as with ASTIS/SCOT regimens as only 1 patient (4%) relapsed by one year, and trends toward improved mRSS and FVC were observed. Long-term efficacy is unknown. We adopted this protocol in 2022 as an alternative for patients for whom the HCT would be relatively contraindicated by conventional criteria.

In Calgary, as of October 2022, we use primarily the modified ASTIS protocol:

## Mobilization

| Day                  | 1                    | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | ≥12 | ≥13 |
|----------------------|----------------------|---|---|---|---|---|---|---|---|----|----|-----|-----|
| Cy*                  | 2.5 g/m <sup>2</sup> |   |   |   |   |   |   |   |   |    |    |     |     |
| GCSF**               |                      |   |   |   |   |   | X | X | X | X  | X  | X   |     |
| Apheresis of MNCs*** |                      |   |   |   |   |   |   |   |   |    |    |     | X   |

\* Cyclophosphamide (2.5 g/m<sup>2</sup> dose dissolved in 500 mL D<sub>5</sub>W and infused over 2 h) with Mesna, hydration (500 mL NS over 1 h before each Cy infusion, and 500-1000 mL NS over 2-4 h after Cy infusion [500 mL over 2 h for <70 kg patient, 1000 mL over 4 h for ≥70 kg patient]) and antiemetics (granisetron + dexamethasone + aprepitant + prn dimenhydrinate + prn metoclopramide + prn prochlorperazine). Mesna, 2500 mg/m<sup>2</sup> IV, should be given in two to three divided doses, the first one concurrently with cyclophosphamide and the second (third) one 4 h (8h) later. The dose of Cy (2.5 g/m<sup>2</sup>) deviates from the ASTIS protocol, which used 4 g/m<sup>2</sup>. This is to keep it standard with our routine mobilization protocol – see SPM chapter on mobilization.

\*\* 300-900 ug per dose depending on weight per SPM chapter on Mobilization; with prn codeine

\*\*\* Start when blood CD34 count is >20 x 10e6/L. Target to collect 8 x 10e6/kg CD34 cells. Perform CD34 selection using CliniMACS. Target 5 x 10e6/kg CD34 cells for infusion. Cryopreserve.

## Conditioning

| Day                | -6       | -5       | -4       | -3        | -2        | -1        | 0 |
|--------------------|----------|----------|----------|-----------|-----------|-----------|---|
| Cy*                | 50 mg/kg | 50 mg/kg | 50 mg/kg | 50 mg/kg  |           |           |   |
| Rabbit ATG**       |          |          |          | 0.5 mg/kg | 2.0 mg/kg | 2.0 mg/kg |   |
| Methylprednisolone |          |          |          | 40 mg bid | 40 mg bid | 40 mg bid |   |
| Stem cell infusion |          |          |          |           |           |           | X |

\* Cyclophosphamide (50 mg/kg ideal body weight in 250 mL D<sub>5</sub>W infused over 2 h) with Mesna (50 mg/kg ideal body weight in 1 L NS over 24 h starting with each Cy dose), hydration (NS at 75 mL/h starting the night before the first Cy dose and continuing till 24 h post the last Cy dose) and antiemetics (ondansetron + dexamethasone [except on day minus 3, when methylprednisolone is given as ATG premedication] + aprepitant + prn dimenhydrinate + prn metoclopramide)

\*\* Thymoglobulin (0.5-2.0 mg/kg in as low volume of NS as possible [0.5 mg/mL] infused over ≥4 h) with premedication (Methylprednisolone 1 mg/kg before each infusion + acetaminophen + diphenhydramine + meperidine prn)

Alternatively (e.g., in patients not meeting conventional HCT eligibility criteria), Burt's low-intensity, fludarabine-based protocol can be used:

## Mobilization

| Day                  | 1                    | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | ≥12 | ≥13 |
|----------------------|----------------------|---|---|---|---|---|---|---|---|----|----|-----|-----|
| Cy*                  | 2.5 g/m <sup>2</sup> |   |   |   |   |   |   |   |   |    |    |     |     |
| GCSF**               |                      |   |   |   |   |   | X | X | X | X  | X  | X   |     |
| Apheresis of MNCs*** |                      |   |   |   |   |   |   |   |   |    |    |     | X   |

\* Cyclophosphamide (2.5 g/m<sup>2</sup> dose dissolved in 500 mL D<sub>5</sub>W and infused over 2 h) with Mesna, hydration (500 mL NS over 1 h before each Cy infusion, and 500-1000 mL NS over 2-4 h after Cy infusion [500 mL over 2 h for <70 kg patient, 1000 mL over 4 h for ≥70 kg patient]) and antiemetics (granisetron + dexamethasone + aprepitant + prn dimenhydrinate + prn metoclopramide + prn prochlorperazine). Mesna, 2500 mg/m<sup>2</sup> IV, should be given in two to three divided doses, the first one concurrently with cyclophosphamide and the second (third) one 4 h (8h) later. The dose of Cy (2.5 g/m<sup>2</sup>) deviates from the Burt protocol, which used 2 g/m<sup>2</sup>. This is to keep it standard with our routine mobilization protocol – see SPM chapter on mobilization.

\*\* 300-900 ug per dose depending on weight per SPM chapter on Mobilization; with prn codeine

\*\*\* Start when blood CD34 count is >20 x 10<sup>6</sup>/L. Target to collect 5 x 10<sup>6</sup>/kg CD34 cells. No product manipulation (no CD34 cell selection). Cryopreserve.

## Conditioning

| Day                | -6     | -5                   | -4                   | -3                   | -2                   | -1        | 0 |
|--------------------|--------|----------------------|----------------------|----------------------|----------------------|-----------|---|
| Fludarabine        |        | 30 mg/m <sup>2</sup> | 30 mg/m <sup>2</sup> | 30 mg/m <sup>2</sup> | 30 mg/m <sup>2</sup> |           |   |
| Rabbit ATG**       |        | 0.5 mg/kg            | 1.0 mg/kg            | 1.5 mg/kg            | 1.5 mg/kg            | 1.5 mg/kg |   |
| Cy*                |        |                      |                      |                      | 60 mg/kg             |           |   |
| Rituximab***       | 500 mg |                      |                      |                      |                      |           |   |
| Methylprednisolone |        | 40 mg bid            | 40 mg bid            | 40 mg bid            | 40 mg bid            | 40 mg bid |   |
| Stem cell infusion |        |                      |                      |                      |                      |           | X |

\* Cyclophosphamide (60 mg/kg ideal body weight in 300 mL D<sub>5</sub>W infused over 2 h) with Mesna (60 mg/kg ideal body weight in 1 L NS over 24 h starting with the Cy dose), hydration (NS at 75 mL/h starting the night before the Cy dose and continuing till 24 h after the dose) and antiemetics (ondansetron + aprepitant + prn dimenhydrinate + prn metoclopramide. As Cy is infused after ATG, the methylprednisolone given before ATG serves also as an antiemetic for Cy). On day -2, hydration (NS at 75 mL/h) should be held while infusing ATG to avoid fluid overload.

\*\* Thymoglobulin (0.5-1.5 mg/kg in as low volume of NS as possible [0.5 mg/ml] infused over ≥4 h) with premedication (Methylprednisolone 1 mg/kg before each infusion + acetaminophen + diphenhydramine + meperidine prn)

\*\*\* Rituximab or a biosimilar. Premedications include acetaminophen + H1 antihistamine + H2 antihistamine + prn steroid/antihistamine/acetaminophen. Rituximab/biosimilar is non-formulary for SSc, so requires STEDT approval or the patient needs to pay (~\$1500). The timing of rituximab deviates from the Burt protocol (day -6 instead of -5). The reason is the difficulty for our inpatient unit to administer fludarabine+ATG+rituximab in one day.

### • Special management notes

- Discontinue DMARDs (e.g., MMF, MTX, cyclophosphamide) 2-4 weeks before mobilization to maximize the likelihood of a high CD34 cell yield. This does not apply to prednisone. If a patient is on prednisone before mobilization, prednisone should be continued peri transplant and tapered by day 37 (as routine – see below). Patients who have been on prednisone long-term may need a slower posttransplant taper.

- Avoid rapid intravascular volume changes, particularly fluid overload, and electrolyte concentration extremes (could trigger CHF or arrhythmia due to subclinical/subechocardiographic myocardial fibrosis)<sup>1,3</sup>
- Avoid hypertension (could trigger renal crisis) – use lisinopril or enalapril
- Supportive care post-transplant
  - Steroids, apart from peri-ATG, are routinely not used.
    - History: In SCOT trial, prednisone was used at 0.5 mg/kg/d from day 6 to day 21, then tapered by day 37, to prevent engraftment syndrome and serum sickness. This was not done in ASTIS trial. We did not use steroids in our first 16 patients who received ASTIS-like conditioning. Neither engraftment syndrome requiring systemic steroids nor serum sickness occurred (per Storek memory, chart review not done). In 2020-2022, in 9 patients, we used SCOT-like conditioning and, to replicate SCOT completely, we included prednisone at 0.5 mg/kg/d from day 6 to day 21, taper by day 37. In 2022, we transitioned back to ASTIS-like conditioning but kept the prednisone, however, changed it to 0.25 mg/kg/d from day 0 to day 21, taper by day 37. Among 7 patients who received the ASTIS-like conditioning with prednisone in 2022-2024, 1 developed renal crisis on or before d35, which responded only partially to captopril and resulted in chronic renal insufficiency with creatinine  $\geq 400$   $\mu\text{mol/L}$ , and 1 patient developed an episode of what appeared to be a mild renal crisis on or before d49 (between ~d20 and d49), which responded near-completely to perindopril. At that point it was decided to abandon the routine use of steroids peri-transplant, except peri-ATG.
  - ACE inhibitor or ARB should be considered, particularly for patients with a cardiac problem or a history of renal crisis. Lisinopril, 10-20 mg qd from start of conditioning to day 60 and targeting systolic BP 90-110 mmHg, was a part of the SCOT protocol. Can be discontinued already around day 30 in patients who need a calcium channel blocker (e.g. nifedipine XL) or a phosphodiesterase-5 inhibitor (e.g., sildenafil) for Raynaud's/digital ulcers, as the combination with lisinopril could be associated with symptomatic hypotension.
  - GCSF from day 7 till engraftment per our SPM
  - Valacyclovir from start of conditioning till 1 day before first dose of VZV vaccine
  - Septra from engraftment till 1 y per our SPM, possibly extension to 2 y if CD4<200/uL at 1 y
  - Levofloxacin from day 0 till engraftment (risk of cardiac mortality with sepsis)
  - Fluconazole from day 1 till day 28 (risk of esophageal candidiasis)
  - EBV and CMV PCR weekly till day 100 (risk of PTLD, particularly with rabbit ATG) and preemptive therapy per SPM
  - Vaccination per the Vaccination chapter of our SPM

## Allogeneic HCT (AlloHCT)

- Case reports suggest efficacy<sup>35-37</sup>
- The only case series is a CIBMTR registry study of 12 cases with follow up of surviving patients of at least 1 year<sup>38</sup>. Of the 12 patients, 6 died, and 6 are alive at 13-60 months posttransplant. SSc status at last follow up was not given. Thus, this report is not informative regarding efficacy but suggests that mortality after alloHCT may be substantial.
- AlloHCT should currently be considered only in patients with concurrent hematologic disease or under a clinical trial.

## CAR T cells

- Case reports and small series of autologous or allogeneic CD19-directed CAR T cells suggest efficacy for skin tightness, Raynaud's, digital ulcers, interstitial lung disease, and myocardial fibrosis.<sup>39,40</sup> Toxicity (CRS, neurotoxicity, hematotoxicity) appears to be less frequent and less severe than in patients with B cell malignancies.
- Currently available only under clinical trials. As of January 2025, none is open in Alberta.



## Pre-Transplant Tests/Appointments – see Table below

| Tests/Appointments*                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    | Rheumatologist to Arrange         | Calgary BMT to Arrange                           |
|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------|--------------------------------------------------|
| PFT, including 6MWT and O <sub>2</sub> sat by forehead probe if low by finger probe (Respirology or PFT lab)                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           | Yes                               |                                                  |
| Chest CT (contiguous & high res), BAL if infection suspected                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           | Yes                               |                                                  |
| VQ scan (Nuclear Medicine)                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             | Yes                               |                                                  |
| ECG +/- Holter (if PVCs or history of palpitation/syncope/falling)                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     | Yes                               |                                                  |
| Echocardiogram including strain                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        | Yes                               |                                                  |
| Cardiac MRI including Gad                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              | Yes                               |                                                  |
| Sperm bank or fertility gynecologist, if relevant                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      | Yes                               |                                                  |
| Blood tests <ul style="list-style-type: none"> <li>○ Labs including CBC+diff, LFTs, Creatinine, Albumin, CRP, ANA, CK, TSH, NTproBNP, Troponin T (Troponin I if elevated CK), IgM, IgG, IgA</li> <li>○ Malabsorption labs: Vitamin A, B12, C, E, folate, ionized Ca, P, Mg, iron studies (TIBC, transferrin saturation, ferritin)</li> <li>○ Serology for HIV, HSV, VZV, CMV, EBV, HepB, HepC</li> <li>○ Pregnancy test (pre-menopausal women only)</li> <li>○ INR, PTT</li> <li>○ Systemic sclerosis associated autoantibodies ("Systemic sclerosis Profile" at Mitogen Advanced Diagnostic Lab, use separate requisition)</li> </ul> | Yes                               |                                                  |
| Urine tests<br>(Urinalysis, urine albumin:creatinine ratio)                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            | Yes                               |                                                  |
| SSc Rheumatologist (confirm Dx, mRSS, quantify contractures, NFCapillaroscopy, baseline for postAHCT decision of relapse, research skin biopsy if patient consents)                                                                                                                                                                                                                                                                                                                                                                                                                                                                    | Yes                               | Yes<br>(if not yet seen by Drs. Durand or Osman) |
| GI doc + EGD (GAVE, esophageal stricture or infection)                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 | If Fe deficient                   | Yes                                              |
| Esophageal manometry (Dr M. Woo)                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       |                                   | Yes                                              |
| Dentist, including X-rays                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              | Encourage pt to see local dentist | Yes                                              |
| RHC with exercise (Dr J.Howlett)                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       |                                   | Yes                                              |
| BMA including flow cytometry & cytogenetics (if MDS suspected)                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         |                                   | Yes                                              |

\* If done >3 months before AHCT, consider repeating, particularly PFT and Echo.

Blood and urine tests should be repeated pre-mobilization unless done within 1 month pre-mobilization.

## Post-Transplant Tests/Appointments (at 6 months, and 1,2,3,4,5 years)

- Rheumatology appointment with Dr. Caylib Durand. This includes capillaroscopy and optionally research skin biopsy. Not needed for patients from Northern AB (taken care of by Dr. Mo Osman)
- GI appointment with Drs. Matt Woo or Dorothy Li (SHC). This includes optional research esophageal manometry
- Echocardiogram
- PFT: Spirometry, DLCO, 6MWT
- Chest CT – 1 year posttransplant only (for non-AB patients, this is just a recommendation)
- Oxygen saturation ideally by forehead probe, if <92%, then ABG
- Labs including CBC+diff, CRP, ANA, CK, TSH, NTproBNP, Troponin T (high sensitivity), IgM, IgG, IgA
- Urinalysis (random)
- Urine albumin:creatinine ratio (from spot urine), if unavailable, then protein:creatinine ratio
- Systemic sclerosis associated autoantibodies (“Systemic sclerosis Profile” at Mitogen Advanced Diagnostic Lab)
- Estradiol and anti-mullerian hormone (females <50-y-old), AM free testosterone (males), FSH and LH (both females <50-y-old and males) – 1 year posttransplant only
- CD4 T cell count – 1 year posttransplant only

## How to Refer for HCT:

- For Alberta Connect Care users: Enter Ambulatory Referral to Blood and Marrow Transplant Team – Internal Referral – To department “CGY ACCC BMT COORDINATION”.
- For non-Connect Care users, including patients from outside of Alberta: Fax referral letter to Calgary Blood and Marrow Transplant Program at 587-231-3994.
- Please refer all patients fitting the above criteria at the outset – not only after they have failed a therapy. In the three randomized studies, most patients received the HCT as first line therapy or after only a few months of immunosuppressive therapy.

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## Appendix 1: Methods and Results of Studies of AHCT for SSc\*

| Study                                              | N  | Patient characteristics                                       | HSC mobilization                        | Conditioning (or control Rx)             | CD34 selection | Med F/U (y) | TRM | Efficacy                                                                                 |
|----------------------------------------------------|----|---------------------------------------------------------------|-----------------------------------------|------------------------------------------|----------------|-------------|-----|------------------------------------------------------------------------------------------|
| <b>Non-randomized</b>                              |    |                                                               |                                         |                                          |                |             |     |                                                                                          |
| Binks M: Ann Rheum Dis 2001 <sup>41</sup>          | 41 | Age 41 (med)<br>Dis.dur. ~2 y<br>mRSS 29<br>FVC <70% in ½ pts | Cy 4 g/m <sup>2</sup> + GCSF (most pts) | Cy 150-200 mg/kg (most pts)              | Yes (most pts) | 1           | 17% | OS at 1 y 73%<br>mRSS improved<br>Lung function stable                                   |
| Farge D: Brit J Haematol 2002 <sup>42</sup>        | 11 | Age 46 (med)<br>Dis.dur. ~2 y<br>mRSS 29<br>FVC 67%           | Cy 4 g/m <sup>2</sup> + GCSF            | Cy 200 mg/kg (most pts)                  | Yes            | 1 ½         | 9%  | OS at 1 ½ y 64%<br>mRSS improved<br>QOL improved                                         |
| Nash RA: Blood 2007 <sup>43</sup>                  | 34 | Age 41 (med)<br>Dis.dur. <4 y<br>mRSS 30<br>FVC 72%           | GCSF                                    | Cy 120 mg/kg + TBI 8 Gy + Atgam 90 mg/kg | Yes            | 5           | 24% | OS at 5 y 64%<br>PFS at 5 y 64%<br>mRSS improved<br>Lung function stable<br>QOL improved |
| Oyama Y: Bone Marrow Transplant 2007 <sup>44</sup> | 10 | Age 46 (med)<br>Dis.dur. ~3 y<br>mRSS 30<br>FVC ~70%          | Cy 2 g/m <sup>2</sup> + GCSF            | Cy 200 mg/kg + Thymoglob. 7.5 mg/kg      | No             | 2           | 0%  | OS at 2 y 90%<br>PFS at 2 y 70%<br>mRSS improved<br>Lung function stable                 |
| Vonk MC: Ann Rheum Dis 2008 <sup>45</sup>          | 26 | Age 42 (med)<br>Dis.dur. ~2 y<br>mRSS 32<br>FVC 76%           | Cy 4 g/m <sup>2</sup> + GCSF            | Cy 200 mg/kg                             | Yes            | 5           | 4%  | OS at 5 y 96%<br>PFS at r y 64%<br>mRSS improved<br>Lung function stable                 |
| Tsukamoto H: Rheumatol 2011 <sup>8</sup>           | 11 | Age 52 (avg)<br>Dis.dur. <5 y<br>mRSS 22<br>FVC 65%           | Cy 4 g/m <sup>2</sup> + GCSF            | Cy 200 mg/kg                             | Yes            | 5           | 0%  | OS at 3 y 91%<br>mRSS improved<br>FVC 65→78%<br>DLCO stable<br>↓ Scl70, TNFa, TGFb       |
| <b>Randomized</b>                                  |    |                                                               |                                         |                                          |                |             |     |                                                                                          |



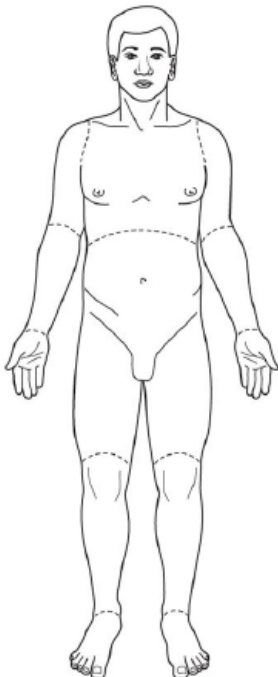
|                                                      |                               |                                                                                                               |                                       |                                                                                                                                            |     |     |                 |                                                                                                                                                                                                                                                           |
|------------------------------------------------------|-------------------------------|---------------------------------------------------------------------------------------------------------------|---------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------|-----|-----|-----------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Burt RK:<br>Lancet<br>2011 <sup>46</sup><br>(ASSIST) | 10<br>vs<br>9<br>ctrls        | Age 45<br>(med)<br>Disease<br>duration<br>~1 y<br>Cy <6 IV<br>doses<br>mRSS ~23<br>FVC ~65%                   | Cy 2<br>g/m <sup>2</sup><br>+<br>GCSF | Cy 200<br>mg/kg<br>+<br>Thymoglob.<br>6.5 mg/kg<br>(w M-pred<br>1 g x 4)<br><b>vs</b><br>Cy 1 g/m <sup>2</sup><br>monthly x 6              | No  | 1   | 0%<br>vs<br>0%  | OS @ 1 y 100% vs 100%<br>Evaluations at BL and at 1<br>y: mRSS 28→15 vs<br>19→22<br>FVC 62→74% vs<br>67→61%<br>QOL (SF36 total score)<br>39→56% vs 50→40%<br>(all differences between<br>groups significant, except<br>for OS)                            |
| Van Laar:<br>JAMA<br>2014 <sup>29</sup><br>(ASTIS)   | 79<br>vs<br>77<br>cntls<br>** | Age 44<br>(avg)<br>Disease<br>duration<br>~1 y<br>Cy <5 g IV<br>total<br>mRSS 25<br>(avg)<br>FVC 81%<br>(avg) | Cy 4<br>g/m <sup>2</sup><br>+<br>GCSF | Cy 200<br>mg/kg<br>+<br>Thymoglob.<br>7.5 mg/kg<br>(w M-pred<br>1 mg/kg x3)<br><b>vs</b><br>Cy 750<br>mg/m <sup>2</sup><br>monthly x<br>12 | Yes | 6** | 10%<br>vs<br>0% | OS @ 4y 86% vs 76%<br>EFS @ 4y 81% vs 74%<br>(event = death or<br>irreversible organ<br>failure)<br>Changes from BL to 2 y:<br>mRSS decrease, 20 vs 9<br>FVC increase, 5 vs -1%<br>QOL (SF36 physical<br>score)<br>increase, 10 vs 4<br>(all significant) |
| Sullivan:<br>NEJM<br>2018 <sup>30</sup><br>(SCOT)    | 36<br>vs<br>39<br>cntls       | Age 18-69<br>Disease<br>Duration<br>≤5 y<br>Cy up to 4-<br>6 mo<br>mRSS 30<br>(avg)<br>FVC 74%<br>(avg)       | GCSF                                  | Cy 120<br>mg/kg<br>+ TBI 8 Gy<br>+ Atgam<br>90 mg/kg<br><b>vs</b><br>Cy 750<br>mg/m <sup>2</sup><br>monthly<br>x12                         | Yes | >5  | 3%<br>vs<br>0%  | OS @ 4 ½ y 91% vs 77%<br>EFS @ 4 ½ y 79% vs<br>50%<br>(event = death or<br>renal/cardiac/pulmonary<br>failure)<br>(all significant)                                                                                                                       |

\* Only studies with ≥10 patients are shown.

\*\* Only 71 vs 57 patients completed treatment, and 8 controls received HCT at ≥2 y. The analyses under Efficacy are intention-to-treat analyses.

**Abbreviations:** Dis.dur., disease duration; med, median; mRSS, modified Rodnan skin score; FVC, forced vital capacity; Cy, cyclophosphamide; GCSF, granulocyte colony stimulating factor (filgrastim); HCT, hematopoietic cell transplantation; F/U, follow up; TRM, transplant related mortality; TBI, total body irradiation; Thymoglob., Thymoglobulin; M-pred, methylprednisolone; pts, patients; OS, overall survival; PFS, progression-free survival; EFS, event-free survival; QOL, quality of life; TNFα, tumor necrosis factor alpha; TGFβ, transforming growth factor beta; BL, baseline;

## Appendix 2: Modified Rodnan Skin Score (mRSS)



|                | Right                                                                                                       |                            |                            |                            | Left                       |                            |                            |                            |
|----------------|-------------------------------------------------------------------------------------------------------------|----------------------------|----------------------------|----------------------------|----------------------------|----------------------------|----------------------------|----------------------------|
| Fingers        | 0 <input type="checkbox"/>                                                                                  | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> | 0 <input type="checkbox"/> | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> |
| Hands          | 0 <input type="checkbox"/>                                                                                  | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> | 0 <input type="checkbox"/> | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> |
| Forearms       | 0 <input type="checkbox"/>                                                                                  | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> | 0 <input type="checkbox"/> | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> |
| Upper Arms     | 0 <input type="checkbox"/>                                                                                  | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> | 0 <input type="checkbox"/> | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> |
| Face           | 0 <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> |                            |                            |                            |                            |                            |                            |                            |
| Anterior Chest | 0 <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> |                            |                            |                            |                            |                            |                            |                            |
| Abdomen        | 0 <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> |                            |                            |                            |                            |                            |                            |                            |
| Thighs         | 0 <input type="checkbox"/>                                                                                  | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> | 0 <input type="checkbox"/> | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> |
| Legs           | 0 <input type="checkbox"/>                                                                                  | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> | 0 <input type="checkbox"/> | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> |
| Feet           | 0 <input type="checkbox"/>                                                                                  | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> | 0 <input type="checkbox"/> | 1 <input type="checkbox"/> | 2 <input type="checkbox"/> | 3 <input type="checkbox"/> |
| Column Totals  |                                                                                                             |                            |                            |                            |                            |                            |                            |                            |
| <b>Total:</b>  |                                                                                                             |                            |                            |                            |                            |                            |                            |                            |
| Key:           | 0 – No Thickening      1 – Mild Thickening      2 – Moderate Thickening      3 – Severe Thickening          |                            |                            |                            |                            |                            |                            |                            |
| <b>Notes:</b>  |                                                                                                             |                            |                            |                            |                            |                            |                            |                            |

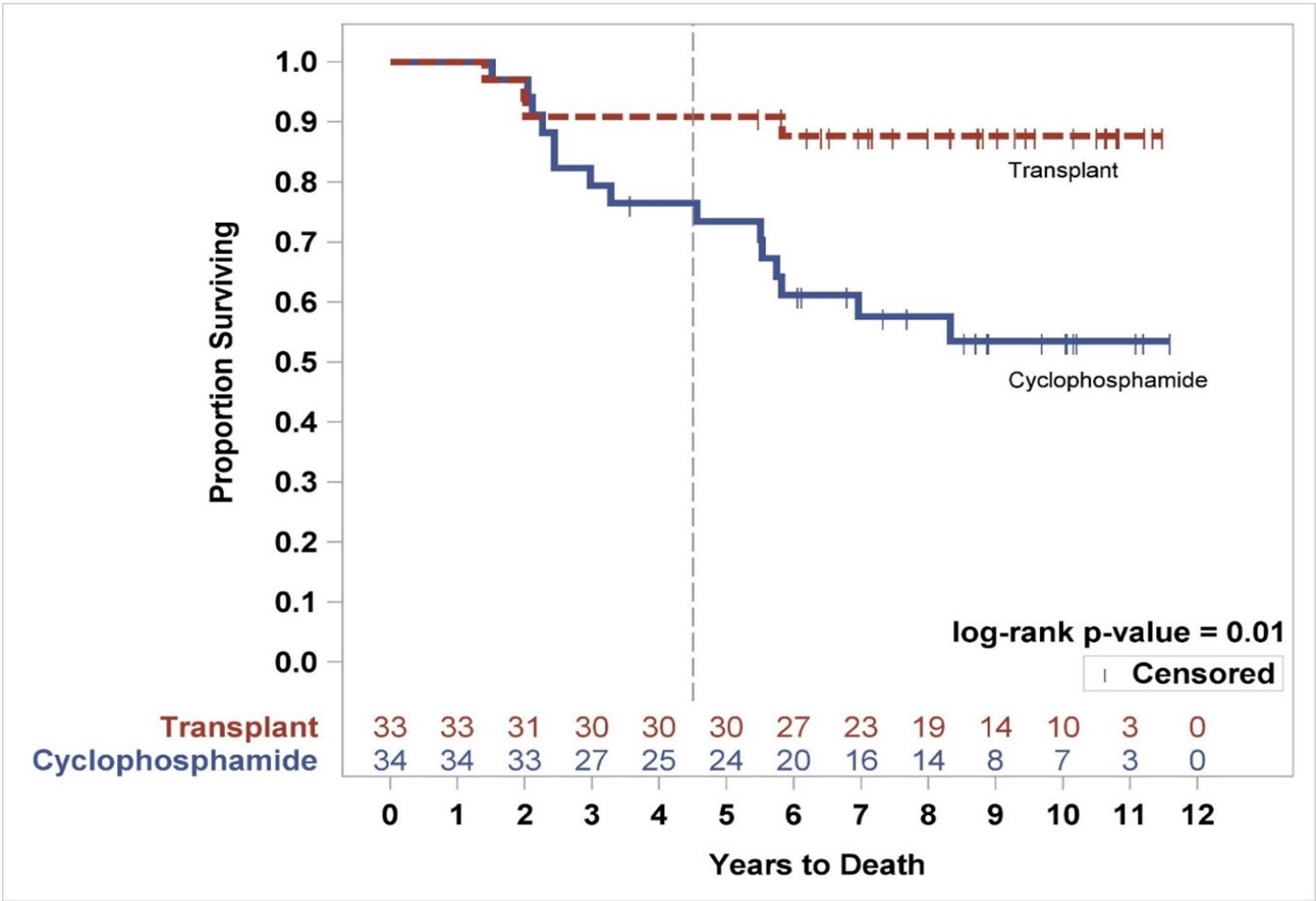
Uninvolved skin = 0, Mild thickening = 1, Moderate thickening = 2, Severe thickening (cannot pinch) = 3.  
 mRSS is the total of points from the above locations (max 51). From Klippel JH: Rheumatology, Mosby 2000.

The total score is out of 51. Ideally the skin scores are evaluated by an experienced Rheumatologist familiar with systemic sclerosis. Tethering is not scored as a 3 but should be noted. Scores are a reflection of the total average for the area of skin tested and not maximum scores as this can be misleading with patches of skin being overrepresented. Care must also be given to lower extremity evaluation because of complications from edema and other comorbidities.

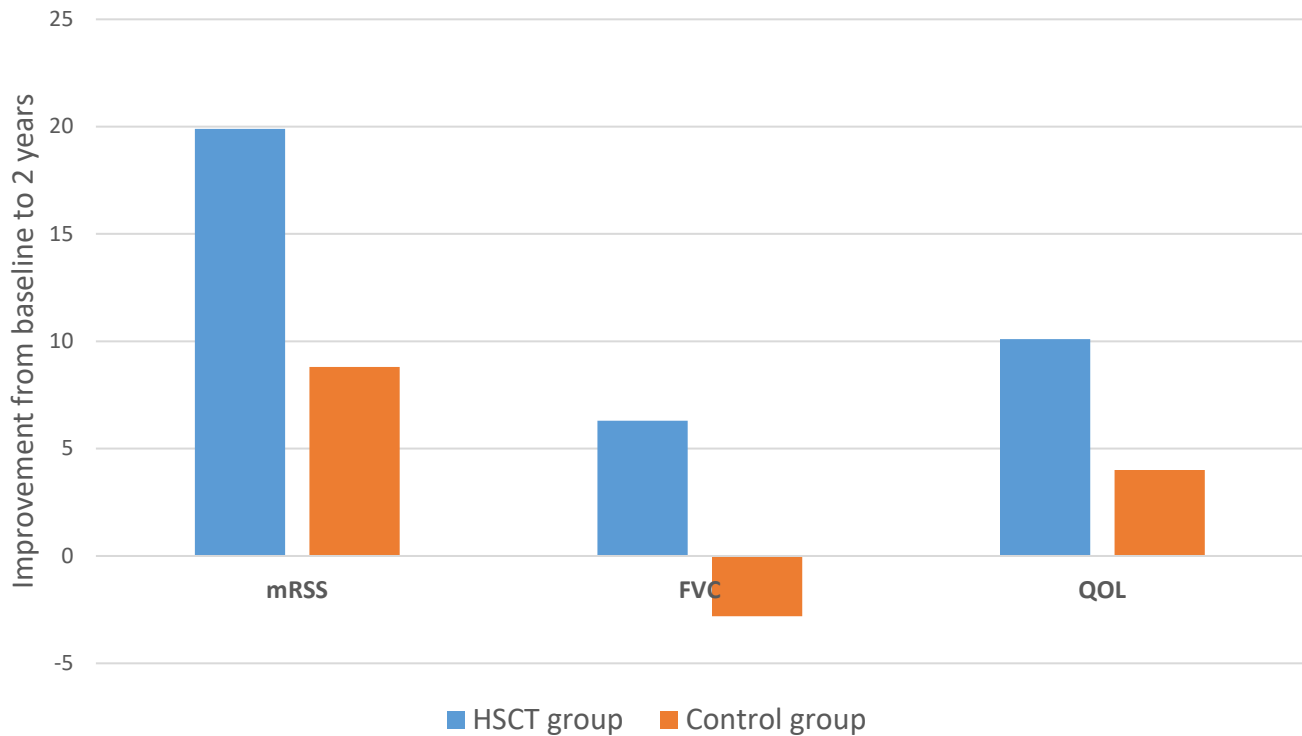
| mRSS |                                                                                                                                                      |                                                                                      |
|------|------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------|
| 0    | <p><b>Fine wrinkles in skin</b></p> <p><b>No skin thickness when rolling skin</b></p>                                                                |    |
| 1    | <p><b>Appreciable skin thickness but still able to make skin fold between 2 fingers</b></p> <p><b>Still able to demonstrate fine wrinkles</b></p>    |    |
| 2    | <p><b>Appreciable skin thickness with difficulty in making skin folds</b></p> <p><b>No fine wrinkles</b></p>                                         |   |
| 3*   | <p><b>Appreciable skin thickness with inability to make skin folds between 2 fingers</b></p> <p><b>Don't count tethering and atrophy! mRSS=0</b></p> |  |

\*No not count scores for tethering or atrophy.

Appendix 3: Survival, mRSS, FVC, and Quality of Life in Randomized Studies of HCT versus conventional therapy (cyclophosphamide).



Overall survival in SSc patients randomized to hematopoietic stem cell transplantation (HSCT) vs 1 year of cyclophosphamide (control). Long-term follow up data on patients from the SCOT study.<sup>31</sup> Presented at American College of Rheumatology 2018 annual meeting by Sullivan KM et al - Abstract 1820.



Modified Rodnan Skin Score (mRSS), forced vital capacity (FVC), and quality of life (QOL) in the ASTIS study. Between baseline and 2 years after start of treatment, mRSS dropped by mean 20 points in the patients randomized to HSCT (hematopoietic stem cell transplantation) vs 9 points in the control patients randomized to 1 year of cyclophosphamide ( $p<0.001$ ). FVC improved by +6 vs -3 percentage points ( $p=0.004$ ), and quality of life (QOL) assessed by Short Form 36 Physical Component improved by 10 vs 4 points ( $p=0.01$ ).<sup>29</sup>

# Appendix 4: Information for Patients

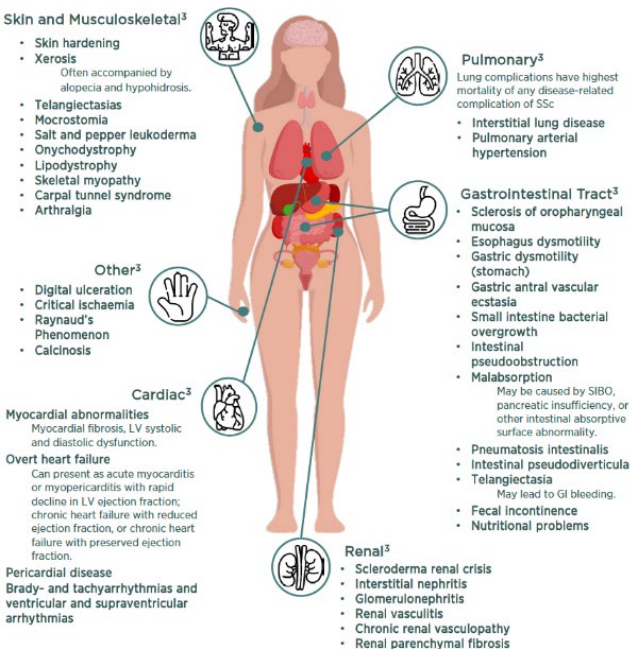
## Understanding Systemic Sclerosis and Autologous Hematopoietic Cell Transplantation in Alberta

### What is Systemic Sclerosis:

Systemic sclerosis (SSc) is a multi-organ, autoimmune, connective tissue disease. It is relatively rare, with the prevalence in Canada among the highest in the world (44/100000). Patients with SSc live an average of 20 years shorter than the general population, and while SSc is 8-9 times more common in women, outcomes in men are significantly worse. There are different types of SSc with one third of all patients with SSc develop the most severe form called diffuse cutaneous systemic sclerosis (dcSSc). Skin tightness or fibrosis is a marker of disease severity and severe skin tightness or fibrosis is a hallmark of dcSSc that has a much higher level of disability and increased risk of death compared to patients with limited cutaneous SSc (lcSSc). Most often SSc is treated with medications that suppress the immune system (e.g., MMF (CellCept)), but they have limited success in the long term management of rapidly progressive dcSSc.

LeRoy and Medsger Classification Criteria<sup>5,11</sup>

| Criteria                             | dcSSc | lcSSc | Sine SSc |
|--------------------------------------|-------|-------|----------|
| Systemic involvement                 | X     | X     | X        |
| Proximal cutaneous changes           | X     |       |          |
| Distal cutaneous changes             | X     | X     |          |
| Raynaud's phenomenon                 | X     | X     | X        |
| SSc-type nail-fold capillary pattern | X     | X     | X        |
| SSc selective autoantibodies         | X     | X     | X        |



**Autologous hematopoietic cell transplant (abbreviated as AutoHCT or HCT) is offered in Alberta for the treatment of dcSSc.**

### What is autologous HCT?

First, your blood-forming stem cells are collected. Next, you receive high dose immunosuppressive chemotherapy that is thought to kill the cells causing SSc. Next, the previously collected stem cells are reintroduced into your body. The stem cells make your immunity against bugs recover faster than if you receive chemotherapy without the stem cells. In the last decade, three randomized trials of HCT (ASSIST, ASTIS and SCOT) were published. Despite differences between patient populations and HCT techniques, all trials demonstrated superiority over conventional medical therapies for dcSSc. One of the most important outcomes for transplant is improved chance of survival. It can also result in less skin



tightness/thickenening/fibrosis along with improvement in other disease outcomes. However, AutoHCT is not for all patients with a summary of eligibility below.

#### Alberta AutoHCT Program Eligibility:

|                         |                                                                                                  |
|-------------------------|--------------------------------------------------------------------------------------------------|
| <b>Age:</b>             | Age up to 65. Younger patients typically do better.                                              |
| <b>Disease Onset:</b>   | Less than 5 years from first non-Raynaud symptom.                                                |
| <b>Type of Disease:</b> | Diffuse systemic sclerosis, severe skin disease (mRSS>20) or less severe skin if lungs involved. |
| <b>Restrictions:</b>    | No features of heart failure or pulmonary hypertension (higher risk of death).                   |
|                         | Only mild or moderate interstitial lung disease (FVC 40-80%, ideally 60-80%).                    |
|                         | No untreated watermelon stomach (risk of bleeding).                                              |

#### Establishing Expectations for AutoHCT:

Rapidly progressive dcSSc has increased risk of death and disability but there are risks associated with AutoHCT as well. First and most important is that there is the risk of transplant-related death of up to 10%, usually due to lung or heart failure with or without an infection. This risk is usually within the first several months post-transplant and is overall still lower than the risk of death associated in the long run with SSc (not treated with AutoHCT). It is also important to understand that it is likely that the AutoHCT will not completely reverse all clinical features of SSc, especially well established fibrosis of the skin or other internal organs. The overall goal is to improve survival and to stop systemic sclerosis progression. Any reversal of other symptoms is hoped for but not always obtained.

#### **If you are offered HCT, there is a net benefit compared to conventional medical therapy.**

There are also other potential risks including a relapse where your disease could return. There is a risk of infection especially within the first 100 days of transplantation. There are possible risks of developing a new (unrelated) autoimmune disease (approximately 10% chance). There may be an increased risk of cancer of any organ, but this is not known for sure. There is a possibility of infertility or premature menopause because of the chemotherapy. This list covers most but not all potential complications of AutoHCT. It is important for each patient to weigh the benefits and risks for transplant before deciding to continue.

#### **What to expect for the AutoHCT evaluation:**

Typically, by the point of seeing the transplant team the patient has been seen and diagnosed by a Rheumatologist with dcSSc and then referred to a Rheumatologist with a focus in systemic sclerosis. The systemic sclerosis specialist then refers to the transplant team for further workup and evaluation. For the AutoHCT evaluation you will meet a Rheumatologist and Hematologist associated with the transplant program.

Multiple tests will be arranged for, including blood/urine tests, pulmonary function test, chest imaging like CAT scan, echocardiogram, cardiac MRI, right heart catheterization, endoscopy of esophagus and stomach.

Your evaluation may also require you to meet with other specialists depending on the organ systems involved but usually include:

- Hematologist, Rheumatologist, Gastroenterologist, Cardiologist, Dentist.

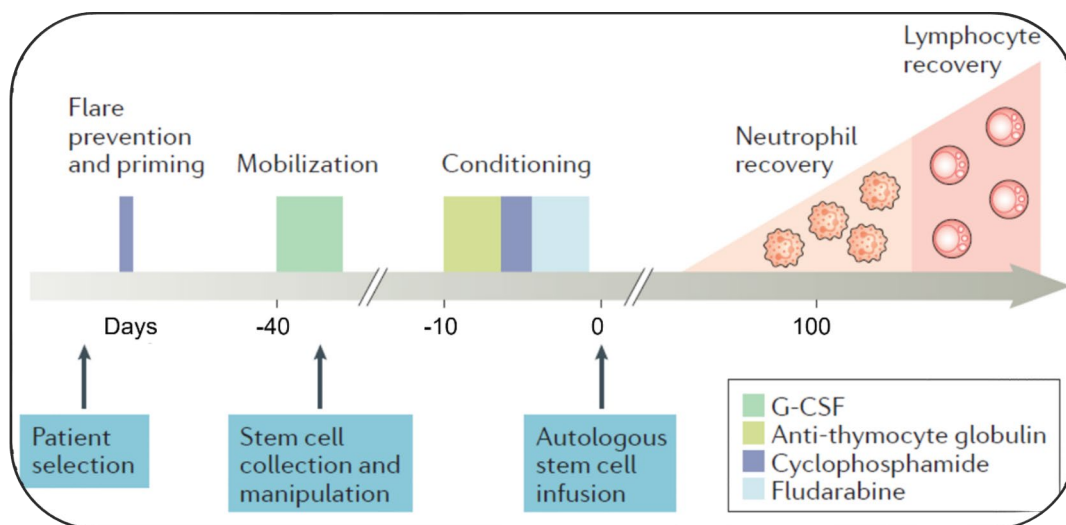
It may also include (less likely):

- Respiriologist, Nephrologist, Dermatologist, Fertility specialist.

After the work up and meetings are completed, the transplant team will decide if there are reasons the transplant should not be considered. If cleared for transplant and you have decided to proceed then the transplant is arranged (usually within 3 months of starting the referral, assuming most of the pretransplant tests have been completed before the referral).

### What to expect during HSCT:

Most likely you will be on immunosuppression medications or medications designed to control the immune system before the transplant. These will need to be stopped before starting the transplant. You will then have your blood-forming stem cells mobilized and collected (outpatient process in Calgary that takes 2-3 weeks), then be admitted to the Arthur Child Comprehensive Cancer Centre in Calgary to get the conditioning chemotherapy and get treatment for the early toxicities of the chemotherapy. Your admission length will vary based on how well you tolerate the transplant, typically one month, sometimes longer. The process is graphically shown in the figure below:



### What Happens After AutoHCT:

At 2-3 months post-transplant you will return home, assuming no significant complications. You will be immunocompromised meaning you will be at increased risk of infection. It will take 1-2

years before your immune system has recovered to normal. You will need to repeat your childhood vaccinations. You will need regular follow up with your primary Rheumatologist and will need to see the Calgary transplant team of specialists at 6 months, 1 year, and then yearly for up to 5 years following your transplant. You will also have yearly investigations and blood work to do to monitor your progress and to watch for complications or relapse.

In case of transplant success, you will be off all immunosuppressive medications. However, there might still be a need for other medications to manage esophageal or intestinal symptoms, Raynaud's, or other organ symptoms which may be lifelong.

In case of transplant success, work or school can be typically started at 6-12 months post-HCT.

### **Patients from outside Calgary or out of Province patients:**

1. Calgary, Alberta is the AutoHCT program for SSc for Western Canada.
2. Your local Rheumatologist will need to refer you to the transplant team for evaluation and transplantation.
3. If accepted you will need to be in Calgary for about 4 months for pretransplant care, the transplant, and posttransplant care.
4. You will need to travel back to Calgary for long-term follow up appointments as not all centers have the specialists needed for the transplant evaluations. It is also important to centralize most of the data collection for the transplant and systemic sclerosis evaluations. You will still follow up with your local Rheumatologist for regular visits. Patients from the Edmonton and surrounding area will follow up for the yearly Rheumatology appointment in Edmonton with Dr Mo Osman but will still need to see the Hematologist in Calgary.
5. After 2-5 years, or if your SSc has relapsed, you will be discharged from the program to follow up with your local Rheumatologist.
6. Important Social and Financial Considerations:
  - a. A caregiver (a relative or a friend) is required to stay with the patient in Calgary for a minimum of one month after hospital discharge. Ideally the caregiver should stay with the patient in Calgary for a total of 4 months (1-2 months pre-transplant and 2-3 months post-HCT)
  - b. Despite the inpatient and outpatient medical care, including inpatient medications, are fully covered, there are important out-of-pocket expenses for patients who live >1 h car ride from Calgary. Here are the estimates:
    - i. Accommodation in Calgary at \$1000-2000/month x 4 months = \$4000-8000
    - ii. In-Calgary transportation at \$1000 (if accommodation is not within walking distance from Arthur Child Comprehensive Cancer Centre)
    - iii. Co-pays for outpatient medications: \$500-3000, assuming 80% is paid by patient's insurance. If patient is on provincial disability benefits, the drugs should be fully covered
    - iv. Transportation from home to Calgary and back. Sometimes covered by a charity (e.g., Hope Air)

- v. Other Expenses that should be considered are meals in Calgary. Meals for caregiver should also be considered. If caregiver has to take a leave from work, s/he can apply for Caregiver Benefits (55% of wage to a max of ~\$600/week in Alberta)
- vi. Total of the out-of-pocket expenses: \$5,500-12,000 plus the Other Expenses, plus transportation from home to Calgary and back.

# Transplantation for Germ Cell Tumours

Presented by: Doug Stewart

Updated by: Kevin Hay

## Summary

- High-dose chemotherapy (HDCT) with autologous stem cell transplantation (ASCT) is indicated in second or third line therapy (i.e. as therapy for 1<sup>st</sup> or 2<sup>nd</sup> relapse) for patients with advanced germ cell tumor. Patients in first relapse who are likely to be cured with conventional dose chemotherapy (CDCT) alone such as TIP include those with low International Prognostic Factor Study Group (IPFSG) scores and those with gonadal or retroperitoneal primary site who have achieved a CR or a marker-negative PR lasting >6 months prior to their first relapse. However, patients in first relapse who are unlikely to be cured with CDCT alone should be considered for HDCT as part of initial salvage therapy, including those with higher IPFSG scores, incomplete response to first-line cisplatin-based therapy, primary platinum refractory disease, or relapse 6 months or less after achieving a marker-negative PR. For patients treated with CDCT in the initial salvage setting, HDCT remains an option in the third-line setting, should subsequent relapses occur.
- Patients with a late relapse >2 years after completing initial chemotherapy or growing teratoma syndrome typically have a chemoresistant disease and are prioritized for surgical resection.
- Stem cell mobilization at our center has typically been performed using the second cycle of salvage bridging chemotherapy; however, salvage CDCT is not required prior to HDCT and some patients can be taken to HDCT using G-CSF alone for mobilization. Typical salvage regimens include TIP (paclitaxel 175 mg/m<sup>2</sup> d1, ifosfamide 1.67g/m<sup>2</sup> d1-3, cisplatin 33 mg/m<sup>2</sup> d1-3, G-CSF 5-10mcg/kg/d starting day 9, and apheresis scheduled days 14-16), or TI (paclitaxel 200 mg/m<sup>2</sup> d1, ifosfamide 2g/m<sup>2</sup> d2-4, GCSF 10mcg/kg from d4 until adequate collection or neutrophil recovery, apheresis on days 11-14). DICEP has also been used successfully at our centre for difficult to mobilize cases. A minimum of 1-2 million CD34+ cells/kg is required for each cycle of HDCT.
- Demonstrating a response to salvage bridging CDCT is not required since HDCT can still achieved durable remissions in a subset of patients with platinum-refractory disease.
- Standard HDCT conditioning for GCT involves tandem transplants using 2 cycles of high-dose Carboplatin 700 mg/m<sup>2</sup>/day plus Etoposide 750 mg/m<sup>2</sup>/day, both given d-5,-4,-3 before ASCT. The second cycle of HDCT is given after recovery of granulocyte and platelet counts, unless there was a grade 4 nonhematologic toxic effect or no response to the first course (based on tumour markers and imaging). In general, the time between day 0 ASCT#1 and day 0 ASCT#2 is only 4-5 weeks.

## Background

Germ cell tumors (GCTs) account for less than 1% of all cancers; however, they represent the most common malignancy in young men between the ages of 15 and 35 years. Approximately 70% of patients with advanced disease are cured with conventional-dose, platinum-based chemotherapy. For patients with advanced disease the current standard first-line therapy is 3-4 cycles of cisplatin, etoposide and bleomycin (BEP). There is no role for high-dose chemotherapy (HDCT) and autologous stem cell transplantation (ASCT) in the first-line treatment of patients with germ cell tumours<sup>1</sup>. Patients who do not achieve long-term remission with initial chemotherapy are still often curable with second- and even third-line treatment strategies. Options include cisplatin and ifosfamide with either paclitaxel (TIP) or vinblastine (VIP) with durable complete response (CR) rates of up to 63% in phase II trials in well-selected patients. Another salvage approach is the use of HDCT and ASCT which has demonstrated long-term remissions primarily in single-arm and retrospective studies.

## Stem Cell Transplantation in GCT

In 2007, Indiana University published a large retrospective evaluation of their experience using high dose carboplatin and etoposide in 184 consecutive patients<sup>2</sup>. Most patients (73%) were treated in the initial salvage setting. The high-dose regimen consisted of two cycles of 700 mg/m<sup>2</sup> of carboplatin plus 750 mg/m<sup>2</sup> of etoposide, both given intravenously 5, 4, and 3 days before ASCT. Four-year PFS was 63% for the study cohort. In a 2017 update of 364 consecutive patients with GCT who progressed after cisplatin-based chemotherapy and received HDCT, the 2-year PFS was 60% and 2-year OS was 66% (Figure 1)<sup>3</sup>. Patients with late relapse of GCT >2 years after previous therapy were excluded. In a multivariable analysis, factors associated with disease progression included use of HDCT as third-line or later therapy, platinum-refractory disease (defined as tumor progression within 4 weeks of cisplatin-based chemotherapy), mediastinal primary tumor site, nonseminoma histology, intermediate- or poor-risk IGCCCG disease at the time of GCT diagnosis, and human chorionic gonadotropin >1,000 mIU/mL at initiation of HDCT. However, durable remissions were still reported in 20-40% of patients with these high-risk features (Table 1). There were 9 (2.5%) treatment-related deaths and 5 (1.3%) patients developed therapy-related leukemia. This multivariable analysis confirmed prior results from the 2007 study, which had developed a prognostic scoring algorithm based on three variables: >2 line (3 points), platinum refractory (2 points), and advanced IGCCCG stage (2 points), Figure 2<sup>2</sup>. This 2007 had not included patients with mediastinal primary tumor site.



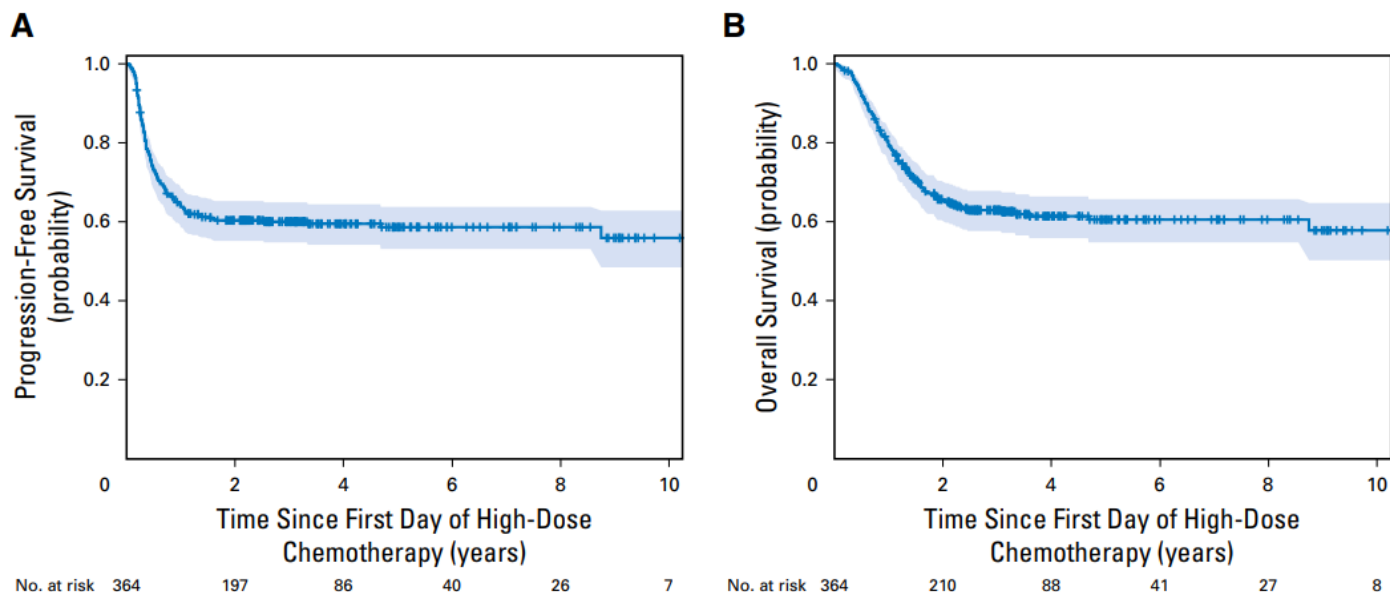


**Figure 1:** Outcomes of HDCT for relapsed GCT at Indiana University

| Variable (no. of patients) | 2-year PFS, % | <i>P</i> * |
|----------------------------|---------------|------------|
| High-dose chemotherapy     |               | 0.03       |
| Second line (303)          | 63            |            |
| Third line or later (61)   | 49            |            |
| Platinum sensitivity       |               | <0.001     |
| Sensitive (242)            | 75            |            |
| Refractory (122)           | 33            |            |
| Location of primary tumour |               | <0.001     |
| Testis/RP (344)            | 63            |            |
| Mediastinum (20)           | 23            |            |
| Tumour histology           |               | <0.001     |
| Seminoma (79)              | 90            |            |
| Nonseminoma (285)          | 52            |            |
| Initial IGCCCG risk        |               | <0.001     |
| Good (151)                 | 82            |            |
| Intermediate (39)          | 58            |            |
| Poor (174)                 | 43            |            |
| Serum AFP, ng/mL           |               |            |
| < 1,000 (336)              | 62            | 0.05       |
| ≥ 1,000 (28)               | 42            |            |
| Serum hCG, mIU/mL          |               |            |
| < 1,000 (274)              | 69            | <0.001     |
| ≥ 1,000 (90)               | 37            |            |
| Age (continuous variable)  | NA            | 0.44       |

**Table 1:** Outcomes of HDCT in different subgroups of patients with relapsed GCT at Indiana University

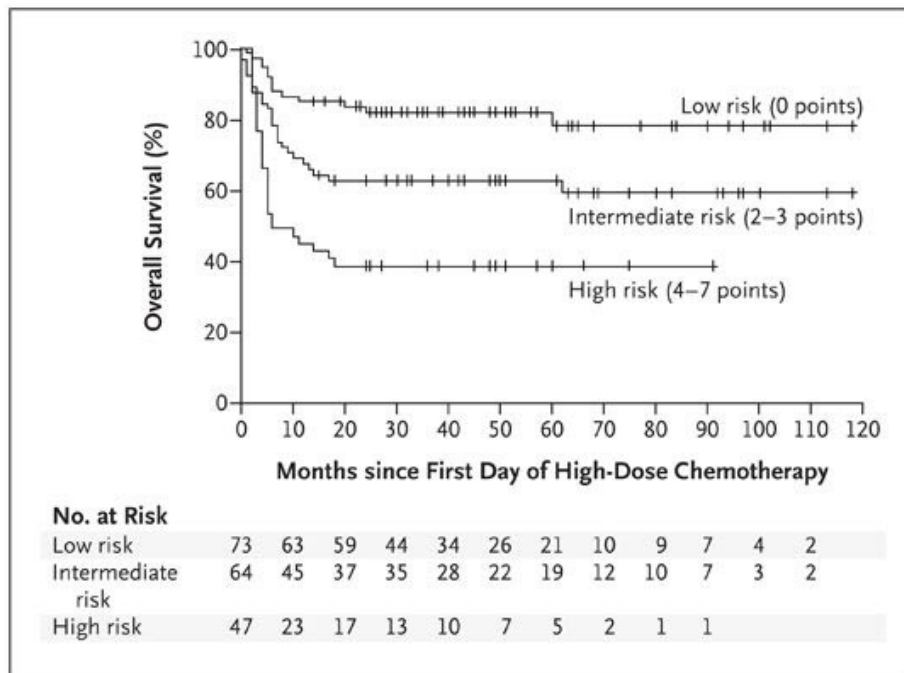


**Figure 2:** Prognostic model for outcomes of HDCT for relapsed GCT at Indiana University

### HDCT vs CDCT for relapsed GCT

The IT-94 randomized Phase III trial compared HDCT to conventional dose chemotherapy (CDCT) in the salvage setting<sup>4</sup>. This multicenter international study enrolled 280 patients from 43 institutions in 11 countries. The trial compared the efficacy of four cycles of CDCT using etoposide/ifosfamide/cisplatin (VIP)/VeIP versus three cycles of the same CDCT followed by one cycle of HDCT using carboplatin (200–550 mg/m<sup>2</sup>), etoposide (1800 mg/m<sup>2</sup>) and cyclophosphamide (200 mg/kg) followed by autologous stem cell rescue. Although no survival benefit was observed for HDCT, the interpretation of the trial is limited by the fact that it lacked sufficient statistical power, patients refractory to first-line platinum-containing chemotherapy were excluded, and that only one cycle of HDCT was provided while those studies which reported an advantage of HDCT over historical results with CDCT included two or more HDCT cycles.

Data from a large multicenter, international retrospective analysis of initial salvage chemotherapy in approximately 1600 subjects were reported in 2011<sup>5</sup>. Approximately equal numbers of patients were treated with CDCT and HDCT. Overall, PFS and OS were found to be superior for patients treated with HDCT as compared with CDCT. On multivariable analysis, important prognostic factors were identified that allowed patient stratification into five well-defined prognostic categories. Within these prognostic categories, PFS and OS remained superior for HDCT in each class with the exception of OS in the low-risk group. Despite the lack of randomized evidence, retrospective studies such as these support the consensus in international guidelines that HDCT is an effective second-line or third-line therapy for patients with relapsed metastatic GCT<sup>6-8</sup>. The Alliance 031102/EORTC 1407 (TIGER)

trial randomized 420 patients with relapsed GCT using the TI-CE approach to HDCT versus conventional TIP chemotherapy ([NCT02375204](#)). TI-CE involves 2 cycles of salvage CDCT TI in 14 day cycles, with apheresis for collection of stem cells, followed by 3 cycles (21 days apart) of HDCT carboplatin AUC targeted to 21 to 24 mg/ml/min total and etoposide 400 mg/m<sup>2</sup>/day. This completed enrollment in December 2022 and is expected to provide a definitive answer to the role for second line HDCT in GCT.

The optimal treatment approach for patients with relapsed metastatic GCT likely varies in accordance with underlying risk factors. Retrospective data suggests that patients with low-risk IPFSG scores may achieve comparable survival rates with CDCT versus HDCT<sup>5</sup>. Indeed, patients with gonadal or retroperitoneal primary site, who have achieved a CR or a marker-negative PR lasting >6 months prior to their first relapse, frequently achieve durable remissions with TIP. For patients treated with CDCT in the initial salvage setting, HDCT remains an option in the third-line setting, should subsequent relapses occur<sup>3</sup>. Conversely, patients with higher IPFSG score and those with incomplete response to first-line cisplatin-based therapy, primary platinum refractory disease, or who relapse 6 months or less after achieving a marker-negative PR, are usually considered for second-line HDCT. Although patients with primary mediastinal NSGCT or very high risk IPFSG scores experience less favorable outcomes with HDCT, the available evidence still supports the consideration of HDCT in these high-risk groups given that durable responses can be achieved in >20-30%<sup>3, 5</sup>. Patients with brain metastases may still benefit from high-dose carboplatin/etoposide conditioning, although surgery and/or radiation may first be required for symptomatic or hemorrhagic brain metastases<sup>9</sup>. In contrast, patients relapsing >2 years after first-line therapy and those with growing teratoma syndrome tend to be chemoresistant and should be prioritized for surgical approaches instead<sup>6</sup>.

### **Stem cell mobilization**

Demonstrating a response to CDCT is not necessarily required prior to HDCT in relapsed GCT, since HDCT can still achieve durable remissions in a subset of patients with platinum-refractory disease<sup>3</sup>. At Indiana University, patients with relapsed GCT typically undergo stem cell mobilization using G-CSF and then proceed directly to HDCT<sup>3</sup>. Bridging chemotherapy is not typically administered for patients with platinum-refractory disease, whereas an optional 1-2 cycles of bridging VIP is considered for patients with platinum-sensitive disease to control symptoms before HDCT. Due to the frequently urgent need for treatment and the resource and logistical constraints at our center, stem cell mobilization has typically been performed using the second cycle of bridging chemotherapy (e.g., TIP: paclitaxel 175 mg/m<sup>2</sup> d1, ifosfamide 1.67g/m<sup>2</sup> d1-3, cisplatin 33 mg/m<sup>2</sup> d1-3), with G-CSF 5-10mcg/kg/d starting day 9 and apheresis scheduled days 14-16. However, difficulties with stem cell mobilization have been observed using TIP<sup>10</sup>. An alternative approach utilized at MSKCC and in the TIGER trial is to collect stem cells using 1-2 cycles of bridging TI chemotherapy (paclitaxel 200 mg/m<sup>2</sup> d1, ifosfamide 2g/m<sup>2</sup> d2-4, GCSF 10mcg/kg from d4 until adequate collection or neutrophil recovery, apheresis on days 11-14) and then proceeding to HDCT. A second cycle of TI may be used to

mobilize additional stem cells if needed. Of note, as few as  $1-2 \times 10^6/\text{kg}$  CD34+ stem cells may be sufficient for each cycle of HDCT in GCT<sup>11</sup>.

### **Sequential HDCT Cycles vs Single HDCT/ASCT for GCT**

German investigators reported the results of a randomized trial that was designed to answer the question of whether multiple sequential HDCT cycles are superior to a single HDCT cycle<sup>12</sup>. Between November 1999 and November 2004, 211 patients with relapsed or refractory GCT were randomly assigned to treatment with either one cycle of conventional-dose VIP plus three additional cycles of high-dose carboplatin  $1,500 \text{ mg/m}^2$  and etoposide  $1,500 \text{ mg/m}^2$  (CE) over 3 days. Treatment in arm B involved three identical conventional dose cycles of VIP plus one additional cycle of high-dose carboplatin  $2,200 \text{ mg/m}^2$ , etoposide  $1,800 \text{ mg/m}^2$ , and cyclophosphamide  $6,400 \text{ mg/m}^2$  (CEC) given over 4 days. The investigators found no statistically significant differences in event-free survival (EFS), progression-free survival (PFS) or OS between the two groups. Toxicity was more severe within the single high-dose CECy arm with 16% treatment-related deaths as compared with 4% in the sequential high-dose CE arm, which led to the premature closure of the trial and a nonsignificant trend toward improvement in OS for the sequential arm. The final conclusion of the study is that 2-3 sequential high-dose cycles remain the standard of care when HDCT is used with curative intent during the treatment of GCT. Restaging tumor markers +/- imaging should be performed after cycle 1 of HDCT to document a response. Patients not responding to the first cycle of HDCT are unlikely to benefit from a second cycle and should be considered for surgical resection or palliative intent standard chemotherapy instead<sup>13</sup>.

### **Post ASCT surgical resection of residual disease**

The management of relapsed/refractory germ cell tumours is multidisciplinary, and the majority of patients with evidence of residual disease after ASCT should be considered for surgical resection if deemed feasible.

### **Prognostic Models**

Lorch and colleagues presented the results of a large retrospective international multicenter analysis conducted by the International Prognostic Factor Study Group to identify prognostic groups for initial salvage therapy independent of regimen intensity<sup>5</sup>. This is the largest series ever reported and included approximately 2000 patients from 38 centers throughout 14 countries in Europe and North America. Seven factors were found to be significant for PFS on multivariate analysis including and overall scores were divided into five groups (Table 2). This is widely considered the standard predictive model in the relapsed/refractory setting.

**Table 2.** Prognostic models: international prognostic factor study group score

| Factors                                                                                                                                                                                                 |                     | Points         |               |
|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---------------------|----------------|---------------|
| Primary site                                                                                                                                                                                            | Gonadal             | 0              |               |
|                                                                                                                                                                                                         | Retroperitoneal     | 1              |               |
|                                                                                                                                                                                                         | Mediastinal (NSGCT) | 3              |               |
| Response to first-line therapy                                                                                                                                                                          | CR/PR-              | 0              |               |
|                                                                                                                                                                                                         | PR+/SD              | 1              |               |
|                                                                                                                                                                                                         | PD                  | 2              |               |
| Progression-free interval after first-line therapy                                                                                                                                                      | > 3 months          | 0              |               |
|                                                                                                                                                                                                         | ≤ 3 months          | 1              |               |
| Serum hCG level                                                                                                                                                                                         | ≤ 1000 IU/l         | 0              |               |
|                                                                                                                                                                                                         | >1000 IU/l          | 1              |               |
| Serum AFP level                                                                                                                                                                                         | Normal              | 0              |               |
|                                                                                                                                                                                                         | ≤1000 ng/ml         | 1              |               |
|                                                                                                                                                                                                         | > 1000 mg/ml        | 2              |               |
| Liver, bone or brain metastases                                                                                                                                                                         | Absent              | 0              |               |
|                                                                                                                                                                                                         | Present             | 1              |               |
| <i>Add points for preliminary score (0-10); regroup into category score: (0):0; (1-2): 1; (3-4): 2; (5 or more): 3 add histology points as below to category score to determine final risk category</i> |                     |                |               |
| Histology                                                                                                                                                                                               | Seminoma            | -1             |               |
|                                                                                                                                                                                                         | NSGCT/mixed         | 0              |               |
| Stratification                                                                                                                                                                                          | Points              | 2-year PFS (%) | 3-year OS (%) |
| Very low risk                                                                                                                                                                                           | -1                  | 75             | 77            |
| Low risk                                                                                                                                                                                                | 0                   | 51             | 66            |
| Intermediate risk                                                                                                                                                                                       | 1                   | 40             | 58            |
| High risk                                                                                                                                                                                               | 2                   | 26             | 27            |
| Very high risk                                                                                                                                                                                          | 3                   | 6              | 6             |

AFP: α-feto protein; CR: Complete response; DFS: Disease-free survival; FFS: Failure-free survival; hCG: Human chorionic gonadotropin; NSGCT: Nonseminomatous germ cell tumour; OS: overall survival; PD: progression of disease; PFS: progression-free survival; PR-: partial response with negative markers; PR+: partial response with positive markers; SD: stable disease.

## Alberta Results

In a study of 43 patients with median age 28 (range 19-56) who received single (n=18) or tandem (n=25) HDCT and ASCT for relapsed metastatic GCT in Alberta between 2000-2018, 2-year PFS was 44% and OS was 65%.

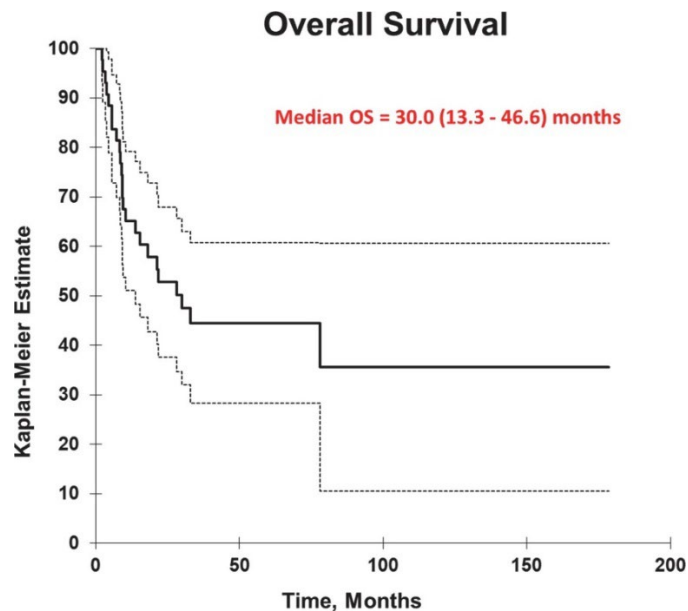


**Figure 3:** Overall survival after HDCT for GCT in Alberta



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# Complications

# Graft-vs-Host Disease

Presented by: Jan Storek

## Summary

- **Prophylaxis**

- For HLA mismatched donor ( $\leq 7/8$ , including haploidentical) HCT:
  - Posttransplant Cyclophosphamide (PTCy), 50 mg/kg on d +3 and 50 mg/kg on d +4,
  - Tacrolimus starting with 0.06 mg/kg on day +5, targeting trough levels of 5-15 ug/L until day 56, and tapering to zero by d 84, and
  - Mycophenolate mofetil (MMF) on d +5 through +35
- For HLA matched donor (8/8 unrelated or sibling) HCT:
  - Antithymocyte globulin (ATG, Thymoglobulin) 4.5 mg/kg,
  - Cyclosporine A (CSA) starting with 2.5 mg/kg IV bid on day -1, targeting trough levels of 200-400 ug/L until day 56, and tapering to zero by day 84, and
  - Methotrexate (MTX) on day 1, 3, 6 and 11.
  - Alternative: PTCy-based prophylaxis can be used with reduced-intensity conditioning (RIC). We use RIC+PTCy routinely for lymphoma/CLL.

- **Diagnosis** is made clinically, with the help of ancillary test like LFT, Histology, Schirmer's, or PFT

- **Acute vs Chronic GVHD** (Appendix 1 and 3):
  - If only skin/GI/liver manifestations of GVHD without a diagnostic sign of cGVHD (at any time since HCT), it is aGVHD.
  - If skin/GI/liver/eye/mouth/lung/genital manifestations of GVHD with a diagnostic sign of cGVHD (at any time since HCT), it is cGVHD.

- **Grading**

- aGVHD is graded per 1994 Consensus criteria
  - Grade 1 = Maculopapular rash covering <50% BSA
  - Grade 2-4 = Maculopapular rash covering >50% BSA or gut or liver involvement
- cGVHD is scored per 2014 NIH criteria
  - Mild = Max 2 organs involved to a mild degree, lungs not involved
  - Moderate-Severe = 3+ organs involved to a mild degree, or any organ involved to moderate or severe degree, or lungs involved (even to a mild degree)

- **Therapy (initial)**

- Grade 1 aGVHD or Mild cGVHD are treated topically or with observation only
- Grade 2-4 aGVHD is treated with prednisone 2–2.5 mg/kg/d or equivalent. If response, taper over 2-3 months.
- Moderate-Severe cGVHD is treated with prednisone 1 mg/kg/d or equivalent. If response, taper over 6-9 months.

- o For patients developing GVHD on a calcineurin inhibitor (CNI), the CNI is continued. For patients off CNI, CNI may be added to the steroid.
- **Next Line Therapy**
  - o Indications for moving from initial to next line therapy:
    - For aGVHD, worsening after 3-5 days, no improvement after 5-7 days, incomplete response after >28 days, or inability to taper methylprednisolone to <0.5 mg/kg/d or prednisone to <0.6 mg/kg/d
    - For cGVHD, worsening after 1-2 weeks, no improvement after 4-8 weeks of prednisone >0.5 mg/kg/d, or two failed attempts at prednisone taper necessitating prednisone dose increase to >0.25 mg/kg/d
  - o Second line therapy: Ruxolitinib, 10 mg bid orally
  - o Third line therapy for aGVHD is undefined. Modalities that can be tried include ECP, sirolimus, vedolizumab (for lower GI), fecal microbiota (e.g., MaaT013, for lower GI). Palliation should be considered.
  - o Third line therapy for cGVHD is undefined. Modalities that can be tried include belumosudil, axatilimab, ECP, rituximab, ibrutinib, sirolimus, imatinib, MMF, or MTX (no particular order of preference). Choose modalities according to their toxicities and other factors listed in Table 10.
  - o Clinical trial is always the preferred option as results with any next line therapy have been suboptimal.

## Background

Despite over 50 years of experience with allogeneic HCT, acute GVHD (aGVHD) remains the main cause of death of patients in remission after HCT. Chronic GVHD (cGVHD) is associated with not only mortality, but mainly poor quality of life long-term.

Risk factors may include HLA disparity, transplantation from an unrelated donor, female-to-male transplants, parity of female donor, recipient or donor age, peripheral blood stem cells (PBSC) vs marrow, or seropositivity for/reactivation of some herpes viruses. However, not all of these historical risk factors may apply to modern GVHD prophylaxis (e.g., PTCy or ATG). For example, per our center analysis of HCT recipients from HLA matched sibling donors (MSD) or 7-8/8 HLA matched unrelated donors (URD) whose GVHD prophylaxis was ATG-based, the only risk factors for aGVHD were donor type other than HLA matched sibling donor (non-MSD), lack of low-dose TBI (non-TBI), CMV donor negative/recipient positive serostatus (D-R+), and HCT for CLL or ALL, and the only risk factors for cGVHD were  $\leq 9/10$  HLA match, donor/recipient sex combination other than male/male, non-TBI, and HCT for myelofibrosis<sup>1</sup>.

Target organs for aGVHD are the skin, the liver, and the gut. Clinical features of skin disease range from localized erythematous rash to bullae and moist desquamation. Liver manifestations range from

mildly abnormal liver enzymes (predominantly cholestatic) to severe hyperbilirubinemia. Gut manifestations range from nausea, vomiting, and diarrhea to severe abdominal pain and ileus. Chronic GVHD may involve virtually any organ, most frequently the skin, the gut, the liver, the eyes, the mouth, the lungs, and genitalia. Details on the clinical manifestations are in Appendix 1.

## Prophylaxis of GVHD

In the 1990's, 2000's, and 2010's, the standard GVHD in most centers was a CNI (CSA or Tacrolimus) plus an antimetabolite (MTX or MMF/MPA). In the 2020's, this was superseded by ATG- or PTCy-based prophylaxis. Rationale:

- For HLA matched
  - ATG (Thymoglobulin) [Grafalon not reviewed here, as results of randomized studies have been contradictory]
    - CTTC trial, comparing ATG(4.5mg/kg)+CNI+antimetabolite vs CNI+antimetabolite
      - Setting:  $\geq 7/8$  URD (mostly 8/8), MAC 67%/RIC 33%, PBSC 88%/BM12%, med age 49 y, DRI not given but 33% late stage
      - 2-y GRFS 45 vs 25% ( $P=.003$ ),  $\uparrow$ QOL, OS 71% vs 53% ( $P=.02$ )
      - Post-hoc subgroup analysis: OS significantly different w vs w/o ATG for MAC, not for RIC (due to small number of patients?)
    - Limitation: Level A1 evidence only available for MAC
  - PTCy
    - US BMT CTN 1703<sup>2,3</sup>, comparing PTCy+Tacrolimus+MMF to Tacrolimus+MTX
      - Setting: Matched related or  $\geq 7/8$  URD (mostly 8/8), RIC, PBSC, med age 67 y, DRI high/very high 31%
      - 2-y GRFS 42% vs 29% ( $P=.001$ ) and  $\uparrow$ QOL. No significant difference in OS, RFS, relapse, or NRM.
    - Dutch (HOVON) trial<sup>4</sup>, comparing PTCy+CSA+MPA to CSA+MPA
      - Setting: MRD or MUD, NMA, PBSC, med age 58, DRI not given but probably not very high
      - 1-y cGRFS 45% vs 21% ( $P<.001$ ). No significant difference in OS, RFS, relapse, or NRM.
    - Limitations: Level A1 evidence only available for RIC. No impact on OS.
      - Phase 2 U Minnesota study of MAC (Flu 160 mg/m<sup>2</sup> + TBI 13.2Gy + Bu 12.8 mg/kg targeted to  $\sim 20,000$  uM.min/L; 32% high risk disease)<sup>5</sup>
        - 2-y GRFS 52%, 2-y OS 74%.
        - Low a/cGVHD and NRM, but relapse 39%.
  - PTCy and ATG appear equal, based on retrospective studies<sup>6</sup> and a prospective study<sup>7</sup>.
- For HLA mismatched
  - For haploidentical related HCT, the CNI+antimetabolite prophylaxis was associated with dismal outcomes. The advent of PTCy and ATG has made haploidentical HCT feasible.

PTCy-based prophylaxis has become the most frequently used option in N.America and Europe, whereas ATG(7.5-10mg/kg)-based prophylaxis in China<sup>8-11</sup>.

- For mismatched unrelated HCT, the CNI+antimetabolite prophylaxis was associated with suboptimal outcomes, and the addition of ATG improved the outcomes only to a minor degree. PTCy-based prophylaxis has been associated with the best outcomes, based on the following studies:
  - EBMT registry comparison of ATG-based with PTCy-based prophylaxis in 9/10 matched PBSCT (N>2000)<sup>12</sup>. 2-y OS was higher with PTCy (HR 0.77, P<0.001).
  - CIBMTR/NMDP multicenter prospective study, 4-7/8 matched URD BMT (N=80, 40 RIC and 40 MAC)<sup>13</sup>. 3-y OS 70% RIC, 62% MAC. 7/8 71% vs 4-6/8 63% (NS)!
  - Italian multicenter prospective single-arm study of PTCy-based prophylaxis in 7/8 matched PBSCT (N=77)<sup>14</sup>. 1-y OS 79%, RFS 55%.
  - “Closed gap between 7/8 and 8/8 URD HCT by PTCy, not so much by ATG
    - CIBMTR registry study (N=613 7/8, 1681 8/8)<sup>15</sup>. OS and GFRFS identical.
    - Albertan unpublished retrospective data on patients who underwent PBSC in 1999-2024 using Flu+Bu+4GyTBI+ATG(4.5mg/kg)+MTX+CSA (N=194 7/8, 759 8/8). OS not different (P=.53) but cGRFS lower in 7/8 (P=.02).

Thus, our standard GVHD prophylaxis is

- ATG + CSA + MTX for MSD and MUD HCT<sup>16</sup>, and
- PTCy + Tacrolimus + MMF for MMUD and haploidentical HCT.
  - The PTCy+Tacro+MMF prophylaxis can also be used for MSD or URD HCT if a patient cannot tolerate ATG (e.g., severe infusional reaction to the first dose) or MTX (e.g., effusions).

Other GVHD prophylaxes, used at other centers or evaluated in research, include

- CNI (CSA or Tacrolimus) + antimetabolite (MTX or MMF)
  - Time-honored GVHD prophylaxis for MSD and MUD HCT, used since 1980's, may still be the “gold standard” if paired with marrow graft (60% cGRFS and 84% OS in US BMT CTN 1301, not inferior to PTCy with marrow graft or CD34 selected PBSCs)<sup>17</sup>.
- Sirolimus + CNI + MTX or MMF
  - Has been evaluated only in the setting of nonmyeloablative HCT, where it was superior to CNI+MMF<sup>18</sup>. We do not use it as busulfan followed by sirolimus can result in a high incidence of liver venoocclusive disease/sinusoid obstruction syndrome<sup>19</sup> or microangiopathic hemolysis<sup>20</sup>.
- CD34 cell enriched graft, without any posttransplant immunosuppressive therapy (IST)
  - Associated with a high mortality due to infections and possibly relapse (retrospectively reviewed by Montoro et al<sup>21</sup>, prospectively studied in US BMT CTN



1301<sup>17</sup>. More studies are needed to determine whether this disadvantage is outweighed by the advantage of reducing aGVHD and cGVHD.

- Naïve T cell-depleted graft
  - Promising given low mortality due to infections and low incidence of cGVHD<sup>22</sup>, but more studies are needed to determine whether this advantage may be outweighed by increased incidence of relapse.
- Alpha/beta T cell-depleted graft, with or without B cell depletion
  - Promising in pediatric HCT<sup>23</sup>, more studies are needed in adult HCT.
- Regulatory T cell enrichment
  - Orca-T (Tregs+hematopoietic cells given on day 0, Tcons on day +2)
  - Preliminary 1 y results of randomized study against Tac+MTX:
    - cGRFS 78 vs 38% (P<.001), OS 94 vs 83% (P=.12)
  - Source: <https://aml-hub.com/medical-information/precision-t-a-phase-iii-trial-of-orca-t-vs-soc-for-cgvhd-prophylaxis-in-advanced-hematologic-malignancies>
- Ruxolitinib+CNI+MTX
  - Randomized study against PTCy+CNI+MMF to be launched (US BMT CTN 2203)
- Itacitinib (Jak-1 inhib) added to PTCy-based conditioning in Haplo
  - In a prospective Phase 2 study of 42 patients, no CRS, no GF, no gr 3-4 aGVHD, mod-sev cGVHD 5%, and 1-y OS 80%<sup>24</sup>.
- Improvements of ATG-based prophylaxis
  - Individualized dosing of ATG to target optimal AUC (pursued by Storek et al)
- Improvements of PTCy-based prophylaxis
  - Reducing Cy dose to 25-40 mg/kg on d 3 and 4
    - St.Antoine (Paris) compared 40 to 50 mg/kg d3&4 in patients >65 y (non-randomized)<sup>25</sup>
      - Faster engraftment, ↓cystitis, no impact on OS.
- Combination of PTCy + ATG (±CNI/other)
  - Meta-analysis of non-randomized studies in Haplo HCT suggested superiority of the combination vs PTCy or ATG in terms of ↓gr2-4 aGVHD, ↑GRFS, and ↑OS<sup>26</sup>.
  - Phase 2 randomized CTTC study (Walker and Paulsen et al, ASH 2024 abstract) showed reduction of gr3-4 aGVHD, but no difference in mod-sev cGVHD, and no difference in OS at 1 y.

Table 1: ATG + CSA + MTX details

| Drug          | Dose                    | Days                                                                                      | Route  |
|---------------|-------------------------|-------------------------------------------------------------------------------------------|--------|
| Cyclosporine* | 2.5 mg/kg every 12 h IV | -1 until oral feasible, then PO** every 12 h until day +56, then taper to zero by day +84 | IV, PO |
| Methotrexate  | 15 mg/m <sup>2</sup>    | Day +1                                                                                    | IV     |
|               | 10 mg/m <sup>2</sup>    | Day +3                                                                                    | IV     |
|               | 10 mg/m <sup>2</sup>    | Day +6                                                                                    | IV     |
|               | 10 mg/m <sup>2</sup>    | Day +11                                                                                   | IV     |
| Thymoglobulin | 0.5 mg/kg               | Day -2                                                                                    | IV     |
|               | 2 mg/kg                 | Day -1                                                                                    | IV     |
|               | 2 mg/kg                 | Day 0***                                                                                  | IV     |

\* Adjust dose to maintain trough plasma level 200 – 400 µg/L

\*\* Convert IV dose resulting in therapeutic trough levels to PO dose by multiplying the IV dose 2.5-times.

\*\*\* If day 0 is postponed by one day from the originally planned day 0 (e.g., because PBSCs are collected over two days instead of one day), the last dose of ATG is infused on the planned day 0, i.e., true day -1.

### Methotrexate Administration and Adjustment Guidelines

The first dose of methotrexate is given on day +1, at least 24 hours following infusion of stem cell product. Dosage adjustments will be made for renal and hepatic function, and for patients with severe mucositis or known fluid collections (pleural effusions or ascites). Dosage reductions between categories are additive: The final dosage reduction is the sum of dosage reductions for renal or hepatic dysfunction, mucositis, and fluid collections (below).

Table 2: Dosage adjustment for hepatic dysfunction\*

| Direct Bilirubin (micromoles/litre) | % Dose Reduction |
|-------------------------------------|------------------|
| < 34                                | 0                |
| 34-50                               | 25               |
| 51-100                              | 50               |
| > 100                               | 100              |

\* Hyperbilirubinemia purely due to CSA (negative abdominal ultrasound, no infection such as bacteremia) is not a reason for dose reduction

Table 3: Dosage adjustment for renal dysfunction

| Creatinine Clearance (mL/minute) | % Dose reduction |
|----------------------------------|------------------|
| >50                              | 0                |
| 40-50                            | 50               |
| <40                              | 100              |

**Mucositis.** Methotrexate should be completely withheld in the presence of severe mucositis defined as impending airway compromise. If no impending airway compromise, no reduction is needed.

**Fluid collection.** If clinically-significant fluid collections are present they should be drained. If they can be successfully drained, reduce methotrexate by 25%. If they cannot be drained, methotrexate should be withheld.

Folinic acid 5 mg IV q 6 h is given 24 hours after each dose of methotrexate, and continued until 12 h before the next dose of methotrexate or, in case of the last dose, until ANC>0.5/nL.

Table 4: PTCy + Tacrolimus + MMF details

| Drug             | Dose                            | Days             | Route            |
|------------------|---------------------------------|------------------|------------------|
| Cyclophosphamide | 50 mg/kg (actual, >30% -> AIBW) | +3, +4           | IV, in ½ L in NS |
| Tacrolimus       | 0.06 mg/kg (ideal) bid          | +5 until +56/84* | PO**             |
| MMF              | 1 g bid*                        | +5 until +35     | PO or IV         |
| MESNA            | 12.5 mg/kg qid (actual/AIBW)    | +3, +4           | IV               |

Abbreviations: AIBW = adjusted ideal body weight; NS = normal saline; IV = intravenous; PO = per oral; QID = 4 times a day.

\* Target trough plasma level of 5-15 ug/L until day 56, then taper to zero by day 84.

\*\* If patient cannot take PO, convert to IV, dividing the PO dose by 3.

\*\*\*If HCT is done for a non-malignant disease, 1 g tid may be used – follow pertinent publication

## Alternative if CNI, MTX, or MMF is not tolerated

### CNI Intolerance:

Up until 2022, our guideline recommended to switch a patient not tolerating CSA to prednisone at the doses given below (next paragraph). A retrospective review of local data in 2022 showed that patients who prematurely discontinued CNI due to intolerance and were switched to corticosteroid prophylaxis had increased risks of grade II-IV and grade III-IV acute GVHD and GVHD-related NRM compared to those who received continuous CNI prophylaxis<sup>27</sup>. Thus, in 2022 the guideline was changed to the following:

CNI prophylaxis should be continued whenever possible – the medical team should tolerate mild/moderate renal dysfunction. A less favored option is to target lower cyclosporine trough levels. This option should ideally be avoided between day 15 and 28, as targeting 200-400 appears important between day 15 and 28 whereas lower target may be acceptable before day 15 or after day 28<sup>28</sup>. For those with severe toxicities which necessitate discontinuation of CNI, MMF at 1 g BID should be added to corticosteroid according to the following schedule:

- Days 7-14 methylprednisolone 0.5 mg/kg IV
- Days 15-28 methylprednisolone 1 mg/kg IV
- Days 19-42 prednisone 0.5 mg/kg
- Days 43-56 prednisone 0.25 mg/kg
- Days 57-84 prednisone taper to zero and MMF taper to zero

Switching CSA to tacrolimus or vice versa is rarely useful. Studies comparing tacrolimus with CSA suggest that efficacy and toxicities are similar, except CSA may be associated with more grade 2-4 aGVHD but possibly less relapse and may be less nephrotoxic. OS is similar<sup>29-32</sup>. The studies have the limitation that the dose/therapeutic level of CSA corresponding to that of tacrolimus is unknown.

### MMF/MTX intolerance:

Switching MTX to MMF is reasonable. In comparative studies OS has been similar. MTX may be associated with less grade 2-4 or 3-4 aGVHD but greater toxicity (e.g., stomatitis) than MMF<sup>33-36</sup>. Toxicities common to both MTX and MMF are gastrointestinal (nausea, diarrhea) and cytopenias, both possibly more common with MMF than MTX but this may be related to the fact that the last day of administration is usually day 11 for MTX and day 35 for MTX. Possible reason of switching MTX to MMF is effusions.

Switching MMF to MTX is not recommended as, in the PTCy setting, the timing and dosing of MTX has not been studied. Taper or discontinuation of MMF (without switching to another drug) may be a better option as MMF in the PTCy setting may be redundant. Alternatively, corticosteroid until day 35 may be given at 50% above doses in addition to continuing the CNI. Similar approach can be considered for patients for whom total MTX dose needs to be reduced to <50%.

### Therapeutic Monitoring and Dosing of CNI

**Cyclosporine A (CSA)** trough plasma level target for GVHD prophylaxis is 200-400 ug/L until day 56, then taper to zero by day 84, providing there is no evidence of GVHD. For non-malignant indications (e.g., aplastic anemia), CSA taper is initiated on day 180.

CSA Neoral<sup>®</sup> for oral use is available as a capsule (10mg, 25mg, 50mg, 100mg) and as an oral solution (100mg/ml). CSA Sandimmune<sup>®</sup> is for IV use. Initial dose is 2.5mg/kg IV q12h or 6.25mg/kg PO q12h if starting on day -1. This dosing typically results in levels higher than 400 ug/L, which requires subsequent dose reduction. If CSA is started later (e.g., after day 84 for new onset grade 2-4 aGVHD or moderate-severe cGVHD), 2.0-2.5 mg/kg PO typically results within three days in the therapeutic level of 200-400 ug/L.

Conversion of IV to oral requires a 2.5 to 3-fold increase in dosage. The following algorithm can be utilized in guiding dose adjustment:

Table 5:

| CSA level    | Adjustment                                           |
|--------------|------------------------------------------------------|
| <200 ug/L    | Increase by 25%                                      |
| 200-400 ug/L | No change                                            |
| 350-400 ug/L | Consider decreasing by 25% if level trending upwards |
| 400-450 ug/L | Decrease by 25%                                      |
| >450 ug/L    | Hold 1-2 doses, decrease by 25-50%                   |

Inpatient, trough levels are drawn three times a week. If infused intravenously, CSA blood specimen should not be drawn from the same line used for administration. Outpatient, levels are drawn weekly, at a minimum. Consider repeating levels after 2-4 doses after a dose adjustment or the initiation/discontinuation of an interacting medication. For cGVHD, once maintenance dose is established for patients on long term CSA, frequency of trough level collection may be decreased to a monthly or as needed basis. In addition to monitoring drug levels, regular monitoring should also include blood pressure, CBC, serum electrolytes (Mg, K), renal function, hepatic function, and CMV and EBV PCR. Lower than the above recommended target levels may be used in case of renal or hepatic impairment, except for pure hyperbilirubinemia due to CSA.

CSA is a substrate and inhibitor of CYP3A4 and P-glycoprotein. Additional monitoring and dose adjustment may be required when starting or stopping drugs that inhibit or induce CYP3A4. Renal function should be closely monitored with co-administration of drugs that might exhibit additive/synergistic nephrotoxicity with CSA.

Patients are reminded to take CSA consistently with or without food to minimize variability. Capsules should be kept in the foil packaging until dose is ready to be taken. Patients are asked to leave capsules open to the air for no more than 15 minutes if needed to tolerate CSA's characteristic smell. Oral solution should be diluted in a glass container. Plastic and styrofoam containers should not be used. Orange juice and apple juice are recommended diluents by the manufacturer. Chocolate milk has also been used. Grapefruit and pomegranate juice should be avoided due to their interaction with the CYP450 system. The provided syringe can be wiped clean, but not washed as it may result in dose variation.

**Tacrolimus** trough plasma level target for GVHD prophylaxis/treatment is 5-15 ug/L. Routine taper (in the absence of GVHD) is the same as for CSA, i.e., from day 56 with the goal of reaching zero by day 84.

Tacrolimus (Prograf®) is available for oral use as an immediate release capsule (0.5mg, 1mg or 5mg) and for IV use. A 1mg/ml oral suspension can also be compounded. Advagraf® extended release capsules are not recommended for HCT setting. Initial dose of Prograf is 0.03 mg/kg/d IV as a continuous infusion or 0.06 mg/kg/day PO q12h if starting on day -1 or in the first several days after HCT. This dosing typically results in levels higher than 15 ug/L, which requires subsequent dose reduction. If tacrolimus is started later (e.g., after day 84 for new onset grade 2-4 aGVHD or moderate-severe cGVHD), 0.02 mg/kg IBW PO q12h typically results within three days in the therapeutic level of 5-15 ug/L.

Conversion of IV to oral requires a 2.5-4 fold increase in dosage.

The following algorithm can be utilized in guiding dose adjustment:

$$\text{New dose} = \frac{(\text{current dose})(\text{target whole blood level})}{(\text{current whole blood level})}$$

As for CSA, tacrolimus blood specimens should not be drawn from the same line used for tacrolimus administration. Levels are drawn three times a week inpatient and at least once weekly outpatient, less frequently after stable levels in case of cGVHD. Consider repeating levels after 2-4 doses after a dose adjustment or the initiation/discontinuation of an interacting medication. In addition to monitoring the levels, regular monitoring should also include blood pressure, blood glucose, CBC, serum electrolytes (Mg, K), renal and hepatic function, and CMV and EBV PCR. Lower than the above recommended target levels may be used in case of renal or hepatic impairment.

Tacrolimus is a substrate of CYP3A4 and p-glycoprotein. Additional monitoring and dose adjustment may be required when starting or stopping drugs that inhibit or induce CYP3A4. Renal function should be closely monitored with co-administration of drugs that might exhibit additive/synergistic nephrotoxicity with tacrolimus.

Patients are reminded to take tacrolimus consistently with or without food to minimize variability. Grapefruit and pomegranate should be avoided due to their interaction with the CYP450 system.

Falsely low tacrolimus levels may be observed in patients with severe anemia<sup>37</sup>. This is because tacrolimus levels are measured in whole blood, but only free (not RBC-bound) tacrolimus is active. Thus, in severe anemia, free tacrolimus concentration may be high despite normal or low whole blood concentration. If a patient with normal or low whole blood levels develops neuro or nephrotoxicity despite normal or low trough whole blood levels, tacrolimus dose should be decreased.



## Grading of aGVHD

For aGVHD grading, we use the 1994 Consensus Conference grading system<sup>38</sup>. To determine the overall grade of aGVHD, organ stages need to be determined first:

Table 6: Organ Staging of aGVHD

| Organ Stage | Skin                                                       | Liver                       | Gut                                                         |
|-------------|------------------------------------------------------------|-----------------------------|-------------------------------------------------------------|
| 0           | No rash                                                    | Total bilirubin < 34 umol/L | No diarrhea                                                 |
| 1           | Maculopapular rash <25% body surface area                  | Total bilirubin 34 to 50    | Diarrhea 500-1000 mL/day or nausea with positive UGI biopsy |
| 2           | Maculopapular rash 25 – 50% body surface area              | Total bilirubin 51 to 100   | Diarrhea 1000- 1500 mL/day                                  |
| 3           | Maculopapular rash > 50% body surface area                 | Total bilirubin 101 to 250  | Diarrhea 1500- 2000 mL/day                                  |
| 4           | Generalized exfoliative, ulcerative, or bullous dermatitis | Total bilirubin ≥250        | Diarrhea >2000 mL/day or severe abdominal pain or ileus     |

Table 7: Grading of aGVHD (determination of overall grade based on organ stages)

| Grade | Stage |     |       |     |       |
|-------|-------|-----|-------|-----|-------|
|       | Skin  |     | Liver |     | Gut   |
| 0     | 0     | And | 0     | And | 0     |
| I     | 1-2   | And | 0     | And | 0     |
| II    | 3     | Or  | 1     | Or  | 1     |
| III   | 0-3   | And | 2 - 3 | Or  | 2 – 4 |
| IV    | 4     | Or  | 4     |     | any   |

**Frequency of aGVHD grading:** At initial diagnosis, at worsening, and generally once a week in the first 3 months postHCT.

## Grading (“Scoring”) of cGVHD

For cGVHD scoring, we use the 2014 NIH scoring system<sup>39</sup>:

Table 8: Organ Scores of cGVHD:

|                            | Score 0                      | Score 1                                                                                                                          | Score 2                                                                                                                                                      | Score 3                                                                                                                                                                                    |
|----------------------------|------------------------------|----------------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>SKIN</b>                | No symptoms                  | <19% BSA with disease signs but NO sclerotic features                                                                            | 19-50% BSA OR involvement with superficial sclerotic features “not hidebound” (able to pinch)                                                                | >50% BSA OR deep sclerotic features “hidebound” (not able to pinch) OR impaired mobility, ulceration, or severe pruritis                                                                   |
| <b>MOUTH</b>               | No symptoms                  | Mild symptoms with disease signs but not limiting oral intake significantly                                                      | Moderate symptoms with disease signs with partial limitation of oral intake                                                                                  | Severe symptoms with disease signs on examination with major limitation of oral intake                                                                                                     |
| <b>EYES</b>                | No symptoms                  | Mild dry eye symptoms not affecting ADL (requiring eyedrops ≤ 3 times daily) OR asymptomatic signs of keratoconjunctivitis sicca | Moderate dry eye symptoms partially affecting ADL (requiring drops >3 times daily or punctal plugs) WITHOUT vision impairment                                | Severe dry eye symptoms significantly affecting ADL (special eyewear to relieve pain) OR unable to work because of ocular symptoms) OR loss of vision caused by keratoconjunctivitis sicca |
| <b>GI TRACT</b>            | No symptoms                  | Symptoms such as dysphagia, anorexia, nausea, vomiting, abdominal pain or diarrhea without significant weight loss (<5%)         | Symptoms associated with mild to moderate weight loss (5-15%)                                                                                                | Symptoms associated with significant weight loss >15%, requires nutritional supplement for most calorie needs OR esophageal dilation                                                       |
| <b>LIVER</b>               | Normal LFT                   | Bilirubin (total) normal<br>ALT 180-300 U/L<br>ALP ≥429 U/L                                                                      | Bilirubin 24-72 umol/L<br>ALT >300 U/L                                                                                                                       | Bilirubin >72 umol/L                                                                                                                                                                       |
| <b>LUNGS</b>               | No symptoms<br><br>FEV1 >80% | Mild symptoms (shortness of breath after climbing 1 flight of steps), or<br><br>FEV1 60-79%                                      | Moderate symptoms (shortness of breath after walking on flat ground), or<br>FEV1 40-59%                                                                      | Severe symptoms (shortness of breath at rest requiring O <sub>2</sub> ), or<br><br>FEV1 ≤39%                                                                                               |
| <b>JOINTS &amp; FASCIA</b> | No symptoms                  | Mild tightness of arms or legs, normal or mild decreased range of motion AND not affecting ADL                                   | Tightness of arms or legs OR joint contractures, erythema thought due to fascitis, moderate decreased range of motion AND mild to moderate limitation of ADL | Contractures WITH significant decrease of range of motion AND significant limitation of ADL (unable to tie shoes, button shirt, dress self)                                                |
| <b>GENITAL TRACT*</b>      | No symptoms/signs            | Mild signs*                                                                                                                      | Moderate signs*                                                                                                                                              | Severe signs*                                                                                                                                                                              |

**\* Genital signs:**

Female genitalia:

- 1) Mild (any of the following); erythema on vulvar mucosal surfaces, vulvar lichen-planus or vulvar lichen-sclerosus.
- 2) Moderate (any of the following); erosive inflammatory changes of the vulvar mucosa, fissures in vulvar folds.
- 3) Severe (any of the following); labial fusion, clitoral hood agglutination, fibrinous vaginal adhesions, circumferential fibrous vaginal banding, vaginal shortening, synechia, dense sclerotic changes, and complete vaginal stenosis.

Male genitalia:

- 1) Mild: lichen planus-like feature;
- 2) Moderate: lichen sclerosus-like feature or moderate erythema;
- 3) Severe: phimosis or urethral/meatal scarring.

Table 9: Global Score of cGVHD:

**Mild cGVHD:**

- $\leq 2$  organs involved with max organ score 1, plus lung score 0

**Moderate cGVHD:**

- $\geq 3$  organs involved with max score 1, or
- $\geq 1$  organ (not lung) with a score of 2, or
- Lung score 1

**Severe cGVHD:**

- $\geq 1$  organ with a score of 3, or
- Lung score 2 or 3

**Frequency of cGVHD scoring:** At initial diagnosis, at worsening, and at least every 3 months in the first year postHCT.

## Treatment of GVHD (initial)

For grade 1 aGVHD and mild cGVHD, we use a topical corticosteroid (e.g., betamethasone 0.1% cream bid). Observation-only is also reasonable. For a list of topical/ancillary therapies, see Appendix 2.

For grade 2-4 aGVHD and moderate-severe cGVHD, systemic corticosteroids remain the cornerstone of treatment. An exception may be sirolimus for aGVHD<sup>40</sup>, which should not be used in patients conditioned with busulfan<sup>19,20</sup>. Only  $\leq 50\%$  patients have a sustained complete or substantial partial response to a corticosteroid. Steroid-refractory or steroid-dependent patients usually have poor outcomes.

Dosing of systemic corticosteroids for initial therapy of grade 2-4 aGVHD and moderate-severe cGVHD has been derived from expert opinions. For aGVHD, we use 2.0-2.5 mg/kg/d prednisone (or equivalent), tapered over 2-3 months. For cGVHD, we use 1.0 mg/kg/d, tapered over 6-9 months. Attempts of lowering the initial dose of prednisone for grade 2-4 aGVHD have been associated with unexciting results. Per a randomized study of starting with 1 vs 2 mg/kg/d, the primary endpoint of lowering the cumulative prednisone dose by >33% was not reached as many patients who started with 1 mg/kg/d needed incrementation to 2 mg/kg/d<sup>41</sup>. However, this may not apply to grade 2a aGVHD.

Grade 2a aGVHD is defined as biopsy-confirmed UGI GVHD (nausea/vomiting/anorexia) with or without mild LGI GVHD (diarrhea <1 L/d), rash <50% BSA, and no hepatic involvement. It can be treated with 1 mg/kg/d prednisone + oral beclomethasone dipropionate 1 mg qid, with or without budesonide 3 mg bid (recommended if diarrhea)<sup>42,43</sup>. If no response in 10 days, increase prednisone to 2.0-2.5 mg/kg. If response in 10 days, taper prednisone over one week.

Adding an immunosuppressive drug to a steroid has been unsuccessful both for aGVHD and cGVHD<sup>44-48</sup>, except possibly for a CNI. The questionable benefit of adding a CNI to a steroid for first line treatment of cGVHD is based on expert opinions<sup>49</sup> and one randomized study of CSA+prednisone vs prednisone alone<sup>50</sup> which showed prednisone-sparing effect of the addition of CSA to prednisone, associated with a lower incidence of avascular necrosis, but no impact on cGVHD-related outcomes or OS. In a small study, tacrolimus had a 39% response rate in steroid-refractory patients<sup>51</sup>. In practice, many patients are already on CNI for prophylaxis. For the patients who are no longer on CNI, in Alberta we use the addition of CNI to prednisone optionally.

## Next-Line Treatment of GVHD (Steroid-Refractory/Dependent)

Favorable outcomes following next-line (post-steroid) therapy have been infrequent with any therapy. Studies of next-line therapy suffer from small patient numbers, short-term follow up, lack of control arm, or cross-over design which makes the evaluation of impact on OS impossible. Thus, whether any additional treatment of steroid-refractory/dependent GVHD has a lasting benefit with acceptable toxicity has not been rigorously determined.

Only patients who failed systemic steroids should proceed to next-line therapy. Failure of systemic steroids has had multiple definitions. We use the EBMT-NIH-CIBMTR Task Force definition<sup>52</sup>, modified to a minor degree in REACH 2 and 3 trials<sup>53,54</sup>, i.e.,

- For aGVHD, worsening after 3-5 days, no improvement after 5-7 days, incomplete response after >28 days, or inability to taper methylprednisolone to <0.5 mg/kg/d (prednisone to <0.6 mg/kg/d)
- For cGVHD, worsening after 1-2 weeks, no improvement after 4-8 weeks of prednisone >0.5 mg/kg/d, or two failed attempts at prednisone taper necessitating prednisone dose increase to >0.25 mg/kg/d

**Ruxolitinib** (Jak1/2 inhibitor) is our first-choice as it is the only post-steroid failure therapy clearly shown to be efficacious in randomized studies (REACH2 for aGVHD, REACH3 for cGVHD)<sup>53,54</sup>. However, the efficacy has been modest. Moreover, it is not known whether ruxolitinib improves OS, as both REACH1 and 2 had a cross-over design. In both studies, about one third of patients randomized to the control arm crossed over to ruxolitinib.

In REACH2, ruxolitinib, 10 mg bid orally, was superior to standard care (other IST) for steroid-refractory grade 2-4 aGVHD. In this trial, 309 patients ≥12 years old were randomly assigned (1:1) to ruxolitinib versus the investigator's choice of therapy (mostly ECP). At day 28, ruxolitinib achieved superior rates of overall response (PR+CR) (62% vs 39%) and CR (34% vs 19%). Superiority of ruxolitinib was maintained at day 56 (40% vs 22% overall response). Treatment was discontinued in 72% patients receiving ruxolitinib and in 85% of patients in the control group; most discontinuation was due to lack of efficacy. The most common grade ≥3 toxicity with ruxolitinib was thrombocytopenia, and anemia, and possibly infections, particularly CMV. Despite impact of ruxolitinib on OS could not be evaluated due to the cross-over design, it is notable that 18-month OS was only ~40% in both arms (Table S6 of the published study<sup>53</sup>).

In REACH3, ruxolitinib, 10 mg bid orally, was superior to standard care (other IST) for steroid-refractory/dependent moderate-severe cGVHD. A total of 329 patients ≥12 years old were randomly assigned (1:1) to ruxolitinib versus the investigator's choice of therapy (mostly ECP). At 6 months, ruxolitinib achieved superior rates of overall response (50% vs 26%) and CR (7% vs 3%). Treatment was discontinued in 50% patients receiving ruxolitinib and in 74% patients in the control group; most discontinuation was due to lack of efficacy. The most common grade ≥3 toxicity with ruxolitinib was thrombocytopenia, neutropenia, and anemia. Despite impact of ruxolitinib on OS could not be evaluated due to the cross-over design, it is notable that 12-month OS was similar (81% in the ruxolitinib arm and 84% in the control arm).

Dose reduction to 5 mg bid or 5 mg qd is required for cytopenias or renal or hepatic impairment.

For patients who have a response, ruxolitinib may be tapered gradually after 2 months (in aGVHD) or after 6 mo (in cGVHD). It is important that ruxolitinib is tapered gradually rather than discontinued abruptly or reduced rapidly, because a "withdrawal syndrome" that resembles systemic inflammatory response syndrome may be seen when ruxolitinib is discontinued in myelofibrosis.

Non-response to ruxolitinib (steroid- and ruxolitinib-refractoriness) is defined as

- For aGVHD, progression (stage/grade or new organ involvement) after 5-10 days, lack of improvement after >14 days, or loss of response after initial improvement<sup>55</sup>.
- For cGVHD, there is no accepted definition. It is reasonable to stop ruxolitinib if progression after 2-4 weeks, lack of improvement after 6-12 weeks, or loss of response after initial improvement.

## Steroid- and ruxolitinib-refractory GVHD

- raGVHD
  - Palliation is a reasonable choice.
    - In a retrospective study of 48 steroid- and ruxolitinib-refractory patients, most of them treated with additional immunosuppressive modalities, median survival was only 21 days.
  - Unproven options include
    - Extracorporeal photopheresis (ECP)
      - ECP has been evaluated only in retrospective studies<sup>56</sup>. In the largest one (n=128)<sup>57</sup>, 77% patients achieved PR or CR. A higher response rate was noted for grade 2 (compared to 3-4) aGVHD. The response rates appear to be the best for skin-only aGVHD<sup>56</sup>.
      - In Alberta, we have not been able to reproduce the relatively high response rate in published studies.
    - Vedolizumab (for lower GI)
      - In a retrospective study (n=20), d56 ORR 25%, 6mo OS 35%<sup>58</sup>.
    - Fecal microbiota (e.g., MaaT013, for lower GI)
      - In Phase II study, GI-ORR 53%<sup>59</sup>.
    - Alpha-1 Antitrypsin
      - In Phase II study of 60 mg/kg iv 2x/wk (n=40), ORR 65%, CR 35%, 6-mo OS 45%<sup>60</sup>.
    - TNFalpha inhibitors (infliximab, etanercept)
      - In retrospective studies, ORR 35-55%, OS 41-69% at ≥6 mo<sup>55</sup>.
    - Sirolimus
      - In a prospective phase 2 study in steroid-refractory (n=31) or intolerant (n=3) patients, PR+CR was achieved in 76% patients<sup>61</sup>. Similar response rate (86%) was described in a small retrospective study (n=22)<sup>62</sup>.
      - Major side effects are hyperlipidemia, headache, poor wound healing, renal dysfunction, edema, cytopenias and hemolytic-uremic syndrome. Hemolytic-uremic syndrome is particularly problematic in patients treated concurrently with CNI.
      - Sirolimus should not be used in the first 1-2 months after transplant if busulfan was used for conditioning and methotrexate was used for aGVHD prophylaxis, due to the high incidence of VOD/SOS and possibly also HUS<sup>19,20</sup>.
      - Dose is initially 1-2 mg qd. The initial dose of sirolimus must be significantly reduced in patients concomitantly treated with azole or macrolide antibiotics. Levels should be drawn 5-7 days after a dose adjustment or the initiation/discontinuation of an interacting medication. Target therapeutic trough level is 5-15 ng/mL. The following algorithm can be utilized in guiding dose adjustment:



$$\text{New dose} = \frac{(\text{current dose})(\text{target whole blood level})}{(\text{current whole blood level})}$$

- cGVHD

- For cGVHD, multiple treatments have been suggested to be efficacious, however, the level of evidence is low or intermediate.
  - Thus, choice of treatment is based on access (e.g., poor access to ECP for patients living outside of major urban areas) and side effects.
- Options are listed in the table below (based on ref<sup>63,64</sup>).

Table 10: Treatment options for steroid-/ruxolitinib-refractory cGVHD:

| Treatment                | Mechanism of action                        | Side Effects (important)                               | Infection risk       | Interactions (notable)   | Dosing (initial)                                   | Comment                                             |
|--------------------------|--------------------------------------------|--------------------------------------------------------|----------------------|--------------------------|----------------------------------------------------|-----------------------------------------------------|
| Belumosudil              | ROCK2 inh*                                 | Edema, GI, ↓P<br>Fatigue                               | ↑                    | None                     | 200 mg qd po                                       |                                                     |
| Axatilimab               | CSF1R inh →<br>macroph depletion           | ↑LFTs, ↓P,<br>GI, Anemia,<br>Myopathy                  | ↑                    | None                     | 0.3 mg/kg<br>q2wk iv                               |                                                     |
| ECP                      | ?                                          | Line clots                                             | Line infections only | None                     | 2-3 sessions per week                              |                                                     |
| Rituximab                | B cell depletion                           | Infusion reactions (IV>PO)                             | Minor ↑              | None                     | 375 mg/m <sup>2</sup> iv x4 (1400 mg sc?)          |                                                     |
| Ibrutinib                | BTK inh*, IL2-inducible kinase inh*        | Fatigue, Diarrhea                                      | ↑                    | Azoles                   | 420 mg qd po                                       |                                                     |
| Sirolimus                | mTOR inh →<br>↓T cell proliferation        | Cytopenia, TMA, ↑TGC, ↑LFTs, Edema, Poor wound healing | ↑ Viral              | Azoles, Macrolides       | 1-2 mg qd po, See under aGVHD Re therap monitoring | Avoid starting w/i 1-2 mo post-Bu (VOD/SOS concern) |
| Imatinib                 | Tyrosine kinase inh*                       | Cytopenia, Edema                                       | Minor ↑              | Statins, CNIs, Sirolimus | 400 mg qd po                                       |                                                     |
| MMF or mycophenolic acid | ↓de novo purine synth on which Ly's depend | Cytopenia, GI                                          | ↑ Viral              | None                     | 1 g bid po                                         |                                                     |
| MTX (low dose)           | Complex                                    | Cytopenia, Cheilitis, Alopecia                         | Minor ↑              | None                     | 7.5 mg/m <sup>2</sup> po weekly                    | Avoid if ↑Cr or effusions                           |
| HCQ                      | Complex                                    | GI, retinal / corneal                                  | Minor ↑              | None                     | 200 mg qd/bid po                                   | Annual eye exam                                     |

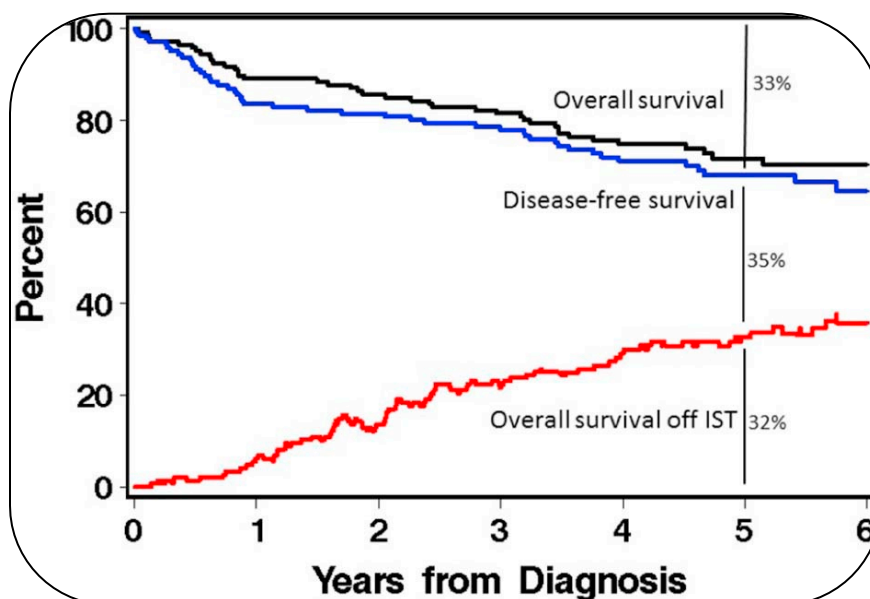
\* Inhibits signal transduction in lymphocytes (Ly's) ± other cells.

## Infection Prophylaxis

Patients with chronic GVHD are immunosuppressed and their treatment with IST makes their immunosuppression even more severe. As these patients are at an increased risk of opportunistic infections, adequate surveillance and prophylaxis is necessary (see sections on infection prophylaxis).

## Prognosis of Chronic GVHD

The prognosis continues to be poor. In a FHCRC study of 250 patients with moderate-severe cGVHD with median follow up >5 years, at 5 years since cGVHD diagnosis about 1/3 patients were dead, 1/3 patients alive and on systemic immunosuppressive therapy (IST), and 1/3 patients alive and off IST (Figure below)<sup>65</sup>.



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## Appendix 1: Signs and Symptoms of Chronic GVHD, including Common Signs/Symptoms for both Acute and Chronic GVHD (which are attributed to aGVHD unless a diagnostic symptom/sign of cGVHD)\*

| Organ/<br>Site                       | Diagnostic<br>(sufficient to establish<br>the diagnosis of<br>cGVHD)                                                                                                                                        | Distinctive<br>(seen in cGVHD but<br>insufficient alone to<br>establish a<br>diagnosis)                                                                                                                              | Other Features                                                                                                                                                                     | Common<br>(seen with both<br>aGVHD and<br>cGVHD)                                                                |
|--------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------|
| Skin<br><br>(see<br>photos<br>below) | <ul style="list-style-type: none"> <li>Poikiloderma</li> <li>Scleroderma / morphea</li> <li>Lichen sclerosus (morphea with overlying hypopigmented, finely wrinkled skin)</li> <li>Lichen planus</li> </ul> | <ul style="list-style-type: none"> <li>Depigmentation</li> <li>Papulosquamous lesions</li> </ul>                                                                                                                     | <ul style="list-style-type: none"> <li>Sweat impairment</li> <li>Ichthyosis</li> <li>Keratosis pilaris</li> <li>Hypo-pigmentation</li> <li>Hyper-pigmentation</li> </ul>           | <ul style="list-style-type: none"> <li>Erythema</li> <li>Maculopapular rash</li> <li>Pruritus</li> </ul>        |
| Nails                                |                                                                                                                                                                                                             | <ul style="list-style-type: none"> <li>Dystrophy</li> <li>Longitudinal ridging, splitting or brittle</li> <li>Onycholysis</li> <li>Pterygium unguis</li> <li>Nail loss (usually symmetric)</li> </ul>                |                                                                                                                                                                                    |                                                                                                                 |
| Scalp &<br>body<br>hair              |                                                                                                                                                                                                             | <ul style="list-style-type: none"> <li>New onset of scarring or nonscarring scalp alopecia (after recovery from chemoradiotherapy)</li> <li>Loss of body hair</li> <li>Scaling</li> </ul>                            | <ul style="list-style-type: none"> <li>Thinning scalp hair, typically patchy, coarse, or dull (not explained by endocrine or other causes)</li> <li>Premature gray hair</li> </ul> |                                                                                                                 |
| Mouth                                | <ul style="list-style-type: none"> <li>Lichen planus</li> </ul>                                                                                                                                             | <ul style="list-style-type: none"> <li>Xerostomia</li> <li>Mucocele</li> <li>Mucosal atrophy</li> <li>Ulcers</li> <li>Pseudomembranes</li> </ul>                                                                     |                                                                                                                                                                                    | <ul style="list-style-type: none"> <li>Gingivitis</li> <li>Mucositis</li> <li>Erythema</li> <li>Pain</li> </ul> |
| Eyes                                 | <ul style="list-style-type: none"> <li>New dry/gritty/painful eyes with Schirmer's test <math>\leq 5</math> mm,</li> <li>Keratoconjunctivitis sicca by slit lamp</li> </ul>                                 | <ul style="list-style-type: none"> <li>New onset of dry, gritty, or painful eyes</li> <li>Cicatricial conjunctivitis</li> <li>Keratoconjunctivitis sicca</li> <li>Confluent areas of punctate keratopathy</li> </ul> | <ul style="list-style-type: none"> <li>Photophobia</li> <li>Periorbital hyper-pigmentation</li> <li>Blepharitis (erythema of the eyelids with edema)</li> </ul>                    |                                                                                                                 |
| Genitalia                            | <ul style="list-style-type: none"> <li>Lichen planus or sclerosus</li> <li>Vaginal scarring/stenosis or clitoral/labial agglutination</li> <li>Phimosis or urethral/meatus scarring or stenosis</li> </ul>  | <ul style="list-style-type: none"> <li>Erosions</li> <li>Fissures</li> <li>Ulcers</li> </ul>                                                                                                                         |                                                                                                                                                                                    |                                                                                                                 |



| Organ/<br>Site            | Diagnostic<br>(sufficient to establish<br>the diagnosis of<br>cGVHD)                                                                                                                                                                          | Distinctive<br>(seen in cGVHD but<br>insufficient alone to<br>establish a<br>diagnosis)                     | Other Features                                                                                                                                                                                                                                      | Common<br>(seen with both<br>aGVHD and<br>cGVHD)                                                                                                                             |
|---------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| GI Tract                  | <ul style="list-style-type: none"> <li>Esophageal web</li> <li>Strictures/stenosis in the upper- to mid-third of the esophagus</li> </ul>                                                                                                     |                                                                                                             | <ul style="list-style-type: none"> <li>Exocrine pancreatic insufficiency</li> </ul>                                                                                                                                                                 | <ul style="list-style-type: none"> <li>Anorexia</li> <li>Nausea/Vomiting</li> <li>Diarrhea</li> <li>Weight loss</li> <li>Failure to thrive (infants and children)</li> </ul> |
| Liver                     |                                                                                                                                                                                                                                               |                                                                                                             |                                                                                                                                                                                                                                                     | <ul style="list-style-type: none"> <li>T.bilirubin or ALT or ALP &gt;2 times UNL</li> </ul>                                                                                  |
| Lung                      | <ul style="list-style-type: none"> <li>Bronchiolitis obliterans diagnosed with lung biopsy</li> <li>Bronchiolitis obliterans syndrome (BOS)* - diagnostic only if at least one distinctive manifestation of cGVHD in another organ</li> </ul> | <ul style="list-style-type: none"> <li>Air trapping / bronchiectasis on CT</li> </ul>                       | <ul style="list-style-type: none"> <li>Cryptogenic organizing pneumonitis</li> <li>Restrictive lung disease</li> </ul>                                                                                                                              |                                                                                                                                                                              |
| Muscles, fascia, joints   | <ul style="list-style-type: none"> <li>Fasciitis</li> <li>Joint stiffness or contractures secondary to sclerosis</li> </ul>                                                                                                                   | <ul style="list-style-type: none"> <li>Myositis or polymyositis (diagnostic if biopsy-confirmed)</li> </ul> | <ul style="list-style-type: none"> <li>Edema</li> <li>Muscle cramps</li> <li>Arthralgia or arthritis</li> </ul>                                                                                                                                     |                                                                                                                                                                              |
| Hemato-poietic and immune |                                                                                                                                                                                                                                               |                                                                                                             | <ul style="list-style-type: none"> <li>Thrombocytopenia</li> <li>Eosinophilia</li> <li>Lymphopenia</li> <li>Hypo- or hyper-gammaglobulinemia</li> <li>Auto-antibodies (AIHA, ITP)</li> </ul>                                                        |                                                                                                                                                                              |
| Other                     |                                                                                                                                                                                                                                               |                                                                                                             | <ul style="list-style-type: none"> <li>Pericardial or pleural effusions</li> <li>Ascites</li> <li>Peripheral neuropathy</li> <li>Nephrotic syndrome</li> <li>Myasthenia gravis</li> <li>Cardiac conduction abnormality or cardiomyopathy</li> </ul> |                                                                                                                                                                              |

\* BOS is defined as all of the following 4 criteria:

1. FEV1/FVC < 0.7
2. FEV1 <75% predicted (even post salbutamol/albuterol), or ≥10% decline over less than 2 years (even post salbutamol)
3. Absence of infection
4. CT evidence of air trapping or airway thickening/bronchiectasis, or PFT evidence of air trapping (RV >120% predicted).

*\*\* Adapted from Jagasia et al: BBMT 2015 (The 2014 Diagnosis and Staging Working Group Report)*

*Abbreviations: GVHD=graft-versus-host disease; ALT=alanine aminotransferase; AST=aspartate aminotransferase; BOOP=bronchiolitis obliterans-organizing pneumonia; PFTs=pulmonary function tests; AIHA=autoimmune hemolytic anemia; ITP=idiopathic thrombocytopenic purpura.*

### **Explanations of uncommon terms for mucocutaneous cGVHD**

**Lichen planus:** A skin eruption characterized in its most typical form by pruritic polygonal purple papules. These small flat-topped papules may show a white lacy network on their surface, Wickham's striae. The oral changes are characteristically erythema with a reticulate lacy pattern on the buccal mucosa. Erosions may also be present. The entire oral cavity may be involved, as can the genitalia of men and women.



**Poikiloderma:** A dermatosis characterized by variegated cutaneous pigmentation, atrophy, and telangiectasia.



**Morphea:** Morphea is a localized sclerosis of the skin. Early lesions typically show evidence of inflammation. A white firm plaque appears at the inflammatory site, surrounded by remaining inflammation. This plaque, over time, spreads peripherally and may become depressed. Telangiectatic vessels may be seen as well as hyperpigmentation.



## Appendix 2. Ancillary Therapy for Chronic GVHD

| Organ/Site                     | Prevention                                                                                                                                                                                                                                                          | Treatment                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      |
|--------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>Skin &amp; Appendages</b>   | <ul style="list-style-type: none"> <li>• photoprotection</li> <li>• surveillance for malignancy</li> </ul>                                                                                                                                                          | <ul style="list-style-type: none"> <li>• Emollients (Glaxal Base)</li> <li>• Corticosteroids (e.g., betamethasone valerate 0.1% cream/ointment, hydrocortisone 1% cream - for face)</li> <li>• Antipruritic agents (e.g., hydroxyzine 25 mg po TID - QID)</li> <li>• Erosions/ulcerations – microbiologic cultures</li> <li>• Topical antimicrobials (e.g. mupirocin)</li> <li>• Wound-care specialist consultation</li> </ul>                                                                                                                                                                                                                                                 |
| <b>Mouth &amp; Oral Cavity</b> | <ul style="list-style-type: none"> <li>• Good oral/dental hygiene</li> <li>• Routine dental cleaning</li> <li>• Surveillance for infection and malignancy</li> <li>• Fluoride (<i>Prevident</i> rinse; prescribed by dentist when there is oral dryness)</li> </ul> | <ul style="list-style-type: none"> <li>• High-potency corticosteroids: Dexamethasone 0.5mg/5mL solution, Budesonide 3mg/10ml solution, or Clobetasol 0.05% solution, swish and spit QID. Clobetasol can be also used as gel or ointment that can be applied to lesions.</li> <li>• Calcineurin inhibitors: cyclosporine 100 mg/mL solution swish + spit, tacrolimus 0.1% ointment</li> <li>• Therapy of oral dryness: <ul style="list-style-type: none"> <li>○ artificial saliva / lubricants (<i>Moi-stir</i>, <i>Oralbalance</i>, <i>Biotene</i>)</li> <li>○ salt water / baking soda or <i>Club soda</i> rinses</li> <li>○ pilocarpine 5-10mg po TID</li> </ul> </li> </ul> |
| <b>Eyes</b>                    | <ul style="list-style-type: none"> <li>• Photoprotection</li> <li>• Surveillance for infection, cataract and increased intraocular pressure</li> </ul>                                                                                                              | <ul style="list-style-type: none"> <li>• Artificial tears (<i>Refresh tears</i>; bottle or individual – preservative-free, <i>Bion tears</i> –one time use, <i>Systane</i>), thicker formulations (<i>Celluvisc</i>, <i>Gentle Gel</i>), artificial tears ointment (<i>Lacrilube</i>, qhs)</li> <li>• Corticosteroids: Prednisone 1% ophthalmic solution – <i>Pred Forte</i></li> <li>• Calcineurin inhibitors: cyclosporin, ophthalmic emulsion 0.05% (<i>Restasis</i>), prescribed by ophthalmologist</li> <li>• Pilocarpine 5-10mg po TID</li> </ul>                                                                                                                        |
| <b>Vulva &amp; Vagina</b>      | <ul style="list-style-type: none"> <li>• Surveillance for estrogen deficiency, infection (HSV, HPV, yeast, bacteria), malignancy</li> </ul>                                                                                                                         | <ul style="list-style-type: none"> <li>• Water-based lubricants (<i>KY jelly</i>, <i>Astroglide</i>, <i>Replens</i>)</li> <li>• Topical estrogens (<i>Premarin</i> - vaginal cream, <i>Vagifem</i> - vaginal tablet)</li> <li>• Corticosteroids: betamethasone – cream or enema</li> <li>• Dilatators</li> <li>• Surgery for extensive synechiae/obliteration</li> <li>• Early gynaecological consultation</li> </ul>                                                                                                                                                                                                                                                          |
| <b>GI tract &amp; liver</b>    | <ul style="list-style-type: none"> <li>• Surveillance for infection (viral, fungal)</li> </ul>                                                                                                                                                                      | <ul style="list-style-type: none"> <li>• Dietary modification</li> <li>• Corticosteroids: <ul style="list-style-type: none"> <li>○ upper GI – beclomethasone dipropionate oral solution 1mg/mL; 1mL po QID</li> <li>○ lower GI – budesonide 3 mg po TID</li> </ul> </li> <li>• Enzyme supplementation: pancreolipase (<i>Cotazym</i>, <i>Pancrease MT</i>, <i>Creon</i>, <i>Ultrase</i>, <i>Viokase</i>)</li> <li>• GI reflux management</li> <li>• Esophageal dilatation</li> <li>• Ursodeoxycholic acid (if pruritus due to cholestasis)</li> </ul>                                                                                                                          |
| <b>Lungs</b>                   | <ul style="list-style-type: none"> <li>• Surveillance for infection (PJP, viral, fungal, bacterial)</li> </ul>                                                                                                                                                      | <ul style="list-style-type: none"> <li>• inhaled corticosteroids: budesonide (<i>Pulmicort</i>), fluticasone (<i>Flovent</i>)</li> </ul>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       |

| Organ/Site              | Prevention                                                                                                                                                                                          | Treatment                                                                                                                                                                                                                                                                                                                       |
|-------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
|                         |                                                                                                                                                                                                     | <ul style="list-style-type: none"> <li>• SABA: salbutamol (<i>Ventolin</i>)</li> <li>• LABA: formoterol (<i>Oxeze</i>), salmeterol (<i>Serevent</i>)</li> <li>• Combo: formoterol + budesonide (<i>Symbicort</i>), salmeterol + fluticasone (<i>Advair</i>)</li> <li>• Anticholinergics: tiotropium (<i>Spiriva</i>)</li> </ul> |
| <b>Musculo-skeletal</b> | <ul style="list-style-type: none"> <li>• Surveillance for decreased range of motion</li> <li>• Bone densitometry</li> <li>• Calcium supplementation</li> <li>• Vitamin D supplementation</li> </ul> | <ul style="list-style-type: none"> <li>• Physical therapy</li> <li>• Treatment of osteoporosis, if present</li> </ul>                                                                                                                                                                                                           |

## Appendix 3. Summary of aGVHD grading, diagnosis of cGVHD (at least one diagnostic sign present), and cGVHD scoring

| aGVHD organ stage   | 1                | 2                          | 3                    | 4                    |
|---------------------|------------------|----------------------------|----------------------|----------------------|
| Skin                | <25% BSA         | 25-50%                     | >50% BSA             | Bullae               |
| Gut                 | 0.5-1 L/d or N/V | 1-1.5 L/d                  | 1.5-2 L/d            | >2 L/d or pain/ileus |
| Liver               | Bili 34-50       | Bili 51-100                | Bili 100-250         | Bili >250            |
| aGVHD overall grade | 1                | 2                          | 3                    | 4                    |
|                     | Skin 1-2         | Skin 3 or Gut 1 or Liver 1 | Gut 2-4 or Liver 2-3 | Skin 4 or Liver 4    |

| Diagnostic signs of cGVHD |                                                                                                              |
|---------------------------|--------------------------------------------------------------------------------------------------------------|
| Skin /fascia              | Poikiloderma, lichen, morphea/sclerosis/fasciitis/contractures                                               |
| Mouth                     | Lichen                                                                                                       |
| Eyes                      | New dry/gritty/painful eyes with Schirmer ≤5 mm, keratoconjunctivitis sicca by slit lamp                     |
| Genital                   | Lichen, ♀: vaginal scarring/stenosis, clitoral/labial agglutination, ♂: phimosis, urethral scarring/stenosis |
| GI                        | Esophageal web/stenosis                                                                                      |
| Lungs                     | Bronchiolitis obliterans diagnosed by biopsy or CT+PFTs                                                      |

| cGVHD organ score  | 1                                      | 2                                                                | 3                                                                |
|--------------------|----------------------------------------|------------------------------------------------------------------|------------------------------------------------------------------|
| Skin/fascia        | BSA <19%, no sclerosis                 | 19-50%, sclerosis but able to pinch, ↓ROM a/w mild impact on ADL | >50%, unable to pinch, ulcer, ↓ROM a/w severe impact on ADL      |
| Mouth              | Mild symptoms                          | Some oral intake limitation                                      | Major oral intake limitation                                     |
| Eyes               | Mild dryness                           | Some impact on ADL, no vision loss                               | Major impact on ADL, vision loss                                 |
| Genital            | Lichen planus (♂♀), Vulvar erythema    | Vulvar erosions or fissures (♀), Lichen sclerosus ♂              | Adhesions, dense sclerosis (♀), Phimosis (♂)                     |
| GI                 | Symptoms with <5% wt loss              | Symptoms with 5-15% wt loss                                      | >15% wt loss, esophag.dilation, suppl.feeding, diarrhea imp. ADL |
| Liver              | Bili normal<br>ALT 180-300<br>ALP ≥429 | Bili 24-72<br>ALT >300                                           | Bili >72                                                         |
| Lungs              | FEV1 60-79%,<br>DOE 1 flight up        | FEV1 40-59%,<br>DOE flat surface                                 | FEV1 <40%,<br>SOB at rest, O <sub>2</sub>                        |
| cGVHD global score | Mild (1)                               | Moderate (2)                                                     | Severe (3)                                                       |
|                    | 1-2 organs w max score 1, no lung      | All in between (including lung score 1)                          | Score 3 in any organ, or lung score 2-3                          |

# CMV, VZV, HSV, HHV6

Presented by: Jan Storek

## Summary

### CMV (cytomegalovirus) Disease Prevention in alloHCT

- *Monitoring and Preemptive therapy:*
  - Plasma CMV DNA by Q-PCR weekly until d100, then monthly until 1 year postHCT.
  - If CMV DNAemia >25,000 IU/mL plasma, treat preemptively with induction valganciclovir, 900 mg bid, until DNAemia has dropped, but for at least one week.
  - Continue valganciclovir at maintenance dose of 900 mg qd until DNAemia <5,000 IU/mL twice, but for at least 2 weeks.
- *Primary prophylaxis with Letermovir* should be used only in seropositive patients with seronegative donor (D-R+). Cord blood is considered D-.
  - Monitoring and preemptive valganciclovir should be used as above, except:
  - Extend weekly DNAemia monitoring until 2-3 months post letermovir discontinuation.
- *Blood products:* Use leukodepleted and irradiated blood products.
- *Donor selection for HCT:* Prefer a CMV serostatus-matched donor. This is of minor importance, except if ATG is used for GVHD prophylaxis in a seropositive patient undergoing alloHCT for a lymphoid malignancy – in this scenario choosing a seropositive donor is of major importance.

### VZV (Varicella Zoster Virus) and HSV (Herpes Simplex Virus) Disease Prevention in allo and autoHCT

- Valacyclovir, 500 mg qd, from start of conditioning until 1 day before the first dose of a live VZV vaccine (alloHCT) or until 1 month after the second dose of a non-live VZV vaccine (autoHCT).
- For *allogeneic* HCT recipients, VZV vaccination with a live vaccine should start at 2 years posttransplant or later. The later start is for patients on prolonged therapy with immunosuppressive drugs – wait until ≥3 mo after discontinuation of immunosuppressive therapy (systemic and topical) and no cGVHD activity.
- For *autologous* HCT recipients, VZV vaccination with a non-live vaccine (Shingrix) should start at 6 months posttransplant or later. The later start is for patients on prolonged maintenance immunosuppressive therapy – wait until ≥6 mo after discontinuation of a lymphodepleting antibody like rituximab.
- For CAR T cell recipients, management is currently identical to that of allogeneic HCT recipients.
- See Vaccination chapter for details.

### HHV6 (Human Herpes Virus 6) Disease Prevention: None.



## Cytomegalovirus (CMV)

**Epidemiology:** Incidences of CMV Reactivation / CMV Disease

- **Seronegative donor → Seronegative patient (D-R-)**

- 70% / 35% with non-CMV safe transfusions (from random donors and not leukodepleted).
- <2% / <2% with CMV safe transfusions
  - In Alberta, 0% CMV reactivation needing preemptive Rx / 0% CMV disease<sup>1</sup>

- **Seropositive donor → Seronegative patient (D+R-)**

- 70% / 35% with non-CMV safe transfusions
- 15% / 5% with CMV safe transfusions, before ganciclovir
- 10-15% / <3% with CMV safe transfusions, since ganciclovir
  - In Alberta, 10% CMV reactivation needing preemptive Rx / 1% CMV disease<sup>1</sup>

- **Seropositive donor → Seropos pt (D+R+), and Seroneg donor → Seropos pt (D-R+)**

- 70% / 35% before ganciclovir.
- 70% / <2%-20% since ganciclovir.
- In Alberta:
  - For MSD/URD with ATG-based GVHD prophylaxis:
    - In D+R+, 31% CMV reactivation needing preempt Rx / 4% CMV disease<sup>1</sup>
    - In D-R+, 64% CMV reactivation needing preempt Rx / 11% CMV disease<sup>1</sup>
      - With letermovir, 8/8 (50%) reactiv w preempt / 2/16 (12%) CMV dis
    - Survival lower in D-R+ than D+R+ patients with lymphoid malignancies (Fig. 1), due at least in part to a higher grade 3-4 aGVHD incidence in D-R+ patients<sup>2</sup>
  - For Haplos with PTCy/MMF-based GVHD prophylaxis (Storek, unpubl, 2023):
    - In D+R+, 16/25 (64%) CMV reactive→preempt Rx / 2/25 CMV dis (non-fatal)
    - In D-R+,
      - Without letermovir, 5/6 CMV reactiv→preempt / 2/6 CMV disease
      - With letermovir, 3/5 CMV reactiv→preempt / 0/5 CMV disease

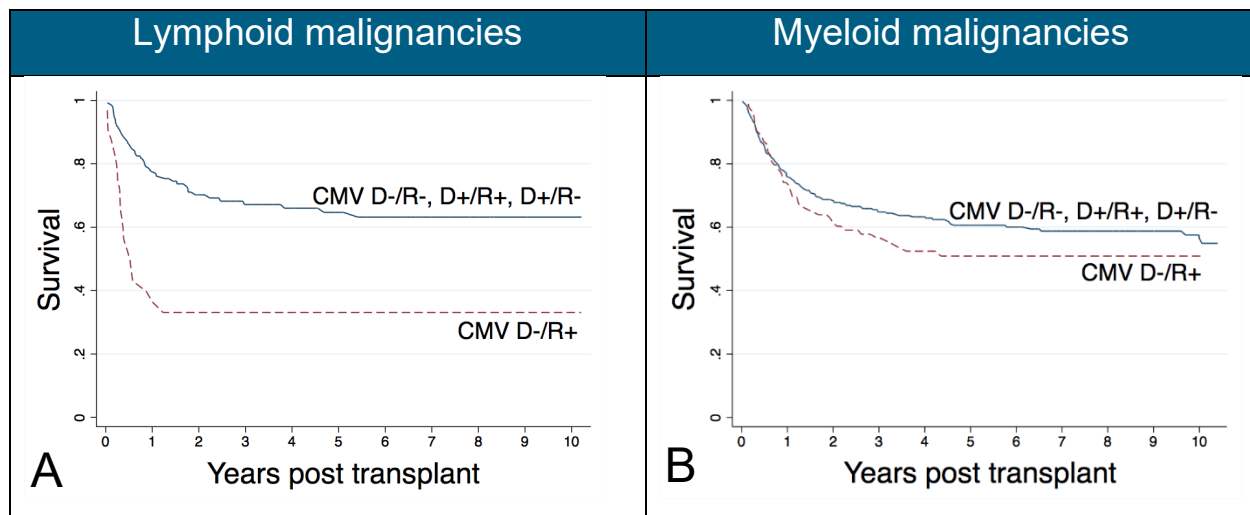


Figure 1. Impact of CMV serostatus on overall survival in patients with lymphoid malignancies (A) and patients with myeloid malignancies (B). In multivariate analysis, the differences were significant in patients with lymphoid malignancies ( $p=.001$ ) but not myeloid malignancies ( $p=0.23$ ).

- **Autologous seropositive patient**
  - 50% reactivation /  $\leq 2\%$  disease
- **Syngeneic seropositive patient**
  - 50% reactivation / 0% disease
- **Healthy individuals**
  - 50% - 80% are infected,  $<5\%$  reactivate (poorly studied), 0% develop CMV disease

#### Risk factors for CMV reactivation and disease:

- Seropositivity of recipient particularly if donor seronegative or cord blood
- T cell depletion, particularly ex vivo
- GVHD / immunosuppressive drugs
- Haploidentical HCT using PTCy+MMF+CNI

#### Clinical manifestations of CMV disease:

- Frequent: Gastroenteritis, Pneumonia
- Less frequent: Retinitis, Encephalitis, Hepatitis, Marrow suppression

## Prevention of CMV Disease

### Transfusions, and Hematopoietic Cell Donor Selection

- All blood products collected in Canada are leuko-depleted at the time of collection (CMV safe). Moreover, blood products for HCT recipients are routinely irradiated prior to transfusion.
- CMV seronegative HCT donor is preferred for CMV seronegative recipient.<sup>3</sup> However, the difference in survival of seropositive patients receiving grafts from seropositive vs negative donors is minor, if any.<sup>1,4</sup>
- CMV seropositive HCT donor is preferred for CMV seropositive recipient.<sup>1,3,5,6</sup>
  - Survival difference (between HCT from seropositive vs seronegative donors) in the setting of ATG-based GVHD prophylaxis is marked in patients with lymphoid malignancies (HR=3.1, p=0.001) whereas minor, if any, in patients with myeloid malignancies (HR=1.2, p=0.23) (Fig. 1).<sup>2</sup>
  - The survival difference is virtually zero in the setting of haploidentical HCT with posttransplant cyclophosphamide based GVHD prophylaxis.<sup>7</sup>
  - In patients with lymphoid malignancies and ATG-based GVHD prophylaxis, if an HLA matched but CMV seronegative sibling donor is available and no HLA matched and CMV seropositive sibling donor is available, search for an HLA matched and CMV seropositive unrelated donor is recommended.<sup>1</sup>
- Recipient CMV serostatus should ideally be determined before blood product transfusions, particularly platelet or plasma transfusions or IVIG. If CMV IgG is transferred from a CMV seropositive blood donor to a CMV seronegative recipient, the recipient may become falsely CMV seropositive.<sup>8</sup>

### Antiviral Prophylaxis or Preemptive Therapy?

- Prophylaxis with val/ganciclovir, foscarnet, brincidofovir, or maribavir decreased the incidence of CMV disease modestly or not at all (if yes, then only during the period of taking the drug) and had no impact on OS due to poor risk:benefit ratio.<sup>9,10</sup>
- Prophylaxis with high dose val/acyclovir
  - May be effective and safe,<sup>11,12</sup> but randomized studies do not exist.
- Prophylaxis with anti-CMV T cells
  - Efficacious and safe, but too costly.
  - Donor-derived anti-CMV T cells cannot be used in D-R+ patients who would most benefit from the prophylaxis.
  - Prophylaxis with 3<sup>rd</sup> party anti-CMV T cells is less efficacious, more data is needed to determine whether the benefit is worth the cost.
- Preemptive Therapy with val/ganciclovir
  - Never evaluated in a randomized study against no prophylaxis/no preemptive Rx.

- In a randomized study of ganciclovir prophylaxis till d100 vs preemptive Rx<sup>13</sup>, CMV disease incidence was similar, toxicity was worse with preemptive Rx (fungal infections, neurotoxicity, no difference in neutropenia), OS was similar.
  - Nevertheless, preemptive Rx has become popular based on single-arm studies showing less CMV disease than expected with no prophylaxis, not more CMV disease than with prophylaxis, and improved safety compared to prophylaxis.
- Prophylaxis with letermovir + Pre-emptive therapy with val/ganciclovir
  - Safe but efficacy is limited – ~halving of % patients with clinically significant (cs) CMV infection (CMV DNAemia requiring preemptive therapy or CMV disease)<sup>14,15</sup>.
  - Pros:
    - Potential reduction of incidence/duration of val/ganciclovir-induced toxicity like neutropenia.
      - This may not be substantial with our preemptive strategy, as in 2012-2022, 256 of 850 alloHCT recipients needed preemptive therapy which lasted only median 27 days (too short to cause clinically significant neutropenia), and only 53 of the 256 recipients needed ≥1 additional preemptive therapy course (Storek, unpublished).
    - Survival benefit?
      - Shown only in a meta-analysis of retrospective studies, and not in a subgroup meta-analysis of US-based studies.<sup>16</sup> Not shown in a randomized study, albeit there was a trend toward improved OS.<sup>15</sup> The lack of clear-cut improvement of OS may have been related to a high incidence of csCMV infection after letermovir discontinuation (after d100).
        - A subsequent randomized study comparing letermovir until d100 vs until d200 did not show any survival benefit (not even a trend – OS 92% vs 92% in patients who took letermovir in the first 100 days and started letermovir vs placebo on ~d100).<sup>17</sup> Interestingly, csCMV infection incidence was similar in both arms, except in the “until d200” arm most csCMV infections occurred after d200 whereas in the “until d100” arm most csCMV infections occurred between d100 and d200.
      - No survival benefit shown in the largest study so far (Japanese registry study of >6,000 patients).<sup>18</sup>
      - In a large single center study (Lille, N=316), there was a trend toward OS benefit in high-risk but not in low-risk patients.<sup>19</sup>
    - GVHD incidence/severity reduction?
      - So far theoretical. Practically reported in only one retrospective study for cGVHD (but not for aGVHD),<sup>20</sup> despite there have been >40 studies comparing letermovir to no letermovir.

- Non-CMV infection incidence/severity reduction? (as CMV seropositivity/infection is a/w [predisposes to?] non-CMV infections, e.g., Covid<sup>21</sup>)
    - So far only theoretical.
- Cons:
  - Cost (\$>20,000 for 3 months)
    - Inpatients require STEDT approval
    - Outpatients are usually covered 70-80% by their insurance.
      - The 20-30% copayment may be too much, but co-pay assistant cards are available.
  - No impact on the incidence of CMV disease<sup>15</sup>
  - No activity against HSV and VZV, so val/acyclovir needed in addition to letermovir
  - CMV DNAemia monitoring with preemptive VGCV needed in addition to letermovir
  - CMV reactivation after end of prophylaxis<sup>22,23</sup> – extended weekly DNAemia monitoring needed (until 2-3 mo post letermovir discontinuation)
- Alberta approach:
  - Preemptive therapy with val/ganciclovir is currently the CMV disease prevention strategy of choice in Alberta
  - Letermovir prophylaxis (primary), together with CMV DNAemia monitoring and preemptive val/ganciclovir, should be considered in D-R+ patients.
    - Rationale
      - D-R+ serostatus is the most important risk factor for csCMV infection and for mortality (clear-cut in pts with lymphoid malignancies, trend in patients with myeloid malignancies) in most AB patients (MSD/URD, ATG-based GVHD prophylaxis)<sup>1,2</sup> and probably also in Haplo setting with PTCy/MMF-based GVHD prophylaxis (Storek, unpublished), so these patients deserve to be offered the potential benefit of letermovir, albeit the benefit is uncertain.
      - For patients other than D-R+, the above uncertain pros do not outweigh the cons.
    - Financial implications should be clarified / discussed with the patient.
    - Weekly CMV DNAemia monitoring should be extended until 2-3 months post letermovir discontinuation.

## Alberta Preemptive approach - Details

- CMV DNA monitoring in plasma from day 0 to day 100 weekly, then monthly to 1 year posttransplant.
- Monitor all patients (including CMV seronegative patients with seronegative donors, as there is a small chance that the CMV IgG test result is falsely negative).
- If 5,000-25,000 IU/mL, repeat DNAemia in 3-7 days.
- If >25,000 IU/mL, start preemptive treatment with ganciclovir or valganciclovir.
  - In 1999-2007 we used a threshold of 10-20 pp65 antigen positive cells per slide (containing ~200,000 granulocytes) and found it to be satisfactory (~2.8% cumulative incidence of CMV disease and no CMV pneumonia in D+R+ patients (ATG-conditioned)). Between 2007 and 2012, we used a ProVLab in-house real time PCR assay and a threshold of 50,000 genome copies/mL plasma, which corresponded to the previous pp65 antigenemia threshold. Since 2012, we have used commercial real time PCR assay (RealStar, Altona) and a threshold of 25,000 IU/mL plasma, which corresponded to the previous in-house real time PCR threshold.
- Induction with valganciclovir 900 mg p.o. BID (or ganciclovir 5 mg/kg IV BID) for at least one week. Continue induction until a down-going trend of CMV DNAemia, then switch to maintenance (QD). For example, if ganciclovir induction was started for 80,000 IU/mL, switch to maintenance after <80,000 IU/mL.
  - If DNAemia has not declined after 2-3 weeks of induction, suspect ganciclovir resistance.
- Maintenance with valganciclovir 900 mg p.o. QD (or ganciclovir 5 mg/kg IV QD) for at least 2 weeks.
  - Treat until <5,000 IU/mL at least twice, but treat for a total of at least 3 weeks (e.g., at least one week of induction and at least 2 weeks of maintenance).
  - Prolonged maintenance/secondary prophylaxis with VGCV can be considered for patients at high risk of recurrent CMV disease (e.g., active GVHD AND history of ≥2 episodes of csCMV infection).
    - Letermovir may be considered for the prolonged maintenance/secondary prophylaxis instead of val/ganciclovir, however, only if CMV DNAemia has become undetectable, as otherwise there is a high likelihood of developing letermovir resistance.<sup>24</sup>
      - Secondary prophylaxis with letermovir is not covered by STEDT or Alberta Blue Cross (except possibly within 100 days postHCT).
- Both ganciclovir and valganciclovir doses need to be adjusted in renal insufficiency.
- If ANC<1.0/nL, give filgrastim. If ANC has not increased to >1.0/nL within 3 days, switch ganciclovir to foscarnet.
  - Alternatively, give maribavir 400 mg bid orally.
    - Efficacy of maribavir is slightly inferior compared to valganciclovir, but it does not appear to cause neutropenia.<sup>25</sup> Main side effect is dysgeusia.



- Maribavir is not active against VZV, so val/acyclovir prophylaxis should be continued.
- If preemptive treatment is given between 3 and 12 months posttransplant, check CMV DNAemia weekly. Resume monthly monitoring after DNAemia has been undetectable at least twice.

### Alberta Letermovir primary prophylaxis – Details

- Use only in high-risk patients (typically only D-R+)
  - Clarify financial implications with patient before starting. Some patients may choose not to start.
- Start on day 9 (the median start in Marty's randomized study<sup>15</sup>)
- Dose: 480 mg qd when off CSA, 240 mg qd orally when on CSA
  - No dose reduction with tacrolimus
- Continue until day 84 (the median in Marty's randomized study was d82)
  - No survival benefit of extending letermovir to d200 shown in a randomized study.<sup>17</sup>
- Stop letermovir when starting pre-emptive therapy or therapy with val/ganciclovir
  - Restarting after val/ganciclovir discontinuation is not recommended. There is a lack of data whether letermovir restarting is of any value.
- CMV DNAemia monitoring weekly should be extended until 2-3 months post letermovir discontinuation!

## Diagnosis and Therapy of CMV Disease

### Diagnosis of CMV disease

- Diagnosis of CMV Enteritis requires histological or immunohistochemical evidence. PCR positivity alone is not sufficient for diagnosis.
- Diagnosis of CMV Pneumonia in the past required positive viral culture of BAL. Viral cultures were discontinued in 2015 and replaced with PCR. PCR has an excellent negative predictive value (>99%) but a poor positive predictive value (cannot distinguish CMV pneumonia from pulmonary CMV shedding). Data on BALs with concurrent viral culture and PCR were analyzed by Dr.R.Tellier of ProVLab in 2015, showing:

| Viral load range (IU/mL)             | Neg (0 to <150) | 150 to 10 <sup>3</sup> | 10 <sup>3</sup> to 10 <sup>4</sup> | 10 <sup>4</sup> to 10 <sup>5</sup> | >10 <sup>5</sup> |
|--------------------------------------|-----------------|------------------------|------------------------------------|------------------------------------|------------------|
| % viral culture positive (pos/total) | 0.33% (1/306)   | 6% (2/33)              | 27% (6/22)                         | 40% (4/10)                         | 100% (4/4)       |

- Based on this data and the fact that pulmonary CMV shedding predisposes to CMV pneumonia, the diagnostic and therapeutic algorithm is as follows:
  - If CMV  $>10^3$  IU/ml BAL, CMV pneumonia is possible/proven. Treat as CMV disease (below).
  - If CMV between 150 (detection limit) and  $10^3$  IU/ml BAL, CMV pneumonia is unlikely. Treat the CMV shedding with 1 week induction and 1-2 weeks maintenance.
  - If CMV undetectable, CMV pneumonia is ruled out.
  - If transbronchial lung biopsy was done at the time of the BAL and is positive for CMV pneumonia by histology or immunohistology, treat as proven CMV pneumonia.

### Therapy of CMV disease

- Induction with ganciclovir 5mg/kg IV twice daily, or Foscarnet 90 mg/kg IV twice daily, for 2-3 weeks. Followed by maintenance ganciclovir/valganciclovir/foscarnet for 3-4 weeks.
- For CMV pneumonia, add IVIG 500 mg/kg every other day for 2 weeks.
- Maribavir, 400 mg bid orally, may be an alternative to foscarnet for val/ganciclovir resistant/refractory CMV disease. In a randomized study, more patients achieved CMV DNAemia clearance by 8 weeks with maribavir than foscarnet and fewer patients developed acute kidney injury.<sup>26</sup> However, effect on CMV disease was not studied.

## Herpes Simplex Virus (HSV) & Varicella Zoster Virus (VZV)

### Background

#### Epidemiology:

##### HSV

- ~70% adults infected
- ~70% adult HCT recipients shed HSV post transplant (typically in the first month) and ~70% of the shedders developed HSV disease in pre-acyclovir era
- <5% pts shed HSV and <<5% pts develop HSV disease post-transplant per literature since acyclovir prophylaxis
  - In Alberta, in adult alloHCT recipients transplanted in 2012-2022 (acyclovir prophylaxis era), in the first posttransplant year, 74/854 (9%) developed HSV disease (Storek, unpubl)
    - 38 pts (4%) mild oro/genital
    - 27 pts (3%) severe oro/genital (requiring or extending peritransplant hospitalization)
    - 9 pts (1%) internal organ, all required/extended hospitalization
      - 3 esophagitis
      - 3 enteritis
      - 1 encephalitis

- 2 pneumonia, both fatal, thus total fatal HSV disease incidence 0.2% (2/854)

## VZV

- >90% adults infected
- 10-50% adult HCT recipients develop VZV disease (typically at 3-9 months post-transplant) without acyclovir prophylaxis
- Similar cumulative VZV disease incidence with acyclovir prophylaxis (without subsequent vaccination), however, the time of onset shifted to after acyclovir discontinuation

**Risk factors:** Seropositivity of recipient, GVHD / immunosuppressive drugs (this may not be a risk factor for any VZV disease, but it probably is a risk factor for severe VZV disease<sup>2</sup>)

**Clinical manifestations of HSV disease:** Painful mucocutaneous lesions of oropharynx/genitalia, internal organs may be involved, (e.g., lungs, GI tract, liver, CNS)

## Clinical manifestations of VZV disease:

- Shingles (typically with reactivation) → neuralgia
- Chickenpox (typically with primary infection)
- Internal organs may be involved, e.g., lungs, GI tract, liver, CNS
  - Visceral VZV disease may be rapidly progressing and fatal

## Prevention of HSV and VZV Disease

### HSV Prevention/Prophylaxis with Valacyclovir:

- Accepted until 1-month post-transplant
- Controversial until 3 months post-transplant – possibly useful for HSV seropositive recipients with HSV seronegative donors, as these patients may develop late HSV disease
- Irrelevant for patients on VZV prophylaxis, who get valacyclovir anyway

### VZV Prevention/Prophylaxis with Valacyclovir:

- Use valacyclovir (VCV) followed by VZV vaccination.
  - In alloHCT, VCV for 2 years (longer if on immunosuppressive therapy) followed by two doses of a live VZV vaccine was effective for zoster incidence reduction and near-elimination of post-herpetic neuralgia (Appendix A).<sup>27</sup>
    - In Alberta, the timing of the start of the vaccination is at 2 years posttransplant or later.
    - The later start is for patients on prolonged therapy with immunosuppressive drugs – wait until ≥3 mo after discontinuation of immunosuppressive therapy (systemic and topical) and no cGVHD activity.

- In autoHCT, VCV for up to 6 months together with two doses of a non-live vaccine composed of recombinant glycoprotein E with an adjuvant (Shingrix) starting at 2 months posttransplant was effective for zoster incidence reduction and substantially reduced post-herpetic neuralgia.<sup>28</sup>
  - In Alberta, the timing of the start of the vaccination is at 6 months posttransplant (to coincide with the start of other vaccines) or later.
  - The later start is recommended for patients on prolonged maintenance immunosuppressive therapy – wait until ≥6 mo after discontinuation of a lymphodepleting antibody like rituximab.
    - It is unclear whether the delay (the later start) is needed – the randomized study<sup>28</sup> did not exclude patients on maintenance bortezomib, rituximab, or other immunomodulatory therapy, but included only 4% of such patients. The reason we recommend the delay is that the immunogenicity of any vaccine is reduced by immunosuppressive/lymphodepleting therapy, and long-term administration of valacyclovir is non-toxic and virtually 100% effective for zoster prevention as long as the patient remains compliant.
      - Lenalidomide maintenance is not considered immunosuppressive.<sup>29</sup>
- Valacyclovir should be started with conditioning.
- Valacyclovir should be discontinued.
  - In alloHCT, 1 day before the first dose of the live vaccine. This is to avoid the killing of the live vaccine with valacyclovir, which could reduce its immunogenicity.
  - In autoHCT, 1 month after the second dose of Shingrix. This approximately mimics the scenario from the randomized study.<sup>28</sup>
- Give valacyclovir to all adult patients. For pediatric patients, refer to Table 1.
- Use valacyclovir 500 mg po once daily (preferred) or acyclovir 400 mg po twice daily (5mg/kg IV twice daily). For children <40 kg with oral intake, use acyclovir suspension 300 mg/m<sup>2</sup> po twice daily.
- If patient is on ganciclovir/valganciclovir/foscarnet/cidofovir, hold acyclovir/valacyclovir.

**Table 1.** Pediatric patients treated with acyclovir/valacyclovir

| HSV Recipient Serostatus | VZV Recipient Serostatus | Start and End of Prophylaxis                                                                          |
|--------------------------|--------------------------|-------------------------------------------------------------------------------------------------------|
| Positive or Negative     | Positive                 | From day 0 until VZV vaccination (24 months or later)                                                 |
| Positive                 | Negative                 | From day 0 until 1 month posttransplant. Consider extending prophylaxis to 3 months posttransplant if |

|          |          |                                                                                            |
|----------|----------|--------------------------------------------------------------------------------------------|
|          |          | donor is HSV-seronegative. Consider immunizing VZV-seronegative contacts with VZV vaccine. |
| Negative | Negative | No prophylaxis. Consider immunizing VZV-seronegative contacts with VZV vaccine.            |

### Exposure prevention for VZV:

- Important for VZV seronegative patients who are not on valacyclovir/acyclovir. Of limited importance for VZV seropositive patients who are not on valacyclovir/acyclovir (they already have the virus, nevertheless, vesicular rash due to a different strain transferred from a contact person has been described). Of probably no importance for patients who are on valacyclovir/acyclovir.
- Consider vaccination of prospective contacts (caregivers, children, related BMT donors) without history of chickenpox or VZV vaccination (seronegative).
- Instruct the patient to avoid skin contact with vaccinees who have developed a rash. Approximately 20% vaccinees develop a rash at 5-42 days post-vaccination.

### Post-exposure prophylaxis of VZV:

- Important only for patients who are not on valacyclovir/acyclovir. Definition of exposure: residing in the same household, playmate (face-to-face), other face-to-face contact with an infectious person.
- If a seronegative patient has been exposed to a person with varicella or zoster, give varicella zoster immune globulin (or intravenous immunoglobulin) and/or treat with treatment dose of acyclovir/valacyclovir for 3 weeks.
- If a seropositive patient has been exposed to a person with varicella or zoster, observe closely.

## Therapy of HSV and VZV Disease

### HSV Disease:

- Valacyclovir 1000 mg twice daily or Acyclovir 400 mg po three times daily (5 mg/kg IV every 8 hours).
- Treat for 7 days or until resolution of lesions, whichever occurs later.

### VZV Disease:

- Acyclovir 10-12 mg/kg every 8 hours for 1-3 days, then (if oral intake possible) switch to acyclovir 800 mg po 5x/d or valacyclovir 1000 mg po three times daily.
- Treat until 2 days after the last new lesion has crusted (generally 10-14 days).
- Hydrate patient to minimize acyclovir/valacyclovir nephrotoxicity.

### Resistance to Acyclovir/Valacyclovir:

- HSV resistance is relatively common in immunocompromised persons (~5%). Resistance should be suspected if lesions progress or do not improve within 7-10 days of oral val/acyclovir therapy.

Documentation of resistance (mutation of thymidine kinase or DNA polymerase) is of unproven benefit but recommended. Treatment of clinically resistant HSV disease is with

- High dose IV acyclovir (10 mg/kg every 8 hours)
  - Foscarnet IV if no improvement of lesions over 7 days on high dose IV acyclovir
  - Pritelivir (oral) is promising, but still under development
    - Helicase-primase inhibitor, thus different mechanism of action than the above nucleoside inhibitors
    - In a small phase 2 trial, which randomized immunocompromised patients with acyclovir-resistant HSV disease to pritelivir vs foscarnet, lesion healing by 28d occurred in 14 out of 15 (93%) in pritelivir vs 4 out of 7 (57%) foscarnet group (Kotton et al: IDWeek Oct 19-22, 2025, not yet published in a peer-reviewed journal).
    - No activity against VZV, so valacyclovir needs to be given concurrently with pritelivir.
  - After resolution of lesions, prophylactic val/acyclovir should be re-started/continued, as recurrent lesions are frequently val/acyclovir-sensitive, and VZV prophylaxis needs to be continued.
- VZV resistance is extremely rare (<0.1%). Other causes of non-resolving zoster like bacterial superinfection should be suspected.

## Human Herpes Virus 6 (HHV6)

- >90% adults infected.
- ~40% adult HCT recipients have HHV6 detectable in blood, typically in the first 2 months.
- <10% adult HCT recipients develop HHV6 disease (encephalitis, ?rash, ?pneumonitis, ?bone marrow suppression/graft failure)
  - In Alberta, only 2/854 adult alloHCT recipients transplanted in 2012-2022 developed HHV6 menig/encephalitis (Storek, unpublished).
- **Prevention:** Insufficient data exist whether prophylaxis or preemptive therapy with ganciclovir or foscarnet is indicated – unlikely in Alberta, where the incidence of HHV6 encephalitis has been very low. Thus, in Alberta, we use no prophylaxis or preemptive therapy.
- **Therapy of HHV6 disease:** Ganciclovir or foscarnet, same dose as for CMV disease.

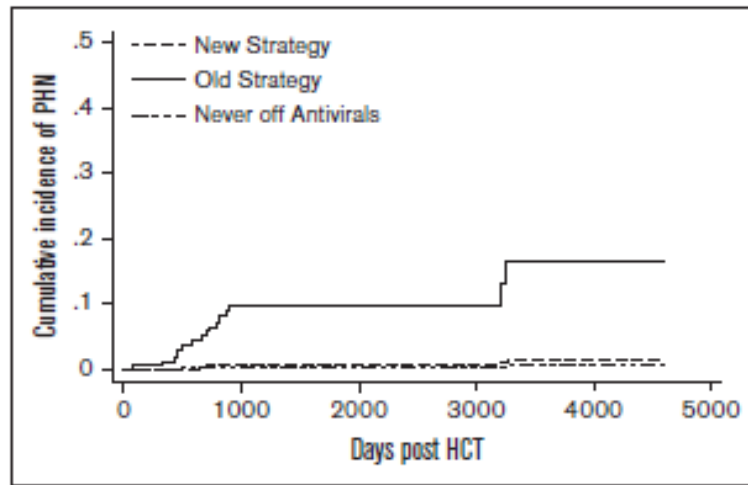


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## Appendix: Cumulative Incidence of Post-Herpetic Neuralgia



Cumulative incidence of post-herpetic neuralgia (PHN) in patients treated with valacyclovir till 2 years followed by vaccination (New Strategy), patients treated with valacyclovir till approximately 1 year without subsequent vaccination (Old Strategy), and patients who continued valacyclovir till the end of follow up (Never off Antivirals). The difference between the New Strategy and the Old Strategy patients was significant ( $p=0.02$ ). From Jamani et al.<sup>27</sup>

# Epstein-Barr virus (EBV) / Posttransplant Lymphoproliferative Disorder

Presented by: Jan Storek

## Summary

### EBV Monitoring

- Use an assay reporting DNAemia in IU/ml blood (the only assay ProvLab offers)
- For allograft recipients, monitor weekly until 3 months and then monthly until 12 months posttransplant.
- For autograft recipients, do not monitor.
- If DNAemia >30,000 IU/mL, watch for symptoms/signs of posttransplant lymphoproliferative disorder (PTLD).
- If DNAemia >300,000 IU/mL, treat PTLD preemptively.

### Preemptive Therapy of PTLD

- Rituximab weekly (1<sup>st</sup> dose 375 mg/m<sup>2</sup> i.v.; 2<sup>nd</sup>-4<sup>th</sup> dose 1400 mg s.c.) until undetectable EBV DNAemia, to a maximum of 4 doses, *and*
- Taper cyclosporine or other immunosuppression to zero over 1-2 weeks (if no GVHD).

### Therapy of PTLD

- Establish PTLD diagnosis by biopsy, or as EBV DNAemia >30,000 IU/mL plus ≥1 of the following:
  - Lymphadenopathy
  - Splenomegaly
  - Mass by imaging
  - B lymphocytosis or kappa/lambda predominance
  - Fever >38.5°C after engraftment, with negative blood cultures, persisting after 48 hours of broad spectrum antibacterials, otherwise unexplained. If fever is the only symptom/sign of PTLD, treat only if EBV DNAemia is >300,000 IU/mL.
- First line therapy: Rituximab and tapering of immunosuppression as for “Preemptive Therapy of PTLD” above. If no response within 2-4 weeks, proceed to second line therapy.
- Second line therapy:
  - If no GVHD and donor is EBV-seropositive:
    - DLI (10<sup>5</sup> T cells/kg), or donor-derived anti-EBV T cells manufactured in our Cellular Therapy Lab (currently unavailable).
    - Consider third-party anti-EBV T cells (less effective but safer than DLI).
  - If no GVHD and donor is EBV-seronegative:
    - Third party anti-EBV T cells – currently available under a trial in Cincinnati (patient has to travel) or from Atara (Tabelecleucel, patient does not have to travel).
    - Consider Blinatumomab or CD19 CAR T cells.
  - If GVHD requiring systemic immunosuppression: No good option. Consider chemo.

- Chemotherapy (eg, CHOP and/or polatuzumab vedotin) may be given while waiting for cellular therapy as a temporizing measure, if PTLD is aggressive. Chemotherapy as definitive therapy is not recommended due to low efficacy and high toxicity.

## Background

### Epstein - Barr virus<sup>1-3</sup>

- EBV is a gamma-herpes virus infecting primarily pharyngeal epithelial cells and B cells.
- Over 90% of adults are infected (seropositive)<sup>4</sup>:
  - EBV is detectable in blood by PCR at one time in 0-16% healthy donors.
  - EBV is detectable in blood by PCR at one of multiple times in 14-83% monitored HCT recipients.
    - In Alberta, with ATG-based GVHD prophylaxis, 86% seropositive HCT recipients reactivate EBV (have EBV detectable in blood by PCR).<sup>5</sup>
      - First reactivation on median day 35
      - Maximum EBV DNAemia: median 33,000 IU/mL
        - 452,000 IU/mL in pts developing PTLD,
        - 23,000 IU/mL in pts not developing PTLD
      - Maximum EBV DNAemia reached on median day 55
- Infected B cells are either quiescent (latent infection) or transformed to proliferate.
- Transformed B cells are eliminated by T cells in immunocompetent hosts.
- PTLD can develop in immunocompromised hosts.
  - Reported incidence after HCT 0.2% - 71%, in Alberta ~10% (using ATG).<sup>5,6</sup>
  - PTLD may be more frequent than clinically appreciated – of 31 retrospectively monitored patients with EBV DNAemia before death due to various causes, PTLD was detected on autopsy in 19/24 patients<sup>6</sup>.

## Risk Factors for Developing EBV PTLD after HCT<sup>7</sup>

|                                  |                                                                                                                                                                                                                                                                                                                                                                                                |
|----------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Major risk factors               | <ul style="list-style-type: none"> <li>• In vivo T cell depletion, particularly using ATG</li> <li>• Ex vivo T cell depletion, particularly without concurrent B cell depletion</li> <li>• Cord blood or marrow graft (compared to unmanipulated blood stem cells)</li> <li>• Donor seropositive with recipient seronegative (D+/R-) serostatus</li> <li>• Second or subsequent HCT</li> </ul> |
| Minor risk factors               | <ul style="list-style-type: none"> <li>• Aplastic anemia (compared to malignancies)</li> <li>• Older recipient</li> <li>• Splenectomy before HCT</li> <li>• Fludarabine in conditioning</li> <li>• Total body irradiation (TBI) in conditioning</li> </ul>                                                                                                                                     |
| Possible risk factor             | <ul style="list-style-type: none"> <li>• RIC or NMAC (compared to myeloablative conditioning)</li> </ul>                                                                                                                                                                                                                                                                                       |
| Major risk-mitigating factors    | <ul style="list-style-type: none"> <li>• Posttransplant cyclophosphamide (without ATG) for GVHD prophylaxis</li> <li>• Rituximab within 2 months before HCT</li> </ul>                                                                                                                                                                                                                         |
| Minor risk-mitigating factors    | <ul style="list-style-type: none"> <li>• Matched sibling donor</li> <li>• Rituximab 2-6 months before HCT</li> </ul>                                                                                                                                                                                                                                                                           |
| Possible risk-mitigating factors | <ul style="list-style-type: none"> <li>• B cell directed CAR T cells or BITEs before HCT</li> <li>• B cell directed antibodies other than rituximab before HCT</li> <li>• Sirolimus for GVHD prophylaxis</li> </ul>                                                                                                                                                                            |

\* The assignment of the attributes of “major”, “minor”, or “possible” is based on our opinion, which is based in part on references<sup>5,8-11</sup>. It takes into account whether a risk factor has been found consistently in multiple studies. In addition, a higher weight is attributed to a risk factor for the mortality due to PTLD than a risk factor for the incidence of PTLD. Even lower weight is attributed to a risk factor for EBV reactivation (EBV DNAemia detectable at all or above a threshold).

## Clinical Manifestations

- Lymphadenopathy
- Splenomegaly
- Mass by imaging
- B lymphocytosis or kappa/lambda predominance
  - Fever >38.5°C after engraftment, with negative blood cultures, persisting after 48 hours of broad spectrum antibacterial(s), otherwise unexplained

## Diagnosis

- Biopsy is the gold standard. Biopsy should include *in situ* hybridization for EBER (EBV-encoded RNA).
- In Alberta, to avoid delay in therapy, we accept for diagnosis at least one of the above clinical manifestations with EBV DNAemia >30,000 IU/mL. However, if fever is the only symptom/sign of PTLD, it should be treated only if EBV DNAemia is >300,000 IU/mL.
  - Rationale for the cutoff of >30,000 IU/mL for diagnosis: This cutoff was originally formulated in 2012, one year after ProVLab’s switching from the DNAemia assay measuring EBV DNA per ug blood DNA to the assay measuring EBV genome copies per mL blood. It was based on a retrospective review of 13 patients with biopsy-proven PTLD occurring in Alberta between 2004 and 2009, who had DNAemia determined within 4 days of onset of symptoms/signs of the PTLD. It included conversion of the old units (genome copies/ug DNA) to the newer units (genome copies/mL, which later turned out to be equivalent to



IU/mL), taking WBC into account. The DNAemia in the 13 cases was 42,383-19,169,040 copies/mL (median 1,633,215). The formulation of the cutoff also took into account data from the first year of EBV monitoring using the assay expressing DNAemia as copies/mL (patients undergoing HCT between Feb 2011 and Jan 2012; only biopsy-proven PTLDs were treated). In that year, 9 PTLDs were diagnosed and all of them were preceded by EBV DNAemia >30,000 copies/mL. This cutoff was further validated in 2015 based on a retrospective review of patients undergoing HCT between May 2012 and Dec 2014 (when EBV DNAemia was monitored weekly and PTLD was treated promptly). In this period, 25 PTLDs were diagnosed and all of them were preceded by EBV DNAemia >30,000 copies/mL. The adequacy of diagnosing PTLD clinically/radiologically in patients with EBV DNAemia >30,000 IU/mL was further proven in 17 patients who also had biopsies.<sup>5</sup>

- Rationale for the cutoff of >300,000 IU/mL when fever is the sole manifestation of PTLD: This cutoff was originally (in 2012) established arbitrarily, by consensus of Calgary transplant physicians, to minimize the likelihood of giving rituximab to patients with fever of etiology other than PTLD. This cutoff was validated in 2015 based on a retrospective review of patients undergoing HCT between Feb 2011 and Dec 2014. In this period, 4 patients died due to PTLD and the diagnosis of all the 4 PTLDs was preceded by EBV DNAemia >300,000 IU/mL.
- Rationale for the conversion of EBV genome copies/mL to IU/mL of 1:1. In mid March 2016, ProvLab started to run 2 EBV DNAemia assays, (1) the in-house assay reporting the EBV DNAemia as copies/mL whole blood, and (2) the RealStar EBV PCR assay (Altona Diagnostics) reporting the EBV DNAemia as IU/mL whole blood. The goal was to transition to running only the RealStar as of June 2016. Between mid March 2016 and mid June 2016, 91 EBV DNAemias above quantitation limit (by both assays) were determined. Results of both assays were near-identical (Kalra et al: submitted).

## **Interventions for Reducing the Incidence or Mortality of PTLD**

Options for reducing the incidence or mortality of PTLD include:

- EBV specific T cells<sup>11-13</sup>
  - 70-100% efficacy
  - No toxicity; however, costly and may be rejected (if from 3<sup>rd</sup> party)
  - Can be given as
    - Prophylaxis (given to all patients early posttransplant)
    - Preemptive therapy (given to patients with high EBV DNAemia in the setting of EBV monitoring)
    - Prompt therapy (given at clinical diagnosis of PTLD in the setting of EBV monitoring)
    - Therapy (given at diagnosis of PTLD in the absence of EBV monitoring)
- Rituximab
  - 60-100% efficacy

- Can be given as prophylaxis, preemptive therapy, prompt therapy, or therapy (without EBV monitoring) – Table 1
- Toxicities/disadvantages of rituximab:
  - Infusion reactions
  - Hypo-IgM/IgG
  - Neutropenia,<sup>12</sup> which may be clinically significant<sup>13-15</sup>
  - Vaccination onset needs to be moved to at least 6 months after the last rituximab dose
- Reduction of immunosuppressive drug(s) preemptively
  - Efficacy and toxicity (GVHD?) in the setting of preemptive or prompt therapy not well studied
  - In the setting of Therapy, reduction of immunosuppression (RI) studied only in addition to rituximab
  - Addition of RI to rituximab ↓'ed mortality due to PTLD & ↑'ed overall survival<sup>16</sup>
- Purging grafts of B cells (theoretical)
- Alemtuzumab instead of ATG
  - PTLD still occurs, though less than with ATG<sup>17-20</sup>
  - Alemtuzumab may be associated with more CMV disease and other non-EBV viral infections.<sup>21</sup> Moreover, impact of alemtuzumab on relapse has not been well studied whereas ATG with myeloablative conditioning has not been associated with increased relapse in 5 randomized studies.<sup>22</sup>
- CD19-directed CAR T cells or Blinatumomab
  - So far only case reports of success.<sup>23-26</sup>
  - Theoretically, blinatumomab needs a minimum number of T cells for efficacy.

**Table 1.** Comparison of four strategies of management of PTLD with rituximab<sup>7</sup>

| Management Strategy            | Number of evaluable patients | Number of controls   | Management strategy in controls            | Efficacy Endpoint   | % patients who achieved the efficacy endpoint | Comment                   | Reference                         |
|--------------------------------|------------------------------|----------------------|--------------------------------------------|---------------------|-----------------------------------------------|---------------------------|-----------------------------------|
| Therapy without EBV Monitoring | 12 w PTLD                    | None                 | n/a                                        | Sustained CR        | 67%                                           |                           | Faye 2001 <sup>27</sup>           |
|                                | 146 w PTLD                   | None                 | n/a                                        | Cure or improvement | 63%                                           |                           | Styczynski 2009 <sup>28</sup>     |
|                                | 144 w PTLD                   | None                 | n/a                                        | Not dying 2° PTLD   | 61%                                           |                           | Styczynski 2013 <sup>16</sup>     |
|                                | 27 w PTLD                    | None                 | n/a                                        | CR                  | 74%                                           |                           | Zhu 2019 <sup>29</sup>            |
|                                | 19 w PTLD                    | None                 | n/a                                        | Sust. regression    | 73%                                           |                           | Kalra 2018 <sup>6</sup>           |
|                                | 70 w PTLD                    | None                 | n/a                                        | CR                  | 69%                                           |                           | Luo 2020 <sup>30</sup>            |
| Prompt therapy                 | 5 w PTLD                     | None                 | n/a                                        | Regression          | 100%                                          |                           | Wagner 2004 <sup>31</sup>         |
|                                | 6 w PTLD                     | None                 | n/a                                        | CR                  | 67%                                           |                           | Kinch 2007 <sup>32</sup>          |
|                                | 6 w PTLD                     | None                 | n/a                                        | Not dying 2° PTLD   | 17%                                           |                           | Sanz 2014 <sup>33</sup>           |
|                                | 87 w PTLD                    | None                 | n/a                                        | ORR                 | 51%*                                          |                           | Garcia-C. 2019 <sup>34</sup>      |
|                                | 266 total, 24 w PTLD         | 199 total, 19 w PTLD | Therapy w/o EBV Monitor.                   | Sust. regression    | 75 vs 73% (N.S.)                              | No impact on OS           | Kalra 2018 <sup>6</sup>           |
|                                |                              |                      |                                            | PTLD incidence      | 11 vs 6% (p=.06)                              |                           |                                   |
|                                |                              |                      |                                            | Mortality 2° PTLD   | 1 vs 1% (N.S.)                                |                           |                                   |
| Preemptive therapy             | 93 w high EBV                | None                 | n/a                                        | EBV undetectable    | 83%                                           | 2 patients died of PTLD   | Garcia-Cadenas 2015 <sup>35</sup> |
|                                | 55 w high EBV                | None                 | n/a                                        | EBV not high        | 91%                                           | 3 patients died of PTLD   | Coppoletta 2011 <sup>36</sup>     |
|                                | 9 w high EBV                 | None                 | n/a                                        | Not dying 2° PTLD   | 44%                                           |                           | Pinana 2016 <sup>37</sup>         |
|                                | 17 w high EBV                | None                 | n/a                                        | Not dying 2° PTLD   | 100%                                          |                           | Ahmad 2009 <sup>38</sup>          |
|                                | 49 total                     | 85 total             | Therapy w/o EBV Monitor.                   | PTLD incidence      | 6 vs 12% (N.S.)                               | Impact on OS not reported | VanEsser 2002 <sup>39</sup>       |
|                                |                              |                      |                                            | Mortality 2° PTLD   | 0 vs 6% (N.S.)**                              |                           |                                   |
|                                | 35 total                     | 30 total             | Therapy w/o EBV Monitor.                   | PTLD incidence      | 6 vs 17% (N.S.)                               | Impact on OS not reported | Blaes 2010 <sup>40</sup>          |
|                                |                              |                      |                                            | Mortality 2° PTLD   | 3 vs 7% (N.S.)                                |                           |                                   |
|                                | 320 total                    | 872 total            | Therapy w/o EBV Monitor. or Prompt Therapy | PTLD incidence      | SHR=1.02 (N.S.)                               | Impact on OS not reported | Kinzel 2022 <sup>5</sup>          |
|                                |                              |                      |                                            | Mortality 2° PTLD   | SHR=0.16 (N.S.)                               |                           |                                   |
| Prophylaxis                    | 55 total                     | 68 total             | Preemptive therapy                         | EBV high            | 14 vs 49% (p<.001)                            |                           | Dominietto 2012 <sup>12</sup>     |

|  |          |           |                   |                   |                    |                                             |                               |
|--|----------|-----------|-------------------|-------------------|--------------------|---------------------------------------------|-------------------------------|
|  |          |           |                   | PTLD incidence    | 0 vs 3% (N.S.)     | No impact on OS or mortality 2° PTLD        |                               |
|  | 51 total | 147 total | Prompt therapy*** | EBV reactivation  | 2 vs 13% (p<.001)  | No impact on NRM. Impact on OS not reported | Van Besien 2019 <sup>41</sup> |
|  |          |           |                   | PTLD incidence    | 0 vs 8% (p=.041)   |                                             |                               |
|  |          |           |                   | Mortality 2° PTLD | 0 vs 3% (N.S.)     |                                             |                               |
|  | 43 total | 43 total  | Prompt therapy*** | EBV reactivation  | 0 vs 53% (p<.001)  | Impact on NRM or OS not reported            | Patel 2023 <sup>42</sup>      |
|  |          |           |                   | PTLD incidence    | 0 vs 14% (p=.02)   |                                             |                               |
|  |          |           |                   | Mortality 2° PTLD | 0 vs 9% (N.S.)**** |                                             |                               |

\* Additional 15 patients with PTLD did not receive rituximab due to poor performance status (n=11) or PTLD diagnosis only post-mortem.

\*\* Significant difference when comparing subgroups of patients with high EBV DNAemia.

\*\*\* Stated in Van Besien 2019, surmised for Patel 2023 (publication from the same group as Van Besien 2019).

\*\*\*\* Significance analyzed by Chi-square or Fisher's exact test based on raw data reported in the paper.

Abbreviations: OS, overall survival; NRM, non-relapse mortality; PTLD, posttransplant lymphoproliferative disorder; Ref, reference; EBV, Epstein-Barr virus; High EBV, high EBV DNAemia; CR, complete remission; ORR, overall response rate (complete or partial remission); Ctrl, control; N.S., not significant; Sust. regression, sustained regression (regression not followed by later progression of PTLD); SHR, subhazard ratio; Mortality 2° PTLD, mortality due to PTLD; Garcia-C., Garcia-Cadenas; w, with; w/o, without; monitor., monitoring; n/a, not applicable.

In Alberta, since September 2015 we use preemptive therapy with rituximab plus taper of immunosuppression. We use the threshold of 300,000 IU/ml. This is a compromise between trying to minimize the number of deaths due to PTLD and to minimize the number of patients exposed to the risks of rituximab/taper of immunosuppression unnecessarily (Table 2 and 3). The addition of the taper of immunosuppression is an extrapolation from the study of Styczynski et al<sup>16</sup> showing overall survival benefit in the setting of therapy (not preemptive therapy). In the setting of preemptive therapy, the taper of immunosuppression on top of rituximab does not appear to cause GVHD.<sup>15</sup>

The use of preemptive therapy in patients whose conditioning includes ATG is in line with EBMT guidelines.<sup>43</sup> If in the future it is shown that all/most patients with PTLD who have failed rituximab plus taper of immunosuppression respond to other treatment like anti-EBV/CD19 T cells or blinatumomab, therapy of established PTLD (without EBV monitoring) may become the management of choice/

### **Length of Interval between Rituximab Doses, and When to Stop Rituximab**

- In the preemptive therapy setting and therapy setting, treatment has been reported once a week (375 mg/m<sup>2</sup> i.v.), until undetectable DNAemia, maximum 4 doses<sup>16,35,36,38,40</sup>. In Alberta, we adopt the weekly dosing given that
  - It is in line with EBMT guidelines<sup>43</sup>.
  - There is no evidence of benefit of more frequent dosing.
  - Weekly dosing saves rituximab, as most patients need only 2-3 doses to achieve undetectable DNAemia<sup>38,40</sup>.
  - One dose only may be sufficient (in preemptive setting)<sup>39</sup>.
- The only exception to the rule of weekly dosing is in a patient whose PTLD manifests with fever and the fever has not abated after 2-3 days following the first rituximab dose (and immunosuppression taper). In this instance twice weekly dosing is reasonable, so that failure of rituximab with immunosuppression taper can be pronounced early and second line therapy organized in 2 weeks after the first rituximab dose.
- Patients who have reached undetectable DNAemia after being treated for PTLD with rituximab have 100% likelihood of sustained clinical regression of PTLD (based on our experience in 15 patients, Kalra et al, ASH 2015 abstract). Thus, rituximab should be stopped when DNAemia has become undetectable.
- Patients who have not reached undetectable DNAemia after being treated for PTLD with 4 doses of rituximab have ~58% likelihood of clinical progression of PTLD (based on our experience in 12 patients, Kalra et al, ASH 2015 abstract). Thus, patients with persistently detectable EBV DNAemia after 4 weekly rituximab doses should be followed closely. Second line therapy should be instituted in case of PTLD progression or new PTLD diagnosis.

**Table 2.** PTLD incidence and mortality according to maximum DNAemia (pre-rituximab, if given).<sup>\*\*</sup>

| EBV DNAemia (max)*                                                                       | Undetectable  | <10,000/mL    | 10,000 – 100,000/mL | 100,000 – 1,000,000/mL | >1,000,000/mL   |
|------------------------------------------------------------------------------------------|---------------|---------------|---------------------|------------------------|-----------------|
| Number of patients with PTLD of total patients in the max DNAemia range (%)              | 0/56<br>(0 %) | 0/43<br>(0 %) | 0/103<br>(0 %)      | 25/82<br>(30 %)        | 18/22<br>(81 %) |
| Number of patients with <i>fatal</i> PTLD of total patients in the max DNAemia range (%) | 0/56<br>(0 %) | 0/43<br>(0 %) | 0/103<br>(0 %)      | 3/82<br>(4 %)          | 2/22<br>(9 %)   |

\* EBV genome copies/mL, which is near-equivalent to IU/mL.

\*\* Data based on 306 Albertan patients who were monitored for EBV DNAemia but not treated preemptively.

**Table 3.** Possible EBV DNAemia thresholds for preemptive therapy.<sup>\*\*</sup>

| Cut off EBV DNAemia (max) | Number of PTLDs avoided by preemptive therapy (%) (assuming 100% efficacy of the preemptive therapy) | % Patients treated with rituximab of total 306 patients | % Patients treated with rituximab necessarily (would develop PTLD) of total 306 patients | % Patients treated with rituximab unnecessarily (would not develop PTLD) of total 306 patients | Number of patients dying of PTLD (% of total 306 patients) |
|---------------------------|------------------------------------------------------------------------------------------------------|---------------------------------------------------------|------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------|------------------------------------------------------------|
| 100,000                   | 43/43 (100 %)                                                                                        | 34 %                                                    | 14.0 %                                                                                   | 20 %                                                                                           | 0/306 (0.0 %)                                              |
| 200,000                   | 39/43 (91 %)                                                                                         | 25.5 %                                                  | 12.7 %                                                                                   | 12.7 %                                                                                         | 1/306 (0.3 %)                                              |
| 300,000                   | 33/43 (77 %)                                                                                         | 16.7 %                                                  | 11 %                                                                                     | 6.2 %                                                                                          | 1/306 (0.3 %)                                              |
| 400,000                   | 31/43 (72 %)                                                                                         | 14.7 %                                                  | 10.1 %                                                                                   | 4.5 %                                                                                          | 1/306 (0.3 %)                                              |
| 500,000                   | 23/43 (53 %)                                                                                         | 11.4 %                                                  | 7.5 %                                                                                    | 3.9 %                                                                                          | 1/306 (0.3 %)                                              |
| 600,000                   | 22/43 (51 %)                                                                                         | 10.7 %                                                  | 7.1 %                                                                                    | 3.6 %                                                                                          | 2/306 (0.65 %)                                             |
| 700,000                   | 22/43 (51 %)                                                                                         | 10.1 %                                                  | 7.1 %                                                                                    | 2.9 %                                                                                          | 2/306 (0.65 %)                                             |
| 800,000                   | 20/43 (46.5 %)                                                                                       | 9.1 %                                                   | 6.5 %                                                                                    | 2.6 %                                                                                          | 3/306 (1.0 %)                                              |
| 900,000                   | 19/43 (44 %)                                                                                         | 7.5 %                                                   | 6.2 %                                                                                    | 1.3 %                                                                                          | 3/306 (1.0 %)                                              |
| 1,000,000                 | 18/43 (42 %)                                                                                         | 7.1 %                                                   | 5.8 %                                                                                    | 1.3 %                                                                                          | 3/306 (1.0 %)                                              |

\* EBV genome copies/mL, which is near-equivalent to IU/mL.

\*\* Data based on 306 Albertan patients who were monitored for EBV DNAemia but not treated preemptively.



## Second Line Therapy<sup>44-46</sup>

- To be used if no response to rituximab with immunosuppression taper in 2-4 weeks.
- If no GVHD and donor is EBV-seropositive:
  - DLI ( $10^5$  T cells/kg), or donor-derived anti-EBV T cells manufactured in our Cellular Therapy Lab (currently unavailable).
    - The main advantage of anti-EBV T cells over DLI is no toxicity. However, this option, even when available, should be discussed on a case-by-case basis as it is resource-intensive and efficacy data are limited (12/16 responders using various donor sources).<sup>47-49</sup>
    - Anti-EBV T cells are manufactured from mononuclear cell apheresis product ( $10^9$  WBCs) in 1-2 days, using IFNg capture system. Our CTL lab uses Prodigy closed system, in which mononuclear cells are incubated overnight with EBV peptides in the presence of a cytokine capture reagent (a bispecific antibody for CD45 and IFNg) and then incubated with anti-IFNg conjugated to a magnetic bead. The magnetic (IFNg secreting, i.e., anti-EBV) T cells are then separated by a magnet.
  - Consider off-shelf third-party anti-EBV T cells (less effective but safer than DLI).<sup>50,51</sup>
    - Currently available under a trial in Cincinnati (patient has to travel) or from Atara (Tabelecleucel, patient does not have to travel).
    - Sustained remission of PTLTD occurs in 70-100% cases after infusion of EBV T cells from the original stem cell donor, but in only 50-60% cases after infusion of EBV T cells from a 3rd party, due in part to rejection of the 3rd party cells.
- If no GVHD and donor is EBV-seronegative:
  - Third party off-shelf anti-EBV T cells
  - Consider Blinatumomab or CD19 CAR T cells.
- If GVHD requiring systemic immunosuppression: No good option. Consider chemo.
- Chemotherapy (e.g., CHOP and/or polatuzumab vedotin (anti-CD79b conjugated to monomethylauristatin E, which damages microtubules)) may be given while waiting for cellular therapy as a temporizing measure, if PTLTD is aggressive. Chemotherapy as definitive therapy is not recommended due to low efficacy and high toxicity (median survival 2 months, range 1-6).<sup>52</sup>
- Future options may include:
  - EBV thymidine kinase inducers, making EBV-infected cells susceptible to ganciclovir.<sup>53</sup>
  - Checkpoint inhibitors like nivolumab.<sup>54</sup>

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# Pneumocystis and Bacterial Prophylaxis

Presented by: Jan Storek

## Summary

- Bacterial prophylaxis peritransplant
  - GCSF – only autologous HCT recipients and cord blood transplant recipients
    - Start on day 7. Discontinue when ANC > 1.0/nl
    - In adults, use 300 micrograms qd sc for <70 kg patients, 480 micrograms qd sc for >70 kg patients
    - In children, use 5 micrograms/kg daily sc
  - No growth factors routinely for allogeneic HCT recipients (except for cord blood)
  - No growth factors routinely for CAR T cell recipients (except if late neutropenia)
  - No antibacterials peri-transplant routinely (both autologous and allogeneic HCT recipients, except some autoimmune diseases (see pertinent chapter))
  - No IVIG routinely. IVIG can be considered for very low IgG (<4g/L), or low IgG (4-6 g/L) associated with severe or recurrent non-neutropenic infections.
- *Pneumocystis jirovecii* and *Streptococcus pneumoniae* prophylaxis
  - Both autologous and allogeneic HCT recipients, and CAR T cell recipients.
  - Start at engraftment. If CD4 ≥ 200/microliter at 12 months, discontinue PJP and *Pneumococcal* prophylaxis. If CD4 < 200/microliter at 12 months, continue until 24 months. Continue/resume prophylaxis when treating GVHD with immunosuppressive drugs, until ≥ 3 months after discontinuation of immunosuppressive therapy (systemic and topical), when cGVHD is inactive.
  - Pretransplant prophylaxis should be considered in patients with substantial immune deficiency, including lymphoma/myeloma patients after mobilization chemotherapy, acute leukemia patients after induction/consolidation chemotherapy, or CLL patients treated with alemtuzumab.
  - Prefer cotrimoxazole (sulfamethoxazole + trimethoprim)
    - In adults, 400/80 mg po qd
    - In children, 375/75 mg/m<sup>2</sup> po qd
  - For cotrimoxazole-intolerant patients (only if intolerance has been well documented), use Dapsone 50 mg po qd every day (1 mg/kg po qd in children), plus Penicillin V 600 mg po qd (150-300 mg po qd in children). Penicillin can be omitted in autologous HCT recipients.
  - In splenectomized patients, give Penicillin (dose as above) indefinitely, except when patient is on cotrimoxazole.

## Background

The literature on bacterial/*Pneumocystis* prophylaxis after HCT contains few randomized trials. Most of the randomized trials on bacterial prophylaxis are of limited value due to the emergence of bacterial resistance to the drug studied in the randomized trial after the follow-up period of the trial. Most of the trials on *Pneumocystis* prophylaxis were performed in HIV patients and recommendations were extrapolated to HCT patients. The literature has been well summarized in international guidelines<sup>1,2</sup>. These recommendations, including Calgary-specific deviations, are summarized below.

## Recommendations for Peritransplant & Early Post-HCT (< 3 month) Period

- Dental consult pretransplant
- Hand washing/sanitizing
- Single-bed rooms and other hospital infection control
- Household contacts and health care workers should be up-to-date with vaccines
- No gut decontamination (resistance, compliance, cost)
- No antibiotic-impregnated central catheters (controversial efficacy, high cost)
- No systemic antibacterials peritransplant
  - Advantage:
    - Low rate of bacterial infection or fever (but no survival benefit)
  - Disadvantages:
    - Resistance
    - *C. difficile*
- Growth factors
  - AutoHCT: G-CSF from day 7. Despite no effect on OS, there is reduction of infections and shortening of hospital stay.
  - AlloHCT: No growth factors routinely, because
    - GVHD may be induced/worsened by GM-CSF or G-CSF<sup>3</sup>
    - T cell reconstitution may be impaired by G-CSF (if ATG used)<sup>4</sup>, which may negatively impact NRM and OS<sup>5</sup>
  - CAR T cells: No growth factors routinely due to insufficient information on risk:benefit
    - OK to use G-CSF in case of late neutropenia.
- No routine IVIG (only a marginal or undetectable reduction in rates of bacterial infections).
  - OK to give IVIG with very low IgG (<4g/L), or low IgG (4-6 g/L) associated with severe or recurrent non-neutropenic infections.
  - This also applies to the late post-HCT (d> 100) period.
- For *Pneumocystis* prophylaxis, see next section.



## Recommendations for Late Post-HCT (d >100) Period

- *Pneumocystis jirovecii* pneumonia (PJP) incidence in pre-prophylaxis era was 4% in the first 3 months and 6% later after allogeneic HCT<sup>6</sup>. When PJP prophylaxis was used until approximately 6 months in allo HCT recipients not getting ATG (typically getting only CNI+MTX for GVHD prophylaxis), PJP incidence was ≤1%. However, with ATG, in Albertan patients using PJP prophylaxis until approximately 6 months, we have noted PJP incidence over 3%<sup>7</sup>. Specifically, in 278 patients without grade 2-4 aGVHD or moderate-severe cGVHD who discontinued PJP prophylaxis at 6 months or soon thereafter, no PJP occurred in the first 6 months, 8 PJPs occurred at 7-12 months, 2 PJPs occurred at 13-24 months, and no PJP at >24 months. As approximately 30% of patients with PJPs need to be treated in the ICU and approximately 15% were fatal, in 2018 we decided to extend PJP prophylaxis until 12 months, and to 24 months in patients with CD4 T cell counts < 200/microliter at 12 mo. CD4 T cell count < 200/microliter is a well-recognized risk factor for PJP (reviewed by Messiaen PE et al.,<sup>8</sup> consistent with Evernden C et al.<sup>7</sup>). Thus:
  - PJP prophylaxis in Alberta is routinely given to patients from engraftment until 12 mo posttransplant. For patients with CD4 T cell count < 200/microliter at 12 months, prophylaxis is continued until 24 months. Patients treated with immunosuppressive drugs for chronic GVHD should continue PJP until ≥3 months after discontinuation of immunosuppressive therapy (systemic and topical), when cGVHD is inactive.
    - Sulfamethoxazole + trimethoprim (cotrimoxazole) is preferred to dapsone, atovaquone and inhaled pentamidine due to highest efficacy (see Tables) and broader antimicrobial spectrum (including *S.pneumoniae*, *Toxoplasma*, *Nocardia*).
    - Patients with documented allergy to cotrimoxazole may be desensitized (see Appendix). Patients who experience non-allergic toxicity to cotrimoxazole (e.g., cytopenia, ↑ALT, ↑creatinine), should be rechallenged with cotrimoxazole prior to being committed to long-term treatment with a second-line agent.
    - Multiple regimens of cotrimoxazole have been found near 100% efficacious for PJP prophylaxis (e.g., 400/80 mg qd, 800/160 mg qd, 800/160 mg 3x a week) (see Tables). In Alberta, 400/80 mg qd is used due to simplicity.
    - For second-line prophylaxis, dapsone 50 mg qd is preferred<sup>7</sup>. Atovaquone as well as inhaled pentamidine have a high breakthrough PJP rate<sup>7,9</sup>.
- *Streptococcus pneumoniae* disease incidence is significantly higher in allogeneic HCT recipients compared to general population (Figure 1). Peak incidence is at 3-24 months posttransplant. Risk factors include:
  - cGVHD (Fig. 1)
  - Splenectomy
  - Hypo-IgG, particularly IgG<sub>2</sub>
- Antibiotics covering *S. pneumoniae* are routinely given to all Albertan HCT recipients from engraftment until the end of PJP prophylaxis, as both cGVHD and low CD4 counts

are risk factors for *S. pneumoniae* disease. In splenectomized patients, *S. pneumoniae* prophylaxis is continued indefinitely.

- In autologous HCT recipients, both PJP and *S. pneumoniae* disease incidences are lower than after allogeneic HCT but higher than in the general population. For simplicity, we use the same PJP/*S. pneumoniae* prophylaxis as for allogeneic HCT recipients. However, for autologous HCT recipients who are cotrimoxazole-intolerant, it is acceptable to give only dapsone and omit penicillin. The rationale is that the incidence of *Pneumococcal* infections after autoHCT is approximately 2-fold lower than after alloHCT (5 vs 12/1000 HCTs,<sup>10</sup> or 5 vs 9/1000 HCTs<sup>11</sup>), which makes *Pneumococcal* prophylaxis less important after auto than alloHCT.
- In CAR T cell recipients, we use the same PJP and *S. pneumoniae* prophylaxis as for allogeneic HCT recipients. The reason is that per limited data available so far, CAR T cell recipients are prone to similar infections and as frequent infections as HCT recipients, albeit possibly not as severe<sup>12</sup>, and reconstitution of CD4 T cell counts to >200/microliter appears to take 1 to 2 years, i.e., similar for ATG-conditioned alloHCT recipients<sup>13,14</sup>. This approach is further supported by current guidelines of EBMT, ASCO, ASTCT, and CARTOX.
  - In a study of 300 CAR T cell recipients (mostly anti-CD19) not prophylaxed against PJP, 8 (2.7%) developed PJP, median day 98 (range, 52-251) (median f/u of total cohort unclear but probably <1 year)<sup>15</sup>. This suggests that prophylaxis may be warranted, but risk:benefit should ideally be established (contribution of cotrimoxazole to post-CART cytopenias?).
- Vaccinate patients against *S. pneumoniae* and with other vaccines per standard schedule (see chapter on Vaccination).

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Appendix, including Figures and Tables

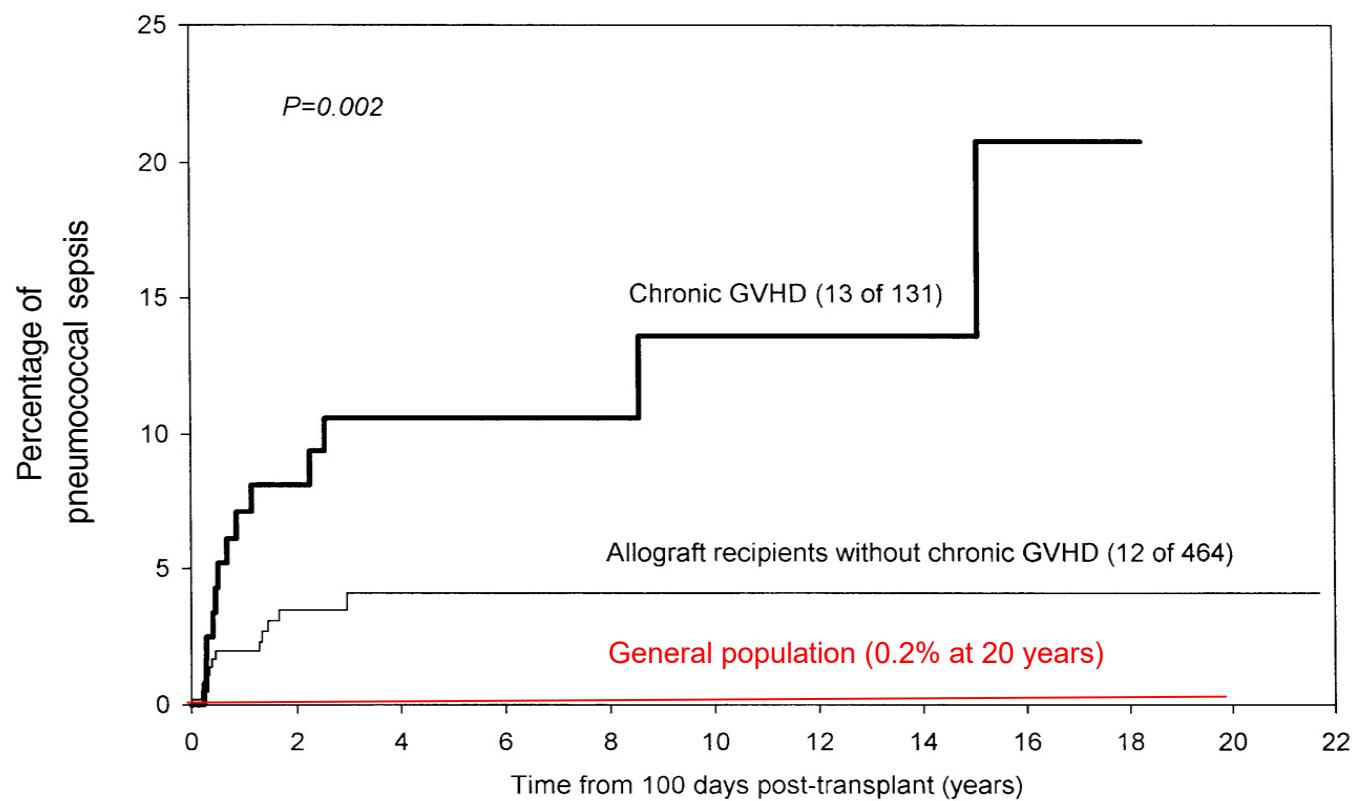
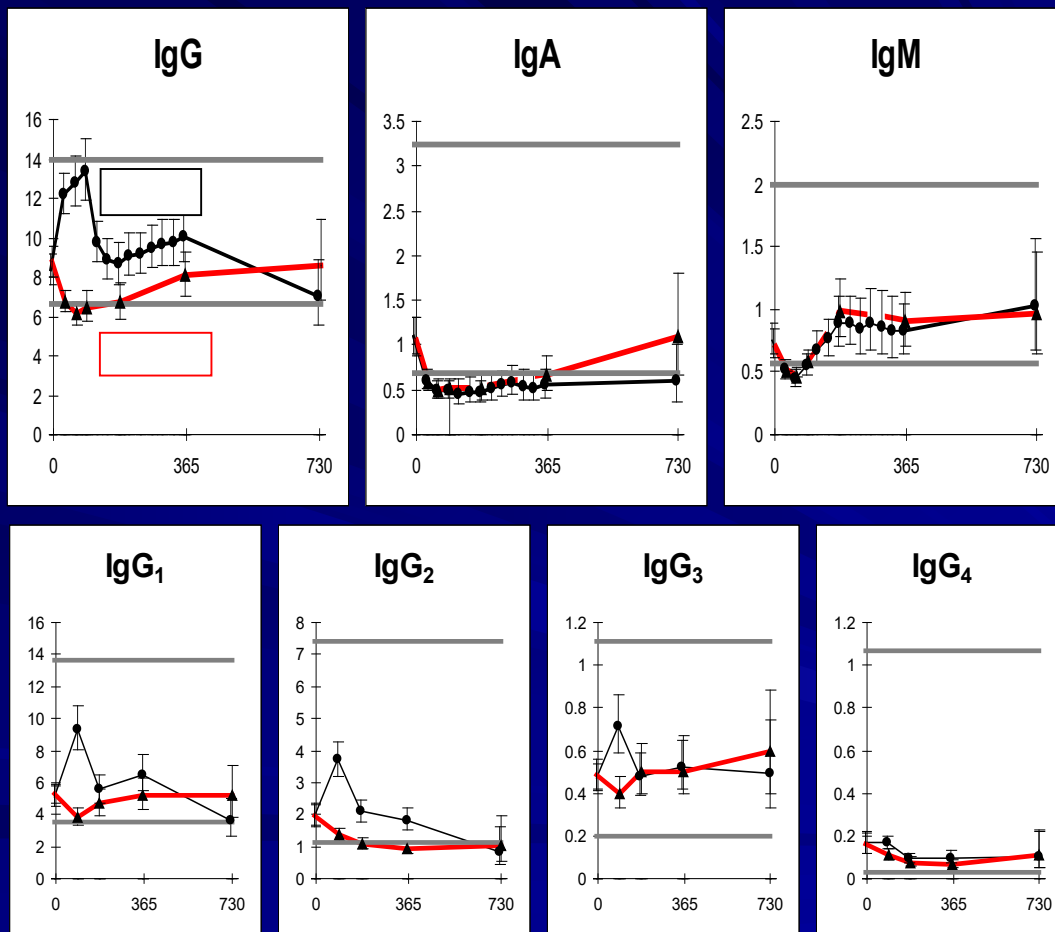


**Fig. 1. Incidence of Pneumococcal Sepsis after alloHCT.** From Kulkarni, S. et al. Blood 2000;95:3683-3686.<sup>16</sup> Red line represents data on general population from Kumar D. et al. BMT 41:743-747, 2008.<sup>17</sup>



Sullivan BBMT 1996

**Fig. 2. Serum Ig concentration in patients not receiving IVIgG (red line) and patients receiving IVIgG in the first 12 months posttransplant, showing that whereas IgG levels were higher in the IVIgG group till 1 year, they were paradoxically lower at 2 years, suggesting that the exogenous IgG hampered reconstitution of the production of endogenous IgG. From Sullivan KM et al. BBMT 2:44-53, 1996.<sup>18</sup>**



## Comparison of prophylactic dosing schedules of Sulfamethoxazole+Trimethoprim and alternative anti-PJP drugs

Schneider MM et al. A controlled trial of aerosolized pentamidine or thrimethoprim-sulfamethoxazole as primary prophylaxis afainst Pneumocystis pneumonia in patient with HIV infection.<sup>19</sup>

|                             | Efficacy<br>(% developing Pneumocystis pneumonia) | Toxicity<br>(% discontinuing drug) |
|-----------------------------|---------------------------------------------------|------------------------------------|
| Pentamidine inhaled monthly | 11%                                               | 3%                                 |
| Sulfa+Trim 800+160 mg daily | 0%                                                | 25%                                |
| Sulfa+Trim 400+80 mg daily  | 0%                                                | 24%                                |

Bozzette SA et al. A randomized trial of three antiPneumocystis agents in patients with advanced human immunodeficiency virus infection.<sup>20</sup>

|                             | Efficacy<br>(% developing Pneumocystis pneumonia per year) | Toxicity<br>(% discontinuing drug) |
|-----------------------------|------------------------------------------------------------|------------------------------------|
| Dapsone 50 mg bid           | 2.6%                                                       | 75%                                |
| Pentamidine inhaled monthly | 5.7%                                                       | 12%                                |
| Sulfa+Trim 800+160 mg bid   | 1.2%                                                       | 79%                                |

Hughes WT et al. Successful intermittent chemoprophylaxis for Pneumocystis pneumonitis (in pts treated with chemotherapy for acute lymphoblastic leukemia).<sup>21</sup>

|                                               | Efficacy<br>(% developing Pneumocystis pneumonia) | Toxicity<br>(% with adverse effect) |
|-----------------------------------------------|---------------------------------------------------|-------------------------------------|
| Sulfa+Trim 800+160 mg daily                   | 0%                                                | 17%                                 |
| Sulfa+Trim 800+160 mg 3x/week (3 consec.days) | 0%                                                | 20%                                 |

Sangiolo D et al. Toxicity and Efficacy of daily dapsone as Pneumocystis jirovecii prophylaxis after HCT: A case-control study.<sup>22</sup>

|                           | Efficacy<br>(% developing Pneumocystis pneumonia) | Toxicity<br>(% discontinuing drug) |
|---------------------------|---------------------------------------------------|------------------------------------|
| Dapsone 50 mg bid         | 1.3%                                              | Not given                          |
| Sulfa+Trim 800+160 mg bid | 0%                                                | Not given                          |

Souza JP et al. High rates of Pneumocystis carinii pneumonia in allogeneic blood and marrow transplant recipients receiving dapsone prophylaxis.<sup>23</sup>

|                                        | Efficacy<br>(% developing Pneumocystis pneumonia) | Toxicity<br>(% discontinuing drug) |
|----------------------------------------|---------------------------------------------------|------------------------------------|
| Dapsone 50 mg bid 3x a week            | 7.2%                                              | Not given                          |
| Sulfa+Trim 800+160 mg bid twice a week | 0.4%                                              | Not given                          |

## Desensitization Protocol for HCT Patients with Sulfa Allergies

(Modified from Purdy et al.<sup>24</sup> and Pyle et al.<sup>25</sup>)

Desensitization should be performed in the clinic (or in hospital), with the patient remaining in the clinic for 30 min after taking a dose. This is more important for a history of anaphylaxis than a history of only rash.

A stock solution (Standard Pediatric Oral Suspension, trimethoprim (TMP) 40 mg plus sulfamethoxazole (SMX) 200 mg per 5 ml) is used, followed by single-strength tablets (80 mg TMP plus 400 mg SMX).

**Sulfa Desensitization Schedule:** (same for adults and children)

On Days 1 through 5: the **stock suspension is diluted:** One (1) ml of stock + 9 ml saline in a 10 ml syringe = 4 mg/ml SMX

Day 1: Take 0.25 ml = 1 mg SMX

Day 2: Take 0.50 ml = 2 mg SMX

Day 3: Take 1 ml = 4 mg SMX

Day 4: Take 2 ml = 8 mg SMX

Day 5: Take 4 ml = 16 mg SMX

On Days 6 through 9, the **stock solution is used full strength:**

Day 6: Take 0.5ml of stock = 20 mg SMX

Day 7: Take 1 ml of stock = 40 mg SMX

Day 8: Take 2 ml of stock = 80 mg SMX

Day 9: Take 4 ml of stock = 160 mg SMX

Day 10: Take 1 single-strength tablet (400 mg SMX).

If no reaction occurs, patient can continue dosing at 1 single-strength tablet once daily. Allergic reaction can occur up to 30 days into this dosing, however, the reaction is usually mild so the doses do not have to be given in the clinic.

**If a mild allergic reaction occurs** or if the desensitization process is interrupted for reasons other than allergic reaction, then give a test dose of half the last dose. If the patient tolerates this test dose, then restart dosing at the last dose.

**If a severe allergic reaction occurs**, administer epinephrine, 0.3-0.5 mL of 1:1000 dilution, subcutaneously every 10-20 min.

If needed, follow by a corticosteroid (eg, 50 mg methylprednisolone IV q 6 h).

If needed, follow by an antihistamine (eg, diphenhydramine 25-50 mg IV or IM or PO q 6 h) and normal saline IV.

# Fungal Prophylaxis

Presented by: Ahsan Chaudhry

## Summary

- Primary prophylaxis
  - Fluconazole 400 mg daily should be given to all allogeneic hematopoietic cell transplant recipients from days 1 to 28. Fluconazole prophylaxis is not routinely accompanied by galactomannan monitoring except in high risk patients.
  - Posaconazole or Isavuconazole can be used in grade 3-4 GVHD, depending on tolerance, drug interactions, and availability.
  - CAR-T cell therapy recipients should receive fluconazole prophylaxis during the neutropenic period (ANC <0.5). If prolonged neutropenia (>14 days) or post-infusion immunosuppression (e.g., corticosteroids, tocilizumab), consider extended anti-mold prophylaxis (Posa or Isavucon) or galactomannan monitoring.
  - No primary prophylaxis or galactomannan screening should be applied to those who develop grade 1-2 acute GVHD, chronic GVHD (irrespective of severity), or to autologous transplant recipients.
- Secondary prophylaxis may be used. It requires consideration of the etiologic agent identified from the previous episode of invasive fungal disease, and of the previous response to antifungal therapy.
- Empiric antifungal treatment is given to patients with persistent neutropenic fever not responsive to at least 4 days of appropriate antibacterial coverage. Micafungin or liposomal amphotericin B is used. The empiric antifungal treatment will be discontinued after 2 days of absolute neutrophil count (ANC)>0.5/nl for afebrile patients and after 4 days of ANC>0.5/nl for those who are persistently febrile.
  - Isavuconazole may be considered in high-risk patients for mold-active prophylaxis, particularly when Mucorales coverage is needed, or where azole toxicity or QTc prolongation is a concern, although it is not currently included as a first-line prophylactic agent in ECIL guidelines.

## Background

Despite the recent development of novel and extended spectrum antifungal antibiotics, invasive fungal infections remain a significant cause of morbidity and mortality in stem cell transplant recipients. Mortality with these infections remains extremely high.

These antifungal standard practice recommendations derive primarily from:

- European Conference on Infections in Leukemia (ECIL 10 guidelines 2025)<sup>1</sup>;
- 2016 Aspergillosis<sup>2</sup> and candidemia<sup>3</sup> treatment guidelines of the Infectious Diseases Society of America;
- Analysis of the important supporting literature; and

- Local considerations (fungal epidemiology, drug availability, ongoing construction projects)

## Primary Prophylaxis

Primary antifungal prophylaxis is indicated for populations at high risk of developing invasive fungal disease, those being leukemic patients receiving chemotherapy, and allogeneic hematopoietic stem cell transplant (HSCT) recipients. Conceptually, prophylactic recommendations for the allogeneic HSCT population have been divided into the early neutropenic and the GVHD phases<sup>1</sup>.

The risk of invasive candidiasis is greatest in the early post-transplant period (phase I) due to the presence of neutropenia, severe mucositis, and central venous catheter use. In the post-engraftment period (phase II and III), few HSCT recipients require prophylaxis against *Candida* species, unless gastrointestinal GVHD or a central venous catheter (CVC) (the main risk factors) are present. Dissemination of endogenous *Candida* species colonizing the gastrointestinal (GI) tract is the usual cause of invasive candidiasis, although more rarely, it may be spread on the hands of healthcare workers. Autologous hematopoietic cell transplant (HCT) recipients have minimal risk for invasive candidiasis once neutropenia and mucositis resolve.

The risk of mold infection, while higher during the GVHD phase, is also relevant during the initial neutropenic phase. During phase I, prolonged neutropenia, active leukemia and prevalence >8% are the main risk factor for mold infection, being higher in bone marrow and umbilical cord blood transplants, and lower in nonmyeloablative and peripheral blood transplants. In phase II and III, cell-mediated immunodeficiency caused by GVHD and its treatment is the main risk factor, especially in those receiving unrelated donor, mismatched or haploidentical transplants.

For these reasons, even though fluconazole is highly recommended in the initial neutropenic phase in low risk populations, it should be used when combined with a mold-directed diagnostic approach (i.e. galactomannan or CT-based) or a mold-directed therapeutic approach (i.e. empiric antifungal therapy) in high risk populations. Of note, a number of prospective and retrospective studies (as cited below) have evaluated various mold-active antifungals versus fluconazole as primary prophylaxis in the neutropenic phase and have failed to demonstrate differences in overall survival.

Primary antifungal prophylaxis in the neutropenic phase at our center is with fluconazole for 28 days due to a low incidence of invasive mold infection (<4% in past 3yrs). It should start from the end of the conditioning regimen. In high risk patients (UCB, active leukemia, prolonged neutropenia, prolonged steroid exposure), serum galactomannan monitoring twice a week during neutropenia will be added or voriconazole/posaconazole prophylaxis can be considered. If galactomannan screening is positive (defined by optical density  $\geq 0.5$  on two consecutive occasions) it will be followed by CT imaging +/- bronchoscopy, followed by anti-aspergillus therapy if proven or probable aspergillosis. Maertens et al. have demonstrated that such a fluconazole plus galactomannan monitoring approach can be highly successful<sup>46</sup>.

## Allogeneic HSCT Recipients, Initial Neutropenic Phase

**Table 1.** ECIL recommendations on primary antifungal prophylaxis in adult allogeneic HSCT recipients: re-engraftment period\*

| Antifungal Agent                                                                                              | Pre-engraftment risk of mold infections |         | ECIL 5-6         |               |
|---------------------------------------------------------------------------------------------------------------|-----------------------------------------|---------|------------------|---------------|
|                                                                                                               | low                                     | high    | low              | high          |
| Fluconazole 400 mg q24h                                                                                       | A-I <sup>a</sup>                        | D-III   | A-I <sup>a</sup> | A-III against |
| Posaconazole tablet 300 mg q24h following a loading dose of 300 mg q12h on day 1 or oral solution 200 mg q8h  | B-II                                    | B-II    | B-II             | B-II          |
| Itraconazole 2.5-7.5 mg/kg q24h i.v. or 200 mg q24h i.v. or p.o.                                              | B-I                                     | B-I     | B-I              | B-I           |
| Voriconazole 6 mg/kg q12h first day then 4 mg/kg q12h i.v. or p.o.                                            | B-I                                     | B-I     | B-I              | B-I           |
| Micafungin 50 mg q24h                                                                                         | B-I                                     | C-1     | B-I              | C-1           |
| Caspofungin and anidulafungin                                                                                 | no data                                 | no data | no data          | no data       |
| Liposomal amphotericin B                                                                                      | C-II                                    |         | C-II             | C-II          |
| Aerosolized liposomal amphotericin B (10mg twice weekly) in combination with systemic fluconazole 400 mg q24h | C-III                                   |         | C-III            | B-II          |
| Isavuconazole 200 mg q24h following a loading dose of 200 mg q8h on days 1 and 2 <sup>b</sup>                 | B-II                                    | B-II    | no data          | no data       |

<sup>a</sup> only when combined with a mold-directed diagnostic approach (biomarker and/or CT scan-based) or a mold-directed therapeutic approach (empirical antifungal therapy).

<sup>b</sup> Isavuconazole can be used as second-line mold active prophylaxis, in case of intolerance to posaconazole/voriconazole, or QTc prolongation. Pre-engraftment risk of mold infection as previously defined: high risk includes active leukemia, cord blood transplantation and unrelated donor. Haplo-identical HSCT using post-transplantation cyclophosphamide should be considered at low risk (B-II) In case of prior IFD, secondary prophylaxis should be tailored according to the previous documentation.

*HSCT hematopoietic stem cell transplantation.*

\* From Pagano L. et al: Primary antifungal prophylaxis in hematological malignancies. Updated clinical practice guidelines by the European Conference on Infections in Leukemia (ECIL)Antimicrob Chemother 2018; 73: 3221–3230. Leukemia 2025.

## Allogeneic HSCT Recipients, GVHD Phase

While fluconazole, itraconazole and voriconazole have been studied through the initial neutropenic phase and into the GVHD phase, posaconazole and fluconazole are the only anti-fungals that have been studied specifically in the setting of significant GVHD (grade 2-4 acute or extensive chronic). This was in a head to head prospective, randomized, placebo-controlled trial which revealed reduced

proven/probable invasive aspergillosis and fewer deaths from invasive fungal infection in the posaconazole group. Overall survival and treatment-related adverse effects were similar<sup>36</sup>.

**In the setting of grades 3-4 acute GVHD (aGVHD), a prophylactic posaconazole strategy can be justified on a number of levels:**

- Patients with aGVHD continue to have risk factors for invasive candidiasis, i.e. central venous catheter, potential GI aGVHD involvement, recently healed/healing conditioning-related mucositis.
- The recent large (1800 patients) Italian prospective observational study demonstrated that grade 2-4 aGVHD remains an independent significant risk factor for invasive fungal infection (IFI) (hazard ratio of 6), predominantly invasive aspergillosis.
- In the two most recent trials of mold-active anti-fungal (posaconazole and voriconazole) vs fluconazole (+galactomannan monitoring) essentially performed equally well. In the voriconazole trial, there was no difference in fungal-free survival or overall survival and a majority of invasive aspergillus infections in the fluconazole arm were picked up by galactomannan screening. While the posaconazole trial demonstrated a reduction in death from fungal infection in the posaconazole arm.
- This will be a relatively small number (19) high risk patient population.

**In the setting of chronic GVHD (cGVHD) requiring immunosuppression there are little data to guide prophylaxis:**

- In the posaconazole trial, the rate of IFI in those with cGVHD was low in both arms and there was no significant benefit of posaconazole (5% in the posaconazole arm vs. 6% in the fluconazole arm). Details of the cGVHD were not provided.
- The prospective Italian study revealed a striking difference in the incidence of IFI in those with de novo cGVHD (3.2%) versus those with cGVHD preceded by acute GVHD (19.4%)
- There are no studies evaluating a galactomannan screening approach in patients with cGVHD and this approach is impractical to apply as these patients do not routinely have weekly lab work/follow-up.
- Patients with cGVHD are likely not at high risk of invasive candida infections and therefore there is likely limited benefit to fluconazole prophylaxis.

**Autologous HSCT recipients**

There is no evidence for primary prophylaxis improving outcomes after autologous transplantation. Therefore, we do not use it routinely. Based upon expert opinion only, prophylaxis may be considered for autologous HCT recipients who have, or are expected to have, the following conditions:

- Prolonged neutropenia and mucosal damage from intense conditioning regimens or graft manipulation
- Receipt of fludarabine or 2-CDA (2-chlorodeoxyadeno-sine) within 6 months of HCT



## CAR T-cell therapy recipients

Invasive fungal infections are rare after CAR-T cell therapy and typically occur in patients with other risk factors such as prolonged neutropenia or additional immunosuppression<sup>62</sup>. There is little evidence to choose between an anti-yeast or anti-mold prophylactic strategy in this population<sup>63</sup>. Consensus guidelines from EBMT, ASTCT, CARTOX, and other groups recommend fluconazole prophylaxis during the severe neutropenic period (ANC <0.5). Anti-mold prophylaxis is recommended in the following patients:

- Recent allogeneic stem cell transplant
- Prior invasive mold infection
- Prolonged neutropenia >14 days
- Prolonged corticosteroid use >72 hours

## Secondary Prophylaxis

Patients who received treatment for suspected or proven invasive fungal infection earlier in their disease course are at high risk of recurrent infection during subsequent treatment. The goal of secondary prophylaxis is to prevent relapse of prior invasive fungal disease, or the occurrence of another invasive fungal disease during a new high risk period (prolonged neutropenia, or a period of severe immunosuppression). No randomized clinical trials exist to guide choice of secondary prophylaxis, and no standard approach exists. Small retrospective studies have been published using liposomal amphotericin B, voriconazole, and caspofungin<sup>37-39</sup>. Benefit from secondary antifungal prophylaxis has been suggested by two large retrospective studies of allogeneic HSCT recipients<sup>40,41</sup>, and a prospective study of voriconazole in this population<sup>42</sup>. No randomized clinical trials have been conducted.

The choice of antifungal agent should be based on: 1) the etiologic agent identified from the previous episode of invasive fungal disease; and 2) the previous response to antifungal agents (ECIL 7)<sup>1</sup>.

Where ongoing antifungal therapy is considered prudent, clinicians must be mindful of drug interactions, especially between azoles, calcineurin inhibitors and QT intervals.

## Empiric Antifungal Therapy during Febrile Neutropenia

Early studies demonstrated that treatment of neutropenic patients with persistent or recurrent fever (variously defined as fever after 4 - 7 days of broad-spectrum antibacterial therapy) with amphotericin B reduced the incidence of documented invasive fungal infection and improved survival<sup>44</sup>. This has led to a strategy of empiric antifungal therapy for patients with persistent fever in neutropenia, and over time the agent of choice has moved away from amphotericin B deoxycholate to less toxic alternatives.

Several principles guide the choice of initial empiric antifungal therapy:

- Liposomal amphotericin B (L ampho B) is as effective as amphotericin B deoxycholate (AMBd), with fewer breakthrough infections at completion of therapy. There are also fewer infusion-related adverse events (IRAEs) and less nephrotoxicity<sup>45</sup>. AMBd receives a D1 grading in the presence of risk factors for renal toxicity and should be avoided<sup>1</sup>.
- Caspofungin is as effective as L ampho B in empiric treatment of suspected invasive fungal infections<sup>48-50</sup>
- Voriconazole actually failed the 10% non-inferiority cut-off when compared with L ampho B for empiric therapy and did not receive FDA approval for this indication. It is included in the table below because it is superior to AMBd for the treatment of IA<sup>52</sup>, effective therapy for candidiasis, and efficacious for prevention of break through invasive fungal disease<sup>53</sup>.
- Fluconazole has no activity against *Aspergillus* species or other molds, and is not approved by the FDA for this indication.
- Only amphotericin B preparations, posaconazole and isavuconazole would be expected to have activity against *Mucorales* species.

The caveat is that empiric antifungal therapy has never been directly compared with placebo or other antifungal strategies. Less desirable aspects of this strategy include over-treatment of patients without invasive fungal disease, with the associated side effects and costs. The strategy is also limited by the fact that fever is a non-specific marker of fungal infection and will miss invasive fungal disease not associated with fever (estimated to be approximately 7% from the preemptive strategy literature<sup>46</sup>).

**Table 2.** Dose and grading of antifungal agents

| Antifungal Agent              | Daily Dose          | ECIL 7 Grading <sup>1</sup> |
|-------------------------------|---------------------|-----------------------------|
| L ampho B <sup>47,48</sup>    | 3-5 mg/kg           | AI                          |
| Caspofungin <sup>48-50</sup>  | 50 mg               | AI                          |
| Itraconazole <sup>51</sup>    | 200 mg i.v.         | BI                          |
| Voriconazole <sup>52,53</sup> | 2 X 6 mg/kg i.v./po | AI                          |
| Posaconazole <sup>60</sup>    | 300x2/300mg         |                             |
| Isavuconazole <sup>61</sup>   | 372q8hx3/372mg      |                             |
| Micafungin <sup>54,55</sup>   | 100mg               | BII                         |
| AMBd <sup>47</sup>            | 0.5 - 1 mg/kg       | BI/DI                       |
| Fluconazole <sup>56</sup>     | 400 mg i.v.         | CI                          |

For patients with prolonged antibiotic resistant fever in neutropenia (>4 days of fever despite appropriate antibacterial coverage and no clinical or radiographic focus of infection) empiric antifungal therapy with L ampho B or Caspofungin/Micafungin will be added. Axial imaging studies (equivalent

to HRCT of chest, and ultrasound/CT abdomen and pelvis) will be carried. If these studies fail to demonstrate a clinical focus, treatment with G-CSF will also be instituted.

Empiric antifungal coverage should be discontinued in afebrile patients once ANC > 0.5 for two days. In patients with persistent fever and no clinical or radiographic focus of infection, empiric treatment with antifungal antibiotics should be discontinued once ANC > 0.5 for four days. Alternative causes including CVC infection, drug fever and GVHD should also be considered.

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# Graft Failure, Poor Graft Function, Chimerism

Presented by: Jan Storek and Adam Bryant

## Summary

- **Primary Graft Failure**, after excluding relapse, medications, or viral infections, is treated by a 2nd HCT, except if palliation is appropriate.
  - Palliation is appropriate in patients with poor prognosis malignancy at 1<sup>st</sup> HCT or poor performance status/elderly.
  - Choose a haploidentical or another suitable donor available ASAP.
    - Can be the same or different donor than for the first HCT.
  - Graft: Filgrastim-mobilized PBSCs
  - Conditioning: Fludarabine 150 mg/m<sup>2</sup> + Cyclophosphamide 29 mg/kg + TBI 2Gy
  - GVHD prophylaxis: Posttransplant cyclophosphamide (PTCy) + tacrolimus + MMF
- **Poor Graft Function or Secondary Graft Failure**, after excluding relapse, medications, and viral infections, can be treated with a TPO agonist, or a filgrastim-mobilized and CD34 cell-enriched PBSC boost.
  - If absent or declining donor chimerism (suggesting graft rejection), PBSC boost should not be given. Instead, consider 2<sup>nd</sup> HCT as for primary GF.
- **Chimerism** of blood T cells (CD3+) and blood malignancy lineage cells (CD13/33+ cells in case of myeloid leukemia or CD19+ cells in case of B cell malignancy) is routinely determined in allotransplant recipients at 3 months. This is to:
  - assess risk for relapse
  - generate baseline values for potential later chimerism testing (when relapse or graft rejection is suspected).
  - No intervention for low donor chimerism alone should be done, as no intervention with level A, B, or C evidence exists, that could decrease the risk of relapse or graft failure.

## Nomenclature

We use the following definitions, endorsed by EBMT, ASTCT, CIBMTR, and APBMT<sup>1,2</sup>:

- Neutrophil recovery: The first of 3 successive days with ANC  $\geq 0.5/\text{nl}$  after post-HCT nadir.
- Platelet recovery: The first of 3 consecutive days with PLT  $\geq 20/\text{nl}$  in the absence of platelet transfusion for 7 consecutive days.
- Graft Failure (GF), which may be due to graft rejection or other causes:
  - Primary GF: Lack of achievement of ANC  $\geq 0.5/\text{nl}$  with associated pancytopenia by day 30 (PBSC or BM) or by day 42 (cord blood).
  - Secondary GF: Decline in hematopoietic function necessitating blood products or growth factor support, after neutrophil and platelet recovery.
- Poor graft function (PGF): Frequent dependence on transfusions and/or growth factors in the absence of other explanations, such as relapse, medications, or infections.
- Donor chimerism
  - Full:  $>95\%$  for both myeloid and lymphoid lineages
  - Mixed (partial): 5-95% for both myeloid and lymphoid lineages
  - Absent:  $<5\%$  for both myeloid and lymphoid lineages

## Graft Failure

- Risk factors<sup>3-8</sup>
  - Non-malignant disease
  - Cell dose (CD34 or TNC)
  - Graft type (PBSC  $<$  BM  $<$  cord blood)
  - Low-intensity conditioning
  - GVHD prophylaxis (PTCy  $>$  CN1+antimetabolite or ATG-based)
  - HLA disparity
  - ABO major mismatch
  - Donor-Specific Antibodies
    - Risk may be reduced by desensitization (eg, plasmapheresis, IVIG, rituximab)
  - Splenomegaly
- Management of primary GF<sup>3</sup>
  - Rule out relapse, medications, or viral infections
  - If responsive to growth factors, treat as PGF (below), unless absent or clearly declining donor chimerism
  - Determine eligibility for 2<sup>nd</sup> HCT. Risk factors for poor outcome include:
    - Poor prognosis underlying disease<sup>7,9</sup>, and
    - Older age / poor performance status/organ dysfunction<sup>9,10</sup>

- Per Japanese registry study<sup>10</sup>, no combination of risk factors resulted in zero OS. Most impactful risk factors were:
  - Age ≥55
  - ECOG 4 (no self-care, confined to bed or chair)
- 2<sup>nd</sup> HCT, using PBSCs:
  - Survival 10-30% irrespective of conditioning, GVHD prophylaxis, and whether original vs new donor. This is per large single-center or registry analyses, usually from pre-haplo/PTCy era<sup>3,5,6,9</sup>.
  - Higher survival has been reported in small, usually single-center or registry studies of salvage haploidentical HCT, allowing short interval from 1<sup>st</sup> to 2<sup>nd</sup> HCT
    - Due to chance or publication bias?
  - Duke Univ group (senior author Rizzieri) reported in 2012 on 11 adult patients receiving fludarabine+cyclophosphamide+2GyTBI+alemtuzumab, all on day -1<sup>11</sup>. With a median F/U of 1 year, 8/11 (73%) survived.
    - However, in their subsequent report on a total of 28 patients with a median follow up of 3 years, survival at 4 years was only ~20%<sup>12</sup>.
    - Median time from 1<sup>st</sup> to 2<sup>nd</sup> HCT was ~42 days.
  - PTCy-based GVHD prophylaxis appears associated with good OS, but based on small studies with short F/U
    - Prata *et al* reported in 2019 on 23 mostly adult patients from French registry receiving fludarabine-based conditioning (usually Flu+Cy+TBI, doses not given) and PTCy-based GVHD prophylaxis. Median F/U was not given. OS at 1 y was 56%.
      - Median time from 1<sup>st</sup> to 2<sup>nd</sup> HCT was 63 days.
    - Albert *et al* reported in 2021 on 12 **pediatric** patients receiving fludarabine+treosulfan+cyclophosphamide+rituximab+alemtuzumab and PTCy-based additional GVHD prophylaxis<sup>13</sup>. With a median F/U of 2 years, 11/12 patients survived.
      - 3 patients had GF diagnosed before d28.
      - Median time from 1<sup>st</sup> to 2<sup>nd</sup> HCT 46 days.
    - Giammarco *et al* reported in 2021 on 19 adult patients receiving fludarabine+cyclophosphamide+2GyTBI with PTCy-based GVHD prophylaxis<sup>14</sup>. With a median F/U of <1 year, 1-year survival was 66%.
      - 14/19 pts surviving but, 3 with GF, so survival at 1 y will be at best 11/19 (58%)
      - Median time from 1<sup>st</sup> to 2<sup>nd</sup> HCT 42 days.
    - Harada *et al* reported in 2022 on 33 adult patients from Japanese registry receiving fludarabine+cyclophosphamide (or melphalan, busulfan, or TBI instead of cyclophosphamide) and PTCy-based

GVHD prophylaxis.<sup>15</sup> With a median F/U of 1 year, 1-year survival was 47% (61% in a subgroup of patients with a regular (not reduced) dose of PTCy of 50 mg/kgx2).

- Median time from 1<sup>st</sup> to 2<sup>nd</sup> HCT 49 days.
- In summary, 2<sup>nd</sup> HCT within 40 to 50 days from 1<sup>st</sup> HCT, thus usually from a haploidentical donor, and using PTCy-based GVHD prophylaxis appears appealing, but F/U so far too short.
- Calgary results with Fludarabine+5GyTBI: 10/11 died within 1 year post-2<sup>nd</sup> HCT, so only 1/11 survived
  - 1/10 who got ATG-based GVHD prophylaxis (in 2001-2023)
  - 0/1 who got PTCy-based GVHD prophylaxis (in 2024)
- New conditionings & GVHD prophylaxis for Calgary (starting July 2024):
  - Fludarabine 50 mg/m<sup>2</sup> days -4, -3, -2
  - Cyclophosphamide 14.5 mg/kg days -4, -3
  - TBI 2 Gy day -1
  - Standard PTCy + tacro + MMF
  - This is based on Giammarco *et al*, but abbreviated, and tacro used instead of CSA. The original Giammarco *et al* (2021) conditioning is:
    - Fludarabine 30 mg/m<sup>2</sup> days -6, -5, -4, -3, -2
    - Cyclophosphamide 14.5 mg/kg days -6, -5
    - TBI 2 Gy day -1
    - Standard PTCy + tacro + MMF
- Management of **secondary** GF has not been adequately studied.
  - If absent or clearly declining mixed donor chimerism, consider 2<sup>nd</sup> HCT as for primary GF.
  - If full chimerism or stable/increasing mixed chimerism, consider PBSC boost as for PGF.

## Poor Graft Function

- Risk factors<sup>3,16</sup>
  - Same as for GF, plus
  - GVHD
  - SOS/VOD
- Management<sup>3,16</sup>
  - Growth factors
    - Thrombopoietin (TPO) agonist
      - In 5 recent case series, each with 11-48 (total 97) adult alloHCT patients with PGF, eltrombopag at doses of 50-150 mg at variable timing post alloHCT (1-14 months) resulted in overall and complete responses of 58-

83% and 50-67%, respectively, without severe toxicity.<sup>17-21</sup> Median time to response varied from 3-10 weeks for platelets and 3-28 weeks for all lineages. Patients who achieved CR maintained responses for the duration of study follow-up (10-43 months) and some were able to stop TPO agonist without reoccurrence of PGF.

- Given a signal for potential efficacy and a good safety profile and despite the studies lacked controls, we recommend a trial of TPO agonist in all patients with PGF.
- See also Cytopenia and Transfusion chapter
- Filgrastim-mobilized and CD34 cell-enriched PBSC boost.
  - Efficacy and toxicity in recent studies is summarized in Table 1.
  - CR was reported in 62% to 92% patients.
    - Onset of rising blood counts in ~2 weeks
    - Maximum response in 1-3 months
  - Toxicity is acceptable – aGVHD in 0-23% patients, usually grade 1-2, and cGVHD in 0-50% patients, usually mild to moderate.
  - Given the probable efficacy and acceptable toxicity and despite the studies lacked controls, we recommend a PBSC boost in patients who failed a TPO agonist.
    - Recommended dose is  $5 \times 10^6$  CD34 cells/kg, infused fresh<sup>22</sup>.
  - Chimerism: All studies in Table 1 used patients with full donor chimerism. In one study (Cuadrado *et al*),<sup>23</sup> patients with mixed chimerism were also included. Their outcome did not differ from those with full chimerism.
    - We recommend giving PBSC boost to patients with full chimerism or stable/increasing mixed chimerism.
    - We do not recommend giving PBSC boost to patients with absent or declining donor chimerism. This is due to the theoretical concern that the PBSC would be rejected.
  - Before PBSC boost, the following should be ruled out: relapse and medication- or viral infection-induced cytopenia.

**Table 1.** Summary of publications describing outcome of CD34-selected stem cell boosts in adults with poor graft function.

| Publication                    | N  | Time <sup>a</sup> | Content                                                            | Response <sup>b</sup> | GVHD <sup>c</sup>      | Survival          |
|--------------------------------|----|-------------------|--------------------------------------------------------------------|-----------------------|------------------------|-------------------|
| Stasia 2014 <sup>24</sup>      | 41 | NR                | CD34 3.45 x 10 <sup>6</sup> /kg<br>CD3 2.5-10 x10 <sup>3</sup> /kg | 76% CR<br>7% PR       | 22% aGVHD<br>7% cGVHD  | 63% 3-y           |
| Haen 2015 <sup>25</sup>        | 20 | NR                | CD34 4.6 x 10 <sup>6</sup> /kg<br>CD3 2 x10 <sup>3</sup> /kg       | 92% CR                | 5% aGVHD<br>NR cGVHD   | 53% 2y            |
| Klyuchnikov 2014 <sup>26</sup> | 32 | 150               | CD34 3.4 x 10 <sup>6</sup> /kg<br>CD3 9 x10 <sup>3</sup> /kg       | 81% CR<br>0% PR       | 19% aGVHD<br>19% cGVHD | 45% 2y            |
| Askaa 2014 <sup>27</sup>       | 18 | 113               | CD34 4.9 x 10 <sup>6</sup> /kg<br>CD3 11 x10 <sup>3</sup> /kg      | 72% CR                | 22% aGVHD<br>50% cGVHD | 48% 2y            |
| Ghobadi 2017 <sup>22</sup>     | 26 | NR                | CD34 5.0 x 10 <sup>6</sup> /kg<br>CD3 NR                           | 62 % CR<br>19% PR     | 23% aGHVD<br>31% cGHVD | 65% 1y            |
| Mohty 2019 <sup>28</sup>       | 10 | 120               | CD34 4.8 x 10 <sup>6</sup> /kg<br>CD3 1.3 x 10 <sup>3</sup> /kg    | 70% CR<br>30% PR      | 0 % aGVHD<br>0% cGVHD  | 80% 1y<br>70% 3y  |
| Cuadrado 2020 <sup>23</sup>    | 62 | 450               | CD34 5.0 x 10 <sup>6</sup> /kg<br>CD3 4.3 x 10 <sup>3</sup> /kg    | 62 % CR<br>13% PR     | 11% aGVHD<br>8% cGVHD  | 70% 1y<br>54% 5 y |

<sup>a</sup> Median time (days) from stem cell transplant to infusion of CD34 selected cells; NR = Not reported

<sup>b</sup> Individual study response definitions varied. Response definitions presented here CR = hematologic improvement in all 3 lineages without transfusion dependence PR = hematologic improvement in 1-2 lineages

<sup>c</sup> aGVHD grade II-IV; cGVHD total (limited + extensive; mild + moderate + severe)

## Chimerism Techniques

Chimerism (% cells of donor versus recipient origin) can be determined using one of the techniques described in Table 2.

In Calgary, we use fluorescent dye-labeled short tandem repeat (STR) polymorphism, multiplexed (a total of 16 polymorphic genomic segments are assayed), and analyzed by capillary electrophoresis. The reasons are the very high quantitation accuracy, which facilitates comparison of current result to previous result(s), high informativeness (which means that chimerism can be reliably determined in >99% donor-recipient pairs), acceptable sensitivity (no data exist suggesting that sensitivity below 1-5% is clinically valuable), and applicability to all donor-recipient pairs (irrespective of sex matching).

The principle of the assay is explained using the following example: in a short tandem repeat segment of genome, the donor has 3 tetranucleotide repeats (GCTG GCTG GCTG) on both paternal and maternal chromosomes whereas the recipient has 4 tetranucleotide repeats (GCTG GCTG GCTG GCTG) on both paternal and maternal chromosomes (simple scenario, as in reality most persons are heterozygous). In a post-transplant patient specimen, the segment of genome is amplified by PCR, using primers for conserved sequences flanking the segment. The PCR product is subjected to electrophoresis, which separates the 3 tetranucleotide repeat amplicons from the 4 tetranucleotide repeat amplicons (the former amplicons move faster). As the amplicons are fluorescent dye-labeled, the ratio of donor to recipient chimerism is determined as the ratio of



fluorescence of the donor (3 repeat) amplicons to the fluorescence of the recipient (4 repeat) amplicons.

**Table 2.** Techniques to determine chimerism (courtesy of F. Khan).

| Technique                                                         | Analytical Sensitivity (%) | Quantitation Accuracy | Informativeness (likelihood of finding alleles different between donor & recipient)                 |
|-------------------------------------------------------------------|----------------------------|-----------------------|-----------------------------------------------------------------------------------------------------|
| Fluorescent dye-labeled STR, multiplex, capillary electrophoresis | 1-5                        | Very High             | High                                                                                                |
| <sup>32</sup> P-labeled STR/VNTR, multiplex, gel electrophoresis  | 1-5                        | Moderate              | High                                                                                                |
| XY Cytogenetics                                                   | 10-20                      | Low                   | Sex-mismatched only                                                                                 |
| XY FISH                                                           | 0.1-0.2                    | Very High             | Sex-mismatched only, potential origin of sex-mismatched cells from transfusion, mother or offspring |
| RFLP                                                              | 5-20                       | Moderate              | Moderate                                                                                            |
| Real time PCR using 'indels' (insertion/deletion polymorphism)    | 0.001-0.1                  | Moderate              | Moderate-High                                                                                       |

## Chimerism Clinical Utility

- Chimerism of blood cells **can** be used for:
  - Diagnosis of graft rejection (with or without autologous reconstitution)
    - Rejection is defined as <5% donor cells among both myeloid cells and T cells, in the absence of relapse.
  - Risk assessment for relapse:
    - In Alberta, we routinely determine chimerism of sorted blood malignancy lineage cells and T cells at 3 months post-transplant (baseline). Subsequent chimerism determination can be done when/if rejection or relapse is suspected. Low % donor cells at 3 mo is only a risk factor for relapse, it is not diagnostic of relapse. Per our analysis of patients undergoing first allo-HCT in Alberta between 2010 and 2020<sup>29</sup>, most of whom received Flu+Bu+ATG+4GyTBI conditioning, PBSCs, and additional GVHD prophylaxis with MTX+CSA,
      - Patients with full T cell chimerism at 3 mo had a lower risk of relapse (22% vs 36%)
        - PPV was only 34% and NPV 81%

- Patients with full chimerism among leukemia lineage cells (eg, CD13/33+ cells in case of AML or CD19+ cells in case of B-ALL) at 3 mo had a lower risk of relapse (20% vs 45%)
  - PPV was only 45% and NPV 79%
- Chimerism of blood cells alone **cannot** be used as an indication for an intervention.
  - No agreement among experts on which intervention, if any, should be used<sup>1</sup>.

**Table 3.** Interpretation of blood chimerism results.

|                                       | % Donor Among Blood                 |                               |
|---------------------------------------|-------------------------------------|-------------------------------|
|                                       | CD3 Cells                           | Leukemic Lineage Cells        |
| Normal                                | >95%                                | >95%                          |
| Benign mixed chimerism                | 5 – 95%, stable or increasing       | 5 – 95%, stable or increasing |
| Graft Rejection**                     | <5%*                                | <5%                           |
| Impending relapse or bonified relapse | Typically >95% or stable/increasing | Decreasing***                 |

\* In Alberta (using MAC, PBSCs, and ATG-based GVHD prophylaxis), <5% donor T cells with >95% donor myeloid cells does not appear to be a risk factor for rejection (Storek, unpublished, 2023).

\*\* With or without autologous hematopoietic reconstitution.

\*\*\* Per preliminary analysis on a small number of Albertan patients transplanted in 2010-2020 (performed by Storek in May 2021), >5% drop of donor chimerism may predict relapse with PPV=79% and NPV=81%.

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# Donor Lymphocyte Infusion (DLI) or Second Allogeneic HCT for Relapse

Presented by: Adam Bryant

## Summary

- Patients who relapse after stem cell transplant have poor prognosis.
- Patients with acute leukemia relapsed after transplant may be considered for donor lymphocyte infusion (typically with chemotherapy), second allotransplant, palliative chemotherapy, targeted immunotherapy, clinical trials, or palliative care.
- Fit patients with relapsed B-acute lymphoblastic leukemia should be considered for CAR T-cell therapy<sup>1, 2</sup>.
- Criteria for eligibility for 2<sup>nd</sup> allo-HCT:
  - >1 year after first transplant for acute leukemia or MDS, and
  - Favorable additional risk factors:
    - Young (preferably age < 40)
    - Fit (KPS ≥ 80 or ECOG 0-1)
    - Disease in remission at time of second transplant
- Selected patients with AML or MDS may be candidates for DLI. These include patients with the following characteristics
  - with hematologic relapse (>5% marrow blasts, peripheral blood blasts, extramedullary disease)
    - < 1 year post alloHCT
    - > 1 year post alloHCT and non-candidate for second alloHCT
  - with minimal residual disease that is
    - defined by disease-identifying cytogenetic, morphologic, molecular, or immunophenotypic means (not incomplete donor chimerism)
    - documented on
      - 3-month post-transplant marrow **AND** demonstrated to be worsening at subsequent time point(s) 4-6 weeks later
      - documented at clinical suspicion of relapse beyond 3 months (and if deemed necessary by clinician discretion) demonstrated to be worsening at subsequent time point(s)
  - without options for clinical trials or targeted chemo/immunotherapy
  - without history of gr3-4 aGVHD or mod-sev cGVHD, and without any active GVHD
  - who are motivated and fit
- When administered without chemotherapy, donor lymphocytes should be administered every 1-2 months based on disease response and the presence or absence of GVHD.

- Donor lymphocytes for DLI should be collected in a single apheresis session and divided into three aliquots of the following cell doses:
  - $1 \times 10^6$  CD3+ cells/kg (infused fresh, others cryopreserved in 10% DMSO)
  - $1 \times 10^7$  CD3+ cells/kg
  - $1 \times 10^8$  CD3+ cells/kg
- Options for patients with relapsed MPN (with < 5% blasts), CML or CLL include novel therapies (later-generation TKIs, B-cell receptor antagonists, BCL-2 inhibitors) if the patient has not previously been exposed to them. Other options include DLI or palliative/supportive care. Second allogeneic transplants will be rare in this population.

## Background

Despite the use of intensive, myeloablative conditioning, relapse remains the most common cause of treatment failure following allogeneic and autologous stem cell transplantation. Selected patients with chemosensitive disease may be considered for repeat transplants. Criteria for patient selection are reviewed below.

### Acute Leukemia

The natural history of acute leukemia that has relapsed following allogeneic bone marrow transplantation has a grim prognosis, with 2-year survival estimates consistently less than 20%<sup>3</sup>.

Aside from conventional chemotherapy, as described above, non-transplant options for acute leukemia patients who relapse after allogeneic transplant include cellular therapy in the form of donor lymphocyte infusion (DLI). While AML is of intermediate sensitivity to DLI (reported response rates vary from 0 – 60%), most patients treated in this way do not experience prolonged survival due to graft-versus-host disease, infection and relapse. Despite the sensitivity of ALL to graft-versus-leukemia effects, responses to DLI in this disease are almost never seen and tend to be short-lived.

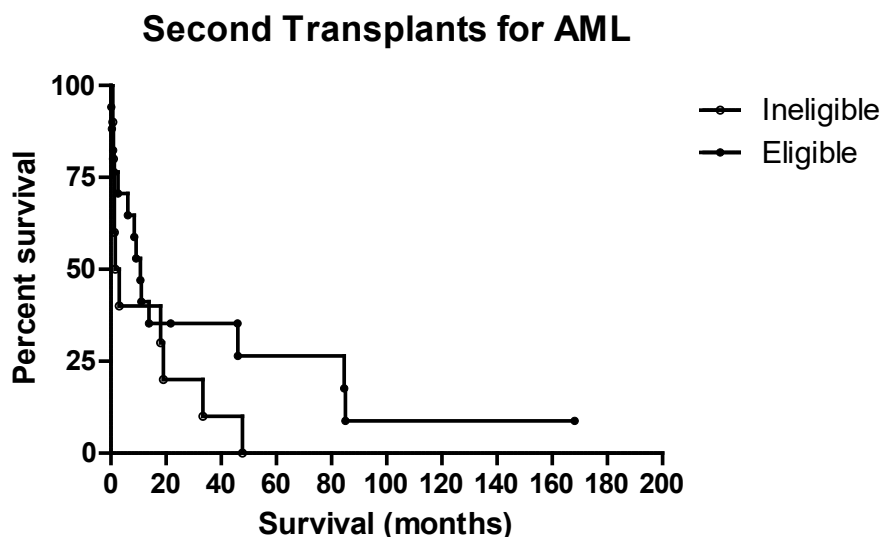
Second allogeneic transplants have been offered to highly selected patients with acute leukemia that has relapsed after a prior transplant. Overall survival following a second allogeneic transplant is limited by high TRM (30-36%) and frequent relapses (44-70%). Most reports describe 2-year overall survival between 18-44%<sup>4-16</sup>. While second transplants may be of benefit to some patients who relapse, it is clear that they are only offered to a minority: in a review of second transplants carried out for the CIBMTR by Eapen *et al.*, only 6% of acute leukemia patients reported to the registry who relapsed received a second transplant<sup>11</sup>. In registry studies and case series, uni- and multivariable analysis of risk factors associated with relapse and survival outcomes have identified common favourable risk factors including

- younger age (variable cutoffs)<sup>6, 7, 13</sup>
- fitness (ECOG 0-1, KPS  $\geq$  80)<sup>4, 8, 11, 14</sup>
- disease remission at second alloHCT<sup>5, 9-14</sup>

- time to relapse from first alloHCT (most >12 mo) <sup>4-8, 12-15</sup>

Several reports have described the negative effect of rapid relapse after allogeneic transplant, many with a cutpoint of 12 months<sup>4-8, 12-15</sup>. Results from our program, shown in the figure below, suggest that patients retransplanted within one year of an allogeneic transplant experience poor outcomes. It is reasonable to take this as a cutoff.

**Figure 1.**



*Outcome of second allogeneic transplants performed in Calgary for patients with AML who have relapsed following a prior allogeneic transplant. Eligible patients (top line) are those who remained in remission for > 1 year following their initial transplant.*

In one 2023 report by Lu *et al.* of 199 acute leukemia cases, risk factors identified in multivariable analysis appeared to be additive. In their report patients at second alloHCT with favourable risk factors including a) MRD -ve CR and b) an HCT-CI score of 0 and c) a second donor with mismatched haplotype, had a 2-year overall survival estimate of 63% compared to 43% in patients having two of the favourable risk factors and 20% in patients with none ( $p = 0.0001$ )<sup>10</sup>.

Given the inconsistency in risk associations reported across series, no one risk factor should necessarily be sufficient to preclude consideration of alloHCT. Based on the combined analysis of available registry studies and case series, and our own local experience, we advocate that patients be selected for second alloHCT if their disease remains in remission at for at least 12 months after transplant, and if on a case by case basis they are deemed more favourable based on the presence and number of favourable risk factors identified above.

The optimal conditioning intensity for second alloHCT remains an unresolved question. A retrospective EBMT registry study involving 215 AML patients undergoing second alloHCT noted no



significant difference in 2 year OS (31 v 40%,  $p = 0.41$ ), 2 year relapse risk (58 v 20% 51%,  $p = 0.24$ ), and 2 year NRM (15 v 28%,  $p = 0.89$ ) in patients who received reduced intensity versus myeloablative conditioning, respectively<sup>11</sup>. In 10 case series and registry studies involving 3,777 patients in which both RIC and MAC regimens were used<sup>4, 5, 7-9, 11-14, 16</sup>, conditioning was associated with survival outcomes in univariable (and not multivariable) analysis in just one 129-patient series<sup>4</sup>. This suggests that decreased TRM expected with reduced intensity conditioning is offset by its increased associations with relapse, ultimately have a net zero impact on overall survival. Barring concerning patient comorbidities, our center favours a myeloablative TBI-based conditioning regimen for second alloHCT, regardless of donor type.

The benefits of using a same or different stem cell donor for second alloHCT has been debated and explored. A recent large retrospective EMBT registry study involving 598 AML patients receiving second alloHCT looked at outcomes comparing recipients of a same MUD or MRD, different MUD/MRD, or different haplo donor. Ultimately they found no differences in 2 year OS (36 v 28 v 23%;  $p = 0.21$ ) relapse rate (51 v 49 v 44%;  $p = 0.90$ ) and NRM (25 v 37 v 34%;  $p = 0.28$ ) in comparing transplant recipients using the same matched donor, different matched donor, or different haplo donor<sup>17</sup>. Given that using a new or repeated donor does not appear to have clear impact on clinical outcomes, our centre favours selecting donors for second alloHCT instead based on availability and other standard donor factors, as summarized in this manual's Donor Selection chapter.

### **Use of Azacitidine and Donor Lymphocyte Infusions to Control Relapsed Leukemia**

The impact of donor lymphocyte infusions (DLI) on relapsed leukemia was first described by Kolb and coworkers in 1990<sup>18</sup>. They demonstrated that patients with cytogenetic or hematological relapse of CML could achieve a second disease-free state after the infusion of lymphocytes from their original stem cell donor. Graft-versus-host disease was observed in most responders. While DLI appeared to induce durable remissions in CML, responses in acute leukemia are uncommon. Augmenting DLI with chemotherapy increases response rates but also adds toxicity, without substantially prolonging survival. One exception may be combining DLI with azacitidine, a low-toxicity alternative to higher dose chemotherapy. Two reports have been published in sufficient detail to understand the outcome of this strategy:

- The German Cooperative Transplant Study Group (Schroeder *et al.*, 2015) described the results of a multicenter retrospective study of azacitidine plus planned DLI in 154 patients with myeloid disorders (AML (n=124), MDS (n=28) or MPN (n=2))<sup>19</sup>. Patients received azacitidine in a five-day 100 mg/m<sup>2</sup> or seven-day 75 mg/m<sup>2</sup> schedule, after which DLI was administered to 105 patients. Reasons for not administering DLI included progressive disease, coexisting GVHD, non-availability of the donor or achievement of CR with azacitidine alone. The overall response rate was 33% (27% CR, 6% PR) and two-year OS 29%. Factors associated with higher likelihood of survival include early relapse (molecular relapse only or fewer than 13% blasts at time of relapse) and a diagnosis of MDS. GVHD occurred in 31%. Given that some patients received the azacitidine+DLI for molecular relapse only and thus some of them may

have been cured ever without the azacitidine+DLI, it is unclear whether this therapy has a meaningful clinical efficacy. However, given the relatively low toxicity and the possibility of meaningful clinical efficacy, we are willing to offer this option to highly motivated patients.

- A subset of 30 patients in the above publication were described in a previous report<sup>20</sup>. These patients received azacitidine 100 mg/m<sup>2</sup>/day for five days every four weeks with escalating DLI after every second cycle. Twenty-two patients received DLI and seven (23%) achieved CR and two (7%) PR. Patients with MDS, AML with MDS-related change, early relapse and high-risk cytogenetics were more likely to show response.

At our center DLI was typically considered for patients with acute leukemia or MDS who have relapsed after alloHCT. Owing to local data describing inconsistent outcomes in patients with minimal residual disease documented on their day +100 bone marrow biopsy, our center originally avoided use of DLI for this subpopulation. In a prospective observation report by Shah *et al.* (BBMT 2018)<sup>21</sup> of 269 AML patients with MRD assessments done by 7-color flow cytometry on bone marrow aspirates at days 30, 100, and 180 post alloHCT, day +30 MRD positivity was strongly predictive of relapse incidence (HR 11, 95% CI 4.3-27; p <0.001) and of shorter OS (HR 4.3, 95% CI 2.0-9.6; p<0.001). These strong prognostic trends were also reported with patients MRD +ve at days 100 and 180. In original Schroder *et al.* report on 105 DLI recipients and a subsequent reports by Rautenberg *et al* using the same protocol<sup>22</sup>, patients with molecular-only relapse has substantially superior survival outcomes compared to those with hematologic relapse (2 year OS 62 v 25%; p=0.003 and 55 v 29%; p<0.0001). Owing to publication bias, retrospective data, varied definitions of molecular relapse (ie. including small proportions of patient with mixed chimerism), and the lack of standardized MRD testing, these observations are challenging to apply directly to practice. They do, however point to a clear trend reported in these studies and elsewhere that those receiving DLI with the smallest amount of disease burden at relapse tend to have better survival outcomes<sup>4, 19, 22-24</sup>. Locally we have given DLI in 2018 and 2019 for molecular relapse in 3 patients with AML, B-ALL, T-ALL. As of September 2023, two are still alive more than 59 (B-ALL) and 60 months (AML) post DLI, with one death (T-ALL) more than two years post DLI (30 months).

To consolidate the above retrospective with local observations that patients with MRD +ve at the 100 day mark have mixed survival outcomes, we will consider DLI for patients meeting the following criteria:

#### Patients with AML or MDS

- who relapse
  - < 1 year post alloHCT
  - > 1 year post alloHCT and non-candidate for second alloHCT
- with relapse defined as

- hematologic relapse: >5% marrow blasts, peripheral blood blasts, extramedullary disease
- molecular relapse that is
  - defined by disease-identifying cytogenetic, morphologic, molecular, or immunophenotypic means (not incomplete donor chimerism)
  - documented on
    - 3 month post-transplant marrow **AND** demonstrated to be worsening at subsequent time point(s) 4-6 weeks later
    - documented at clinical suspicion of relapse beyond 3 months (and if deemed necessary by clinician discretion) demonstrated to be worsening at subsequent time point(s)
- without options for clinical trials or targeted chemo/immunotherapy
- without a history of gr3-4 aGVH or mod-sev cGVH
- who are motivated and fit

Patients with B- or T-ALL could be considered for DLI if meeting the above criteria and options.

Given the alternatively dismal outcomes for relapsed leukemia after allotransplant, and somewhat favourable survival and toxicity outcomes above, in 2017 our center adopted and has used the Azacitidine and DLI schedule described in the above report, which involves 8 total cycles of Azacitidine, with 4 escalating DLI doses, each DLI occurring after 2 Azacitidine cycles. After more than 5 years using this protocol, and more than 25 years using DLI altogether, Calgary DLI recipients have rarely received more than 2 DLI doses (Table 1). Common reasons for not proceeding beyond one dose included disease progression, GVHD, other toxicity, or death.

| Max DLIs Received | No. Pts (%) |
|-------------------|-------------|
| 1                 | 67 (70)     |
| 2                 | 19 (20)     |
| 3                 | 7 (7)       |
| 4                 | 3 (3)       |
| 96                |             |

**Table 1:** Maximum DLI Doses Received by 96 DLI Recipients in Calgary, 1996-2023

- Harvesting, aliquoting, and freezing to target 4 different DLI doses in Calgary has proven restrictively labour intensive. Owing to both total harvest yield and loss of cell viability in the freeze-thaw process, it is also rare that our center achieves target viable cells for infusion at all doses, particularly at the highest concentrations. Given these challenges and the rarity with which our center proceeds with a 4th DLI dose, we will adapt the regimen above to target 6

cycles of Azacitidine and 3 DLI doses, while keeping the dose intensity of the 4th dose increment described in the original Schroeder < 1 year post alloHCT

- > 1 year post alloHCT and non candidate for second alloHCT report.

We will use the following adapted schedule:

- Week 1: Azacitidine 100 mg/m<sup>2</sup>/d x 5
- Week 5: Azacitidine 100 mg/m<sup>2</sup>/d x 5
- Week 6: DLI #1 (1 x 10<sup>6</sup> T cells/kg)
- Week 9: Azacitidine 100 mg/m<sup>2</sup>/d x 5
- Week 13: Azacitidine 100 mg/m<sup>2</sup>/d x 5
- Week 14: DLI #2 (1 x 10<sup>7</sup> T cells/kg)
- Week 17: Azacitidine 100 mg/m<sup>2</sup>/d x 5
- Week 21: Azacitidine 100 mg/m<sup>2</sup>/d x 5
- Week 22: DLI #3 (1 x 10<sup>8</sup> T cells/kg)

This protocol was adopted by the Alberta Blood and Marrow Transplant Program for treatment of relapsed AML and MDS in 2017. As of March 2021 we have treated 13 patients with mixed results. Median follow-up of surviving patients is 720 days, range 102-1202 days. Median overall survival for the entire cohort is 185 days with 43% of patients alive at one year. Three-year overall survival is 16%. As of September 2023 Calgary has performed DLI in 29 patients using this protocol. Of the 22 patients with at least 24 month follow-up, 2 year OS is 41%. Of the 9 patients surviving more than two years post DLI, 7 remain alive and are approaching or well past 4 years since their infusion.

### **Indolent Diseases: Chronic Myelogenous Leukemia (CML) and Chronic Lymphocytic Leukemia (CLL)**

In the era of numerous effective TKIs and accumulating targeted therapies, Calgary rarely performs alloHCT, let alone DLI for CML and CLL: since 2014, Calgary has performed DLI for two relapsed CML cases and none for CLL. The short review below summarizes experience with DLI in these conditions.

In CML and CLL the risk of recurrence is related to the status of the disease at the time of transplantation. Outcomes of transplantation for CML beyond first chronic phase or for CLL that has transformed to aggressive lymphoma remain inferior to those of less advanced disease. Outcomes of transplantation for CLL with adverse cytogenetics (17p-, 11q-), advanced stage at diagnosis or that is fludarabine-refractory are inferior to those of patients without these features<sup>25</sup>.

The management of relapsed CML and CLL after transplant should take into account prior therapies the patient has received and the existence of newer therapies that the patient may not have been exposed to prior to undergoing transplant. In CML, later-generation tyrosine kinase inhibitors (dasatinib, nilotinib, bosutinib and ponatinib) may not have been available to the patient prior to

transplant. Similarly, in CLL, patients may not have received B-cell receptor antagonists (ibrutinib or idelalesib) or a BCL-2 inhibitor (venetoclax). It is reasonable to use these agents in the post-transplant relapse setting if the patient has not previously been exposed.

The existence of an immunological graft-versus-leukemia effect in these diseases is well described. In both diseases relapses are more common using T-cell depleted grafts, relapses are less common once chronic GVHD develops, responses are delayed and tend to deepen over time. Donor lymphocyte infusions (DLI) are a practical way of exploiting this graft-versus-leukemia effect, although they are not without significant toxicities of their own. The majority of patients who respond to DLI develop some degree of acute or chronic GVHD and 8% of patients treated with DLI develop aplasia and may require retransplantation.

The table below summarizes the response of relapsed CML to DLI. While responses are seen in the majority of patients with early relapse, responses in accelerated phase disease are more the exception than the rule. Similar results are observed in CLL, although large series have not been published to date. The existing literature suggests that 44 – 86% of patients with relapsed CLL will respond to DLI, and this response may be enhanced by the addition of rituximab<sup>26, 27</sup>. In the case of CML, the addition of TKI's or interferon may enhance response to DLI, while in CLL chlorambucil or rituximab may be used to delay progression of disease until a graft-versus-leukemia effect occurs<sup>28</sup>. Fludarabine should not be given within 48 hours of DLI as it may abrogate the allogeneic T-cell responses necessary for a graft-versus-leukemia effect to take place.

**Table 1.** Response of Relapsed CML to DLI

|            | Molecular or Cytogenetic Relapse | Chronic Phase | Accelerated Phase | Total       |
|------------|----------------------------------|---------------|-------------------|-------------|
| Van Rhee   | 11/11                            | 8/14          | 1/5               | 20/30 (66%) |
| Collins    | 3/3                              | 25/34         | 5/18              | 33/42 (78%) |
| Drobyski   | —                                | —             | 6/8               | 6/8 (75%)   |
| Porter     | —                                | 6/8           | 0/3               | 6/11 (54%)  |
| Kolb       | 14/17                            | 39/53         | 1/14              | 54/84 (64%) |
| MacKinnon  | 8/8                              | 9/10          | 2/4               | 19/22 (86%) |
| Bacigalupo | —                                | —             | —                 | 10/18 (55%) |
| Alyea      | 15/19                            |               | 0/5               | 15/24 (62%) |
| Verdonck   | —                                | 9/9           | 4/5               | 13/14 (93%) |
| Sehn       | NS                               | NS            | NS                | 19/23 (82%) |

*Response to DLI in relapsed CML by phase at relapse. Adapted from Dazzi et al.<sup>19</sup>*

The literature is surprisingly silent on the topic of repeat transplantation for relapsed CML or CLL. While such transplants have no doubt taken place, they are likely restricted to the small number of patients whose disease fails to respond to DLI and whose performance status, comorbidities and

disease status permits. It remains uncertain what additional benefit is to be derived from retransplantation in the setting of disease that fails to respond to the graft-versus-leukemia effect engendered by DLI.



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# Neutropenic Fever

Presented by: Ahsan Chaudhry

## Summary

- Febrile neutropenia is a medical emergency and should be treated rapidly. The initial evaluation should include blood cultures drawn peripherally and through a central line (if present). Further investigations should be carried out based on foci identified on clinical examination.
- Empiric antibiotics should be administered within one hour of presentation.
- Empiric therapy for **stable patients, without a clinical focus**:
  - Piperacillin/tazobactam 4.5 g IV stat and every 6 hours, OR ceftazidime/cefepime 2 g IV every 8 hours.
  - Gentamicin 7 mg/kg (AIBW for obese) IV q24-36h if beta lactam resistance is suspected, or for probable gastrointestinal (GI) source.
  - Above doses assume normal renal function.
- Additional empiric therapy for **unstable patients**:
  - Gentamicin 7mg/kg (AIBW for obese) IV q24-36h for probable GI source
  - Vancomycin 1 gram (or 25mg/kg) IV loading dose for CVC or Pulmonary source.
  - IV fluids, oxygen, early ICU support.
- If blood cultures positive, adjust coverage based on organism and sensitivity.
- If blood cultures negative but persisting fever or patient clinically unwell, continue antibacterials until ANC  $\geq 0.5/\text{nl}$  for 2 consecutive days.
- If blood culture negative, afebrile, and *clinically well* after 72 hours, discontinue antibacterials.
- Empirical anti-fungal therapy should be considered in patients who have persistent or recurrent fever after 4-7 days of treatment with broad spectrum antibacterials. (See chapter on Fungal prophylaxis).

## Definitions

**Fever:** single core temperature of  $\geq 38.5^{\circ}\text{C}$  (or oral  $> 38.3$ ), or a core temperature of  $\geq 38.3^{\circ}\text{C}$  (or oral  $> 38.0$ ) sustained over a 1 hour period.

**Neutropenia:** an absolute neutrophil count of  $< 0.5/\text{nl}$ , or an ANC that is expected to decrease to  $< 0.5/\text{nl}$  during the next 48 hours.

## Investigations

In addition to a focused history, review of systems and physical examination, all patients with fever in neutropenia should be investigated as follows:

1. Routine blood cultures drawn through central line and peripheral vein.

2. Chest X-ray (posterior-anterior (PA) and lateral views) if clinically indicated.
3. Culture specimens from other sites of suspected infection should be obtained if clinically indicated
4. If fevers persist then repeat blood cultures should be drawn every 48 hours from central line only.
5. If cultures are positive then should repeat cultures until negative.

## Empiric Therapy

Both ASCO (American Society of Clinical Oncology) and Surviving Sepsis campaigns recommend TTA (time to antibiotic) of < 60minutes.

### Stable Patients

1. Piperacillin/tazobactam at 4.5 grams every 6 hours
2. Ceftazidime/Cefepime 2 grams q8h is given to patients who may have allergy to penicillin, recognizing that 5% of patients may still cross react.
3. True **penicillin** anaphylaxis likely requires an Infectious Diseases consult, but consider:
  - Aztreonam 2 grams IV every 6 hours (only gram-negative coverage) + vancomycin 1 gram IV every 12 hours (gram-positive coverage), OR
  - Tobramycin 6mg/kg/day + Levofloxacin 500 mg IV daily, OR
  - *Ciprofloxacin 400 mg IV every 12 hours + clindamycin 600mg IV every 8 hours*

### Unstable Patients

Severe sepsis is a syndrome defined by evidence for SIRS (systemic inflammatory response syndrome) (defined by  $\geq$  two of the following criteria):

- body temperature  $> 38^{\circ}\text{C}$  or  $< 36^{\circ}\text{C}$ ,
- heart rate  $> 90$  beats/minute,
- respiratory rate  $> 20$ /minute,
- $\text{Pa CO}_2 < 32$  mmHg,
- an alteration in the total leukocyte count to  $> 12 \times 10^9/\text{L}$  or  $< 4 \times 10^9/\text{L}$ , or the presence of  $>10\%$  band neutrophils in the leukocyte differential
- PLUS evidence of infection and end-organ dysfunction (altered mental status, hypotension (systolic blood pressure  $< 90$  mmHg, mean arterial pressure  $< 70$  mmHg, or systolic blood pressure decrease of  $> 40$  mmHg,) elevated serum lactate  $>4$  mmol/L, oliguria (urine output  $<0.5$  mL/kg/hour), and/or hypoxia.

Patients with sepsis or pneumonia with bacteremia have mortality  $>50\%$  despite prompt antibiotics. Aggressive fluid resuscitation, oxygen and early physiological goal directed therapy, including ICU support, is critical.

1. Meropenem 500 mg q6h, Gentamicin 7 mg /kg/day (AIBW) given every 24-36 hours, or Ciprofloxacin (750mg po bid/400mg IV q12) may be initiated if antimicrobial resistance or GI source is suspected.
2. Vancomycin may be added empirically for SIRS, hospital acquired pneumonia (HAP), gram-positive bacteremia, endocarditis, meningitis, or osteomyelitis. Vancomycin loading dose (25-30mg/kg ABW) should be considered if practical for HAP or SIRS (although TTA may be more important). Maintenance dosing (15mg/kg ABW) is then continued q12h until discontinuation or dose adjustment per trough level.
  - If vancomycin is used only empirically, it should generally be discontinued after 24-48 h.
  - Vancomycin trough levels should be considered if plasma creatinine >40 mmol/L above baseline, BMI >40, age >60, or treatment duration >7d. Trough target level is 15-20 mg/L for HAP/MRSA and 10-15 mg/L for empiric therapy. First trough level should be determined at steady state (pre 4<sup>th</sup> or 5<sup>th</sup> dose) and repeated after adjustment in new steady state, every 7-10 d or if concurrent nephrotoxic drugs.
  - Vancomycin may be added **temporarily** in the case of blood cultures showing gram-positive organisms, although in this case one set of blood cultures each should be collected peripherally and centrally to confirm persistent bacteremia and exclude a false-positive (i.e. contaminated) blood culture.
  - There is no proven advantage to adding vancomycin empirically in the setting of persistent or recrudescent fever and neutropenia in an otherwise asymptomatic hemodynamically stable patient. If vancomycin was added empirically at the outset of therapy for neutropenic fever, it should be stopped if blood cultures have incubated for 48 hours and demonstrated no pathogenic gram-positive organisms.

## Re-Assessment

Patients are reassessed for response to treatment daily. Antibacterial coverage is adjusted to ensure coverage of organisms grown in culture, preferably based on *in vitro* sensitivity testing.

## De-Escalation

Two randomized controlled trials compared de-escalation strategies in hematologic malignancy patients undergoing chemotherapy or HCT with FN without microbiologic diagnosis (Table 3). The How Long study compared empiric antibiotic therapy (EAT)-free days among 157 neutropenic patients randomized to de-escalation using ECIL-4 criteria versus standard of care based on hospital-specific protocols. EAT-free days was significantly lower in the experimental group, with an absolute difference of 6.4 days. There was no difference in crude mortality (0 in both groups), and mean days of fever did not differ significantly between groups. Another randomized trial compared short (minimum 72 hours) versus extended carbapenem in 281 patients with fever of unknown origin with median 14 days of neutropenia. Carbapenem treatment duration significantly differed: 3 versus 8 days. Primary composite outcome was treatment failure (carbapenem-sensitive infection, fever recurrence,

septic shock, respiratory insufficiency, or death) from Day 4 to neutropenia recovery, and was nonsignificantly higher in the short duration group (19% vs. 15%), which met criteria for non-inferiority.

There have also been several observational studies that compare clinical outcomes in patients undergoing early de-escalation of empiric broad-spectrum antibiotics versus continuation through ANC recovery, with the six adopting an ECIL-4-based de-escalation strategy.

**Table 1.** Reassessment criteria for patients

| Persistent fever after 3 to 5 days of treatment:                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          | Afebrile after initial antimicrobial treatment with no etiology identified:                                                                                                                                                                                                                                                                                                                                                                                                                       | Positive blood cultures/focus:                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <ol style="list-style-type: none"> <li>1. Repeat blood cultures and other investigations as indicated above.</li> <li>2. Imaging on day 5 – enhanced CT of chest+abdomen+pelvis if good renal function. If poor renal function, then nonenhanced CT of chest plus US of abdomen.</li> <li>3. Empirical antifungal treatment as indicated (see chapter on Fungal prophylaxis).</li> <li>4. Add vancomycin for 48hrs if criteria are met, e.g. skin and soft tissue infection, catheter related infection, pneumonia or hemodynamic instability.</li> </ol> | <ol style="list-style-type: none"> <li>1. High risk patients should continue antibiotics until ANC greater than 500 cells/mm<sup>3</sup> for 2 consecutive days.</li> <li>2. Antimicrobials are stopped for ATG (antithymocyte globulin) related fevers if afebrile and blood culture is negative after 48 hours.</li> <li>3. Discontinuation or simplification of anti pseudomonal antibiotics possible after 72 h or more of apyrexia plus clinical recovery while still neutropenic</li> </ol> | <ol style="list-style-type: none"> <li>1. Treat according to sensitivities if available.</li> <li>2. For blood culture positive for gram positive microorganism, repeat another set of blood culture centrally and peripherally before starting Vancomycin to rule out possibility of contamination.</li> <li>3. For documented infection with positive culture, the duration of antimicrobial therapy depends on the type, site and source of infection.</li> <li>4. Consider central line source if &gt; 2hr difference in TTP (time to positivity).</li> <li>5. Investigate focus appropriately and treat according to common pathogens.</li> </ol> |



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# Central Venous Access Device (CVAD) and its Complications

Presented by: Ahsan Chaudhry and Robin Pommier

## Summary

- **Line Type Preferences**
  - **Recipients (Autologous transplant, Allogeneic transplant, Chimeric antigen receptor T-Cells (CAR T-cells):**
    - The recommended catheter for recipients is the *Trifusion™ 12F triple lumen carbothane apheresis catheter*
    - If a tunneled line or apheresis is not necessary, a silicone double lumen Groshong peripherally inserted central catheter (PICC) can be considered.
      - CAR T-cells that require gravity infusion (e.g. tisagenlecleucel) cannot be infused through a valved PICC, such as a Groshong.
  - **Donors:**
    - Peripheral venous access is preferred for collection from all donors. Two large-bore antecubital lines will be inserted just prior to apheresis.
    - If large bore antecubital lines cannot be inserted, a *double-lumen temporary Quinton Mahurkar* (8 French diameter, length 15 cm) will be inserted under image guidance and removed prior to the patient leaving the hospital.
- **Prevention of CVAD Infections**
  - The central venous catheter care clinical bundle (including hand hygiene, maximal barrier precautions, and chlorhexidine skin antisepsis) will be used for placement and maintenance of all CVADs.
    - Rigorous attention to hand hygiene and aseptic technique is essential before inserting, removing, or manipulating the CVAD.
    - Prepare clean skin with a >0.5% chlorhexidine preparation with alcohol before CVAD insertion and during dressing changes.
    - Use sterile gauze or sterile, transparent, semi permeable dressing on CVAD insertion site. For tunneled CVADs, dressings may be removed as per unit policy and procedure.
  - Promptly remove CVAD lines that are no longer being used.
  - Insertion of CVAD on right side is preferred. Avoid femoral vein.
  - Remove CVAD if not used or used infrequently.
- **Treatment of CVAD Infections**
  - **Empiric Treatment:**
    - Collect bacterial cultures from CVAD entrance/exit site and blood (central and peripheral) prior to initiating treatment.
    - Vancomycin to cover gram positive organisms.

- In systemically ill patients, add Tazocin, ceftazidime or meropenem to cover gram negatives.
- **Treatment of Proven or Complicated Infection:**
  - Treat according to IDSA guidelines as described in main text below.
- **Prevention of Line Occlusion**
  - Diligent care and maintenance as outlined in nursing vascular access device infusion therapy Clinical Care Topic (CCT)
  - CVADs are typically flushed/locked with 0.9% sodium chloride.
    - **Exception:** CVADs inserted for the purpose of apheresis are locked with 4% sodium citrate for the lifetime of the catheter.
- **Treatment of Line Occlusion (Thrombotic or Mechanical)**
  - Thrombotic occlusion will be treated with r-tPA.
  - Unless mechanical occlusion is suspected radiographic imaging is not necessary prior to r-tPA instillation.
  - If a mechanical issue is suspected or signs of tip malposition, an x-ray and/or dye study will be carried out.
  - For chemical occlusion consult Advanced Venous Access Service (AVAS) for assessment.
- **Treatment of Line Related Venous Thrombosis**
  - There is insufficient evidence to recommend routine removal of clinically necessary, functioning and non-infected CVAD's in the setting of catheter-related thrombosis. If anticoagulation is not feasible then line removal is indicated.

## Background

Multi-lumen catheters are placed prior to transplant to facilitate transfusions, blood draws and medication administration and are preferably tunneled to decrease infection risk.

### Line Type Preferences

#### **Autologous Transplant and CAR T-cell Recipients:**

- For autologous transplant and CAR T-cell recipients, a rigid line is needed for apheresis. The current recommended catheter used prior to apheresis is the *Trifusion™ 12F triple lumen carbothane apheresis catheter* and is to remain in place until after autologous transplant.
- CVAD lines inserted for the purpose of apheresis are locked with 4% sodium citrate for the life time of the catheter.

- If a patient has had a previously installed implanted vascular access device (IVAD), it need not be removed prior to transplant but another catheter may also be placed.
- If a peripherally inserted central catheter (PICC) will be used for transplant instead of a tunneled CVAD a Bard Groshong silicone PICC line is preferred. Other valved PICCs have been found to hemolyze blood samples.
  - CAR T-cells that require gravity infusion (e.g. tisagenlecleucel) cannot be infused through a valved PICC, such as a groshong.

### **Allogeneic Transplant:**

- In allogeneic transplantation, a large bore, triple lumen catheter is preferred for transfusions and medication administration. The current CVAD recommendation is the same line used for autologous transplant.
- Since the CVAD is not inserted for the purpose of apheresis it is standardly flush/locked with 0.9% sodium citrate.
- If a PICC line needs to be inserted pre transplant a Bard Groshong silicone line is preferred.

### **Donors:**

- Two large bore antecubital lines are to be inserted.
- If large bore antecubital line insertion is not possible or donor is unwilling a double lumen temporary Quinton Mahurkars (8 French diameter), length 15 cm, is inserted the day of collection to facilitate apheresis and then removed post apheresis.
- A patient or donor with a temporary non-tunneled CVAD (e.g. Quinton Mahurkars) must remain in hospital until the line is removed.

## **Complications Associated with Central Venous Catheters**

### **Bleeding Following Insertion**

- The bleeding risk associated with insertion of a tunneled central line is variable and depends on coagulative function as well as operator experience and skill.
- To minimize bleeding risk for line insertion, ensure platelets >50 and INR <1.4 prior to line insertion, or as specified by radiologist.
- Avoid high dose heparin.
- Bleeding can be managed with local pressure to site, hemostatic dressing, reversal of anticoagulation (i.e. heparin from line, PT and PTT must be checked), clotting factors if necessary, tranexamic acid.
- Rarely, surgical intervention may be required to repair site.

*Abbreviations: INR = international normalized ratio; PT = prothrombin time; PTT = partial thromboplastin time.*

## Catheter-Related Infections

Catheter-related infections are important causes of morbidity, mortality and health care costs, with an infection rate of approximately 5 per 1000 catheter days in the critical care population. In a meta-analysis of 2573 catheter-related blood infections, case-mortality rate was 14% with 19% of deaths due to catheter-related infection.<sup>1</sup> Mortality was the highest with *Staph. aureus* at 8.2% and lowest with coagulase negative *Staph.* at 0.7%<sup>1</sup>.

Skin organisms predominate in the first few weeks as they migrate into the catheter tract and cause tip infections. In long term catheters, hub infections become a more common source. Line infections can also result from hematogenous seeding from other sites.

- Peripheral IV – 0.5/1000 catheter days
- Cuffed Tunneled CVAD – 1.6/1000
- PICC – 2.1/1000
- Temporary non cuffed CVAD – 2.7/1000

Catheters made of Teflon, silicone elastomer, or polyurethane are less likely to cause infection than catheters of polyvinyl chloride or polyethylene<sup>2,3</sup>. Surface irregularities enhance the microbial adherence of some organisms (i.e. coagulase negative *Staph.*, *Acinetobacter calcoaceticus*, *Pseudomonas aeruginosa*). Some catheters are also more thrombogenic, which can contribute to subsequent infections. Host factors can be important; for example *Staph. aureus* adheres to proteins such as fibronectin that are commonly present on catheters and this can make infection difficult to clear. In addition, coagulase negative *Staph.* adheres well to polymer surfaces and can produce an extra cellular polymer “slime” which allows it to withstand host defenses by killing neutrophils and acting as a barrier to antibiotics and phagocytes. *Candida* can also produce slime in presence of glucose-containing fluids, which may contribute to increased fungal infections in people on total parenteral nutrition. The most common organisms cultured from patients with central line infections are as follows<sup>4</sup>:

- Coagulase negative *Staphylococcus* (50%)
- Gram negative organisms (20%)
  - Increasing third generation cephalosporin resistance in *E.coli* and *Klebsiella*, increasing imipenem and ceftazidime resistance among *pseudomonas aeruginosa*
- *Staphylococcus aureus* (30%)
  - Rare MRSA

## History Suggesting Catheter-Related Infection:

Components of the patient history supporting the presence of a catheter-related infection include continuous or persistent bacteremia, sepsis after infusing through a line, blood cultures of organisms known to colonize/infect lines, catheter thrombosis, clinical improvement with catheter removal, and the lack of another clinical source of infection. Physical exam findings are unreliable but can include fever, shock or inflammation/purulence at the exit, entrance or tunnel site.

## Diagnostic Tests:

If a catheter-related infection is suspected, the following tests should be ordered:

- Gram stain and culture of exudate if present
- Culture of line tip if removed (best if plated at bedside)<sup>5</sup>
  - Positive result when >15 colony-forming units present on tip
- Central and peripheral blood cultures drawn prior to antibiotics (min 10 mL/bottle, yield increases 3% per additional mL blood up to 20 mL)
  - A difference in the time to positivity of ≥120 minutes between centrally- and peripherally-drawn blood cultures is 91% sensitive, and 94% specific for catheter infection<sup>5</sup>
  - Negative predictive value for central line infection when negative culture drawn from central line prior to antibiotics: 99%<sup>6</sup>
  - Cultures of *Staph. aureus*, coagulase negative *Staph.* and *Candida* are most suggestive of central line-related infection

## Prevention of CVAD Infections (Adapted from IDSA Guidelines)<sup>2</sup>

- Rigorous attention to hand hygiene and aseptic technique is essential before inserting, removing, or manipulating the CVAD.
- Prepare clean skin with a >0.5% chlorhexidine preparation with alcohol before CVAD insertion and during dressing changes.
- Evaluate the catheter site daily by palpation through the dressing for tenderness and by inspection if transparent dressing; if opaque dressing this does not need to be removed.
- Consider removal of CVAD if intraluminal catheter thrombosis cannot be corrected
- Promptly remove CVAD lines that are no longer being used, non-functional or bulging.
- Diligent care and maintenance as outlined in nursing vascular access device infusion therapy Clinical Care Topic (CCT)

## Treatment of CVAD Infections

Definite indications for tunneled catheter removal are as follows<sup>7</sup>:

- Complicated infections (septic thrombosis, endocarditis, osteomyelitis, possible metastatic seeding).
- Tunneled catheter pocket infections or port abscess.
- Persistently positive cultures or persistent fever (>72 hours) while on treatment for a known line infection
- Relapse after antibiotics are discontinued.
- Fungal catheter-related blood infection, candida, mycobacteria, *Pseudomonas aeruginosa*, *Staph. Aureus*.



There should be a low threshold for catheter removal with catheter related blood stream infections including *Burkholderia cepacia*, *Actinobacter baumannii*, *Stenotrophomonas* species, *Bacillus* species, and *Corynebacterium* species. For coagulase negative *Staph.* bacteremia, recurrence by 12 weeks was seen in 20% of patients with line salvage versus 3% with line removal; another study found *Staph. aureus* patients were 6.5 times more likely to relapse or die of infection without line removal (studies were done without antibiotic lock therapy)<sup>7,8</sup>. Reinsertion of central lines should be postponed until after serial negative blood cultures are obtained; although not always practical, this is ideally done after negative blood cultures are obtained 5-10 days after completion of antibiotics.

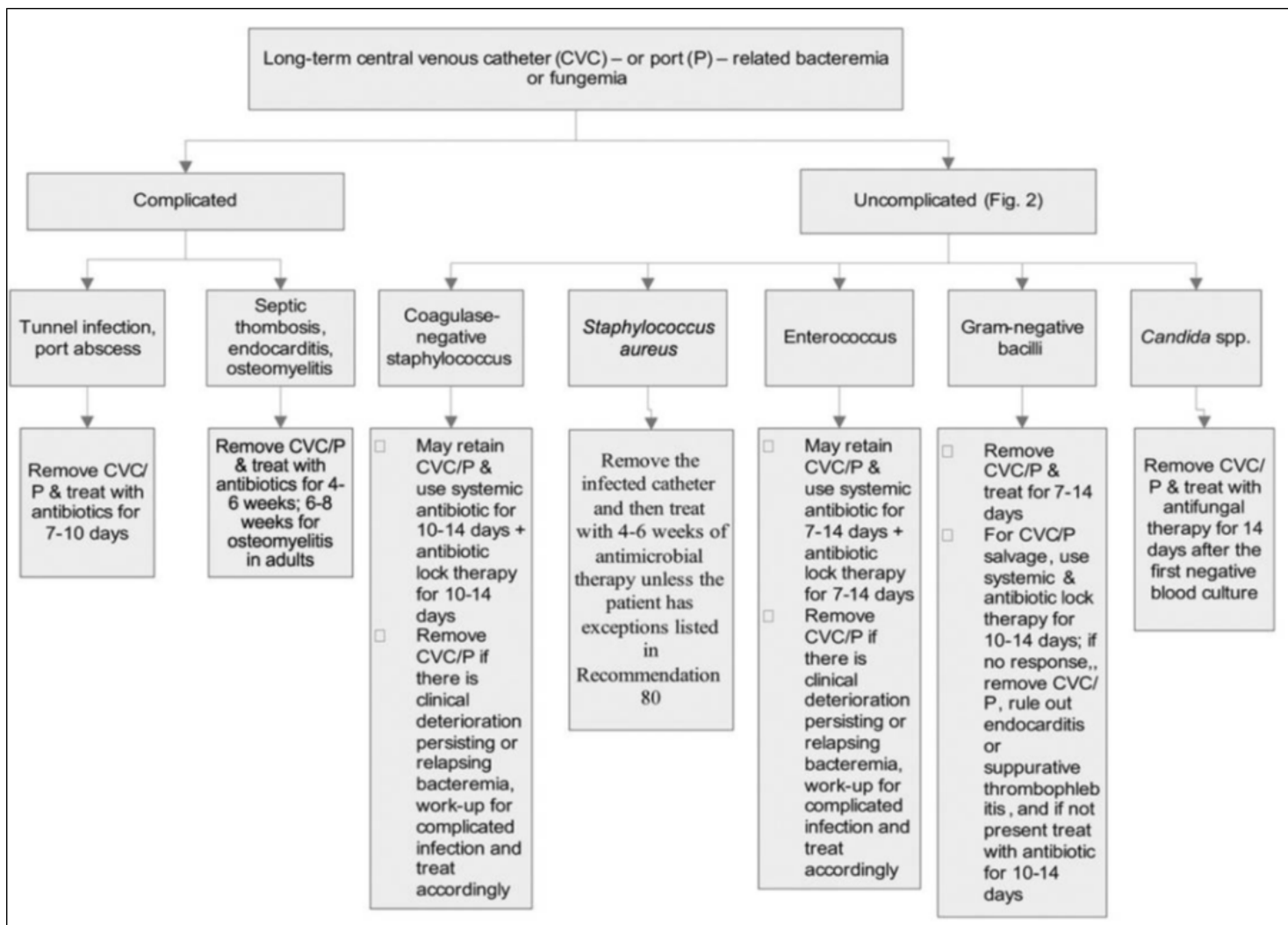
There are limited prospective randomized controlled trials examining the optimal treatment choices and duration of therapy for CVAD infections. Based largely on published guidelines, the following empiric therapy is suggested<sup>7</sup>:

- Vancomycin in hospitals/areas with MRSA; if resistance to vancomycin is seen, daptomycin is the alternative and linezolid is not indicated as empiric therapy for catheter-related bloodstream infection (CRBSI)
  - Covers *Staph. aureus*, coagulase negative *Staph.* and *Enterococci*
- Gram negative bacilli coverage (including *Pseudomonas*) in neutropenic/markedly immunocompromised or severely ill patients
  - Tazocin
  - Alternatives could include meropenem or Third or fourth generation antipseudomonal cephalosporin (i.e., cefepime, ceftazidime)
- Empiric fungal coverage in high risk patients/suspected fungal disease, patients on TPN or with prolonged use of antibiotics, known candida colonization
- Step down antibiotics once organisms/ sensitivities are known
- Avoid use of topical antibiotic ointment or cream at insertion sites

The optimal duration of therapy remains controversial. General guidelines include the following<sup>7</sup>:

- If prompt antibiotic response, treat 10-14 days for pathogens other than coagulase negative *Staph.* (7 days plus antibiotic lock therapy or 10-14 days) if no valvular heart disease or intravascular prosthetic device
- 4-6 weeks antibiotics should be considered if persistent bacteremia or fungemia after catheter removal (>72 hours post catheter removal), endocarditis, septic thrombosis
- 6-8 weeks of therapy for the treatment of osteomyelitis
- For complicated infections, consultation with Infectious Diseases is suggested

**Figure 1.** Approach to the treatment of a patient with long-term central venous catheter (CVAD) or a port (P)-related bloodstream infection<sup>7</sup>.



## Antibiotic Lock Therapy

Antibiotic lock therapy, with pharmacologic doses of antibiotics instilled into the lumen of a line daily for hours, could be considered in uncomplicated tunneled CVAD infections (i.e., no tunnel infection or abscess) with *Staph. aureus*, coagulase negative *Staph.*, and gram negative bacilli. This method is not effective in fungemia, and responses with coagulase negative *Staph.* have been better than with *Staph. aureus* and *Pseudomonas*. When data from four trials were pooled, antibiotic lock therapy plus IV antibiotics were associated with clearance of an organism in 138/167 (82%) of catheter infections compared to pooled data from 14 trials showing clearance of 342/514 (66.5%) with IV antibiotics alone (response rate (RR) of catheter salvage 1.24)<sup>9</sup>.

Two weeks of antibiotic lock therapy can be considered in CVAD infections with coagulase negative *Staph.* and gram negative bacilli and in uncommon situations with *Staph. aureus* where line removal is not feasible.

- Ethanol locks have also been associated with decreased primary catheter related bloodstream infections. WARNING: Alcohol should not be used to soak or declog polyurethane catheters because alcohol is known to degrade polyurethane catheters over time with repeated and prolonged exposure. (e.g. Trifusion™ catheter).

## Specific Management Challenges

### Staphylococcus aureus:

- *Staph. aureus* bacteremia is associated with a high risk of metastatic infections and provides a management challenge (25% - 32% occult endocarditis in patient with staph aureus bacteremia), hematogenous complications in 25-30%.
- Beta-lactam drugs (cloxacillin or cefazolin) are preferred therapy if the *Staph. aureus* is sensitive.
- If the bacteremia is not cleared by 72 hours after antibiotics, long-term therapy is required (minimum 4 weeks).<sup>5</sup>
- Non-tunneled catheters should be removed.
- Tunneled catheters should be removed and must be removed in the presence of abscess or tunnel site infection.
- Search for metastatic infection is indicated, starting with a TTE (transthoracic echocardiography) if there are no contraindications, and clinical monitoring for osteomyelitis, septic arthritis, and other sites of infection.
- ID consultation will likely be needed.

### Enterococcus:

- Ampicillin is treatment of choice +/- gentamicin; vancomycin in cases of ampicillin resistance.
- Daptomycin in cases of VRE (vancomycin-resistant *Enterococcus*) based on susceptibility.
- Line removal is preferred. Lines should be removed in the case of vancomycin resistant species.

### Fungal infections:

- If there is documented catheter-related fungal infection, the CVAD should be removed.<sup>7,10</sup>
- Antifungal therapy should continue until 14 days after last positive blood cultures and signs/symptoms resolved.

### Septic thrombophlebitis:

- The most common organisms implicated in septic thrombophlebitis are *Staph. aureus*, *Candida* species and gram negative bacilli; the presence of thrombus greatly increases the risk of CVAD-related infections.
- In the presence of septic thrombophlebitis, the catheter should be removed.

- Surgical consultation is indicated in the case of suppurative thrombophlebitis, infection persists on antibiotics or there is pseudo aneurysm formation.
- Routine anticoagulation of patients with septic thrombophlebitis is not recommended. It can be considered for selected patients, such as those who are highly symptomatic of their thrombosis.
- Thrombolysis is not indicated. Infectious disease consultation is suggested.

## Catheter-Related Thrombosis or Mechanical Occlusion

### Line Occlusion

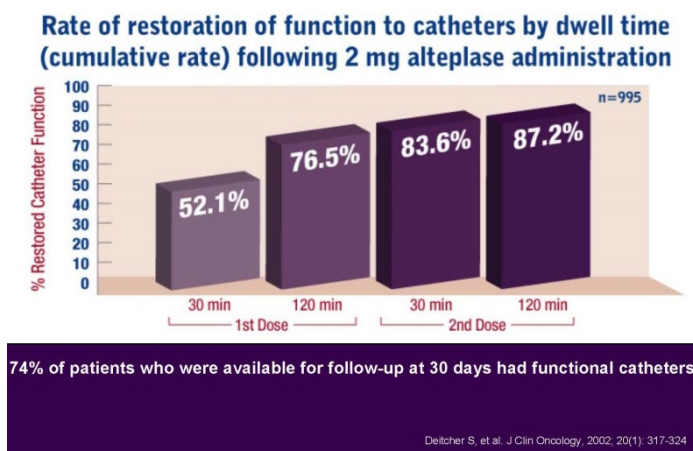
#### Prevention of Line Occlusion

- Diligent care and maintenance as outlined in nursing vascular access device infusion therapy Clinical Care Topic (CCT)
- CVADs are typically flushed/locked with 0.9% sodium chloride.
  - **Exception:** CVADs inserted for the purpose of apheresis are locked with 4% sodium citrate for the lifetime of the catheter.
- 0.9% sodium chloride flush/locks are standardly used over other locking agents as there is insufficient evidence to recommend one lock solution over the other <sup>16</sup>.
- CVADs inserted with the purpose of apheresis are locked with 4% sodium citrate instead of heparin due to the higher risk of HIT<sup>17</sup>

#### Thrombotic occlusions:

- Occluded CVADs should be treated with r-tPA.
- Unless mechanical occlusion is suspected radiographic imaging is not necessary prior to tPA instillation.
- 2 mg alteplase (Cathflo) is reconstituted with 2.2mL sterile water by a certified RN. As much as possible up to 2mg is instilled into the blocked CVAD lumen and as per nursing procedure.
- Place r-tPA into lumen for 2-24 hours then aspirate. R-tPA can be aspirated after 30 minutes if line access is urgent.
- Can be repeated x1 if unsuccessful; tPA can be left in situ overnight.

**Figure 2.** Rate of restoration of function to catheters by dwell time (cumulative rate) following 2 mg alteplase administration. Note: subjects with occluded, no dialysis CVADs were enrolled, not specifically neutropenic patients.



### Mechanical occlusions:

- If line patency is not restored, consider consulting interventional radiology (line stripping, TPA drip in IR). If this is unsuccessful the line is to be removed as soon as safe to do so.

### Catheter-Related Venous Thrombosis

The incidence of symptomatic catheter-related deep vein thrombosis (DVT) in patients with malignancies is approximately 3-4%, although ultrasound surveillance documents clots in about 12% of patients<sup>11</sup>. A small series in bone marrow transplant patients showed an incidence as high as 50% although the majority were asymptomatic<sup>12</sup>. Risk factors include malplacement of the catheter, >1 insertion attempt, a previous CVAD, placement of the catheter on the left-hand side and malignancy.

Symptoms that suggest an upper extremity DVT include erythema and swelling (which may be exercise-dependent or gravity-dependent), and pain or tenderness at the base of the neck, superclavicular fossa, arm or shoulder. Collateral blood flow often develops and vessels may be visible. Embolization is the major cause of morbidity and mortality, and pulmonary embolism (PE) occurs in up to 20% of patients with symptomatic thrombosis. The following tests may confirm the diagnosis:

- Ultrasound or venogram of extremity
- If symptoms of respiratory compromise/pulmonary embolism, workup requires a PE protocol CT scan or V:Q scan; rarely pulmonary angiogram is indicated

### Prophylaxis of CVAD-related Thrombosis and Deep Venous Thrombosis

- DVT prophylaxis should be carried out as per established guidelines for the medical patient in the absence of significant bleeding, coagulopathy or thrombocytopenia (platelets < 50)<sup>12</sup>. Options for

thromboprophylaxis include low-dose unfractionated heparin, low molecular weight heparin or mechanical prophylaxis.

- Mobilization should be encouraged
- Use of anticoagulation for routine prophylaxis of catheter-related thrombosis is not recommended<sup>13-15</sup>.

### **Treatment of CVAD-related Thrombosis and Deep Venous Thrombosis**

- Anticoagulation should be continued for the duration of line placement if removal is not feasible.
- Anticoagulation duration is controversial and catheter-related thrombosis should be treated as per established guidelines for provoked DVT<sup>12,15,19,20</sup>.
  - Catheter-related thrombosis should be treated as a provoked thrombosis and treated with anticoagulation for a total of 3 months.
  - Patients whose lines have been removed and who experience bleeding complications while on anticoagulation may be taken off of anticoagulation before completing 3 months of treatment provided symptoms of catheter-related thrombosis have resolved. They should be reimaged in 10-14 days to exclude propagation of venous thrombus if anticoagulation is discontinued early.
  - Patients with active malignancy should receive anticoagulation.
  - Tinzaparin 175 IU/kg once daily or DOAC may provide easier and more reliable anticoagulation compared with warfarin in patients taking multiple interacting medications, antibiotics and/or with unpredictable dietary intake. Caution should be exercised when using low molecular weight heparins in individuals with impaired renal function.
  - Awareness of drug interactions with DOAC's.

## **Catheter Care**

- Patients shall be educated about their own catheter care in preparation for outpatient therapy.
- Written instructions for catheter care shall be given to patients prior to discharge (e.g. Adult Cellular Therapy: Important Information When Leaving the Hospital)
  - Include CVAD Emergency Kit (gauze, large transparent dressing (Tegaderm), plastic clamp, needleless connector, alcohol swabs, chlorhexidine sticks)

## **Catheter Removal**

- With all central line removals, informed consent shall be obtained, and sterile technique maintained.
- Central line removals should be done in the supine position during exhalation to minimize air embolus risk.
- All patients shall have their central lines removed once they are no longer using it regularly.
- All patients shall have central lines removed if they are eating/drinking well and not requiring transfusions or IV medications.



- A new line should be inserted if it is needed (i.e. second transplant).
- Prior to line removal, platelets should ideally be  $>50$  and INR  $<1.4$ . Send catheter for review if mechanical issue/infection potential suspected during line removal.
- If accidental CVAD removal (falls out, pulled out) pressure and dressing shall be immediately applied, ongoing assessment of patient and vital signs for signs and symptoms of air embolism or other complications. Inspect catheter to ensure it is intact.

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#### **Additional Resources**

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# Hepatic Complications and Viral Hepatitis

Presented by: Mona Shafey

## Summary

### Viral Hepatitis

- All recipients and donors will be screened for hepatitis B and C, with further viral load/nucleic acid testing (NAT) required for those with a positive screening test.
- Hepatology referral for assessment and peri-transplant management is required for patients with chronic active hepatitis B or positive hepatitis C serology, and donors who are HBV NAT positive.
- Long-term risks of developing cirrhosis and HCC appear to be similar to non-HSCT population with HBV.
- HCV infection in HSCT population is associated with increased risk of morbidity (e.g. early cirrhosis, GVHD, VOD/SOS) and mortality (e.g. fatal VOD/SOS, excess bacterial infection, fibrosing hepatitis) compared to non-HSCT population.
- Antiviral prophylaxis should be given to all HBV-DNA or HBsAg-positive patients, and for patients who are HBsAg negative but anti-Hbc-positive and undergoing SCT, CAR T-cell therapy, or post-transplant depleting therapies. HCV can be treated concomitantly with chemotherapy if treatment of the malignancy is urgent, and though the presence of HCV-RNA positive in the recipient does not constitute a contraindication to HCT, consider postponing the HCT to allow completion of a treatment course
- Use of mycophenolate mofetil (MMF) has been linked to developing fibrosing cholestatic hepatitis and should not be used in HCV-infected patients.

### Veno-Occlusive Disease/Sinusoidal Obstruction Syndrome (VOD/SOS)

- Patients with risk factors for VOD/SOS development should have a fibroscan as part of pre-transplant investigations to assess for baseline liver fibrosis/dysfunction.
- Ursodiol 15-20 mg/kg/day will be given for the first 80 days post-transplant for prophylaxis of VOD/SOS in allogeneic HCT recipients, regardless of conditioning used for transplant
- Serial doppler ultrasounds post transplant can be considered in patients with risk factors for VOD/SOS or for those with established mild VOD/SOS to monitor for progression
- VOD/SOS should be suspected clinically in the patient with weight gain, hyperbilirubinemia, hepatomegaly, +/- ascites early post transplant. The diagnosis should be confirmed by ultrasound, liver biopsy or measurement of hepatic vein wedge pressure gradient if possible.
- Standard treatment of VOD/SOS is supportive, with careful attention to fluid balance and renal perfusion and elimination of hepatotoxic medications. Defibrotide 25 mg/kg/day should be considered for patients with severe or very severe VOD/SOS, and considered pre-emptively in patients with moderate VOD/SOS

## Hepatotropic Viruses

In patients with hematologic malignancies, including stem cell transplant and cell therapy recipients, changes in the immune status can profoundly alter the course of viral hepatitis, leading to reactivation or hepatic and extrahepatic complications. Additionally, allogeneic stem cell donors with past or current viral hepatitis infections may transmit the virus to recipients. The summary below incorporates the updated recommendations from the 9<sup>th</sup> European Conference on Infections in Leukemia (ECIL-9) for the management of hepatitis virus infections in patients undergoing hematopoietic cell transplantation (Mikulska 2025).

### Hepatitis B Virus (HBV)

HBV reactivation is a potentially life-threatening complication of chemotherapy or immunosuppressive therapies, including depleting agents (e.g. anti-CD20 and anti-CD38 antibodies), cancer chemotherapy, checkpoint inhibitors, bone marrow and stem cell therapies, immunosuppressive therapies for transplant recipients, and CAR T-cell recipients. HBV is a frequent infection worldwide, with approximately 254 million people (3.2% of the global population) living with the virus reported in 2022 (WHO 2024). Consequently, HBV infection in HCT candidates is a frequent situation. An infection due to HBV does not preclude an HCT or CAR T-cell therapy, but the liver disease caused by this virus can.

Reactivation is defined by one of the following: detection of HBsAg in individuals previously HBsAg-negative, detection of HBV-DNA in patients previously negative for HBV-DNA, or a ten-fold increase in the HBV-DNA levels in patients with previously detectable HBV-DNA. The most important practical issue with HBV in HCT is the prevention of severe or fatal hepatitis due to HBV (Siyahian 2018).

In HCT candidates with chronic hepatitis based on biopsy or positive HBsAg or high levels of HBV-DNA, the transplant procedure should be delayed *when possible*, and antiviral therapy given for 3-6 months before conditioning. Antiviral prophylaxis should be given to all HBV-DNA or HBsAg-positive patients, and for patients who are HBsAg negative but anti-HBc-positive and undergoing SCT, CAR T-cell therapy, or post-transplant depleting therapies. Lamivudine is no longer recommended. Tenofovir or entecavir are the drugs of choice, starting before the conditioning regimen and maintained for at least 1 year after HCT, and longer if GVHD is present or immunosuppression. After stopping antiviral prophylaxis, HBV monitoring every 3 months is recommended as late reactivation with fulminant hepatitis has been described (Lee 2019). **Treatment of patients with acute or chronic HBV hepatitis should be done in collaboration with hepatology.**

Studies have reported HBV-DNA positivity in blood donors who are HBsAg-negative and anti-HBc positive, ranging from 3.3% in regions of low HBV endemicity to up to 38% in high endemic regions (Silva 2005, Yotsuyanagi 2001). A HBsAg-positive donor transmits HBV to the recipient in 22-44% of cases. Antiviral therapy of the donor and prophylaxis for the recipient is recommended if such a donor

is required. Sustained clearance has been reported in HBsAg+ recipients receiving transplants from donors with natural immunity to HBV (Hui 2005, Lau 1997). Vaccination of the donor should be considered if this is in the best interest of the donor.

Vaccination of anti-HBc-negative and anti-HbSAg negative patients after HCT is recommended.

**Table 1.** Hepatitis B Serology and BMT (adapted from Strasser *et al.* Blood 1999)<sup>9</sup>

| Patient Result | Donor Result   | Interpretation                                                                                                               |
|----------------|----------------|------------------------------------------------------------------------------------------------------------------------------|
| Anti-HBs       | Anti-HBs       | Exposed or vaccinated                                                                                                        |
| Anti-HBc       |                | Exposed. Risk of reactivation present.                                                                                       |
|                | Anti-HBc       | Exposed. Negligible risk of transmission if HBV DNA negative                                                                 |
| HBsAg positive |                | Active infection: Liver biopsy and start treatment if HBV DNA+                                                               |
|                | HBsAg positive | Assess donor for liver disease. Consider different donor or antiviral treatment pre-harvest. Monitor recipient HBV DNA post. |

## Hepatitis C Virus (HCV)

The hepatitis C virus is a single-stranded RNA virus The hepatitis C virus is a blood borne virus, with transmission of HCV by blood transfusion being an exceptional event in developed countries. The incubation period is 6 – 12 weeks, followed by a generally mild, self-limiting hepatitis. 85% develop chronic infection, and of these 20% will develop cirrhosis and 5% will die of liver failure or hepatocellular carcinoma (HCC). In 2022, an estimated 50 million individuals worldwide (0.6% of the global population) were living with chronic HCV (WHO 2024). In HCT, the most important aspect is the detection of HCV infection in order to implement treatment to avoid liver damage.

The treatment of HCV with direct-acting antivirals (DAAs) can obtain cure in over 90% of patients. Thus antiviral treatment should be considered for all HCV-RNA positive patients with hematologic cancers. HCV can be treated concomitantly with chemotherapy if treatment of the malignancy is urgent, and though the presence of HCV-RNA positive in the recipient does not constitute a contraindication to HCT, consider postponing the HCT to allow completion of a treatment course. As before, **consultation with an expert hepatologist is required for these patients.**

The course of hepatitis C after BMT appears to be generally mild. While some reports suggest an increased rate of severe veno-occlusive disease (VOD) in BMT recipients who are positive for the hepatitis C virus (HCV), most suggest that the risk is not substantially higher than in HCV-negative recipients. *Strasser et al.* reported the results of BMT in patients who were HCV-positive at the time of transplant, and they found the risk of severe VOD in HCV-positive patients was 48%, compared with 14% in HCV-negative control patients (Strasser, Hepatology 1999)The risk of VOD was only increased in this report if patients had elevations of ALT at the time of BMT. Most other reports suggest that the rate of VOD in patients with HCV is approximately 8%, roughly that seen in HCV-negative recipients (Locasciulli) Over the long term, patients with hepatitis C do show features of mild,

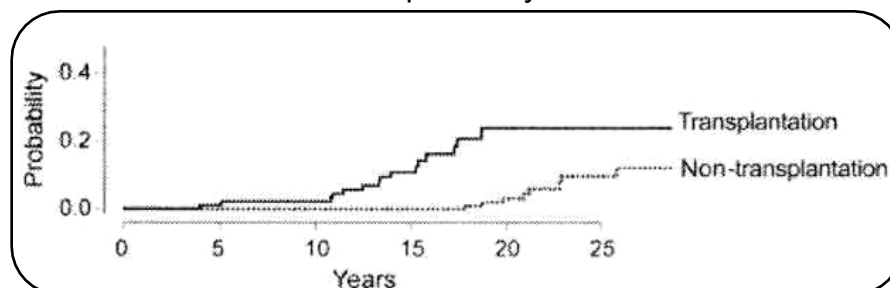
chronic hepatitis after BMT. AST levels are generally higher for 5 to 10 years, although the risk of fulminant hepatic failure (FHF) is not increased. There does not appear to be excessive mortality in long-term (> 3 years) survivors who are HCV+.

Patients with hepatitis C who undergo BMT do appear to be at higher risk of developing cirrhosis than similar patients who do not undergo BMT. As shown in Figure 1, in one series measuring time to progression to cirrhosis (from time of infection with HCV), the median time to cirrhosis was 18 years in BMT recipients versus 40 years in non-transplant patients (Peffault de Latour 2004) The cumulative incidence of cirrhosis in transplanted patients was 24% at 20 years.

A European prospective trial of 195 patients who had undergone stem cell transplant (134 allogenic, 61 autologous hematopoietic HCT) demonstrated an overall survival probability of 82% and 6.1% death rate due to liver disease. The rate of decompensated liver disease and death was 12% at 20 years post-transplant. HCV infection was associated with increased risk of morbidity and mortality while treatment was associated with improved outcomes (Ljungman 2012)

In both the liver transplant and HCT settings, use of mycophenolate mofetil has been linked to development of fibrosing cholestatic hepatitis C, thus this drug should not be used in HCV-infected patients (Evans 2015).

**Figure 1.** Cumulative Incidence of Cirrhosis Reported by Peffault de Latour *et al.*<sup>4</sup>



## Hepatitis E Virus (HEV)

HEV is a positive-stranded RNA virus that is associated with increasing numbers of infections, estimated at 20 million cases annually, including 3.3 million symptomatic cases (Rein 2012). It is gaining attention in developed countries due its zoonotic transmission and seroprevalence rates, ranging from 0.6% to 52.5% in European studies from 2003-2015 (Hartl 2016). The main route of transmission of HEV in high-income countries is foodborne. Transfusion or transplant-transmitted HEV infections have been observed sporadically, with one report from the EU confirming 1 in 3019 blood donations were found to be HEV positive (Boland 2019). In one UK study, it was estimated that the transfusion risk of HEV infection exceeds the annual dietary risk when more than 13 individual donor components are transfused (Tedder 2017).



HEV infection can manifest as either acute or chronic hepatitis and last more than 3-6 months with possible extrahepatic manifestations (Pischke 2017). HEV is now acknowledged as a cause of cirrhosis and end-stage liver disease in individuals who are immunocompromised, including hematological patients in the EU (von Felden 2019). The prevalence of HEV infection in HCT recipients varies between 0 and 3.4% in countries in Europe, and up to 3.9% in China, with a pooled prevalence of positive HEV-RNA of 3.0% (95% CI 2.3-4.0) (Mikulska 2025). Delays in diagnosis can occur due to inadequate awareness and cases of HEV-related liver failure and death have also been reported, particularly in patients whose hepatitis was attributed to other causes (e.g. GVHD) and HEV testing was delayed (Furfaro 2020).

Since HEV-specific antibodies might be absent in up to 75% of viremic donors and HEV viremia might be persistent for more than 3 months (Wen 2018), HEV RNA screening of donors has been proposed and is universal in some countries (e.g. Ireland, UK, Netherlands, Switzerland, Japan). Currently, Health Canada does not require testing of HEV in blood products, but it should be noted that province of Quebec has a prevalence of anti-HEV and HEV RNA positivity similar to some European countries (Fearon 2017). There have been two reported cases of detecting HEV RNA on the day of stem cell donation in asymptomatic donors, in which cases the stem cells were not infused (O'Donghaile 2017, Koenecke (2014). A single case of successful transplantation from a viremic donor in which the recipient was preventively treated with ribavirin has been reported (Boriskina 2020). It is currently not possible to establish the risk-to-benefit ratio of systematically testing stem cell donors for HEV RNA or delaying transplantation to search for an alternative donor.

Immunocompromised patients should avoid consumption of undercooked meat (pork, wild boar and venison) and shellfish (Dalton 2018). Donors should follow dietary advice within 6 weeks of stem cell harvesting to prevent acute HEV infection.

Treatment of chronic HEV infection in patients who are immunocompromised includes the reduction of immunosuppression and the use of oral ribavirin. In regard to ribavirin, high efficacy and low rate of side effects were observed overall in the HCT setting (Xhaard 2019). Suppression of HEV RNA under the limit of detection can guide cessation of ribavirin treatment after 3 months of negativity. Interferons have also been shown to be effective against chronic HEV infection in the hematologic setting (Mallet 2017).

## Veno-Occlusive Disease/ Sinusoidal Obstruction Syndrome (VOD/SOS)

Hepatic veno-occlusive disease (VOD), increasingly referred to as sinusoidal obstruction syndrome (SOS), is a well-recognized complication of all stem cell transplantations, irrespective of the stem cell source, type of conditioning therapy, or underlying disease. Although the incidence has decreased in recent years, it is still between 5 and 15% for myeloablative transplants and up to 5% after RIC alloHCT. The table below describes the patient, disease, and transplant factors associated with SOS.

**Table 2.** Patient, disease, and transplant factors associated with SOS

| Patient Factors        | Disease Factors                     | Transplant Factors             |
|------------------------|-------------------------------------|--------------------------------|
| Prior liver disease    | Advanced disease                    | Ablative conditioning          |
| Active hepatitis       | Malignant disease                   | Non T-cell depleted transplant |
| Age $\geq$ 20 years    | Prior SCT                           | High dose TBI                  |
| Prior fungal infection | Abdominal radiation                 | Oral or High Busulfan AUC      |
| Hepatitis C infection  | Prior hepatotoxic chemotherapy      | Unrelated or mismatched donor  |
| Iron overload          | Gemtuzumab or inotuzumab ozogamicin | Sirolimus GVHD prophylaxis     |
| HFE C282Y genotype     |                                     | Norethistrone use              |
| Steatohepatitis        |                                     |                                |

It is recommended that patients who have risk factors for SOS, particularly those with history of liver disease, iron overload, hepatitis (viral or steato-) or past hepatotoxic therapy, undergo a fibroscan to assess baseline fibrosis/cirrhosis risk, and consider hepatology referral if abnormal.

Prior to assuming SOS however, it is important to consider and rule out: congestive heart failure, fungal or viral liver infections, sepsis- or drug-induced cholestasis, and tumour infiltration of the liver.

The diagnosis of SOS has traditionally used either the Seattle Criteria<sup>1</sup> or the Baltimore Criteria<sup>2</sup> (see Appendix 1). The Baltimore criteria is more specific and less discrepant when corroborated with histopathology and hemodynamic studies, particularly for classic VOD/SOS (onset <21 days). Late onset VOD/SOS does not require hyperbilirubinemia and should be considered when at least two other clinical manifestations are present along with hemodynamic and/or ultrasound evidence of VOD/SOS is confirmed. In an effort to improve early diagnosis and promote early intervention, the European Group for Blood and Marrow Transplantation (EBMT) have revised these diagnostic criteria for adults (Table 3).

**Table 3.** EBMT criteria for VOD/SOS diagnosis in adults<sup>3, 4</sup>

| Classical VOD/SOS                                                                                                                                                                                                              | Late-onset VOD/SOS (>21 days after HSCT)                                                                                                                                                                                                                                                      |
|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| In the first 21 days after HSCT, hyperbilirubinemia (>34 $\mu\text{mol/L}$ ) AND 2 of the following <ul style="list-style-type: none"> <li>• Painful hepatomegaly</li> <li>• Weight gain &gt; 5%</li> <li>• Ascites</li> </ul> | (a) Classical VOD/SOS beyond day 21<br>(b) Histologically proven VOD/SOS<br>(c) Two or more of the following PLUS hemodynamic and/or ultrasound evidence of VOD/SOS <ul style="list-style-type: none"> <li>• Painful hepatomegaly</li> <li>• Weight gain &gt;5%</li> <li>• Ascites</li> </ul> |

The Cairo/Cooke revised diagnostic criteria<sup>5</sup> for VOD/SOS in children and adults has also been proposed that is not dependent on timing post HSCT and can be diagnosed with (a) Any two of the following – elevated bilirubin (>34  $\mu\text{mol/L}$ ), unexpected weight gain >5%, excessive platelet transfusions consistent with refractory thrombocytopenia post HSCT, hepatomegaly for age or increased size over pre-HSCT, RUQ pain, ascites confirmed by exam or imaging studies, reversal of portal venous flow by doppler ultrasound, OR (b) hepatic biopsy consistent with VOD/SOS, OR (c) unexplained elevated portal venous wedge pressure. The primary goal of these proposed criteria was to incorporate more signs and increase awareness for earlier detection and more rapid institution of therapy in hopes of improving survival.

Neither the EBMT criteria nor the Cairo/Cooke criteria have been prospectively validated in clinical trials.

Ultrasound features associated with SOS include: increased GB thickness, elevated hepatic artery resistive index (SV-DV/SV), decreased portal flow, and ascites. However, ultrasound results generally have low sensitivity and specificity. Ascites generally shows a high serum-albumin ascites gradient (>11.1 gm/l). Serial doppler ultrasounds post transplant can be considered in patients with risk factors for VOD/SOS or for those with established mild VOD/SOS to monitor for progression.

The use of transvenous liver biopsy has been shown to confirm diagnosis or reveal an alternate diagnosis in the majority of cases of early posttransplant liver disease. Shulman *et al.* reviewed 60 BMT patients with liver dysfunction who underwent transvenous liver biopsy and measurement of the hepatic venous pressure gradient<sup>4</sup>. The wedged hepatic venous pressure gradient  $\geq 10$  mmHg correlated with a histologic diagnosis of SOS ( $p = 0.001$ ), and this gradient value provided 91% specificity and 86% positive predictive value. Bleeding complications were reported in 11 cases, and there were 3 procedure-related deaths.

In order to promote earlier therapeutic intervention, the EBMT also proposed criteria for severity of grading of VOD/SOS once the diagnosis was made<sup>3</sup> (Table 4). Four stages of severity (mild, moderate, severe, and very severe) are based on five parameters: time since first clinical manifestation of VOD/SOS, bilirubin level and kinetics, transaminase level, weight gain, and renal

function. Importantly, in the presence of two or more risk factors, patients are classified in the upper grade. These criteria were validated in a study of 203 patients with VOD/SOS<sup>6</sup>: 5.9% were mild, 12.8% moderate, 18.2% severe, and 63.1% very severe; the day 100 OS of these groups were 83.3%, 84.3%, 94.6%, and 58.6%, respectively. The day 100 TRM was significantly higher in very severe VOD/SOS at 36.7%, compared to 8.3% in mild, 8.0% in moderate, and 2.7% in severe ( $p<0.0001$ )<sup>6</sup>.

**Table 4.** EBMT criteria for severity grading of suspected SOS/VOD in adults<sup>4</sup>

|                                               | Mild*                        | Moderate*                             | Severe                              | Very Severe<br>(multi-organ<br>dysfunction/failure)  |
|-----------------------------------------------|------------------------------|---------------------------------------|-------------------------------------|------------------------------------------------------|
| Time since first clinical symptoms of SOS/VOD | >7 days                      | 5-7 days                              | ≤4 days                             | Any time                                             |
| Bilirubin (umol/L)                            | ≥34 and <51                  | ≥51 and <85                           | ≥85 and <136                        | ≥136                                                 |
| Bilirubin kinetics                            |                              |                                       | Doubling within 48h                 |                                                      |
| Transaminases                                 | ≤2x normal                   | >2 and ≤5x normal                     | >5 and ≤8x normal                   | >8x normal                                           |
| Weight Increase                               | <5%                          | ≥5% and <10%                          | ≥5% and <10%                        | ≥10%                                                 |
| Renal function                                | <1.2x baseline at transplant | ≥1.2 and <1.5x baseline at transplant | ≥1.5 and <2x baseline at transplant | ≥2 baseline at transplant, or other signs of MOD/MOF |

Patients belong to the category that fulfills two or more criteria, classified into the most severe category if criteria achieved in more than one category

\*Patients with two or more *risk factors* for SOS/VOD (Table 2) are classified in the upper grade

## Treatment of VOD/SOS

Defibrotide (DF) is the only agent with proven efficacy for the treatment of severe/very severe VOD/SOS. Defibrotide is a single-stranded polydeoxyribonucleotide that has anti-inflammatory and antithrombotic properties. *Richardson et al.* reported on the use of defibrotide in 88 patients who developed severe SOS and multisystem organ failure after stem cell transplantation<sup>7</sup>. The patients ranged in age from 8 to 62 years (mean 35 years), and were assessed according to the Baltimore Criteria. Defibrotide was administered IV in doses ranging from 5 to 60 mg/kg per day for a median of 15 days. Complete resolution of SOS was reported in 32 patients (36%), with 35% survival at day +100. There was no worsening of clinical bleeding or attributable grade III or IV toxicity noted in the patients. Grade I/II toxicities included hypotension, fever, abdominal cramping, and hot 75 patients on 40 mg/kg/day of defibrotide. The 141 evaluable patients ranged in age from 0.5 to 63 years (mean 36 years), and 99% of patients were in multisystem organ failure. Complete resolution of SOS was reported in 65 patients (46%), with an overall survival rate of 42% at day +100. There was no difference in response rates between the 2 doses, but the higher dose was associated with more grade III and IV toxicities, as well as a greater risk of bleeding. Early stabilization or lower bilirubin was associated with better outcome<sup>9</sup>. The final results from a defibrotide treatment-IND study for 1000 patients with hepatic VOD/SOS demonstrated Day +100 survival was 58.9% overall; 67.9% in pediatric patients and 47.1% in adult patients, and higher in the subgroup of patients without multi-organ dysfunction (MOD)<sup>9</sup>. Similarly, a systematic review of 17 defibrotide studies in the treatment of

VOD/SOS demonstrated that among those treated with 25 mg/kg/day dosing the Day+100 survival rate was 56%, higher in patients without MOD at 71% vs. 44% with MOD<sup>10</sup>.

Given the mortality associated with severe and very severe VOD/SOS, it is mandatory to treat these patients promptly, with initiation of DF as soon as possible. Standard dosing for defibrotide is 25 mg/kg/day with recommended duration of at least 21 days and until resolution of all VOD/SOS symptoms. No dose adjustments are required with renal failure, while in obese patients corrected body weight should be used for dose calculation. In patients where such resolution happens before 21 days, it is possible to stop DF earlier (e.g. to facilitate patient discharge) with close monitoring for the rare possibility of recurrence.

The indication of DF treatment in patients with mild or moderate VOD/SOS is more questionable. In expanded/compassionate access treatment protocols, an earlier treatment initiation after VOD/SOS diagnosis was associated with higher day +100 OS ( $p < 0.001$ )<sup>11</sup>. Up to 32% mortality was seen despite treatment, which while favorable compared to severe/very severe VOD/SOS, it remains significant. The EBMT recommends that patients who fulfill the diagnosis criteria and whose severity grading is moderate should be considered for preemptive DF and closely followed. In patients with mild VOD/SOS, supportive care should be intensified and severity criteria monitoring should be strictly applied to allow immediate DF initiation in case of deterioration.

### Supportive Care Measures

- Careful management of fluid and sodium balance to limit third-space fluid and maintain renal perfusion.
- Limit hepatotoxic medications; wherever possible, antifungal azoles should be substituted for echinocandins
- Transjugular intrahepatic portosystemic shunt (TIPS) may improve fluid balance and symptom control with no benefit on survival.
- While on DF, discontinue any other agents that may increase risk of bleeding.
- Nutritional support is also important, and enteral nutrition should be favored to prevent patient's malnutrition; parental nutrition is associated with fluid overload, infectious complications, and hepatotoxicity and should be avoided.

### Preventative Therapy

Results of a randomized controlled trial of ursodiol for SOS prophylaxis were reported by Essell *et al.*<sup>12</sup> The patients were 67 consecutive recipients of allogeneic BMT, and they all received a busulfan plus cyclophosphamide conditioning regimen. Patients were randomly assigned to receive ursodiol, 300 mg twice daily (or 300 mg in the morning and 600 mg in the evening if body weight was > 90 kg), or placebo until day +80. The incidence of SOS was 40% (13 of 32 patients) in placebo recipients and 15% (5 of 34 patients) in ursodiol recipients ( $p = .03$ ). The authors concluded that ursodiol prophylaxis seemed to decrease the incidence of hepatic complications after allogeneic BMT. A larger

randomized controlled trial involving 242 patients reported no significant impact of ursodiol on the incidence of SOS, but did report significantly lower incidences of grades III and IV acute GVHD, stage II and IV liver and intestinal GVHD, and stage III and IV skin GVHD<sup>13</sup>. In addition, among the patients given ursodiol, the survival at 1 year was significantly better, (71% versus 55%,  $p=.02$ ), and the non-relapse mortality rate was lower (19% versus 34%,  $p=.01$ ), when compared to the control group. In a long-term (10 year) follow-up of this study, the difference in survival and NRM in favor of the ursodiol-treated group was maintained<sup>14</sup>. A systematic review of three RCTs, including the two mentioned above, of ursodiol as compared to placebo demonstrated a reduced risk of SOS on ursodiol; RR 0.34, 95% CI 0.17-0.66 although no significant difference in survival<sup>15</sup>. It is recommended that all patients undergoing allogeneic SCT, regardless of conditioning used, to proceed with ursodiol prophylaxis.

Potentially there may be a role for defibrotide as prophylaxis for SOS; a systematic review of 1230 patients from one RCT, 4 cohort studies and 8 case series studies showed an incidence of about 5% with defibrotide versus controls (14%) with a relative risk of 0.46 (95% CI 0.31-0.73)<sup>16</sup>. British guidelines suggest giving defibrotide at 6.25 mg/kg IV q.i.d. for prophylaxis in adults undergoing allogeneic stem cell transplant with a history of pre-existing liver disease, second myeloablative transplant, allogeneic transplant for leukemia beyond second relapse, conditioning with busulfan-based regimens, past treatment with gemtuzumab or inotuzumab ozogamicin, diagnosis of primary hemophagocytic lymphohistiocytosis, adrenoleucodystrophy or osteopetrosis<sup>17</sup>. However, defibrotide did not show a benefit in the prophylaxis of SOS in a randomized phase 3 trial in pediatric and adult patients ( $n=372$ ) undergoing allo or auto SCT<sup>18</sup>. Given that current studies in this area are limited, as well as the considerable cost and lack of access of defibrotide, further research is needed and routine use of defibrotide is not routinely recommended.



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## Appendix

### Comparison of Seattle and Baltimore Diagnostic Criteria for SOS

| Seattle Criteria <sup>22</sup>                                                              | Baltimore Criteria <sup>23</sup>                                                                 |
|---------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------|
| Development of 2 of the following within 20 days of transplant:                             | Hyperbilirubinemia (> 34 micromolar) within 21 days of transplant <i>and</i> 2 of the following: |
| <ul style="list-style-type: none"> <li>• Hyperbilirubinemia (&gt; 34 micromolar)</li> </ul> | <ul style="list-style-type: none"> <li>• Ascites</li> </ul>                                      |
| <ul style="list-style-type: none"> <li>• Tender hepatomegaly</li> </ul>                     | <ul style="list-style-type: none"> <li>• Hepatomegaly (may be painful)</li> </ul>                |
| <ul style="list-style-type: none"> <li>• Weight gain (&gt; 2%)</li> </ul>                   | <ul style="list-style-type: none"> <li>• Weight gain (&gt; 5%)</li> </ul>                        |

# Management of Cytokine Release Syndrome and Immune Effector Cell-Associated Neurotoxicity Syndrome

Presented by: Robert Puckrin

## Summary

- Cytokine Release Syndrome (CRS) and Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS) are common after immune effector cell therapy and may present with fever, hypotension, hypoxia, organ dysfunction, and neurological symptoms during the first weeks after infusion
- The ASTCT Consensus Grading system is used to determine severity of CRS based on changes in vital signs (temperature, hypotension, and hypoxia) and of ICANS based on changes in ICE score, level of consciousness, seizure, motor findings, elevated intracranial pressure, or cerebral edema.
- Management of CRS consists of antipyretics, intravenous fluids, supplemental oxygen, tocilizumab, corticosteroids, and treatment of concurrent infection. Corticosteroid prophylaxis may be considered for patients at high risk of CRS.
- Management of ICANS consists of supportive care and corticosteroids. Tocilizumab may be given if there is concurrent CRS. Careful evaluation for possible metabolic, medication-associated, and infectious causes is essential, including neuroimaging and CSF evaluation when appropriate. Anti-epileptic drug prophylaxis should be given to all patients with ICANS and considered from the time of CAR-T cell infusion for those at increased risk of ICANS. Anakinra should be considered for patients with steroid-refractory and/or severe CRS or ICANS.

## Background

A range of unique toxicities has been observed in patients treated with CAR T-cell therapy, including Cytokine Release Syndrome (CRS), Immune Effector Cell-Associated Neurotoxicity Syndrome (ICANS), cytopenias, infections, B-cell aplasia, and secondary malignancies (1). Table 1 summarizes the rate of CRS and ICANS observed in published trials.

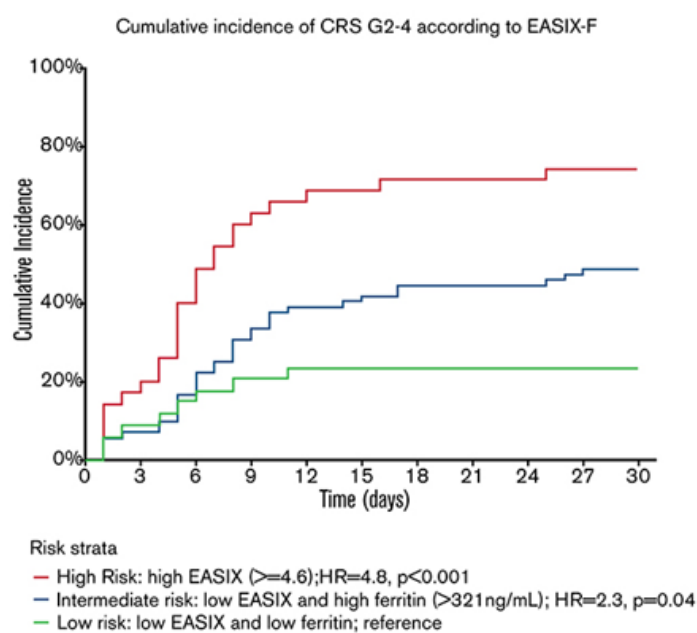
**Table 1.** Frequency of CRS and ICANS observed in key clinical trials.

| Trial                  | Product  | CRS | Grade ≥3 CRS* | ICANS | Grade ≥3 ICANS |
|------------------------|----------|-----|---------------|-------|----------------|
| <b>DLBCL</b>           |          |     |               |       |                |
| ZUMA-1 <sup>2</sup>    | Axi-cel  | 93% | 13%           | 64%   | 28%            |
| JULIET <sup>3</sup>    | Tisa-cel | 58% | 23%           | 20%   | 11%            |
| TRANSCEND <sup>4</sup> | Liso-cel | 42% | 2%            | 30%   | 10%            |
| ZUMA-7 <sup>5</sup>    | Axi-cel  | 92% | 6%            | 60%   | 21%            |
| BELINDA <sup>6</sup>   | Tisa-cel | 61% | 5%            | 10%   | 3%             |
| TRANSFORM <sup>7</sup> | Liso-cel | 49% | 1%            | 12%   | 4%             |

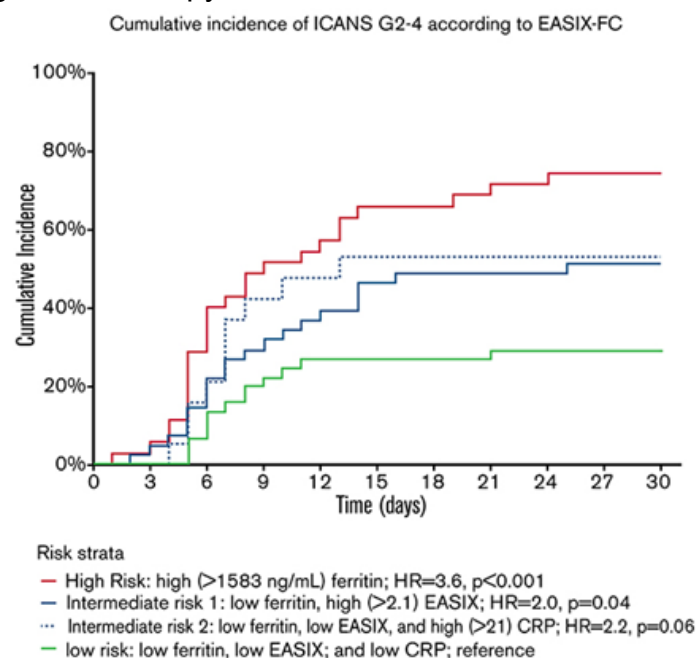
| Mantle cell               |           |     |         |     |     |
|---------------------------|-----------|-----|---------|-----|-----|
| ZUMA-2 <sup>8</sup>       | Brexu-cel | 91% | 15%     | 63% | 31% |
| Follicular lymphoma       |           |     |         |     |     |
| ZUMA-5 <sup>9</sup>       | Axi-cel   | 82% | 7%      | 59% | 19% |
| ELARA <sup>10</sup>       | Tisa-cel  | 49% | 0%      | 37% | 4%  |
| B-ALL                     |           |     |         |     |     |
| ELIANA <sup>11</sup>      | Tisa-cel  | 77% | 47% ICU | 40% | 13% |
| ZUMA-3 <sup>12</sup>      | Brexu-cel | 89% | 24%     | 60% | 25% |
| Multiple myeloma          |           |     |         |     |     |
| KaRMMA <sup>13</sup>      | Ide-cel   | 84% | 6%      | 18% | 3%  |
| CARTITUDE-1 <sup>14</sup> | Cilta-cel | 95% | 5%      | 21% | 9%  |

\* Interstudy comparisons of CRS grading is difficult due to variations in the grading systems used

The risk of grade 2-4 CRS or ICANS can be predicted by the modified EASIX score proposed by Greenbaum et al.<sup>15</sup>, which incorporates the EASIX score ( $[\text{Creatinine}/88 \times \text{LDH}]/\text{platelets}$ ), ferritin, and CRP measured at the start of lymphodepleting chemotherapy:



| Alberta CAR-T recipients | Grade 2-4 CRS |
|--------------------------|---------------|
| Low EASIX-F (n=6)        | 16%           |
| Int./high EASIX-F (n=17) | 65%           |



| Alberta CAR-T recipients  | Grade 2-4 ICANS |
|---------------------------|-----------------|
| Low EASIX-FC (n=7)        | 14%             |
| Int./high EASIX-FC (n=16) | 31%             |

## Clinical Features of Cytokine Release Syndrome

CRS occurs in 50–95% of patients receiving CAR T-cell therapy. Risk factors include high disease burden at infusion, baseline inflammatory state, and CAR construct, with CRS being more common with CD28-based products than with 4-1BB constructs. Most patients present with fever which may reach 40°C and is often accompanied by malaise, myalgias, and nausea. In ZUMA-1, JULIET, and TRANSFORM, median CRS onset was 2 days after axi-cel, 3 days after tisa-cel, and 5 days after liso-cel, with a median duration of 5–8 days. Severe CRS is characterized by hypotension and vasoplegic shock; early vasopressor support is critical. CRS can also cause organ-specific toxicities, including tachycardia, arrhythmias, stress cardiomyopathy, pulmonary edema, renal injury, coagulopathy, and multi-organ dysfunction that may be fatal. It can often be difficult to distinguish fever from infection in this context and it is recommended that patients with CRS should undergo careful screening for infection and receive treatment with antibiotics appropriate to their clinical presentation.

## General Care of the CAR T-Cell Recipient

Frequent physician and nursing assessment is essential for safe CAR T-cell management. Most cases of CRS and ICANS develop within the first 2 weeks post-infusion. Vital signs should be monitored closely, and medical staff notified of any fever ( $\geq 38^{\circ}\text{C}$ ), hypotension (SBP  $\leq 90$  mmHg), tachycardia (HR  $\geq 120$  bpm), hypoxia ( $\text{SpO}_2 \leq 90\%$ ), or new organ toxicity. Any neurological change (declining ICE score, abnormal movements, altered alertness, or visual symptoms) should be reported promptly. CRS and ICANS grade should be documented in the medical record at least once daily.

A daily physical exam and review of systems should be completed. ICANS screening should be performed at least every 12 hours using a standardized tool (e.g., ICE). Daily labs should include CBC, electrolytes, creatinine, calcium, magnesium, phosphate, uric acid, liver enzymes, PTT/INR, fibrinogen, CRP, and ferritin, with more frequent testing as clinically indicated. Patients with significant tumour burden require tumor lysis prophylaxis and monitoring.

## Grading Cytokine Release Syndrome

Several CRS grading systems have been used in the clinical management of CAR T-cell recipients, and the ASTCT grading system is currently the most widely used<sup>16</sup> (Table 2).

**Table 2.** ASTCT cytokine release syndrome grading (per Lee et al.<sup>16</sup>).

|             | Grade 1                | Grade 2                    | Grade 3                 | Grade 4                         |
|-------------|------------------------|----------------------------|-------------------------|---------------------------------|
| Fever*      | Yes, $\geq 38^{\circ}$ | Yes, $\geq 38^{\circ}$     | Yes, $\geq 38^{\circ}$  | Yes, $\geq 38^{\circ}$          |
|             |                        | with                       |                         |                                 |
| Hypotension | None                   | Not requiring vasopressors | Requiring a vasopressor | Requiring multiple vasopressors |



|           |      |                                                             | with or without vasopressin                                                               | (excluding vasopressin)                                                          |
|-----------|------|-------------------------------------------------------------|-------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------|
|           |      |                                                             | with                                                                                      |                                                                                  |
| Hypoxia** | None | Requiring low-flow nasal cannula ( $\leq 6$ LPM) or blow-by | Requiring high-flow nasal cannula ( $> 6$ LPM), facemask, non-rebreather, or Venturi mask | Requiring positive pressure (CPAP, BiPAP, intubation and mechanical ventilation) |

Organ toxicity may be graded according to CTCAE Version 5.0 (2017)

[https://ctep.cancer.gov/protocolDevelopment/electronic\\_applications/ctc.htm](https://ctep.cancer.gov/protocolDevelopment/electronic_applications/ctc.htm) but does not change grade.

\* Fever is defined as temperature  $\geq 38$  degrees not attributable to other cause. In patients who have CRS then receive antipyretic or anticytokine therapy such as tocilizumab or steroids, fever is no longer required to grade CRS. In this case CRS grading is driven by hypotension and/or hypoxia.

\*\* Hypoxia should not be explained by other causes i.e. rigors or sedation in order to meet the definition of hypoxia in CRS.

\*\*\* Low-flow is defined as oxygen delivered at  $\leq 6$  LPM. Low flow also includes blow-by oxygen delivery, sometimes used in pediatrics. High-flow nasal cannula is defined as oxygen delivered at  $> 6$  LPM.

## Management of Cytokine Release Syndrome

Cytokine release syndrome should be suspected in patients who develop new fever ( $\geq 38^\circ\text{C}$ ), hypotension ( $\text{SBP} \leq 90$  mmHg), hypoxia ( $\text{SpO}_2 \leq 90\%$ ) or organ toxicity. Careful clinical evaluation, including blood cultures, viral studies (respiratory virus panel, CMV and EBV PCR) and imaging tests should be carried out as appropriate to assess for other causes of these findings. Infection, septic or cardiogenic shock, venous thromboembolism, alveolar hemorrhage, tumor lysis syndrome and other syndromes may resemble CRS in their initial presentations and these diagnoses should be either treated empirically or excluded through appropriate investigations. CRS should be managed according to grade as described in Table 3, which is a harmonized summary of the management guidelines proposed by EBMT, ASCO, NCCN, CARTOX groups and ZUMA-1 safety cohorts 4 and 6<sup>17-21</sup>.

Principles of CRS management include the following:

1. The anti-IL6 receptor antibody tocilizumab and corticosteroids are the mainstays of CRS management. Their use does not appear to negatively impact important outcomes of CAR T-cell therapy such as progression-free and overall survival<sup>19, 20, 22, 23</sup>. However, high-dose or prolonged corticosteroid use has been associated with poor outcomes in some studies<sup>24</sup>, hence corticosteroids should be rapidly tapered upon improvement and resolution of CRS.
2. Hypotension that persists after 1-2 liters of 0.9% normal saline is unlikely to respond to further fluid resuscitation, due to the presence of vascular leak, vasoplegia and occult stress cardiomyopathy. Hypotension that fails to respond to two fluid challenges and tocilizumab should be treated with vasopressors.

3. In general CRS that develops shortly (<24-72 hours) after CAR T-cell infusion has a more aggressive course and requires more intensive treatment than that arising more gradually. In addition, patients with comorbid medical conditions may experience a more complicated course with CRS and so early initiation of anticytokine therapy is warranted in this population.
4. The results of ZUMA-1 safety cohort 6 showed that low-dose corticosteroid prophylaxis may reduce the risk of severe CRS and severe ICANS compared to historical controls<sup>20</sup>. However, interpretation of this study is limited by its small sample size, single-arm design, and baseline differences in tumor burden and inflammatory state. Nevertheless, corticosteroid prophylaxis is reasonable for patients receiving axi-cel or brexu-cel who have  $\geq 1$  risk factor for severe CRS or ICANS, such as age >65, high tumor burden, and/or elevated baseline inflammatory markers.

## Pharmacotherapy of CRS

- Tocilizumab
  - Anti-IL6 receptor antibody with most extensive track record in CRS
  - Dose 8 mg/kg (not to exceed 800 mg per dose) IV over one hour
  - Half-life of tocilizumab is 13 days in rheumatoid arthritis. Some guidelines suggest repeat dosing q4-6 hours based on response, up to 3 doses in 24 hours and 4 doses in total
  - May premedicate with Benadryl or Tylenol if not recently given.
- Steroids
  - Dexamethasone 10 mg IV may be given for grade 2 CRS especially if high risk and/or refractory to tocilizumab
  - Higher doses of dexamethasone (e.g. 10mg IV q6h) are used for grade 3 CRS
  - If life-threatening and no response to tocilizumab may give up to Methylprednisolone 500mg IV BID x 3 days
- Other monoclonal antibodies for steroid refractory toxicities
  - Anakinra (preferred for steroid-refractory CRS/ICANS or overlap with HLH)
  - Siltuximab

**Table 3.** Management of CRS after CAR T-cell therapy.

| Grade           | Recommendations                                                                                                                                                                                                                                                                                                                                                                                    |
|-----------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Prophylaxis     | <ul style="list-style-type: none"> <li>▪ Consider dexamethasone 10mg on morning of infusion and on days +1 and +2 if receiving axi-cel or brexu-cel with age &gt;65, intermediate or high modified EASIX, or high tumor burden</li> </ul>                                                                                                                                                          |
| Supportive care | <ul style="list-style-type: none"> <li>▪ Anti-pyretics (e.g. acetaminophen, NSAID, external cooling) as needed</li> <li>▪ Order infectious work-up and start antibiotics if neutropenic, unstable, or infection suspected</li> <li>▪ Provide supplemental oxygen for hypoxia and IV fluids for hypotension with early consideration for vasopressors if hypotension persists after 1-2L</li> </ul> |

|         |                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             |
|---------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Grade 1 | <ul style="list-style-type: none"> <li>▪ Consider tocilizumab x1 dose if fever lasting &gt;24-72 hours, especially if early onset CRS, comorbidities, or high risk for severe CRS</li> </ul>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                |
| Grade 2 | <ul style="list-style-type: none"> <li>▪ Tocilizumab x1 dose → repeat up to 3 doses per day PRN for up to 4 total doses</li> <li>▪ Dexamethasone 10mg x1 dose if high risk and/or refractory to tocilizumab → re-assess in 6 hours</li> </ul>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               |
| Grade 3 | <ul style="list-style-type: none"> <li>▪ Tocilizumab up to 3 doses per day PRN for up to 4 total doses</li> <li>▪ Dexamethasone 10mg q6h → 20mg q6h if refractory</li> </ul>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                |
| Grade 4 | <ul style="list-style-type: none"> <li>▪ Tocilizumab up to 3 doses per day PRN for up to 4 total doses</li> <li>▪ Methylprednisolone 500mg IV BID x 3 days then rapid taper (e.g. 250mg IV BID x 2 days then 125mg IV BID x 2 days then 60mg IV BID until Grade 1 or lower then taper over 1-2 weeks; note: taper can be omitted if symptoms resolve to grade 1 within 1 week)</li> <li>▪ For refractory symptoms lasting &gt;24 hours or rapid deterioration, start anakinra 100mg IV q6h until resolved to grade 1 or lower</li> <li>▪ For ongoing refractory symptoms, consider ruxolitinib, siltuximab, or dasatinib</li> <li>▪ Consider CAR-T ablation with cyclophosphamide or ATG as measure of last resort for life-threatening symptoms</li> </ul> |

## Management of ICANS

Neurologic toxicity is relatively common after CAR T-cell therapy. Risk factors include age >65, CD28-based constructs, high tumor burden, elevated baseline inflammation, and early or severe CRS. ICANS often presents with tremor, inattention, dysgraphia, mild expressive aphasia (e.g., difficulty naming objects), and somnolence. More severe manifestations include ataxia, aphasia, encephalopathy, and seizures. Severe ICANS is characterized by motor weakness, obtundation, increased intracranial pressure, and cerebral edema, which can rarely be fatal. ICANS typically develops a few days after CRS onset. In ZUMA-1, JULIET, and TRANSFORM, median ICANS onset was 5 days after axi-cel, 6 days after tisa-cel, and 9 days after liso-cel, with a median duration of 11–17 days.

Early detection is supported by standardized screening tools such as the ASTCT Immune Effector Cell-associated Encephalopathy (ICE) score (0–10), which can be administered at least twice daily by trained nursing or medical staff.

### ICE tool:

- Orientation to
  - Year – 1 point
  - Month – 1 point
  - City – 1 point
  - Hospital – 1 point

- Naming 3 objects – *up to 3 points*
- Following Commands (eg, Show me two fingers) – *1 point*
- Writing a short sentence – *1 point*
- Attention: Count backwards from 100 by ten – *1 point*

The ICE tool has been integrated into a comprehensive ASTCT ICANS grading system (Table 4). ICANS management according to grade is outlined in Table 5. The mainstay of treatment is supportive care along with corticosteroids. Tocilizumab does not appear to have a significant role in the treatment of ICANS but can be considered if there is concurrent CRS.

**Table 4.** ASTCT Grading Immune Effector Cell-Associated Neurological Syndrome<sup>16</sup>

| Symptom or sign             | Grade 1               | Grade 2          | Grade 3                                                                                                                   | Grade 4                                                                                                                              |
|-----------------------------|-----------------------|------------------|---------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------|
| ICE score                   | 7-9 (mild)            | 3-6 (moderate)   | 0-2 (severe)                                                                                                              | Unable to perform                                                                                                                    |
| Level of Consciousness      | Awakens spontaneously | Awakens to voice | Awakens to touch                                                                                                          | Unarousable or requires vigorous or repeated stimuli to arouse. Stupor or coma                                                       |
| Seizure                     | NA                    | NA               | Any clinical seizure focal or generalized that resolves rapidly or non-convulsive seizure that resolves with intervention | Life-threatening or prolonged seizure (> 5 minutes) or repetitive clinical or electrical seizures without return to baseline between |
| Motor Findings              | NA                    | NA               | NA                                                                                                                        | Deep focal motor weakness such as hemiparesis or paraparesis                                                                         |
| Elevated ICP/Cerebral edema | NA                    | NA               | Focal/local edema on neuroimaging                                                                                         | Diffuse cerebral edema on neuroimaging; decerebrate or decorticate posturing; or CN VI palsy; or papilledema or Cushing's            |

**Table 5. Management of ICANS**

|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                |
|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>Prophylaxis</b> <ul style="list-style-type: none"> <li>Consider dexamethasone 10mg on morning of infusion and on days +1 and +2 if receiving axi-cel or brexu-cel with age &gt;65, intermediate or high modified EASIX, or high tumor burden</li> <li>Consider levetiracetam 500mg BID prophylaxis from day 0 to +14 for patients receiving axi-cel or brexu-cel or who have other risk factors for ICANS. Duration of levetiracetam should be extended for patients who develop seizures or have ongoing ICANS, in conjunction with the neurology service.</li> </ul>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      |
| <b>Grade 1</b> <ul style="list-style-type: none"> <li>Aspiration precautions, intravenous hydration</li> <li>Withhold oral intake of food, medicines, and fluids and assess swallowing</li> <li>Convert all medications and nutrition to IV if swallowing is impaired</li> <li>Avoid sedating medications</li> <li>Low-dose lorazepam (0.25-0.5 mg IV q8h) or haloperidol (0.5 mg IV q6h) for agitated patients</li> <li>Fundoscopy exam for papilledema</li> <li>Consider MRI of the brain with and without contrast +/- MRI spine if focal neurological deficits</li> <li>Consider diagnostic LP if other causes of encephalitis suspected</li> <li>Consider tocilizumab 8 mg/kg IV if ICANS occurs in setting of CRS</li> <li>Dexamethasone 10 mg IV x 1 dose and reassess in 6 hours</li> <li>Start levetiracetam 500 mg bid prophylaxis if not already given</li> </ul>                                                                                                                                                                                                                                   |
| <b>Grade 2</b> <ul style="list-style-type: none"> <li>Workup and supportive care as described above</li> <li>Tocilizumab 8 mg/kg IV if ICANS occurs in setting of CRS</li> <li>Dexamethasone 10 mg IV q6-12h. Once ICANS improves to grade 1 or less taper and/or discontinue steroids if clinically appropriate</li> <li>Start levetiracetam 500 mg bid prophylaxis if not already given</li> </ul>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           |
| <b>Grade 3</b> <ul style="list-style-type: none"> <li>Workup and supportive care as described above</li> <li>ICU transfer</li> <li>Neurology consultation</li> <li>Tocilizumab 8 mg/kg IV if ICANS occurs in setting of CRS, if not administered previously</li> <li>Dexamethasone 10-20 mg IV q6h for most patients. Consider treating focal brain edema with methylprednisolone 500mg IV BID x 3 days</li> <li>Increased intracranial pressure should be treated according to standard guidelines with acetazolamide 1000 mg IV followed by 250-1000 mg q12h (based on renal function and acid/base balance), elevate HOB.</li> <li>Diagnostic LP if no improvement with treatment or if other causes of encephalitis suspected</li> <li>Consider repeat neuroimaging every 2-3 days</li> <li>Antiepileptic drugs as prescribed by neurology (avoid phenytoin and lacosamide due to cardiotoxicity)</li> </ul>                                                                                                                                                                                               |
| <b>Grade 4</b> <ul style="list-style-type: none"> <li>Supportive care and workup as described above</li> <li>ICU monitoring and mechanical ventilation for airway protection</li> <li>Tocilizumab 8 mg/kg IV if ICANS occurs in setting of CRS, if not administered previously</li> <li>Treat with methylprednisolone 500mg IV BID x 3 days then 250mg IV BID x 2 days then 125mg IV BID x 2 days then 60mg IV BID until Grade 1 or lower then taper over 1-2 weeks (Note: taper can be omitted if symptoms resolve to grade 1 within 1 week)</li> <li>For refractory symptoms lasting &gt;24 hours or rapid deterioration, start anakinra 100mg IV q6h until resolved to grade 1 or lower</li> <li>For ongoing refractory symptoms, consider intrathecal chemotherapy (e.g. hydrocortisone 50mg + methotrexate 12mg ± cytarabine 50mg), siltuximab, ruxolitinib, or dasatinib.</li> <li>CAR-T ablation with cyclophosphamide or ATG may be considered as measure of last resort for life-threatening symptoms</li> <li>For convulsive status epilepticus treat according to established guidelines</li> </ul> |

- Cerebral edema should be treated as per established guidelines, including hyperventilation, hyperosmolar therapy, frequent metabolic profiling and neurosurgical or anesthesia consultation for burst-suppression pattern EEG
- Antiepileptic drugs as prescribed by neurology (avoid phenytoin and lacosamide due to cardiotoxicity)

## Immune Effector Cell-Associated HLH-like Syndrome (IEC-HS)

IEC-HS is a rare hyperinflammatory syndrome distinct from CRS and ICANS that resembles Hemophagocytic Lymphohistiocytosis / Macrophage Activation Syndrome (HLH/MAS) and occurs in <5% of CAR T-cell recipients. It is characterized by new or worsening cytopenias, hyperferritinemia, hypofibrinogenemia, and/or transaminitis. While symptoms may overlap with severe CRS, the key distinguishing feature of IEC-HS is its delayed onset, typically emerging as CRS is resolving or has resolved. ASTCT diagnostic criteria and grading have been established (25). First-line treatment is anakinra (e.g., 100 mg IV q6–12h) ± dexamethasone 10 mg IV q6h. Severe or refractory cases may require ruxolitinib, low-dose etoposide (e.g., 50-100 mg/m<sup>2</sup> weekly as needed), or emapalumab.



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# Transfusions and Management of Cytopenias Early Post-HSCT

Presented by: Jason Tay

## Summary

1. RBC transfusion threshold: We recommend the provision of 1 unit of red cells based on daily CBC demonstrating a hemoglobin  $<70\text{g/L}$  (15 mL/kg to a maximum of 1 unit for pediatrics). The use of a higher red cell transfusion threshold would be at the discretion of the clinician based on clinical judgement such as symptoms of anemia.
2. PLT transfusion threshold: We recommend the provision of 1 unit of pooled platelets based on daily CBC demonstrating platelets  $<10 \times 10^9/\text{L}$  (10 mL/kg, maximum of 1 unit for pediatrics). The use of a higher platelet transfusion threshold would be at the discretion of the clinician based on clinical judgement such as active bleeding.
3. We recommend the routine use of prophylactic platelets in patients undergoing allogeneic HCT.
4. It is reasonable to consider a therapeutic-only (only if clinically relevant bleeding) in patients undergoing autologous HCT. It may be also reasonable to consider their use in exceptional circumstances in both allogeneic or autologous HCT settings e.g., Jehovah's Witnesses.
5. We utilize irradiated cellular blood products from the Canadian Blood Services (universally leukoreduced and CMV Safe) for patients who have received a Hematopoietic Cell Transplant (HCT). There will be NO need to use irradiated red cells for red cells that are older than 14 days.
6. We do NOT recommend choosing blood products based on duration of storage.
7. We recommend the routine use of G-CSF post-autologous HCT starting on Day 7 until  $\text{ANC} > 0.5 \times 10^9/\text{L}$ .
8. We do NOT recommend the routine use of G-CSF post-allogeneic HCT. G-CSF can be considered for non-malignant pediatric disorders.
9. We do NOT recommend the routine use of erythropoietin post-HCT. It may be reasonable to consider its use in exceptional circumstances e.g., Jehovah's Witnesses.
10. It is reasonable to consider on a case-by-case basis, the use of Thrombopoietin receptor agonists (e.g. eltrombopag) in circumstances of prolonged isolated thrombocytopenia post-HCT or secondary failure of platelet recovery.

11. For stem cell boost or 2nd HCT, see chapter on Graft Failure / Poor Graft Function.

## Background

High dose chemotherapy with or without radiation followed by hematopoietic stem cell (HSC) rescue, typically suppress the production of blood cells by the bone marrow for 7 to 14 days. This results in aplasia, during which the patient is likely to require prophylactic or therapeutic transfusions of red cells and platelets. This is particularly salient in the allogeneic setting where longer periods of transfusion support may be required, when recovery is complicated by delayed engraftment, acute graft-versus-host disease (GvHD) or severe sepsis.

There is a paucity of randomized controlled trial data to guide transfusion practice in the HCT setting<sup>1</sup>. Indeed, guidelines for transfusion support are often extended to the hematopoietic cell transplant (HCT) setting from general oncology/medicine despite the lack of strong clinical trial evidence (insert guidelines)<sup>2-6</sup>.

## Transfusion Utilization

It has been estimated that up to 39% of the total blood transfused in the USA in 2017 was used by inpatient medicine (including hematology/oncology) patients<sup>7</sup>, while an estimate of 27.1% in hematology/oncology was noted in the UK in 2014<sup>8</sup>. The frequency of transfusion support is the highest post-conditioning chemoradiotherapy and decreases significantly after the 1<sup>st</sup> 30 days consistent with the timing of stem cell engraftment. For instance, Xenocostas et al. reported that the mean number of red cell units transfused per patient from 0 to 60 days was 6.8 +/- 6.4; 61 to 120 days, 3.2 +/- 5.5; and 121 to 180 days, 2.0 +/- 4.6<sup>9</sup>. Factors that are associated with avoidance of transfusion after HCT include male sex ( $p = 0.0013$ ), diagnosis, specifically plasma cell dyscrasias ( $p < 0.0001$ ), early-stage disease ( $p = 0.006$ ), and higher baseline hemoglobin (Hb) at time of transplant ( $p < 0.0001$ )<sup>10</sup>. The following Table summarizes selected clinical studies reviewing transfusion utilization in the HCT setting.

**Table 1:** Selected clinical studies reviewing transfusion utilization in the HCT setting

|                                     | Setting               | Timeline     | Red Cell Utilization (units)       | Platelet Utilization (units)       |
|-------------------------------------|-----------------------|--------------|------------------------------------|------------------------------------|
|                                     |                       |              | Mean (SD)                          | Mean (SD)                          |
| <b>Observational Studies</b>        |                       |              |                                    |                                    |
| Xenocostas 2003                     | Allo                  | Day 0 to 60  | 6.8 ± 6.4                          |                                    |
| Sohl 2011                           | Allo                  | Day 0 to 60  | 5.2 (95% CI 3.7-6.7)               | 12.9 (95% CI 9.4-16.4)             |
|                                     | Cord Allo             | Day 0 to 60  | 7.8 (95% CI 6.7-8.9)               | 25.2 (95% CI 22.1-28.2)            |
| Kekre 2012                          | Auto & Allo           | Day 0 to 30  | 4.7 ± 4.5                          |                                    |
| Christou 2015                       | Auto & Allo           | Day 0 to 60  |                                    | 7.5 (95% CI 6.7-8.4)               |
| LeViellez 2015                      | Allo                  | Day 0 to 60  | Median 4                           | Median 4                           |
| Leahy 2017                          | Induction AML or Allo | Day          | 3.7                                | 4.1                                |
| Gastecki 2019                       | Allo                  | Day          | Median 19                          |                                    |
| Konuma 2019                         | Cord Allo only        | Day 0 to 30  | Median 12 (range 4-66)             |                                    |
| <b>Randomized Controlled Trials</b> |                       |              |                                    |                                    |
| Wandt 2012                          | Induction AML or Auto | Day 0 to 30  | 2.85 (95% CI 2.58-3.12) in St. Arm | 2.44 (95%CI 2.22-2.67) in St. arm  |
|                                     |                       |              | 3.14 (95% CI 2.81-3.46) in Exp arm | 1.63 (95% CI 1.42-1.83) in Exp arm |
| Stansworth 2013                     | Chemotherapy or HCT   | Day 0 to 30  | 3.0 ± 3.4 in St. arm               | 1.7 ± 2.6 in St. arm               |
|                                     |                       |              | 2.8 ± 3.1 in Exp. arm              | 3.0 ± 3.2 in Exp. arm              |
| Tay 2020                            | Auto & Allo           | Day 0 to 100 | 5.02 ± 6.13 in St. arm             |                                    |
|                                     |                       |              | 2.73 ± 4.81 in Exp. arm            |                                    |

## Association with post-HCT Outcomes

While confounded by indication, lower hemoglobin levels and/or transfusion support have been associated with various negative post-HCT outcomes. However, it remains unclear if such “risk factors” are modifiable and whether they would lead to improved post-HCT outcomes.

For instance, Xenoscostas et al. in a retrospective study of 519 consecutive patients receiving allogeneic HCT between January 1995 and March 2000 demonstrates an increased mortality during the 6-month period after HCT was associated with lower pre-HCT hemoglobin levels. Similar findings have been reported in the cord blood HCT setting where RBC transfusion  $\geq 18$  units by day 30 was significantly associated with higher overall mortality (hazard ratio, 1.86;  $P = 0.018$ )<sup>11</sup>. A more recent retrospective study by Hosoba et al. of 322 consecutive patients receiving an allogeneic bone marrow or granulocyte colony-stimulating factor-mobilized blood stem cell graft for a hematologic malignancy<sup>12</sup>. This study demonstrated that transfusion of more than the median number of RBC units (HR, 2.1; 95% CI, 1.1 to 3.7;  $P = .02$ ) were independently associated with greater risk of grade III-IV acute GVHD in a multivariable analysis model and transfusion of more than the median number (5 units within 27 days) of RBC units (HR, 1.4; 95% CI, 1.0 to 2.0;  $P = .054$ ) was associated with inferior overall survival.

Vande Vusse et al. examined the associations between platelet transfusions and idiopathic pneumonia syndrome (IPS) among 914 individuals who underwent myeloablative allogeneic HCT between 1997 and 2001. They identified 77 IPS cases (8.4%), where each additional platelet unit transfused in the prior week was associated with 16% higher IPS risk with a HR 1.16; 95%CI (1.09-1.23)<sup>13</sup>. Likewise, Christou et al. confirmed that the number of platelet transfusion events was associated with increased 100-day non relapse mortality ( $p < 0.01$ ), post-HCT length of hospital stay ( $p < 0.01$ ), need for intensive care unit admission ( $p < 0.01$ ), and number of organs affected by severe toxicity ( $p < 0.01$ )<sup>14</sup>.

## Blood Products

The Alberta Bone Marrow and Blood Cell Transplant Program relies on the Canadian Blood Services for transfusion products. The collection, testing, processing, contents and distribution of these blood products will follow the Canadian Blood Services procedures<sup>15</sup>. Blood products are issued by Transfusion Medicine, Alberta Precision Laboratories.

### Red Cell Transfusion

#### Threshold:

Different strategies have been developed for RBC transfusions. A restrictive transfusion strategy seeks to maintain a lower hemoglobin level (usually between 70 g/L to 90 g/L) with a trigger for transfusion when the hemoglobin drops below 70 g/L), whereas a liberal transfusion strategy aims to maintain a higher hemoglobin (usually between 100 g/L to 120 g/L, with a threshold for transfusion when hemoglobin drops below 100 g/L). There are very few randomized studies examining red cell transfusion thresholds in the oncology setting as summarized by a recent Cochrane review in 2017<sup>16</sup>. The review suggests that a restrictive RBC transfusion policy has little or no effect on mortality at 30 to 100 days, bleeding, or hospital stay. However, there is absence of data on health-related quality of life, arterial or venous thromboembolic events, length of intensive care admission or readmission to hospital.

A recent noninferiority randomized controlled trial in four different Canadian centers evaluated 300 patients with hematologic malignancies requiring HCT between 2011 and 2016<sup>17</sup>. Patients were randomly assigned to either a restrictive (hemoglobin [Hb] threshold  $< 70$  g/L) or liberal (Hb threshold  $< 90$  g/L) RBC transfusion strategy between day 0 and day 100 with a primary outcome of health-related quality of life measured by FACT-BMT. Clinical outcomes of transplantation-related mortality, length of hospital stay, intensive care unit admissions, acute graft-versus-host disease, Bearman toxicity score, sinusoidal obstruction syndrome, serious infections, WHO Bleeding Scale, transfusion requirements and reactions to therapy were collected. The authors demonstrate that the number of RBC units transfused was lower in the restrictive-strategy group than in the liberal-strategy group (mean, 2.73 units [standard deviation, 4.81 units] v 5.02 units [standard deviation, 6.13 units];  $P =$



.0004). After adjusting for transfusion type and baseline FACT-BMT score, the restrictive-strategy group had a higher FACT-BMT score at day 100 (difference of 1.6 points; 95% CI, -2.5 to 5.6 points), which was noninferior compared with that of the liberal-strategy group. Additionally, there were no significant differences in clinical outcomes between the transfusion strategies. Another Canadian Blood and Marrow Transplant Group trial compared red blood cell transfusion thresholds of 120 g/L in the experimental arm and 70 g/L in the control arm. The study was stopped early due to excess sinusoidal obstructive syndrome in the experimental group<sup>18</sup>. Observational/real world data would also support a 70g/L threshold strategy (<https://doi.org/10.1111/tme.12952>). In the pediatric population, there is an ongoing study that is evaluating to restrictive transfusion thresholds of 65 g/L (restrictive) vs. 80 g/L (liberal) for HCT days 0–100. (<https://doi.org/10.1111/bjh.20051>)

### **Number of Red Cell Units per Transfusion:**

With the advent of the Choosing Wisely initiatives<sup>19</sup>, many clinicians are using one-unit transfusions as opposed to using two units at a time. Observational studies support this safe strategy with decreases in the use of red cell units<sup>20-22</sup> although the results are inconsistent<sup>23</sup>. A Systematic review evaluating single vs. double unit transfusion concludes that a single unit transfusion “is associated with lower hemoglobin levels at discharge and a reduction in the total number of red cell units used per admission, with no significant difference in terms of length of hospital stay, severe bleeding risk, transfusion-related adverse events and mortality.” (<https://doi.org/10.1016/j.tmr.2024.150862>) Overall, it remains unclear if a one-unit transfusion leads to better patient satisfaction with trade-offs of improved health care utilization.

### **Duration of Storage of Red Cell Products:**

Laboratory evidence coupled with early clinical observational studies suggest that there might be concerns of using older red cell products, *i.e.* storage lesion<sup>24</sup>. However, numerous randomized studies in a varied clinical settings, summarized by systematic reviews have found no convincing evidence that the age of red blood cells have an adverse effect on patient outcomes<sup>25-29</sup>. Moreover, retrospective reviews in the general cancer<sup>30</sup> and HCT<sup>31</sup> settings would further affirm this absence of association.

### **Red Cell Transfusion- Summary:**

Taken together, the clinical evidence supports the use of a restrictive red cell transfusion threshold in patients undergoing HCT. However, it is reasonable to consider an individualized higher red cell threshold depending on clinical judgement such as the presence of symptoms of anemia. There is no evidence to support preference for “younger” red cell units over “older” units. It is reasonable and practical to provide 1-unit red cell unit per transfusion.

## **Platelet Transfusion**

### **Thresholds:**

Clinically significant bleeding related to thrombocytopenia occurs in nearly 50% of all patients undergoing HCT. Over the last 2 decades, there have been randomized controlled trials examining platelet transfusion thresholds (where the primary endpoint is WHO grade 2 or higher bleeding), summarized by the following table:

**Table 2:** RCTs examining platelet transfusion thresholds

| RCT                         | Setting                              | Patients (N) | Interventions | Key findings                                                                                                                                                                                                                                                                                                                                                                           |
|-----------------------------|--------------------------------------|--------------|---------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Heckman 1997 <sup>32</sup>  | Induction therapy for acute leukemia | 78           | <10 vs. <20   | <ol style="list-style-type: none"> <li>&lt;10 group received more platelet transfusions for bleeding.</li> <li>&lt;20 group arm received more platelet transfusions for prophylactic indications.</li> <li>No difference in RBC transfusion requirements, febrile days, days hospitalized, days thrombocytopenic, need for HLA-matched platelets, remission rate, or death.</li> </ol> |
| Rebulla 1997 <sup>33</sup>  | Induction for acute myeloid leukemia | 255          | <10 vs. <20   | <ol style="list-style-type: none"> <li>&lt;10 group had 21.5 percent fewer platelet transfusions.</li> <li>No difference in risk of major bleeding.</li> <li>No difference in survival, absence of major bleeding or length of stay</li> </ol>                                                                                                                                         |
| Zumberg 2002 <sup>34</sup>  | Autologous or Allogeneic HCT         | 159          | <10 vs. <20   | <ol style="list-style-type: none"> <li>No differences in bleeding incidence or severity.</li> <li>More transfusions were given above the assigned transfusion threshold.</li> <li>No difference in transfusion utilization.</li> </ol>                                                                                                                                                 |
| Diedrich 2005 <sup>35</sup> | Allogeneic HCT                       | 166          | <10 vs. <30   | <ol style="list-style-type: none"> <li>&lt;10 group had fewer platelet transfusions.</li> <li>No difference in bleeding, bacteremia, engraftment, GVHD, hospital stay, death, and survival.</li> <li>No difference in RBC transfusions.</li> </ol>                                                                                                                                     |

The results from these studies have informed the care of patients with hematologic malignancies receiving high dose chemotherapy, where the current standard practice is to transfuse platelets prophylactically when the daily platelet count is  $< 10 \times 10^9/L$  to prevent bleeding<sup>3,6,36-38</sup>.

More recently, the 2025 AABB guideline proposes (2025;334;(7):606-617.

doi:10.1001/jama.2025.7529):

- Allogeneic HCT: Strongly recommends a prophylactic threshold of  $< 10 \times 10^9/L$  for non-bleeding patients.
- Autologous HCT: Issued a conditional recommendation against routine prophylactic platelet transfusion for non-bleeding adults, suggesting a therapeutic-only (transfuse only if bleeding) or more restrictive approach may be appropriate.

**Platelet Dose:**

Additionally, there have been 5 RCTs evaluating the efficacy of different platelet doses<sup>39-43</sup> summarized by a Cochrane review<sup>44</sup>. For instance, the more recent and adequately powered study by Slichter et al. randomized (n=1,272) patients undergoing hematopoietic stem-cell transplantation or chemotherapy for hematologic cancers or solid tumors to receive prophylactic platelet transfusions at a low dose, a standard dose, or a high dose ( $1.1 \times 10^{11}$ ,  $2.2 \times 10^{11}$ , or  $4.4 \times 10^{11}$ ) platelets per square meter of body-surface area, respectively<sup>43</sup>. They demonstrate that low doses of platelets administered as a prophylactic transfusion led to fewer platelets transfused per patient, but an increased number of transfusions given. At doses between  $1.1 \times 10^{11}$  and  $4.4 \times 10^{11}$  platelets per square meter, the number of platelets in the prophylactic transfusion had no effect on the incidence of bleeding.

Taken together, there is no evidence to suggest that a low-dose platelet transfusion policy is associated with an increased bleeding risk compared to a standard-dose or high-dose policy, or that a high-dose platelet transfusion policy is associated with a decreased risk of bleeding when compared to a standard-dose policy<sup>44</sup>. However, a low-dose platelet transfusion strategy leads to an increased number of transfusion episodes compared to a standard-dose strategy. A high-dose platelet transfusion strategy does not decrease the number of transfusion episodes per participant compared to a standard-dose regimen, and it may increase the number of transfusion-related adverse events.

**Duration of Storage of Platelets:**

Platelets are usually stored for up to 5 days prior to transfusion, although in some blood services the storage period is extended to 7 days. During storage, changes occur in both platelets and storage medium, which may lead to platelet activation and dysfunction. There have been several observational studies, mostly in the critical care setting that evaluated the potential impact of platelet storage and clinical outcomes. A recent systematic review of such studies suggests that storage time does not appear to be associated with clinical outcomes, including bleeding, sepsis or mortality<sup>27</sup>. However, the freshest PLTs (less than 3 days) were associated with a better CCI, although there was no impact on bleeding events. Taken together, the duration of storage of transfused platelets likely has no appreciable impact on post-HCT outcomes.

**Platelet Transfusion Alternatives:**

However, the true benefit of prophylactic platelet transfusions in the autologous HCT setting is unclear; subgroup analyses from 2 recent studies suggest that a therapeutic strategy (transfusing platelets only to treat bleeding) may be as effective as a prophylactic transfusion strategy<sup>45,46</sup>. There has been interest in the use of prophylactic TXA to prevent bleeding in patients with chemotherapy-related hypoproliferative thrombocytopenia<sup>47,48</sup>. This includes 3 ongoing RCTs, with 2 examining the addition of prophylactic TXA to prophylactic platelet transfusions in patients with in patients with hematological malignancies with severe thrombocytopenia (NCT03136445<sup>49</sup> and NCT02578901) and another examining prophylactic tranexamic acid with prophylactic platelets in patients undergoing autologous HCT (NCT04448184)<sup>50</sup>.

### Platelet Transfusion-Summary:

Taken together, the clinical evidence supports the use of  $< 10 \times 10^9/L$  as a platelet transfusion threshold. However, it is reasonable to consider an individualized higher platelet threshold and/or dose, depending on clinical judgement such as the presence of persistent bleeding. There is a paucity of high-quality evidence to routinely use therapeutic over prophylactic platelet transfusions. There is no evidence to support preference for “younger” platelet units over “older” units.

### Platelet Refractoriness:

Platelet transfusion refractoriness is defined as the repeated failure to achieve satisfactory responses to platelet transfusions from random donor<sup>51</sup>. There have been various formulae that have been proposed to determine platelet refractoriness<sup>52</sup>. However, it is argued that such calculations are not clinically useful. Instead, a pragmatic definition of failure of the immediate post-transfusion (10 to 60 minutes) platelet increment to exceed the transfusion trigger or a rise of less than  $10 \times 10^9/L$  20 to 24 hours after transfusion with unsatisfactory responses to two or more transfusions<sup>53</sup>. The following table summarizes some potential causes for platelet refractoriness:

**Table 3:** Potential causes for platelet refractoriness

| Immune                                                                  | Non-Immune                                              |
|-------------------------------------------------------------------------|---------------------------------------------------------|
| Platelet alloantibodies:<br>HLA, ABO, HPA                               | Infection                                               |
| Other antibodies:<br>Platelet autoantibodies, Drug dependent antibodies | Fever                                                   |
| Immune complexes                                                        | Anti-microbials (vancomycin, fluoroquinolones, Ampho B) |
|                                                                         | Graft versus host Disease                               |
|                                                                         | Veno-occlusive Disease                                  |
|                                                                         | Splenomegaly                                            |
|                                                                         | Disseminated Intravascular Coagulation                  |

One recent observation study suggests that splenomegaly (odds ratio [OR] = 26.98,  $p < .001$ ) and JAK mutation (OR = 17.32,  $p = .024$ ) were independent risk factors for PTR and was also associated with worse overall survival (hazard ratio = 2.794, 95% confidence interval = 1.083–7.207,  $p = .034$ ) (<https://doi.org/10.1080/09537104.2023.2229905>)

While universal leukoreduction has decreased the incidence of platelet refractoriness from HLA antibodies<sup>54</sup>, it does not address other causes. It is likely that there are multiple competing etiologies for platelet refractoriness. Moreover, it would be impossible to determine their relative contributions to platelet refractoriness. Beyond, optimizing specific and potential etiologies, it is practical to consider the following measures where the risk of bleeding is considered high or unacceptable. These measures should be continued until the risk of bleeding is deemed low.

1. Prophylactic HLA matched platelets

- a. Prophylactic HLA matched platelets can be considered in patients where the platelet counts are expected to 'drop" given the underlying clinical scenario and care trajectory. It may be also reasonable to otherwise withhold its use. The use of clinical judgement is warranted in this circumstance.
  - b. Antibodies against HLA-A and –B antigens are the only clinically relevant HLA antibodies that cause platelet refractoriness.
2. Prophylactic Tranexamic acid 1gram TID PO/IV

The following are additional considerations:

1. It is reasonable to provide both prophylactic standard daily prophylactic non-HLA matched platelets and tranexamic acid measures in the absence of a satisfactory response with HLA matched platelets or if HLA matched platelets are not available.
2. In cases where the risk of bleeding is deemed low, it is reasonable to solely provide prophylactic tranexamic acid.
3. There are advocates for platelet cross-matching, but it is not routinely available in Canada and not recommended over HLA matched platelets.
4. Intravenous immunoglobulin is not effective in the management of platelet refractoriness.

## Cellular Therapy

A form of immunotherapy known as chimeric antigen receptor (CAR) T cell treatment was created from adoptive T cell transfer, which normally employs patients' own immune cells to treat cancer. CAR-T cells are equipped with certain antibodies that enable them to identify antigens in self-tumor cells and cause cytotoxic effects. CAR-T cell therapy has had exceptional success treating hematologic malignancies in recent years.

**Table 4:** Randomized control trials showing the effectiveness of Car T cell treatment in multiple myeloma, lymphoma and leukemia.

| Disease | Ref            | Year of Pub | Sample Size                          | Product                              | Anemia                                       | Thrombocytopenia                                                                                              | Neutropenia                                                                              | Leukopenia                                   | Comments                                                                                                                                            |
|---------|----------------|-------------|--------------------------------------|--------------------------------------|----------------------------------------------|---------------------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------|----------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------|
| Myeloma | Berdeja et al. | 2021        | 113 (enrolled); 97 (infused cohort)  | Ciltacabtagene autoleucel (Carvykti) | Grade 3-4: 68% after day 1 of cilta-cel      | Grade 3-4: after day 1 of cilta-cel →reduced to grade 2 or lower by day 30 in (59%)                           | Grade 3-4: 95% after day 1 of cilta-cel →reduced to grade 2 or lower by day 30 in (70%)  | Grade 3-4: 61% after day 1 of cilta-cel      | Pts received supportive measures to treat cytokine release syndrome or associated symptoms, most commonly tocilizumab, corticosteroids and anakinra |
|         | Munshi et al.  | 2021        | 140 (enrolled); 129 (infused cohort) | Idecabtagene vicleucel (Abecma)      | Grade 3-4: 60% within 8 weeks after infusion | Grade 3-4: 52% within 8 weeks after infusion →recovery to grade 2 or lower occurred at a median of 2.1 months | Grade 3-4: 89% within 8 weeks after infusion →recovery to Grade 2 or lower occurred at a | Grade 3-4: 39% within 8 weeks after infusion | Pts received supportive measures to treat cytokine release syndrome or associated symptoms, most commonly                                           |



|          |                 |      |                                          |                                      |                                                                                                       |                                                                                                                                                    |                                                                                                                                                       |                                                     |                                                                                                                                      |
|----------|-----------------|------|------------------------------------------|--------------------------------------|-------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------|
|          |                 |      |                                          |                                      |                                                                                                       |                                                                                                                                                    | median of 1.9 months                                                                                                                                  |                                                     | tocilizumab and corticosteroids                                                                                                      |
| Lymphoma | Neelapu et al.  | 2017 | 111 (total cohort); 101 (infused cohort) | Axicabtagene cilio-leucel (Yescarta) | Grade 3-4: 43% (unspecified duration) →at 3 months, grade 3 or higher was reported (3%)               | Grade 3-4: 38% (unspecified duration) →at 3 months, grade 3 or higher was reported (7%)                                                            | Grade 3-4: 78% (unspecified duration) →at 3 months, grade 3 or higher was reported (11%)                                                              | Grade 3-4: 29% (unspecified duration)               | Pts received tocilizumab and glucocorticoids for supportive measures to treat cytokine release syndrome, neurologic events, or both. |
|          | Schuster et al. | 2022 | 167 (total cohort); 115 (infused cohort) | Tisagenlecleucel (Kymriah)           | Grade 3-4: 39% after infusion (unspecified duration)                                                  | Grade 3-4: 28% after infusion (unspecified duration) →unresolved grade 3-4 at day 28 in 41% of pts →unresolved grade 3-4 at 3 months in 38% of pts | Grade 3-4: 33% after infusion (unspecified duration) →unresolved grade 3-4 at day 28 in 24% of pts →At 3 months, no patients had unresolved grade 3-4 | Grade 3-4: 31% after infusion (unspecific duration) | 17% of pts received systemic anti-cytokine therapy, 16% received tocilizumab, and 10% received corticosteroids                       |
|          | Abramson et al. | 2020 | 344 (total cohort); 269 (infused cohort) | Lisocabtagene maraleucel (breynzi)   | Grade 3-4: 37% after 29 days →after 90 days recovery to grade 2 occurred in 82%                       | Grade 3-4: 27% after 29 days →after 90 days, recovery to grade 2 occurred in 62 %                                                                  | Grade 3-4: 60% after 29 days →After 90 days, recovery to grade 2 occurred in 84%                                                                      | Grade 3-4: 14% after 29 days                        | Pts received tocilizumab, corticosteroids, vasopressors, siltuximab, and anakinra for supportive measure of cytokine release         |
| Leukemia | Maude et al.    | 2018 | 75 (infused cohort)                      | Tisacel (Kymriah)                    | Grade 3-4: 4% (unspecified duration)                                                                  | Grade 3-4: 7% (unspecified duration) →By last assessment, 71% of pts has Grade 2 or lower                                                          | Grade 3-4: 11% (unspecified duration) →By last assessment, 80% of patients has Grade 2 or lower                                                       | Grade 3-4: 9% (unspecified duration)                | Tocilizumab was given to pts experiencing cytokine release syndrome                                                                  |
|          | Shah et al.     | 2021 | 54 (enrolled); 45 (infused cohort)       | Brexu-cel (Tecartus)                 | Grade 3-4: 49% with median duration of 9 days →At day 30, grade 3 or higher was reported by 7% of pts | Grade 3-4: 30% with median duration of 9 days →At day 30, grade 3 or higher was reported by 18% of pts                                             | Grade 3-4: 27% with median duration of 9 days →At day 30, grade 3 or higher was reported by 25% of pts                                                | Grade 3-4: 23% with median duration of 9 days       | Steroids and tocilizumab were used fir cytokine release syndrome and neurologic events                                               |

**Table 5:**

| Lineage             | Grade 1                        | Grade 2                       | Grade 3                       | Grade 4                       |
|---------------------|--------------------------------|-------------------------------|-------------------------------|-------------------------------|
| Neutrophils         | <LLN to 1,500/mm <sup>3</sup>  | 1,000-1,500/mm <sup>3</sup>   | 500-1,000/mm <sup>3</sup>     | <500/mm <sup>3</sup>          |
| Platelets           | <LLN to 75,000/mm <sup>3</sup> | 50,000-75,000/mm <sup>3</sup> | 25,000-50,000/mm <sup>3</sup> | <25,000/mm <sup>3</sup>       |
| Hemoglobin          | <LLN to 10 g/dL                | 8.0-10.0 g/dL                 | <8.0 g/dL                     | Life-threatening consequences |
| Lymphocytes (total) | <LLN to 800/mm <sup>3</sup>    | 500-800/mm <sup>3</sup>       | 200-500/mm <sup>3</sup>       | <200/mm <sup>3</sup>          |

Neutropenia, thrombocytopenia, anemia, and lymphopenia were determined from the complete blood count after chemotherapy, and the lowest count was used for calculating grade of toxicity. All patients with sustained fever of >100.4°F in the midst of chemotherapy-induced grade 4 neutropenia received a first course of IV antibiotics in hospital. Taken from National Cancer Institute Common terminology Criteria for Adverse Events (NCI CTCAE, version 3.0). LLN, lower limit of normal.

**Table 6:** Time-frame for hematopoietic toxicity from the day of CAR T-cell infusion as early (<30days), short-term (30-90 days), and prolonged

|                              | Early (<30 days)                                                                                                                                                          | Short-term (30-90 days)                                                                | Prolonged (>90 days)                                                |
|------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------|---------------------------------------------------------------------|
| Possible etiologies          | Lymphodepletion chemotherapy                                                                                                                                              | Lymphodepletion chemotherapy                                                           | Disease relapse                                                     |
|                              | CRS/ICANS<br>CAR-HLH<br>Infections<br>Antibody-mediated autoimmune<br>Cytopenias                                                                                          | CAR-HLH<br>Disease relapse<br>Infections<br>Antibody-mediated autoimmune<br>Cytopenias | MDS                                                                 |
| Possible factors associated  | Grade 3-4 CRS/ICANS                                                                                                                                                       | CAR construct                                                                          |                                                                     |
|                              | CAR construct<br>Peak CRP/ferritin<br>Baseline cytopenia<br>Number of prior lines of therapy<br>Prior lines of therapy<br>Prior allo-MBT<br>Marrow tumor burden           |                                                                                        |                                                                     |
| Consideration for management | Evaluation for CAR-HLH including ferritin, triglycerides, coagulopathy, bone marrow biopsy to assess for hemophagocytosis. Treatment with tocilizumab, anakinra ±steroids | Supportive care with transfusions, growth factors, TPO agonists                        | Marrow biopsy for primary disease or another malignancy such as MDS |
|                              | Evaluation for hemolysis, immune mediated thrombocytopenia                                                                                                                | Disease evaluation for relapse                                                         |                                                                     |
|                              | Evaluation and treatment of infections                                                                                                                                    | Marrow biopsy for disease or another malignancy such as MSD                            |                                                                     |

## Prevention of Transfusion- Associated GVHD

While recipient anti-donor responses are usually able to eliminate donor leukocytes, settings in which the recipient anti-donor responses are impaired permits unabated donor anti-recipient responses (which can occur in the HCT setting), resulting in transfusion-associated graft-versus-host disease (TA-GVHD)<sup>55</sup>. There are 2 standard complimentary approaches to reduce the risk of TA-GVHD: 1) Leukoreduction, and 2) Irradiation of the blood product.

The ability to deplete leukocytes from the blood product reduces the incidence of TA-GVHD<sup>56,57</sup>. Blood products from the Canadian Blood Services are universally leukodepleted, a practice that is supported by randomized trial data<sup>54,58,59</sup>. However, leukodepletion may not be fully protective<sup>60</sup>. The results of *in vitro* studies led to the adoption of a dose of 25–30 Gy γ-irradiation as a standard for the inactivation of T lymphocytes in blood products<sup>61,62</sup>. This led to routine irradiation of blood products, especially in settings in where patients are at risk for developing TA-GVHD. Reports from Japan (population at higher risk of TA-GVHD) indicate that no further cases of TA-GVHD were detected once universal irradiation was implemented<sup>63,64</sup>.

It should be kept in mind that prolonged storage of pre-irradiated red blood cells has been associated with high potassium levels, in vitro hemolysis and decreased post-transfusion recovery<sup>65</sup>. Irradiation

of red blood cell products should occur as near as possible to the time of transfusion, and no longer than 14 days prior to transfusion. This is particularly important in infants, where life threatening hyperkalemia can occur following transfusion of irradiated RBC. In patients at risk of hyperkalemia, it is recommended red cells be transfused within 24 hours of irradiation. If freshly irradiated RBC are not available, the product should undergo centrifugation and supernatant plasma removal prior to transfusion.

There are other laboratory techniques such as psoralen/UVA light treatment (primary purpose of pathogen reduction) that can inactivate T cells in blood products and potentially reduce the risk of transfusion-associated graft-versus-host disease (TA-GVHD), but this has not been routinely adopted by Canadian Blood Services. Importantly, pathogen reduced platelets increase the risk of platelet refractoriness and the platelet transfusion requirement<sup>66</sup>.

### Prevention of CMV Transmission

Cytomegalovirus (CMV) infection continues to be a serious complication following HSCT<sup>67,68</sup>. Most CMV infections may be due to a reactivation of the virus from a previous infection rather than due to the acquisition of a new strain<sup>69</sup>. However, CMV antibody-negative persons are at risk for developing a transfusion-transmitted *de novo* CMV infection. There are 2 standard complimentary approaches to reduce this risk: 1) Use of CMV-antibody negative blood, and 2) leukoreduced components.

As discussed, blood products from the Canadian Blood Services undergo universal leukoreduction. This practice is further supported by studies demonstrating that leukoreduced components are as effective as antibody-negative components in the prevention of transfusion-transmitted CMV infection<sup>70-73</sup>. However, early consensus guidelines supported the use of both leukoreduction and provision of CMV-antibody negative blood in at risk population (belt and suspenders approach)<sup>74,75</sup>.

However, this practice has been challenged<sup>71,76</sup>. For instance, a single Canadian HCT institution before-after study reviewed 186 patients who were CMV negative and received an allogeneic HCT from a CMV-negative donor between October 1, 1999 and June 30, 2012<sup>77</sup>. Of these, 89 patients received an HCT before January 2007, during the time when patients received leukoreduced and CMV-negative blood products. Seventy-seven patients received an HCT after this time, receiving only leukoreduced blood products. CMV viremia was detected in 3 patients who received CMV-negative leukoreduced blood products (3.37%) and in 1 patient who received only leukoreduced blood products (1.30%,  $P = .6244$ ). Of the patients who developed CMV viremia, 2 developed suspected CMV disease. Both of these patients were transfused with CMV-negative blood products.

This suggests that in the era of universal leukoreduction of blood products, that testing for CMV-negative blood products is not needed for HCT recipients. Indeed, the Canadian National Advisory Committee on Blood and Blood Products (NAC) subcommittee has deemed that CMV-safe

leukoreduced cellular blood products are equivalent in safety to CMV-seronegative and leukoreduced blood products for transfusion in all patient populations except for intrauterine transfusion.

### **Transfusion in ABO- or RhD-Incompatible HCT**

Please refer to Chapter entitled ABO Incompatible Graft and Recipient in the Alberta Bone Marrow and Blood Cell Transplant Program: BMT Standard Practice Manual.

### **Growth Factor Support**

#### **Granulocyte Colony-Stimulating Factor:**

The use of G-CSFs post-HCT is supported by decreased time to engraftment ranging between 1–6 days, with some studies demonstrating savings in duration of hospitalization, infections and survival<sup>78</sup>. The timing of administration of G-CSF has been the subject of investigation – early or delayed approaches. Such studies have not demonstrated any clear disadvantage of delaying G-CSF for up to 10 days post-HCT in the autologous setting<sup>79-92</sup>. G-CSF has been used in the allogeneic setting, but there is theoretical concern that T cell reconstitution may be impaired by G-CSF, and GVHD may be induced or worsen with its use. Although G-CSF is often administered post-HCT to accelerate neutrophil recovery, there appears to be no long-term benefit or disadvantage of giving G-CSF after HCT to promote hematopoietic recovery<sup>93</sup>.

Taken together, it is reasonable to provide routine post-autologous HCT G-CSF starting on Day 7 (until ANC >0.5) to accelerate neutrophil recovery which in principle may improve infectious complications and assist with earlier hospital discharge. We do not recommend the routine use of G-CSF in the allogeneic setting. This recommendation may not apply to most pediatric non-malignant conditions.

#### **Erythropoiesis-stimulating agents:**

The use of erythropoiesis-stimulating agents to manage anemia raises hemoglobin levels, reduces the need for RBC transfusions, but increases the risk of thromboembolic events<sup>94</sup>. In the setting of HCT, there have been 6 randomized studies addressing the utility of erythropoietin. The most recent study, which might inform current practice, was conducted by Jaspers et. al in 2014<sup>95</sup>. The authors randomized 131 patients between no treatment (control arm) or erythropoietin at 500 U/kg per week. They demonstrate that erythropoietin results in a higher proportion (63.1%) of hemoglobin  $\geq 13$  g/dL before Day 126 as compared with 8.1% in the control arm. Additionally, Hb levels were higher and transfusion requirements decreased in the erythropoietin arm. There was no difference in rates of thromboembolic events or other complications between the 2 arms. There was no impact on long term survival<sup>96</sup>.

The following table from Christou et al. summarizes and provides a scoping review of the 6 randomized trials evaluating the use of erythropoietin post-HCT<sup>1</sup>.

**Table 7: 6 RCTs evaluating the use of erythropoietin**

| Reference          | Year of Publication | Sample Size | EPO treatment arm                                                            | Control arm  | Hb level | Bleeding | Days to PLT engraftment | Days to neutrophil engraftment | RBC utilization | PLT utilization | Hospital LOS | GVHD | Infection | Transfusion reaction | Overall survival |
|--------------------|---------------------|-------------|------------------------------------------------------------------------------|--------------|----------|----------|-------------------------|--------------------------------|-----------------|-----------------|--------------|------|-----------|----------------------|------------------|
| Steegmann et al.   | 1992                | 27          | 100 U/kg/day IV Day 0-Day 7 then 150 U/kg/day IV Day 7-Day 30                | No injection | ↔        |          | ↓                       | ↔                              | ↓               | ↓               |              |      |           |                      | ↔                |
| Link et al.        | 1994                | 329         | 150 U/kg/day IV continuous infusion Day 0-Day 42 or transfusion independence | Placebo      |          | ↔        | ↔                       | ↔                              | ↓               | ↔               |              | ↔    |           |                      |                  |
| Klaesson et al.    | 1994                | 50          | 200 U/kg/day IV Day 0-Day 28, then 2X/week IV Day 29-Day 48                  | Placebo      | ↑        | ↔        | ↔                       | ↔                              | ↓               | ↔               | ↔            |      | ↔         |                      | ↔                |
| Biggs et al.       | 1995                | 91          | 300 U/kg/day IV 3x/week Day 0- Day 42                                        | No injection | ↑        |          | ↔                       | ↔                              | ↔               | ↔               | ↔            | ↔    | ↔         |                      | ↔                |
| Vanstraelen et al. | 2006                | 60          | 500 U/kg/day SC weekly starting at Day 0 or Day 30                           | No injection | ↑        |          | ↔                       | ↔                              | ↓               |                 |              |      |           |                      |                  |
| Jaspers et al.     | 2014                | 131         | 500 U/kg SC weekly starting at Day 0 or Day 28                               | No injection | ↑        |          | ↔                       | ↔                              | ↓               |                 |              | ↔    | ↔         |                      | ↔                |

\* arrows indicate whether any significant increase (↑), decrease (↓), or no change (↔) was reported for the outcomes listed in the table. LOS=Length of stay; SC= subcutaneously

Taken together, the use of erythropoietin post-HCT can improve hemoglobin levels and assist with red cell utilization. However, the absence of quality of life data from the available trials tempers its use. Moreover, enthusiasm for the use of erythropoietin in the HCT setting has waned, especially with the recognition of serious adverse events in several patient populations<sup>97,98</sup>.

### Thrombopoietin receptor agonists:

Prolonged thrombocytopenia after HCT is a strong risk factor for transplantation-related mortality, and efforts to improve platelets may lead to improve post-HCT outcomes and decrease platelet utilization<sup>99,100</sup>. The literature categorizes these patients into 2 types—prolonged isolated thrombocytopenia (PIT) and secondary failure of platelet recovery (SFPR)—according to the timeline of presentation.

Prolonged isolated thrombocytopenia (PIT) can be attributed to:

1. Delayed platelet engraftment, often defined as persistent severe thrombocytopenia with a platelet count  $<20 \times 10^9/L$  beyond 35 days after HSCT,
2. Primary graft failure, defined as failure to achieve initial engraftment by day +28 post-HSCT or
3. Poor graft function, usually defined as persistent thrombocytopenia (platelet count  $\leq 20 \times 10^9/L$ ) with neutropenia (absolute neutrophil count ANC  $\leq 5 \times 10^9/L$ ) and/or hemoglobin  $<7$  g/dL for at least 3 consecutive days by 28 days after HSCT, with transfusion dependence associated with hypoplastic-aplastic bone marrow and complete donor chimerism without concurrent GVHD or disease relapse.

Secondary failure of platelet recovery (SFPR) refers to thrombocytopenia that develops after initial platelet engraftment and is not due to graft rejection or relapse. SFPR is defined as a decline in platelet count of  $<20 \times 10^9/L$  for 7 consecutive days or requiring transfusion support after achieving a sustained platelet count  $\geq 50 \times 10^9/L$  without transfusion for 7 consecutive days after HCT<sup>101</sup>.

Thrombopoietin receptor agonists (TPOs) are novel treatments for patients with chronic ITP aimed at increasing platelet production through interactions with the TPO receptor on megakaryocytes<sup>102</sup>. Beyond potential improvements in platelet counts, there is laboratory and clinical evidence to support the use of TPO to overcome depletion of HSCs and progenitor cells in aplastic anemia<sup>103</sup>. Invariably, these observations and experiences have led to the use of TPOs in patients with either persistent thrombocytopenia or general hypoplasia post-HCT<sup>104,105</sup>.

A recent systematic review comprising of 25 reports (case series and reports) suggests that patients with prolonged post-HCT thrombocytopenia may respond to both eltrombopag (overall response rate [ORR], 70%) and romiplostim (ORR, 82%), with no evidence of serious adverse effects<sup>106</sup>. The authors note that most of the studies reported initiating treatment at a lower dosage range and escalating the dosage based on the response to treatment. Patients were treated for variable durations, ranging from 2 weeks to 1 year, depending on the severity of thrombocytopenia and the response to treatment.

Taken together, it is reasonable to initiate the use of eltrombopag starting at 25 to 50mg, increasing to 150mg. Start at a maximum dose of 25 mg daily in patients of East Asian ethnicity (Pediatric dosing: initiate at 25 mg/day, increase weekly by 25 mg to a maximum of 75 mg daily based on platelet count. Not licensed for patients under 1 year of age.) for prolonged isolated thrombocytopenia (PIT) or secondary failure of platelet recovery. It is reasonable to gradually wean off TPO over weeks once stable hematic parameters are achieved.



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# Other Topics



# Conditioning for Allogeneic HCT

Presented by: Jan Storek

## Summary

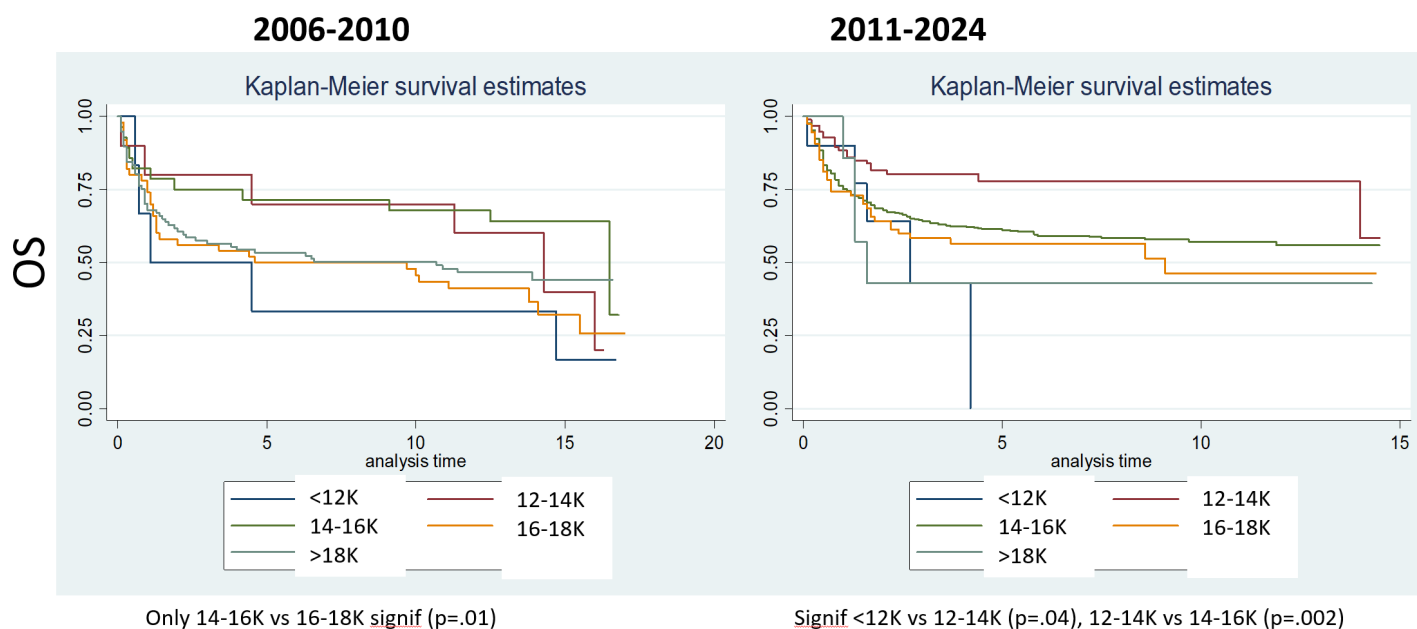
- Ideal conditioning is unknown. A uniform conditioning is a prerequisite for an academic HCT program wishing to produce consistent results and evaluate the effects of practice changes. Since the 2000's, the uniform conditioning in Alberta has been fludarabine (200-250 mg/m<sup>2</sup>) + busulfan (targeting 15000 uM.min till 2025, 13500 uM.min since 2026) + TBI (4 Gy) (FluBuTBI). This is a myeloablative conditioning (MAC). There is controversy whether similar survival can be achieved with a reduced intensity conditioning (RIC). In Alberta, RIC consisting of fludarabine (120 mg/m<sup>2</sup>) + treosulfan (30 g/m<sup>2</sup>) (FluTreo) has been used since the early 2020's for patients with a high risk of transplant-related mortality (TRM) and patients with lymphoma/CLL.
- FluBuTBI should be used for myeloid malignancies and acute lymphoblastic leukemia (ALL) up to age 70.
- FluTreo should be used for lymphoma/CLL irrespective of age, and for myeloid malignancies/ALL if age >70 or if high TRM risk, particularly due to poor hepato/pulmonary function.
- For alloHCT for non-malignant diseases and for autologous HCT, conditioning regimens are specified in disease-specific chapters, except in 2025-2027 when they may appear in this chapter.

## Introduction

- Despite intense research since the 1970's/1980's, the best conditioning is unknown.<sup>1</sup>
- Comparisons of high vs low intensity conditioning, including randomized prospective studies,<sup>2-5</sup> have yielded conflicting results.
  - Most of the randomized studies were flawed, eg, by comparing different drugs rather than different doses of the same drug(s), differences in GVHD prophylaxis, or by insufficient accrual or short follow up. The best study (largest N, no major flaws) by Scott et al showed superiority of MAC over RIC.<sup>6</sup> This may apply only to MRD positive patients, as MRD negative patients had similar outcomes with MAC and RIC.<sup>7</sup>
  - Studying only MDS and/or AML has been a limitation of the randomized and most retrospective studies.
- A common theme emerged from the multiple randomized and retrospective studies: The higher the intensity of conditioning, the higher NRM and the lower the incidence of relapse. This is exemplified by a large EBMT registry study that categorized conditioning regimens as low, intermediate, or high intensity. With increasing intensity, NRM increased, relapse decreased, and OS was similar in all 3 groups.<sup>8</sup>

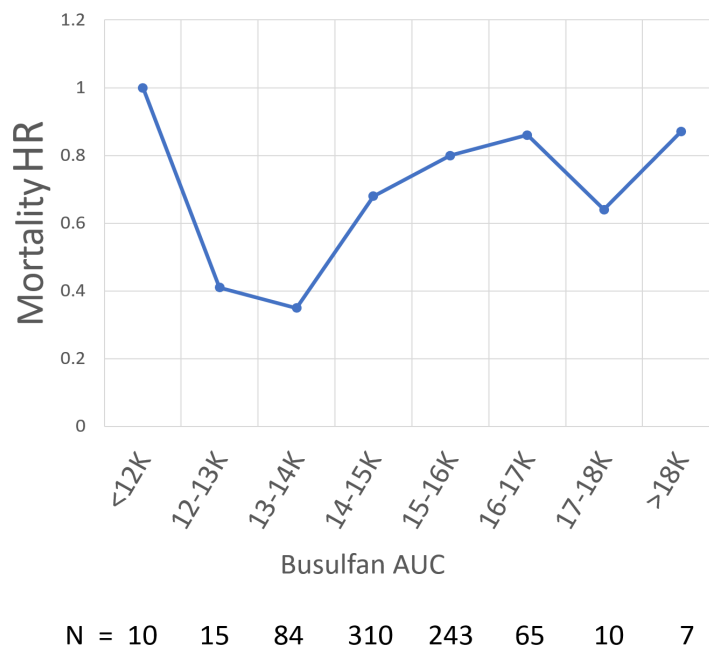
- Studies of busulfan AUC have shown that high AUC is associated with high mortality due to high NRM, low AUC with high mortality due to primarily relapse, and intermediate AUC with the lowest mortality (best OS).<sup>9</sup>
- Given the above controversies and inconsistencies, in ~2010 our center chose one main MAC, to be used in most patients, and in ~2021 one RIC, to be used in patients with a high risk of TRM and since ~2023 in all lymphoma/CLL patients (see Lymphoma chapter for rationale).
  - Our main MAC (FluBuTBI) consisted of fludarabine (250 mg/m<sup>2</sup> for matched related and unrelated HCT and 200 mg/m<sup>2</sup> for haploidentical HCT because haploidentical HCTs were started as part of a multi-center trial that mandated 200 mg/m<sup>2</sup>) + busulfan (in 2006-2010 avoiding total AUC of >24,000, in 2011-2025 targeting 15000 uM.min, the latter achieved by median total dose of 9.5 mg/kg) + TBI (4 Gy in two fractions). This regimen has been used for acute leukemia since the early 2000's and for most hematologic malignancies since ~2010. This regimen will undergo two adjustments, starting in 2026:
    - Fludarabine dose unification to 200 mg/m<sup>2</sup>. The reasons are simplicity/uniformity, the fact that 250 mg/m<sup>2</sup> is a much higher dose than used in published conditioning regimens (90-190 mg/m<sup>2</sup>), that based on an EBMT registry study doses <135 mg/m<sup>2</sup> were associated with a trend toward better OS compared to higher doses,<sup>10</sup> and to avoid one day of hospitalization for matched related/unrelated HCT recipients (the 250 dose was given as 50 mg/m<sup>2</sup> on day -6, -5, -4, -3, and -2, the 200 dose is given as 50 mg/m<sup>2</sup> on day -5, -4, -3, and -2, busulfan given on days -5, -4, -3, and -2, and TBI on day -1).
    - Busulfan target AUC reduction to 13500 uM.min. This is based on a retrospective analysis of 745 patients with myeloid malignancies or ALL who in 2011-2024 underwent PBSCT from matched related or 7-8/8 unrelated donors using Flu 250 mg/m<sup>2</sup> + Bu targeted to a total AUC of 15000 uM.min + TBI 4 Gy together with Thymoglobulin 4.5 mg/kg + MTX + CNi GVHD prophylaxis (Storek and Kangaroo, unpublished). The best survival / the lowest mortality hazard ratio (HR) was associated with total AUC of 13000-14000 uM.min (Figure on next page).

# Ideal Bu AUC in 2006-2010 vs 2011-2024



Conclusion: Ideal target AUC in 2006-2010 was 14-16K, in 2011-2024 was 12-14K

## Mortality in 2011-2024 by AUC



- The busulfan target range is 13500  $\pm$ 20%, ie, 10800-16200 uM.min. The rationale for  $\pm$ 20% rather than a narrower range is that
  1. We need to have a sufficient range of AUCs to be able to detect a potential shift of the ideal AUC in the future (as occurred between 2006-2010, when the ideal AUC was 15000, and 2011-2024, when the ideal AUC was 13500).
  2. There may be a concern that some patient's AUC will be <12000, which appears associated with high mortality, however, that is based on only 10 patients – confirmation is needed.
  3. The logistics of targeting a narrower range would be more challenging. For example, with  $\pm$ 10% target a second adjustment of Bu dose (based on day -5 PK) would be needed in ~14% patients (compared to only ~3% patients with  $\pm$ 20% target). This would make the dose adjustment based on the day -5 PK imperative, which would make the determination of Bu levels by PK lab prior to day -3 imperative, which might be difficult in case of holidays or personnel shortage in the PK lab.
- RIC is thought to give better outcomes for patients at a high risk of TRM, eg, patients with hepato/pulmonary compromise (who may not tolerate myeloablative busulfan) and elderly (>70-yo?) patients.
  - The age cutoff of 70 is arbitrary, as no comparisons of high vs low intensity conditioning for elderly patients exist. The best MAC vs RIC randomized study (Scott et al<sup>6</sup>) had an upper age limit of 65. Thus, we could take 65 as the age limit for RIC. On the other hand, per our analysis of >65-y-o patients who received our conventional MAC in 2011-2024 and achieved busulfan AUC of 12000-14000 uM.min (N=18, median age 67, range 66-72, median FU 3.4 y), 14/18 (78%) survived, which is similar to <65-y-o patients (who achieved Bu AUC of 12000-14000).
  - Our main RIC (FluTreo) consists of fludarabine and treosulfan. The reason is that fludarabine (150 mg/m<sup>2</sup>) + treosulfan (30 g/m<sup>2</sup>) was superior in OS to fludarabine (150 mg/m<sup>2</sup>) + busulfan (6.4 mg/kg) in a large phase 3 study. The superior OS was due to lower NRM, whereas relapse incidence was similar.<sup>11</sup> This applied not only to the whole cohort (age 18-70) but also to older patients (age 50-70, median 60); 3-y OS was 67% in the treosulfan arm vs 50% in the busulfan arm.<sup>12</sup>
    - In 2026, we changed fludarabine from 30 mg/m<sup>2</sup> on days -6, -5, -4, -3, and -2 (total 150 mg/m<sup>2</sup>) to 30 mg/m<sup>2</sup> on days -5, -4, -3, and -2 (total 120 mg/m<sup>2</sup>). The rationale was to save one day of hospitalization. The outcome should not be compromised as in an EBMT registry study of fludarabine dose in various RIC regimens, patients with a total dose of <135 mg/m<sup>2</sup> had a trend toward better OS than patients with higher doses.<sup>10</sup>

## Drugs Used for Conditioning

## Busulfan

Busulfan is an alkylating agent believed to act through alkylation and cross-linking of DNA strands. Busulfan is cell-cycle non-specific and induces prolonged cytopenias when used alone or in combination with other agents. The liver converts busulfan to inactive metabolites, which are then excreted in the urine. Very little busulfan is excreted unchanged.

Busulfan is available as oral 2 mg tablets and as a 6 mg/mL solution for IV administration. When used in conditioning, the IV form is preferred due to unpredictable absorption and metabolism of the oral form. When busulfan is administered for myeloablative conditioning within the ABMTP together with total body irradiation (TBI), we target an exposure (total AUC) of 15,000 uM.min in 2011-2025 / 13,500 uM.min since 2026 (ref<sup>13-15</sup> and unpublished data of Storek/Kangaroo shown on page 3). In rare cases in which TBI is contraindicated, an exposure of 18,000 uM.min is targeted.<sup>16</sup>

**ABMTP Protocol for Busulfan Dosing:** A test dose of 0.8 mg/kg ideal body weight (IBW) is given typically on day -8, the first dose on day -5, second dose on day -4, third dose on day -3, and fourth dose on day -2. All doses are infused at a constant rate of 80 mg/h. The test dose is given ideally on day -8. Nevertheless, it can be given on an earlier day (e.g. day -9) as long as Bu PK is expected to stay unaltered (e.g., no liver function change, no starting/discontinuing of an azole). Alternatively, the test dose can be given on a later day (e.g. day -7), but it is not preferred as it puts a strain on our PK lab. The area under the time-concentration curve (AUC) expected to be achieved with the first dose (day -5 dose) is determined from post-test dose PK plasma specimens, collected within 5 min and at 1, 3, 5, and 7 h post ending the infusion. The first and second doses (day -5 and -4 doses) are adjusted to achieve the desired total AUC. On day -5, first dose AUC is determined from post-first dose PK plasma specimens (collected within 5 min and at 1, 3, 5, and 7 h post-infusion end). If the estimated total AUC is >20% over or under the target total AUC, the third and fourth doses (day -3 and -2 doses) are adjusted. Based on our 2011-2024 experience, the second adjustment (of the third and fourth doses based on the first dose AUC) is needed in only 3% patients. The formula for dose adjustment is the same for the first adjustment (based on the test dose AUC) and the second adjustment (based on the first dose AUC):

$$\text{Adjusted Dose (mg)} = \text{Intended Dose (mg)} \times (\text{Target AUC (uM.min)} / \text{Observed AUC (uM.min)})$$

The intended first/second/third/fourth (main) dose in 2006-2025 was 3.2 mg/kg because it was assumed that the target total AUC of 15000 uM.min would be achieved by a total dose of 13.0 mg/kg (0.8 mg/kg test dose + 3.2 mg/kg main dose x 4). In reality, the adjusted main dose was virtually always lower than 3.2 mg/kg, as the mean total dose needed to achieve the total AUC of 15000 uM.min was 9.6 mg/kg (0.8 mg/kg test dose + 2.2 mg/kg main dose x 4). As of 2026, we continue to use the intended main dose of 3.2 mg/kg, despite the adjusted dose is estimated to be on average ~2.0 mg/kg (total dose ~8.8 mg/kg, ie, 0.8 mg/kg test dose + ~2.2 mg/kg main dose x 4). The reason

is to avoid changing the way the adjusted dose is calculated, which could lead to mistakes by staff used to calculate the adjusted dose based on the intended dose of 3.2 mg/kg.

In case of Bu-Cy preparative regimen, the busulfan test dose is given on day -10, and busulfan treatment doses on days -8 to -5 are adjusted analogous to above, however, there is no target AUC. The goal is to avoid total AUC of >24000 uM.min.

Common side effects of intravenous busulfan include nausea, vomiting, abdominal pain, anorexia, skin rash, hyperbilirubinemia (grade 3/4 in 30%), electrolyte disturbances, dizziness, headache and insomnia. Serious adverse reactions include hemorrhagic cystitis, male infertility, ovarian failure, venoocclusive disease of the liver, and pneumonitis. Seizures may also occur, and busulfan is always administered with anticonvulsant medications. The ABMTP uses lorazepam 1 mg po qid until 24 hours after the last dose of busulfan for seizure prophylaxis as other anticonvulsant medications show significant drug interactions.

### **Treosulfan**

Treosulfan is an alkylating agent that is structurally and pharmacodynamically similar to busulfan. However, treosulfan may be less toxic than busulfan despite similar antileukemic activity. This has been documented only in the RIC setting. In a randomized study of treosulfan (30 g/m<sup>2</sup>) + fludarabine vs busulfan (6.4 mg/kg) + fludarabine in >50-y-old patients with AML or MDS, treosulfan was associated with a higher OS.<sup>11,12</sup> In this study, busulfan was administered at 0.8 mg/kg i.v. every 6 hours x 8 doses. No PK-adjustment was done. Given the results of this study, we adopted FluTreo for RIC. However, we have not adopted treosulfan for MAC, as it is unclear whether in combination with fludarabine and low-dose TBI, treosulfan is superior to targeted busulfan given daily. Also, the dose of treosulfan in the MAC setting is unknown. Of note, the above RIC randomized study started with 42 g/m<sup>2</sup> and later the dose was changed to 30 mg/m<sup>2</sup> due to “concerns about prolonged neutropenia and subsequent serious infectious complications in the treosulfan group”.<sup>11</sup> On the other hand, the 42 g/m<sup>2</sup> dose has been used with good results in some RIC regimens with fludarabine and 2Gy TBI.<sup>5,17</sup>

An advantage of treosulfan over busulfan is more predictable pharmacokinetics, so no therapeutic drug monitoring with PK-guided dose adjustment is needed.

Common side effects of treosulfan are nausea/vomiting, stomatitis, rash, musculoskeletal pain.

### **Fludarabine**

Fludarabine phosphate (F-Ara-AMP) is a highly-immunosuppressive nucleoside analog with a profound impact on T-Lymphocytes. It is actively dephosphorylated to F-Ara-A in peripheral blood and rephosphorylated to F-Ara-ATP after intracellular transport. It inhibits DNA polymerase alpha,



ribonucleotide reductase and DNA primase, thereby inhibiting DNA synthesis. It also interferes with RNA transcription and translation, and induces apoptosis.

Fludarabine is licensed for the treatment of chronic lymphocytic leukemia. Off-label indications include acute myelogenous leukemia, follicular lymphoma, certain T-cell lymphomas and membranous glomerulonephritis. Within the context of stem cell transplantation fludarabine is used for its immunosuppressive properties and is typically given with an alkylating agent.

Side effects of fludarabine include nausea, vomiting, diarrhea and immune system dysfunction. The latter include incidents of autoimmune cytopenias, hemolysis, hemophagocytic syndrome and opportunistic infection (PJP, progressive multifocal leukoencephalopathy, cryptococcal infection). Herpes zoster, Cytomegalovirus and Epstein-Barr virus reactivations may occur. Overdosage may be associated with neurological effects, including blindness, coma, convulsions and death.

In the ABMTP the last dose of fludarabine is given at least 48 h prior to graft infusion, as the presence of residual fludarabine at the time of graft infusion is associated with a two-day difference in the time to neutrophil engraftment<sup>18</sup>. Approximately 40% of fludarabine clearance is renal; dosage adjustments are recommended for patients with compromised renal function. Patients with normal renal function (creatinine clearance > 60 ml/minute) should receive full dose, while those with moderate renal impairment (creatinine clearance 45-60 ml/minute) should receive a 30% dose reduction. Patients with severely impaired renal function (creatinine clearance <45 ml/minute) should receive a 70% dose reduction. This information is also contained in the BMT protocol data sheets maintained by pharmacy on Unit 105/119.

### **Etoposide (VP-16)**

Etoposide is a topoisomerase-II inhibitor which acts at the premitotic phase to inhibit DNA synthesis. It is cell-cycle specific with maximum activity in the S and G2 phases of cell division. Etoposide has been licensed by the US FDA for treatment of small cell lung and testicular cancers. A long list of off-label uses includes acute myeloid and acute lymphoblastic leukemia and Hodgkin and non-Hodgkin lymphomas.

Etoposide is administered at concentrations no higher than 0.4 mg/mL as it may precipitate. It is given over 4 hours as hypotension may occur with more rapid infusions. Anaphylaxis should be treated with methylprednisolone 250 mg IV) +/- epinephrine 0.2-0.5 mg (0.2-0.5 mL of 1:1000 solution) subcutaneously or intramuscularly.

Common side effects of etoposide include nausea, vomiting, diarrhea and severe mucositis. A rare but important side effect is anaphylaxis.

## Melphalan

Melphalan is an alkylating agent that acts primarily through the alkylation and cross-linking of DNA. It is not cell cycle dependent. Melphalan is detoxified by chemical hydrolysis in plasma. The primary metabolites are inactive and dosage adjustment is not required in renal failure.

FDA has licensed melphalan for palliative treatment of multiple myeloma and ovarian carcinoma. A black box warning indicates that severe myelosuppression may occur with melphalan. Its use has also been associated with development of chromosomal damage and leukemia, although this effect has been only rarely observed with the use of single-agent melphalan conditioning.

In addition to severe cytopenias, high-dose melphalan causes severe mucositis in transplant recipients.

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## Appendix A Table 1. Conditioning regimen, GVHD prophylaxis, and graft preferences by diagnosis.

| Disease                                               | Conditioning                                                                                                          | GVHD Prophylaxis                                        | Graft  |
|-------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------|---------------------------------------------------------|--------|
| <b>Allogeneic HCT</b>                                 |                                                                                                                       |                                                         |        |
| <i>Myeloid Malignancies / ALL</i>                     |                                                                                                                       |                                                         |        |
| MAC up to age 70                                      | <b>Flu</b> (200mg/m <sup>2</sup> ) + <b>Bu</b> (13500uM.min) + <b>TBI</b> (2Gy x2)                                    | ATG+CNI+MTX if HLA-matched*<br>PTCy+Tacro+MMF if HLA-MM | PBSC   |
| RIC if >70 or high TRM risk                           | <b>Flu</b> (120mg/m <sup>2</sup> ) + <b>Treo</b> (30g/m <sup>2</sup> )                                                | ATG+CNI+MTX if HLA-matched*<br>PTCy+Tacro+MMF if HLA-MM | PBSC   |
| Second allogeneic transplant for relapse (same donor) | <b>Etoposide</b> (60mg/kg) + <b>TBI</b> (5Gy x1)                                                                      | CNI                                                     | PBSC   |
| Second allogeneic transplant for relapse (new donor)  | <b>Etoposide</b> (60mg/kg) + <b>TBI</b> (5Gy x1)                                                                      | CNI+MTX                                                 | PBSC   |
| Second allogeneic transplant for graft failure        | <b>Flu</b> (150mg/m <sup>2</sup> ) + <b>Cy</b> (29 mg/kg) + <b>TBI</b> (2Gy x1)                                       | PTCy+Tacro+MMF                                          | PBSC   |
| <i>Lymphoma/CLL</i>                                   | <b>Flu</b> (120mg/m <sup>2</sup> ) + <b>Treo</b> (30g/m <sup>2</sup> )                                                | PTCy+Tacro+MMF                                          | PBSC   |
| <i>Aplastic Anemia</i>                                |                                                                                                                       |                                                         |        |
| Matched sib                                           | <b>Flu</b> (120mg/m <sup>2</sup> ) + <b>Cy</b> (120mg/kg)                                                             | ATG+CNI+MTX                                             | Marrow |
| Haploidentical or Unrelated                           | <b>Flu</b> (150mg/m <sup>2</sup> ) + <b>Cy</b> (29mg/kg) + <b>TBI</b> (2 or 4Gy x1)**                                 | ATG+PTCy+Tacro+MMF                                      | Marrow |
| <i>Hemoglobinopathy</i>                               |                                                                                                                       |                                                         |        |
| Matched sib                                           | <b>TBI</b> (3Gy x1)                                                                                                   | Alemtuzumab+Sirolimus                                   | PBSC   |
| Haploidentical or Unrelated                           | <b>Flu</b> (150mg/m <sup>2</sup> ) + <b>Cy</b> (29mg/kg) + <b>Thiotepa</b> (10mg/kg) + <b>TBI</b> (2Gy)               | ATG+PTCy+Sirolimus+MMF                                  | Marrow |
| <b>Autologous HCT</b>                                 |                                                                                                                       |                                                         |        |
| <i>Multiple myeloma</i>                               | <b>Melphalan</b> 200 mg/m <sup>2</sup> ***                                                                            | NA                                                      | PBSC   |
| <i>Aggressive NHL (DLBCL, PTCL)</i>                   | ( <b>R</b> ) + <b>Bu</b> (13500uM.min) + <b>Mel</b> (140mg/m <sup>2</sup> )                                           | NA                                                      | PBSC   |
| <i>Indolent NHL (FL, MZL, LPL, CLL/SLL)</i>           | ( <b>R</b> ) + <b>Mel</b> (180mg/m <sup>2</sup> ) + <b>TBI</b> (5Gy x1)                                               | NA                                                      | PBSC   |
| <i>Mantle cell lymphoma</i>                           | ( <b>R</b> ) + <b>Mel</b> (180mg/m <sup>2</sup> ) + <b>TBI</b> (5Gy x1)                                               | NA                                                      | PBSC   |
| <i>Hodgkin lymphoma</i>                               | <b>Gemcitabine</b> 1500 mg/m <sup>2</sup> + <b>Melphalan</b> 200 mg/m <sup>2</sup>                                    | NA                                                      | PBSC   |
| <i>Primary CNS lymphoma</i>                           | ( <b>R</b> ) + <b>Thiotepa</b> (600mg/m <sup>2</sup> ) + <b>Bu</b> (13500 uM.min)                                     | NA                                                      | PBSC   |
| <i>Secondary CNS lymphoma</i>                         | ( <b>R</b> ) + <b>Thiotepa</b> (500mg/m <sup>2</sup> ) + <b>Bu</b> (13500uM.min) + <b>Mel</b> (100mg/m <sup>2</sup> ) | NA                                                      | PBSC   |

\* Tacro+MMF (no ATG) if cord blood.

\*\* 2 Gy if previous immunosuppressive therapy, 4 Gy (in a single fraction) if no previous immunosuppressive therapy.

\*\*\* 140-180 mg/m<sup>2</sup> if impaired renal function.

**Abbreviations:** CNI = calcineurin inhibitor (Cyclosporine A or Tacrolimus)

## Appendix B. Conditioning Protocol Details

### **Flu(200mg/m<sup>2</sup>) + Bu(13500uM.min) + TBI(2Gy x2)**

Fludarabine 50 mg/m<sup>2</sup>/day on days -5 to -2

Busulfan 0.8 mg/kg test dose (on ~day -8) and PK-determined dose on days -5 to -2 (approximately 2 mg/kg/day) to achieve total busulfan exposure of 13500 uM.min.

TBI 4 Gy delivered to midplane in two divided doses on day -1 or 0 (before graft infusion), at least 6 hours apart.

### **Flu(250mg/m<sup>2</sup>) + Bu(18000uM.min)**

Fludarabine 30 mg/m<sup>2</sup> on days -5 to -2

Treosulfan 10 mg/m<sup>2</sup> on days -4 to -2

### **Cy(120mg/kg) + Bu(<24000uM.min)**

Cyclophosphamide 60 mg/kg IV on days -8 and -7

Busulfan 3.2 mg/kg IV on days -5, -4, -3, -2. Avoid daily AUC >6,000 umol.min/L (avoid overall AUC >24,000 umol.min/L).

*The reason for using Cy+Bu instead of “conventional” Bu+Cy is a randomized study that showed borderline superiority of Cy+Bu due to lower hepatotoxicity and lower NRM.<sup>19</sup>*

*This conditioning should be used only in case of fludarabine shortage. Bu+Cy has been found in two randomized studies to result in similar overall survival as Flu+Bu,<sup>20,21</sup> however, whether it is equivalent to Flu+Bu+4GyTBI is not known. Moreover, the busulfan toxic AUC of >24,000 uM.min has been determined in combination with fludarabine; it is not known whether it applies also to combination with cyclophosphamide. Moreover, in the randomized studies, daily busulfan dose was divided into two or four doses. We use only once daily dose as an extrapolation from our Flu-Bu experience.*

### **Etoposide(60mg/kg) + TBI(5Gy x1)**

Etoposide 60 mg/kg on day -4

TBI 500 cGy delivered to midplane in a single fraction on day 0 (before graft infusion)

### **Flu(120mg/m<sup>2</sup>) + Cy(120mg/kg)**

Fludarabine 30 mg/m<sup>2</sup>/day on days -6 to -3

Cyclophosphamide 60 mg/kg on days -4 and -3

### **Flu(150mg/m<sup>2</sup>) + Cy(29mg/kg) + TBI(2 or 4Gy x1) (Baltimore SAA regimen)**

Fludarabine 30 mg/m<sup>2</sup>/day on days -6 to -2

Cyclophosphamide 14.5 mg/kg on days -6 and -5

TBI 2 or 4 Gy delivered to midplane in a single fraction on day -1 (2 Gy if previous immunosuppressive therapy, 4 Gy if no previous immunosuppressive therapy).  
Note: ATG is given as 0.5 mg/kg on day -9, 2 mg/kg on day -8, and 2 mg/kg on day -7

***TBI(3Gy x1) (NIH SCD regimen)***

TBI 300 cGy delivered to midplane in a single fraction on day -2  
Note: Alemtuzumab is given as 0.03 mg/kg on day -7, 0.1 mg/kg on day -6, and 0.3 mg/kg on days -5, -4, and -3..

***Flu(150mg/m<sup>2</sup>) + Cy(29mg/kg) + Thiotepa(10mg/kg) + TBI(2Gy) (Vanderbilt SCD regimen)***

Fludarabine 30 mg/m<sup>2</sup>/day on days -6 to -2  
Thiotepa 10 mg/kg on day -7  
Cyclophosphamide 14.5 mg/kg on days -6 and -5  
TBI 2 Gy delivered to midplane in a single fraction on day -1  
Note: ATG is given as 0.5 mg/kg on day -9, 2 mg/kg on day -8, and 2 mg/kg on day -7

***Historical pre-alloHCT Regimens***

***Flu(250mg/m<sup>2</sup>) + Bu(18000uM.min) (MAC for pts with too much previous radiation)***

Fludarabine 50 mg/m<sup>2</sup>/day on days -6 to -2  
Busulfan 3.2 mg/kg/day on days -5 to -2, adjusted based on pharmacokinetics in order to achieve total busulfan exposure of 18000 µmol·min/L

***Flu(120mg/m<sup>2</sup>) + Mel(140mg/m<sup>2</sup>) (RIC)***

Fludarabine 30 mg/m<sup>2</sup> days -5 to -2  
Melphalan 140 mg/m<sup>2</sup> day -1

***Flu(250mg/m<sup>2</sup>) + TBI(5Gy x1)***

Fludarabine 50 mg/m<sup>2</sup> on days -6 to -2  
TBI 500 cGy to midplane on day -1 or 0

***Pre-autoHCT Regimens (will be removed from here to appear in disease-specific chapters)***

***Gem-Mel***

Gemcitabine 75mg/m<sup>2</sup> bolus then 1425mg/m<sup>2</sup> infusion on day -1  
Melphalan 200 mg/m<sup>2</sup> on day -1

***Mel 200***

Melphalan 200 mg/m<sup>2</sup> on day -1



Note: 140-180 mg/m<sup>2</sup> if impaired renal function

**(R) + Etoposide(60 mg/kg) + Mel(180mg/m<sup>2</sup>)**

(Rituximab 375 mg/m<sup>2</sup> IV day -4)

Etoposide 60mg/kg day -4

Melphalan 180mg/m<sup>2</sup> day -2

**(R)BEAM** (not used since 2018 due to carmustine becoming too expensive)

(Rituximab 375 mg/m<sup>2</sup> IV on day -6)

Carmustine (BCNU) 300 mg/m<sup>2</sup> on day -6

Etoposide 100 mg/m<sup>2</sup> q12h x 8 doses on days -5 to -2

Cytarabine 200 mg/m<sup>2</sup> q12h x 8 doses on days -5 to -2

Melphalan 140-160 mg/m<sup>2</sup> on day -1

**Thiotepa(600mg/m<sup>2</sup>) + Bu(13500uM.min) for Primary CNS Lymphoma**

Thiotepa 300 mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age 18-60years

270mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age 61-65years

240mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age 66-70years

210mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age >70years

Busulfan (only 3.2 mg/kg/day days -4 to -2 targeted to achieve busulfan AUC <13500 uM.min)

3.2 mg/kg (Ideal weight) IV daily on days -4 to -2 if age 18-60years

2.9mg/kg (Ideal weight) IV daily on days -4 to -2 if age 61-65years

2.55mg/kg (Ideal weight) IV daily on days -4 to -2 if age 66-70years

2.25mg/kg (Ideal weight) IV daily on days -4 to -2 if age >70years

PK testing is run on the first dose and the third dose is adjusted (if needed) to avoid a total AUC of over 13500 uM.min

**(R) + Thiotepa(500mg/m<sup>2</sup>) + Bu(<13500uM.min) + Mel(100 mg/m<sup>2</sup>) for Secondary CNS Lymphoma**

(Rituximab 375 mg/m<sup>2</sup> IV on day -7)

Thiotepa 250 mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age 18-64years

225mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age 65-69years

200mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if over age 70years

Busulfan (only 3.2 mg/kg/day days -4 to -2 targeted to achieve busulfan AUC <13500 uM.min)

3.2 mg/kg (Ideal weight) IV daily on days -4 to -2 if age 18-64years

2.9mg/kg (Ideal weight) IV daily on days -4 to -2 if age 65-69years

2.55mg/kg (Ideal weight) IV daily on days -4 to -2 if over age 70years

Melphalan 100 mg/m<sup>2</sup> on day -1 if age 18-64years

90mg/m<sup>2</sup> IV on day -1 if age 65-69years

80mg/m<sup>2</sup> IV on day -1 if over age 70years

PK testing is run on the first dose and the third dose is adjusted to target total AUC of 13500 uM.min +/- 20%

**(R) + Bu(1350uM.min) + Mel(140mg/m<sup>2</sup>)**

Rituximab 375 mg/m<sup>2</sup> IV on day -5

Busulfan 3.2 mg/kg/day on days -4 to -2 targeted to achieve busulfan AUC of 13500 uM.min

Melphalan 140 mg/m<sup>2</sup> on day -1

***Mel 200***

Melphalan 200 mg/m<sup>2</sup> on day -1

Note: 140-180 mg/m<sup>2</sup> if impaired renal function

***(R) + Etoposide(60 mg/kg) + Mel(180mg/m<sup>2</sup>)***

(Rituximab 375 mg/m<sup>2</sup> IV day -4)

Etoposide 60mg/kg day -4

Melphalan 180mg/m<sup>2</sup> day -2

***(R)BEAM*** (not used since 2018 due to carmustine becoming too expensive)

(Rituximab 375 mg/m<sup>2</sup> IV on day -6)

Carmustine (BCNU) 300 mg/m<sup>2</sup> on day -6

Etoposide 100 mg/m<sup>2</sup> q12h x 8 doses on days -5 to -2

Cytarabine 200 mg/m<sup>2</sup> q12h x 8 doses on days -5 to -2

Melphalan 140-160 mg/m<sup>2</sup> on day -1

***Thiotepa(600mg/m<sup>2</sup>) + Bu(13500uM.min) for Primary CNS Lymphoma***

Thiotepa 300 mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age 18-60years

270mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age 61-65years

240mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age 66-70years

210mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age >70years

Busulfan (only 3.2 mg/kg/day days -4 to -2 targeted to achieve busulfan AUC <13500 uM.min)

3.2 mg/kg (Ideal weight) IV daily on days -4 to -2 if age 18-60years

2.9mg/kg (Ideal weight) IV daily on days -4 to -2 if age 61-65years

2.55mg/kg (Ideal weight) IV daily on days -4 to -2 if age 66-70years

2.25mg/kg (Ideal weight) IV daily on days -4 to -2 if age >70years

PK testing is run on the first dose and the third dose is adjusted (if needed) to avoid a total AUC of over 13500 uM.min

***(R) + Thiotepa(500mg/m<sup>2</sup>) + Bu(<13500uM.min) + Mel(100 mg/m<sup>2</sup>) for Secondary CNS Lymphoma***

(Rituximab 375 mg/m<sup>2</sup> IV on day -7)

Thiotepa 250 mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age 18-64years  
225mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if age 65-69years  
200mg/m<sup>2</sup> (Ideal BSA) on days -6 and -5 if over age 70years  
Busulfan (only 3.2 mg/kg/day days -4 to -2 targeted to achieve busulfan AUC <13500 uM.min)  
3.2 mg/kg (Ideal weight) IV daily on days -4 to -2 if age 18-64years  
2.9mg/kg (Ideal weight) IV daily on days -4 to -2 if age 65-69years  
2.55mg/kg (Ideal weight) IV daily on days -4 to -2 if over age 70years  
Melphalan 100 mg/m<sup>2</sup> on day -1 if age 18-64years  
90mg/m<sup>2</sup> IV on day -1 if age 65-69years  
80mg/m<sup>2</sup> IV on day -1 if over age 70years  
PK testing is run on the first dose and the third dose is adjusted to target total AUC of  
13500 uM.min +/- 20%

**(R) + Bu(1350uM.min) + Mel(140mg/m<sup>2</sup>)**

Rituximab 375 mg/m<sup>2</sup> IV on day -5

Busulfan 3.2 mg/kg/day on days -4 to -2 targeted to achieve busulfan AUC of 13500 uM.min

Melphalan 140 mg/m<sup>2</sup> on day -1

Appendix C. Results using our standard conditioning (Flu250 + Bu15000 + TBI4Gy) in combination with PBSC graft and ATG+CSA+MTX GVHD prophylaxis, using matched sib or 7-8/8 HLA matched unrelated donor

Figures 1-5 are based on Ousia et al: Clin Transplant 2020.<sup>22</sup>  
 Figure 6 is based on patients who underwent HCT in 2011-2024 (unpublished).

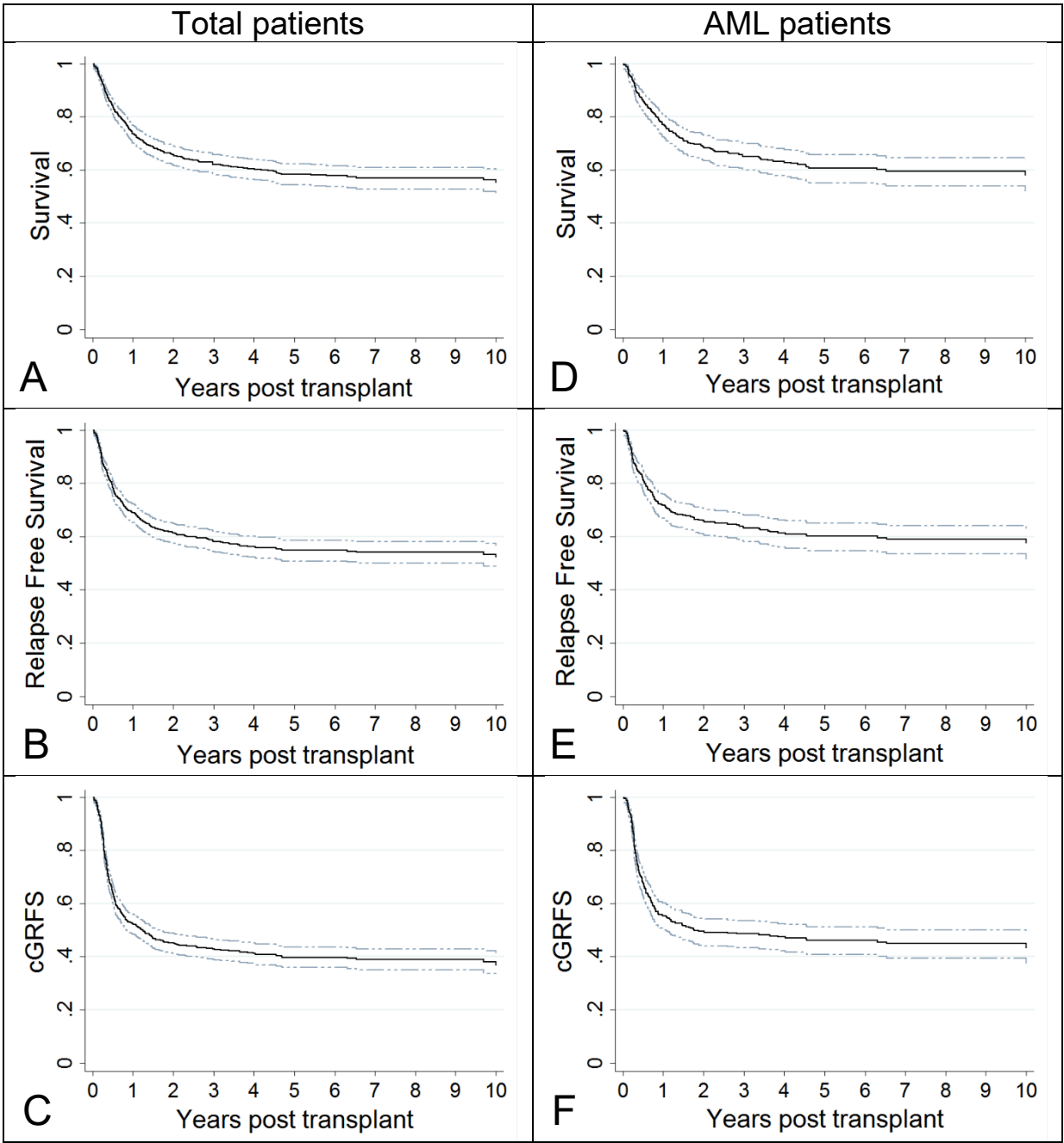


**Figure 1.** Overall survival (A), relapse-free survival (B), and chronic graft versus host disease- and relapse-free survival (cGRFS) (C) in the total cohort of 700 patients (any hematologic malignancy). Overall survival (D), relapse-free survival (E), and cGRFS (F) in patients with acute myeloid leukemia (AML). The dot-dash lines show 95% confidence intervals.

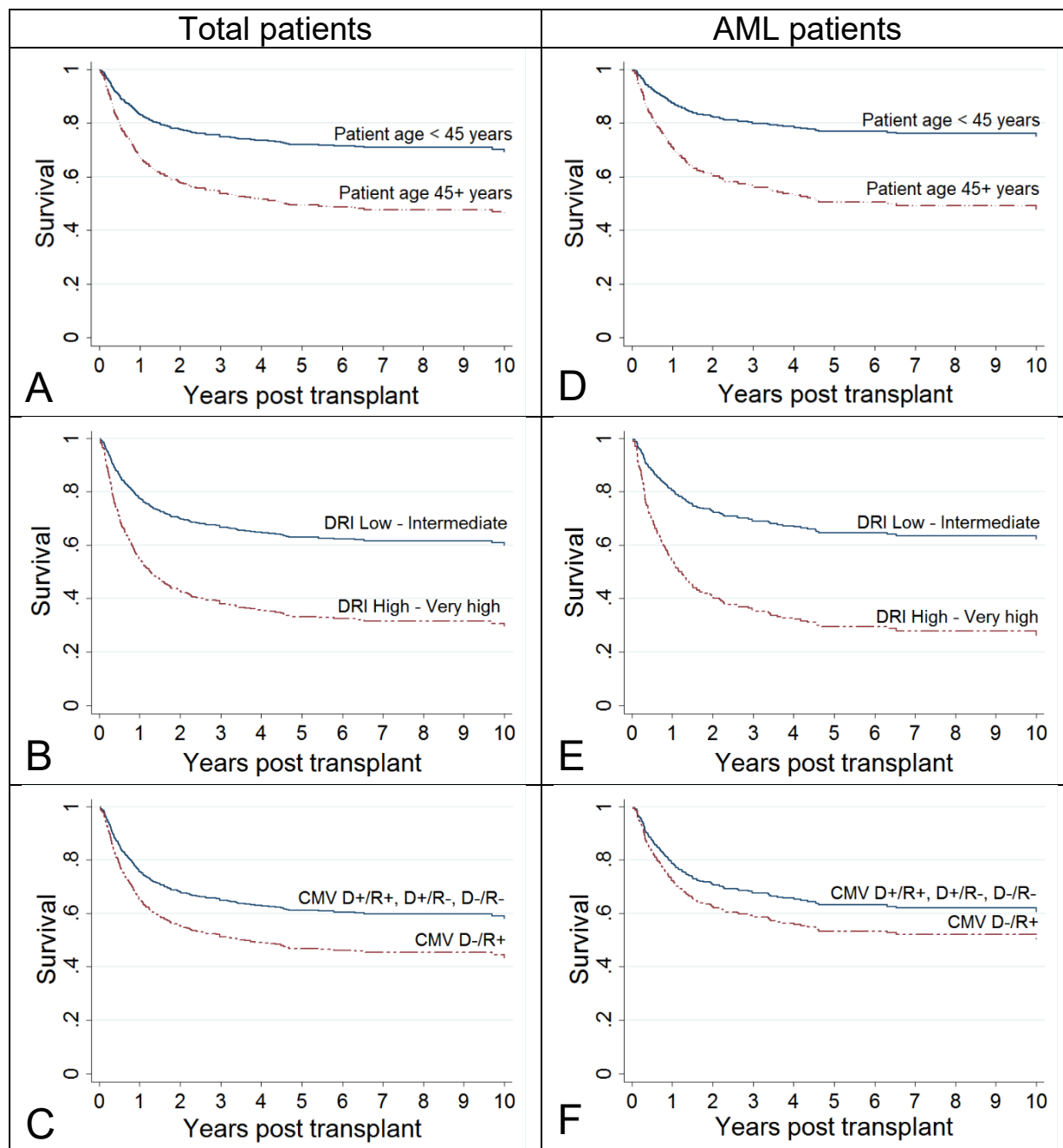


**Figure 2.** Impact of patient age, disease risk index (DRI), and CMV serostatus on overall survival in total patients (A, B, C) and patients with acute myeloid leukemia (D, E, F). All the differences were significant in multivariate analysis, except for the CMV serostatus in AML patients. DRI = disease risk index per Armand et al.<sup>23</sup>

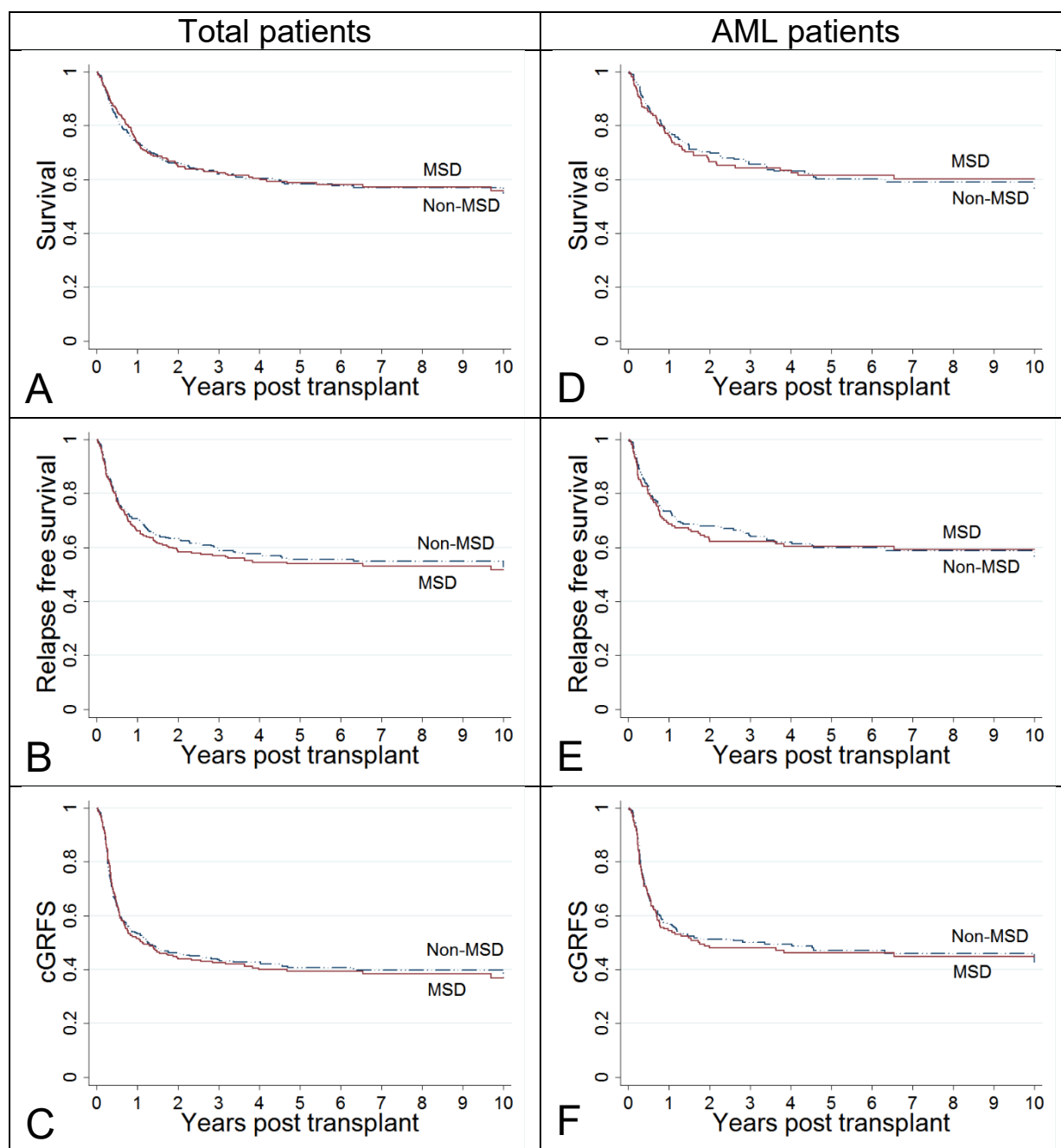


**Figure 3.** Virtually no impact of HLA matched sibling donor (MSD, solid red curve) vs non-MSD (mostly 7-8/8 HLA allele-matched unrelated donor, dashed blue curve) on OS, RFS, or cGRFS. None of the differences was significant in multivariate analysis.



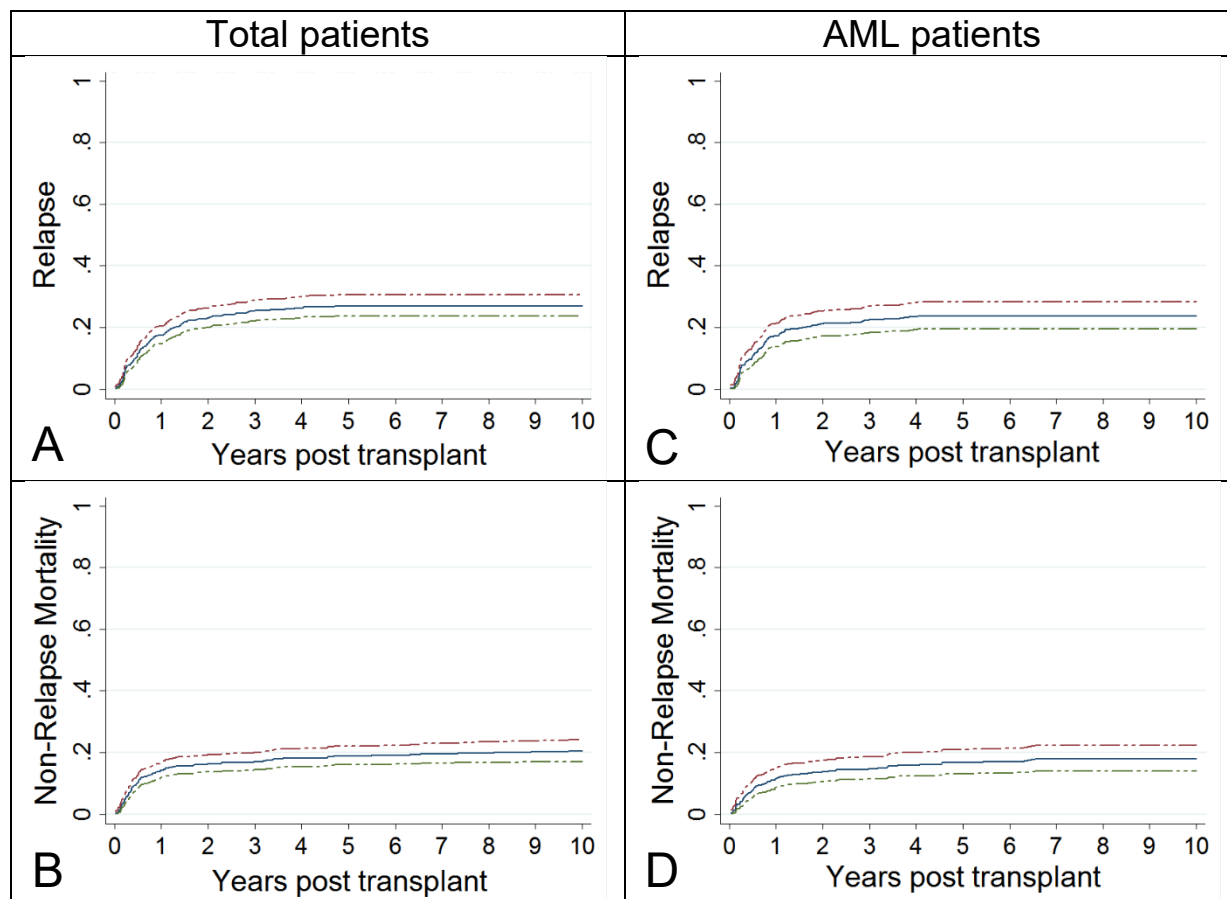


**Figure 4.** Relapse and non-relapse mortality in total patients (the whole cohort of 700 patients) (A, B) and patients with acute myeloid leukemia (C, D). The dot-dash lines show 95% confidence intervals.

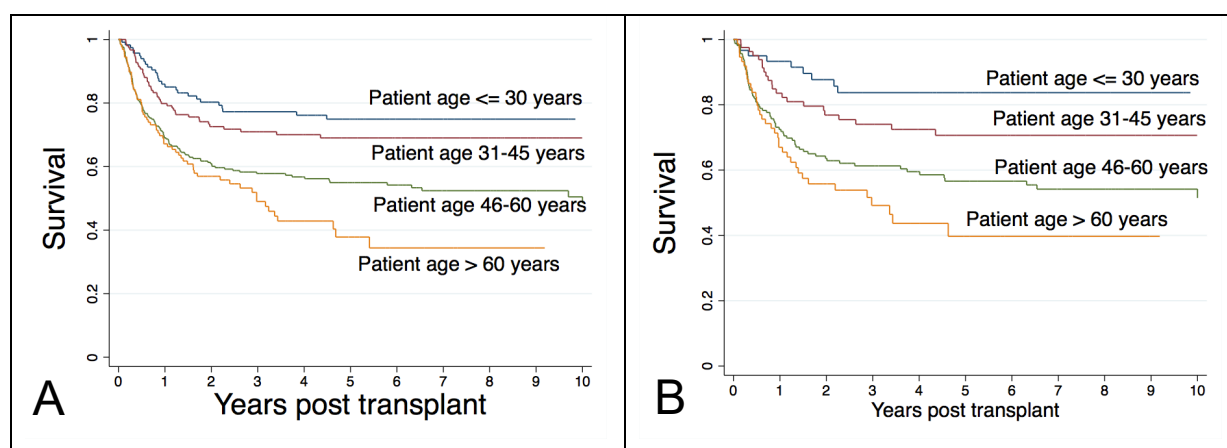
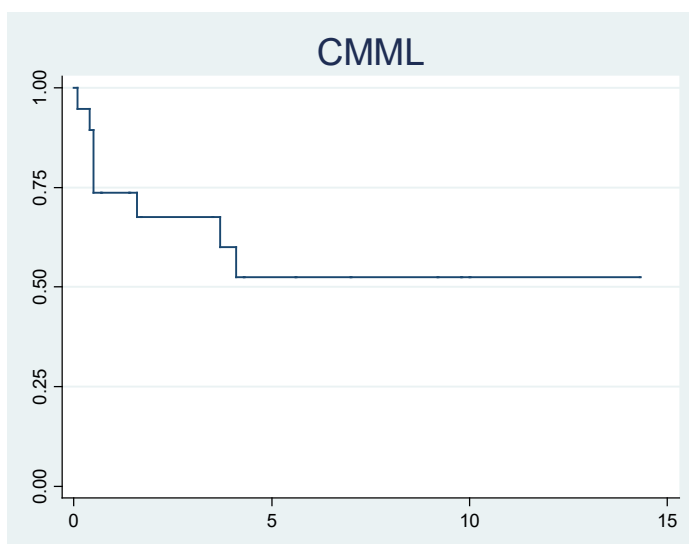
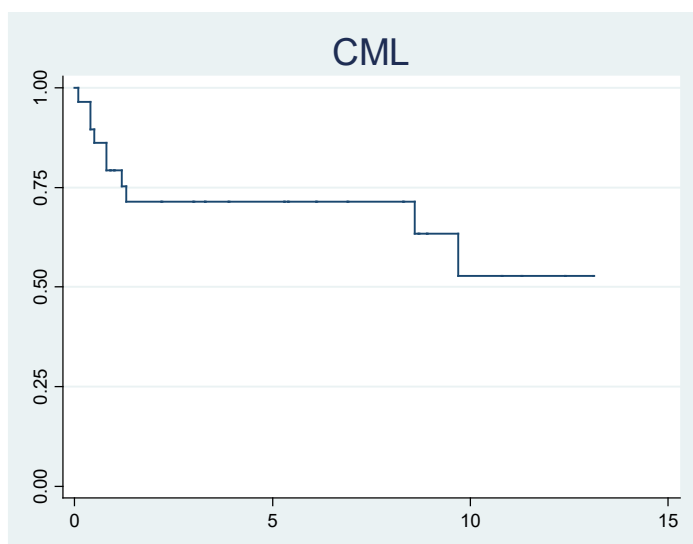
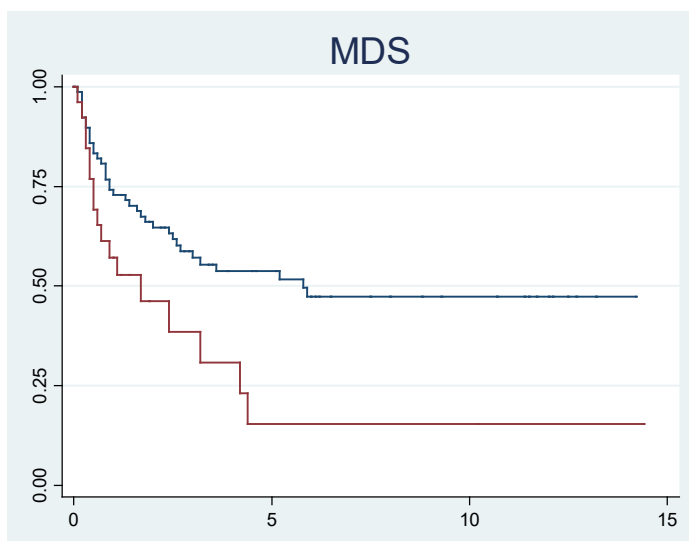
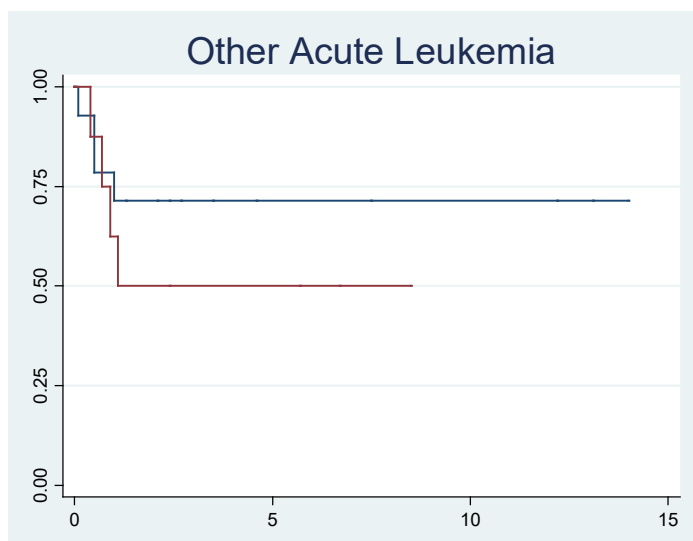
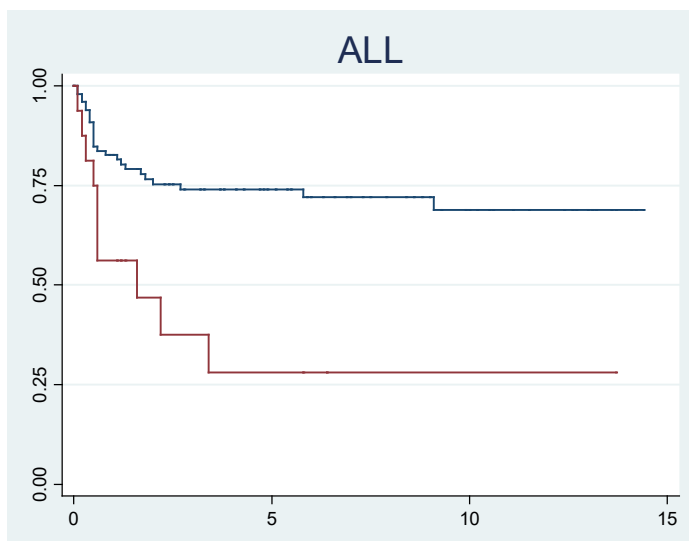
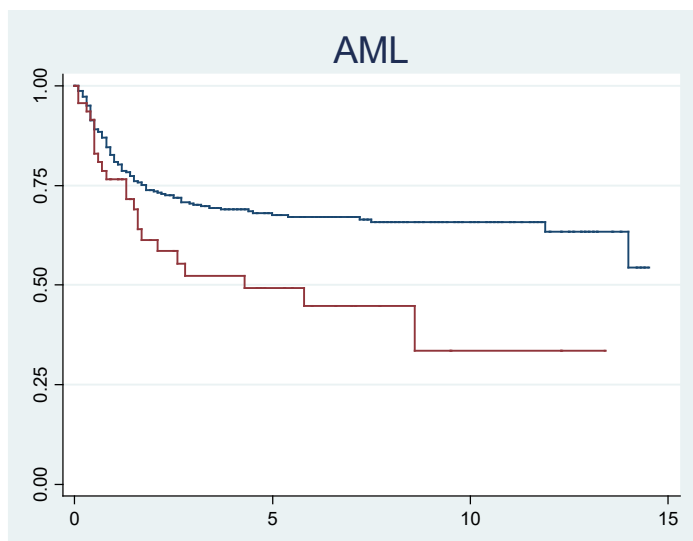


**Figure 5.** Overall survival by patient age among total patients (any hematologic malignancy) (A) and AML patients (B).



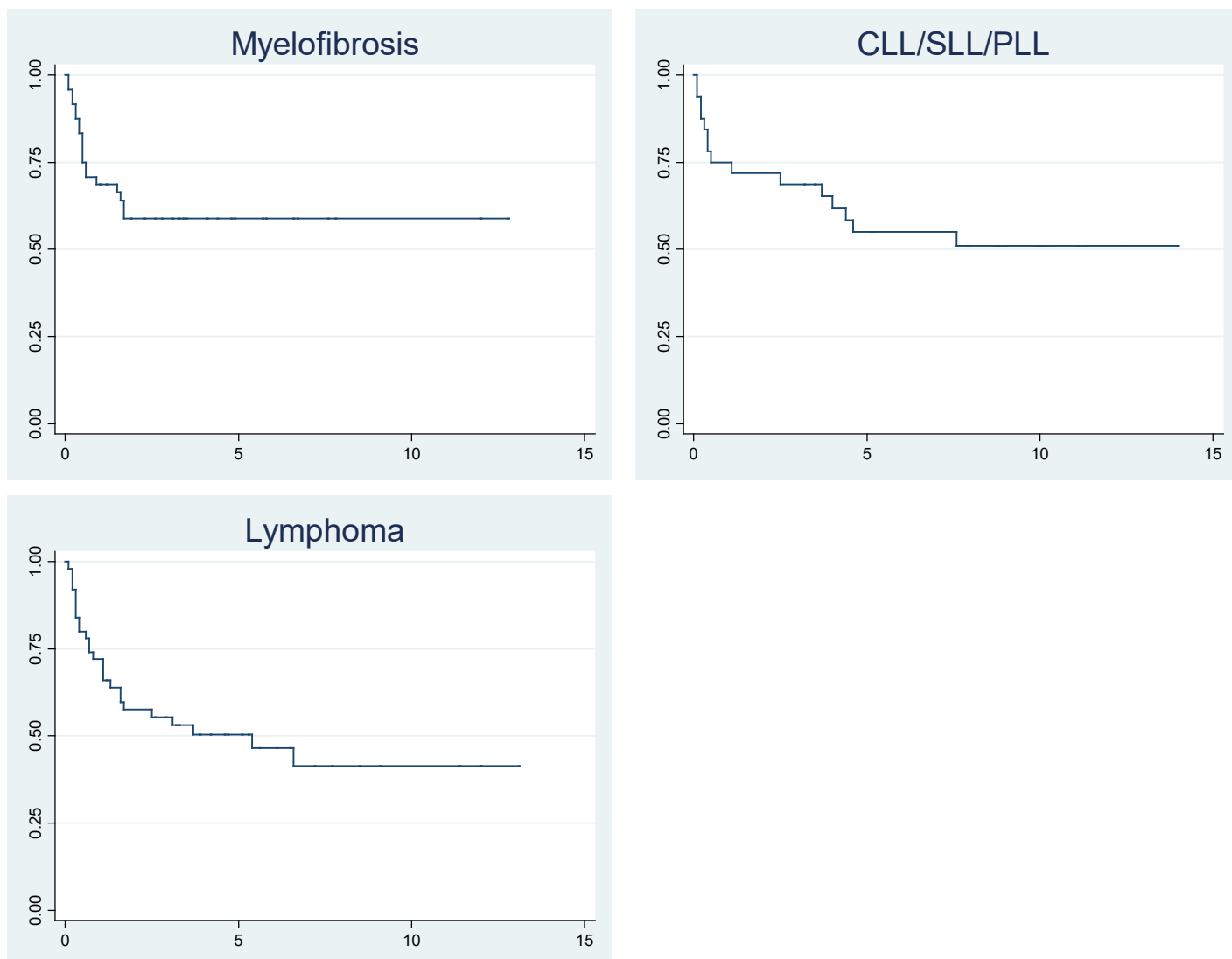


**Figure 6.** Overall survival of alloHCT recipients in 2011-2024 (N=829) according to their disease. In graphs in which 2 curves are given, the top curve pertains to low or intermediate disease risk index (DRI) and the bottom curve to high or very high DRI.<sup>23</sup>

For AML and other acute leukemia, high or very high DRI refers to non-CR or CR with complex cytogenetics ( $\geq 5$  abnormalities).

For ALL, high or very high DRI refers to beyond 1CR.

For MDS, high or very high DRI refers to  $>5\%$  blasts or  $<5\%$  blasts with chromosome 7 abnormality or complex cytogenetics ( $\geq 3$  abnormalities).

For CLL/SLL/PLL and Lymphoma, the graphs are of only historical interest, as we currently use FluTreo RIC.

# Transplantation Eligibility Assessment: Patient Factors

Presented by: Jason Tay

## Summary

1. We recommend the routine documentation and use of the Hematopoietic cell transplantation (HCT) specific comorbidity index (HCT-CI) and its components as part of the pre-transplant evaluative process.
2. A Frailty & Functionality Assessment should be performed for all patients undergoing allogeneic HCT, regardless of age, and the HCT frailty score should be calculated. A more in-depth Geriatric Assessment can be considered in select patients, particularly those over age 65, to better aid decision making. It would also be good practice in the autologous HCT setting but should be individualized.
3. The following are RELATIVE contraindications for HCT. A referral to appropriate subspecialty services is indicated if HCT is being considered for a patient who does not meet any of these minimal thresholds:
  - a. Age >65
  - b. Karnofsky performance score (KPS) <60
  - c. FEV1 or DLCO <60% predicted
  - d. LVEF <45% or arrhythmia
  - e. Bilirubin/ALT/ALP >2x upper normal limit (UNL)
  - f. Creatinine >2x ULN
  - g. Uncontrolled infection, including dental
  - h. HCT frailty scale Score  $\geq 5.5$
4. The following are ABSOLUTE contraindications for HCT.
  - a. Active second malignancy
  - b. Cirrhosis of the liver
  - c. Pregnancy
  - d. HCT-CI  $\geq 3$  plus one abnormal IADL (ability to use phone, laundry, shopping, mode of transportation, food preparation, responsibility for own medications, housekeeping, ability to handle finances) in patient's >65 years.
5. Early (ideally at diagnosis of malignancy) referral of the patient with mental illness or other psychosocial concerns to psychology, social work, psychiatry as appropriate is important. When psychosocial factors severely impair functioning and/or adherence to treatment plan, or place the patient at immediate safety risk (e.g., actively psychotic, suicidal, substance dependent, extreme poverty, high degree of family conflict), HCT may be deferred in order to prioritize stabilization of psychosocial concerns.
6. The ultimate decision to proceed to HCT is an interdisciplinary team-based decision paying attention to recipient characteristics and their perceived "trade-offs" with disease and donor characteristics.

7. Eligibility for chimeric antigen receptor T-cell therapy (CAR-T) will follow the same principals for assessment as in standard HCT.

## Background

Hematopoietic cell transplantation (HCT) is a potentially curative therapy for a variety of malignant and nonmalignant hematological disorders. The decision to recommend and proceed with HCT is complex and multi-faceted. Prior to recommendation a throughout assessment of 1) Disease characteristics, 2) Patient characteristics – Physical and Psychosocial, and 3) Donor characteristics (allogeneic setting) is required.

The relative contributions of these characteristics (potentially overlapping) to HCT success is not and unlikely to be clearly defined. In part, evaluations of individual characteristics within observational studies variably consider other pertinent characteristics. Moreover, secular trends in HCT technology and supportive care would suggest the relative contributions would be “fluid”. *The ultimate decision to proceed to HCT is an interdisciplinary team-based decision paying attention to these characteristics and their perceived “trade-offs”.*

While it is important to acknowledge that patient characteristics are associated with post-HCT outcomes, there is no clear and/or consistent evidence that modification of these characteristics clearly attenuates post-HCT outcomes –modifiable risk factor(s). This review focuses on **Patient** characteristics and draws attention to assessments and variables that might influence the decision to proceed with HCT.

## Physical Assessment(s)

A detailed history, physical examination complemented with investigative diagnostics is a crucial 1<sup>st</sup> step in documenting and assessing comorbidities<sup>1-3</sup>. The rationale is presented in the following sections:

### Age

An ideal HCT candidate should be in excellent physical and physiologic health at the time of HCT. There is a movement to consider physiologic age over chronologic age in the determination of HCT eligibility. However, the chronologic age could be considered a simple variable that embraces a multitude of patient characteristics.

In the autologous setting, data on the impact of age on outcomes post HCT is predominantly in myeloma setting. Data from randomized controlled trials would support autologous HCT up to a biologic age of 65 years. In contrast, there is no randomized controlled trial data supporting autologous HCT in patients aged >65 years with multiple myeloma. However, there are indeed observational studies that suggest that autologous HCT can be safely performed in older patients in myeloma<sup>4-8</sup>. There are fewer studies examining its impact in the lymphoma setting<sup>9-11</sup>. The paucity of

data would suggest that biologic age should not be sole criteria used to determine eligibility for autologous HCT. Rather, one may need to consider attenuating the dosing of the conditioning regimen to compensate for age among other factors as discussed.

In the allogeneic setting, CIBMTR registry data suggests that the median age of HCT has increased to up to 75 years over the last few decades<sup>12</sup>. Indeed, a retrospective study from EBMT suggests that there is no significant association between age and relapse or non-relapse mortality in a cohort of 1333 patients (age 50-74 years)<sup>13</sup>. Further, a similar analysis from CIBMTR in 1080 patients (>40 years) receiving a reduced intensity conditioning found that chronologic age did not impact rates of non-relapse mortality, relapse or GVHD<sup>14</sup>. Finally, a review of 372 patients aged 60-75 enrolled in prospective clinical trials of a reduced intensity conditioning determined that age did not appear to influence GVHD, PFS or OS but older individuals had increased bacterial infections and hospitalization<sup>15</sup>.

Given potential and inherent biases in the assessment of physiologic age, there is increasing interest in utilizing biomarkers of physiologic age<sup>16</sup>. Indeed, there are numerous candidate markers including: p16<sup>INK4A</sup>, Leukocyte telomere length, DNA methylation, miRNA, Immunosenescence, SASP, Anemia, IL-6, CRP, NT-proBNP, Albumin, D-dimer, TNF and sICAM-1. Further, various geriatric assessment scales have also been used<sup>17,18</sup>. Among these, it appears that p16<sup>INK4A</sup> may be a leading biomarker candidate – a molecular marker of cellular senescence<sup>19-21</sup>.

*Observational health outcomes research evaluating age is inherently confounded by indication that might suggest a more conservative approach in utilizing age in determining HCT eligibility. Taken together, it is reasonable to consider using a cautious and extensive evaluation for older (e.g. >65years) or frailer patients.*

## **Performance Status**

With respect to performance status assessment, we prefer the Karnofsky Performance Score (KPS) score over Eastern Cooperative Oncology Group (ECOG) score as it allows a more “granular” range to base one’s assessment. Moreover, the assessment of performance status is subjective, and a wider scoring range may improve the quality of the assessment. Given concerns that performance status is clinician assigned with overestimation<sup>22</sup>, a geriatric assessment (GA) has its proponents in older patients<sup>23,24</sup>. See section on Geriatric assessments.

*We suggest that it is reasonable to proceed with HCT if the KPS score >60 and consider utilizing CGA in individuals who are >65 years of age to better guide decision making. In those with a KPS of <60, a more extensive evaluation would be warranted. Further, we suggest that HCT be not offered in the presence of a HCT-CI score of ≥3 and one abnormal ADL.*



## Pulmonary Evaluation

Post-HCT pulmonary complications such as therapy related lung toxicity, pulmonary GVHD and its variants, TRALI and infectious complications can occur. Pre-existing lung disease as measured by pulmonary function tests (PFTs) can increase the risk and morbidity of post-HCT pulmonary complications with up to 3% and 24% of autologous and allogeneic HCT patients developing severe pulmonary complications requiring mechanical ventilation<sup>25</sup>. Indeed, an abnormal PFT pre-HCT is associated with poorer post-transplant outcomes<sup>26-29</sup>. Further, smoking pre-HCT is independently associated with poor outcomes<sup>30</sup>.

The proposed cutoff for eligibility in HCT in clinical trials is typically a corrected DLCO >50% although a true cutoff is unknown. This cutoff which may be dependent on the planned conditioning chemotherapy<sup>31</sup>. In the allogeneic setting, a higher threshold of DLCO >60% has been used. Moreover, the PAM score (described later) uses a DLCO cutoff of 60%<sup>32</sup>. The correlation between FEV<sub>1</sub> and DLCO pre-HCT is poor, with pre-HCT FEV<sub>1</sub> independently predictive of early respiratory failure<sup>33,34</sup>.

*Taken together, it is optimal to consider HCT in an individual with a DLCO >60% and a FEV<sub>1</sub> >60%. In all other scenarios, the case should be discussed at an individual basis.*

## Cardiac evaluation

In general, individuals with poor cardiac reserve with a LVEF <40%, uncontrolled arrhythmia or coronary artery should not proceed with HCT<sup>35</sup>. Overall, the rate of major or life-threatening cardiac events post-HCT has been estimated to be <1%<sup>36</sup>.

Cardiac injury can occur post-HCT, and it is assumed to be more serious in individuals with less cardiac reserve. A higher LVEF threshold maybe warranted when cardiotoxic conditioning (e.g., cyclophosphamide or TBI) is contemplated. However, it may be reasonable to accept a LVEF of >45% in most circumstances<sup>37,38</sup>. Separately, there is also an association between prolonged QT and QT dispersion noted on routine EKG with post-HCT morbidity from heart failure<sup>39,40</sup>. Further, it would be important to optimize cardiac risk factors prior to HCT<sup>41</sup>.

*Taken together, it is optimal to consider HCT in an individual with a LVEF >45% with a normal EKG. In all other scenarios, the case should be discussed at an individual basis.*

## Hepatic Evaluation

Baseline elevations of serum transaminases and alkaline phosphatase are associated with an increased risk of sinusoidal obstruction syndrome (SOS) post-HCT in the allogeneic setting<sup>42</sup>.

Serum hyperferritinemia is also associated with increased risk of SOS, disease free and overall survival<sup>43-47</sup>. Given these associations, it may be reasonable to consider chelation therapy for iron

overload prior to HCT, in particular patients with multiple red cell transfusion supports; however, whether the chelation improves outcomes has not been reported.

*Taken together, it is reasonable to proceed with HCT if the liver function tests as measured by (Bilirubin, AST, ALT or ALP) are < 2 times upper limit of the normal reference range.*

Seropositivity for Hepatitis B, C or HIV should not preclude HCT, recognizing that it affects peri-transplant care, where viral prophylaxis or optimization of anti-viral therapy would be required. Unsurprisingly, viral hepatitis is associated with increased risk of reactivation, SOS, liver disease post-HCT and non-relapse mortality<sup>48-50</sup>. The use of Transient Elastography (Fibroscan) is suggested if there is clinical concern of cirrhosis<sup>51</sup>.

*In general, it is reasonable to exclude patients with frank cirrhosis from HCT.*

### **Nutritional Evaluation**

There is a paucity of evidence to suggest a specific nutritional state that would preclude HCT. However, is notable that patients that medically obese have similar post- autologous HCT outcomes as patients with a normal BMI<sup>52-55</sup>. Interestingly, obesity is associated with higher non-relapse mortality<sup>56</sup> but a lower relapse rate, resulting in similar overall survival in the allogeneic setting<sup>57</sup> and low BMI is associated with poor HCT outcome<sup>56,58</sup>. In contrast, a Chinese study would suggest that overweight and obese patients had a superior overall survival when compared with underweight and normal-weight patients (HR=0.60; 95% CI: 0.38-0.95)<sup>59</sup>.

*Consequently, it may be reasonable to attempt to increase BMI before HCT, but it has not been reported whether this intervention improves outcome.*

### **Renal Function Evaluation**

Renal dysfunction is associated with a higher morbidity and mortality in patients undergoing autologous HCT for myeloma<sup>60-62</sup>. Importantly, the value of autologous transplants studied in a randomized fashion only included patients with good renal function. In contrast, there is a paucity of data in the autologous setting in lymphoma given that traditional conditioning chemotherapy was not administered in patients with a serum creatinine >177micromol/L.

A similar argument applies in the allogeneic setting and maybe more pertinent given that acute renal injury can occur 15-18% of patients receiving allogeneic HCT<sup>63</sup>. Further, there is some evidence to support an increased risk of non-relapsed mortality in patients with renal impairment pre-HCT<sup>64</sup>. Indeed, long-term follow-up data suggests that the more severe the acute renal injury peri-HCT, the higher the likelihood of chronic kidney disease<sup>65</sup>. Interestingly, the risk of acute renal injury could be anticipated using the HCT-CI (see discussion later)<sup>66</sup>.

*Overall, it is reasonable to proceed with HCT where the Creatinine is < 177micromol/L and are < 2 times upper limit of the normal reference range. All other scenarios should be individualized.*

## **Dental Evaluation**

The goal of pre-HCT dental assessments is to identify potential sources of infection during the peri-HCT period<sup>67,68</sup>. This appears to be good practice but there has been no clear evidence to support an association between radiographic periodontal disease and infections/mortality post-HCT<sup>69,70</sup>.

## **Active Infections**

It appears to be good practice that HSCT should be deferred if there is active infection or infection(s) that are not responding to therapy to decrease the odds of morbidity and mortality peri-HSCT.

## **Comorbidity Indices**

There are multiple standardized co-morbidity indices in clinical use that aims to aid pre-HCT assessments<sup>71</sup>. The purpose would be to incorporate and assign differing weights to characteristics considered in the above sections. However, it is important to note that not all characteristics are considered or considered in the same fashion in the derivation studies.

## **Kaplan-Feinstein Scale**

The Kaplan-Feinstein Scale (KFS) was originally developed to evaluate the impact of survival in patients with diabetes based on comorbid conditions and involves assigning Grades (range 1-3) to 12 comorbid conditions<sup>72</sup>. Artz et al. evaluated 105 consecutively enrolled patients who underwent HCT, receiving reduced intensity conditioning with fludarabine, melphalan, and alemtuzumab. A simple scale combining the Kaplan-Feinstein Scale (KFS) and Eastern Cooperative Oncology Group Performance Status (PS) scale PS enabled separation of high- from low-risk patients, with 6-month cumulative incidences 50% and 15%, respectively for transplant-related mortality (P = .001)<sup>73</sup>.

## **Pre-transplant Assessment of Mortality Score- PAM Score**

This risk score was developed at the Fred Hutchinson center and incorporates 8 pre-transplantation clinical variables: patient age, donor type, disease risk, conditioning regimen, FEV1, carbon monoxide diffusion capacity, serum creatinine level, and serum alanine aminotransferase concentration<sup>32</sup>. This score is useful for predicting the risk for death within the first 2 years after HCT.

The authors re-evaluated the PAM score using a contemporary cohort (2003-2009) to update and recalibrate its predictive capability<sup>74</sup> and the score was also validated in non-Caucasians<sup>75</sup>. Importantly, the score was modified where carbon monoxide diffusing capacity, serum alanine aminotransferase, and serum creatinine concentrations were no longer significantly associated with 2-year mortality, whereas patient and donor cytomegalovirus serology was associated with mortality. However, there is also literature to support an assertion that the PAM score may not be useful in all

allogeneic<sup>76,77</sup> or autologous<sup>78</sup> settings. The following is a link to an online calculator:

<http://pamscore.org/>

### **EBMT Score**

The EBMT risk score incorporates both recipient and disease variables. It evaluates five factors: age of patient, disease stage, time interval from diagnosis to transplant, donor type and donor recipient sex combinations. The current EBMT risk score is an extension of the “old” CML risk score. This scoring system explains 63% of the post-transplant outcomes in the EBMT registry<sup>79</sup>. More recently, the EBMT was re-evaluated in patients with primary or secondary myelodysplasia undergoing an allogeneic transplant where the EBMT score predicts overall survival and transplant related mortality but did not correlate with relapse risk<sup>80</sup>. Similarly, the EBMT score has utility in the autologous setting<sup>81</sup>.

### **Hematopoietic cell transplantation specific comorbidity index (HCT-CI)**

Using the Charlson Comorbidity Index<sup>82</sup> as a template, Sorror et al. re-developed this tool as a prognostic tool to better gauge post-allogeneic transplant survival outcomes – HCT-CI<sup>83</sup>. This index embraces the variables discussed in Section 2. This index has been validated and is independent of disease characteristics. Importantly, the variables that were considered in this model are predominantly physical with little to no evaluation of mental or psychosocial variables. The use of the HCT-CI allows an estimation of the transplant-related mortality (see appendix 1). The following is web link to facilitate score calculations: <http://www.hctci.org/Home/Calculator>

### **HCT-CI in clinical settings and comparisons with other scoring systems**

The HCT-CI has been evaluated and deemed prognostically useful in a variety of allogeneic transplant settings with modifications to incorporate combinations of age, remission status and performance status<sup>84-89</sup>. Further, modifications of the HCT-CI have been used in the autologous setting<sup>90-93</sup>. On reviewing our local data on 700 patients who received allogeneic HCT, HCT-CI did not appear to be associated with post-HCT outcomes<sup>94</sup>.

Others have attempted to compare the accuracy of EBMT Score and the HCT-CI. For instance, Michaelis et al., in a single centre retrospective analysis using regression modeling suggest that a modified Pre-Transplant EBMT Risk Score is superior to the HCT-CI Score in predicting overall survival and non-relapse mortality after allogeneic HCT in patients with acute myeloid leukemia<sup>95</sup>. Separately and similarly, Terwey et al. evaluated HCT-CI and modified EBMT Risk score in the adult patients with ALL within a single European center and suggests that the EBMT risk score may be preferable over the HCT-CI<sup>96</sup>.

The PAM score was compared with the HCT CI at a single institution and suggests the HCT-CI was more predictive of overall survival<sup>93</sup> but the conclusions are inconsistent<sup>97</sup>.

Taken together, there is no clear co-morbidity index that clearly embraces all aspects of recipient and/or disease variables. Moreover, the accuracy of prediction tools is likely dependent on local variables that are either known or unknown. However, *it is reasonable to adopt the HCT-CI as the default index as it is the most widely used tool for pre-HCT comorbidity assessment. The routine use of this tool would allow within-center and cross-center outcome comparisons.* Moreover, it has been adopted by the CIBMTR.

## Geriatric Assessments

In patients  $\geq 65$  years receiving chemotherapy, geriatric assessment (GA) should be used to identify vulnerabilities that are not routinely captured in oncology assessments. There are many variants of GAs with different domains. In principle, it would include domains of functional status, physical performance and falls, comorbid medical conditions, depression, social activity/support, nutritional status, and cognition.

The comprehensive geriatric assessment (CGA) include domains of functional status, cognitive function, comorbidities & geriatric syndromes, polypharmacy, psychological status, social support and nutritional status and is suggested in the practice guidelines developed by the National Comprehensive Cancer Network<sup>98</sup> as well as ASCO<sup>99</sup>. The use of such tools are able to predict adverse outcomes (including chemotherapy toxicity and mortality), which can help inform shared discussion making with the patient. The ASCO guidelines<sup>99</sup> suggest a Minimum Data Set for Practical Assessment of Vulnerabilities in Older Patients (>65 years) With Cancer. Specifically, they recommend:

1. Predict chemotherapy toxicity (if clinically applicable): Cancer and Aging Research Group (CARG) or Chemotherapy Risk Assessment Scale for High-Age Patients tools
  - a. *The CARG tool takes < 5 minutes to complete and is freely available online for use on the CARG website*
  - b. [https://www.mycarg.org/?page\\_id=934](https://www.mycarg.org/?page_id=934)
2. Estimate (noncancer) life expectancy (if clinically applicable): ePrognosis<sup>100</sup>, Project Big Life<sup>101</sup>
  - a. <https://eprognosis.ucsf.edu/leeschonberg.php>
  - b. <https://www.projectbiglife.ca/life-expectancy-home>
3. Functional assessment: instrumental activities of daily living
  - a. *The Lawton Instrumental Activities of Daily Living. (IADL) Scale is commonly used where any dependence on any task signifies impairment<sup>102</sup>.*
  - b. <https://neurotoolkit.com/lawton-iadl-scale/>
4. Comorbidity assessment: medical record review or validated tool
  - a. *See Section on HCT-CI*
5. Screening for falls, one question: how many falls or falls with an injury have you had in the previous 6 months (or since your last visit)?

- a. *A simple one-item can be useful: “How many falls have you had in the previous 6 months (or since your last visit)?”*
6. Screening for depression: Geriatric Depression Scale<sup>103</sup> or other validated tool
  - a. *GDS 15 item: a score of 5 suggests depression and requires follow-up*
  - b. <https://neurotoolkit.com/geriatric-depression-scale/>
7. Screening for cognitive impairment: Mini-Cog<sup>104</sup> or Blessed Orientation-Memory-Concentration test<sup>105</sup>
  - a. *The Mini-Cog consists of two components, a 3-item recall test for memory and a simply scored clock drawing test*
  - b. <https://mini-cog.com/mini-cog-instrument/standardized-mini-cog-instrument/>
8. Screening for malnutrition: weight loss/body mass index
  - a. *Assess for Unintentional weight loss; 10% weight loss from baseline weight); BMI <21 kg/m<sup>2</sup>.*

*This link from the Cancer and Aging Research Group provides online calculators for patient and healthcare providers for Geriatric Assessments: [https://www.mycarg.org/?page\\_id=898](https://www.mycarg.org/?page_id=898)*

The use of GA was able to identify older patients with inferior survival undergoing allogeneic HCT<sup>106</sup>. Specifically, limitations in instrumental activities of daily living (HR 2.38, 95%CI: 1.59–3.56; P<0.001), slow walk speed (HR 1.80, 95%CI: 1.14–2.83; P=0.01), high comorbidity by hematopoietic cell transplantation-specific comorbidity index (HR 1.56, 95%CI: 1.07–2.28; P=0.02), low mental health by short-form-36 mental component summary (HR 1.67, 95%CI: 1.13–2.48; P=0.01), and elevated serum C-reactive protein (HR 2.51, 95%CI: 1.54–4.09; P<0.001) were significantly associated with inferior overall survival. Further, it is notable that the 2-year overall survival was zero in patients >60 years in the presence of one abnormal Instrumental Activity of Daily Living (IADL) by Lawton, and a HCT-CI score of ≥3.

More recently, Deschler et al.<sup>107</sup> performed a concurrent GA by Up-and- Go and quality of life by the European Organization for Research and Treatment of Cancer Quality of Life Questionnaire (EORTC QLQ) C-30 assessments in addition to disease-specific data in 106 older (median age 66 years) patients undergoing allogeneic HCT. Collecting data at 4 time-points (before HCT and days +30, +100 and +180), they demonstrate that negative prognostic factors for PFS were age (HR 1.084) and Comorbidity index by HCT-CI (HR 1.13), while the negative prognostic factors for OS were age (HR 1.08), performance status by Karnofsky Index (HR 0.97), quality of life by EORTC QLQ C- 30 fatigue (HR 1.09) and GA by Up-and-Go (HR 3.26). Geriatric assessments have also been utilized the autologous HCT setting for myeloma, where impairments in geriatric domains are common even among those considered to have a good performance states<sup>108</sup> and can identify patients who are at a greater risk for morbidity<sup>109</sup>. Such studies suggest and emphasize that CGA is strongly associated with post-HCT outcomes and has a higher HR, even when other traditional patient and disease variables are factored in.



## Frailty & Functionality Assessment

The concepts that encompass frailty has been recently summarized by an excellent review article<sup>110</sup>. The authors suggest that the historic use of chronologic age, comorbidities and performance may be too simplistic to capture a full sense of vitality. They and others have suggested that the regular incorporation of frailty measurements such as CGA may be useful in evaluating fitness for HCT<sup>111</sup>.

Implementation of routine GAs is impractical, however, as it requires qualified specialists to perform these evaluations (not readily available), and the time needed to perform them is often unavailable or too-taxing for patients. A pilot prospective study of a “clinic-friendly” Frailty & Functionality assessment in routine clinical assessment in allogeneic SCT has been published and involved performing the assessment successfully in approximately 5 minutes<sup>111</sup>. It can be used to calculate an HCT “frailty” score that identify patients as fit, “pre-frail”, and frail, which has correlated with overall survival<sup>112</sup>.

### HCT Frailty Scale

| Evaluated Parameter             | Score – Normal 0, Abnormal – as indicated                         |
|---------------------------------|-------------------------------------------------------------------|
| Clinical Frailty Score $\geq 3$ | 1.5                                                               |
| IADL Score $\geq 1$ limitation  | 1                                                                 |
| Timed Up & Go Test              | 1.5                                                               |
| Grip Strength                   | 1 (Abnormal is $<16\text{kg}$ (female) and $<26\text{kg}$ (male)) |
| Self-rated Health Questionnaire | 1                                                                 |
| Fall in last 6 months           | 1                                                                 |
| Albumin level                   | 1.5                                                               |
| C-reactive protein              | 2                                                                 |

A video on how to perform this assessment is available at the following link:

<https://youtu.be/RPrCnWothIY>

A total score is calculated, with fit  $\leq 1$ , pre-frail  $>1$  and  $<5.5$ , and frail  $\geq 5.5$ . The estimated probabilities of 1-year OS in each group of frailty, were, respectively: 83.7%, 75.6%, and 52.8% ( $p < 0.001$ ) in the training cohort and 90.3%, 69.5%, and 46.2% ( $p < 0.001$ ) in the validation cohort<sup>113</sup>. Additionally, the estimated 1-year NRM of fit, pre-frail, and frail patients were 6.4%, 14.8%, and 31.2%, respectively. Frailty also did not necessarily correlate with performance status and HCT-CI score. Although, a clear cut-off score has not been established as to which patients should not be offered HCT, this should be considered an independent factor in the decision-making process for proceeding with allogeneic HCT.

Salas *et al.* externally validated their score via sixteen Spanish Institutions<sup>113</sup>. Between 2021 and 2023, 341 candidates for allogeneic HCT participated. They state: “Ninety-four (27.6%) adults were classified as fit, 203 (59.5%) as pre-frail, and 44 (12.9%) as frail. Frail patients were more likely to have a KPS  $< 90\%$  (OR 2.80,  $p < 0.01$ ) and an abnormal result of the Mini-Cog test ( $< 3$ ) (OR 8.21,

P<0.001). The probability of being frail was independent of age (continuous) (p=0.654), sex (p=0.323), and comorbidities (HCT-CI>3) (p=0.196) (multivariate binary regression analysis)". They demonstrate that 1-year OS was higher with frailty measured at the time of HCT admission; respectively, 20%, 66.8% and 78.9%. (p<0.001). Interestingly, they suggest that a pre-transplant rehabilitation (pre-hab) program may be able to modify this risk factor. In contrast, the value of frailty assessments in the autologous setting is less clear<sup>114</sup> where pre-HCT frailty did not correlate with prolonged hospitalization or overall survival. Others have reported that persistence of frailty determines poor outcomes<sup>115</sup>.

Taken together, there appears to a complex interplay between frailty, disease status and type of HCT that determines post-HCT outcomes. Additionally, it is possible that frailty is a modifiable risk factor for post-HCT outcomes with pre-habilitative interventions.

## Patient Eligibility for chimeric antigen receptor T-cell therapy (CAR-T)

The eligibility for CAR-T cells will be predominantly assessed on disease-based factors (see appropriate chapters for discussion). The principals as described in the preceding paragraphs for standard allogeneic and autologous HCT would apply. Recognizing there is a paucity of evidence, there is a sense that lower thresholds for eligibility from the perspective of patient factors may apply given that there is no high-dose therapy.

There are areas of uncertainty on whether a history of malignancy, allogeneic HCT, prior treatment with BiTE antibodies or CNS involvement represented an exclusion criteria<sup>116</sup>. The following table summarizes the eligibility criteria from large RCTs and the recommendations from EBMT/JACIE<sup>117</sup>. Our center would support these pragmatic guidelines.

| Characteristics | ELIANA (ALL Kymriah™)                                                                                                    | JULIET (CLMCL Kymriah™)                                                   | ZUMA-1 (High-grade B-cell NHL Yescarta™)                                  | EBMT recommendations | Comment                                                                                                 |
|-----------------|--------------------------------------------------------------------------------------------------------------------------|---------------------------------------------------------------------------|---------------------------------------------------------------------------|----------------------|---------------------------------------------------------------------------------------------------------|
| Age limit (NHL) | N/A                                                                                                                      | ≥ 18 years<br><br>SPC-No data are available on children < 18 years of age | ≥ 18 years<br><br>SPC-No data are available on children < 18 years of age | No upper age limit   | Decision should be based on physical condition rather than age                                          |
| Age limit (ALL) | 'Age 3 years at the time of screening to age 21 years at the time of initial diagnosis'<br><br>SPC-up to 25 years of age | N/A                                                                       | N/A                                                                       | **Follow SPC         | Ability to collect sufficient cells by apheresis can be a limiting factor in infants and small children |

|                                                                     |                                                                                                                                                |                                                                                                                                                                                                                                         |                                                                                                                                                                                        |                                                                                                                                                                                  |                                                                                        |
|---------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------|
| ECOG PS Performance Status                                          | Karnofsky (age ≥ 16 years) or Lansky (age < 16 years) PS ≥ 50 at screening                                                                     | ECOG PS of either 0 or 1 at screening                                                                                                                                                                                                   | ECOG PS of 0 or 1                                                                                                                                                                      | >2 not recommended<br>Note, however, that real-world data with Yescarta™ included patients with ECOG PS > 2                                                                      | Prognosis may be less poor in the decline in PS is due to active disease               |
| History of malignancy                                               | No prior malignancy, except carcinoma <i>in situ</i> of the skin or cervix treated with curative intent and with no evidence of active disease | No previous or concurrent malignancy except adequately treated BCC or SCC, <i>in situ</i> cancer of the breast or cervix treated and without recurrence for 3 years, primary malignancy resected and in remission for more than 5 years | No history of malignancy other than nonmelanoma skin cancer or carcinoma <i>in situ</i> (e.g. cervix, bladder, breast) or follicular lymphoma unless disease free for at least 3 years | Absence of history of malignancy other than carcinoma <i>in situ</i> (e.g. cervix, bladder, breast) unless disease-free and off therapy for at least 3 years                     |                                                                                        |
| Prior allo-HCT                                                      | Not excluded; however, excluded if grade II-IV acute or extensive chronic GvHD                                                                 | Excluded                                                                                                                                                                                                                                | Excluded                                                                                                                                                                               | Not a contraindication                                                                                                                                                           | Active GvHD is listed as a reason to delay treatment in the Kymriah™ and Yescarta™ SPC |
| Prior anti-CD19/ anti-CD3 BiTE antibodies or any other CD19 therapy | Excluded<br>Not a contraindication as per SPC                                                                                                  | Excluded                                                                                                                                                                                                                                | Excluded if prior CD19 targeted therapy                                                                                                                                                | Not a contraindication                                                                                                                                                           |                                                                                        |
| Previous CAR-T-cell therapy                                         | Not applicable in trials<br>Not in SPC                                                                                                         | Not applicable in trials<br>Not in SPC                                                                                                                                                                                                  | Excluded                                                                                                                                                                               | Not a contraindication                                                                                                                                                           | Further Car T-cell therapy outside of clinical trials is to be avoided                 |
| History of autoimmune disease                                       | Not an exclusion criterion                                                                                                                     | Not an exclusion criterion                                                                                                                                                                                                              | Not an exclusion criterion                                                                                                                                                             | Not recommended in active autoimmune disease resulting in end-organ injury or requiring systemic immune-suppression or systemic disease-modifying agents within the last 2 years | Individualized risk-benefit assessment required                                        |
| Current systemic immune-suppressive treatment                       | Any GvHD therapy must be stopped more than 4 weeks prior to enrollment to confirm that                                                         | Any immunosuppressive medication must be stopped more than 4 weeks prior to enrollment                                                                                                                                                  | Any immunosuppressive medication must be stopped more than 4 weeks to enrollment                                                                                                       | Contraindication                                                                                                                                                                 | Intermittent topical, inhaled or intranasal corticosteroids are allowed                |

|                                                                    |                                                                                                                                                              |                                                                                                                                                                                    |                                                                                                                                                                               |                                                                             |                                                                               |
|--------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------|-------------------------------------------------------------------------------|
|                                                                    | GvHD recurrence in not observed                                                                                                                              |                                                                                                                                                                                    |                                                                                                                                                                               |                                                                             |                                                                               |
| Existing or suspected fungal, bacterial, viral, or other infection | Active or latent HBV or HCV (test within 8 weeks of screening) or any uncontrolled infection at screening                                                    | Uncontrolled active or latent HBV or active HCV; Uncontrolled acute life-threatening bacterial, viral or fungal infection (e.g. blood cultures positive < 72 h prior to screening) | Known history of HIV, HBV (HepBs Ag positive) or HCV (anti-HCV); Clinically significant active infection, or currently receiving IV antibodies or within 7 days of enrollment | Relative contra-indication; individualized risk-benefit assessment required | Active infection should be controlled and on treatment prior to leukapheresis |
| History of CNS disease                                             | CNS involvement by malignancy defined as CNS-3 as per NCCN guidelines excluded; however, those with history of effectively treated CNS disease were eligible | Active CNS involvement by malignancy excluded                                                                                                                                      | Subjects with detectable CSF malignant cells, or brain metastases, or with history of CSF malignant cells or brain metastases excluded                                        | Relative contra indication; individualized risk-benefit assessment required | Caution required as higher risk of neurological toxicity                      |

ALL: acute lymphoblastic leukemia; DLBCL: diffuse large B-cell lymphoma; NHL: non-Hodgkin lymphoma; EBMT: European Society for Blood and Marrow Transplantation; N/A: not available; SPC: summary of product characteristics; ECOG: Eastern Cooperative Oncology Group; PS: performance status; BCC: basal cell carcinoma; SCC: squamous cell carcinoma; allo-HCT: allogeneic hematopoietic cell transplantation; GvHD: graft-versus-host disease; BiTE: bispecific monoclonal antibodies; CAR: chimeric antigen receptor; HBV: hepatitis B virus; HCV: hepatitis C virus; HIV: human immunodeficiency virus; CNS: central nervous system; NCCN: National Comprehensive Cancer Network.

## Psychosocial Assessment

Psychosocial assessment(s) forms an important piece in pre-HCT evaluation, performed by different clinicians –physicians, psychologists, social workers and nurses. A dedicated program and staff is preferred to ensure consistency and expertise.

The following observations could suggest that measures (complex interventions) that broadly support and improve psychosocial health may lead to improve post-HCT psychosocial, patient reported outcomes as well as traditional medical post-HCT outcomes (e.g. survival). A recent cross-sectional sample of 351 HCT recipients at NCI-designated centers suggest that only 14% of patients presenting with pre-HCT psychosocial evaluation utilized psychotherapy services – patients who utilized services reported lower levels of distress, depression and anxiety<sup>118</sup>.

## Psychosocial Uncertainties

Foster et al. performed a survey of HCT professionals in 2006 using 17 case vignettes each representing a different psychosocial issue to which respondents indicated whether or not they would recommend proceeding with allogeneic HCT. In six vignettes, at least 64% indicated do not proceed: suicidal ideation (86.8%) uses addictive illicit drugs (81.7%), history of noncompliance (80.5%), no lay caregiver (69.3%), alcoholic (64.8%), and mild dementia/Alzheimer's (64.4%). In 10 vignettes, at least 73% indicated proceed. On four vignettes, professional subgroups differed in their recommendation on whether or not to proceed with allogeneic BMT<sup>119</sup>.

Interestingly, a follow-up survey of 62 chairpersons of the hospital ethics committees (HEC) with an accredited HCT program elicited whether they would recommend HCT in the 6 scenarios (as above) where the majority HCT clinicians would not. Opinions regarding transplant differed in one case only, in a patient with mild dementia; 27% of HEC chairpersons recommended not proceeding with BMT, which was significantly lower than that of nurses (68%,  $P < 0.001$ ), physicians (63.5%,  $P < 0.001$ ) and social workers (51.9%,  $P = 0.05$ )<sup>120</sup>.

## Psychosocial pre-HSCT Assessment Tools

Although a Gestalt approach to assessment is feasible, a formal validated tool is preferred. Indeed, there are numerous general screening tools including distress screening tools<sup>121-123</sup>, but may not be specific to the HCT population.

Garcia et al. developed a psychosocial structured interview to assess candidates for HCT with the interview averaging 50 minutes to complete<sup>124</sup>. In the absence of a comprehensive and validated tool, we suggest using this structured interview tool to ensure consistent history taking. The elements garnered from this structured interview could potentially be utilized to subsequently populate other scales and questionnaires.

## Psychosocial assessment of candidates for transplantation (PACT)

This scale captures information in four domains (social support, psychological health, lifestyle factors, and patients understanding of the transplant process) with eight subscales, each rated on a 5-point scale<sup>125</sup>. This scale was originally developed for clinical decision-making in psychosocial screening of organ transplant (heart and liver) candidates<sup>126</sup>. The use of PACT rating at a single institution study was associated with non-relapse mortality (HR 0.82 per point increase [95% CI, 0.69-0.98],  $P = 0.03$ ), but not with overall survival (HR 0.91 [95% CI, 0.79-1.05],  $P = 0.18$ ). There was no association between final PACT rating and neutrophil or platelet engraftment, acute or chronic GVHD, or relapse<sup>127</sup>. In contrast, data from Japan suggests that lower PACT scores in the domain of compliance with medications and medical advice were significantly associated with poorer OS (HR = 1.75,  $P = 0.03$ )<sup>128</sup>.

### **Patient Health Questionnaire (PHQ)**

In a small randomized study<sup>129</sup>, the Patient Health Questionnaire (PHQ) was used to assess for depressive disorders, anxiety, substance abuse, and problems in occupational or interpersonal functioning (functional disruption) and was provided to patients before meeting with their medical provider (n = 50; experimental group) or afterwards (n = 51; control group). The prevalence of clinically significant depression (21%), anxiety (14%), or suicidal ideation (8%) did not differ between the 2 groups. Patients in the experimental group were likely to have discussion of psychological symptoms than the control group (68% versus 49%,  $P = .05$ ). Medical providers were significantly more satisfied with the management of psychological issues for the experimental group ( $P < .001$ ). Patients with depression or anxiety were significantly more likely to prefer the PHQ be used at future visits ( $P = .02$  and  $P = .001$ , respectively).

### **The Transplant Evaluation Rating Scale (TERS)**

The TERS score is a compilation of scores on 10 weighted factors: psychiatric history of Axis I disorder, psychiatric history of Axis II disorder, substance abuse, health behaviors, compliance, quality of family/social support, history of coping, coping with disease and treatment, quality of affect, and mental status. The TERS was prospectively used at a single institution<sup>130</sup>; where patients in the high-risk TERS group had significantly longer hospital stays during the first 180 days and 1 year post-allogeneic HSCT compared with the low-risk group (16 vs 13 and 21 vs 16 days;  $P = .05$  and  $.02$ , respectively). In their multivariable analysis, intermediate- and high-risk TERS scores predicted for inferior OS, similar DFS, and higher NRM compared with low-risk TERS score. In a subset analysis of patients with low/intermediate risk per Disease Risk Index, multivariable analysis showed that high- and intermediate-risk TERS scores predicted for significantly worse OS, worse DFS, higher NRM, and similar relapse rates compared with low-risk TERS score. Additionally, poor TERS scores have also been associated with higher readmission rates<sup>109</sup>.

### **Distress**

Distress is a complex term that is utilized to embrace multiple aspects mental health states. This broad concept has been evaluated in the context of HCT using different scales (validated and unvalidated). Consequently, it is challenging to given firm conclusions on its association with post-HCT outcomes.

Cancer and treatment specific distress pre-allogeneic HCT is associated with post-traumatic stress disorder (PTSD). Specifically, uncertainty, appearance and sexuality as well as health burden were concepts associated with PTSD<sup>131</sup>. It has been estimated that PTSD occurs in 3.3% of HCT recipients; interestingly PTSD in HCT recipient caregivers is estimated at 6.6% ( $p=0.02$ )<sup>132</sup>. Pre-HCT psychological distress as measured with an unvalidated Likert-like scale was unrelated to survival in a single centered study<sup>133</sup>.

Taken together, the presence of pre-transplant patient distress may have psychologic consequences post-transplant but does not clearly influence survival. Moreover, the management of distress of peri-



HCT is not well-defined with a recent systematic review suggesting psychological interventions (cognitive behavioral or emotional processing methods) may provide some benefit in alleviating distress in HCT but conclusions remain tentative in light of methodological limitations and risk of bias in their included studies<sup>134</sup>.

Interestingly, a recent RCT randomly assigning 160 patients undergoing HCT to receive inpatient palliative care integrated with transplant care (n = 81) or transplant care alone (n=79) suggest that intervention participants reported lower depression symptoms, lower PTSD symptoms but no difference in HRQOL or anxiety<sup>135</sup>. The intervention was composed of a palliative care clinician reviewing care at least twice a week broadly addressing topics of nausea, pain and mucositis, fatigue, insomnia, bowel problems, and psychological distress.

### **Anxiety and Depression**

A recent prospective observational study in six German allogeneic HCT centers evaluated the prevalence of depression and anxiety using the Hospital Anxiety and Depression Scale (HADS), comparing it to gender- matched controls (reference). They demonstrate that the rates of HADS-defined depression increased from 12% before HCT to up to 30% at 5 years after HCT, while anxiety was highest before HCT and settles to control baseline after HCT.

Pre-HCT clinical depression is associated with lower overall survival and higher acute GVHD among allogeneic transplant recipients<sup>136</sup>. Further, it is associated with fewer days alive and out of hospital within the 1<sup>st</sup> 100 days after autologous and allogeneic setting. This could suggest routine screening for depression and providing pre-emptive pharmacologic and/or psychologic therapies to mitigate this risk factor.

### **Non-Compliance**

Compliance has been defined as the extent to which a person's behavior (in terms of medication, following diets, or executing lifestyle changes) coincides with medical or health advice<sup>137</sup>. The prevalence on non-compliance is unknown in the HCT population<sup>138</sup>. Rate of adherence to oral medications ranged from 33% to 94.7%<sup>139,140</sup> where it has been associated with increased risk of infections in the pediatric HCT setting<sup>141</sup>. Overall, there is a paucity of research that evaluates the consequences of noncompliance in adult HCT patients, nor the predictive value of pre-transplant compliance in determining post-transplant behavior. Further, the impact of compliance on therapeutic outcomes and the interventions that effectively increase compliance are all unknown.

Mumby et al. in a study of 151 autologous HCT patients suggests 80% of patients were deemed non-compliant with an aspect of the transplant on  $\geq 1$  day<sup>142</sup>. Non-compliance was defined as refusal of oral hygiene, prescribed exercise programs, oral nutrition and/or prescribed medications. In a multivariate analysis, the predictors of non-compliance in their cohort of patients were 1) gender - men, 2) presence of depression, 3) global distress and 4) nausea and vomiting severity. Interestingly,

a small and older study of 92 HCT patients did not identify compliance as predictive of post-HCT outcomes<sup>143</sup>.

More recently, a single centre study from Japan retrospectively assigning the PACT scale to 119 HCT recipients demonstrate that lower PACT scores in the domain of compliance with medications and medical advice were significantly associated with poorer OS (HR = 1.75, P = 0.03)<sup>128</sup>. Similarly, the Stanford Integrated Psychosocial Assessment for Transplantation (SIPAT) scale was able to predict nonadherence defined as at least 1 life-threatening nonadherence event in the first 6 months post-transplant<sup>144</sup>.

It has been suggested that the following considerations may improve compliance<sup>138</sup>: 1) Provision of clear and consistent information with specific information on why consistent compliance is beneficial, 2) simplify treatment, 3) prioritize environmental precautions and health behaviors, 4) suggest ways to assist with forgetfulness, and 5) tailor the regimen as much as possible to the lifestyle of the patient.

*Due to the paucity of data, non-compliance should not be an absolute contraindication to HCT.*

### **Substance Abuse**

Lifetime substance abuse appears to be associated with adverse outcomes post-HCT<sup>145</sup>. In a single center case –control study, Chang et al. identified 17 individuals with lifetime substance abuse where with alcohol (71%), marijuana (30%), and opiates (30%) were identified as the principal substances of abuse. They identified controls, matching for disease and stage, type of transplant, pre-HCT conditioning regimen, and age. Survival analysis demonstrated reduced survival times for patients with substance abuse (p = .0022)<sup>146</sup> with 15 of 17 patients dying within the first year. Interestingly, a follow-up study did show this association<sup>147</sup>. Graf et al. suggest that a history of alcohol use disorder was associated with non-relapse mortality (HR 2.17; p=0.004) in an analysis of 754 patients undergoing autologous HCT for lymphoma at a single centre<sup>148</sup>.

*Due to the paucity of data, substance abuse should not be an absolute contraindication to HCT.*

### **Other Psychological Functioning and Coping Styles**

Data on aspects of psychological functioning/coping style is sparse. However, a recent study evaluating 332 recipients of HCT using measures of illness perceptions (beliefs about cancer consequences and course, personal and treatment control over cancer, and understanding of one's cancer) suggest that greater personal and treatment control was associated with a healthier diet and reported greater well-being. Further, a better understanding of their cancer also was associated with a healthier diet, less depression, less anxiety, and greater well-being<sup>149</sup>. In addition, greater personal resilience (composite variable of self-esteem, mastery, and optimism) may promote better psychologic adjustment via improvements in depressive symptoms with decreased use of

maladaptive meaning-making (searching for a reason for one's illness) was associated with less PTSD symptoms<sup>150</sup>. Similar findings were noted by Barata *et al.*<sup>151</sup>.

### **Financial/Socioeconomic Status**

The socioeconomic status (SES) of the recipient is associated with poor HCT outcomes due to multiple interrelated factors<sup>152,153</sup>. Specifically, lower attained education was associated with increased distress ( $P = .002$ ), lower income was related to worse physical functioning ( $P = .005$ ) and increased distress ( $P = .008$ ), lack of employment before transplantation was associated with worse physical functioning ( $P < .01$ )<sup>154</sup>.

Further, low SES is also associated with higher risks of all-cause mortality (hazard ratio (HR) 1.98,  $P=0.012$ ) and non-relapse mortality (NRM) (HR 2.22,  $P=0.028$ ), but similar risks of relapse mortality (HR 1.01,  $P=0.97$ ) compared with high SES patients. A trend toward better survival and lower NRM for high SES patients with no chronic GVHD was observed; low SES patients without GVHD had similar survival as patients with chronic GVHD<sup>155</sup>. Similar results were noted by Silla *et al*<sup>156</sup>. In contrast, Hamilton *et al.* suggest that SES was not associated with chronic GVHD outcomes<sup>157</sup>.

Interestingly, Knight *et al.* suggests that low SES effects are modulated through upregulation of conserved transcriptional response to adversity (CTRA)<sup>158</sup>. From a psychologic perspective, it has been suggested that the effects of “objective” SES is modulated through the individual’s “subjective” SES<sup>159</sup>.

The influence of SES is less clear in the autologous setting<sup>160,161</sup> and it is likely that other patient and/or disease factors are more important in this setting.

### **Caregiver Considerations**

There is an increasing reliance on caregivers as HCT programs shift towards an outpatient setting. In general, outpatient HCT are performed when there is a consistent caregiver. This invariably shifts the burden of care on to the caregiver. Indeed, caregivers are emotionally vulnerable than patients before HCT where cancer-related distress was the strongest correlate of anxiety and depression in both patients and caregivers<sup>162</sup>. Acknowledging, understanding and supporting the caregiving role would be prudent from the perspective of the HCT program.

The consistent presence of a caregiver is independently associated with superior post-allogeneic HCT overall survival<sup>125,143,163</sup>, but the results are mixed<sup>164,165</sup>. The optimal caregiver(s) and the quality of the caregiver remains unclear, however there is evidence to suggest the quality of the caregiver may matter more than caregiver consistent presence<sup>166</sup>. Caregivers commonly report elevated distress pre-HCT and both patient and caregiver distress were associated with patient HRQOL, with patients’ physical well-being a significant contributor to caregiver well-being<sup>167</sup>. Interestingly, patient’s perception of over-benefiting within a dyadic relationship was associated with patient distress, but not

the patient’s self-perceived burden<sup>168</sup>. Separately, there is evidence to suggest that unmarried status is associated with worse sleep in the allogeneic setting<sup>154</sup>.Overall, there is a paucity of evidence to guide practice as summarized by a systematic review<sup>169</sup>.

## Integrating Psychosocial Factors

Due to lack of definitive evidence, none of the psychosocial factors discussed above represent absolute contraindications to HCT. However, it needs to be recognized that there is also a lack of safety data for patients who exhibit severe psychiatric illness (e.g. major depression, suicidal ideation/planning, psychotic illness with delusions/hallucinations, etc.), active abuse of alcohol or street drugs, or those who demonstrate profound degrees of non-compliance. Prior to accepting that such patients are eligible to proceed with HCT, they require early referral (ideally at diagnosis of malignancy) to psychology, social work, or psychiatry as appropriate. If the patient does not demonstrate engagement and compliance with psychosocial services, or if psychosocial concerns are not stabilized, transplant may be deferred in order to prioritize patient safety. This would be considered especially when psychosocial factors severely impair functioning and/or adherence to treatment plans or place the patient at immediate safety risk (e.g., actively psychotic, suicidal, substance dependent, extreme poverty, high degree of family conflict).

**Table 1:** Optimal physiologic parameters for transplant eligibility<sup>1</sup>

| Physiologic Parameters                          | Optimal “Cut-Offs” |
|-------------------------------------------------|--------------------|
| <b>Age and Performance Status</b>               |                    |
| Age (years)                                     | ≤65                |
| KPS                                             | >60                |
| <b>Pulmonary</b>                                |                    |
| FEV <sub>1</sub> (% of predicted value)         | >60                |
| DLCO (% of predicted value)                     | >60                |
| <b>Cardiac</b>                                  |                    |
| LVEF (%)                                        | >45                |
| Heart rhythm                                    | Normal             |
| <b>Hepatic</b>                                  |                    |
| Serum Bilirubin                                 | <2 x normal        |
| ALT/AST/ALP                                     | <2 x normal        |
| <b>Renal</b>                                    |                    |
| Serum Creatinine                                | <2 x normal        |
| <b>Second active malignancy</b>                 | Absent             |
| <b>Pregnancy test</b>                           | Negative           |
| <b>Uncontrolled Infections including dental</b> | Absent             |

KPS=Karnofsky performance Status; FEV1=force expiratory volume in 1 second; DLCO=diffusion capacity; ALT/AST/ALP=alanine aminotransferase/aspartate aminotransferase/alkaline phosphatase; LVEF= Left ventricular ejection fraction

**Table 2: Hematopoietic cell transplantation specific comorbidity index (HCT-CI)<sup>83</sup>**

| <b>Co-morbidity</b>           | <b>Definition/compartments</b>                                                                                                   | <b>Yes</b>                                                                                                     | <b>Score</b>                   |
|-------------------------------|----------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------|--------------------------------|
| 1. Arrhythmia                 | -Atrial fibrillation*<br>-Atrial flutter*<br>-Sick sinus syndrome*<br>-Ventricular arrhythmia*                                   | <input type="checkbox"/><br><input type="checkbox"/><br><input type="checkbox"/><br><input type="checkbox"/> } | <input type="text" value="1"/> |
| 2. Cardiovascular             | -Coronary artery disease*<br>-Congestive heart failure*<br>-Myocardial infarction*<br>-Ejection fraction ≤50%§                   | <input type="checkbox"/><br><input type="checkbox"/><br><input type="checkbox"/><br><input type="checkbox"/> } | <input type="text" value="1"/> |
| 3. Inflammatory bowel disease | -Crohn's disease*<br>-Ulcerative colitis*                                                                                        | <input type="checkbox"/><br><input type="checkbox"/> }                                                         | <input type="text" value="1"/> |
| 4. Diabetes                   | -Treated with insulin or oral hypoglycemic drugs§                                                                                | →                                                                                                              | <input type="text" value="1"/> |
| 5. Cerebro-vascular           | -Transient ischemic attacks*<br>-Cerebro-vascular ischemic or hemorrhagic stroke*                                                | <input type="checkbox"/><br><input type="checkbox"/> }                                                         | <input type="text" value="1"/> |
| 6. Depression/anxiety         | -Requiring psychological consultation and/or specific treatments§                                                                | →                                                                                                              | <input type="text" value="1"/> |
| 7. Hepatic - mild             | -Chronic hepatitis§<br>-Bilirubin >ULN- 1.5 X ULN§<br>-AST/ALT >ULN- 2.5 X ULN§                                                  | <input type="checkbox"/><br><input type="checkbox"/><br><input type="checkbox"/> }                             | <input type="text" value="1"/> |
| 8. Obesity                    | -Body mass index >35 (adults)§<br>-Body mass index-for-age ≥95% percentile (children)§                                           | <input type="checkbox"/><br><input type="checkbox"/> }                                                         | <input type="text" value="1"/> |
| 9. Infection                  | -Requiring anti-microbial treatment before, during, and after the start of conditioning§                                         | →                                                                                                              | <input type="text" value="1"/> |
| 10. Rheumatologic             | -Requiring Treatment*                                                                                                            | →                                                                                                              | <input type="text" value="2"/> |
| 11. Peptic ulcer              | -Confirmed by endoscopy and requiring treatment*                                                                                 | →                                                                                                              | <input type="text" value="2"/> |
| 12. Renal                     | -Serum creatinine >2mg/dl (or >177 μmol/L)§<br>-On dialysis§<br>-Prior renal transplantation*                                    | <input type="checkbox"/><br><input type="checkbox"/><br><input type="checkbox"/> }                             | <input type="text" value="2"/> |
| 13. Pulmonary - Moderate      | -DLco corrected for hemoglobin 66-80% of predicted§<br>-FEV1 66-80% of predicted§<br>-Dyspnea on slight activity§                | <input type="checkbox"/><br><input type="checkbox"/><br><input type="checkbox"/> }                             | <input type="text" value="2"/> |
| 14. Pulmonary - Severe        | -DLco corrected for hemoglobin ≤ 65% of predicted§<br>-FEV1 ≤ 65% of predicted§<br>-Dyspnea at rest or requiring oxygen therapy§ | <input type="checkbox"/><br><input type="checkbox"/><br><input type="checkbox"/> }                             | <input type="text" value="3"/> |
| 15. Heart valve disease       | -Except asymptomatic mitral valve prolapse§                                                                                      | →                                                                                                              | <input type="text" value="3"/> |
| 16. Prior solid malignancy    | -Treated with surgery, chemotherapy, and/or radiotherapy, excluding non-melanoma skin cancer*                                    | →                                                                                                              | <input type="text" value="3"/> |
| 17. Hepatic - moderate/severe | -Liver cirrhosis§<br>-Bilirubin > 1.5 X ULN§<br>-AST/ALT > 2.5 X ULN§                                                            | <input type="checkbox"/><br><input type="checkbox"/><br><input type="checkbox"/> }                             | <input type="text" value="3"/> |
| <b>Total Score</b>            |                                                                                                                                  |                                                                                                                | <input type="text"/>           |

\*Diagnosed at any time in the patient's past history

§Detected at the time of pretransplant assessment - ULN indicates upper limit of normal; DLco, diffusion capacity of carbon monoxide; FEV1, forced expiratory volume in one second; AST, aspartate aminotransferase; and ALT, alanine aminotransferase

The HCT-CI is able to classify patients into three risk groups:

|       | Non-Relapse Mortality |          | Overall Survival |          |
|-------|-----------------------|----------|------------------|----------|
|       | HR (95% CI)           | 2-year % | HR (95% CI)      | 2-year % |
| Score |                       |          |                  |          |
| 0     | 1.0                   | 14       | 1.0              | 71       |
| 1 - 2 | 1.42 (0.8-2.7)        | 21       | 1.31 (0.8-2.0)   | 60       |
| >3    | 3.54 (2.0-6.3)        | 41       | 2.69 (1.8-4.1)   | 34       |



**Table 4:** Psychosocial Assessment Interview of Candidates for Hematopoietic Stem cell Transplantation (PAIC-HSCT)<sup>124</sup>.

## 1. IDENTIFICATION, SOCIAL AND DEMOGRAPHIC INFORMATION

- 1.1. Name: \_\_\_\_\_
- 1.2. Gender: \_\_\_\_\_ (1-Male / 2-Female)
- 1.3. Date of birth: \_\_\_\_\_ / \_\_\_\_\_ / \_\_\_\_\_
- 1.4. Marital status: \_\_\_\_\_ (1-Single / 2-Married / 3-Widowed / 4-Divorced)
- 1.5. Instruction level: \_\_\_\_\_ years
- 1.6. Do you have any difficulties to read? \_\_\_\_\_ (1-Yes / 2-No)
- 1.7. Occupation: \_\_\_\_\_
- 1.8. Current job status: \_\_\_\_\_ (1-Employed / 2-Unemployed / 3-Retired / 4-Sick leave)
- 1.9. Job contract: \_\_\_\_\_ (1-Formal / 2-Unofficial)
- 1.10. What is the longest period you remained in a job? \_\_\_\_\_ years
- 1.11. Monthly income: \_\_\_\_\_
- 1.12. Ethnicity: \_\_\_\_\_ (1-Caucasian, 2-Black, 3-Asian, 4-Brown)
- 1.13. Religion: 1. \_\_\_\_\_ 2. None
- 1.14. How often do you visit temples or participate at church meetings? \_\_\_\_\_ times/month
- 1.15. Children: \_\_\_\_\_
- 1.16. Home address: \_\_\_\_\_
- 1.17. Telephone number: \_\_\_\_\_
- 1.18. Person who takes care of you: \_\_\_\_\_
- 1.19. Your family relationship with this person: \_\_\_\_\_
- 1.20. His/her telephone number: \_\_\_\_\_
- 1.21. Donor: \_\_\_\_\_
- 1.22. Family relationship with your donor: 1. \_\_\_\_\_ 2. None

## 2. COMPREHENSION OF THE ILLNESS

- 2.1. How have you discovered you are sick?
- 2.2. Do you know what your illness is? ☐Y ☐N ☐Partially
- 2.3. Do you know any possible causes of this illness? ☐Y ☐N ☐Partially
- 2.4. Do you know consequences and treatments of your illness? ☐Y ☐N ☐Partially
- 2.5. Have you got any previous medical treatment?
- 2.6. What medicines do you currently take? ☐Y ☐N ☐Partially

### 3. COMPREHENSION OF THE TRANSPLANTATION

- 3.1. What is bone marrow ☐Y ☐N ☐Partially
- 3.2. What is a hematopoietic stem cell transplant and how can it help your health? ☐Y ☐N ☐Partially
- 3.3. Considering your clinical condition, what are the advantages and disadvantages of the HSCT? ☐Y ☐N ☐Partially
- 3.4. Do you know why you have been chosen to undergo a HSCT? ☐Y ☐N ☐Partially
- 3.5. Can you tell me what you know about what will happen during the transplant once you are in hospital? ☐Y ☐N ☐Partially
- 3.6. Can you tell me what you know about the period following your discharge from hospital? ☐Y ☐N ☐Partially
- 3.7. What are the possible side effects of the medicines used during the transplantation? ☐Y ☐N ☐Partially
- 3.8. Do you think you understand all the risks of the treatment you are going to go through? ☐Y ☐N ☐Partially
- 3.9. What are the possible complications and late effects of a HSCT? ☐Y ☐N ☐Partially
- 3.10. Did you have the chance to meet somebody who has already undergone a HSCT? ☐Y ☐N ☐Partially
- 3.11. How was this meeting?
- 3.12. Do you believe you have received enough information to make a decision about HSCT? ☐Y ☐N ☐Partially

### 4. MEDICAL COMPLIANCE

- 4.1. In previous medical treatments did you miss consultations? Did you refuse to take prescribed drugs or did you stop taking them without medical consent? Did you refuse to follow medical advices or restrictions? Did you refuse to do any exams prescribed by your doctor? ☐Y ☐N ☐Partially
- 4.2. Have you ever interrupted a medical treatment before the scheduled end? ☐Y ☐N ☐Partially

(Questions 4.3 - 4.5 are about the pre-transplant procedures)

- 4.3. Did you miss any consultations with your doctor? If yes, tell us why. ☐Y ☐N
- 4.4. Did you refuse to follow medical advices or restrictions or did you refuse to do any exams prescribed by your doctor? If yes, tell us why. ☐Y ☐N ☐Partially
- 4.5. Did you refuse to attend the psychosocial assessment? If yes, tell us why. ☐Y ☐N ☐Partially

(Question 4.6 should be answered by the interviewer)

- 4.6. Is the patient against the psychosocial evaluation? ☐Y ☐N ☐Partially

## 5. LIFE STYLE

- 5.1. Do you practice physical exercises regularly or did you use to do it before the illness? ☐Y ☐N ☐Partially
- 5.2. Do you have a healthy eating pattern? ☐Y ☐N ☐Partially
- 5.3. BMI<sup>1</sup>: \_\_\_\_\_ Weight: \_\_\_\_\_ kg Height: \_\_\_\_\_ m
- 5.4. Do you usually have spare moments or meetings with friends? ☐Y ☐N ☐Partially
- 5.5. Are you satisfied with your sexual performance? ☐Y ☐N ☐Partially
- 5.6. Has the illness affected your sexual performance? ☐Y ☐N ☐Partially
- 5.7. After the transplant you will need to change your way of living. Do you agree with this? ☐Y ☐N ☐Partially
- 5.8. Are you satisfied with the your quality of life ☐Y ☐N ☐Partially

### Smoking:

- 5.1. Do you smoke? (if you have stopped, please answer the next items 5.2 and 5.4) ☐Y ☐N
- 5.2. How long did you smoke? \_\_\_\_\_ years
- 5.3. How long did you stop smoking? \_\_\_\_\_ years
- 5.4. The fact of being ill has affected your decision of stopping smoking? ☐Y ☐N ☐Partially

### Alcoholism:

- 5.5. Have you ever felt you should cut down on your drinking? ☐Y ☐N
- 5.6. Have people annoyed you by criticizing your drinking? ☐Y ☐N
- 5.7. Ever felt bad or guilty about your drinking? ☐Y ☐N
- 5.8. Have you ever had a drink first thing in the morning to steady  
Your nerves or get rid of a hangover? ☐Y ☐N

## 6. COPING STRATEGIES

- 6.1. How do you usually behave in difficult situations?
- 6.2. How did you face the fact of being sick when your illness was diagnosed?
- 6.3. How have you changed your life due to the illness?
- 6.4. How do you face the fact that you need to undergo to transplantation?

As you respond to each of the statements below, please keep in mind the moment when your doctor told you would need to undergo to a Hematopoietic Stem Cell Transplantation. Read each statement carefully and indicate to what extent you used it in the situation, by putting on a circle in front of the response. (0 – Does not apply or not used; 1 – Used somewhat; 2 – Used quite a bit; 3 – Used a great deal)

- |                                                                                                       |   |   |   |   |
|-------------------------------------------------------------------------------------------------------|---|---|---|---|
| 6.5. I took it out on other people .....                                                              | 0 | 1 | 2 | 3 |
| 6.6. I expressed anger to the person(s) who caused the problem .....                                  | 0 | 1 | 2 | 3 |
| 6.7. I made light of the situation and refused to get too serious about it .....                      | 0 | 1 | 2 | 3 |
| 6.8. I refused to believe that it had happened .....                                                  | 0 | 1 | 2 | 3 |
| 6.9. I tried to keep my feelings to myself .....                                                      | 0 | 1 | 2 | 3 |
| 6.10. I looked for the silver lining, so to speak; I tried to look on the bright side of things ..... | 0 | 1 | 2 | 3 |
| 6.11. I asked a relative or friend I respected for advice .....                                       | 0 | 1 | 2 | 3 |
| 6.12. I talked to someone about how I was feeling .....                                               | 0 | 1 | 2 | 3 |
| 6.13. I made a promise to myself that things would be different next time .....                       | 0 | 1 | 2 | 3 |
| 6.14. I criticized or lectured myself .....                                                           | 0 | 1 | 2 | 3 |
| 6.15. I wished that the situation would go away or somehow be over with .....                         | 0 | 1 | 2 | 3 |
| 6.16. I fantasized or wished about how things could turn out .....                                    | 0 | 1 | 2 | 3 |
| 6.17. I knew what had to be done, so I doubled my efforts to make things work .....                   | 0 | 1 | 2 | 3 |
| 6.18. I am making a plan of action and following it .....                                             | 0 | 1 | 2 | 3 |
| 6.19. I rediscovered what is important in life .....                                                  | 0 | 1 | 2 | 3 |
| 6.20. I changed or grew as a person in a good way .....                                               | 0 | 1 | 2 | 3 |

## 7. MENTAL STATUS EXAMINATION

- 7.1. Memory disorders ☐Y ☐N \_\_\_\_\_
- 7.2. Attention or concentration disorders ☐Y ☐N \_\_\_\_\_
- 7.3. Sleep disorders ☐Y ☐N \_\_\_\_\_
- 7.4. Appetite disorders ☐Y ☐N \_\_\_\_\_
- 7.5. Energy level change ☐Y ☐N \_\_\_\_\_
- 7.6. Loss of interest in activities ☐Y ☐N \_\_\_\_\_
- 7.7. Panic attack ☐Y ☐N \_\_\_\_\_
- 7.8. Speech disturbance ☐Y ☐N \_\_\_\_\_
- 7.9. Impulsiveness ☐Y ☐N \_\_\_\_\_

### 7.10. BPRS (Brief Psychiatric Rating Scale)<sup>2</sup> This form consists of 18 symptom constructs, each to be rated in a 7-point scale of severity ranging from 'not present' to 'extremely severe'. If a specific symptom is not rated, mark 'NA' (not assessed). Circle the number headed by the term that best describes the patient's present condition

|                                  |                                                           |
|----------------------------------|-----------------------------------------------------------|
| Somatic concern .....            | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Anxiety.....                     | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Emotional Withdrawal .....       | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Conceptual disorganization ..... | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Guilt .....                      | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Tension.....                     | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Mannerisms and posturing .....   | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Grandiosity .....                | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Depression.....                  | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Hostility.....                   | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Suspiciousness.....              | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Hallucinations .....             | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Motor retardation .....          | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Uncooperativeness .....          | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Unusual thought content .....    | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Blunted affect .....             | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Excitement.....                  | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |
| Disorientation .....             | 0 ----- 1 ----- 2 ----- 3 ----- 4 ----- 5 ----- 6 ----- 7 |

## 8. PSYCHIATRIC HISTORY

- 8.1. Psychotic disorders ☐Y ☐N \_\_\_\_\_
- 8.2. Depressive disorders ☐Y ☐N \_\_\_\_\_
- 8.3. Anxiety disorders ☐Y ☐N \_\_\_\_\_
- 8.4. Eating disorders ☐Y ☐N \_\_\_\_\_
- 8.5. Suicide attempts ☐Y ☐N \_\_\_\_\_
- 8.6. Psychiatric hospitalizations ☐Y ☐N \_\_\_\_\_
- 8.7. Use of psychotropic drugs (What of them) ☐Y ☐N \_\_\_\_\_
- 8.8. Use of home-made teas or beverages with calming effects? ☐Y ☐N \_\_\_\_\_
- 8.9. Use of alcohol (Duration and intensity) ☐Y ☐N \_\_\_\_\_
- 8.10. Use of prohibited or illegal drugs (Kind, duration of use motivation to quit) ☐Y ☐N \_\_\_\_\_
- 8.11. Violent behavior ☐Y ☐N \_\_\_\_\_
- 8.12. Problems with the police ☐Y ☐N \_\_\_\_\_

## 9. FAMILY HISTORY

- 9.1. Are there in your family any relatives who have or had any psychiatric problems (treatments, hospitalizations, suicide, and use of calming drugs or antidepressants)? ☐Y ☐N
- 9.2. Has anyone in your family used illegal drugs? ☐Y ☐N
- 9.3. Did anyone in your family die in the past six months? ☐Y ☐N
- 9.4. Has any relative or friend of yours had cancer?  
If yes, could you please tell me how this experience was? ☐Y ☐N

## 10. SOCIAL AND FAMILY SUPPORT

- 10.1. In some of the stressful situations you have been through, who has given you emotional support? \_\_\_\_\_
- 10.2. In financial difficulty, who has given you economic support? \_\_\_\_\_
- 10.3. Since the beginning of your disease, who has given you emotional support? \_\_\_\_\_
- 10.4. Since the beginning of your disease, who has given you financial support? \_\_\_\_\_
- 10.5. Who will take care of you (caregiver) during your hematopoietic stem cell transplant (HSCT)? \_\_\_\_\_
- 10.6. Did your caregiver attend consultations with you? Do you think he/she was well informed about the care you will need during your recovery? \_\_\_\_\_
- 10.7. Please, tell me about the relationship between you and the caregiver? \_\_\_\_\_



## 11. EXPECTATION OF THE TRANSPLANTATION

- 11.1. How do you think this treatment will be?
- 11.2. Do you worry about the failure of this treatment?
- 11.3. How do you think your hospitalization time will be? And the recovery time after HSCT?
- 11.4. Do you believe you will recover your previous health status after the transplant? If you believe it, how long you think it will take you to be recovered? ☐ Y ☐ N ☐ Partially
- 11.5. Considering your answers above tell us about your plans for the future?

(Make the following questions only at the end of the interview, after all other questions are answered)

While you answered these questions you had the opportunity to think about many aspects of this moment of your life: your understanding about the illness and about your transplant, your expectations, your emotional feelings, the way you face crisis situations, the way you follow medical prescriptions, your lifestyle, how your family is and how you can count or rely on it. Furthermore, you had the opportunity to think about how you enjoy your life:

- Do you think this interview is too long or boring? ☐ Y ☐ N ☐ Partially
- Do you think this interview helped you get prepared for the transplant? ☐ Y ☐ N ☐ Partially
- Do you think this interview helped you think about aspects concerning your illness or your transplant which you had not considered before? ☐ Y ☐ N ☐ Partially
- Would you like to make any comments? ☐ Y ☐ N ☐ \_\_\_\_\_

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# Criteria for Donor Selection

Presented by: Kareem Jamani

## Summary

- Donor selection will be based on human leukocyte antigen (HLA)-match and important non-HLA factors that influence transplant outcomes (e.g., age of donor, urgency of transplant, cytomegalovirus (CMV) serostatus) – see Figures 1-3.
- Syngeneic donors may be preferred for aplastic anemia and other diseases with minimal reliance on graft-versus-tumor effect.
  - If a twin of the same sex is considered to be a donor, short tandem repeat polymorphism (“chimerism”) test should be done to determine whether the donor is syngeneic vs allogeneic.
- With HLA-mismatched donors, graft failure has been reported in patients with donor-specific HLA antibodies (DSA). HLA antibody testing should be performed prior to transplantation.
- For donor eligibility (acceptable health), refer to ABMTP Donor Eligibility and Suitability Standard Operating Procedures for allogeneic and cord blood donations, located on ABMTP Sharepoint, or the WMDA website  
<https://share.wmda.info/display/DMSR/WMDA+Donor+Medical+Suitability+Recommendations>

## Selecting an Allogeneic Donor: Guiding Concepts

- Donor characteristics associated with improved overall survival are prioritized.
- When available, local data is incorporated into the selection algorithm.
- For patients with high-risk disease, time to transplant may influence physician choice of donor.

The donor selection algorithms presented in this chapter are mostly consistent with recently published CIBMTR/NMDP guidelines for selection of unrelated donors<sup>1</sup>, with differences related to local data with respect to CMV matching.

## Role of HLA Matching

Large registry studies have provided insight into the role of HLA matching in unrelated donor selection. Most recently, the NMDP/CIBMTR examined ~8000 donor-recipient pairs from almost 200 transplant centers<sup>1</sup>. Importantly, this analysis reflected modern transplant technique: patients were transplanted in 1999-2011 with predominantly peripheral blood stem cells for AML, ALL, MDS and CML. Further, ABMTP transplant technique was well represented in the study: all patients received myeloablative conditioning with ~50% receiving non-TBI based conditioning and ~30% receiving in-vivo T-cell depletion. Important findings included:

- Mismatch at HLA-A, -B, -C or –DR was associated with increased risk of death, grades 2-4 and grades 3-4 aGVHD, cGVHD and treatment-related mortality.



- HLA DQB1 and DPB1 mismatches: among 8/8 matched cases, DQB1 mismatches were only associated with grades 2-4 aGVHD (RR1.2) and DPB1 mismatches were associated with grades 2-4 aGVHD (RR 1.4), grades 3-4 aGVHD (RR 1.5), and decreased relapse (RR 0.71). Importantly, DPB1 non-permissive mismatches were associated with a higher risk of death as compared to DPB1 permissive mismatches. There was no difference in survival between those receiving DPB1 permissive mismatch versus those receiving DPB1 matched donors, however DPB1 permissive mismatched donors were associated with an increased risk of grades 2-4 (RR 1.3) and 3-4 aGVHD (RR 1.4) and a lower risk of relapse (RR 0.7) as compared to DPB1 matched donors. These findings with respect to DPB1 replicated those of an earlier study in a less recent cohort of patients receiving primarily bone marrow grafts after conditioning with predominantly TBI-based myeloablative conditioning with only 16% receiving in-vivo T-cell depletion<sup>2</sup>.
- In contrast, regarding HLA-DPB1 matching, local data did not demonstrate significant differences in relapse and GVHD outcomes among donor-recipient pairs matched or permissive/non-permissive mismatched at HLA-DPB1 in a small cohort without multivariable adjustment<sup>4</sup>. The local data requires further follow-up in a larger cohort with multivariable modelling.

## Older Matched Sib (MSD) vs. Younger MUD Donors

Among MUDs, younger age has been repeatedly found to be the most important donor characteristic associated with recipient overall survival. Thus, investigators have sought to evaluate outcomes of allo-HCT with a young MUD (typically defined as <35 y.o.) vs an older MSD (typically defined as >50 y.o.). Retrospective analyses seeking to answer this question can be summarized as follows:

- AML/MDS: In two separate CIBMTR analyses MUDs were associated with a lower risk of relapse and improved disease-free survival vs. MSDs, but this was countered by an increased risk of acute and chronic GVHD and non-relapse mortality. No difference in overall survival was noted. (Guru Murthy 2022 PMID 35024768 & Abid 2023 PMID 37406882). These results could not be replicated in a more contemporary large single-centre analysis where no difference in any outcome was noted (Kim 2024 PMID 38703824).
- B-cell ALL: In a CIBMTR analysis, MUDs were associated with a lower risk of relapse vs. MSDs, but a greater risk of cGVHD and NRM, leading to no difference in overall- or leukemia-free-survival (Abid 2023 PMID 37481243).

Summary: Older MSDs and young MUDs are associated with similar outcomes. Select the donor that allows for fastest time to transplant.

## HLA-Mismatched Donors

Per unpublished 2025 analysis of Albertan data on patients with hematologic malignancies who underwent PBSCT in 1999-2014 with conditioning of fludarabine (250 mg/m<sup>2</sup>) + busulfan(12.8 mg/kg,



PK adjusted) + 4GyTBI, OS and cGRFS were virtually identical for 7/8 HLA-matched unrelated donors with ATG(4.5 mg/kg)-based GVHD prophylaxis and haploidentical related donors with PTCy(100 mg/kg)-based GVHD prophylaxis (Storek, unpublished).

Haploidentical donor is the preferred donor in Alberta when an HLA matched donor is not available. The rationale is that haploidentical related donors are widely and quickly available and excellent outcomes have been reported in the literature using the PTCy platform. Also, there is some cost saving when a haploidentical related donor is used instead of a 7/8 matched unrelated donor. Umbilical cord HCT has not been a typical choice for alternative donor HCT for adults in Alberta. Umbilical cord HCT may be pursued in circumstances where proceeding to allo-HCT is felt to be critically important and 7/8 unrelated or haploidentical donors are unavailable.

In both 7/8 unrelated donor HCT and haploidentical HCT, it is critical to identify donor-specific antibodies (DSAs) against the locus/loci of mismatch. DSAs have been strongly associated with graft failure, poor graft function and reduced overall survival<sup>6</sup>. Donors without the corresponding HLA antigens should be selected. If the latter are unavailable, recipients should undergo desensitization prior to HCT.

With respect to selecting a 7/8 HLA matched unrelated donor, the recent CIBMTR/NMDP analysis noted the following<sup>3</sup>:

- When comparing locus of HLA mismatch, there were no large differences in outcomes.
- The exception to this was mismatch at HLA-C 03:03/03:04, which was associated with similar survival to 8/8 matched transplants, a finding that replicated that of a previous study<sup>7</sup>.
- Among 7/8 donors, no significant effect of mismatch at DQB1 or DPB1 mismatch were observed. This stands in contrast to findings of other studies, i.e. the literature does not provide clear guidance with respect to matching DQB1 and DPB1 in 7/8 matched donors. However, an as of yet unreplicated single study suggests that >2 mismatches of low expression HLA loci (HLA-DP, -DQ, -DRB3/4/5) in 7/8 transplants is associated with increased mortality (HR 1.45), predominantly in the form of treatment-related mortality (HR 1.68)<sup>7</sup>.

Selection of donors for haploidentical HCT has been reviewed by the EBMT along with the publication of consensus recommendations<sup>8</sup>. Overall, the literature regarding haploidentical donor selection with the PTCy platform is sparse as compared to the literature regarding unrelated donor selection. The following haplo HCT donor characteristics have been associated with improved survival in >1 published study<sup>8</sup>:

- Younger donor age (particularly less than 40 y.o.).
- Sibling or offspring donor as opposed to parent donor (however, this was not replicated in the large CIBMTR study discussed below).
- Avoid major ABO incompatibility only in bone marrow transplantation (no effect of ABO mismatch in peripheral blood stem cell transplant).

The following donor characteristics may be associated with improved recipient outcomes but require further verification. These characteristics may be considered if the above donor criteria are met and there remains >1 suitable haploidentical donor choice<sup>8</sup>:

- Male donor for male recipient.
- If using a parent donor, father donor preferred over mother.

The following donor characteristics have not consistently been shown to affect recipient outcome in haploidentical HCT<sup>8</sup>:

- Degree of HLA mismatch.

A recent CIBMTR study examined the largest cohort to date (>1400) of haploidentical HCTs transplanted with a PTCy platform and provided significant insight with respect to haploidentical HCT donor selection, particularly with respect to the role of HLA matching<sup>9</sup>. Major findings included:

- Improvement in overall survival was associated with younger donor (particularly age <40), match at HLA-B leader, nonpermissive mismatch at HLA-DPB1 and CMV seronegative donor for seronegative recipient.
- Improvement in disease-free survival was associated with match at HLA-B leader, non-permissive mismatch at HLA-DPB1, CMV seronegative donor for CMV seronegative recipient and mismatch at HLA-DRB1 in conjunction with match at HLA-DQB1.
- Donor-recipient relationship was not associated with any major outcome studied.
- Because of the difficulty of simultaneously considering clinical and HLA factors, the authors developed a disease-free survival calculator that considers all factors associated with disease-free survival noted above. The calculator can be used to compare multiple donor options. The calculator is located at: <http://haplodonorselector.b12x.org/v1.0/>
- Importantly, no further HLA typing is required to determine HLA-B leader matching as leaders are known for most HLA-B alleles and may be sought out by entering donor-recipient HLA-B alleles in the following calculator: <https://bleader.nmdp.org>

A suggested approach is to type all potential haploidentical donors under age 40 given the recurrently found strong association between donor age and overall survival. If there are >1 potential donors under age 40, the above calculator may be used to select the donor that will result in the best disease-free survival. If there are no potential donors under age 40 but >1 over age 40, the calculator may be used to select the donor that may result in the best disease-free survival.

### **Donor-Specific Antibodies (DSA)**

For recipients with potential HLA-mismatched unrelated donors and haploidentical donors, HLA antibody screening will be performed. If HLA antibody screening is positive then Single Antigen Bead analysis will be performed to assess the presence of donor-specific HLA antibodies (HLA-DSA). In recipients positive for HLA-DSA, a T and B cell flow crossmatch will be performed using donor lymphocytes and recipient serum.

## Non-HLA Factors

Three modern studies have confirmed that age is indeed the only non-HLA donor factor that is consistently associated with recipient survival. The first study examined two cohorts of several thousand patients each with the intention of producing a donor selection score: three donor characteristics predicted recipient mortality in the first cohort: CMV mismatch (negative impact for D-R+, HR 1.14), non-permissive DPB1 mismatch (negative impact, HR 1.13) and older donor age (negative impact, HR 1.07 per decade increase in age). In the second cohort, only donor age predicted recipient survival (negative impact, HR 1.11 per decade increase in age). In absolute terms, choosing a donor 2 years older was associated with a 1% decrease in 2 year survival, 5 years older a 2% decrease, 10 years older a 3% decrease and 20 years older a 7% decrease<sup>11</sup>.

In a second large study, donor age was the only non-HLA donor factor that was associated with survival. As in the former study, donor age was continuously associated with recipient survival: for every 10 year increment in donor age, there was a 5.5% increase in hazard ratio for overall mortality<sup>12</sup>.

In a third study, a machine learning model using ~12 000 patients reported to CIBMTR provided further insight into the degree of donor age that leads to an optimal outcome: donors aged 30 or younger lead to optimal overall survival with little additional benefit going further. A male donor leads to optimal event-free survival. All other donor characteristics were not associated with EFS or OS. Thus, male donors aged 30 or younger are preferred. (Spellman 2024 PMID 39368807)

**CMV serostatus.** Selecting a CMV seronegative donor for a CMV seronegative recipient is a commonly accepted practice based on multiple reports of worse survival of seronegative recipients receiving grafts from seropositive donors<sup>13,14</sup>, but this is generally based on studies in which GVHD prophylaxis did not include rabbit ATG (anti-thymocyte globulin). Kalra *et al.* published the outcomes in 928 Alberta patients who underwent myeloblastic HSCT in hematological malignancies between 1999 and 2014 who received ATG as part of the conditioning regimen, and focused on the impact of donor and recipient CMV serostatus on transplant outcomes<sup>15</sup>. In this study, donor CMV serostatus had no impact on recipients who were CMV sero-negative, whereas there was a substantially lower survival in the D-R+ patient group versus D+R+ (41% vs. 59% at five years, p=0.001). Survival rates were also lower in D-R+ HLA-matched sibling transplant recipients compared with D+R+ HLA matched unrelated donor transplant recipients (44% vs. 66%) at 5 years, p=0.009). The differences in survival were being attributed to higher non-relapse mortality. The conclusion from this study was that, when using ATG for patients with malignancies, choosing a CMV seropositive donor for a CMV seropositive recipient is important, even if this requires an unrelated graft. In an updated analysis, the difference in survival between the the D-R+ group versus D+R+ group appeared to be limited to those with lymphoid malignancies<sup>4</sup>. Thus, CMV matching for R+ recipients need only be considered for those undergoing allo-HCT for lymphoid malignancies.

Thus, after matching for factors that influence recipient survival: HLA-A, -B, -C and –DR matching, CMV serostatus in those with lymphoid malignancies, younger age of donor, avoidance of HLA-DPB1 non-permissive mismatching, and other less important factors (e.g., gender, ABO compatibility, donor size, in no particular order) can be considered.

## Syngeneic Donors

There is no need for GVHD prophylaxis. These transplants are associated with a higher relapse rate compared to matched sibling transplants in malignancies where a graft-versus-leukemia effect is important; such as acute myeloid leukemia (AML) (52 versus 16%), CML (40 versus 7%), and acute lymphoblastic leukemia (ALL) (36 versus 26% at 3 years)<sup>10</sup>. However, this disadvantage may be outweighed by the advantage of no GVHD and thus lower NRM, at least for AML patients undergoing HCT in 1CR.<sup>16</sup>

No graft-versus-lymphoma effect has been seen in non-Hodgkin lymphoma syngeneic versus allogeneic registry data, and syngeneic transplant may be a good option for lymphomas or benign disorders.

Short tandem repeat polymorphism (same test as we use for chimerism) is used to determine whether a same-sex twin is monozygotic (syngeneic) or dizygotic (allogeneic).<sup>17</sup>

## Donor Eligibility and Suitability

According to the Health Canada lymphohematopoietic cells for transplant standards manual, the following donors are **not suitable** for peripheral blood stem cell or bone marrow donation:

- persons with prion-related disease
- persons with a potentially transmittable neurological disease of an unestablished etiology
- persons with active encephalitis or meningitis of infectious or unknown etiology
- persons with rabies or persons who, within the past 6 months, were bitten by an animal and treated as if the animal was rabid
- persons with a history of infection with HIV, clinically active HCV or HBV
- persons with a family hx of Creutzfeldt-Jakob disease
- persons who have received human-derived pituitary growth hormone
- persons who have received dura matter
- persons with HTLV-I or HTLV-II, active WNV, or syphilis
- persons who have known or suspected sepsis at the time of donation
- persons previously diagnosed with a hematologic malignancy or with melanoma
- persons previously diagnosed with other types of cancer, unless they have been evaluated and deemed suitable to donate by a physician

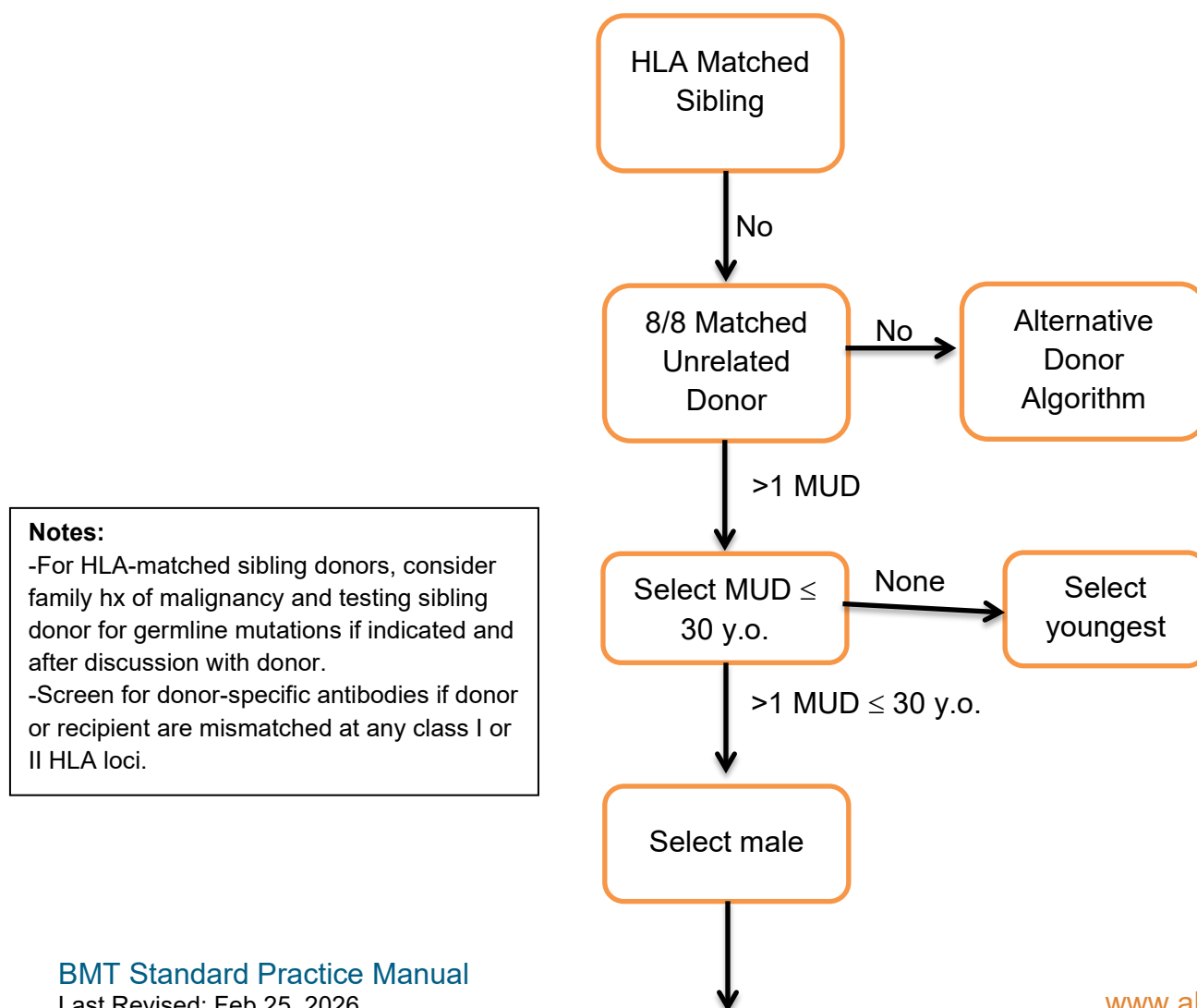
Further, the WMDA Donor Medical Suitability Wiki can be used as a resource regarding further suitability issues, particularly with respect to donor medical conditions. This can be found on WMDA's website at

<https://share.wmda.info/display/DMSR/WMDA+Donor+Medical+Suitability+Recommendations>

Finally, potential donors may screen positive on the ABMTP Donor Personal History Questionnaire (for example, related to travel, habits, sexual history etc.). In the case of a positive screen, donor clearance will be at the donor physician's discretion. If a clinically relevant risk of disease transmission is identified, an exceptional release may be requested by the donor physician in case of urgent medical need. Further guidance with respect to positive screens on the ABMTP Donor Personal History Questionnaire and their implications are found in the Donor Personal History Questionnaire Guidance Document found here:

#### Donor Questionnaire Guidance Document

**Figure 1.** Algorithm for Selection of a Donor for Recipients with Non-Lymphoid Malignancies or Recipients with Lymphoid Malignancies who are CMV Seronegative.



>1 male MUD  $\leq 30$  y.o.

In no particular order:  
 Avoid non-permissive HLA-DPB1 mismatch  
 -Match HLA-DQB1  
 -Donor size  
 -ABO compatibility  
 -CMV serostatus match

**Figure 2.** Algorithm for Selection of a Donor for CMV Seropositive Recipients with Lymphoid Malignancies and Receiving ATG-based GVHD Prophylaxis

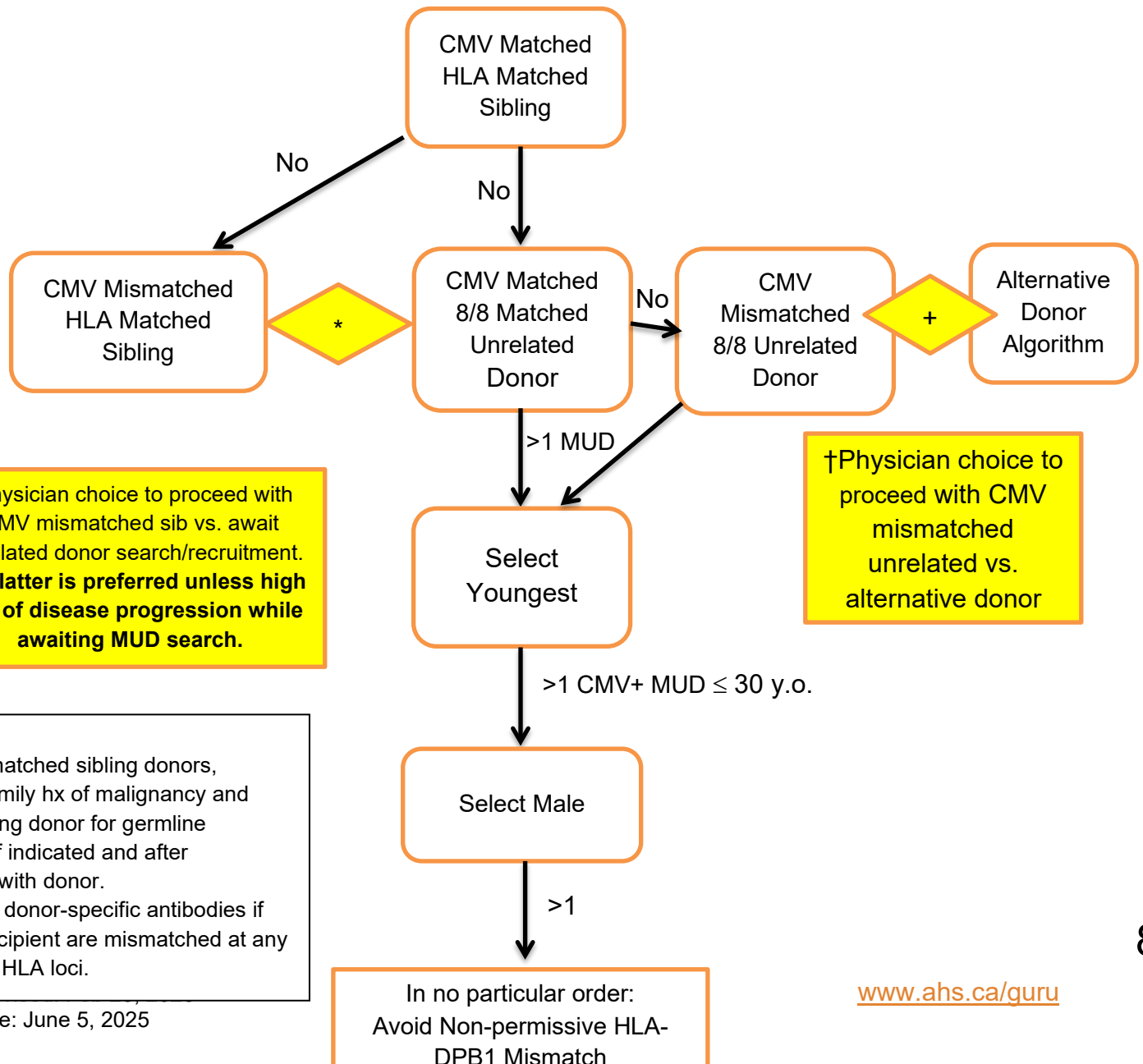
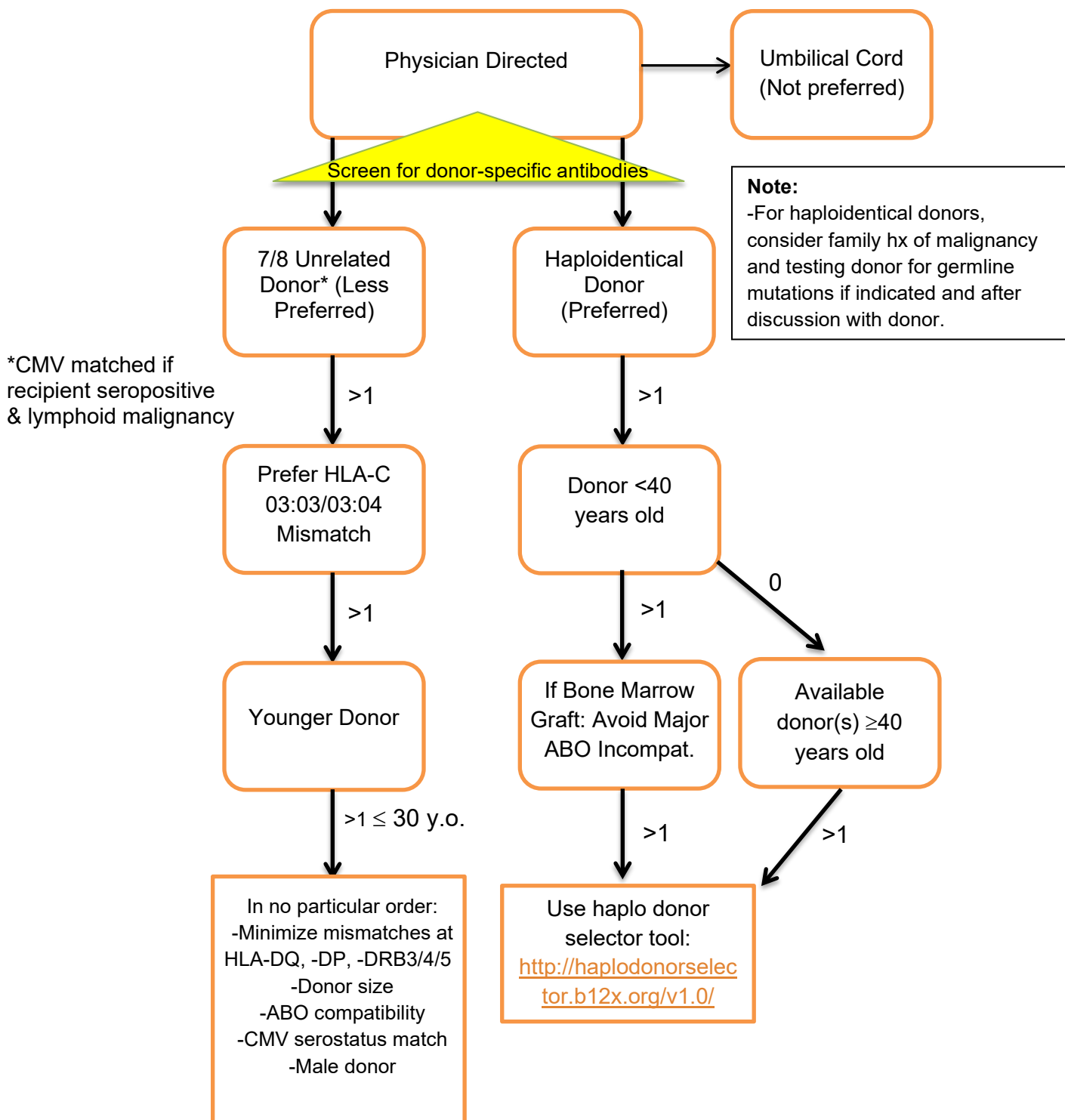




**Figure 3. Algorithm for Selection of an Alternative Donor**



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# Stem Cell Mobilization

Presented by: Peter Duggan

## Summary

### Autologous Stem Cell Collections

- For autologous stem cell collection, mobilization options include G-CSF alone (for patients who have not had prior chemo- or radio-therapy), plerixafor plus G-CSF, or combined chemotherapy and G-CSF mobilization (for those who have had prior chemo- or radio-therapy).
- Filgrastim or biosimilars may be used for autologous stem cell mobilization.
- Plerixafor is indicated for patients who are at risk for poor mobilization, those who have failed a previous mobilization attempt, for salvage during a suboptimal mobilization attempt, and for planned collection in combination with G-CSF.
- Peripheral blood is the recommended source of stem cells for autologous transplantation. Bone marrow harvests are not recommended.
- Ex-vivo purging of malignant cells from autologous stem cell products (eg, by CD34 enrichment) in patients with malignancies is not recommended.
- For CD34 enriched autografts (for autoimmune diseases), CD34 collection targets will be set at  $8 \times 10^6/\text{kg}$  and collected in a single full day apheresis. Plerixafor may be required if a sufficient peripheral blood CD34 count is not achieved with G-CSF alone.

### Allogeneic Stem Cell Collections

- Allogeneic donors undergoing mobilization will receive G-CSF daily for five days. Additional dose(s) should be given on day 5 and/or day 6 if fewer than  $3 \times 10^6$  CD34+ cells/kg are collected, and a second day of collection should be arranged. Filgrastim or biosimilars may be used.
- The Canadian Blood Services Stem Cell Donor Registry has indicated that *plerixafor* should not be given to unrelated donors who fail to mobilize with G-CSF. In the case of unrelated donors who are not mobilizing as expected, early communication to CBS is essential. The transplant center should be contacted in order to confirm that bone marrow would be acceptable prior to arranging a bone marrow harvest in these cases.
- Related allogeneic donors who mobilize poorly with G-CSF alone (blood CD34 count 5-10 per microliter) will be offered off-label plerixafor. Plerixafor may also be considered prior to a second day of apheresis if the target CD34 yield was not achieved on the first day of apheresis. Those refusing plerixafor will undergo surgical bone marrow harvest on day +6 of G-CSF. Those who fail mobilization (CD34 count  $< 5$ ) will undergo surgical marrow harvest on day +6.

## Autologous Stem Cell Mobilization Options

The preferred stem cell source for autologous SCT is mobilized peripheral blood stem cells collected by apheresis. This is based upon small RCTs that demonstrated improved quality of life, shorter engraftment times, decreased blood and platelet transfusions, decreased hospital stays, and reduced

costs relative to traditional bone marrow harvests<sup>1-8</sup>. Chemotherapy (a salvage regimen or cyclophosphamide 2-4g/m<sup>2</sup>) plus G-CSF 5-10 mcg/kg/day is an acceptable standard method of stem cell mobilization<sup>9-19</sup>. Predictors of poor mobilization include: advancing age, prior treatment with chlorambucil, fludarabine, melphalan, radiotherapy to >25% of bone marrow, or repeated cycles of chemotherapy plus G-CSF within the past 6 months, as well as those with low blood platelet counts prior to mobilization treatment, or those who have experienced prior failure of stem cell mobilization.

A chemotherapy free regimen using GCF plus plerixafor has been utilized. There are no randomized controlled trials comparing chemotherapy based mobilization regimens to mobilization with plerixafor and growth factors. Retrospective comparisons do show higher total and day one CD34+counts following chemotherapy based mobilization, but this is associated with higher risk of febrile neutropenia. There is no significant difference in the number of patients able to proceed to transplant, the number of apheresis days and transplant outcomes<sup>33-35</sup>. Those mobilized with plerixafor and GCSF are almost always collected on their planned day of apheresis. This approach is most widely described in the setting of multiple myeloma, and can be considered when disease targeted chemotherapy is not required and the collection target is modest (e.g. single transplant). It may not be a successful approach if collection for multiple transplants is planned.

*The role of graft purging of malignant cells in patients with malignancies.* Registry (CIBMTR and EBMT) data suggest there may be a role for purging based on the extremely low relapse rates following syngeneic SCT, followed by higher relapse rates with purged autologous SCT and then significantly higher relapse rates with unpurged autologous SCT<sup>20</sup>. This data, however, is potentially biased, and randomized controlled trials evaluating ex-vivo autograft tumor purging techniques have not been reported in the setting of autologous transplantation for lymphoma. In addition, autograft purging results in stem cell loss and delays hematopoietic and immunological engraftment<sup>21-23</sup>. Because of these facts, routine ex-vivo autograft purging is not recommended.

## **Option 1. Granulocyte Colony Stimulating Factor (G-CSF) Alone, or G-CSF with Plerixafor.**

### **Indications:**

- Mobilization of peripheral blood stem cells for autologous stem cell transplant patients who *have not* had prior chemotherapy or radiotherapy.
- G-CSF with or without plerixafor is preferred over chemotherapy + growth factors for mobilizing patient with multiple myeloma and amyloidosis. Chemotherapy may be considered for those who are expected to mobilize poorly, who have a high CD34 target, or who have significant residual myeloma in marrow.
- Plerixafor may be given “on demand” for those with a low CD34 count after 4-5 days of GCSF alone, or planned on day 4, prior to apheresis the following day.

### **G-CSF dosing - autologous donor:**

- G-CSF 5-10 µg/kg/day for 4 days, rounded to nearest vial size and fewest injections (see Table 1).

### **Plerixafor dosing**

- 0.24mg/kg/day, to maximum dose of 24mg/day.

## **Option 2. Combined Chemotherapy and G-CSF Stem Cell Mobilization**

### **Indications:**

- Mobilization of peripheral blood stem cells for autologous stem cell transplant patients who *have* had prior chemotherapy or radiotherapy.

### **Standard intensity regimen indications:**

- Myeloma, when growth factor only mobilization is not preferred.
- Germ cell tumours
- Lymphoma with largest tumour mass less than 5 cm and negative marrow biopsies
- Most other miscellaneous indications

### **Standard intensity regimens include:**

- Cyclophosphamide 2.5 g/m<sup>2</sup> day 1 OR standard dose regimen such as DHAP, high dose cytarabine, VIP, or TIP
- Add Rituximab 1400mg sc on first day chemotherapy for CD20+ B-cell lymphomas
- G-CSF starting on Wednesday of the following week (~day 7-9)
- Apheresis scheduled for Monday-Wednesday 2 weeks after chemo (~days 12 to 14)

### **High intensity regimen indications:**

- Lymphoma with mass greater than 5 cm, bone marrow involvement, or refractory disease

### **High intensity regimen example:**

- DICEP regimen
  - Cyclophosphamide 1.75 g/m<sup>2</sup>/day x 3 days
  - Etoposide 350 mg/m<sup>2</sup>/day x 3 days
  - Cisplatin 35 mg/m<sup>2</sup>/day x 3 days
- Add Rituximab 375mg/m<sup>2</sup> IV (or 1400mg sc) for CD20+ B-cell lymphomas (decrease dose of all chemo by 20% for patients >60yrs)
- G-CSF starting day 14
- Apheresis ~days 19 to 21

### G-CSF dosing:

Patients without risk factors for poor mobilization should receive G-CSF 5-10 µg/kg/day, rounded to nearest vial size and fewest injections, beginning on the day indicated in the protocol and continuing until completion of apheresis (see Table 1)

**Table 1.** Dosing for granulocyte colony stimulating factor based on weight for autologous stem cell mobilization

| Donor Weight (kg) | G-CSF Dose (µg)* |
|-------------------|------------------|
| < 60kg            | 300              |
| 60 - 75           | 480              |
| 75.1 – 100        | 600              |
| > 100             | 780              |

\*dose ranges yield 5 to 8 mcg/kg

### Option 3. Plerixafor for Stem Cell Mobilization

#### Risk Factors for poor mobilization:

1. Advanced disease (≥2 lines of chemotherapy)
2. Extensive BM involvement or cellularity <30% at time of mobilization
3. Age >60
4. Prior radiotherapy to >25% of bone marrow surface area
5. Prior treatment with fludarabine and other purine analogues, lenalidomide, melphalan
6. Platelets less than 100x 10<sup>9</sup>/L prior to mobilization
7. Prior failed mobilization attempt

#### Plerixafor use should be considered in the following settings:

1. Preemptively for patients predicted to mobilize poorly based on the risk factors above. It should be used in combination with G-CSF with or without chemotherapy.
2. For salvage immediately prior to apheresis for patients with suboptimal mobilization. Plerixafor should be given if the post nadir WBC count is >5 x 10<sup>9</sup>/L and CD34 count is >5 but <20 x 10<sup>6</sup>/L or if <50% of the target CD34 yield was achieved on the first day of apheresis.
3. Re-mobilization for patients with a prior failed attempt at mobilization with G-CSF with or without chemotherapy.
4. As part of a planned chemotherapy-free collection strategy using plerixafor with GCSF.

#### Plerixafor dosing:

- The recommended dose of Plerixafor is 0.24 mg/kg body weight (0.16 mg/kg for creatinine clearance <50 mL/min) by subcutaneous injection, injection with the dose capped at 24mg, given the day before apheresis is planned, and then daily until apheresis is complete.



**Apheresis:**

- Performed on the day when the post-chemotherapy nadir blood counts have recovered to:
  - Platelet greater than  $30 \times 10^9/\text{L}$  and hemoglobin  $>75 \text{ g/L}$
  - CD34+ count greater than  $20 \times 10^6/\text{L}$
- Plan for large volume apheresis ( $\geq 3$  blood volumes, approximately 15 L) using a central venous catheter for autologous donors. Minimum apheresis volume of 8L.
- Target CD34+ collection:
  - Minimum target all patients:  $2 \times 10^6$  CD34+ cells/kg/transplant
  - Ideal target 5 to  $10 \times 10^6$  CD34+ cells/kg/transplant (preferred)

**Mobilization and Collection for CD34 Enriched Autologous Transplants**

For patients requiring CD34 enriched autologous stem cell products, such as those undergoing ASCT for autoimmune disease, CD34 collection targets will be higher in order to compensate for losses during processing and to obtain a final product containing  $5 \times 10^6$  CD34 cells. This target is to be collected on a single day of apheresis to avoid needing more than one day of CD34 selection. Based on these requirements, the following collection parameters are recommended:

1. Requested CD34 cell dose for collection will be set at  $8 \times 10^6/\text{kg}$ .
2. Total CD34 cell target must be collected in a single full day apheresis.
3. Sufficient peripheral blood CD34 count is necessary to meet these parameters. Plerixafor may be required if a sufficient peripheral blood CD34 count is not achieved with G-CSF alone.
  - a. If PB CD34  $\geq 150$ , regardless of target, proceed with collection.
  - b. If PB CD34  $\geq 60$  and CD34 collection target is  $\leq 700$ , proceed with collection.
  - c. If PB CD34  $\geq 60$  and  $<150$  but CD34 collection target  $\geq 700$ , arrange for plerixafor and collect on the following day.
  - d. If PB CD34  $\leq 60$ , arrange for plerixafor and collect the following day. Exceptions to this may include those with CD34  $< 60$  but a low collection target based on weight that is expected to be collected based on the current CD34 count without the addition of plerixafor.

## Allogeneic Stem Cell Transplant Donors

**G-CSF dosing - allogeneic donor**

- For donors weighing more than 48 kg, G-CSF 8-10  $\mu\text{g/kg/day}$ , rounded to nearest vial size (see table). Individualize dosing for donors weighing  $< 48 \text{ kg}$  or  $> 120 \text{ kg}$ , irrespective of vial sizes. Doses are given daily for five days (days 1 to 5) with apheresis collection on day 5.

**Table 2.** Dosing for granulocyte colony stimulating factor based on weight for allogeneic donor stem cell mobilization

| Donor Weight (kg) | G-CSF Dose (µg) |
|-------------------|-----------------|
| < 48              | 300             |
| 48-60             | 480             |
| 60.1-78           | 600             |
| 78.1-100          | 780             |
| 100.1-120         | 960             |

**Additional doses of G-CSF for allogeneic donors:**

- Order additional doses of G-CSF to be administered after collection and the following morning if fewer than  $3 \times 10^6$  CD34+ cells/kg were collected in the first apheresis session.

**Apheresis:**

- Plan large volume apheresis ( $\geq 3$  blood volumes) on day 5 using peripheral venous access. A second collection day may be required if the minimum dose is not reached on one day of apheresis.
  - Donor apheresis using central venous access devices (CVAD) will be used only in exceptional circumstances. In the case of unrelated donors, the need for CVAD use will be discussed with CBS.
- Minimum of 8L apheresis
- Target CD34+ Collection:  $5 - 10 \times 10^6$ /kg recipient weight
- Minimum target:  $3 \times 10^6$  CD34+ cells/kg recipient weight

**Failed Mobilization of an Allogeneic Donor** Failure to collect a sufficient number of stem cells to transplant an allogeneic recipient has very significant implications for that recipient. Options for the transplant center include approaching a backup donor or approaching an alternative family member for haploidentical donation; in most cases, however, the recipient has already been conditioned and approaching a second donor may result in significant delays and prolonged aplasia. In Alberta, if a related donor mobilizes poorly (blood CD34 count 5-10 per microliter) after five days of G-CSF they will be offered a dose of plerixafor off label and will undergo collection by apheresis on day 6 if CD34 count increases to  $\geq 15$ . Donors who decline to receive plerixafor or whose blood CD34 count is  $< 5$  per microliter on the fifth day of G-CSF will be asked to undergo surgical bone marrow harvest urgently on day 6. In cases where a related donor completes apheresis on day 5 but the target CD34 yield was not achieved on the first day of apheresis, plerixafor may also be considered prior to a second day of apheresis.

For unrelated donors, in the event that there is a failed mobilization, the collection center must have the ability to collect marrow as an emergency procedure. The collection center must inform the CBS if the donor expresses concerns or would not be willing and/or eligible for marrow HPC as a contingency collection. The CBS Stem Cell Registry has advised that plerixafor should not be

administered to unrelated donors. The WMDA also does not recommend the use of plerixafor but recognizes that there may be situations where its use is appropriate. In the case of failed mobilization of an unrelated donor, early notification of the transplant center is essential. The collection center will work closely with CBS to determine the next steps in the collection plan.

### **Cytokine-Stimulated Bone Marrow**

The use of G-CSF stimulated bone marrow for hematopoietic cell transplantation was proposed as a way of providing a product with the rapid engraftment potential of G-CSF mobilized peripheral blood grafts but with the low risk of GVHD associated with bone marrow. Studies have shown that GVHD rates are lower with bone marrow (including G-CSF stimulated marrow) but no consistent advantage of G-CSF stimulated marrow over unstimulated marrow has been demonstrated either in terms of overall- or progression-free survival<sup>24</sup>. The ABMTP will not routinely administer G-CSF to bone marrow donors.

### **Donors from Vulnerable Groups**

#### **Minor Donors: <18 years of age may be selected if:**

- a. There is no equivalent histocompatible adult donor who is willing and readily available for donation
- b. It is deemed the recipient will benefit from transplantation
- c. The clinical, emotional and psychological risks to the donor are minimized and are reasonable in relation to the benefits expected to the donor and the recipient as outlined in the pediatric donor eligibility and suitability evaluation SOP (BMTS20005)
- d. Following a psychological evaluation, the staff has deemed that there is a strong personal and emotionally positive relationship between the donor and recipient as outlined in the Pediatric Donor Eligibility and Suitability Evaluation SOP (BMTS20005)
- e. Parental permission/consent and child assent will be obtained as per Pediatric Blood and Marrow Transplant Consent Procedure (BMTS20009)
- f. A donor advocate trained in pediatrics will be assigned as outlined in the Pediatric Donor Eligibility and Suitability Evaluation SOP (BMTS20005)
- g. The donor must weigh a sufficient amount to safely undergo collection

#### **Older Donors: >65 years of age:**

- a. Must be able to complete standard donor testing outlined in Standard Protocol Allogeneic Donor Collection Workup (BMTW34092)
- b. Must meet suitability and eligibility criteria as defined in Donor Eligibility and Suitability SOP (BMTS10212)
- c. Must have general good health as determined by physician assessment
- d. Comorbidities are identified and evaluated by donor physician

- e. Must have a performance status that will permit the safe collection of cells as determined by physician assessment
- f. Resources will be provided for disabilities, including the visual or hearing impairments
- g. Donors may access a third party advocate as they feel appropriate as per “Interaction Between Alberta Health Services and Third Party Advocates PRR-04”

## **Hemoglobinopathies:**

G-CSF is not advised for mobilization in donors with sickle cell disease.

Hemoglobinopathy assessment is required for all donors (autologous and allogeneic) as administration of mobilization agents such as G-CSF may pose a risk to the donor as it was associated with morbidity (e.g. veno- occlusive crisis) and mortality in donors with sickle cell disease (HBSS), (HPSC), and also with compound hemoglobinopathies such as sickle-beta-thalassemia (S/β Thal). Patients with hemoglobinopathy will be identified based on their medical history. Testing of donors with no history of hemoglobinopathy and with a normal hemoglobin is not required, but is an acceptable method.

Of note, medical literature reports that donors with sickle cell trait have been safely mobilized and collected<sup>32-34</sup>. Although donors with the sickle cell trait did have higher symptom scores than control donors, there were no symptoms suggestive of sickle crisis. Thus, in this group the risk is limited.

## **Repeat Donations**

The Alberta BMT Program permits donors to donate on more than one occasion, provided the risk of donating is justified by the condition of the recipient. Donors will only be permitted to donate stem cells (bone marrow and/or G-CSF stimulated peripheral blood stem cells) twice, although the program will not limit the number of donations of non-mobilized cells such as donor lymphocyte infusions.

## **Weekend Apheresis**

If a Weekend Apheresis Collection is probable or confirmed, the most responsible physician shall contact personnel in Flow Cytometry, Apheresis and Cellular Therapy Laboratory by Friday at noon.

- Apheresis Manager: (403) 944-5471
- Flow Cytometry, Tech III: (587) 231-3904
- Cellular Therapy Lab: (585) 231-5203

## **Use of G-CSF Biosimilars**

Biosimilars are approved biologics with comparable quality, safety, and efficacy to a reference product for which patent protection has expired. Biosimilar regulatory approval is provided on the basis of a robust comparability exercise demonstrating similarity with the original product, rather than on the need

to show a positive risk-benefit assessment, which it is assumed has already been proven. The degree of clinical similarity required to achieve biosimilar status is considered on a case-by case basis by the regulatory authorities.

Biosimilars of G-CSF, based on the original filgrastim product Neupogen, have been available for a number of years and are now widely used, often exceeding the use of the original product. For the currently approved biosimilar G-CSFs (e.g. Nivestym), extrapolation to all indications of the reference product has been granted, given the comparable receptor site kinetics and mode of action. This includes mobilization of peripheral blood stem cells in patients undergoing autologous stem cell transplantation as well as for stem cell mobilization in patients and healthy donors.

For autologous stem cell mobilization, the overall effectiveness of biosimilar G-CSF has been evaluated in several open-label studies, some of which have include the reference product as a comparator. All of these studies have shown no significant differences in efficacy (e.g. median number of CD34+ cells collected, number of G-CSF injections required, apheresis days, etc.), and safety, with similar incidence and severity of common adverse events such as bone or muscles pain and headache, and no severe or unexpected AEs<sup>25-27</sup>. There are a few reports of biosimilar G-CSF use for PBSC mobilization in healthy donors that suggest these agents are effective and well tolerated, with similar mobilization outcomes in comparison to Neupogen, with no clinically significant differences between groups<sup>28-30</sup>. There is an ongoing long-term safety study over 10 years, which will contribute data for up to 2000 person-years and add to the cumulative assessment of the long-term safety of G-CSF as a mobilizing agent<sup>31</sup>.

As of 2025, the WMDA recommendation on the use of filgrastim biosimilars has been revised as follows: Stem cell donor registries and affiliated organisations may use filgrastim biosimilars for CD34-positive cell mobilisation in healthy donors.

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# Vaccination

Presented by: Kareem Jamani

## Summary

- Transplant recipients should be immunized according to the Guidelines of Community and Population Health Division (“Public Health”), Alberta Health and Wellness, posted at <https://www.alberta.ca/alberta-immunization-policy.aspx#toc-4> (under ‘Special situations for immunization’).

### Highlights of the Schedule:

- 6 mo posttransplant, start non-live vaccines (given at 6, 7, 8, 12, 14 and 24 mo)
- 24 mo posttransplant, start live vaccines (given at 24 and 27 mo) – contraindicated in patients with relapse or active cGVHD – wait until  $\geq 3$  mo after discontinuation of immunosuppressive therapy (systemic and topical) and no cGVHD activity. Discontinue valacyclovir 1 day before first VZV vaccine dose.
- Autologous HCT recipients will be vaccinated for varicella/zoster with Shingrix (rather than live VZV vaccine) at 6 and 7 months post-HCT. Valacyclovir will be discontinued 1 month after the second Shingrix dose. Continuation of VCV can be considered in cases of immunosuppressive therapy during/after the Shingrix vaccination.
- CAR-T recipients will be vaccinated according to the post allogeneic HCT vaccination schedule.
- 36 mo posttransplant, check antibody levels to tetanus, hepatitis B, measles and rubella, and order boosters if needed.

## Background

The Albertan Guidelines were developed based on international guidelines,<sup>1</sup> keeping simplicity in mind. For example, the same schedule was developed for autologous and allogeneic transplant recipients, and similar schedule was developed for children and adults. The reason for simplicity is to minimize confusion that could arise from the fact that many parties are involved in the vaccination process, including the transplant physician, the hematologist/oncologist to whom an autologous transplant recipient is referred after autologous transplantation, the Public Health vaccination clinic administering the vaccines and, in special scenarios, Infectious Disease specialist, Public Health specialist (“Medical Officer of Health”) or Travel Clinic physician.

### Practical Considerations

- Antibody levels to vaccine-preventable diseases decline during 1-10 years posttransplant if the recipient is not revaccinated.
  - The decline is more substantial in allogeneic compared to autologous HCT recipients. Therefore, and because influenza, pneumococcal disease and shingles are less frequent

after autologous than allogeneic HCT, vaccination is less important after autologous than allogeneic HCT.

### **Why Vaccinate?**

- Let transplant recipients enjoy the same protection from vaccine-preventable diseases as the general population.
  - Haemophilus influenzae type b
  - Neisseria meningitidis
  - Diphtheria
  - Tetanus
  - Pertussis
  - Poliomyelitis
  - Hepatitis B
- Protect against infectious diseases that occur more frequently in transplant recipients than in the general population, or are more severe in transplant recipients, in particular:
  - Influenza virus
  - Covid-19
  - Streptococcus pneumoniae
  - Varicella zoster virus

### **When to Revaccinate?**

- Depends on multiple considerations, which were taken into account when creating the schedule and should be taken into account by clinicians when adjusting the schedule to a specific patient
- B cell counts recover to normal at 3-6 mo, memory B cells later
  - In case of B cell depleting antibodies (eg, rituximab), B cell counts are near-zero for 6 mo after last dose. If a patient was treated with a B cell depleting antibody posttransplant, delay start of vaccination till at least 6 mo after the last antibody dose.
- CD4 T cell counts recover to normal at >1 year, but T cell responses are detectable earlier
  - In case of T cell depleting antibodies (eg, rabbit ATG for GVHD), T cell counts are very low for 6 mo after last dose. If a patient was treated with a T cell depleting antibody posttransplant, delay start of vaccination till at least 6 mo after the last antibody dose.
- Antigen consideration
  - Ab responses to recall protein antigens (eg, diphtheria toxoid, tetanus toxoid) recover early
  - Ab responses to neoantigens (eg, hepatitis B vaccine in individuals not vaccinated and not infected pre-transplant) and to polysaccharides (eg, pneumococcal polysaccharide vaccine [Pneumovax]) recover late, particularly late in patients with GVHD
    - For polysaccharides, the response occurs earlier and even in patients with GVHD if conjugated to a recall protein (eg, pneumococcal polysaccharide-protein conjugate vaccine [Prevnar])

- Need for immediate vs long-term immunity
  - The later the start of immunization, the higher and probably more durable Ab responses
  - On the other hand, low response early postHCT may be better than no response to confer at least some protection against influenza, Covid-19, and S.pneumoniae.
- Live vaccine consideration
  - Safety documented in patients at 2 y posttransplant
    - If no relapse
    - If no active GVHD
    - Off of immunosuppressive drugs for at least 3 mo
    - Off of IVIG for 7 months (efficacy of live vaccines is decreased with IVIG; wash-out of 3 months is probably sufficient; however, Public Health official recommendation is to wait 7-11 months)
  - Probably safe as early as 1 year posttransplant, so could be used during outbreak
- GVHD status consideration
  - Patients with active GVHD and/or treated with systemic immunosuppressive drugs mount lower antibody responses to vaccines than patients without GVHD/off of immunosuppressive drugs. However, even the low response is thought to protect at least some patients from influenza, Covid-19, or pneumococcal disease. Given that protection against influenza, Covid-19, and pneumococcus is more important in these patients (compared to patients without GVHD/off of immunosuppressive drugs), immunization with non-live vaccines should not be delayed due to GVHD/immunosuppressive therapy. Live vaccines are contraindicated.
- Malignancy status consideration
  - Patients with relapsed original malignancy or second malignancy treated with chemotherapy, radiation or comfort measures only should not get any vaccine. Live vaccines are contraindicated and non-live vaccines are probably ineffective and/or futile.
- Maintenance therapies consideration
  - Patients on post-transplant maintenance therapies may receive non-live & live vaccines at the discretion of attending physician.
    - For maintenance rituximab or T-cell depleting antibodies, start of vaccination with non-live vaccines should be delayed until  $\geq 6$  months after the last dose of rituximab, daratumomab or T-cell depleting antibodies.<sup>2</sup>
    - Live vaccines can be started at  $\geq 12$  months after the last dose of rituximab (opinion, no data exist).
    - Maintenance lenalidomide and bortezomib are not contraindications to vaccination
      - Non-live vaccine safety and efficacy is not jeopardized by lenalidomide.<sup>3</sup>
      - Live vaccines are safe (if given  $\geq 2$  y postHCT and no relapse) but no data exist on efficacy.<sup>4</sup> Given that multiple myeloma patients are always at risk of

relapse, it is recommended to continue valacyclovir indefinitely and forego live vaccines.

#### **Donor Vaccination:**

- Theoretically useful and possibly practical only for
  - Pneumococcal Conjugate Vaccine and Influenza Vaccine, unknown for Covid-19
  - Related donors
  - If vaccine can be given at least 10 days before stem cell collection
  - Consider immunizing the donor if recipient at high risk of GVHD

#### **Close Contact Vaccination (eg, Vaccination of Family Members):**

- Important for influenza and Covid-19
- Recommended for VZV if no history of chickenpox or shingles or vaccination, or for seronegative family members; however, practicability is limited
  - If a family member or a health care worker vaccinated with a VZV vaccine (live) develops a vesicular rash, there is a small chance of transmitting the virus and, theoretically, causing VZV disease in the immunocompromised patient. Thus, it may be prudent to advise VZV vaccinees that if they develop a rash within 6 weeks post-vaccination, they should avoid contact with immunocompromised patients, particularly VZV seronegative immunocompromised patients.

#### **Non-Routine Vaccines:**

- Funding
  - If used for medical/occupational reason, funded by Alberta Public Health. Examples:
    - Hepatitis A for illicit drug users or patients with chronic liver disease
    - Rabies for handlers of potentially rabid animals
    - Salmonella typhi for close contacts of carriers or lab workers
  - If used for travel reason, NOT funded by Alberta Public Health. Examples:
    - Hepatitis A
    - Salmonella typhi
    - Tick-borne encephalitis
    - Japanese encephalitis
    - Yellow fever (live)
- Timing
  - Non-live vaccines can be given already at 6-24 mo posttransplant; however, immunogenicity is limited. If travel is planned at 2 ½ y posttransplant or later, vaccinate at 24 mo. In case of GVHD, wait until at least 3 mo after immunosuppressive drugs have been discontinued and GVHD inactive.
  - Live vaccines (yellow fever) can be given at 24 mo (if off of immunosuppressive & maintenance therapy drugs)
    - Disclaimer: Probably safe, however, data is limited.

### **Shingrix (Recombinant Zoster Vaccine):**

- Based on the results of a large, international randomized placebo-controlled trial, Shingrix is safe and effective in autologous-HCT recipients when given 50-70 days post HCT, with a second dose 1-2 months later.<sup>6</sup> In Alberta, Shingrix will be given at 6 and 7 months post-HCT to align with current non-live vaccine clearance practices. Those receiving post-HCT maintenance therapies for multiple myeloma (ex. bortezomib/lenalidomide) were included in the trial, although detailed outcomes for these patients have not been presented. Non B-cell or T-cell depleting maintenance therapies, therefore, are not a contraindication to vaccination with Shingrix.
- While Shingrix appears to be safe after allo-HCT, data on efficacy are mixed and there are no data on safety or efficacy in recipients of CAR-T (Reynolds, doi: [10.1111/tid.14109](https://doi.org/10.1111/tid.14109))..
- Allo-HCT and CAR-T recipients late (≥3 years) post-transplant/therapy may choose to receive Shingrix per general population guidelines. This is probably safe but the incremental efficacy of receiving Shingrix after completing the ABMTP VZV vaccination strategy is unknown. If patients choose to receive Shingrix, Shingrix should be administered at least 1 year after completing live Varicella vaccination per the ABMTP vaccination schedule. As in the general population, Shingrix in this setting is not publicly funded.

### **Recommendations for HCT Recipients that have Missed Post-HCT Vaccinations:**

- Those who had vaccination delayed due to ongoing immunosuppressive therapy and/or chronic GVHD should receive all vaccines per the post-HCT immunization schedule.
- Those who missed vaccination for other reasons (for example, receipt of HCT before routine immunization protocol developed (pre-2008) or non-compliance/missed appointments with public health) should receive routine vaccinations per the general population:
  - 2 Td and 1 dTap
  - 2 MMR
  - 2 live attenuated Varicella
  - Hep B series if born in 1981 or later
  - One Pneumo-P for 65 years and over
  - Covid-19 per current Public Health recommendations

### **Special Topics:**

#### ***Vaccination for COVID and Influenza Earlier than 6 months Post-HCT***

- COVID and Influenza infections are associated with morbidity and mortality, especially in the first 6 months post-HCT.<sup>8</sup> Similar humoral and cellular immune responses to COVID mRNA vaccines have been noted in those vaccinated at <4 months versus 4-12 months post allo-HCT.<sup>9</sup> Responses to influenza vaccination have also been documented as early as 3 months



post-HCT.<sup>10</sup> With both vaccines, thus, there is support to begin vaccination at 3 months post-HCT depending on current season (for influenza) or current epidemiology (for COVID). Physicians will have discretion to clear auto/allo-HCT and CAR-T recipients as early as 3 months post-transplant for these vaccines.

### ***Vaccination in the Setting of an Outbreak.***

- The resurgence of a vaccine-preventable illness in the community is possible. In the case of an outbreak, vaccination for the relevant pathogen may be considered earlier than recommended in the routine post-transplant schedule. For example, MMR vaccination at 1 year post-transplant has been found to be safe in retrospective cohorts of HCT recipients in the context of measles outbreak.<sup>11</sup> The decision to vaccinate earlier than planned should be made as a program and in consultation with public health and infectious diseases.

### ***Vaccination after CAR-T Cell Therapies***

Developing recommendations for vaccination after CAR-T therapies is challenging because:

- There are no studies detailing the immunogenicity, efficacy, or safety of vaccination after CAR-T therapy.
- There are a lack of consensus guidelines and limited expert opinion.
- Patients have received varying pre-CAR-T therapies that will affect post-CAR-T risk of infection as well as immunogenicity, efficacy and safety of vaccines (i.e. allogeneic-HCT vs. autologous-HCT vs no HCT before CAR-T cell therapy).
- Patients will receive varying post-CAR-T therapies that will affect risk of infection as well as immunogenicity, efficacy and safety of vaccines (i.e. no therapy vs. allogeneic-HCT).
- Varying CAR-T cell constructs/targets (i.e. CD19 vs. BCMA vs. others) will lead to varying quantity/quality of immunodeficiency and varying paces of immune reconstitution. Relevant but limited data are available for CD19 CAR-T therapies and no data are available for other constructs/targets.
- Specifically, after CD19 CAR-T therapy, nearly all responding patients will rapidly develop B-cell aplasia which may persist >1 year post-therapy. However, ~50% of recipients will experience B-cell recovery beginning at 6-12 months post-therapy.<sup>12</sup>
- Despite B-cell aplasia, pre-existing humoral immunity to vaccine-related antigens may be preserved in CD19 CAR-T recipients due to the persistence of CD19 negative plasma cells.<sup>13</sup>

Despite the limitations noted above, infections remain the leading cause of late mortality after CAR-T<sup>14</sup> and it is reasonable to vaccinate patients empirically, in accordance with expert opinion/guidelines, until further data become available. In Alberta, we will vaccinate patients according to the schedule developed for allo-HCT recipients.

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# Umbilical Cord Blood Transplantation

Presented by: Adam Bryant

## Summary

- For adults, cord blood as a stem cell source can be considered when no HLA-matched or haploidentical donor is available.
- Total nucleated dose and degree of HLA-matching are the most important factors when selecting units for cord blood transplantation
- For single umbilical cord blood transplantation in malignant conditions:
  - TNC at freezing must be  $\geq 2.5 \times 10^7/\text{kg}$
  - In comparing available units with  $\text{TNC} \geq 3.0 \times 10^7/\text{kg}$ , prioritize highest degree of HLA match. Increased TNC dose beyond this threshold is not associated with improved outcomes
  - In comparing smaller dose units ( $\text{TNC} < 3.0 \times 10^7/\text{kg}$ ) prioritize dosing over HLA match
  - HLA matches must be 4/6, 5/6, or 6/6 at HLA-A or -B antigen and -DRB1 allele level matching
  - Where possible, higher TNC doses (ie  $\geq 3.5 \times 10^7$ ) are preferred for units with 4/6 HLA disparity
  - HLA-A or -B mismatch is preferable over DRB1 mismatch
  - Donor specific antibodies are preferred to be absent
  - Other factors to consider if multiple units available (see text)
    - Unit quality based on collection center characteristics
    - High-resolution allele-level HLA-matching
    - Availability of RBC-depleted units
    - Unit CD34+ count
- For non-malignant conditions
  - Higher TNC doses are required ( $\geq 5.0 \times 10^7/\text{kg}$ )
  - Wherever possible a high degree of HLA matching is preferred
  - Absence of HLA-antibodies likely has greater importance than in transplant for malignant disease
- Double unit cord blood transplantation is feasible if no adequate single unit is available. Selection of DUCBT units involves
  - Selection two best available cord blood units, each with minimum  $\text{TNC} \geq 1.5 \times 10^7/\text{kg}$  (preferred  $\geq 2.0 \times 10^7/\text{kg}$ ) and best HLA match to recipient
  - Unit-unit HLA match should not be considered in selection of double unit graft since there is no association with sustained engraftment or speed of neutrophil engraftment
- Conditioning for UCBT is myeloablative and uses our center's preferred regimen (Flu/Bu/low dose TBI). This and many other myeloablative regimens have been reported without clear superiority of one regimen over another

- GVHD prophylaxis after UCBT is with mycophenolate mofetil (~15 mg/kg bid) to day 35 and cyclosporine to day 84 and. Standard dose methotrexate is not used as it has been associated with delayed engraftment/graft failure. Our center may however consider exploring low dose methotrexate regimens in the future based on reports of promising outcomes with respect to engraftment and toxicity (see text). ATG is not used as it has been associated with prohibitively high rates of posttransplant infection.
- Red blood cell replete units will be thawed and washed to remove cellular debris prior to infusion. Buffy coat and red blood cell depleted units will be thawed and diluted. DMSO content for thawed and diluted products will not exceed 5 mL/kg of 20% DMSO solution per day.

## Background

The first successful umbilical cord blood transplantation was performed in 1989, on a 5-year old male with severe Fanconi's anemia who received cord blood stem cells from an HLA-identical sibling<sup>1</sup>. Decades later his graft remains durable with no evidence of disease. Since that time, umbilical cord blood stem cells have become a well-established source of hematopoietic stem cells for allogeneic stem cell transplantation. More than 40,000 patients have undergone UCBT for malignant and non-malignant conditions. Owing to the increased use of haploidentical transplants donors as stem cell sources the use of umbilical cord transplants has been on a steady decline in Calgary and North America since the early 2010s.

When selecting a donor source for hematopoietic stem cell transplantation, haploidentical related donors and mismatched unrelated donors are feasible options for those without matched related or unrelated donors, as outlined in the AHS BMT Standard Practice Manual chapter on Donor Selection. In cases where a search for matched related, unrelated, or haploidentical donors is or is suspected to be unfruitful a simultaneous cord blood search should be performed, especially if transplantation is urgent.

Advantages of umbilical cord blood transplantation include:

- Rapid availability – median 25-36 days sooner than unrelated volunteer marrow/blood stem cells
- Larger donor pool – tolerance of 1-2/6 HLA mismatches (i.e. 4-6/6 HLA-A, -B antigen, and DRB1 allele)
- Lower incidence and severity of acute graft-versus-host-disease (GVHD)
- Lower incidence of chronic GVHD
- Lower risk of viral transmission (e.g. CMV, EBV)
- Lack of donor attrition
- Lack of risk to donor

Disadvantages of umbilical cord blood transplantation include:

- Lower number of progenitor cells and stem cells – higher risk of graft failure, delayed engraftment
- Delayed immune reconstitution – increased risk of infection leading to death
- Not possible to obtain more cells for future treatment (e.g. donor lymphocyte infusion, second transplant)
- Genetic history of donor unknown

## Umbilical Cord Blood Transplantation

There is little randomized clinical trial data comparing transplantation of umbilical cord blood (UCB) vs. related or unrelated marrow or peripheral blood stem cell donors. The best data available comes from retrospective single center and registry data available for both children and adults.

In adults, the large retrospective EBMT/CIBMTR (European Group for Blood and Marrow Transplantation / Center for International Blood and Marrow Transplant Research) study compared leukemia-free survival for umbilical cord blood, peripheral blood progenitor cell, and marrow transplantation in 1525 patients aged 16 or older<sup>5</sup>. When compared to 7-8/8 allele-matched peripheral blood or marrow (matched unrelated [MUD] and mismatched unrelated donor [MMUD]) transplantation, UCB transplantation had comparable leukemia-free survival, higher transplant-related mortality, and lower rates of graft-vs-host disease. The authors concluded that data support UCB transplantation (UCBT) for adults with acute leukemia when no HLA-matched donor is available for urgent transplants. Similarly a large 2020 Japanese registry study including 4150 adult patients comparing 7/8 MMUD (n = 488) to UCB (n= 3662) acute myeloid leukemia (AML) transplant recipients from 2008 to 2017 reported comparable OS (46 v 54 %; HR 1.01; p= 0.89), NRM (HR 1.16; p=0.16), and relapse rates (HR 0.85; p = 0.08) in both groups, and decreased grade II-IV aGVHD (HR 0.76; p<0.001) cGVHD (HR 0.77, p = 0.002) in UCBT patients<sup>27</sup>.

Other comparisons have reported advantages of MMUD transplant over UCBT. A 2019 retrospective EBMT registry study including 2963 total MUD, MMUD, and UCB transplant recipients recently reported lower relapse incidence (HR = 0.8, p = 0.02) lower NRM (HR = 0.7, p = 0.008), improved GRFS (HR = 0.8, p = 0.01), and improved OS (HR = 0.7, p < 0.001) in 9/10 MMUD (n=677) compared to UCB recipients (n=285)<sup>28</sup>. Given the absence of consistent data favoring MMUD over UCB transplant, selection between these donor sources should be done on a case by case basis factoring in donor, disease, and patient characteristics.

Retrospective comparisons, meta-analyses, and small prospective trials have recently reported data supporting improved transplant outcomes with haploidentical (haplo) donor hematopoietic cell transplant (HCT) when compared to UCBT<sup>29-33</sup>. In two meta-analyses from 2020<sup>29</sup> and 2019<sup>32</sup> with overlapping study inclusion reporting on 2,793 (1,432 haplo; 1,361 UCB) and 3,434 (1,759 haplo, and

1,675 UCB) mostly adult transplant recipients for hematologic malignancies, haplo HCT was associated with decreased rates of acute GVHD, increased rates of chronic GVHD, decreased non-relapse mortality, improved relapse-free survival, and improved overall survival.

Umbilical cord allotransplantation at our center is thus considered in cases where a suitable matched related, unrelated, or haploidentical donor is not available. Mismatched unrelated donors are preferred over cord blood sources, but this selection should nonetheless be reviewed on a case-by-case basis.

## Selection of Cord Blood Unit for Single Unit Cord Blood Transplantation

### Cell Dose & HLA Match

Both the total nucleated cell dose and degree of HLA-match of the umbilical cord blood unit in single cord blood transplantation have a strong impact on survival via effect on transplant-related mortality. In a large 2010 retrospective single center analysis of 1061 predominantly pediatric recipients of single-unit myeloablative UCBT for the treatment of hematological malignancies from 1993 to 2006, the best transplantation outcomes were in recipients of 6/6 units regardless total nucleated cell (TNC) dose, though median dose was notably  $4.0 \times 10^7/\text{kg}$ <sup>6</sup>. Recipients of 4/6 HLA-matched units required a TNC  $\geq 5.0 \times 10^7/\text{kg}$  to achieve comparable TRM and RFS to that of recipients of 5/6 units with a TNC of  $\geq 2.5 \times 10^7/\text{kg}$ . This study may suggest that the greater the degree of HLA disparity, the higher the required TNC dose to ensure transplantation survival<sup>6</sup>.

More recent reports have suggested that beyond a minimum TNC threshold, HLA disparity may not have adverse impact of NRM or other transplant outcomes. A 2019 Japanese registry study of 1,355 adults receiving UCB allogeneic HCT (alloHCT) between 2003 and 2016, including only those with minimum TNC dose of  $2 \times 10^7/\text{kg}$  compared pairings above or below the median TNC dose ( $2.68 \times 10^7/\text{kg}$ ) and with either 1 or 2 mismatches<sup>34</sup>. No advantage was seen in those with the least degree of mismatch and higher cell dose when compared to other pairings with greater mismatch, smaller cell dose, or both. TNC in the highest quartile was associated with improved ANC engraftment rates when compared to the lowest quartile, but higher TNC dose quartiles were otherwise not associated with improved engraftment, relapse, or survival outcomes, suggesting against benefit of increased TNC dosing beyond a given minimum threshold. In the large 2014 EBMT and Eurocord study addressing the impact of allele-level matching on outcomes, in multivariate analysis, the effect of a TNC count  $\geq 3 \times 10^7/\text{kg}$  on NRM was independent of HLA disparity, and held true when patients were analyzed as pediatric and adult subcohorts<sup>35</sup>. In a phase 3, 320-patient multicenter randomized controlled trial (RCT) comparing single UCB (SUCB) to double UCB (DUCB) alloHCT, single units were eligible if above a TNC of  $2.5 \times 10^7/\text{kg}$  regardless of HLA mismatch, included 39% 4/6 mismatches, and demonstrated similar outcomes with respect to neutrophil engraftment, relapse, PFS, and OS when compared to DUCBTs<sup>36</sup>. The 2019 NMDP/CIBMTR Guidelines on cord unit selection suggest a



minimum  $2.5 \times 10^7/\text{kg}$  minimum dose per single unit and do not specify higher minimum doses for differing degrees of HLA disparity<sup>37</sup>.

Other studies<sup>7-12</sup> consistently demonstrate cell dose to be the most important factor on survival outcomes, and the EBMT and NMDP/CIBMTR guidelines have recommended using  $\geq 2.5-3 \times 10^7$  total nucleated cells/kg at collection for patients with malignant disease and  $\geq 5.0 \times 10^7$  nucleated cells/kg for those with non-malignant disease<sup>7, 37</sup>.

An increasing number of HLA mismatches is associated with delayed engraftment, higher treatment-related mortality, higher rates of chronic GVHD, and in malignant diseases, decreased relapse rates<sup>7</sup>. As greater HLA disparity is not associated with improved relapse outcomes in nonmalignant diseases, optimal HLA matching is important in these cases. When choosing between multiple cord units that meet the minimum cell dose requirement, optimal HLA matching should be prioritized. Memorial Sloan-Kettering Cancer Center (MSKCC) guidelines for single UCBT suggest a minimum nucleated cell dose of  $2.5 \times 10^7/\text{kg}$  with 1 or 2 mismatches at the HLA-A, -B antigen, or -DRB1 allele<sup>13</sup>. There is no data to guide dosing of TNC by actual versus ideal or adjusted body weight, thus the dose should be based on the patient's actual weight at time of transplantation.

Based on the above data, at our center selection of a single umbilical cord blood transplantation in malignant conditions requires a minimum TNC at freezing of  $\geq 2.5 \times 10^7/\text{kg}$ . When comparing available units with TNC  $\geq 3.0 \times 10^7/\text{kg}$ , the highest degree of HLA match should be prioritized, as increasing TNC dose beyond this threshold is not associated with improved outcomes. In comparing smaller dose units (TNC  $< 3.0 \times 10^7/\text{kg}$ ) we prioritize dosing over HLA match. Where possible higher TNC doses (i.e.  $\geq 3.5 \times 10^7$ ) are preferred for units with greater (i.e. 4/6) HLA disparity.

HLA matching in UCBT is based on HLA antigen typing for -A and -B, and allelic typing for HLA-DRB1. A single institution retrospective analysis of 79 adults with AML who received single unit UCBT was analyzed for the impact of directional donor-recipient HLA disparity using allele-typing at HLA-A, -B, -C, and DRB1<sup>14</sup>. With the extended high-resolution typing, the donor-recipient compatibility ranged from 2/8 to 8/8, but this did not have a negative impact on non-relapse mortality, GVHD or engraftment. The 5-year cumulative incidence of relapse was 44% vs. 22% for patients receiving UCB units matched  $\geq 6/8$  or  $< 6/8$ , respectively ( $p=0.01$ ). On multivariate analysis higher HLA-disparity in the GVH direction and first complete remission at time of transplantation were the only variables significantly associated with an improved DFS. The effect of allele-level matching on non-relapse mortality in 1568 single umbilical cord blood transplantations for hematological malignancy was published in 2014<sup>15</sup>. Only 7% of donor-recipient pairs were matched at HLA-A, -B, -C, and DRB1; 15% were mismatched at one, 26% at two, 30% at three, 16% at four, and 5% at five alleles. Only 54% of units matched at HLA-A, -B, and -DRB1 were actually matched at the allele-level at all loci. Non-relapse mortality was higher with units mismatched at one (26%), two (26%), three (34%), four (37%), or five alleles (41%) compared to HLA-matched units (9%). Cell dose  $< 3.0 \times 10^7/\text{kg}$  was

associated with higher NRM independent of HLA-match. Neutrophil recovery was lower with mismatches at 3-5 alleles but not at 1 or 2 alleles. These data support allele-level HLA-matching in the selection of single UCB units whenever possible.

### **Donor Specific Antibodies (DSA)**

Since most UCBT are performed with HLA-mismatched UCB units, the presence of anti-HLA donor-specific antibodies in the patients against the UCB can result in failure or delay of engraftment. Anti-HLA antibodies before transplant may occur due to alloimmunization to HLA through blood transfusions, pregnancy, and also in some unexposed individuals. In the UCBT setting, few studies with controversial results are available on the impact of DSA on outcomes. One analysis showed an increased risk of graft failure and lower survival for patients with positive DSA undergoing single (n=386) or double (n=73) UCBT<sup>16</sup>. Another report showed no association between the presence of DSA and transplant outcomes in 126 double UCBT recipients<sup>17</sup>. Presence of DSA was found to be associated with higher 1-year TRM (46 v 32%; p=0.06) and lower engraftment (44% vs. 81%, p=0.006) in patients with or without antibodies, respectively<sup>18</sup>. Based on these data, whenever possible, it is important to avoid selecting a unit when the patient has donor specific anti-HLA antibodies.

### **Other factors to consider in selecting cord blood units:**

- Unit quality, measured by practices of the bank with from which it originates, is associated with increased unit potency including post-thaw CD34+ cell recovery and viability. Where possible it is desirable to obtain cord blood units that were more recently cryopreserved, from banks that are FACT-accredited, that store RBC-depleted cord units, at adequate volume ( $\geq 25$  cc thawed volume per unit), and that are closer in location<sup>37</sup>.
- CD34+ cell count can be considered when choosing between multiple cord units that are otherwise similar from the same bank. NMDP/CIBMTR guidelines suggest a minimum CD34+ count of  $\geq 1.5 \times 10^5$  kg<sup>37</sup>. Given variation in and non-standardization of CD34+ count and viability measurements, interpretation of reported CD34+ count should be done cautiously, particularly if it appears discrepant from TNC dose<sup>37,38</sup>.
- Red blood cell (RBC) content of the unit. Buffy coat enriched and RBC depleted units should be considered over RBC replete units. RBC replete units contain red cell debris and free hemoglobin, which can be associated with infusion reaction and washing of these RBC replete units can result in progenitor cell loss.
- Natural killer cell immunoglobulin-like receptor mismatch, non-inherited maternal antigens and inherited paternal antigens may influence decisions about which units to select in the future

## Double Unit Umbilical Cord Blood Transplantation

The use of single unit UCBT is limited since many adults do not have access to a single cord blood unit with the recommended TNC dose. Strategies for ex-vivo expansion of UCB units are being actively explored in the literature with some early phase I/II trials showing promise<sup>39-42</sup>. These strategies have yet to be standardized or widely adopted and warrant further exploration.

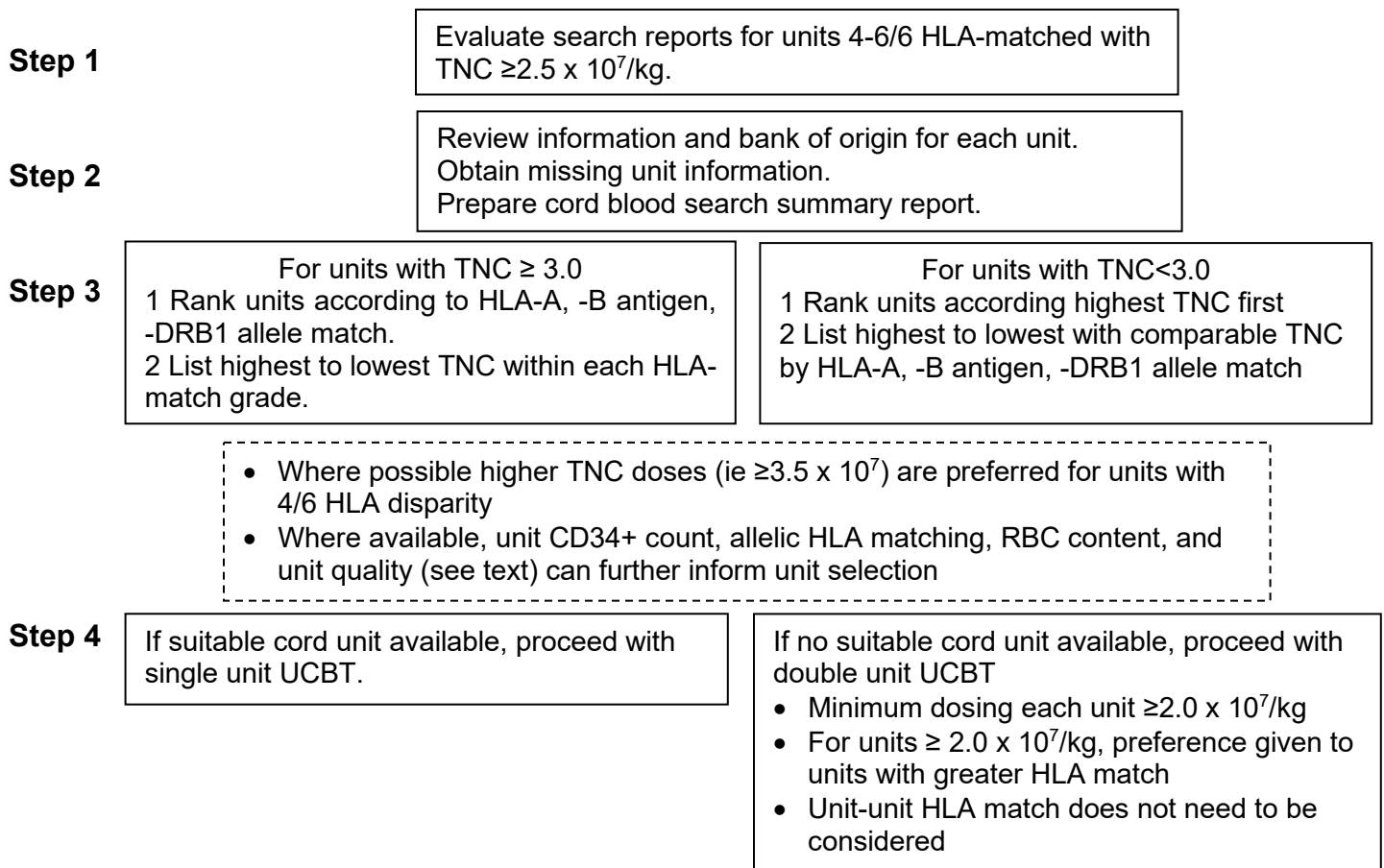
Double unit UCBT as a strategy to augment cell dose of the graft has been successful with improved sustained donor engraftment and post-transplantation survival compared with historic SUCBT controls<sup>19</sup>. Sustained hematopoiesis is accounted for by only one of the two units, with demonstration of dominance as early as Day +21 post-transplant. Higher CD3<sup>+</sup> cell dose and percentage of CD34<sup>+</sup> cell viability was associated with unit dominance<sup>19</sup>. The exact biological mechanism responsible for single-donor predominance after double unit CBT remain incompletely understood. Accumulating reports support the use of DUCBT to overcome the cell dose barrier in adults. In one study with 177 patients who underwent myeloablative UCBT, DUCBT recipients experienced lower relapse rates (19 vs 34% at 5-years), higher rates of GVHD (acute GVHD 48 vs 29%; chronic GVHD 18 vs 10%), and higher RFS (51 vs 40%) when compared to SUCBT recipients<sup>20</sup>.

Given that either unit may engraft after a double unit UCBT, each unit of a double-unit graft is equally important and the same unit selection principles should apply to both units. Optimal strategies for balancing cell dose against HLA match are not well established. Of note there is no relationship between unit-unit HLA match and the likelihood of sustained donor engraftment<sup>19</sup>. When analyzing HLA-A or -B antigen or -DRB1 allele mismatches or 10/10 HLA-allele matches in 84 recipients of double unit UCBT, there was no association between unit-unit HLA match and sustained engraftment, speed of neutrophil engraftment, or unit dominance<sup>19</sup>. Thus, unit-unit HLA match should not be considered in the selection of a double-unit graft, particularly at the expense of available grafts with higher cell doses. MSKCC guidelines recommend each DUCBT have a minimum TNC of  $\geq 2.0 \times 10^7$  /kg, with preference for HLA matching above this TNC threshold, and unit-unit HLA match is not considered<sup>13</sup>. NMDP/CIBMTR guidelines and most DUCBT studies in adult and pediatric populations recommend or require a minimum TNC dose of  $\geq 1.5 \times 10^7$ /kg for DUCBT, suggesting a less stringent TNC threshold may be safe and effective<sup>37,43</sup>.

Whether DUCBT is preferable SUCBT when the cell dose in one unit is acceptable is unknown in the adult setting. In The Blood and Marrow Transplant Clinical Trials Network randomized trial (BMT CTN 0501) was a phase III trial that randomized patients to receive a single (n=113) or double (n=111) UCBT, with median pre-cryopreserved TNC dose of 4.8 and 8.9  $\times 10^7$ /kg, respectively<sup>21</sup>. The results of this study showed no survival advantage after DUCBT compared to SUCBT in children with hematological malignancies (1 year OS 65% vs. 71%, p=0.13). Except for a higher risk of grade III-IV acute GVHD in recipients of a double UCBT, all outcomes were similar between the two groups. Similar findings were reported in a large 2019 systematic review comparing DUCBT to SUCBT that included the above RCT and 24 additional retrospective and prospective reports. A total of 6571 total

patients were included, with 12 of 25 studies enrolling adults exclusively<sup>43</sup>. DUCBT was associated with higher incidence of acute and extensive cGVHD but with lower relapse incidence, and outcomes were ultimately similar between DUCBT and SUCBT for NRM, RFS, and OS. Authors suggested the increased GVH and presumed GVL effect reported here may suggest that DUCBT could be considered in patients with higher risk disease, but this possibility has yet to be explored in a prospective randomized fashion.

## Schema for Unrelated Cord Blood Unit Selection



## Infusion of Cord Blood Units

Cord blood units are processed and infused according to established standard operating procedures. Processing requirements for Cord Blood Units are determined by transplant physician, in consultation with the Cellular Therapy Laboratory (CTL), prior to planned infusion. The following considerations are taken into account when determining processing requirements:

### 1. Red Cell Content

- Buffy coat and Red Cell Depleted units are typically thawed and diluted for Adult Recipients<sup>22</sup>
- Red Cell Replete units are thawed and washed for Adult Recipients<sup>22</sup>

### 2. DMSO content

- DMSO content in thawed & diluted products should not exceed 5 mL/kg/day of 20% DMSO

### 3. Infusion of Double Cords for transplant

- For double cord blood transplants, the first unit must be thawed, processed and administered safely prior to thaw and processing of the second unit.

## Conditioning

Non-relapse mortality has historically been an important contributor to adverse outcomes in UCBT. In the aim of reducing NRM, many studies have explored reduced intensity conditioning regimens. The largest and most recent comparison of conditioning approaches was reported in a 2016 EBMT Eurocord registry study involving adults transplanted for AML with SUCBT or DUCBT from 2004 to 2013<sup>44</sup>. A total 894 adults who underwent myeloablative (MAC, n = 479) or reduced intensity conditioning (RIC, n = 415) were included. MAC regimens were varied and included 40 (8%) patients receiving Flu/Bu based ablative conditioning. Compared to MAC UCBMT recipients, RIC recipients had similar neutrophil engraftment and rates of acute and chronic GVHD. RIC recipients also had a higher incidence of disease relapse, but lower NRM, ultimately resulting in comparable PFS, GRFS, and OS between both groups. MAC regimens were not compared amongst each other.

Given these findings, our center prefers a myeloablative Flu/Bu/low dose TBI conditioning platform, the same regimen as that used for MRD, MUD, MMUD and haploidentical alloHCT at our center, and a regimen with which we have extensive experience.

## GVHD Prophylaxis

Use of ATG has been associated with decreased survival primarily due to infections<sup>23-25</sup>. Use of methotrexate has been associated with delayed engraftment and graft failure<sup>26</sup>. Some evidence suggests that delayed or failed engraftment is lower with a mycophenolate mofetil/calcineurin inhibitor (MMF/CNI) based regimen than with methotrexate<sup>45</sup>. Exploration of reduced-dose methotrexate, often less than half the standard dose, has in some reports been also associated with lower NRM, infectious complications, and improved engraftment<sup>45,46</sup> and in one retrospective three arm study,

comparable engraftment to that seen with MMF/CNI based prophylaxis<sup>47</sup>. Further comparative and prospective studies are required to establish the optimal GVHD prophylaxis regimen in UCBT.

The most frequent GVHD prophylaxis reported in UCBT (including in the US BMT CTN study 1101) remains MMF/Cyclosporine (CSA), which in the absence of clear evidence pointing to an optimal regimen, is the prophylaxis regimen we prefer at our center (MMF to day 35, CSA to day 84).

### **Calgary Results**

Among 22 pts who received UCB between 2004 and 2013 (using CSA+ATG GVHD prophylaxis), 8 patients are alive at ≥5 y posttransplant. Of the 14 patients who died, 4 died due to an infection (not associated with GVHD), 3 due to GVHD, 3 due to relapse and 4 due to other (typically multi-organ failure).



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# ABO Incompatible Graft and Recipient

Presented by Nicole Prokopishyn

## Summary

- Donor/recipient pairs of different blood groups may exhibit major ABO incompatibility (the recipient has pre-formed hemagglutinin antibodies reactive against donor red blood cells), minor ABO incompatibility (the donor has pre-formed hemagglutinin antibodies reactive against recipient red blood cells), or bidirectional (the donor and recipient both have hemagglutinin antibodies reactive against the other).
- There is no consistent evidence that ABO incompatibility unduly influences clinically relevant outcomes (e.g., survival, GVHD).
- An ABO compatible donor is preferred over an ABO incompatible donor to minimize the risk of non-lethal complications like hemolytic anemia or pure red cell aplasia.
- Major ABO incompatibility, including bidirectional incompatibility:
  - For adult recipients, if the red cell volume is >30mL, the product is split into aliquots with no greater than 30mL red cells per unit. If the initial incompatible red cell volume is <30mL, no further action is taken. No more than 30mL of incompatible red blood cells should be infused in a 6-hour period.
  - For pediatric recipients, the accepted range for ABO incompatible blood volume transfused is 0.2 to 0.5 mL/kg. The transplant physician will be contacted with the volume of incompatible red blood cells and will direct Cellular Therapy Lab (CTL) on desired final red blood cell content per infusion bag. CTL will aliquot and/or red cell reduce product as necessary for infusion into the patient.
  - For products with very large volumes of red cells, where dividing into several aliquots is not practical, red cell reduction using the Spectra Optia can be considered.
- Minor ABO incompatibility:
  - No action is taken as local validation data at CTL has indicated no adverse reactions associated with minor ABO incompatibilities.

## Background

Up to 50% of related and 50% of unrelated donor transplants involve an ABO incompatible donor and recipient, not including differences between minor red cell antigens.<sup>1, 2, 3</sup> Donor- recipient pairs with the same ABO blood type are said to be compatible. Minor incompatibility occurs when the donor has antibodies against recipient ABO antigens, and major incompatibility occurs when the recipient carries antibodies against donor red cells. When both occur in the same donor-recipient pair, a bi-directional incompatibility is present, as shown in Table 1 below.<sup>3, 4</sup> Major incompatibility can result in acute hemolytic transfusion reaction at the time of stem cell infusion and delayed red cell engraftment. Minor incompatibility rarely causes hemolysis at the time of transplant due to the infusion of incompatible donor plasma, but can result in a delayed transfusion reaction 7-14 days post

transplant from production of isohemagglutinins by lymphocytes infused with the graft (passenger lymphocyte syndrome). ABO antigens are the primary concern in graft compatibility, though non-ABO antigens such as Rh and Kidd have been reported to cause post transplant hemolysis.<sup>5, 6</sup>

**Table 1.** Donor-recipient ABO compatibility. <sup>3, 4</sup>

| Mismatch Type | ABO Blood Type |       | Potential Clinical Consequence                                | Etiology                                                                                               |
|---------------|----------------|-------|---------------------------------------------------------------|--------------------------------------------------------------------------------------------------------|
|               | Recipient      | Donor |                                                               |                                                                                                        |
| Major         | O              | A,B   | -Acute hemolytic episode                                      | -Transfusion of incompatible red blood cells                                                           |
|               |                | AB    | -Delayed RBC engraftment                                      | -Recipient anti-donor isohemagglutinins                                                                |
| Major         | A              | AB    | -Pure red blood cell aplasia                                  | -Loss of immature stem cells from processing with ABO antigens expressed on granulocytes and platelets |
| Major         | B              | AB    | -Delayed granulocyte and platelet engraftment                 |                                                                                                        |
| Minor         | A              | O     | -Acute hemolytic episode                                      | -Donor plasma with elevated isohemagglutinin titers/small blood volume recipient                       |
| Minor         | B              | O     | -Delayed hemolysis secondary to passenger lymphocyte syndrome |                                                                                                        |
| Minor         | AB             | O,A,B |                                                               | -Passenger lymphocytes producing isohemagglutinins                                                     |
| Bidirectional | A              | B     | -Combination of major and minor consequences                  | -Combination of major and minor etiologies                                                             |
| Bidirectional | B              | A     |                                                               |                                                                                                        |

## Consequences of ABO Incompatible Transplant

The relative importance and discordant consequences of ABO incompatible transplants (as described below) is dependent on the era of transplants, underlying disease, type of transplants (Haplo-identical, Cord), graft source (marrow, PBSC etc.), conditioning (e.g. reduced intensity) as well as the availability of superior supportive measures.

Historically, there have been a number of single center reports as well as four large registry reports<sup>2,7-9</sup> describing the impact of ABO incompatibility on transplant outcomes. Overall the results are inconsistent though some show a negative effect on neutrophil engraftment,<sup>2,7</sup> acute graft versus host disease,<sup>2, 7</sup> non-relapse mortality,<sup>2,8</sup> and overall survival.<sup>2,8</sup> Moreover, an individual patient data-based meta-analysis conducted in 2009 suggests that there is no adverse association between any ABO mismatching and survival.<sup>10</sup>

## **Acute Hemolytic Reaction**

Acute hemolytic reactions occur in 15% of transplants with major ABO incompatibility<sup>11</sup>, and in almost half of those receiving a high volume (>50mL) of incompatible red cells<sup>12</sup> resulting in renal failure and even death in some patients. Transplants with minor ABO compatibility will rarely cause acute hemolysis from the transfusion of donor isoagglutinins against recipient red cells.

## **Delayed Red Cell Engraftment and Pure Red Cell Aplasia (PRCA)**

Recipient antibodies directed against donor red cells (isoagglutinins) are usually cleared rapidly following transplant, with the only consequence being a slight increase in transfusion requirements compared to ABO compatible grafts.<sup>13</sup> Isoagglutinins disappear more rapidly following unrelated donor compared to related donor transplants,<sup>1, 14</sup> and in those with graft versus host disease,<sup>1, 15</sup> and more slowly following non-myeloablative transplant.<sup>16</sup> Persistent anti-donor red cell isoagglutinins can cause delayed red cell engraftment that may persist for months or even years following transplant. In some cases, bone marrow biopsy will show normal erythroid precursors up to the point of expression of the incompatible antigen, with absence of precursors beyond that point reflecting the expression of ABO antigens at different stages of red cell development.<sup>17</sup> There is an increase in transfusion requirements contributing to iron overload.

## **Delayed Transfusion Reaction**

Infusion of grafts with minor ABO incompatibility has rarely resulted in a delayed transfusion reaction, thought to be due to production of anti-host red cell antibodies by donor B-cells infused with the graft. These have mostly occurred 7-10 days after the transplant in red cell group A recipients of group O grafts<sup>18</sup>. Almost all patients had GVHD prophylaxis consisting of cyclosporine without methotrexate.

## **Neutrophil and Platelet Engraftment**

It is not clear if ABO incompatibility can affect neutrophil and platelet engraftment, or contribute to graft failure. Major incompatibility was associated with delayed neutrophil engraftment in 3 registry studies,<sup>2, 7, 19</sup> but was not observed in several other studies, including a recent large CIBMTR/NMDP evaluation of donor characteristics.<sup>8, 9, 20-23</sup> Even if there is a difference, a median 1-2-day delay in engraftment is not likely to be clinically relevant. One registry study suggests delayed platelet recovery with major incompatible grafts.<sup>2</sup> Some single center studies have reported both platelet and neutrophil engraftment issues, but the majority of studies find no impact of incompatibility.<sup>18</sup> A significantly higher rate of graft failure was reported in major or bidirectional incompatible transplants (6/83 vs 0/141 compatible transplants),<sup>24</sup> though one or more HLA mismatches was also present in 3 of the 6 cases. Two small series also suggested a risk of graft failure that was not seen in a number of other reports.<sup>18</sup>



## **Graft Versus Host Disease (GVHD)**

Red blood cell membranes are rich in proteins of great structural diversity. Polymorphisms of these antigens, incompatible ABO antigens, and allelic variations of ABO antigens could serve as minor histocompatibility antigens influencing rates of GVHD. Expression of similar antigens on endothelial and epithelial tissues could serve targets for the donor immune system, inciting a GVH response.

### *Acute GVHD*

Increased rates of grade II-IV aGVHD were reported in two cohort studies<sup>25, 26</sup> as well as two registry studies,<sup>2, 7</sup> but were not seen in most other reports.<sup>18, 23, 27-30</sup>

Interesting, bi-directional mismatching (but not major mismatch) was associated with increased risk of grade II-IV acute graft-versus-host disease in a recent EBMT registry study evaluating leukemia patients undergoing haplo-identical transplants with a HR 2.387; 95% CI: 1.22-4.66; P=0.01.<sup>19</sup> However, the same authors note that patients with minor mismatching transplanted with bone marrow grafts experienced increased grade II-IV acute graft-versus-host disease rates (HR 2.03; 95% CI: 1.00-4.10; P=0.04).<sup>19</sup> In contrast, the effect of ABO mismatch on transplant outcomes and transfusion requirements in 594 patients undergoing reduced-intensity conditioned (RIC) HSCT with alemtuzumab was evaluated in three UK transplant centres and did not demonstrate any association with aGVHD risk.<sup>31</sup> Further, a registry study from CIBMTR evaluating 1,013 AML patients who underwent MMURD transplantation between 2005 and 2014 suggest that the incidence of grade II-IV acute graft versus host disease was marginally lower in patients with major ABO mismatching (HR 0.7, 95% CI, 0.5-1; P = .049).<sup>21</sup> In the absence of clear biologic plausibility and conflicting evidence, such positive associations maybe due to chance.

### *Chronic GVHD*

There are minimal studies that link chronic GVHD with ABO incompatibility. In the before mentioned UK study, the incidence of extensive chronic GVHD was higher in patients with minor and major mismatch compared with those who were ABO matched (hazard ratio (HR) 1.74, P=0.032 for minor, HR 1.69 P=0.0036 for major mismatch).<sup>31</sup>

## **Relapse, Non-Relapse Mortality, and Survival**

There is little evidence to suggest an influence of ABO incompatibility on relapse. None of the four registry studies found this association. One case series reported a decrease in relapse when minor or bidirectional incompatible grafts were used compared to major incompatible or ABO matched grafts on univariate analysis, but this association was not significant on multivariate analysis.<sup>32</sup> By contrast, cohort and registry studies have found an increase in NRM and decrease in overall survival,<sup>2, 8, 18, 33, 34</sup> though these findings were not confirmed by other studies.<sup>7, 20-23, 27-29, 31, 35</sup>

More recently, Kollman et al.<sup>20</sup> re-examined the association of donor characteristics associated with post-HSCT outcomes in the modern HSCT era using data from CIBMTR/NMDP. Utilizing 2

independent datasets: 1988 to 2006 (N = 6349; training cohort) and 2007 to 2011 (N = 4690; validation cohort), they noted a potential association of ABO compatibility with survival in HSCT prior to 2007 with ABO minor mismatch conferring a HR 1.10 (95%CI 1.01-1.18) and ABO major mismatch a HR 1.13 (95%CI 1.05-1.21). However, this association was not seen in the HSCT after 2007 (validation cohort) where the mortality risks associated with minor and major ABO mismatched transplants were HR, 1.09 (95%CI, 0.98-1.23) and HR 1.09 (95% CI, 0.91-1.21) respectively. They also considered the effect of ABO match separately for bone marrow and peripheral grafts and did not see a significant effect of ABO mismatching on overall mortality. Further, ABO compatibility was not associated with NRM, Relapse Mortality, acute or chronic GVHD.

Similarly, the EBMT evaluated the influence of ABO compatibility in 837 patients who underwent haploidentical transplantation and did demonstrate differences in Non-relapse mortality, relapse incidence, leukemia-free survival, overall survival, and chronic graft-versus-host disease rates between ABO-matched and -mismatched patients. However, patients with *major ABO mismatching and bone marrow grafts* had decreased survival (HR=1.82; CI 95%: 1.048 - 3.18; P=0.033). This finding was not observed in a CIBMTR study evaluating the impact of ABO mismatch on transplant outcomes with various graft types.<sup>22</sup>

In contrast, the Chinese developed a risk score utilizing data from 1199 consecutive subjects receiving transplants from an HLA-haplotype-matched relative using granulocyte colony-stimulating factor and anti-thymocyte globulin (n=685) or an HLA-identical sibling (n=514). They suggest that ABO mismatch was 1 of 3 (others were older donor/recipient age, female-to-male transplants) independent risk factors that conferred risk of TRM and LFS.<sup>36</sup>

## Summary

An ABO compatible donor is preferred over ABO incompatible donor, but priority is given to HLA matching, donor age. The relative importance of ABO compatibility over CMV status and female gender/parity is less clear with respect to post-HSCT outcomes.<sup>37</sup>

## Management

The red cell content of the graft is partially dependent on whether the graft is from bone marrow or peripheral blood collection by apheresis. In the later, the red cell content is normally <10ml per collection while is higher and more variable with bone marrow.

The safe volume of transfused incompatible red cells has not been established in large studies. In one case series, sixteen of 36 patients receiving over 50 mL of incompatible red cells experienced signs or symptoms of an acute hemolytic reaction, 10 had renal failure, and 6 died, compared to no deaths, no renal failure, and only 3 hemolytic reactions in 12 patients transfused less than 50 mLs.<sup>12</sup> Thresholds of 20mL and 30mL have been reported as associated with

minimal toxicity. The risk of acute hemolytic reactions can be reduced by decreasing the 1) red cell content of the graft, or 2) the isoagglutinin titers of the recipient.

Red cell depletion of the HPC product can reduce the total nucleated cell count.<sup>38</sup> It has been suggested that this may be of importance if the HPC content is low or if additional cells are not readily available, as with cord blood units or volunteer unrelated donor grafts.<sup>39</sup> In addition, because unrelated donor HPC products can come from anywhere in the world, and prolonged intervals between collection and infusion into the recipient are associated with decreased likelihood of engraftment and increased mortality<sup>40</sup>, further product manipulation in these circumstances could be undesirable.

An alternative approach to red cell depletion is to consider isohemagglutinin reduction by plasma exchange for major ABO-incompatible bone marrow grafts. Sheppard et. al report their single centre experience suggests that engraftment times, transfusion requirements, incidence and severity of graft-versus-host disease, and 100-day treatment-related mortality did not differ between the patients with a major ABO donor mismatch and those with an ABO-compatible donor. Further, no hemolytic transfusion reactions were observed during product infusion.<sup>41</sup> This approach has been counter-challenged<sup>42</sup> as antibody titering is a laboratory technique shown to be difficult to standardize across institutions.<sup>43, 44</sup> Therefore, it may be difficult to determine whether a concentration of incompatible antibody can be universally considered to be protective against a hemolytic reaction.

## The Calgary Approach

In Calgary, the Alberta Bone Marrow Transplant Program (ABMTP) work-up obtains donor and recipient blood type information prior to selection of suitable donor for transplant. The transplant physician reviews the donor and recipient blood type information and is responsible for determining compatibility and indicating on the order for stem cell collection the compatibility status of the donor product. Compatibility is determined based on Table 27-1 in AABB Technical Manual (Table 1 above). The Cellular Therapy Laboratory will determine the product compatibility at the time of receipt of a cellular therapy product. If there is major incompatibility, the red cell volume is then determined (SOP: CTL.705 Product Preparation and Process Overview).

1. For pediatric recipients, the accepted range for ABO incompatible blood volume transfused is 0.2 – 0.5 mL/kg. The transplant physician will be contacted with the volume and will direct CTL on desired final RBC content per infusion bag (based on hydration status and renal function of the recipient). CTL will aliquot and/or red cell reduce product as necessary.

2. For adult recipients, less than or equal to 30 mL +/- 1 mL of incompatible red cells will be allowed per infusion bag of apheresis product (HPC(A)). If product contains greater than 31 mL of incompatible red cells the product will be split into aliquots. HPC(M) will be red cell reduced to achieve  $\leq 30$  mL/infusion aliquot. If the initial incompatible red cell volume is  $< 30$  mL, no further action is taken.
3. For products with very large volumes of red cells, where dividing into aliquots is not practical, red cell reduction using the Spectra Optia can be considered (CTL.762 Red Cell Reduction Using the Spectra Optia ®Apheresis System). Note this protocol requires a minimum of 125 mL of red cells.
4. For plasma incompatible transplants (minor incompatibility), no action is taken for any recipient as it has been determined by CTL validation studies that there is no association with adverse infusion reactions and minor incompatibility of products.

Following transplants with minor ABO incompatible grafts, the appropriate red cell type to be transfused cannot be determined by the usual blood bank techniques. Blood bank is notified about these transplants in order to provide appropriate blood product support (see Table 2).<sup>3</sup>

There is little evidence to guide the management of pure red cell aplasia (PRCA) beyond transfusion support until red cell engraftment occurs. There have been case reports of improvement following administration of erythropoietin,<sup>45-47</sup> though this was unsuccessful in other reports.<sup>16, 48</sup> There are also case reports of successful treatment of PRCA with rituximab,<sup>49, 50</sup> plasma exchange,<sup>48, 49</sup> anti-thymocyte globulin,<sup>51-53</sup> bortezomib<sup>54</sup> and donor lymphocyte infusion.<sup>55, 56</sup> There is insufficient evidence to support the routine use of these treatments for PRCA following ABO incompatible transplant.

There is a suggestion that methotrexate based GVHD prophylactic regimens will result in fewer cases of delayed transfusion reactions. However, given that this is so rare, its clinical impact is negligible compared to that of GVHD. The choice of GVHD regimen should therefore reflect optimal management/prevention of graft versus host disease.

**Table 2.** Recommended blood products for compatible and incompatible transplant recipients.

| Recipient | Donor | Compatibility | 1 <sup>st</sup> Choice red cells | 2 <sup>nd</sup> Choice red cells | 1 <sup>st</sup> choice platelets | 2 <sup>nd</sup> choice platelets | FFP           |
|-----------|-------|---------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|---------------|
| A         | A     | Compatible    | A                                | O                                | A                                | O                                | A, AB         |
| B         | B     | Compatible    | B                                | O                                | B                                | O                                | B, AB         |
| AB        | AB    | Compatible    | AB                               | A, B, O                          | AB                               | O                                | AB            |
| O         | O     | Compatible    | O                                | N/A                              | O                                | N/A                              | O, A, A, B, B |
| A         | O     | Minor inc     | O                                | N/A                              | O                                | O                                | A, AB         |
| B         | O     | Minor inc     | O                                | N/A                              | O                                | O                                | B, AB         |
| AB        | O     | Minor inc     | O                                | N/A                              | O                                | O                                | AB            |
| AB        | A     | Minor inc     | A                                | O                                | A                                | O                                | AB            |
| AB        | B     | Minor inc     | B                                | O                                | B                                | O                                | AB            |
| O         | A     | Major inc     | O                                | N/A                              | A                                | O                                | A, AB         |
| O         | B     | Major inc     | O                                | N/A                              | B                                | O                                | B, AB         |
| O         | AB    | Major inc     | O                                | N/A                              | AB                               | O                                | AB            |
| A         | AB    | Major inc     | A                                | O                                | AB                               | O                                | AB            |
| B         | AB    | Major inc     | B                                | O                                | AB                               | O                                | AB            |
| A         | B     | Bidirectional | O                                | N/A                              | B                                | O                                | AB            |
| B         | A     | Bidirectional | O                                | N/A                              | A                                | O                                | AB            |
| Rh+       | Rh-   |               | Rh-                              | N/A                              | N/A                              | N/A                              | N/A           |
| Rh-       | Rh+   |               | Rh-                              | N/A                              | N/A                              | N/A                              | N/A           |

\* Alberta Precision Labs issues psoralen-treated, platelet additive solution-E (PAS-E) containing platelet units obtained from Canadian Blood Services. These are considered “low titer” for anti-A/anti-B isohemagglutinins due to plasma reduction and replacement with PAS-E. This represents a low risk for hemolysis due to ABO incompatibility. In the HSCT setting, the first choice platelet issued by transfusion medicine is the donor’s blood type, the second choice is a group O platelet and the third choice is the 1st expiring unit of any other blood group. This approach seeks to balance plasma compatibility while avoiding anti-A/anti-B isohemagglutinin-mediated platelet clearance.

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# Long-Term Follow-Up

Presented by: Kareem Jamani

## Summary

### Frequency of Follow-Up

- The recommended follow-up interval for allo-HCT recipients between day 80 and 1 year post-HCT is every 4-6 weeks.
- Long-term follow-up visits for allo-HCT recipients should be at least annually.

### Cardiovascular Disease

- All patients should be counselled with respect to lifestyle modifications that reduce the risk of cardiac events, such as tobacco avoidance, adequate physical activity, maintaining a healthy weight and a healthy diet.
- All patients should have yearly evaluation of blood pressure with treatment per established Canadian guidelines (CHEP; <http://guidelines.hypertension.ca/>, essentially target <135/85 for most patients and <130/80 for those with diabetes).
- HCT recipients with established arterial disease should resume secondary prevention as soon as possible after HCT (i.e. ASA, statins, ACE inhibitors)
- For all allo-HCT recipients and selected auto-HCT recipients (those with a history of chest irradiation or cumulative dose of anthracycline  $\geq 250$  mg/m<sup>2</sup>): patients should have a baseline lipid panel and fasting glucose/haemoglobin A1C at 1 year post-HCT and those  $\geq 30$  years old should have these repeated at 2-3 year intervals with calculation of Framingham risk score. Those who are intermediate or high risk by Framingham or who have LDL  $>4$ -5 mmol/L should be initiated on a statin to target lipid values as outlined by the Canadian guidelines (CCS; <https://www.ccs.ca/en/guidelines/guidelines-library>).
- For all other auto-HCT recipients: screening and management per the general population guidelines.
- Lipid and glucose screening may begin earlier than age 30 for those with one or more risk factors for cardiovascular disease including obesity, smoking, family history of early cardiovascular disease, diabetes, chest radiation or history of GVHD requiring systemic immunosuppression.
- For patients who received HCT in childhood, the CCSS cardiovascular risk calculator (<https://ccss.stjude.org/tools-and-documents/calculators-and-other-tools/ccss-cardiovascular-risk-calculator.html>) may supplement risk estimates obtained with Framingham-type calculators.
- Those with a history of cumulative dose of anthracycline  $\geq 250$  mg/m<sup>2</sup> should have yearly history and physical exam for signs/symptoms of CHF and aggressive management of cardiovascular risk factors as outlined above. Echocardiograms at yearly intervals for 5 years post-HCT may be considered for those at highest risk of CHF (i.e. those with one or more risk

factors in addition to anthracycline exposure, including younger age at anthracycline exposure, female sex, chest radiation, hypertension or diabetes).

## Bone Health

- All patients should be counselled regarding lifestyle modifications for bone health including calcium intake 1200 mg/day from all sources, vitamin D 1000 IU/day, smoking cessation, limiting alcohol use & regular weight-bearing exercise.
- At 1 year post-HCT or at the onset of cGVHD requiring systemic therapy (whichever occurs earlier), patients should have an assessment of BMD by dual X-ray absorptiometry (DXA) and subsequently a 10 year probability of fracture calculated with the FRAX clinical assessment tool (<https://www.sheffield.ac.uk/FRAX/>) (except patients with multiple myeloma who are already being treated with bisphosphonates).
- For those who are no longer on immunosuppression at 1 year post-HCT, the Alberta Toward Optimized Practice (TOP) guidelines for osteoporosis ([http://www.topalbertadoctors.org/cpgs/?sid=18&cpg\\_cats=81](http://www.topalbertadoctors.org/cpgs/?sid=18&cpg_cats=81)) should be followed.
- For those who remain on immunosuppression at 1 year post-HCT, therapy should be offered to: 1) those with established osteoporosis (BMD T-score  $\leq -2.5$ ) or history of fragility fracture, and 2) those with a moderate to high probability of fracture by FRAX as outlined by the Alberta TOP guidelines (>10% probability).
- In those who remain on chronic glucocorticoid therapy (prednisone equivalent dose >5mg/day) beyond 1 year post-HCT and who do not initiate therapy, annual DXA measurement should be considered.
- For those who initiate therapy, repeat DXA should be obtained at 3-5 years on therapy.
- First line therapy is typically with oral bisphosphonates. If identified, treatment of hypogonadism also could be considered in men and pre-menopausal women (after evaluation and discussion of risks/benefits of hormone replacement). Referral to an endocrinologist may be considered for alternate or second-line therapy options.
- The need for ongoing bisphosphonate therapy should be reassessed at the end of immunosuppressive therapy and/or at 5 years on therapy.
- In those with osteoporosis, a workup for secondary or contributing causes should be undertaken (for example, hypogonadism, hyperthyroidism & hyperparathyroidism).

## Subsequent Malignancy Screening

- All patients should be counselled regarding smoking cessation and cutaneous solar protection.
- All patients should have a yearly history and physical exam that includes oral cavity, thyroid, and a complete skin exam.
- All patients should visit their dentist for oral/dental examination and cleaning; at least yearly for most patients and every 6 months for those with oral chronic GVHD.
- Screening for breast (see modifications below for women who have received TBI), cervical, colorectal (see modifications below for those who have received high dose TBI or localized

radiation) and prostate cancer should follow established Albertan/Canadian guidelines ([www.screeningforlife.ca](http://www.screeningforlife.ca) & for prostate cancer [www.canadiantaskforce.ca](http://www.canadiantaskforce.ca)).

- For women who have received any dose of TBI: screening mammography starting at age 25 or 8 years after radiation exposure, whichever occurs later but no later than age 40.
- For patients who received high dose TBI ( $\geq 6$  Gy) or abdominal, pelvic, lumbar, sacral, or whole spine irradiation, colorectal cancer screening with colonoscopy should begin at 5 years post-transplant or age 30, whichever occurs later.
- HCT recipients who have received radiation within a particular field, for example chest and those with familial cancer syndromes will require an individualized plan for malignancy screening.
- There are no proven screening measures for t-MN after auto-HCT; yearly CBC with early workup of cytopenias for up to 10 years post-transplant may be considered.

### Renal Disease

- All HCT recipients should have at least yearly monitoring of creatinine.
- Allo-HCT recipients should a spot urine albumin/creatinine ratio yearly.
- All patients should have at least yearly blood pressure evaluation and treatment per Canadian (CHEP) guidelines (essentially target BP  $<140/90$  for most patients).
- Basic medical management of CKD includes initiation of an ACE inhibitor or ARB for proteinuria, tight glycemic control for diabetics, aggressive management of cardiovascular risk factors and avoidance of nephrotoxins.
- Referral to a nephrologist should be considered when estimated GFR is  $<30$  mL/min, for management of CKD with proteinuria or for workup of CKD of unknown etiology.

### Pulmonary Disease

- The approach to prevention of late pulmonary infections are outlined in the bacterial/pneumocystis prophylaxis, fungal prophylaxis and vaccination chapters of these guidelines.
- For autologous-HCT recipients: PFT at 6 weeks post-HCT for those who received potentially pulmonary toxic conditioning. For all, at least yearly history and physical exam for signs and symptoms of pulmonary disease is recommended.
- For allogeneic-HCT recipients: routine PFTs for all patients every 3 months for the first year post-HCT followed by yearly PFTs until 5 years post-HCT. For those with active cGVHD beyond 1 year post-HCT, continued every 3 month PFTs should be strongly considered. History and physical exam should accompany PFTs.
- Abnormal PFTs or new respiratory symptoms should be worked up promptly with CXR +/- NP swab and sputum culture. For clinical presentations that are not consistent with upper respiratory tract infection or community-acquired pneumonia, CT chest and referral to the BMT pulmonary clinic are suggested.



## Endocrine Disease

- Yearly thyroid examination and TSH measurement for all HCT recipients.
- A slow terminal taper of corticosteroids is required for those receiving prolonged courses (>3 weeks) for treatment of GVHD.
- A high index of clinical suspicion for adrenal insufficiency should be maintained when tapering patients from prolonged courses of corticosteroids.
- Management of diabetes and hyperlipidemia within the context of global cardiovascular risk as outlined in the “Cardiovascular disease” section.
- Workup and management of gonadal dysfunction and infertility as published<sup>1,2</sup>.

## Chronic Pain

- HCT recipients with chronic pain should be managed within a multi-disciplinary team that includes HCT providers, pharmacists and the psychosocial team within the Alberta Blood and Marrow Transplant Program with low threshold for referral to the palliative care team at the Tom Baker Cancer Centre.

## Transfusion

- Red cell and platelet transfusion thresholds should be individualized based on clinical circumstances.
- The appropriate blood-group products for transfusion after ABO-incompatible allo-HCT per the “ABO Incompatibility” chapter.
- For allo-HCT recipients: irradiated blood products should be used from start of conditioning until the later of: 1 year post-HCT, end/“burn out” of chronic GVHD or discontinuation of immunosuppression.
- For auto-HCT recipients: irradiated blood products for 7 days prior to stem cell collection and from start of conditioning until 3 months post-HCT, or 6 months post-HCT if TBI was part of conditioning.
- All auto- and allo-HCT recipients should receive standard leukoreduced (“CMV safe”) blood products.
- Transitioning from Pediatric to Adult Post-HCT Care
- Survivors of pediatric HCT are typically followed into adulthood and indefinitely in the Alberta Children’s Hospital long-term follow-up/survivorship clinic.

## Mental Health

- Long-term follow up visits should include screening for mental health concerns. If concerns are identified, patients should be referred to mental health providers and/or community resources as appropriate.



## Background

Survival after hematopoietic cell transplant (HCT) has improved<sup>3</sup>. Survivors, however, face significant health challenges that contribute to morbidity and mortality even late after transplant. Among HCT survivors, the 15 year cumulative incidence of a severe or life threatening chronic health condition, such as stroke, myocardial infarction, diabetes and subsequent neoplasm, is approximately 40%: the cumulative incidence does not differ significantly between recipients of autologous (auto) and allogeneic (allo) HCT<sup>4</sup>. As a result, the risk of death after both allogeneic and autologous HCT, remains significantly higher than that of the general population even many years post HCT<sup>5,6</sup>. Therefore, it is imperative to have a structured long-term follow-up plan for survivors of HCT. This document will summarize the current literature with respect to late effects after HCT and will provide guidelines for clinical practice. The following important aspects of post-HCT survivorship care have already been reviewed in detail elsewhere in the ABMTP standard practice manual and can be found in their respective chapters:

- Diagnosis and management of chronic graft-versus-host disease (cGVHD)
- Management of post-HCT relapse
- Reproductive system complications
- Infection prophylaxis and vaccination

### Frequency of Follow-up

Due to the potential for onset of cGVHD, the recommended follow-up interval for allo-HCT recipients between day 80 and 1 year post-HCT is every 4-6 weeks. Those suffering from GVHD, infection, relapse or other toxicity may need to be evaluated more frequently. Follow-up of allo-HCT recipients beyond 1 year post-HCT and of auto-HCT recipients may be individualized. However, long-term follow-up visits for allo-HCT recipients should be at least annually.

### Transitioning from Pediatric to Adult Post-HCT Care

In Alberta, transition from the pediatric to adult HCT centre for ongoing survivorship/long-term follow-up care occurs when there is agreement between the pediatric survivorship provider, patient and family, and adult survivorship provider. A comprehensive treatment and medical history are forwarded from the pediatric to the adult centre. A shared electronic medical record further enhances continuity of care.

## Cardiovascular Disease

Cardiovascular disease is a major cause of late non-relapse mortality in survivors of HCT. Compared to the general population, HCT survivors have a significantly increased cumulative incidence of cardiovascular death (incidence rate difference 3.6 per 1000 person years) and a significantly higher incidence of cardiovascular risk factors, such as diabetes and hypertension, when compared to age and sex matched controls.<sup>7</sup> Cardiovascular disease after HCT can be conceptualized as arterial disease (cerebrovascular, peripheral arterial and coronary artery disease) and cardiac disease

(particularly congestive heart failure but also constrictive pericarditis and valvular disease) with allogeneic HCT survivors being at higher risk of the former and autologous HCT survivors being at higher risk of the latter<sup>8</sup>.

The cumulative incidence of arterial events among allo-HCT recipients is in excess of 20% at 20 years and the median age at first myocardial infarction is approximately 53 years, which is at least a decade earlier than that of the general population<sup>8</sup>. In a large single centre study, the cumulative incidence of  $\geq 2$  cardiovascular risk factors (of hypertension, dyslipidemia and diabetes) at 10 years after HCT was ~40% for allo-HCT survivors and 26% for auto-HCT survivors<sup>9</sup>. Older age and obesity at HCT, TBI ( $>2$  Gy) and grades 2-4 aGVHD were risk factors for acquisition of cardiovascular risk factors post-HCT. In keeping with these findings, the prevalence of the metabolic syndrome in allo- and auto-HCT recipients is double that of the age-matched general population<sup>10</sup>. Risk factors for occurrence of cardiovascular disease after HCT encompass both traditional risk factors in addition to chest irradiation, GVHD, and exposure to anthracycline chemotherapy<sup>8,9</sup>. Healthy lifestyle choices such as physical activity and fruit/vegetable intake are associated with a lower risk of cardiovascular disease after HCT<sup>11</sup>. Current Canadian guidelines for the general population recommend measurement of a lipid panel and glucose in women and men  $\geq$  age 40 every 5 years<sup>12</sup>.

In a large single centre review, the cumulative incidence of late congestive heart failure (CHF) in auto-HCT survivors was approximately 10% at 15 years post-HCT-a 4.5 fold increased risk over that of the general population<sup>13</sup>. Pre-HCT anthracycline exposure, particularly cumulative dose  $\geq 250$  mg/m<sup>2</sup> is the primary driver of CHF risk, although significant modifiers that increase this risk further include younger age at anthracycline exposure, female sex, chest radiation, hypertension and diabetes<sup>4,8</sup>.

## Recommendations:

- All patients should be counselled with respect to lifestyle modifications that reduce the risk of cardiac events, such as tobacco avoidance, adequate physical activity, maintaining a healthy weight and a healthy diet.
- All patients should have yearly evaluation of blood pressure with treatment per established Canadian guidelines (CHEP; <http://guidelines.hypertension.ca/>, essentially target  $<135/85$  for most patients and  $<130/80$  for those with diabetes).
- HCT recipients with established arterial disease should resume secondary prevention as soon as possible after HCT (i.e. ASA, statins, ACE inhibitors)
- For all allo-HCT recipients and selected auto-HCT recipients (those with a history of chest irradiation or cumulative dose of anthracycline  $\geq 250$  mg/m<sup>2</sup>): patients should have a baseline lipid panel and fasting glucose/haemoglobin A1C at 1 year post-HCT and those  $\geq 30$  years old should have these repeated at 2-3 year intervals with calculation of Framingham risk score. Those who are intermediate or high risk by Framingham or who have LDL  $>4.5$  mmol/L should

be initiated on a statin to target lipid values as outlined by the Canadian guidelines (CCS; <https://www.ccs.ca/en/guidelines/guidelines-library>).

- For all other auto-HCT recipients: screening and management per the general population guidelines.
- Lipid and glucose screening may begin earlier than age 30 for those with one or more risk factors for cardiovascular disease including obesity, smoking, family history of early cardiovascular disease, diabetes, chest radiation or history of GVHD requiring systemic immunosuppression.
- Those with a history of cumulative dose of anthracycline  $\geq 250 \text{ mg/m}^2$  should have yearly history and physical exam for signs/symptoms of CHF and aggressive management of cardiovascular risk factors as outlined above. Echocardiograms at yearly intervals for 5 years post-HCT may be considered for those at highest risk of CHF (i.e. those with one or more risk factors in addition to anthracycline exposure, including younger age at anthracycline exposure, female sex, chest radiation, hypertension or diabetes).

## Bone Health

Loss of bone density after HCT is well described and typically occurs in the first 6-12 months post-transplant<sup>12</sup>. Beyond one year post-HCT, recovery of bone mineral density (BMD) to a variable degree may occur if patients do not experience additional risk factors for bone loss. Additional risk factors for osteoporosis include prolonged exposure to corticosteroids and calcineurin inhibitors (i.e. ongoing treatment of cGVHD), major weight loss, malnutrition, older age at HCT and female gender<sup>14,15</sup>. In a recent study, the prevalence of osteoporosis and osteopenia in patients experiencing moderate-severe cGVHD was 17% and 60%, respectively<sup>16</sup>. The estimates in the literature of the incidence of osteoporosis and osteopenia after HCT in those without cGVHD vary widely; however, both auto and allo-HCT recipients, particularly females and older males, have a marked increase in risk of fracture compared to that of the general population<sup>15</sup>. At least two studies have revealed a higher risk of fracture after auto-HCT versus allo-HCT<sup>15,17</sup>. Finally, degree of bone loss after HCT does not seem to directly correlate with risk of fracture,<sup>18</sup> highlighting the potential importance of using clinical assessment tools for fracture risk (such as FRAX) which take into account BMD and clinical factors. The Alberta Toward Optimized Practice (TOP) guidelines for osteoporosis provide recommendations for therapy and follow-up DXA based on the 10 year probability of fracture as calculated by FRAX. While FRAX takes into account corticosteroid exposure in general, it may underestimate the fracture risk for those who are receiving long courses of moderate to high doses of corticosteroids (i.e. cGVHD) and those who have more severe bone loss at the spine versus the hip (as it does not take into account BMD at the lumbar spine). There are, however, no universally agreed upon adjustments to fracture risk estimates for these variables. The recommendations for those with cGVHD below are generally in agreement with the 2017 American College of Rheumatology guidelines for the management of corticosteroid-induced osteoporosis<sup>19</sup>.

## Recommendations:

- All patients should be counselled regarding lifestyle modifications for bone health including calcium intake 1200 mg/day from all sources, vitamin D 1000 IU/day, smoking cessation, limiting alcohol use & regular weight-bearing exercise.
- At 1 year post-HCT or at the onset of cGVHD requiring systemic therapy (whichever occurs earlier), patients should have an assessment of BMD by dual X-ray absorptiometry (DXA) and subsequently a 10 year probability of fracture calculated with the FRAX clinical assessment tool (<https://www.sheffield.ac.uk/FRAX/>) (except patients with multiple myeloma who are already being treated with bisphosphonates).
- For those who are no longer on immunosuppression at 1 year post-HCT, the Alberta Toward Optimized Practice (TOP) guidelines for osteoporosis ([http://www.topalbertadoctors.org/cpgs/?sid=18&cpg\\_cats=81](http://www.topalbertadoctors.org/cpgs/?sid=18&cpg_cats=81)) should be followed.
- For those who remain on immunosuppression at 1 year post-HCT, therapy should be offered to:
  - those with established osteoporosis (BMD T-score  $\leq -2.5$ ) or history of fragility fracture, and
  - those with a moderate to high probability of fracture by FRAX as outlined by the Alberta TOP guidelines (>10% probability).
- In those who remain on chronic glucocorticoid therapy (prednisone equivalent dose >5mg/day) beyond 1 year post-HCT and who do not initiate therapy, annual DXA measurement should be considered.
- For those who initiate therapy, repeat DXA should be obtained at 3-5 years on therapy.
- First line therapy is typically with oral bisphosphonates. If identified, treatment of hypogonadism also could be considered in men and pre-menopausal women (after evaluation and discussion of risks/benefits of hormone replacement). Referral to an endocrinologist may be considered for alternate or second-line therapy options.
- The need for ongoing bisphosphonate therapy should be reassessed at the end of immunosuppressive therapy and/or at 5 years on therapy.
- In those with osteoporosis, a workup for secondary or contributing causes should be undertaken (for example, hypogonadism, hyperthyroidism & hyperparathyroidism).

## Subsequent Malignancy Screening

Both auto- and allo-HCT recipients are at increased risk of secondary solid tumours. The risk of secondary solid tumours increases over time post-HCT with a cumulative incidence of about 15% at 25 years post-HCT—a two to three fold increased risk versus age and sex matched controls<sup>20,21</sup>. HCT recipients are particularly at risk (standardized incidence ratio >1) of the following malignancies: all skin including melanoma, thyroid, oropharyngeal, esophageal, liver, bone, central nervous system

and connective tissue<sup>22</sup>. Chronic GVHD and duration of immunosuppression > 2 years are major risk factors for skin, oropharyngeal, cervical and esophageal cancers, while conditioning with total body irradiation (TBI), particularly myeloablative TBI, is a risk factor for skin, thyroid, liver and breast cancers<sup>22</sup>. There is little data to guide second malignancy screening practices in HCT survivors: expert recommendations generally suggest screening similar to the general population with some additions as will be described below<sup>22,23</sup>. Those who have received radiation within specific fields, such as chest or cranial, or those with a cancer predisposition syndrome (ex. Fanconi anemia) may require individualized enhanced screening measures.

Survivors of autologous-HCT are at risk of therapy-related myeloid neoplasms (t-MN) (predominantly myelodysplastic syndrome and AML). The cumulative incidence of t-MN is about 7% at 15 years post auto-HCT<sup>20</sup>. Major risk factors for t-MN are alkylator therapy (5-7 year latency from HCT) and topoisomerase II inhibitor therapy (6 month-5 year latency from HCT)<sup>20</sup>. Unfortunately, outcomes of those who develop t-MN is poor<sup>24</sup>.

### **Recommendations:**

- All patients should be counselled regarding smoking cessation and cutaneous solar protection.
- All patients should have a yearly history and physical exam that includes oral cavity, thyroid, and a complete skin exam.
- All patients should visit their dentist for oral/dental examination and cleaning; at least yearly for most patients and every 6 months for those with oral chronic GVHD.
- Screening for breast (see modifications below for women who have received TBI), cervical, colorectal (see modifications below for those who have received high dose TBI or localized radiation) and prostate cancer should follow established Albertan/Canadian guidelines ([www.screeningforlife.ca](http://www.screeningforlife.ca) & for prostate cancer [www.canadiantaskforce.ca](http://www.canadiantaskforce.ca)).
- For women who have received any dose of TBI: screening mammography starting at age 25 or 8 years after radiation exposure, whichever occurs later but no later than age 40.
- For patients who received high dose TBI ( $\geq 6$  Gy) or abdominal, pelvic, lumbar, sacral, or whole spine irradiation, colorectal cancer screening with colonoscopy should begin at 5 years post-transplant or age 30, whichever occurs later.
- HCT recipients who have received radiation within a particular field, for example chest and those with familial cancer syndromes will require an individualized plan for malignancy screening.
- There are no proven screening measures for t-MN after auto-HCT; yearly CBC with early workup of cytopenias for up to 10 years post-transplant may be considered.

## **Renal Disease**

The definition of chronic kidney disease (CKD) encompasses both decreased kidney function (glomerular filtration rate (GFR) <60 mL/minute) and kidney damage other than decreased GFR (ex.

albuminuria) with duration  $\geq 3$  months<sup>25</sup>. Estimates of the cumulative incidence of CKD in the months and years after HCT vary widely at 7 to 48%<sup>26</sup>. In one report, 4% of long-term survivors of allo-HCT (19% of the long-term survivors who had developed CKD) developed end-stage renal disease<sup>27</sup>. In both the general population<sup>28</sup> and the post-HCT population<sup>29</sup>, CKD is independently associated with increased mortality, particularly cardiovascular mortality. Risk factors for CKD after HCT include history of acute kidney injury (AKI), occurrence of acute & chronic GVHD, age  $\geq 45$  at HCT, pre-HCT baseline GFR  $<90$  mL/minute, hypertension and exposure to high dose total body irradiation<sup>26</sup>. While CKD may arise from a number of clinicopathologic entities after HCT; the best described being thrombotic microangiopathy, viral nephropathies and nephrotic syndrome; it is most commonly idiopathic or as a result of incomplete recovery from acute kidney injury early post-HCT<sup>30</sup>. These pathologies have been recently reviewed and are summarized from these sources in Table 1<sup>26,30,31</sup>.

**Table 1.** Etiologies of chronic kidney disease after hematopoietic cell transplant

| Clinicopathologic Entity                                   | Incidence                       | Risk Factors                                                        | Clinical Characteristics                                                                                             |
|------------------------------------------------------------|---------------------------------|---------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------|
| Idiopathic                                                 | Most patients with CKD post HCT | -AKI after HCT<br>-aGVHD & cGVHD<br>-High dose TBI<br>-Hypertension | -None specific                                                                                                       |
| Thrombotic microangiopathy                                 | 2-21%                           | -TBI<br>-Calcineurin inhibitor use<br>-aGVHD & cGVHD                | -Microangiopathic hemolysis<br>-Acute kidney injury, often with incomplete recovery of renal function leading to CKD |
| Nephrotic Syndrome (66% membranous and 19% minimal change) | 1%                              | -cGVHD                                                              | -Associated with cGVHD<br>-Proteinuria $>3.5$ g/24 hours<br>-Hypoalbuminemia<br>-Edema<br>-Hyperlipidemia            |
| BK Nephropathy                                             | Rare                            | -Immunosuppression                                                  | -BK viremia                                                                                                          |

Proteinuria, even microalbuminuria, particularly after allo-HCT, is increasingly recognized as a prognostic marker. Specifically: 1) those with albuminuria at day 100 post-HCT have a significantly higher risk of non-relapse mortality by one year post-HCT (predominantly due to GVHD and infection), and 2) those with albuminuria at any point between day 100 and one year post-HCT have an increased risk of developing CKD<sup>24,30</sup>.

**Recommendations:**

- All HCT recipients should have at least yearly monitoring of creatinine.
- Allo-HCT recipients should a spot urine albumin/creatinine ratio yearly.



- All patients should have at least yearly blood pressure evaluation and treatment per Canadian (CHEP) guidelines (essentially target BP <140/90 for most patients).
- Basic medical management of CKD includes initiation of an ACE inhibitor or ARB for proteinuria, tight glycemic control for diabetics, aggressive management of cardiovascular risk factors and avoidance of nephrotoxins.
- Referral to a nephrologist should be considered when estimated GFR is <30 mL/min, for management of CKD with proteinuria or for workup of CKD of unknown etiology.

## Pulmonary Disease

HCT recipients are at risk of both late infectious and non-infectious pulmonary diseases. Late infectious pulmonary complications include recurrent sinopulmonary infections, Pneumocystis and fungal infections. The approach to late pulmonary infections is addressed elsewhere in these guidelines (bacterial/pneumocystis prophylaxis, fungal prophylaxis and vaccination chapters). Late onset non-infectious pulmonary complications (LONIPCs) mainly affect allo-HCT recipients. LONIPCs are very rare after autologous-HCT-the vast majority of non-infectious pulmonary complications after auto-HCT occur in the peri-engraftment period<sup>33</sup>. The most common LONIPCs are summarized in Table 2 and include bronchiolitis obliterans syndrome (BOS) (a manifestation of cGVHD) and interstitial lung disease (the best defined being organizing pneumonia (OP), but diffuse alveolar damage, non-specific interstitial pneumonia and lymphoid interstitial pneumonia have also been described)<sup>34-36</sup>. Idiopathic pneumonia syndrome, diffuse alveolar haemorrhage and pulmonary veno-occlusive disease most often occur early (day 0-30) post-HCT, but rarely occur as a late toxicity<sup>37</sup>. In a recent prospective study, all LONIPCs were associated with cGVHD and were found to occur predominantly in the first 2 years after allo-HCT with a cumulative incidence of 20% at 3 years<sup>38</sup>. The LONIPCs consisted of BOS (40%), interstitial lung disease (22%), venous thromboembolic disease (16%) and restrictive lung disease with no interstitial lung or pleural disease (including cGVHD with cutaneous sclerosis) (15%). Those who experienced a LONIPC were at increased risk of death (HR 2.2); the main causes of death included relapse followed by respiratory causes and GVHD. Importantly, lower respiratory tract infection in the first 100 days after HCT, pre-HCT chest irradiation and low FEF25-75% at day 100 were risk factors for the development of a LONIPC. Evaluation of risk factors for BOS after allo-HCT have variably found older age, sex-mismatched HCT, history of aGVHD, busulfan-based conditioning, unrelated donor and peripheral blood stem cell graft to be associated with the development of BOS, while T-cell depletion is protective<sup>36</sup>. Finally, it should be noted that several chemotherapeutic agents (such as BCNU, bleomycin, busulfan and methotrexate) may contribute to or cause pulmonary toxicity<sup>22</sup>.

**Table 2.** Late-onset non-infectious pulmonary complications after allogeneic-HCT

| Entity                   | Time of Onset             | CT Imaging Features                                                                 | PFT Features                                              | Clinical Features                                                                                                      | Therapy                                            |
|--------------------------|---------------------------|-------------------------------------------------------------------------------------|-----------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------|
| Bronchiolitis Obliterans | 3 months-2 years post-HCT | -Air trapping<br>-Bronchial thickening<br>-Bronchiectasis<br>-Centrilobular nodules | -Obstructive<br>-Diagnosis per NIH criteria <sup>37</sup> | -Extra-pulmonary cGVHD usually present<br>-Asymptomatic early<br>-Cough, dyspnea, wheezing                             | -Systemic and topical therapy per cGVHD guidelines |
| Organizing Pneumonia     | Median 3 months post-HCT  | -Diffuse consolidation or ground glass opacity                                      | -Restrictive > Normal > Obstructive > Mixed               | -“Non-resolving infectious pneumonia”<br>-Often in the setting of taper of immunosuppression for acute or chronic GVHD | -1 mg/kg prednisone with slow taper                |

Because the onset of LONIPCs is often insidious, particularly for BOS, with the potential for significant loss of lung function before symptoms develop, post-HCT screening pulmonary function tests (PFTs) are essential. The approach to abnormal PFTs begins with history and physical exam to elucidate recent or current infections. The investigation is guided by history, physical exam, and pattern of abnormal PFT, but generally begins with chest x-ray (CXR) and non-invasive infectious workup such as nasopharyngeal (NP) swab for respiratory viruses and sputum culture. If no clear etiology is found or empiric therapy fails, the next steps are guided by acuity of the presentation but generally include obtaining a CT chest and referral to the BMT pulmonary clinic with consideration of bronchoscopy with bronchoalveolar lavage +/- lung biopsy.

### Recommendations:

- The approach to prevention of late pulmonary infections are outlined in the bacterial/pneumocystis prophylaxis, fungal prophylaxis and vaccination chapters of these guidelines.
- For autologous-HCT recipients: PFT at 6 weeks post-HCT for those who received potentially pulmonary toxic conditioning. For all, at least yearly history and physical exam for signs and symptoms of pulmonary disease is recommended.
- For allogeneic-HCT recipients: routine PFTs for all patients every 3 months for the first year post-HCT followed by yearly PFTs until 5 years post-HCT. For those with active cGVHD beyond 1 year post-HCT, continued every 3 month PFTs should be strongly considered. History and physical exam should accompany PFTs.
- Abnormal PFTs or new respiratory symptoms should be worked up promptly with CXR +/- NP swab and sputum culture. For clinical presentations that are not consistent with upper respiratory tract infection or community-acquired pneumonia, CT chest and referral to the BMT pulmonary clinic are suggested.

## Endocrine Disease

### Thyroid Function:

Hypothyroidism is relatively common after HCT, occurring in up to 30% of long-term survivors<sup>40</sup>. Risk factors include younger age at HCT, radiation (neck, mediastinal or total body) and exposure to busulfan and cyclophosphamide<sup>20,40</sup>. Symptoms of hypothyroidism are non-specific and include fatigue, cold intolerance, weight gain, constipation and dry skin. Hypothyroidism is also a secondary cause/contributor to hyperlipidemia.

### Hyperlipidemia and Diabetes:

As discussed in the cardiovascular disease section above, both autologous and allogeneic HCT recipients acquire cardiovascular risk factors such as hyperlipidemia and diabetes faster and more frequently than the general population. While GVHD & immunosuppressive therapy are well known risk factors for hyperglycemia and hyperlipidemia<sup>41</sup>, it should be noted that HCT survivors at least five years post-transplant without active GVHD and not on immunosuppressive therapy had double the risk of developing metabolic syndrome versus the age-matched population and this risk was independent of allo- versus auto-HCT<sup>10</sup>. As discussed in the cardiovascular disease section above, management of diabetes should follow standard practice for that of the general population and management of hyperlipidemia should be guided by global cardiovascular risk.

### Adrenal Insufficiency:

A single centre study found that the cumulative incidence of adrenal insufficiency after allo-HCT was 13%, while it was 1% after auto-HCT<sup>42</sup>. Those who are treated with long courses of corticosteroids for GVHD are particularly at risk. An ACTH stimulation test may be used to confirm the diagnosis of adrenal insufficiency. Management of adrenal insufficiency includes initiation of physiologic corticosteroid dosing followed by a very slow taper. Weak data and expert opinion suggest that an alternate day tapering regimen may reduce the risk of adrenal insufficiency<sup>40</sup>. Additionally, a medical alert bracelet or information card should be worn or carried and patients should be alerted to seek immediate medical attention if they develop signs or symptoms of adrenal insufficiency (ex. nausea/vomiting/abdominal pain/postural hypotension).

### Gonadal Dysfunction and Fertility:

Gonadal dysfunction and infertility are reviewed elsewhere<sup>1,2</sup>.

### Recommendations:

- Yearly thyroid examination and TSH measurement for all HCT recipients.
- A slow terminal taper of corticosteroids is required for those receiving prolonged courses (>3 weeks) for treatment of GVHD.
- A high index of clinical suspicion for adrenal insufficiency should be maintained when tapering patients from prolonged courses of corticosteroids.

- Management of diabetes and hyperlipidemia within the context of global cardiovascular risk as outlined in the “Cardiovascular disease” section.
- Workup and management of gonadal dysfunction and infertility as published<sup>1,2</sup>.

## Management of Chronic Pain

Survivors of HCT may experience chronic pain related to a number of treatment-related complications such as GVHD, peripheral neuropathy and non-specific cramping/muscle spasm among others. Management of chronic pain requires a multi-disciplinary approach that includes HCT providers, palliative care providers, pharmacists, and psychosocial providers.

### Recommendations:

- HCT recipients with chronic pain should be managed within a multi-disciplinary team that includes HCT providers, pharmacists and the psychosocial team within the Alberta Blood and Marrow Transplant Program with low threshold for referral to the palliative care team at the Tom Baker Cancer Centre.

## Transfusion

In general, most HCT recipients do not require transfusion in the post-engraftment period. However, if transfusion is required, thresholds for transfusion of red cells and platelets should be individualized based on the specific clinical circumstances (ex. symptoms, co-morbidities, underlying disease etc.). Product attributes for transfusion as recommended below (i.e. irradiation and CMV status) are in agreement with established Canadian guidelines<sup>43,44</sup>. Appropriate blood-group products for ABO-incompatible allo-HCT are reviewed in the “ABO Incompatibility” chapter of these guidelines.

### Recommendations:

- Red cell and platelet transfusion thresholds should be individualized based on clinical circumstances.
- The appropriate blood-group products for transfusion after ABO-incompatible allo-HCT per the “ABO incompatibility” chapter.
- For allo-HCT recipients: irradiated blood products should be used from start of conditioning until the later of: 1 year post-HCT, end/“burn out” of chronic GVHD or discontinuation of immunosuppression.
- For auto-HCT recipients: irradiated blood products for 7 days prior to stem cell collection and from start of conditioning until 3 months post-HCT, or 6 months post-HCT if TBI was part of conditioning.
- All auto- and allo-HCT recipients should receive standard leukoreduced (“CMV safe”) blood products.

## Mental Health

A significant minority of HCT survivors will experience persistent anxiety, depression and/or post-traumatic stress disorder in the years after transplant<sup>45</sup>.

### **Recommendations:**

- Long-term follow up visits should include screening for mental health concerns. If concerns are identified, patients should be referred to mental health providers and/or community resources as appropriate.

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# Nutrition Support in Hematopoietic Cell Transplant

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## Summary

- Hematopoietic cell transplant patients are at risk of malnutrition before, during, and after transplant.
- Malnutrition risk may be mitigated through initiation of nutrition support, preferably through the use of enteral nutrition (tube feeding).
- Nutrition support should be initiated for hematopoietic cell transplant patients if: inadequate intake (less than 60% of requirements) for 5 days or likely to remain inadequate for >5 days, or weight loss in a euvolemic patient of >5% in one month or >2% in one month if BMI <20kg/m<sup>2</sup>, except for those patients meeting these criteria during the expected period of mucositis and for 7-10 days after resolution of mucositis.

## Background

Patients undergoing hematopoietic cell transplant (HCT) are at risk of malnutrition due to the gastrointestinal (GI) side effects of the treatment which impede oral intake, including, but not limited to, mucositis, anorexia, dysgeusia, nausea, vomiting, and diarrhea<sup>1,2</sup>. As a result, patients often are unable to consume enough food to adequately meet their nutritional needs during transplant, with some consuming less than 60% of their estimated requirements for one to two weeks<sup>3-5</sup>, which is the defined criteria in the literature for inadequate oral intake among cancer patients. HCT patients are often strongly advised to gain weight prior to their transplant as pre-transplant weight loss is related to increased relapse risk and shorter disease-free survival post-transplant and non-relapse mortality<sup>6,45</sup>. Body mass index (BMI) less than 18.5kg/m<sup>2</sup>, underweight<sup>7</sup> and weight loss were historically used to identify malnutrition, but are no longer considered effective markers on their own<sup>5</sup>. Assessment and diagnosis of malnutrition is not always straightforward and requires dietitian involvement early and throughout the transplant process<sup>43-45</sup>. Malnutrition assessments that account for sarcopenia—defined as low muscle mass and poor muscle strength—categorize patients into mild, moderate and severe malnutrition<sup>45,50</sup>. Individuals who have more than two of the following are considered severely malnourished: obvious muscle wasting, loss of subcutaneous fat, decreased functional status, weight loss of > 2% in 1 week, 5% in 1 month, or 7.5% in 3 months<sup>8</sup> or consume less than 60% of recommended nutritional intake for 2 weeks or more<sup>3-5</sup>. Muscle loss comprises a large proportion of short-term weight loss, wherein immobile or bedridden hospitalized patients can lose up to 0.5% of total body muscle mass per day<sup>9</sup>, with considerable muscle loss occurring during a prolonged hospital stay<sup>10</sup>. Malnutrition is associated with lower health-related quality of life<sup>11,12</sup>, impaired functional ability<sup>12,13</sup>, higher rates of infection<sup>14</sup>, impaired wound healing<sup>15</sup>, longer hospital length of stay<sup>16,17</sup>, increased health care costs<sup>16</sup>, and higher mortality<sup>18,19</sup>.

Both enteral and parenteral nutrition can be utilized to provide nutrition support to HCT patients who are at risk of malnutrition and whose oral intake is impeded by the gastrointestinal side effects of treatment. Standard of clinical practice regarding use of enteral and/or parenteral nutrition varies significantly among institutions due to differences in country of practice and knowledge, availability of nutrition experts, and clinician perception of invasiveness and poor tolerance though there is consensus that nutrition support during HCT is required<sup>47,48</sup>.

Enteral nutrition can be delivered via nasogastric or nasoduodenal/nasojejunal tubes (short term, ideally less than 4 weeks) or via endoscopically, surgically or radiologically placed percutaneous gastric/jejunal tubes (long term)<sup>24</sup>. Naso-enteric feeding tubes can contribute to both physical and psychological discomfort for the patient, or possibly more serious complications of nasopharyngeal lesions, sinusitis or reflux esophagitis, tube dislocation and/or movement within the nares and throat, and the tube presence being a reminder of illness<sup>25</sup>. Clinicians may be concerned about risk of bleeding during naso-enteric feeding tube insertion. Prophylactic platelet infusion (i.e. transfuse if platelet count  $<30 \times 10^9$ ) pre-insertion may help to mitigate this risk, with one study demonstrating no association between thrombocytopenia and increased risk of bleeding after feeding tube insertion among critically ill oncology patients<sup>26</sup>.

Nasogastric tubes may be advantageous over nasoduodenal/nasojejunal tubes due to: (1) decreased cost, as they can be inserted at the bedside by a nurse, rather than by a physician in fluoroscopy, (2) ability to infuse either intermittent (bolus) or continuous tube feeds and (3) more physiological for motility and hormones<sup>27</sup> and medication administration<sup>28</sup>. Nasoduodenal/nasojejunal feeds require a pump for continuous infusion and may be preferred over nasogastric tubes in settings of intractable vomiting and to reduce risk of aspiration, although this risk is not entirely eliminated<sup>27</sup>. However, a continuous tube feed may impact patient quality of life as it can limit activity when a patient is connected to the pump all day<sup>28</sup>. Bolus feeding (gastric feeds only) may contribute to increased risk of aspiration and to increased incidence of diarrhea, depending on rate of formula infusion<sup>28</sup>.

Parenteral nutrition may be indicated when a patient has a non-functional gastrointestinal tract (i.e. bowel obstruction or paralytic ileus) or in the presence of high gastrointestinal losses (diarrhea, vomiting, short bowel syndrome)<sup>20</sup>. Total parenteral nutrition (TPN) requires central venous access such as a percutaneous non-tunneled central catheter, a tunneled cuffed catheter, a peripherally inserted central catheter (PICC) or an implanted port prior to initiation<sup>21</sup>. Peripheral parenteral nutrition (PPN) may be used when enteral nutrition is contraindicated and the patient has a peripheral access but no central venous access, usually as a short-term bridge to total parenteral nutrition or enteral nutrition. Complications of parenteral nutrition may include refeeding syndrome, infection, thromboembolic events, hyperglycemia, cholestasis, hypertriglyceridemia, metabolic bone disease<sup>21,22,46</sup> and acalculous cholecystitis<sup>23</sup>. Although parenteral nutrition may be delivered more consistently than enteral nutrition due to interruptions related to gastrointestinal intolerance of the enteral formula, enteral nutrition is advantageous for a number of reasons including maintenance of the gut function, integrity, and barrier which protects against bacterial translocation, fewer infections, decreased length of stay and decreased financial cost<sup>27,29,30,46</sup>. Enteral nutrition may be discontinued

for a number of reasons including nausea, vomiting, diarrhea, psychological intolerance, tube blockage or displacement<sup>32</sup>.

Both the American Society of Parenteral and Enteral Nutrition (A.S.P.E.N.) and the European Society for Clinical Nutrition and Metabolism (E.S.P.E.N.) recommend enteral nutrition over parenteral nutrition for nutrition support among HCT patients<sup>33,34</sup>. A.S.P.E.N. recommends enteral nutrition in HCT patients with a functioning gastrointestinal tract, and nutrition support in malnourished patients anticipated to be unable to ingest and/or absorb adequate nutrients for a prolonged period of time<sup>33</sup>. E.S.P.E.N. recommends initiation of enteral nutrition except in situations of severe mucositis, intractable vomiting, ileus, severe malabsorption, protracted diarrhea or symptomatic gastrointestinal graft versus host disease (GVHD), when parenteral nutrition could be provided instead<sup>34</sup>.

Tolerance to enteral nutrition in allogeneic HCT patients was demonstrated in a large RCT by Andersen *et al.* who found that over a median of 8 days, patients received a median of 73% of goal calories and protein through enteral nutrition. The authors suggested that enteral nutrition use over parenteral use could decrease costs and minimize parenteral nutrition-related complications<sup>32</sup>. A variety of retrospective and prospective studies found that enteral nutrition in HCT was associated with reduced non-relapse mortality, lower incidence of acute GVHD of the GI tract, decreased risk of infection, improved neutrophil and platelet engraftment, and improved overall survival<sup>32,35-37</sup>. In contrast, parenteral nutrition was associated with increased early mortality and delayed platelet engraftment<sup>37</sup>. Several studies and systematic reviews concluded that enteral nutrition, in conjunction with medications to improve tolerance (antiemetics, antidiarrheals, analgesia), may be associated with a decreased risk of infectious complications and prevention of GVHD compared to parenteral nutrition<sup>36-41</sup>.

Studies and reviews that have emerged since 2022 examine the benefits of enteral nutrition over parenteral nutrition on the modulation of the gut microbiota and suggest that maintaining gut stimulation has a role in retention of muscle mass, lower rates of acute GVHD, and lower risk of bloodstream infections. As chemotherapy, radiation therapy, and antibiotic treatments disrupt the mucosal barrier and cause microbiota dysbiosis (defined as decrease microbiota diversity, loss of commensal bacteria, expansion of pathogenic bacteria), nutrition being a main driver of species diversity and function in the gut microbiome can be a primary defense<sup>44,46,48</sup>. Oral or enteral nutrition stimulates metabolite production such as short-chain fatty acids that have a role in anabolism and muscle energy expenditure, and may prevent intestinal domination by a single bacterium (a single bacterial taxon comprising of > 30% of the gut microbiome)<sup>44,48</sup>. Stein-Thoeringer *et al.* found *Enterococcus spp.* domination in up to 65% of patients after an allogeneic transplant, and this was associated with reduced overall survival and increased risk of acute GVHD<sup>49,49</sup>. Conversely, parenteral nutrition cannot counter the gut atrophy that occurs inevitably in intestinal fasting if it is the sole source of nutrition<sup>46</sup>. Literature suggests that the longer patients fast after completing

conditioning chemotherapy, the higher the risk of GVHD, as well as a reduction in gut microbiota diversity<sup>46,48</sup>.

## 2021-2024 Data on Use of Enteral Nutrition at the Foothills Medical Centre

The nutrition support algorithm (Figure 1) was implemented in 2021 and from August that year to September 2024, data was collected on the use of enteral and parenteral nutrition in the HCT population at the Foothills Medical Centre. During this period, approximately 22% of the 612 patients admitted for HCT were screened out and offered early enteral nutrition (36% of all allogeneic HCTs and 8% of all autologous HCTs). Of those offered enteral nutrition, 56% of allogeneic HCT patients (60 patients) and 50% of autologous HCT patients (13 patients) accepted and started enteral nutrition during their transplant admission. Several patients who started enteral nutrition on a readmission are not reflected in this data, as the focus of this standard practice was on initiation of early enteral nutrition during the transplant admission.

Reasons patients do not start enteral nutrition despite being screened for it include: patient perception of invasiveness and fear of discomfort, fear of worsening GI symptoms like nausea or diarrhea, fear of epistaxis, and underestimating the barrier that mucositis will pose on oral intake as the admission progresses. Despite these barriers, patients have been increasingly more accepting of supplemental enteral nutrition as part of their HCT treatment process due to earlier introduction to the option of enteral nutrition from the physician in pre-transplant assessments and from the dietitians and nurses throughout the patient's admission as they reinforce the benefits of EN or answer questions patients may have.

Collected data shows the tolerance of enteral nutrition is high, with allogeneic HCTs tolerating 100% median or 83% average, and autologous HCT tolerating 80% median or 69% average of their total prescribed calories and protein. The currently preferred enteral feeding route for allogeneic HCTs is with a nasoduodenal or nasojejunal feeding tube, a calorically dense (2.0kcal/mL) formula, and feeding during the daytime only, all with the goal of lowering aspiration risk. The most important consideration to lower aspiration risk regardless of enteral tube type is maintaining a head of bed of at least 30 degrees. Use of hypercaloric formulas may improve patients' tolerance by reducing volume administered<sup>46</sup>.

Instances of complications with enteral feeding were tracked in allogeneic HCT patients: 18% (11 patients) had nausea and vomiting induced tube dislodgement, 0% had tube related aspiration events, 3% (2 patients) had tube clogging, and 2% (1 patient) had worsened epistaxis. After enteral nutrition was attempted, 15% of allogeneic HCT (9 patients) required a transition to parenteral nutrition without supplemental enteral nutrition, either due to loss of feeding tube and inability or refusal to reinsert, or due to gut-related complications that required gut rest. Patients are often transitioned back to oral nutrition prior to discharge and very rarely are discharged home on the

Home Enteral Nutrition Program. More commonly, patients may be readmitted with failure to thrive and prolonged poor intake post-transplant requiring supplemental enteral nutrition and discharge with the Home Enteral Nutrition Program.

Allogeneic HCT patients who received early enteral nutrition had a median weight loss of 6.1% (4kg) compared to 7.8% (6.2kg) for screened patients who did not start enteral nutrition. This difference was statistically significant ( $p=0.00988$ ) however this median difference of 2kg may not be clinically significant. Fluid retention, edema, pre-transplant weight or BMI, and length of stay are all confounding variables. Though data was collected to compare length of stay, it was observed that patients with more complications resulting in longer lengths of stay may eventually require enteral nutrition despite initially refusing. Therefore, the longer average length of stay for HCT patients who received enteral nutrition is likely affected by other confounding factors.

Though data on incidences of GVH, infection, and 100-day survival were not explicitly collected for patients screened for enteral nutrition, literature explores and proposes mechanistic explanations for the benefits of enteral nutrition on these outcome measures.

## Standard Practice in Alberta for Adults

The standard of practice in Alberta currently is for practitioners, primarily Registered Dietitians who are consulted for every HCT patient admitted, to use the Nutrition Support Algorithm (Figure 1) to recommend initiation of enteral nutrition when appropriate. As literature varies in recommendation and use of feeding route, duration, and formula type, the Registered Dietitian should make a clinical decision alongside the medical team with close monitoring for tolerance and adjusting as indicated. If enteral nutrition is tried and unsuccessful, parenteral nutrition should be considered after an assessment of benefits against risk with the medical team. Regardless of use of enteral or parenteral nutrition, maintenance of oral nutrition even in small amounts should be recommended as tolerated to support the gut barrier and prevent gut atrophy among various other benefits.



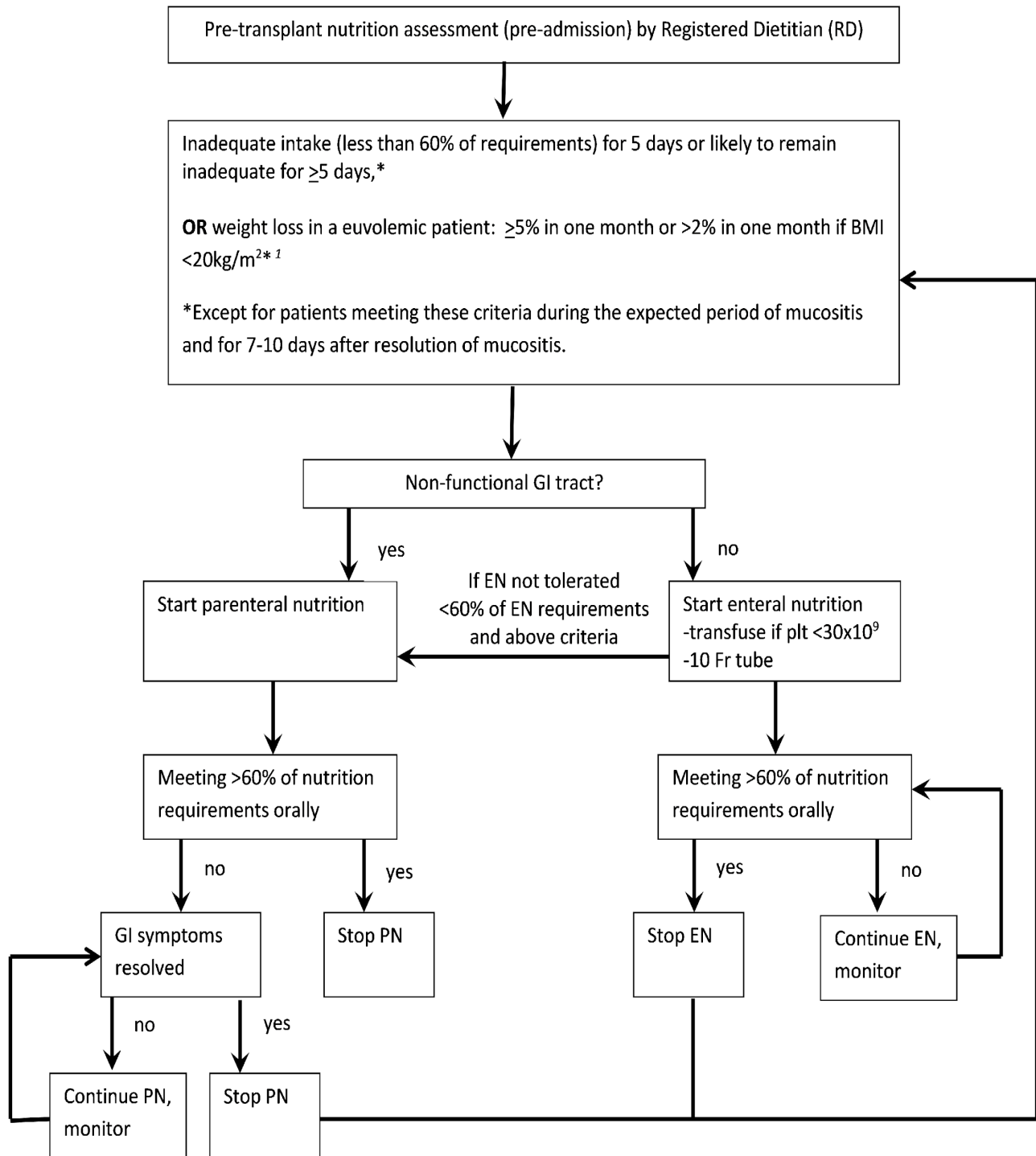


**Figure 1.** Nutrition support algorithm for hematopoietic cell transplant (HCT) patients in Calgary, AB. Adapted from *Beckerson et al.*<sup>35</sup> and *Andersen et al.*<sup>42</sup>  
plt = platelets

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# Microbially Contaminated or Non-Conforming Cellular Therapy Products

Presented by: Andrew Daly

## Summary

**Upon notification of a potentially or confirmed microbially-contaminated or otherwise non-conforming cellular therapy product the *recipient's transplant physician* will:**

- Notify the recipient of the non-conformance and ensure the recipient receives follow up care/treatment to reduce risks associated with the non-conforming product. This will be documented in the recipient's medical record.
- Notify the donor transplant physician.
- Notify the Program Quality Manager.
- Notify Cellular Therapy Lab (if not yet notified). CTL will initiate non-conforming product investigation according to applicable SOP's.
- In the case that the donor is an unrelated donor the physician will contact the Canadian Blood Services Stem Cell Registry Case Manager on call at (613) 260-6800. Registry personnel must notify the transplant centre of the non-conformance.

**Upon notification of a potentially or confirmed microbially-contaminated or otherwise non-conforming cellular therapy product the *donor's transplant physician* will:**

- Notify the donor of the positive microbial result. Ensure the donor receives follow up care if applicable. This discussion shall be documented in the donor's medical record and the donor's regular physician should be advised.

## Background

Despite rigorous quality control and adherence to good manufacturing practices, cellular therapy products (CTPs) may occasionally fail to meet the high standards set for cellular therapy. These products may still be suitable for use, and in most cases are the most appropriate products for the patient. The purpose of these guidelines is to ensure notification and appropriate follow-up of the donor and recipient of these products, notification of the donor and recipient physicians and to ensure notification of regulatory agencies. These guidelines are also intended to standardize the management of patients receiving non-conforming products, in accordance with the foundation for accreditation of cellular therapy (FACT) standards.

**Non-conforming products include but are not limited to products with the following types of deficiencies:**

1. Those with potential or proven microbial contamination
  - Positive microbial testing
  - Cracked or damaged storage bag
  - Improper transport or storage
  - CTP variance at time of infusion
2. Those with increased potential for infusion-related adverse events
  - Failed release criteria (clots, clumps, abnormal colour)
  - Deficiencies or errors in processing
3. Those that increase risk of engraftment failure
  - Low cell dose
  - Improper storage or handling

**The identification of any of the above situations will require the following protocol(s) to be followed:**

1. *For cellular therapy products with potential or proven microbial contamination:*
  - a. A non-conforming product investigation will be initiated by the Cellular Therapy Laboratory.
  - b. The recipient and donor transplant physicians shall be informed of the positive culture result or a potentially contaminated product, and this discussion shall be documented in the medical record.
  - c. In the case of allogeneic cellular therapy products with positive microbial cultures, the donor physician shall be advised of the positive result in order that he or she can arrange appropriate follow-up of the donor.
  - d. All products will have aerobic, anaerobic and fungal cultures drawn and kept in culture for 5-14 days to allow isolation of fastidious organisms. This should be indicated on the requisition.
  - e. Patients should receive a dose of Vancomycin before infusion of the product, with further doses based upon results of repeat cultures, likelihood of falsely positive cultures and the patient's clinical status.
  - f. Daily blood cultures will be drawn from the patient for a minimum of 3 days after infusion of the cellular therapy product.
  - g. Fevers should be managed according to appropriate guidelines, with repeat blood cultures drawn according to guidelines for management of febrile neutropenia or based on advice of the infectious disease consultant.

- h. The potential for infusion of a microbially- or endotoxin-contaminated cellular therapy product should be considered in patients with flushing, high fever (> 2 degree C rise from baseline), rigors, confusion or circulatory collapse shortly after infusion and appropriate management instituted. Appropriate antibiotic treatment should be initiated and an infectious disease consult called as needed.
  - i. Canadian Blood Services Stem Cell Registry must be informed immediately of positive microbial test results on products collected for distribution outside the ABMTP. They can be reached by calling the Registry On Call Case Manager at (613) 260-6800.
- 2. *For cellular therapy products with increased potential for infusion-related adverse events:*
  - a. A non-conforming product investigation should be initiated by CTL for products that fail to meet release criteria or when a deficiency or error occurs during processing.
  - b. The patient should be advised of the product variance and of any action to mitigate risk (such as increased premedication or monitoring post-infusion). This should be documented in the patient's medical record.
- 3. *For cellular therapy products with higher risk of engraftment failure:*
  - a. Inform the Cellular Therapy Laboratory and Workup Nurse of the deficiency.
  - b. Inform the patient and the transplant physician of the risk of engraftment failure and any action that may be taken to decrease the risk (such as early infusion of a new cellular therapy product or enhanced monitoring for engraftment failure). Document this discussion in the patient's medical record.



# Appendices

| TEST                                                                                                                                                                                                                                                                                                       | 2 wk                                                                                                                                                                                   | 4 wk | 6 wk | 8 wk               | 11 wk | 12 wk |
|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------|------|--------------------|-------|-------|
| <b>Standard post auto SCT blood work:</b><br>CBC & Diff, EP, Creatinine, ALT, Alk Phos, LDH, Bili, Mg                                                                                                                                                                                                      | Frequency to be done in accordance with physician clinic assessment                                                                                                                    |      |      |                    |       |       |
| SPEP and serum Immunofixation (MM pt.'s only)                                                                                                                                                                                                                                                              |                                                                                                                                                                                        |      |      |                    | X     |       |
| Quantitative Immunoglobulins (IgG, IgA, IgM)                                                                                                                                                                                                                                                               |                                                                                                                                                                                        |      |      |                    |       | X     |
| PFT<br>*Do at 6 weeks post-SCT if patient has received BCNU                                                                                                                                                                                                                                                |                                                                                                                                                                                        |      | X    | Physician directed |       |       |
| Bone Marrow Asp/Bx <b><u>as ordered by physician.</u></b><br>Myeloma: Restage between Day + 80-90 (11 wk)                                                                                                                                                                                                  |                                                                                                                                                                                        |      |      |                    | X     |       |
| IRS Study (Storek Lab) Autoimmune patients only<br><br>Notify BMT Clinical Research RN if pt. relapses                                                                                                                                                                                                     | See Connect Care story board for "Research Participant" banner. If banner noted, see "SnapShot" found under "Chart Review" for research time points and specific dates of blood draws. |      |      |                    |       |       |
| Myeloma only: 24-hr urine testing for: creat clearance, T. protein, UPEP, Immunofixation <b><u>as ordered by physician.</u></b><br>Myeloma: Restage between Day + 80-90 (11 wk). Do not repeat at 12 wk.                                                                                                   |                                                                                                                                                                                        |      |      |                    | X     |       |
| Myeloma only: Free Light Chain (Serum)<br>If non secretory myeloma or light chain disease only                                                                                                                                                                                                             |                                                                                                                                                                                        |      |      |                    | X     |       |
| Skeletal Survey: Myeloma Only <b><u>as ordered by physician.</u></b>                                                                                                                                                                                                                                       |                                                                                                                                                                                        |      |      |                    | X     |       |
| AFP, Beta HCG (Testicular)<br>Germ Cell Tumors                                                                                                                                                                                                                                                             |                                                                                                                                                                                        |      |      |                    | X     |       |
| Amyloid <ul style="list-style-type: none"> <li>Serum testing for: Troponin T, FLC, alk phos, albumin</li> <li>24hr urine testing for: UPEP, SPEP, creat clearance</li> <li>Other: echocardiogram, Bone Marrow Asp/Bx At 1y, 2y, 3y, 5y, and then every 5 years post transplant (10y, 15y, etc).</li> </ul> |                                                                                                                                                                                        |      |      |                    | X     |       |
| VRE/MRSA surveillance swabs                                                                                                                                                                                                                                                                                | Frequency of swabs directed as per Infection Prevention and Control however ordered under the BMT Attending physician                                                                  |      |      |                    |       |       |

| TEST                                                                                           | 2<br>wk | 3<br>wk | 4<br>wk | 5<br>wk | 6<br>wk | 7<br>wk | 8<br>wk | 9<br>wk | 10<br>wk | 11<br>wk | 12<br>wk | 4<br>mo | 5<br>mo | 6<br>mo | 7<br>mo | 8<br>mo | 9<br>mo | 10<br>mo | 11<br>mo | 12<br>mo |
|------------------------------------------------------------------------------------------------|---------|---------|---------|---------|---------|---------|---------|---------|----------|----------|----------|---------|---------|---------|---------|---------|---------|----------|----------|----------|
| CD4, TSH, lipid panel, A1C<br><b>Females:</b> FSH, LH, estradiol<br><b>Males:</b> testosterone |         |         |         |         |         |         |         |         |          |          |          |         |         |         |         |         |         |          |          | x        |
| <b>Autoimmune Only</b> (i.e. Multiple Sclerosis, Scleroderma, Myasthenia Gravis)               |         |         |         |         |         |         |         |         |          |          |          |         |         |         |         |         |         |          |          |          |
| CMV PCR (Prov lab)                                                                             | x       | x       | x       | x       | x       | x       | x       | x       | x        | x        | x        |         |         |         |         |         |         |          |          |          |
| EBV PCR (Prov lab)                                                                             | x       | x       | x       | x       | x       | x       | x       | x       | x        | x        | x        |         |         |         |         |         |         |          |          |          |
| CBC & Diff, EP, Creatinine, ALT, Alk Phos, LDH, Bili                                           | x       | x       | x       | x       | x       | x       | x       | x       | x        | x        | x        |         |         |         |         |         |         |          |          |          |

| TEST                                                                                                                                                                                                                                                                                                                                                                                                 | Monthly | 6<br>mo                                                                                                                                                           | 12<br>mo | Every<br>3 mo | Every<br>6 mo | Yearly | 24<br>mo | 3<br>Years | q5<br>Years |
|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------|---------------|---------------|--------|----------|------------|-------------|
| CBC & Diff, EP<br>Weekly during RT.                                                                                                                                                                                                                                                                                                                                                                  |         | X                                                                                                                                                                 | X        |               | X             |        |          |            |             |
| Creatinine, ALT, Alk Phos, LDH, Bili                                                                                                                                                                                                                                                                                                                                                                 |         | X                                                                                                                                                                 | X        |               |               |        |          |            |             |
| Quantitative Immunoglobulins (IgG, IgA, IgM)                                                                                                                                                                                                                                                                                                                                                         |         | X                                                                                                                                                                 | X        | X             |               |        |          |            |             |
| SPEP<br><br>*Patients with M-protein detected in past, SPEP and CBC & Diff should be done every 3 months                                                                                                                                                                                                                                                                                             |         | X                                                                                                                                                                 | X        |               | X             |        |          |            |             |
| Myeloma Only: SPEP, UPEP, Serum Free Light Chains, Calcium                                                                                                                                                                                                                                                                                                                                           |         |                                                                                                                                                                   |          | X             |               |        |          |            |             |
| Bone Density<br>Six to twelve months post ASCT. Start bisphosphonate therapy if significant bone loss                                                                                                                                                                                                                                                                                                |         |                                                                                                                                                                   | X        |               |               |        |          |            |             |
| CT (Lymphoma) <u>as ordered by physician.</u>                                                                                                                                                                                                                                                                                                                                                        |         | X                                                                                                                                                                 |          |               |               |        |          |            |             |
| <b>Non-live vaccinations and COVID-19 and influenza vaccinations due</b><br>(Alberta Health Wellness Guidelines)<br>- Send referral to Population and Public Health<br>- Provide approval letter to patient and instruct them to book appointment.                                                                                                                                                   |         | X                                                                                                                                                                 |          |               |               |        |          |            |             |
| <b>Live vaccine clearance due</b><br>(Alberta Health Wellness Guidelines)<br>- Provide approval letter to patient and instruct them to book appointment.                                                                                                                                                                                                                                             |         |                                                                                                                                                                   |          |               |               |        | X        |            |             |
| <b>Other Immunization Serology</b><br>(Prov Lab)<br>- Anti-tetanus, measles (ALL pts)<br>-, Rubella (only in women in childbearing potential years)<br>- Anti-HBs (only if pt. is a health care worker or has lifestyle risks)<br><br>Monitor immunization serology for booster requirements and obtain order from primary physician.<br>Provide patient with <i>Booster Dose(s) Required letter</i> |         |                                                                                                                                                                   |          |               |               |        |          | X<br><br>X |             |
| <b>IRS Study (Storek Lab) - Autoimmune patients only</b><br>Notify BMT Clinical Research RN if pt. relapses                                                                                                                                                                                                                                                                                          |         | See EMR for "Research Participant" banner. If banner noted, see "SnapShot" found under "Chart Review" for research time points and specific dates of blood draws. |          |               |               |        |          |            |             |

## Alberta Blood and Marrow Transplant Program Follow-up Guidelines: Post-Allogeneic Transplant

| TEST                                                                                                                                                        | WEEK |   |   |   |   |                          |    |    |    |   | MONTHLY |   |   |   |   |    |    |    | YEARLY |     |     |     |  |
|-------------------------------------------------------------------------------------------------------------------------------------------------------------|------|---|---|---|---|--------------------------|----|----|----|---|---------|---|---|---|---|----|----|----|--------|-----|-----|-----|--|
|                                                                                                                                                             | 4    | 5 | 6 | 7 | 8 | 9                        | 10 | 11 | 12 | 4 | 5       | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 24     | q1y | q3y | q5y |  |
| CBC, Differential<br>Note: Retics (physician directed)                                                                                                      | x    | x | x | x | x | x                        | x  | x  | x  | x | x       | x | x | x | x | x  | x  | x  | x      | x   | x   | x   |  |
| If pt on CSA (Levels to be stopped once taper has begun). Begin taper @ Day +56 - off by Day +84, except for Aplastic Anemia. (Physician directed)          | x    | x | x | x | x | As directed by physician |    |    |    |   |         |   |   |   |   |    |    |    |        |     |     |     |  |
| If pt on Tacrolimus (Levels to be stopped once taper has begun).). Begin taper @ Day +56 - off by Day +84, except for Aplastic Anemia. (Physician directed) | x    | x | x | x | x | As directed by physician |    |    |    |   |         |   |   |   |   |    |    |    |        |     |     |     |  |
| EP, Creat, Liver panel 1, Mg, Random Glucose (if on steroids)                                                                                               | x    | x | x | x | x | x                        | x  | x  | x  | x | x       | x | x | x | x | x  | x  | x  | x      | x   |     |     |  |
| CMV PCR (Prov Lab)                                                                                                                                          | x    | x | x | x | x | x                        | x  | x  | x  | x | x       | x | x | x | x | x  | x  | x  |        |     |     |     |  |
| EBV PCR (Prov Lab)                                                                                                                                          | x    | x | x | x | x | x                        | x  | x  | x  | x | x       | x | x | x | x | x  | x  | x  |        |     |     |     |  |
| Ferritin, Transferrin saturation,                                                                                                                           |      |   |   |   |   |                          |    |    | x  |   |         |   |   |   |   |    |    | x  |        |     |     |     |  |
| IgG, IgM, IgA                                                                                                                                               |      |   |   |   |   |                          |    |    | x  |   |         | x |   |   |   |    |    | x  | x      |     | x   |     |  |
| Hemoglobin A1C and lipid profile                                                                                                                            |      |   |   |   |   |                          |    |    |    |   |         |   |   |   |   |    |    | x  |        |     |     |     |  |
| TSH                                                                                                                                                         |      |   |   |   |   |                          |    |    |    |   |         |   |   |   |   |    |    | x  | x      | x   |     |     |  |

## Alberta Blood and Marrow Transplant Program Follow-up Guidelines: Post-Allogeneic Transplant

| TEST                                                                                                                                                                                                                                                                                                                                                                                                                                    | WEEK |   |   |   |   |   |    |    |    |   | MONTHLY |   |   |   |   |    |    |    | YEARLY |     |     |     |   |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------|---|---|---|---|---|----|----|----|---|---------|---|---|---|---|----|----|----|--------|-----|-----|-----|---|
|                                                                                                                                                                                                                                                                                                                                                                                                                                         | 4    | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 4 | 5       | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 24     | q1y | q3y | q5y |   |
| <b>Non-live vaccinations and COVID-19 and influenza vaccinations due</b><br>(Alberta Health Wellness Guidelines)<br>-Send referral to Population and Public Health<br>- Provide approval letter to patient and instruct them to book appointment.                                                                                                                                                                                       |      |   |   |   |   |   |    |    |    |   |         | x |   |   |   |    |    |    |        |     |     |     |   |
| <b>Live vaccine clearance due</b><br>(Alberta Health Wellness Guidelines)<br>-Provide approval letter to patient and instruct them to book appointment                                                                                                                                                                                                                                                                                  |      |   |   |   |   |   |    |    |    |   |         |   |   |   |   |    |    |    | x      |     |     |     |   |
| <b>Immunization serology</b><br>(Prov Lab)<br>Order the following blood work:<br>-Anti-tetanus, measles (all pts.)<br>- Rubella (only women in child bearing potential years)<br>- Anti-HBs (only if pt. is a health care worker or has lifestyle risks)<br><br>Monitor immunization serology for booster requirements, if required obtain order from primary physician and provide patient with <i>Booster Dose(s) Required letter</i> |      |   |   |   |   |   |    |    |    |   |         |   |   |   |   |    |    |    |        |     | x   |     | x |
| <b>Hormones testing</b><br>Males : Free Testosterone<br>(sex hormone binding globulin. To be done between 7 and 10am.)<br><br>Females: FSH/LH/estradiol                                                                                                                                                                                                                                                                                 |      |   |   |   |   |   |    |    |    |   |         |   |   |   |   |    |    | x  |        |     |     |     |   |



## Alberta Blood and Marrow Transplant Program Follow-up Guidelines: Post-Allogeneic Transplant

| TEST                                                                                                                                                                                                                | WEEK                                                                                                                                                                                         |   |   |   |   |   |    |    |    |   | MONTHLY |   |   |   |   |    |    |    | YEARLY |     |     |     |  |
|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---|---|---|---|---|----|----|----|---|---------|---|---|---|---|----|----|----|--------|-----|-----|-----|--|
|                                                                                                                                                                                                                     | 4                                                                                                                                                                                            | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 4 | 5       | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 24     | q1y | q3y | q5y |  |
| Pharmacist –post-transplant medication review                                                                                                                                                                       |                                                                                                                                                                                              |   |   |   |   |   |    |    | x  |   |         |   |   |   |   |    |    | x  |        |     |     |     |  |
| Cardiovascular Risk Assessment                                                                                                                                                                                      |                                                                                                                                                                                              |   |   |   |   |   |    |    |    |   |         |   |   |   |   |    |    | x  |        |     |     |     |  |
| IRS and Predictors of Relapse (Storek Lab)                                                                                                                                                                          | See EMR for Research Participant banner. If patient flagged as “Research participant”, see “SnapShot” found under “Chart Review” for research time points and specific dates of blood draws. |   |   |   |   |   |    |    |    |   |         |   |   |   |   |    |    |    |        |     |     |     |  |
| Notify BMT Clinical Research RN if:<br>- pt. has relapsed *sample to be collected at time of relapse<br>- pt. has graft failure<br>- pt. has PTLD (only if requires an admission)<br>- pt. has secondary malignancy |                                                                                                                                                                                              |   |   |   |   |   |    |    |    |   |         |   |   |   |   |    |    |    |        |     |     |     |  |

## Alberta Blood and Marrow Transplant Program Follow-up Guidelines: Post-Allogeneic Transplant

| TEST                                                                                                                                                                       | WEEK |   |   |   |   |   |    |    |                |                | MONTHLY |   |   |   |   |    |    |                | YEARLY |     |     |     |  |
|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------|---|---|---|---|---|----|----|----------------|----------------|---------|---|---|---|---|----|----|----------------|--------|-----|-----|-----|--|
|                                                                                                                                                                            | 4    | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12             | 4              | 5       | 6 | 7 | 8 | 9 | 10 | 11 | 12             | 24     | q1y | q3y | q5y |  |
| Bone Density (Spine and Hip)<br>*Physician directed, consider if patient has been on steroids                                                                              |      |   |   |   |   |   |    |    |                |                |         |   |   |   |   |    |    | x <sup>*</sup> |        |     |     |     |  |
| Ophthalmology Follow-up<br>*Calgary pts: RN to instruct pt. to arrange                                                                                                     |      |   |   |   |   |   |    |    | x              |                |         |   |   |   |   |    |    |                |        |     |     |     |  |
| Schirmer's Testing                                                                                                                                                         |      |   |   |   |   |   |    |    | x              |                |         | x |   |   | x |    |    | x              |        |     |     |     |  |
| Gynecology Follow-up<br>*Calgary pts. :RN to instruct pt. to arrange                                                                                                       |      |   |   |   |   |   |    |    |                |                |         | x |   |   |   |    |    |                |        |     |     |     |  |
| Dental<br>*Dental: TBB @3mths for Calgary pts.,<br>TBB@4mths for Edmonton pts.<br>*Calgary pts.: RN to instruct pt. to arrange                                             |      |   |   |   |   |   |    |    | x <sup>*</sup> | x <sup>*</sup> |         |   |   |   |   |    |    |                |        |     |     |     |  |
| Bone Marrow Aspirate +/- Biopsy<br><br>Order as directed by physician<br><br>Do NOT order chimerism on marrow<br>(routine chimerism is performed on blood,<br>not marrow). |      |   |   |   |   |   |    |    | x              |                |         |   |   |   |   |    |    |                |        |     |     |     |  |

## Alberta Blood and Marrow Transplant Program Follow-up Guidelines: Post-Allogeneic Transplant

| TEST                                                                                                                                                                                               | WEEK                                                                                                                  |   |   |   |   |   |    |     |    |   | MONTHLY |   |   |   |     |    |     |    |                              |             | YEARLY |     |  |  |  |  |
|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------|---|---|---|---|---|----|-----|----|---|---------|---|---|---|-----|----|-----|----|------------------------------|-------------|--------|-----|--|--|--|--|
|                                                                                                                                                                                                    | 4                                                                                                                     | 5 | 6 | 7 | 8 | 9 | 10 | 11  | 12 | 4 | 5       | 6 | 7 | 8 | 9   | 10 | 11  | 12 | 24                           | q1y         | q3y    | q5y |  |  |  |  |
| <b>Blood for Molecular Studies</b><br>Order as directed by physician<br><br>Example: bcr/abl, PML/RAR, FLT3, and<br>*Suggested interval for testing however<br>frequency may be physician directed |                                                                                                                       |   |   |   |   |   |    | x * |    |   | x *     |   |   |   | x * |    | x * |    | x *                          | Then yearly |        |     |  |  |  |  |
| <b>Blood for Chimerism</b><br>(CD3 cells, and malignancy<br>lineage/phenotype cells)                                                                                                               |                                                                                                                       |   |   |   |   |   |    |     | x  |   |         |   |   |   |     |    |     |    |                              |             |        |     |  |  |  |  |
| CT chest/abdomen/pelvis – only<br>lymphoma or CLL<br>*if no prior CR confirmed                                                                                                                     |                                                                                                                       |   |   |   |   |   |    |     | X* |   |         |   |   |   |     |    |     | x  |                              |             |        |     |  |  |  |  |
| CXR (if abnormal PFT)                                                                                                                                                                              |                                                                                                                       |   |   |   |   |   |    |     |    | x |         |   |   |   |     |    |     |    |                              |             |        |     |  |  |  |  |
| PFT<br>*ONLY required if history of acute or chronic<br>GVHD*<br>*Edmonton patients booked @ CCI on return<br>visit (unless delayed return)                                                        |                                                                                                                       |   |   |   |   |   |    |     | x* |   | x       |   |   |   | x   |    |     | x  | Yearly for the first 5 years |             |        |     |  |  |  |  |
| VRE/MRSA surveillance swabs                                                                                                                                                                        | Frequency of swabs directed as per Infection Prevention and Control however ordered under the BMT Attending physician |   |   |   |   |   |    |     |    |   |         |   |   |   |     |    |     |    |                              |             |        |     |  |  |  |  |
| <b>Multiple Myeloma:</b><br>Calcium, SPEP, UPEP, Free Light Chain                                                                                                                                  |                                                                                                                       |   | x |   |   |   |    | x   |    |   | x       |   |   | x |     |    | x   |    | x                            |             |        |     |  |  |  |  |
| <b>Multiple Myeloma:</b><br>Skeletal survey                                                                                                                                                        |                                                                                                                       |   |   |   |   |   |    | x   |    |   |         |   |   |   |     |    | x   |    |                              | x           |        |     |  |  |  |  |

| TEST / ASSESSMENT / PROCEDURE                                                                                                                                                                                                                                                                                                                                                                                                        | 3 wk                                                                                                                 | 4 wk |
|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------|------|
| <b>Standard post CAR T-cell blood work:</b><br>CBC and differential, electrolyte panel, calcium, magnesium, creatinine, ALT, alkaline phosphate, albumin, LDH, total bilirubin, uric acid, c-reactive protein, ferritin                                                                                                                                                                                                              | Frequency done in accordance with physician clinic assessment                                                        |      |
| Weekly assessment by ABMTP physician <ul style="list-style-type: none"> <li>History and physical</li> <li>Labs (as above)</li> <li>Cytokine Release Syndrome (CRS) and Neurotoxicity Assessment (<b>BOTH NURSING AND PHYSICIAN SECTIONS</b>)               <ul style="list-style-type: none"> <li>CRS/Neurotoxicity Assessment Flowsheet</li> <li>Neurotoxicity Monitoring BMTF70007</li> </ul> </li> </ul>                          | To be completed at minimum weekly.                                                                                   |      |
| Additional non-MD nursing visits (1-2x/week at ABMTP physician discretion) <ul style="list-style-type: none"> <li>Labs (as above)</li> <li>Cytokine Release Syndrome (CRS) and Neurotoxicity assessment (<b>NURSING SECTION ONLY</b>)               <ul style="list-style-type: none"> <li>Cytokine Release Syndrome (CRS) and Neurotoxicity Monitoring Flowsheet</li> <li>Neurotoxicity Monitoring BMTF70007</li> </ul> </li> </ul> | Physician directed                                                                                                   |      |
| Antimicrobial prophylaxis - Septra & Valtrex                                                                                                                                                                                                                                                                                                                                                                                         | As per Standard Practice Manual for Hematopoietic Cell Therapy (HCT)                                                 |      |
| Growth factor support                                                                                                                                                                                                                                                                                                                                                                                                                | Physician directed                                                                                                   |      |
| Bone marrow aspirate & biopsy (morphology, flow cytometry and cytogenetics)<br>Leukemia disease Surveillance ( <i>can be done day 28-32</i> )                                                                                                                                                                                                                                                                                        |                                                                                                                      | X    |
| Computerized Tomography (CT) Scan<br>Lymphoma disease surveillance ( <i>can be done day 28-32</i> )                                                                                                                                                                                                                                                                                                                                  |                                                                                                                      | X    |
| MRSA surveillance swabs                                                                                                                                                                                                                                                                                                                                                                                                              | Frequency of swabs directed as per Infection Prevention and Control (IPC), however ordered under the ABMTP physician |      |

| TEST / ASSESSMENT / PROCEDURE                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       | 2 mo                                                          | 3 mos | Q 1-3 mo  | 12 mo |
|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---------------------------------------------------------------|-------|-----------|-------|
| <b>Labs:</b> CBC and differential, electrolyte panel, calcium, magnesium, creatinine, ALT, alkaline phosphate, albumin, LDH, total bilirubin, uric acid, Immunoglobulins (IgG, IgA, and IgM),                                                                                                                                                                                                                                                                                                                                                                                                                       | X                                                             | X     | X         | X     |
| CD4, TSH, lipid panel, A1C<br>Females: FSH, LH, estradiol<br>Males: testosterone                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    |                                                               |       |           | X     |
| B-cell aplasia bloodwork <b>*Leukemia disease ONLY*</b> <ul style="list-style-type: none"><li>Flow cytometry (peripheral blood) - Immunodeficiency screening panel</li></ul>                                                                                                                                                                                                                                                                                                                                                                                                                                        |                                                               | X     | Mos 6 & 9 | X     |
| Physician clinic assessment – minimum recommendation as per schedule <ul style="list-style-type: none"><li>History and physical</li><li>Labs (as above)</li></ul>                                                                                                                                                                                                                                                                                                                                                                                                                                                   | X                                                             | X     | X         | X     |
| Cytokine Release Syndrome (CRS) and neurotoxicity assessment <ul style="list-style-type: none"><li>To be done in accordance with physician clinic assessment</li><li>The following methods are appropriate options for CRS/Neurotoxicity assessment as per physician direction.<ul style="list-style-type: none"><li>Connect Care Cytokine Release Syndrome (CRS) and Neurotoxicity Monitoring Flowsheet</li><li>Neurotoxicity Monitoring – Handwriting monitoring (see appendix for example)</li><li>American Society for Transplantation and Cellular Therapy (ASTCT) Practice Guidelines App</li></ul></li></ul> | Physician directed<br><br>No longer recommended after 8 weeks |       |           |       |
| Growth factor support                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               |                                                               |       |           |       |
| Antimicrobial prophylaxis - Septra & Valtrex                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        | As per Standard Practice Manual for HCT                       |       |           |       |
| Intravenous immunoglobulin (IVIg) replacement                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       | Physician directed                                            |       |           |       |
| Positron emission tomography/ computerized tomography (PET/CT) scan: disease status assessment lymphoma – to be ordered by ABMTP physician                                                                                                                                                                                                                                                                                                                                                                                                                                                                          |                                                               | X     |           |       |
| Bone marrow aspirate & biopsy (morphology, flow cytometry and cytogenetics):<br>Disease status assessment Leukemia - (ONLY required if abnormal at 4 week period)                                                                                                                                                                                                                                                                                                                                                                                                                                                   |                                                               | X     |           |       |
| <b>COVID-19 and Influenza vaccination Due</b><br>(Alberta Health Wellness Guidelines)<br>- Provide approval letter to patient and instruct them to book appointment                                                                                                                                                                                                                                                                                                                                                                                                                                                 |                                                               | X     |           |       |

| TEST / ASSESSMENT / PROCEDURE                                                                                                                                                                              | 6 mo                                    | Yearly | 24 mo | 3 Years | Years 2-15 |
|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------|--------|-------|---------|------------|
| Assessment as directed by physician (minimum 1/year)                                                                                                                                                       |                                         | X      |       |         |            |
| Antimicrobial prophylaxis <ul style="list-style-type: none"> <li>Septra &amp; Valtrex</li> </ul>                                                                                                           | As per Standard Practice Manual for HCT |        |       |         |            |
| <b>Immunization Serology in All Patients</b><br>(Prov Lab)<br>- Anti-tetanus<br>- Measles<br>- Rubella<br>- Anti-HBs                                                                                       | X                                       |        | X     |         |            |
| <b>Non-live vaccinations due</b><br>(Alberta Health Wellness Guidelines)<br>- Send referral to Population and Public Health<br>- Provide approval letter to patient and instruct them to book appointment. | X                                       |        |       |         |            |
| <b>Live vaccine clearance due</b><br>(Alberta Health Wellness Guidelines)<br>- Provide approval letter to patient and instruct them to book appointment.                                                   |                                         |        | X     |         |            |
| Diseases Surveillance - Discretion of Physician                                                                                                                                                            |                                         |        |       |         | X          |



# Appendix



Patient Label

## Alberta Blood and Marrow Transplant Program Neurotoxicity Monitoring – Handwriting Assessment

**Instructions:**  
Handwriting assessment for neurotoxicity shall be completed as part of the **Neurotoxicity 10 Point Assessment (ICE)** within the CRS/Neurotoxicity Assessment flowsheet within the patient’s Electronic Medical Record (EMR).  
  
An alternate sentence may be used with patients when English is not their first language or utilize Neurotoxicity Monitoring – Modified Handwriting Assessment BMTF70033 for persons with literacy issues.

Monitoring Frequency: As per physician orders

Write the following sentence: **The leaf on the Canadian flag is red.**

|      |      |  |                |
|------|------|--|----------------|
| Date | Time |  | RN<br>Initials |
|      | Time |  | RN<br>Initials |
|      | PRN  |  | RN<br>Initials |

|      |      |  |                |
|------|------|--|----------------|
| Date | Time |  | RN<br>Initials |
|      | Time |  | RN<br>Initials |
|      | PRN  |  | RN<br>Initials |

|      |      |  |                |
|------|------|--|----------------|
| Date | Time |  | RN<br>Initials |
|      | Time |  | RN<br>Initials |
|      | PRN  |  | RN<br>Initials |

|      |      |  |                |
|------|------|--|----------------|
| Date | Time |  | RN<br>Initials |
|      | Time |  | RN<br>Initials |
|      | PRN  |  | RN<br>Initials |

|      |      |  |                |
|------|------|--|----------------|
| Date | Time |  | RN<br>Initials |
|      | Time |  | RN<br>Initials |
|      | PRN  |  | RN<br>Initials |

## Abbreviations

2-CDA, 2-chlorodeoxyadeno-sine; AAIPI, age-adjusted international prognostic index; ABG, arterial blood gas; ABMTP, Alberta Bone Marrow Transplant Program; ABMTR, Alberta Bone Marrow Transplant Registry; ABVD, adriamycin + bleomycin + vinblastine + dacarbazine; ABW, actual body weight; ACA, additional cytogenetic abnormalities OR anti-centromere antibody (depending on section); ACE, angiotensin-converting enzyme; ACTH, adrenocorticotrophic hormone; ADL, activities of daily living; aGVHD, acute graft-versus-host-disease; AIBW, adjusted ideal body weight; AILD, angioimmunoblastic lymphadenopathy with dysproteinemia; AITL, angioimmunoblastic T-cell lymphoma; AL, amyloid light-chain; ALCL, anaplastic large cell lymphoma; Alem, alemtuzumab; Alk p, alkaline phosphatase; ALL, acute lymphoblastic leukemia; alloSCT, allogeneic stem cell transplant; ALT, alanine transaminase / alanine aminotransferase; ALT, adult T-cell lymphoma; AMBd, amphotericin B deoxycholate; AML, acute myeloid leukemia; ANA, antinuclear antibody; ANC, absolute neutrophil count; AP, accelerated phase; APC, antigen presenting cells; aPML, acute promyelocytic leukemia; ARR, annual relapse rate; ARS, antigen recognition site; ASBMT, American Society for Blood and Marrow Transplantation; ASCO, American Society of Clinical Oncology; ASCT, allogeneic stem cell transplantation OR autologous stem cell transplantation (dependent upon section); AST, aspartate aminotransferase OR alanine aminotransferase (dependent upon section); ATG, antithymocyte globulin; AUC, area under the curve; autoSCT, autologous stem cell transplant; BAL, bronchoalveolar lavage; BAT, best-available therapy; BC, blast crisis; BCLL, B-cell chronic lymphocytic leukemia; BCR, B-cell receptor; BCT, blood cell transplantation; BEAC, BCNU + etoposide + Ara-C + cyclophosphamide; BEAM, BCNU + etoposide + AraC + melphalan; BEAU, BCNU + etoposide + Ara-C + cyclophosphamide; BL, Burkitt lymphoma; BM, busulfan + melphalan; BMA, bone marrow aspirate; BMD, bone mineral density; BMT, bone marrow transplantation; BOOP, bronchiolitis obliterans organizing pneumonia; BP, blast phase; BSA, body surface area; Bu, busulfan; BuCy, busulfan + cyclophosphamide; CA, cytogenetic abnormalities; CAP, cyclophosphamide + doxorubicin + prednisone; CAR-T, chimeric antigen receptor T-cells; CBC, complete blood count; CBF, core binding factor; CCR or CCyR or CCgR, complete cytogenetic response; CDC, Centers for Disease Control and Prevention; CEB or CBV, cyclophosphamide + etoposide + carmustine; cGVHD, chronic graft-versus-host-disease; CHF, congestive heart failure; CHF, congestive heart failure; CHOP, cyclophosphamide + adriamycin + vincristine + prednisone; CHR, complete hematologic response; CI, confidence interval; CIBMTR, Center for International Blood and Marrow Transplant Research; CK, creatine kinase; CLL, chronic lymphocytic leukemia; CML, chronic myeloid leukemia; CMML, chronic myelomonocytic leukemia; CMR, complete molecular response; CMV, cytomegalovirus; CN, cytogenetically normal; CNI, calcineurin inhibitor; CNS, central nervous system; CP, chronic phase; CR, complete remission/response; CR1, 1st complete remission; CR2, second complete response; CRBSI, catheter-related bloodstream infection; CRe, early complete remission; CRel, correctly released; CREST, calcinosis of skin, Raynaud's

phenomenon, esophageal dysmotility, sclerodactyly, teleangiectasia; CRS, cytokine release syndrome; Csa, cyclosporine; CSF, cerebrospinal fluid; CT, computerized tomography; CVAMP, cyclophosphamide + vincristine + doxorubicin + methylprednisolone; CVC, central venous catheter; Cy, cyclophosphamide; CyA-Mtx, cyclosporine + methotrexate; Cy-ATG, cyclosporine + antithymocyte globulin; CyR, cytogenetic response; CyTBI or VPCyTBI, cyclophosphamide and possible etoposide (VP-16); DEXA scan, dual energy X-ray absorptiometry; DFS, disease-free survival; DHAP, dexamethasone + Ara-C (cytarabine) + cisplatin; DICEP, dose-intensified cyclophosphamide 5.25 g/m<sup>2</sup>, etoposide 1.05g/m<sup>2</sup>, and cisplatin 105 mg/m<sup>2</sup>; DIPSS, Dynamic International Prognostic Scoring System; DLBCL, diffuse large B-cell lymphoma; DLCO, diffusing capacity of lung for carbon monoxide; DLI, donor lymphocytic infusion; DMT, disease modifying therapy; DSA, donor specific antibodies; DVT, deep vein thrombosis; EBER, EBV-encoded RNA; EBMT, European Group for Blood and Marrow Transplantation; EBMT, European Bone Marrow Transplant Registry; EBV, Epstein-Barr virus; ECG, electrocardiogram; ECOG, Eastern Cooperative Oncology Group; ECP, extracorporeal photopheresis; ECTRIMS, European Committee for Treatment and Research in Multiple Sclerosis; ED, erectile dysfunction; EDSS, expanded disability status scale; EFS, event-free survival; EGD, upper endoscopy; ELN, European LeukemiaNet; ENT, ear, nose, and throat; ESR, erythrocyte sedimentation rate; ET, essential thrombocythemia; EVIL, European Conference on Infections in Leukemia; FAB, French-American-British; FCA, fludarabine + Cy-ATG; FCR, fludarabine + cyclophosphamide + rituximab; FEV1, forced expiratory volume in one second; FFS, freedom from second failure OR failure-free survival (depending on chapter); FHF, fulminant hepatic failure; FISH, fluorescence in situ hybridization; FL, follicular lymphoma; FLIPI, follicular lymphoma international prognostic index; FLU, fludarabine; FLUBU, fludarabine + busulfan; FND, fludarabine + mitoxantrone + dexamethasone; FSH, follicle stimulating hormone; FSS, functional system scores; FTBI, fractionated total body irradiation; FVC, forced vital capacity; GB, gallbladder; GCSF, granulocyte colony stimulating factor; GD, gadolinium; GDP, gemcitabine + dexamethasone + cisplatin; GI, gastrointestinal; GM-CSF, granulocyte-macrophage colony-stimulating factor; GnRH, gonadotropin-releasing hormone; GVH or GVHD, graft-versus-host or graft-versus-host disease; GVL, graft-versus-leukemia; HAMA, human anti-mouse antibodies; HAP, hospital acquired pneumonia; hATG, horse ATG; Hb, hemoglobin; Hb S, sickled hemoglobin; HBeAg, hepatitis B viral protein; HBsAg, hepatitis B surface antigen; HBV, hepatitis B virus; HCC, hepatocellular carcinoma; HCT/HSCT, hematopoietic stem cell transplantation; HCT-CI, Hematopoietic cell transplantation comorbidity index; HCV, hepatitis C virus; HDCT/HDT, high-dose chemotherapy; HHV6, human herpes virus 6; HiDAC, high dose Ara-c; HLA, human leukocyte antigens; HLA-DSA, donor-specific HLA antibodies; HMPV, human metapneumovirus; HR, hazard ratio; HRT, hormone replacement therapy; HSV, herpes simplex virus; HTLV-1, Human T-cell lymphoma virus type 1; IBMTR, International Bone Marrow Transplant Registry; IBW, ideal body weight; ICE, ifosfamide + carboplatin + etoposide; ICSI, intracytoplasmic sperm injection; ICU, intensive care unit;

IFI, invasive fungal infection; IL-2, interleukin-2; INR, international normalized ratio; IPI, international prognostic index; IPS, idiopathic pneumonia syndrome; IPSS, International Prognostic Scoring System; IRAEs, infusion-related adverse events; IST, immunosuppressive therapy; ITD, internal tandem duplication; ITT, intent-to-treat; IV, intravenous; IVIMG or IVIG, intravenous immunoglobulin; JAK, Janus Kinase; KGF, keratinocyte growth factor; KIR, killer-cell immunoglobulin like receptor; KPS, Karnofsky Performance Scale; KSAV, Kaposi's sarcoma-associated virus; L ampho B, liposomal amphotericin B; LBCL, large B-cell lymphoma; LBL, lymphoblastic lymphoma; LDH, lactate dehydrogenase test; LDHD, lymphocyte depletion Hodgkin disease; LES, lower esophageal sphincter; LFS, leukemia-free survival OR lung function score (dependent upon section); LH, luteinizing hormone; LMWH, low molecular weight heparin; LP, lumbar puncture; LPHD, lymphocyte predominant Hodgkin disease; LPL, lymphoplasmacytic lymphoma; LPS, Lansky Performance Status; LRCHD, lymphocyte-rich classical Hodgkin disease; LVEF, left ventricular ejection fraction; MALT, mucosa-associated lymphoid tissue; MBL, metallo-beta-lactamase; MCHD, mixed cellularity Hodgkin disease; MCL, mantle cell leukemia/lymphoma; MCR or MCyR, major cytogenetic response; MDS, Myelodysplasia / myelodysplastic syndrome; MEL, melphalan; MeVPTBI, melphalan + etoposide with or without TBI; MHC, major histocompatibility complex; miHAs, minor histocompatibility antigens; MK, monosomal karyotype; MM, mismatched; MMF, mycophenolate mofetil; MMR, major molecular response; MMRD, mismatched related donor; MP, melphalan + prednisone; MPD, methylprednisolone; MPT, melphalan + prednisone + thalidomide; MRC, Medical Research Council; MRD, minimal residual disease; MRel, mistakenly released; MRI, magnetic resonance imaging; MRSA, methicillin-resistant *Staphylococcus aureus*; mRSS, modified Rodnan skin score; MS, Multiple Sclerosis; MSD, matched sibling donor; MSKCC, Memorial Sloan-Kettering Cancer Center; mTOR, mammalian target of rapamycin; MTX, methotrexate; MUD, matched unrelated donor; MUGA, multiple gated acquisition scan; MZL, marginal zone lymphoma; NBTE, nonbacterial thrombotic endocarditis; NCCN, National Comprehensive Cancer Network; NHL, non-Hodgkin lymphoma; NK, natural killer cells; NMDP, National Marrow Donor Program; NOS, not otherwise specified; NR, no response; NRM, non-relapse mortality; NSAID, nonsteroidal anti-inflammatory drugs; NSHD, nodular sclerosis Hodgkin disease; NST, non-myeloablative conditioning; NT-proBNP, N-terminal propeptide brain-type natriuretic peptide; OR, odds ratio; ORR, overall response rate; OS, overall survival; PA, posterior-anterior; PAH, pulmonary arterial hypertension; PAP, pulmonary arterial pressure; PBPC, peripheral blood progenitor cells; PBSC, peripheral blood stem cells; PCNSL, primary central nervous system lymphoma; PCP, *Pneumocystis jirovecii* pneumonia; PCR, Polymerase chain reaction; PDGFR, platelet-derived growth factor receptor gene; PE, pulmonary embolism; PET, positron-emission tomography; PFS, progression-free survival; PFT, pulmonary function test; PHN, post-herpetic neuralgia; PICC, peripherally inserted central catheter; PK, Pharmacokinetics; PMF, primary myelofibrosis; PML, progressive multifocal leukoencephalopathy; PMLCL, primary mediastinal large B-cell lymphoma; PNH, paroxysmal nocturnal hemoglobinuria;

PPI, proton pump inhibitor; PR, partial response/remission; Pred, Prednisone; PT, prothrombin time; PTCL, peripheral T-cell lymphoma; PTCy, post-transplantation cyclophosphamide; PTLD, Post-transplant lymphoproliferative disorder; PTT, partial thromboplastin time; PUD, peptic ulcer disease; PV, polycythemia vera; QID PO, four times a day per os (orally); QOL, quality of life; Q-PCR, quantitative polymerase chain reaction; R, rituximab; rATG, rabbit ATG; R-CHOP, rituximab + cyclophosphamide + Adriamycin + vincristine + prednisone; RCT, randomized controlled trial; RCVP, rituximab + cyclophosphamide + vincristine + prednisone; R-DHAP, rituximab + dexamethasone + Ara-C (cytarabine) + cisplatin; RFCM, rituximab + fludarabine + cyclophosphamide + mitoxantrone; RFND, rituximab + fludarabine + mitoxantrone + dexamethasone; RFS, relapse-free survival; RI, reduction of immunosuppression; RIC, reduced intensity conditioning; R-ICE, rituximab + ifosfamide + carboplatin + etoposide; RIST, Reduced intensity SCT; RIT, radioimmunoconjugate therapy; RPLS, reversible posterior leukoencephalopathy syndrome; RR, response rate; RRMS, relapsing-remitting multiple sclerosis; RSV, respiratory syncytial virus; SAA, severe aplastic anemia; SAAIPI, Salvage Age Adjusted International Prognostic Index; SCA, sickle cell anemia; SCD, sickle cell disease; SCID, severe combined immunodeficiency; SCT, stem cell transplant; SDCT, standard-dose chemotherapy; SDH, subdural hematoma; siPI, second-line International Prognostic Index; SIRS, systemic inflammatory response syndrome; SLL, small lymphocytic leukemia; SOS, sinusoidal obstruction syndrome; SSc, systemic sclerosis; STPR, skin thickness progression rate; STR, short tandem repeat; SWOG, Southwest Oncology Group; T2/FLAIR, T2-weighted-fluid-attenuated inversion recovery; Tac, tacrolimus; t-AML, therapy-related AML; TBC, thiotepa + busulfan + cyclophosphamide; TBI, total body irradiation; TEE, transesophageal echocardiography; TGF- $\beta$ , transforming growth factor beta; TIPS, transjugular intrahepatic portosystemic shunt; TKI, tyrosine kinase inhibitor; TMA, thrombotic microangiopathy; TMP-SMX, co-trimoxazole (Septra®, Bactrim®); TNC, Total nucleated cell; TRM, transplant-related mortality or treatment-related mortality; Trv, tricuspid regurgitant velocity; TSH, thyroid stimulating hormone; TTA, time to antibiotic; TTP, time to positivity; UCBT, umbilical cord blood transplantation; UGI, upper gastrointestinal series (test); ULN, upper limit of normal; URTI, upper respiratory tract infection; UV, ultraviolet; VAD, vincristine + adriamycin + dexamethasone; VGPR, very good partial response; VOD, veno-occlusive disease; VP-16, etoposide; VRE, vancomycin-resistant *Enterococcus*; VRSA, vancomycin-resistant *Staphylococcus aureus*; VUD, volunteer unrelated donor; VZV, varicella zoster virus; WBC, white blood cell; WBRT, whole-brain radiotherapy; WHO, World Health Organization; WMUD, 'well matched' unrelated donor; WPSS, World Health Organization Prognostic Scoring System; -X or -Y, deleted X or Y chromosome.

## Disclaimer

The recommendations contained in this guideline are a consensus of the Alberta Bone Marrow and Blood Cell Transplant Program and are a synthesis of currently accepted approaches to management, derived from a review of relevant scientific literature. Clinicians applying these guidelines should, in consultation with the patient, use independent medical judgment in the context of individual clinical circumstances to direct care.

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[Outpatient Cancer Drug Benefit Program Master List](#).

## Conflict of Interest Statements

*Generated using the standard COI form.*

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| Date          | Topic Title                                                                          | Action  |
|---------------|--------------------------------------------------------------------------------------|---------|
| July 6, 2011  | Severe Aplastic Anemia: Indications for Stem Cell Transplantation                    | Updated |
| Sept 12, 2011 | Management of Relapse after Stem Cell Transplantation                                | Updated |
| Sept 14, 2011 | BCR-ABL1-Negative Myeloproliferative Neoplasms                                       | New     |
| Oct 4, 2011   | Transplantation for CLL                                                              | Updated |
| Oct 5, 2011   | Management of Chronic Graft Versus Host Disease                                      | New     |
| Oct 25, 2011  | Chimerism and Its Uses                                                               | New     |
| Dec 13, 2011  | MDS and Secondary AML: Indications for Transplantation                               | Updated |
| Dec 13, 2011  | Acute GVHD: Prevention and Treatment                                                 | Updated |
| Dec 13, 2011  | Catheter-Related Complications                                                       | Updated |
| Dec 13, 2011  | Head and Neck Complications, Including Mucositis                                     | Updated |
| Dec 13, 2011  | Donor Management, Including Mobilization                                             | New     |
| Jan 26, 2012  | Fungal Infections Before, During and After Transplant                                | Updated |
| Jan 26, 2012  | CMV and other Herpes Viruses                                                         | New     |
| Mar 16, 2012  | Acute GVHD: Prevention and Treatment                                                 | Updated |
| Mar 16, 2012  | Workup and Treatment of Fever Post-Transplant                                        | Updated |
| Mar 16, 2012  | Cardiac Complications of Transplant                                                  | Updated |
| Mar 16, 2012  | Umbilical Cord Blood Transplantation                                                 | New     |
| July 27, 2012 | CMV and other Herpes Viruses                                                         | Updated |
| Oct 17, 2012  | Criteria for Donor Selection                                                         | Updated |
| Oct 17, 2012  | GI Complications of Transplant                                                       | Updated |
| Oct 17, 2012  | Urinary and Renal Complications                                                      | New     |
| Oct 17, 2012  | Epstein Barr Virus/Posttransplant Lymphoproliferative Disorder                       | New     |
| Oct 17, 2012  | Poor Graft Function and Engraftment Failure                                          | New     |
| Oct 17, 2012  | Vaccination                                                                          | New     |
| Oct 17, 2012  | Hepatic Complications and Viral Hepatitis                                            | Updated |
| Oct 17, 2012  | Therapeutic Drug Monitoring                                                          | New     |
| Nov 12, 2012  | Management of the ABO-Incompatible Graft and Recipient                               | New     |
| Feb 5, 2013   | CMV and other Herpes Viruses                                                         | Updated |
| Feb 5, 2013   | Management of Chronic GVHD                                                           | Updated |
| Feb 5, 2013   | Pneumocystis & Bacterial Prophylaxis                                                 | New     |
| May 28, 2013  | Donor Management, Including Mobilization                                             | Updated |
| June 20, 2013 | Vaccination                                                                          | Updated |
| Aug 21, 2013  | Catheter-Related Complications                                                       | Updated |
| Mar 13, 2014  | Criteria for Donor Selection                                                         | Updated |
| Mar 14, 2014  | Cord blood transplants                                                               | Updated |
| Mar 14, 2014  | Acute GVHD, Prevention and Treatment                                                 | Updated |
| Mar 14, 2014  | Criteria for Donor Selection                                                         | Updated |
| May 28, 2014  | CMV, VZV, HSV, HHV6 (formerly CMV and other Herpes viruses)                          | Updated |
| Jul 22, 2014  | Management of Chronic Graft Versus Host Disease                                      | Updated |
| Oct 17, 2014  | Distribution of Microbially-Contaminated or Non-Conforming Cellular Therapy Products | New     |



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| Oct 17, 2014   | Pretransplant Conditioning                                        | New     |
| Nov 10, 2014   | Autologous Hematopoietic SCT for Active Multiple Sclerosis        | New     |
| Nov 10, 2014   | Transplantation for Acute Lymphoblastic Leukemia                  | Updated |
| Feb 18, 2015   | Pneumocystis and Bacterial Prophylaxis                            | Updated |
| Feb 26, 2015   | Pretransplant Conditioning                                        | Updated |
| Feb 26, 2015   | BCR-ABL1 Negative Myeloproliferative Neoplasms                    | Updated |
| Feb 26, 2015   | Acute Myeloid Leukemia: Indications for Stem Cell Transplant      | Updated |
| Feb 26, 2015   | Hodgkin and Non-Hodgkin Lymphoma: Indications for Transplantation | Updated |
| Mar 17, 2015   | Transplantation for Germ Cell Tumours                             | Updated |
| April 28, 2015 | Hepatic Complications and Viral Hepatitis                         | Updated |
| Sep 8, 2015    | CMV, VZV, HSV, HHV6                                               | Updated |
| Sep 15, 2015   | Epstein-Barr Virus/Posttransplant Lymphoproliferative Disorder    | Updated |
| Nov 24, 2015   | Pretransplant Conditioning                                        | Updated |
| May 17, 2016   | Transplantation for Scleroderma/Systemic Sclerosis                | New     |
| June 20, 2016  | Epstein-Barr Virus / Posttransplant Lymphoproliferative Disorder  | Updated |
| Nov 25, 2016   | Hemoglobinopathies                                                | New     |
| Dec 9, 2016    | Autologous Hematopoietic SCT for Active Multiple Sclerosis        | Updated |
| Jan 3, 2017    | Hematopoietic Cell Transplantation for Severe Aplastic Anemia     | Updated |
| Jan 16, 2017   | Acute GVHD: Prevention and Treatment                              | Updated |
| Jan 16, 2017   | Acute Myeloid Leukemia                                            | Updated |
| Jan 24, 2017   | Chronic Lymphocytic Leukemia                                      | Updated |
| Feb 8, 2017    | Vaccination                                                       | Updated |
| Feb 8, 2017    | Chronic GVHD                                                      | Updated |
| Feb 8, 2017    | Pneumocystis & Bacterial Prophylaxis                              | Updated |
| Feb 9, 2017    | CMV, VZV, HSV, HHV6                                               | Updated |
| Aug 2, 2017    | Epstein Barr Virus / Posttransplant Lymphoproliferative Disorder  | Updated |
| Aug 2, 2017    | Fungal Prophylaxis                                                | Updated |
| Aug 2, 2017    | Criteria for Donor Selection                                      | Updated |
| Aug 2, 2017    | Neutropenic Fever                                                 | Updated |
| Aug 2, 2017    | Multiple Sclerosis                                                | Updated |
| Sept 21, 2017  | Patient Eligibility                                               | New     |
| Oct 2, 2017    | Conditioning                                                      | Updated |
| Oct 10, 2017   | EBV/PTLD                                                          | Updated |
| Jan 23, 2018   | Chimerism                                                         | Updated |
| Feb 6, 2018    | Therapeutic Drug Monitoring                                       | Updated |
| Feb 27, 2018   | Hodgkin and Non-Hodgkin Lymphoma: Indications for Transplantation | Updated |
| May 8, 2018    | Relapse of Leukemia after Transplant                              | Updated |
| May 8, 2018    | Acute Lymphoblastic Leukemia                                      | Updated |
| May 8, 2018    | Graft Failure and Poor Graft Function                             | Updated |
| May 8, 2018    | ABO Incompatible Graft and Recipient                              | Updated |
| May 8, 2018    | Chronic Graft Versus Host Disease                                 | Updated |
| May 8, 2018    | Myelodysplastic Syndromes                                         | Updated |
| May 8, 2018    | Reproductive System Complications Post-transplant                 | Updated |



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| May 8, 2018    | Hepatic Complications and Viral Hepatitis                              | Updated |
| May 8, 2018    | BCR-ABL1-Negative Myeloproliferative Neoplasms                         | Updated |
| June 22, 2018  | Transplantation for Chronic Myelogenous Leukemia                       | Updated |
| June 22, 2018  | Scleroderma / Systemic Sclerosis                                       | Updated |
| June 22, 2018  | Transplantation for Germ Cell Tumours                                  | Updated |
| June 22, 2018  | Pretransplant Conditioning                                             | Updated |
| June 22, 2018  | Central Venous Catheter (CVC)-Related Complications                    | Updated |
| June 22, 2018  | Vaccination                                                            | Updated |
| June 22, 2018  | Umbilical Cord Blood Transplantation                                   | Updated |
| June 22, 2018  | Long-Term Follow-Up                                                    | New     |
| Sep 28, 2018   | Acute GVHD: Prevention and Treatment                                   | Updated |
| Oct 15, 2018   | Pneumocystis & Bacterial Prophylaxis                                   | Updated |
| Jan 15, 2019   | CAR T Cell Toxicity                                                    | New     |
| Jan. 24, 2019  | Vaccination                                                            | Edited  |
| April 18, 2019 | Stem Cell Mobilization (formerly Donor Management)                     | Updated |
| April 18, 2019 | Multiple Sclerosis                                                     | Updated |
| April 18, 2019 | CLL                                                                    | Updated |
| April 18, 2019 | Hemoglobinopathies                                                     | Updated |
| July 10, 2019  | Transplant AML                                                         | Updated |
| July 10, 2019  | Patient Eligibility                                                    | Updated |
| July 10, 2019  | Fungal Prophylaxis                                                     | Updated |
| July 10, 2019  | Infusion of Microbially-Contaminated or Non-Confirming Products        | Updated |
| July 26, 2019  | Infusion of Microbially-Contaminated or Non-Confirming Products        | Updated |
| July 31, 2019  | Hemoglobinopathies                                                     | Updated |
| July 31, 2019  | CMV, VZV, HSV, HHV6                                                    | Updated |
| July 31, 2019  | Hematopoietic Cell Transplantation for Severe Aplastic Anemia          | Updated |
| July 31, 2019  | Pretransplant Conditioning                                             | Updated |
| Aug. 1, 2019   | Scleroderma / Systemic Sclerosis                                       | Edited  |
| Aug. 1, 2019   | Hepatic Complications and Viral Hepatitis                              | Updated |
| Aug. 7, 2019   | CAR T Cell Toxicity                                                    | Updated |
| Oct. 7, 2019   | Chronic GVHD                                                           | Updated |
| Oct. 24, 2019  | Pretransplant Conditioning                                             | Updated |
| Dec, 16, 2019  | Donor Selection                                                        | Updated |
| Jan. 9, 2020   | Systemic Sclerosis                                                     | Updated |
| Jan. 23, 2020  | Nutritional Support                                                    | New     |
| Jan. 23, 2020  | Lymphoma                                                               | Updated |
| July 30, 2020  | CML                                                                    | Updated |
| July 30, 2020  | Acute GVHD                                                             | Updated |
| July 30, 2020  | Transfusions and Management of Cytopenias Early Post-HSCT              | New     |
| Sept. 23, 2020 | Vaccination                                                            | Updated |
| Sept. 29, 2020 | ALL                                                                    | Updated |
| Oct. 20, 2020  | Transfusions and Management of Cytopenias Early Post-HSCT              | Updated |
| Dec. 2, 2020   | Symptomatic Myeloma and AL Amyloidosis, formerly Plasma Cell Disorders | Updated |
| Dec. 2, 2020   | Pneumocystis and Bacterial Prophylaxis                                 | Updated |

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| Dec. 2, 2020   | Acute GVHD                                                                           | Updated |
| Jan. 8, 2021   | EBV/PTLD                                                                             | Updated |
| Jan. 8, 2021   | CAR T Cell Toxicity                                                                  | Updated |
| Jan. 12, 2021  | Umbilical Cord Blood Transplant                                                      | Updated |
| Jan. 18, 2021  | Scleroderma/ Systemic Sclerosis (SSc)                                                | Updated |
| Jan. 19, 2021  | Multiple Sclerosis                                                                   | Updated |
| Feb. 3, 2021   | CLL                                                                                  | Updated |
| Mar. 16, 2021  | DLI/2 <sup>nd</sup> HCT Relapse, formerly Relapse                                    | Updated |
| Mar. 23, 2021  | Donor Selection                                                                      | Updated |
| Mar. 30, 2021  | Lymphoma                                                                             | Updated |
| Apr. 12, 2021  | Stem Cell Mobilization                                                               | Updated |
| Apr. 13, 2021  | CVAD Complications, formerly CVC-Related Complications                               | Updated |
| April 21, 2021 | ALL                                                                                  | Updated |
| May 27, 2021   | Vaccination                                                                          | Updated |
| May 27, 2021   | Fungal Prophylaxis                                                                   | Updated |
| May 27, 2021   | Hepatic Complications and Viral Hepatitis                                            | Updated |
| May 27, 2021   | Hemoglobinopathies                                                                   | Updated |
| May 27, 2021   | Chimerism                                                                            | Updated |
| June 8, 2021   | CMV, VZV, HHV6                                                                       | Updated |
| June 15, 2021  | AML                                                                                  | Updated |
| July 5, 2021   | Reproductive System Complications                                                    | Removed |
| July 7, 2021   | Stem Cell Mobilization                                                               | Edited  |
| July 7, 2021   | Umbilical Cord Blood Transplant                                                      | Edited  |
| July 12, 2021  | Myelodysplastic Syndromes (MDS)                                                      | Updated |
| July 13, 2021  | BCR-ABK-Negative Myeloproliferative Neoplasms                                        | Updated |
| Aug 10, 2021   | Distribution of Microbially-Contaminated or Non-Conforming Cellular Therapy Products | Updated |
| Aug 24, 2021   | Transplantation Eligibility Assessment: Patient Factors                              | Updated |
| Sept 14, 2021  | Long-Term Follow up                                                                  | Updated |
| Oct 12, 2021   | Graft vs Host Disease                                                                | New     |
| Nov 2, 2021    | Conditioning (formally Pretransplant Conditioning)                                   | Updated |
| Nov 4, 2021    | Acute GVHD                                                                           | Removed |
| Nov 4, 2021    | Chronic GVHD                                                                         | Removed |
| Nov 4, 2021    | Therapeutic Drug Monitoring                                                          | Removed |
| Dec. 8, 2021   | Germ Cell Tumours                                                                    | Updated |
| Dec. 14, 2021  | Fever                                                                                | Updated |
| Dec. 17, 2021  | Vaccination                                                                          | Edited  |
| Feb. 8, 2022   | Donor Selection                                                                      | Updated |
| Mar. 2 2022    | Vaccination                                                                          | Edited  |
| Apr. 26, 2022  | Scleroderma/ Systemic Sclerosis (SSc)                                                | Updated |
| Apr. 26, 2022  | Vaccination                                                                          | Updated |
| May 3, 2022    | CML                                                                                  | Updated |
| May 25, 2022   | Scleroderma/ Systemic Sclerosis (SSc)                                                | Edited  |
| May 25, 2022   | Donor Selection                                                                      | Edited  |
| July 26, 2022  | Fungal Prophylaxis                                                                   | Updated |

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| Sept. 13, 2022 | Nutritional Support                                                  | Updated |
| Oct. 4, 2022   | Pneumocystis and Bacterial Prophylaxis                               | Updated |
| Oct. 11, 2022  | ALL                                                                  | Updated |
| Oct. 25, 2022  | Symptomatic Myeloma and AL Amyloidosis                               | Updated |
| Dec. 6, 2022   | GVHD                                                                 | Updated |
| Dec. 12, 2022  | Scleroderma/ Systemic Sclerosis (SSc)                                | Edited  |
| Jan. 24, 2023  | CAR T Cell Toxicity                                                  | Updated |
| Jan. 31, 2023  | Scleroderma/ Systemic Sclerosis (SSc)                                | Edited  |
| Jan. 31, 2023  | Conditioning                                                         | Updated |
| Feb. 1, 2023   | CLL                                                                  | Updated |
| Mar. 21, 2023  | Aplastic Anemia                                                      | Updated |
| May 16, 2023   | GVHD                                                                 | Updated |
| May 23, 2023   | ABO Incompatible Graft and Recipient                                 | Updated |
| June 27, 2023  | Transfusions and Management of Cytopenias Early Post-HSCT            | Updated |
| Aug 3, 2023    | Management of Transfusion and Cytopenias Post-HSCT                   | Edited  |
| Aug 22, 2023   | Transplantation Eligibility Assessment: Patient Factors              | Updated |
| Sept. 26, 2023 | DLI/2 <sup>nd</sup> HCT                                              | Updated |
| Oct. 3, 2023   | Hemoglobinopathies                                                   | Updated |
| Oct. 19, 2023  | Stem cell Mobilization                                               | Updated |
| Nov. 7, 2023   | Germ Cell Tumours                                                    | Updated |
| Nov. 21, 2023  | Microbially-Contaminated or Non-Conforming Cellular Therapy Products | Updated |
| Nov. 22, 2023  | Conditioning                                                         | Edited  |
| Dec. 5, 2023   | Hepatic Complications and Viral Hepatitis                            | Updated |
| Jan. 9, 2024   | CMV, VZV, HHV6                                                       | Updated |
| Jan. 30, 2024  | Transplantation Eligibility Assessment: Patient Factors              | Updated |
| Feb. 6, 2024   | Multiple Sclerosis                                                   | Updated |
| Feb. 13, 2024  | AML                                                                  | Updated |
| Mar. 5, 2024   | Vaccination                                                          | Updated |
| Mar. 5, 2024   | Long-Term Follow Up                                                  | Updated |
| Apr. 2, 2024   | EBV/PTLD                                                             | Updated |
| May 7, 2024    | CVAD Complications                                                   | Updated |
| May 28, 2024   | Lymphoma                                                             | Updated |
| June 18, 2024  | Graft Failure and Poor Graft Function                                | Removed |
| June 18, 2024  | Chimerism                                                            | Removed |
| June 18, 2024  | Graft Failure, Poor Graph Function, Chimerism                        | New     |
| June 25, 2024  | Hemoglobinopathies                                                   | Updated |
| July 15, 2024  | Scleroderma/ Systemic Sclerosis (SSc)                                | Edited  |
| July 23, 2024  | Multiple Sclerosis                                                   | Updated |
| Aug. 6, 2024   | Scleroderma/ Systemic Sclerosis (SSc)                                | Edited  |
| Sept. 3, 2024  | CML                                                                  | Updated |
| Sept. 17, 2024 | Symptomatic Myeloma and AL Amyloidosis                               | Updated |
| Sept. 24, 2024 | Symptomatic Myeloma and AL Amyloidosis                               | Edited  |
| Oct. 1, 2024   | Donor Selection                                                      | Updated |
| Oct. 1, 2024   | Umbilical Cord Blood Transplantation                                 | Updated |

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| Oct. 15, 2024 | Nutrition Support                                                   | Updated |
| Nov. 5, 2024  | Pneumocystis and Bacterial Prophylaxis                              | Updated |
| Nov. 12, 2024 | Fever                                                               | Updated |
| Dec. 13, 2024 | Stem Cell Mobilization                                              | Edited  |
| Jan. 21, 2025 | Autologous Hematopoietic Cell Transplant in Systemic Sclerosis(SSc) | Updated |
| Feb. 26, 2025 | Germ Cell Tumours                                                   | Edited  |
| Apr. 15, 2025 | ALL                                                                 | Updated |
| May 5, 2025   | Conditioning                                                        | Edited  |
| May 6, 2025   | Fungal Prophylaxis                                                  | Updated |
| June 5, 2025  | Donor Selection                                                     | Edited  |
| June 5, 2025  | GVHD                                                                | Updated |
| Sept 9, 2025  | Myelodysplastic Syndromes (MDS)                                     | Updated |
| Oct. 21, 2025 | Stem cell Mobilization                                              | Updated |
| Oct. 28, 2025 | Aplastic Anemia                                                     | Updated |
| Nov. 25, 2025 | Germ Cell Tumours                                                   | Updated |
| Jan 6, 2026   | Transfusions and Management of Cytopenias Early Post-HSCT           | Updated |
| Jan 13, 2026  | Hepatic Complications and Viral Hepatitis                           | Updated |
| Jan 20, 2026  | Conditioning                                                        | Updated |
| Jan 27, 2026  | CAR T Cell Toxicity                                                 | Updated |
| Feb 10, 2026  | CMV, VZV, HSV, HHV6                                                 | Updated |
| Feb 24, 2026  | Donor Selection                                                     | Edited  |
| Mar 06, 2026  | AML                                                                 | Updated |
| Apr, 2026     | Vaccination                                                         | Updated |
| Apr 21, 2026  | Aplastic Anemia                                                     | Edited  |
| May 12, 2026  | ABO Incompatible Graft and Recipient                                | Updated |