



# Hematopoietic Stem Cell Transplantation

## Clinical Guidelines

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# Guideline Application

For medical necessity clinical coverage criteria for Medicare Advantage plans, refer first to the Medicare Coverage Database for NCDs and LCDs/LCAs, then the Medicare Benefit Policy Coverage Manual.

## Introduction

Hematopoietic stem cell transplants, including peripheral blood, bone marrow, and cord blood transplants are used most often to treat cancers affecting the blood or immune system. There are two main types of stem cell transplant: autologous and allogeneic. Autologous stem cells come from the person who will be receiving the transplant and are mainly used to treat leukemias, lymphomas, and multiple myeloma as well as other cancers such as testicular cancer and neuroblastoma. Autologous stem cell transplants are also used to treat certain childhood cancers. Allogeneic stem cells come from another individual. They can be from a matched related or unrelated donor or a donor without a complete match. Allogeneic stem cells are most commonly used to treat leukemias, lymphomas or non-malignant inherited disorders. An allogeneic transplant provides the advantage of a graft vs. cancer effect but occurs with the potential risk of graft vs. host disease. The need to balance these two outcomes makes this a more complicated procedure.

The purpose of this guideline is to identify the indications and contraindications for hematopoietic stem cell transplant as well as provide helpful reference tools to better understand a request for transplant.

## General Information

- “Back-up” autologous harvesting for patients in complete remission (CR) with no evidence of marrow involvement by malignancy is appropriate. For example, bone marrow or peripheral blood progenitor cell harvesting is appropriate for patients with multiple myeloma in CR and who might be transplanted in the future. Consult benefit document.
- While the development of chimeric antigen receptor T-cell (CAR T) therapy has introduced a new field of therapeutic possibilities for patients with certain hematological malignancies, hematopoietic stem cell transplantation remains a cornerstone of care.
- Donor lymphocyte infusion (DLI) following allogeneic stem cell transplant is appropriate for incomplete chimerism and/or disease relapse in the setting of incomplete chimerism. This is not a second stem cell transplant. There is not a standardized approach to the use of DLI and can come at various times following the initial transplant (Castagna et al., 2016).
- Repeat stem cell transplant is appropriate for primary and secondary failure to engraft and disease relapse.
- Primary failure is the failure to reach three consecutive days with a neutrophil count (absolute neutrophil count/ANC)  $> 500 \mu\text{l}$  ( $0.5 \times 10^9/\text{liter}$ ) after SCT, while secondary failure is associated with a successful SCT graft where neutrophils increase to  $> 500 \mu\text{l}$  ( $0.5 \times 10^9/\text{liter}$ ) for at least three consecutive days and subsequently decrease to a lower level until additional treatment is given to obtain engraftment. (There can be a loss of an allogeneic graft with normal blood cell counts due to autologous reconstitution. This can be confirmed with chimerism studies).
- Stem cell boost is a hematopoietic stem cell infusion (HSCI) provided to a transplant recipient to assist with hematopoietic recovery or declining donor chimerism. It is not preceded by a preparative regimen and is not considered a new transplant event. Stem cell boost is a non-standardized term and has been used interchangeably with terms such as reinfusion, support, and rescue. For the purposes of this guideline, we endorse use of the term “boost” based on the recommendation of the task force set up by the American Society for Blood and Marrow Transplantation in collaboration with National Marrow Donor Program (LeMaistre et al., 2013) and the existence of a CPT code for the term boost (CPT 38243).
- Autologous stem cell transplant with or without a second autologous transplant (tandem transplant) is considered a standard of care for the treatment of multiple myeloma although controversy does exist

particularly in the era of newer and more effective chemotherapy agents such as bortezomib, lenalidomide and thalidomide (Blade, 2010; Harousseau & Moreau, 2009; Bashey, 2008; Kumar, 2009). As the primary and salvage treatment for multiple myeloma has become increasingly successful in recent years, it is likely that, going forward, multiple factors will need to be considered prior to making decisions regarding the use of transplantation procedures, e.g., risk stratification, age, comorbidities, etc. and that the role of transplantation may decrease for certain subgroups.

- During a tandem transplant, a patient receives two sequential courses of high-dose chemotherapy with stem cell transplant. Peripheral blood hematopoietic stem cells (HSCs) are collected either during recovery of a cycle of induction chemotherapy or after filgrastim mobilization. The patient receives a second preparative regimen, along with hematopoietic progenitor cells (HPCs) collected during the initial mobilization. Both transplantations are planned and typically are performed a few weeks to a few months apart (LeMaistre et al., 2013).
- Tandem stem cell transplants require review by the Medical Director **except** for the following conditions:
  - Multiple myeloma
  - Testicular and other germ cell tumors
  - Neuroblastoma
  - Pediatric brain tumors
  - Other conditions as part of an IRB-approved clinical trial
- Third stem cell transplants require Medical Director review.
  - If part of a sequence of high-dose chemotherapy followed by rescue stem cell infusion as is the case with some neuroblastoma, medulloblastoma and testicular germ cell tumor protocols, the entire course may be approved initially.
- The stem cell transplant expert panels have confirmed that the treatment of any pediatric patient under a Children's Oncology Group (COG) protocol should be considered Standard of Care.
- Patients who have undergone stem cell transplant have altered immune systems post-transplant. In the case of allogeneic stem cell transplant, the immune system may never fully recover. These patients have unique care needs in the post-transplant period and will require lifelong follow-up and management (Optum Expert Panel, 2015).
- To improve outcomes of blood and marrow transplantation, the use of maintenance therapy has become standard over the past few years. Maintenance therapy is considered an important component of the transplant event and therefore is covered if supported by adequate clinical evidence.
- Minimal/Measurable Residual Disease (MRD) is a measure of persistent disease which has emerged as a powerful tool in determining prognosis and informing treatment decisions for patients with hematologic malignancies. MRD detection is measured using flow cytometry, real-time quantitative polymerase chain reaction (RQ-PCR) or next-generation sequencing assays (Short, 2017). MRD is part of the standard evaluation of response to HSCT for several underlying disorders and is considered medically necessary.

## Veno-Occlusive Disease (VOD)

Traditionally the treatment of veno-occlusive disease (VOD) has been supportive and the outcomes poor. In March 2016, FDA gave approval to the new drug defibrotide for the treatment of active VOD. At the present time there is not an approved indication for its use in a prophylactic manner which is commonly done overseas in Europe.

- Defibrotide is covered for the treatment of adult and pediatric patients with active hepatic VOD with renal or pulmonary dysfunction following hematopoietic stem cell transplant.
- Defibrotide is not covered for the prevention of VOD

## Umbilical Cord Blood Stem Cell Transplantation

- Single unit umbilical cord blood stem cell transplants are standard of care for children in many programs. Children who receive a single cord blood unit may experience prolonged time to engraftment and other post-

transplant complications; therefore, a calculation of  $2.5 \times 10^7$  nucleated cells per kilogram may improve response (de Lima, 2006).

- Umbilical cord blood and haploidentical donor cells are appropriate stem cell sources (Brunstein et al., 2007; Klingebiel et al., 2010) and can be used at the discretion of the treating team.

Delayed engraftment is a significant limitation of umbilical cord blood transplantation (UCBT) due in part to low cell doses and a defect in the cord blood cells' ability to home to the bone marrow (Popat et al., 2015). In April 2023, the FDA approved Omidubicol only (Omisirge<sup>®</sup>) for use in adults and pediatric patients 12 years and older with hematologic malignancies who are planned for umbilical cord blood transplantation following myeloablative conditioning to reduce time to neutrophil engraftment and the incidence of infection. Omidubicol is an ex vivo expanded hematopoietic progenitor cell and nonexpanded myeloid and lymphoid cell product derived from a single umbilical cord blood unit. Omidubicol is prepared as a cell suspension for intravenous infusion. A single dose consists of a Cultured Fraction (CF): a minimum of  $8.0 \times 10^8$  total viable cells of which a minimum of 8.7% is CD34+ cells and a minimum of  $9.2 \times 10^7$  CD34+ cells, and a Non-cultured Fraction (NF): a minimum of  $4.0 \times 10^8$  total viable cells with a minimum of  $2.4 \times 10^7$  CD3+ cell. Horwitz et al. (2021) reported the outcomes of a phase 3 trial evaluating the efficacy of Omidubicol vs standard UCBT in 125 patients aged 13 to 65 years with hematologic malignancies. Patients were randomly assigned to Omidubicol or standard UCBT. All patients received myeloablative conditioning and prophylaxis with a calcineurin inhibitor and mycophenolate mofetil for graft-versus-host disease (GVHD). The primary endpoint was time to neutrophil engraftment. In the Omidubicol arm, median time to neutrophil engraftment was 12 days (95% CI, 10-14 days) and 22 days (95% CI, 19-25 days) for the control arm. The cumulative incidence of engraftment was 96% for patients receiving Omidubicol and 89% for patients receiving standard UCBT. Additionally, the Omidubicol arm experienced faster platelet recovery (55% vs 35% recovery by 42 days;  $P = .028$ ), had lower incidence of first grade 2 to 3 bacterial or invasive fungal infection (37% vs 57%;  $P = .027$ ) and spent more time out of the hospital during the first 100 days after transplant (median, 61 vs 48 days;  $P = .005$ ) than controls. Of note, the logistical complexity of producing Omidubicol led to a median 2-week delay in time from randomization to transplantation. The authors report this delay did not result in increased risk of pretransplant relapse in the 52 patients transplanted with Omidubicol. It should also be noted that Lin et al. (2023) reported donor-derived myeloid neoplasm were an adverse event of special interest. Donor-derived myelodysplastic syndrome (MDS) occurred in one patient receiving Omidubicol in the fourth year post-transplant while two patients developed post-transplant lymphoproliferative disorder (PTLD) in the second year post-transplant.

**Omidubicol** only (Omisirge<sup>®</sup>) may be considered medically necessary when all of the following are met:

- Patient is 12 years of age or older
- Has been diagnosed with a hematologic malignancy
- Is a candidate for myeloablative allogeneic HSCT
- Has no readily available matched sibling or matched unrelated donor
- The medication is being requested to reduce the time to neutrophil recovery and incidence of infection
- Following a comprehensive evaluation of the clinical condition it is felt this is the best choice of cell source for this particular patient

# Indications

## Leukemias

Leukemia more often is a cancer affecting white blood cells. Leukemia occurs most often in adults older than 55, however it is also the most common cancer in children younger than 15 years (National Cancer Institute, 2024). There are four main types of leukemia: acute lymphoblastic leukemia (ALL), acute myeloid leukemia (AML), chronic lymphocytic leukemia (CLL), and chronic myeloid leukemia (CML). The rate at which the disease progresses and how leukemic cells replace the normal blood and marrow cells differs with each type of leukemia. There also exists several uncommon or rare forms of leukemia including, but not limited to, hairy cell leukemia, B-cell prolymphocytic leukemia, and T-cell prolymphocytic leukemia (Leukemia and Lymphoma Society, 2024). There is a lack of data supporting autologous stem cell transplantation for CLL; however, the availability of new agents such as idelalisib and ibrutinib, which are highly effective against this condition will likely change how stem cell transplantation is used in this disease. A history of prior treatment should be obtained with every transplant request.

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary for the following types of leukemia:

- ALL (McNeer et al., 2019)
- Intermediate and high-risk AML ([See Appendix A](#)) (Dohner et al., 2017) including but not limited to:
  - CR1 with poor-risk cytogenetic or markers
  - Following relapse
  - CR2 and beyond
- CLL
- CML
- Prolymphocytic leukemia (Kalaycio et al., 2010; Krishnan et al., 2010)

**Allogeneic** hematopoietic stem cell transplantation is considered unproven and not medically necessary for all other leukemia types.

**Autologous** hematopoietic stem cell transplantation may be considered medically necessary for the following types of leukemia:

- Prolymphocytic leukemia (Kalaycio et al., 2010; Krishnan et al., 2010)
- ALL (McNeer et al., 2019)
  - May be indicated in certain adults when there is no suitable allogeneic donor. Refer to Medical Director.
  - In pediatric patients certain cytogenetic features including hypodiploid (<44 chromosomes and iAMP21) are associated with high-risk disease and may influence the decision to transplant in first complete response (CR1). Refer to Medical Director.
- AML (Dohner et al., 2017)
  - May be indicated in certain adults when there is no suitable allogeneic donor. Refer to the Medical Director.

**Autologous** hematopoietic stem cell transplantation is considered unproven and not medically necessary for all other leukemia types.

## Myelodysplastic Syndromes & Mixed Myelodysplastic/Myeloproliferative Neoplasms

Myelodysplastic syndromes (MDS) are a collection of myeloid malignancies distinguished by one or more blood cytopenias. In the United States, MDS is diagnosed in a little over 10,000 cases annually, translating to roughly 4.4 cases per 100,000 individuals (National Cancer Institute, 2024). The World Health Organization (WHO) classifies CMML as a myelodysplastic/myeloproliferative neoplasm.

The sole curative therapy for myelodysplastic syndrome (MDS) is allogeneic hematopoietic cell transplantation (HCT). Recent prospective trials have confirmed that allogeneic HSCT confers survival benefits in patients with advanced or high-risk MDS compared with nontransplantation approaches, and the use of HSCT is increasing in older patients with good performance status. However, patients with high-risk cytogenetic or molecular mutations remain at high risk for relapse. It is unknown whether administration of novel therapies before or after transplantation may decrease the risk of disease relapse in selected populations. Ongoing and future studies will investigate revised approaches to disease risk stratification, patient selection, and post-transplantation approaches to optimize allogeneic HCT outcomes for patients with MDS (DeFilipp et al., 2023).

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary for the following:

- Myelodysplastic syndromes (MDS)
- Juvenile myelomonocytic leukemia/juvenile chronic myelogenous leukemia (JMML/JCML)
- Chronic myelomonocytic leukemia (CMML) (Swerdlow et al., 2017)

**Autologous** hematopoietic stem cell transplantation is considered unproven and not medically necessary for the following:

- Myelodysplastic syndromes (MDS)
- Juvenile myelomonocytic leukemia (JMML/JCML)
- Chronic myelomonocytic leukemia (CMML) (Swerdlow et al., 2017)

**Note:** On March 6, 2024, CMS issued a [final decision](#) under National Coverage Determination (NCD) 110.23 to expand Medicare coverage for allogeneic hematopoietic stem cell transplant using bone marrow, peripheral blood or umbilical cord blood stem cell products for Medicare patients with MDS dependent on prognostic risk scores.

[See Appendix B](#)

## Myeloproliferative Disorders

Myelofibrosis is a rare disorder that impairs normal blood cell formation, resulting in splenomegaly, anemia, and fibrosis. Myelofibrosis develops when a mutation occurs in a hematopoietic stem cell, the genetic material in the cell starts to proliferate and disrupts normal blood synthesis, leading to the development of myelofibrosis. The identification of adverse karyotypes is evolving. New clinical molecular scoring systems may become useful in determining post-transplant prognosis. Myelofibrosis occurs in approximately 0.25 cases per 100,000 (Manning, 2022).

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary in the following:

- Primary myelofibrosis and related conditions (e.g., polycythemia vera) in patients with Intermediate-2 or High-Risk score using the Dynamic International Prognostic Scoring System Plus (DIPSS) (Gagelman et al., 2019). [See Appendix C](#)
- Secondary myelofibrosis
  - Allogeneic transplant evaluation approved for patients with polycythemia vera or essential thrombocythemia.

**Autologous** hematopoietic stem cell transplantation is considered unproven and not medically necessary for the following:

- Primary myelofibrosis and related conditions (Gagelman et al., 2019)
- Secondary myelofibrosis



## Brain Tumors

Brain tumors comprise 85% to 90% of primary central nervous system (CNS) tumors. The combined incidence of brain and other CNS tumors in the United States is 6.2 per 100,000 people per year, with a mortality rate of 4.4 deaths per 100,000 people per year (National Cancer Institute, 2024). The main types of brain tumors include anaplastic astrocytoma, brain stem glioma, ependymoma, germinoma, glioblastoma multiforme, medulloblastoma, oligodendroglioma, pineoblastoma, embryonal tumors with multi-layered rosettes (formerly known as primitive neuroectodermal tumor).

**Allogeneic** hematopoietic stem cell transplantation is considered unproven and not medically necessary for brain tumors including but not limited to the following:

- Anaplastic astrocytoma
- Brain stem glioma
- Ependymoma
- Germinoma
- Glioblastoma multiforme (GBM)
- Medulloblastoma
- Oligodendroglioma
- Pineoblastoma
- Embryonal tumors with multi-layered rosettes (ETMR)

**Autologous** hematopoietic stem cell transplantation may be considered medically necessary for the following brain tumors:

- Medulloblastoma
- Oligodendroglioma
- Pineoblastoma
- Embryonal tumors with multi-layered rosettes (ETMR)
- Glioblastoma multiforme (GBM)
  - May be considered in infants. Refer to Medical Director.

**Autologous** hematopoietic stem cell transplantation is considered unproven and not medically necessary for all other brain tumors including but not limited to the following:

- Anaplastic astrocytoma
- Brain stem glioma
- Ependymoma
- Germinoma

## Germ Cell Tumors

Germ cell tumors arise from primordial germ cells. Testicular and sacrococcygeal GCTs arising during early childhood characteristically have deletions at chromosome arms 1p and 6q and gains at 1q, and they lack the isochromosome 12p that is highly characteristic of malignant GCTs of adults. Testicular GCTs also may demonstrate loss of imprinting. Ovarian GCTs from older females characteristically have deletions at 1p and gains at 1q and 21. Dysregulation of microRNAs have been linked to GCTs. Because GCTs may contain benign and mixed malignant elements in different areas of the tumor, extensive sectioning is essential to confirm the correct diagnosis. The many histologically distinct subtypes of GCTs include teratoma (mature and immature), endodermal sinus tumor, and embryonal carcinoma. Teratomas occur in many locations, presenting as masses. They are not associated with elevated markers unless malignancy is present. The sacrococcygeal region is the most common site for teratomas. Sacrococcygeal teratomas occur most commonly in infants and may be diagnosed in utero or at birth, with most found in girls. The rate of malignancy in this location varies, ranging from <10% in children younger than 2 months to >50% in children older than 4 months. Germinomas occur intracranially, in the mediastinum, and in the gonads. In the ovary, they are called dysgerminomas, and in the testis, they are called seminomas. They usually are tumor-marker-negative masses despite being malignant. Endodermal sinus or yolk sac tumor and choriocarcinoma appear highly

malignant by histologic criteria. Both occur at gonadal and extragonadal sites. Embryonal carcinoma most often occurs in the testes. Choriocarcinoma and embryonal carcinoma rarely occur in the pure form and are usually found as part of a mixed malignant GCT. Complete surgical excision of the tumor usually is indicated, except for patients with intracranial tumors, for whom the primary therapy consists of radiation therapy and chemotherapy. High-dose chemotherapy followed by autologous stem cell rescue may also be considered (Herzog & Huh, 2025).

**Allogeneic** hematopoietic stem cell transplantation is considered unproven and not medically necessary for germ cell tumors including but not limited to the following:

- Testicular germ cell tumor
- Extragonadal germ cell tumor
- Seminoma
- Choriocarcinoma
- Embryonal carcinoma
- Mixed germ cell tumors
- Teratoma
- Yolk-sac tumor (endodermal sinus tumor)
- Germ cell tumor of the ovary

**Autologous** and tandem autologous hematopoietic stem cell transplantation may be considered medically necessary for the following:

- Testicular germ cell tumor
- Extragonadal germ cell tumor
- Seminoma
- Choriocarcinoma
- Embryonal carcinoma
- Mixed germ cell tumors
- Teratoma
- Yolk-sac tumor (endodermal sinus tumor)
- Germ cell tumor of the ovary

## Multiple Myeloma

Multiple myeloma (MM) is a plasma cell malignancy in which monoclonal plasma cells proliferate in bone marrow, resulting in an overabundance of monoclonal paraprotein (M protein), destruction of bone, and displacement of other hematopoietic cell lines. MM is part of a spectrum of diseases including monoclonal gammopathy of undetermined significance (MGUS) and plasma cell leukemia. Nearly all patients with multiple myeloma evolve from MGUS. Since MGUS is asymptomatic, over 50% of individuals who are diagnosed with MGUS have had the condition for over 10 years prior to the clinical diagnosis. MGUS progresses to multiple myeloma or related malignancy at a rate of 1% per year. In some patients, an intermediate asymptomatic but more advanced pre-malignant stage referred to as smoldering multiple myeloma (SMM) can be recognized clinically. SMM progresses to multiple myeloma at a rate of approximately 10% per year over the first 5 years following diagnosis, 3% per year over the next 5 years, and 1.5% per year thereafter. This rate of progression is influenced by the underlying cytogenetic type of disease; patients with t(4;14) translocation, del(17p), and gain(1q) are at a higher risk of progression from MGUS or SMM to multiple myeloma (Rajkumar, 2022).

Allogeneic stem cell transplant for multiple myeloma is controversial either as a single allogeneic transplant as initial therapy with curative intent or as the second stage of a planned tandem transplant preceded by an autologous transplant. The following recommendations are consistent with the evolving practice and recognize the expertise of treating physicians within network programs. These recommendations may change as additional experience is gained with the newer disease modifying agents for the treatment of myeloma and as more experience is gained with reduced intensity allogeneic stem cell transplant for this disease.

**Note:** Refer all requests for allogeneic stem cell transplant in multiple myeloma to the Medical Director.

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary for multiple myeloma in the following circumstances:

- Initial therapy in newly diagnosed patients with high-risk disease and in otherwise good health
  - High risk myeloma has been defined by the International Myeloma Working Group (IMWG) based on cytogenetics [presence of at least one of the following: del(17p), t(4;14) or t(14;16) determined by FISH]
  - The Mayo Clinic classification adds hypoploidy and t(14;20) to the IMWG definition. Regardless of the source of definition, the requestor should present evidence of sufficient factors that cause the case to be considered high risk.
- Early relapse (less than 24 months) after primary therapy that included an autologous stem cell transplant or with high-risk features (i.e., cytogenetics, extramedullary disease, plasma cell leukemia, or high lactate dehydrogenase) if they respond favorably to salvage therapy. (Giralt, 2015)
- Reduced intensity matched related donor (MRD) and matched unrelated donor (MUD) allogeneic SCT as the second transplant of a planned tandem transplant. (Bruno, 2009; Rotta, 2009)

**Autologous** hematopoietic stem cell transplantation may be considered medically necessary in multiple myeloma in the following circumstances:

- Single autologous
- Tandem (autologous followed by autologous)
- Tandem (autologous followed by allogeneic) in early relapse [less than 24 months] after primary therapy that included an autologous SCT or with high-risk features (i.e., cytogenetics, extramedullary disease, plasma cell leukemia, or high lactate dehydrogenase) if they respond favorably to salvage therapy (Giralt, 2015).

## Plasma Cell Disorders

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary in the following:

- Waldenstrom macroglobulinemia
- May be appropriate in a clinical trial for AL- Amyloidosis

**Allogeneic** hematopoietic stem cell transplantation is unproven and not medically necessary in all other circumstances including but not limited to:

- Monoclonal gammopathy of uncertain significance (MGUS)
- Polyneuropathy organomegaly endocrinopathy monoclonal gammopathy skin defects syndrome (POEMS)
- Solitary plasmacytoma

**Autologous** hematopoietic stem cell transplantation may be considered medically necessary for the following:

- AL-Amyloidosis
- POEMS (D'Souza et al., 2012; Ji et al., 2012; Li et al., 2013)
- Waldenstrom macroglobulinemia

**Autologous** hematopoietic stem cell transplantation is unproven and not medically necessary in all other circumstances including but not limited to:

- MGUS
- Solitary plasmacytoma

## Monoclonal Gammopathy of Renal Significance (MGRS)

Monoclonal gammopathy of renal significance (MGRS) is a clonal proliferative disorder that produces a nephrotoxic monoclonal immunoglobulin and does not meet previously defined hematological criteria for treatment of a specific malignancy. Monoclonal immunoglobulin-related diseases show higher rates of recurrence after kidney transplantation (often > 80%) than their non-monoclonal counterparts. They are poorly responsive to conventional immunosuppression (Leung et al., 2019). Targeting the underlying B-cell clone with chemotherapy, although it is not a malignant clone per se, is the only available treatment option for MGRS. High-dose melphalan (HDM) supported by autologous SCT may be a therapeutic option in some patients (Ferland et al., 2013).

**Autologous** hematopoietic stem cell transplantation may be considered medically necessary for MGRS in patients who meet the following criteria:

- Have failed chemotherapy targeting the underlying B-cell clone
- AND
- Have sufficient renal function to tolerate high-dose chemotherapy

**Allogeneic** hematopoietic stem cell transplantation is unproven and not medically necessary in MGRS.

## Hodgkin Lymphoma

Hodgkin lymphoma (HL) is an uncommon malignancy of B-cell origin. Most patients are diagnosed between ages 15 and 30 years, followed by another peak in adults  $\geq 55$  years. In 2024, an estimated 8570 people will be diagnosed with HL in the United States and 910 people will die from the disease (American Cancer Society, 2024). The World Health Organization (WHO) classification divides HL into two main types: classic Hodgkin lymphoma (CHL) and nodular lymphocyte-predominant Hodgkin lymphoma (NLPHL). In Western countries, CHL accounts for 95% and NLPHL accounts for 5% of all HL. While the WHO has maintained the term NLPHL, the International Consensus Classification (ICC) has now replaced the term NLPHL with the term nodular lymphocyte predominant B-cell lymphoma (NLPBL) based on biological and clinical differences with CHL. The past few decades have seen significant progress in the management of HL. The advent of more effective treatment options has improved the 5-year survival rates, which have been unmatched in any other cancer over the past 4 decades. It is among the most curable of malignancies with modern treatments, and newly diagnosed HL has a very high likelihood of being cured with appropriate management. In fact, cure rates for HL have increased so markedly that overriding treatment considerations often relate to long-term toxicity. Clinical trials still emphasize improvement in cure rates for patients with advanced disease, but the potential long-term effects of treatment remain an important consideration (Campo et al., 2022).

**Autologous** hematopoietic stem cell transplantation is considered medically necessary for Hodgkin lymphoma.

**Allogeneic** hematopoietic stem cell transplantation is considered medically necessary for Hodgkin lymphoma.

## Non-Hodgkin Lymphoma (NHL)

Non-Hodgkin lymphoma (NHL) is a heterogeneous group of malignancies of the lymphoreticular system subdivided into indolent, aggressive, and highly aggressive disease. The therapeutic regimen varies with specific lymphoma subtype and pathologic stage. There are approximately 60 different NHL subtypes. NHL is the seventh most common neoplasm in the U.S. with >80,000 new cases annually. Incidence increases with age with the majority of patients older than 60 years. In the U.S. and Europe, diffuse large B-cell lymphoma (DLBCL) is the most common subtype (30% of the cases) followed by follicular lymphoma (FL) comprising 25% of cases. In patients with HIV, NHL is the most common tumor followed by Kaposi sarcoma. The presence of B symptoms such as unexplained weight loss, fever, fatigue, and night sweats are seen typically in aggressive lymphomas. Aggressive lymphomas have acute or subacute presentation with increasing size of the mass and B symptoms. Indolent lymphomas have a more chronic course, with asymptomatic lymphadenopathy and/or slowly progressive cytopenias. Initial laboratory evaluation may

be entirely normal. Elevated serum LDH may be seen in aggressive lymphoma or in indolent lymphoma with high disease burden. In cases of highly aggressive NHL (e.g., Burkitt lymphoma), spontaneous tumor lysis syndrome (TLS) may be seen, characterized by hyperkalemia, hyperuricemia, hypocalcemia, hyperphosphatemia, and acidosis. TLS can be life threatening and is considered a medical emergency (Alaggio et al., 2022).

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary in the following:

- Small B-cell lymphocytic lymphoma
- Follicular lymphoma (Epperala et al., 2018; Oliansky et al., 2010; Sureda et al., 2018)
- Lymphoplasmacytoid lymphoma/immunocytoma
- Marginal zone lymphoma (mucosa-associated lymphoid tissue, splenic, nodal)
- Burkitt lymphoma
- Diffuse, large cell lymphoma (mediastinal large cell, primary effusion) (Oliansky et al. 2011)
- Mantle cell lymphoma
- Precursor B-cell leukemia/lymphoma
- T-cell lymphoma

**Autologous** hematopoietic stem cell transplantation may be considered medically necessary for the following:

- Follicular lymphoma (Epperala et al., 2018; Oliansky et al., 2010; Sureda et al., 2018)
- Lymphoplasmacytoid lymphoma/immunocytoma
- Marginal zone lymphoma (mucosa-associated lymphoid tissue, splenic, nodal)
- Burkitt lymphoma
- Diffuse, large cell lymphoma (mediastinal large cell, primary effusion) (Oliansky et al. 2011)
- Mantle cell lymphoma
- Precursor B-cell leukemia/lymphoma
- T-cell lymphoma

For autologous hematopoietic stem cell transplants, tumors must be chemosensitive which is defined as a complete or partial response based on the Cheson criteria. [See Appendix D](#)

**Note:** Autologous hematopoietic stem cell transplantation is not considered standard of care for small lymphocytic lymphoma and should be treated in the same manner as chronic lymphocytic leukemia (CLL). Refer requests to Medical Director.

## Other Malignancies

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary in the following:

- Blastic plasmacytoid dendritic cell neoplasm (Dietrich et al., 2014)
- Rhabdomyosarcoma/soft tissue sarcoma may be appropriate as part of a clinical trial (Stiff et al., 2010). Refer to Medical Director.

**Allogeneic** hematopoietic stem cell transplantation is unproven and not medically necessary in all other circumstances including but not limited to the following:

- Atypical teratoid rhabdoid tumors (Nikolaides et al., 2010)
- Epithelial ovarian cancer
- Ewing tumor (Ewing sarcoma)
- Neuroblastoma
- Osteogenic sarcoma
- Renal cell carcinoma
- Retinoblastoma
- Supratentorial ependymoma (Venkatramani et al., 2013)

- Wilms tumor (Brown et al., 2010; Campbell et al., 2004).

**Autologous** hematopoietic stem cell transplantation may be considered medically necessary for the following:

- Ewing tumor (Ewing sarcoma)
- Neuroblastoma
- Retinoblastoma
- Supratentorial ependymoma (Venkatramani et al., 2013)
- Atypical teratoid rhabdoid tumors (Nikolaides et al., 2010)
  - Tandem auto may be indicated as part of a clinical trial
- Rhabdomyosarcoma/soft tissue sarcoma (Stiff et al., 2010)
  - May be appropriate as part of a clinical trial. Refer to Medical Director
- Wilms tumor (Brown et al., 2010; Campbell et al., 2004)
  - May be appropriate in relapsed disease as part of a clinical trial. Refer to Medical Director

**Autologous** hematopoietic stem cell transplantation is considered unproven and not medically necessary for all other malignancies including but not limited to:

- Blastic plasmacytoid dendritic cell neoplasm (Dietrich et al., 2014)
- Epithelial ovarian cancer
- Osteogenic sarcoma
- Renal cell carcinoma

## Hematological Disorders

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary in the following:

- Aplastic anemia
- Diamond-Blackfan syndrome
- Chronic granulomatous disease
- Congenital agranulocytosis (Kostmann syndrome)
- Congenital amegakaryocytic thrombocytopenia
- Dyskeratosis congenita
- Fanconi anemia
- Paroxysmal nocturnal hemoglobinuria (PNH)
- Shwachman-Diamond syndrome
- Sickle cell disease (SCD) (Kanter et al., 2021)
- Thalassemia major

**Autologous** hematopoietic stem cell transplantation is unproven and not medically necessary in hematologic disorder including but not limited to the following:

- Aplastic anemia
- Diamond-Blackfan syndrome
- Chronic granulomatous disease
- Congenital agranulocytosis (Kostmann syndrome)
- Congenital amegakaryocytic thrombocytopenia
- Dyskeratosis congenita
- Fanconi anemia
- Paroxysmal nocturnal hemoglobinuria (PNH)
- Shwachman-Diamond syndrome
- Sickle cell disease (SCD)
- Thalassemia major.

## Immunodeficiency Syndromes

Primary immunodeficiency diseases (PIDDs) are a collection of >200 rare disorders involving the absence or malfunction of integral parts of the immune system. The etiologies of PIDDs are not communicable in the infectious sense but vertically transmissible via transfer of inherited genetic defects. Although some defects affect a single part of the immune system, others cause multicomponent breakdown of combined innate and cellular immune responses and suggest the interplay of epigenetic influences. To be considered a PIDD, the cause must not be secondary to another disease, drug/chemical, or environmental exposure. Primary immunodeficiency disease onset may occur at birth, as most do, or at any subsequent developmental stage and may affect anyone, with only specific regard to gender or ethnicity (Ferri, 2024).

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary in the following:

- CD40 ligand deficiency
- Chediak-Higashi syndrome
- Hemophagocytic lymphohistiocytosis (HLH) (same as familial erythrophagocytic lymphohistiocytosis - FEL)
- Leukocyte adhesion deficiency
- Omenn syndrome
- Wiskott-Aldrich syndrome
- X-linked lymphoproliferative syndrome
- Lysosomal storage diseases (Heese, 2008)
- Severe combined immunodeficiency disease (SCID)
  - In addition to classical SCID, there are a variety of severe mixed (B- and T- cell) immune deficiency syndromes, with or without defined genetic abnormalities, which can be treated with allogeneic stem cell transplant.
  - As new genetic abnormalities are identified that can result in immunodeficiency syndromes, allogeneic transplantation may be appropriate treatment.
- Gaucher disease type I (Pastores et al., 2004; Charrow et al., 2004; Peters & Steward, 2003; Jmoudiak & Futerman, 2005)
  - Patients with the non-neuropathic type may benefit from a stem cell transplant following failed enzyme replacement therapy or if significant bone pain exists despite enzyme replacement therapy.
- Niemann-Pick type B (Schuchman, 2009)
  - In a non-cerebral form, transplantation may effectively diminish the impact of the accumulation of metabolic byproducts in lung and liver. These patients die from lung and liver disease and are candidates for stem cell transplantation.
- Fucosidosis (Miano et al., 2001; Vellodi et al., 1995)
  - There is little experience with transplantation for fucosidosis, a very rare entity among rare entities, but reports indicate that stem cell transplantation performed early effectively ameliorates disease progression.

**Autologous** hematopoietic stem cell transplantation is unproven and not medically necessary for immunodeficiency syndromes including but not limited to the following:

- CD40 ligand deficiency
- Chediak-Higashi syndrome
- Hemophagocytic lymphohistiocytosis (HLH) (same as familial erythrophagocytic lymphohistiocytosis - FEL)
- Leukocyte adhesion deficiency
- Omenn syndrome
- Severe combined immunodeficiency disease (SCID)
- Wiskott-Aldrich syndrome
- X-linked lymphoproliferative syndrome
- Gaucher disease type I (Pastores et al., 2004; Charrow et al., 2004; Peters & Steward, 2003; Jmoudiak & Futerman, 2005)
- Niemann-Pick type B (Schuchman, 2009)

- Fucosidosis (Miano et al., 2001; Vellodi et al., 1995)
- Lysosomal storage diseases (Heese, 2008)

## Autoimmune Diseases

Autoimmune diseases result when the immune system is overactive, causing it to attack and damage one's own body tissues. More than 100 different autoimmune diseases have been identified which together affect over 24 million people in the U.S. Treatment usually focuses on reducing immune system activity. While not a standard of care, hematopoietic stem cell transplantation, both allogeneic and autologous, continue to be studied in clinical trials for a number of diseases.

**Allogeneic** hematopoietic stem cell transplantation is unproven and not medically necessary in the following autoimmune disease (not an exhaustive list):

- Multiple sclerosis
- Systemic sclerosis (scleroderma)

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary in the following:

- Crohn's disease
  - Not a standard of care
  - Must be performed under a clinical trial and would only be considered for approval if the member's benefit plan supports participation in a clinical trial.
- Rheumatoid arthritis
  - Not a standard of care.
  - Must be performed under a clinical trial and would only be considered for approval if the member's benefit plan supports participation in a clinical trial.
- Systemic lupus erythematosus (SLE)
  - Not a standard of care
  - Must be performed under a clinical trial and would only be considered for approval if the member's benefit plan supports participation in a clinical trial

**Autologous** hematopoietic stem cell transplantation may be considered medically necessary in the following:

- Crohn's disease
  - Not a standard of care
  - Must be performed under a clinical trial and would only be considered for approval if the member's benefit plan supports participation in a clinical trial
- Multiple sclerosis (Cohen et al., 2019; Hooper, 2011; Kurtzke, 1982; National Multiple Sclerosis Society 2011)
  - Patient must meet the definition of relapsing-remitting (RR) or secondary progressive (SP) multiple sclerosis. [See Appendix E](#)
  - Expanded Disability Status Scale (EDSS) score between 2.0 and 6.0
  - Patient has failed treatment with one or more disease-modifying therapies (DMT)
  - Evidence of either of the following while being treated with DMT:
    - Two or more clinical relapses at separate times but within the previous 12 months
    - One relapse and a magnetic resonance imaging (MRI) gadolinium-enhancing lesion(s) at a separate time than the relapse but within the previous 12 months
- Rheumatoid arthritis
  - Not a standard of care.
  - Must be performed under a clinical trial and would only be considered for approval if the member's benefit plan supports participation in a clinical trial.
- Systemic lupus erythematosus (SLE)
  - Not a standard of care.
  - Must be performed under a clinical trial and would only be considered for approval if the member's benefit plan supports participation in a clinical trial.



- Systemic sclerosis (scleroderma) (Host et al., 2017; Sullivan et al., 2018) Adult patients at least 18 years of age with rapidly progressive systemic sclerosis (scleroderma) at risk of organ failure with either:
  - Pulmonary involvement with active interstitial lung disease and both:
    - Consistent bronchoalveolar cell composition or ground-glass opacities on CT of the chest
    - Either a forced vital capacity (FVC) or a diffusing capacity of the lung carbon monoxide (DLCO) of less than 70% of the predicted value
  - Renal involvement
- AND
- Patient does not have ANY of the following:
  - DLCO < 40% of predicted value
  - FVC < 45% of predicted value
  - Creatinine clearance < 40 ml/min
  - Pulmonary arterial hypertension
  - Left ventricular ejection fraction < 50%
- Patient is felt to be an appropriate candidate for autologous transplant by the treating facility

## Inherited Metabolic Disorders

**Allogeneic** hematopoietic stem cell transplantation may be considered medically necessary in the following:

- Adrenoleukodystrophy
- Epidermolysis bullosa
- Globoid cell leukodystrophy (Krabbe disease)
- Hurler syndrome (MPS I)
- Hunter syndrome (MPS II)
- Mannosidosis
- Maroteaux-Lamy syndrome (MPS VI)
- Metachromatic leukodystrophy
- Mitochondrial neurogastrointestinal encephalopathy (MNGIE) (Filosto et al., 2012; Halter et al., 2011)
- Osteopetrosis
- Rett syndrome

**Autologous** hematopoietic stem cell transplantation is unproven and not medically necessary for inherited metabolic disorders including but not limited to the following:

- Adrenoleukodystrophy
- Epidermolysis bullosa
- Globoid cell leukodystrophy (Krabbe disease)
- Hurler syndrome (MPS I)
- Hunter syndrome (MPS II)
- Mannosidosis
- Maroteaux-Lamy syndrome (MPS VI)
- Metachromatic leukodystrophy
- Mitochondrial neurogastrointestinal encephalopathy (MNGIE) (Filosto et al., 2012; Halter et al., 2011)
- Osteopetrosis
- Rett syndrome

## Cardiac Conditions

**Allogeneic and autologous** hematopoietic stem cell transplantation are unproven and not medically necessary for cardiac conditions. Stem cell transplantation would only be considered for approval under a clinical trial if the members benefit plan supports participation in a clinical trial.

## Additional Conditions

**Allogeneic** stem cell transplantation may be considered medically necessary in rare and unusual conditions. [See Appendix F](#)

**Note:** If there is a condition found within Appendix F that is not included above, refer to Medical Director.

## Relative Contraindications

The following list contains potential contraindications for hematopoietic stem cell transplant. While the conditions listed below would not be absolute contraindications for treatment they need to be addressed prior to transplant.

- Infections
  - Systemic or uncontrolled infection including sepsis.
- Significant uncorrectable life-limiting medical conditions.
- Severe end-stage organ damage that would have an impact on patient survival.
- Irreversible, severe brain damage.
- Social and psychiatric issues — It is expected that a patient has demonstrated adherence to all treatment plans and scheduled appointments and there is documentation of a support system and/or caregiver available to provide necessary care. A case should be referred for psychosocial evaluation and/or psychiatry consultation for guidance in any of the following circumstances:
  - Emotional instability, significant depression or other psychiatric illness that cannot be controlled and that would impact ability to comply with a complex evaluation process, surgical procedure and post-transplant plan of care and/or ability to give informed consent (and does not have a representative/guardian/conservator).
  - Limited cognitive ability (memory loss, dementia, etc.) that would impact ability to comply with a complex evaluation process, surgical procedure and post-transplant plan of care and/or ability to give informed consent (and does not have a representative/guardian/conservator).
  - Lack of psychosocial support as indicated by either no identified caregiver or an uncommitted caregiver. This would include the lack of transportation to and from transplant related appointments, patient and/or caregiver is unable to adhere to the requirements of transplant related treatment plan. A care contract may be needed.
  - Lack of sufficient financial means to purchase post-transplant medications.
  - History of non-adherence that has not been successfully remediated.
  - Inability to give informed consent. If the patient has an authorized representative/guardian/conservator or parent in the case of a minor, that individual must understand and support the ongoing health care needs of the patient.
- Limited irreversible rehabilitative potential (Bunnapradist, 2007).

## Special Considerations

Additional consultation and/or evaluation may be indicated in these situations. **Note:** Refer to Medical Director if questions remain.

- HIV infection
  - Patients should have a formal infectious disease consult indicating adequate treatment and proper assessment of risks related to this transplant.
  - Patients with known HIV infection must be on a HAART regimen and there must be documented evidence of viral load suppression.
- Serum creatinine < 2.5 mg/dL ( $\leq 1.5$  mg/dL in children) or GFR > 50ml/min.
  - Serum creatinine may be higher in patients with multiple myeloma or other plasma cell dyscrasias. Patients with multiple myeloma with reduced renal function are not prohibited from undergoing autologous SCT when the decreased renal function is related to the multiple myeloma (myeloma kidney). This includes patients on hemodialysis with no other contraindications.
- Active untreated or untreatable malignancy in patients undergoing stem cell transplantation for non-malignant indications. **Note:** Refer to Medical Director
- Patients with post-transplant lymphoproliferative disease (PTLD), having failed other conventional therapies, must have no active disease as demonstrated by negative positron emission tomography (PET) scan and resolved adenopathy on computed tomography (CT) and/or magnetic resonance imaging (MRI) (Blaes, 2009; Khedmat, 2009, Panagiotidis, 2014).

# Transplant Consultation Timing Guidelines

The following guidelines were developed jointly by the National Marrow Donor Program® (NMDP®)/Be the Match® and the American Society for Transplant and Cellular Therapy (ASTCT), and are based on current clinical practice, medical literature, National Comprehensive Cancer Network® (NCCN®) Guidelines for the treatment of cancer and evidence-based reviews.

The guidelines identify appropriate timing of consultation for autologous or allogeneic hematopoietic cell transplantation (HCT) based on disease characteristics. Evaluation and coordination of timing of HCT for eligible patients is determined in collaboration with the transplant center. Early referral is a critical factor for optimal transplant outcomes. In many situations, there may be a narrow window of opportunity to proceed to transplant and delays might preclude transplant or impair transplant outcomes. Research data comparing outcomes by disease status can be found at: <https://bethematchclinical.org/transplant-indications-and-outcomes/disease-specific-indications-and-outcomes/>

## Adult Leukemias and Myelodysplasia

### Acute Myeloid Leukemia (AML)

*High-resolution HLA typing is recommended at diagnosis for all patients*

HCT consultation should take place early after initial diagnosis for all patient with AML, including:

- Primary induction failure
- Measurable (also known as minimal) residual disease after initial therapy
- CR1- except favorable risk AML [defined as: t(8:2109q22;q22.1); RUNX1-RUNX1T1, inv(16)(p13.1q22) or t(16;16)(p13.1;q22); CBFβ-MYH11, mutated NPM1 without FLT3-ITD, biallelic mutated]. Transplant consultation may be reasonable for favorable-risk AML patients with unusual or adverse co-mutations or cytogenetic alterations. Early referral for allogeneic HCT should also be considered for any AML patients in CR1 who are 60 years old or older; regardless of cytogenetic or genomic information.
- Antecedent hematological disease (e.g., myelodysplastic syndrome [MDS]), either based on prior clinical diagnosis or suggested by the presence of secondary-type somatic mutations on molecular testing
- Treatment-related leukemia
- First relapse
- CR2 and beyond, if not previously evaluated

### Acute Lymphoblastic Leukemia (ALL)

(adult defined as ≥ 40 years)

*High-resolution HLA typing is recommended at diagnosis for all patients*

HCT consultation should take place early after initial diagnosis for all patients with ALL, including:

- Primary induction failure
- Measurable (also known as minimal) residual disease after initial therapy
- CR1
- First relapse
- CR2 and beyond, if not previously evaluated

### Myelodysplastic Syndromes (MDS)

*High-resolution HLA typing and referral to HCT consultation is recommended at diagnosis for all patients with:*

- Any intermediate or high IPSS or IPSS-R score
- Any MDS with poor prognostic features, including:
  - Treatment-related MDS
  - Refractory cytopenias
  - Adverse cytogenetics and molecular features
  - Transfusion dependence
  - Failure of hypomethylating agents or chemotherapy
  - Moderate to severe marrow fibrosis

## Chronic Myeloid Leukemia (CML)

- Inadequate hematologic or cytogenetic/molecular response to tyrosine kinase inhibitor (TKI) therapies
- Disease progression
- Intolerance to TKI therapies
- Accelerated phase
- Blast crisis (myeloid or lymphoid)
- T3151 mutation

## Myeloproliferative Neoplasms (MPN)

Including primary myelofibrosis (PMF) and essential thrombocythemia or polycythemia vera that has progressed to MF (secondary MF)

*High-resolution HLA typing and referral to HCT consultation is recommended at diagnosis for all patients with:*

- DIPSS or DIPSS Plus Intermediate-1 (INT-1) or higher
- MIPSS70/MIPSS 70 plus version 2.0 intermediate-risk or higher
- Cytopenic subtype
- Young age
- High-risk features including high-risk mutations (ASXL1, TP53), triple negative (lack of a driver mutation such as JAK2, MPL or CALR)
- Patients failing JAK inhibitor therapy

HCT is recommended upfront for patients with:

- DIPSS or DIPSS Plus Intermediate-2 (INT-2) and high-risk disease
- MIPSS70/MIPSS 70 plus version 2.0 high-risk disease
- Patients with DIPSS INT-1 or MIPSS70/MIPSS 70 plus version 2.0 intermediate-risk, cytopenic subtype, young age, high-risk features, including high-risk mutations (ASXL1, TP53), triple negative (lack of a driver mutation such as JAK2, MPL, or CALR) and those failing JAK inhibitor therapy should be considered for upfront HCT balancing patient preferences and clinical trial options

## Chronic Lymphocytic Leukemia (CLL)

- Resistance or intolerance to BTK inhibitors and/or BCL2 inhibitors
- Richter's transformation

## Pediatric Acute Leukemias and Myelodysplasia

### Acute Myeloid Leukemia (AML)

*High-resolution HLA typing is recommended at diagnosis for all patients*

HCT consultation should take place early after initial diagnosis for patients with AML including:

- Age < 2 years at diagnosis
- Primary induction failure
- Measurable (also known as minimal) residual disease after initial therapy
- CR1 – except favorable risk AML [defined as: t(8;21)(q22;q22.1); *RUNX1-RUNX1T1*, inv(16)(p13.1q22) or t(16;16)(p13.1;q220); *CBFB-MYH11*, mutated *NPM1* without *FLT3-ITD* or with *FLT3-ITD*<sup>low</sup>, biallelic mutated *CEBPA*]
- Monosomy 5 or 7
- Treatment-related leukemia
- First relapse
- CR2 and beyond, if not previously evaluated

### Acute Lymphoblastic Leukemia (ALL) (age ≤ 39 years)

- Infant at diagnosis
  - Unfavorable genetics

- Age < 3 months with any WBC, or < 6 months with WBC > 300,000 at presentation or any infant with measurable (also known as minimal) residual disease (MRD)+ after consolidation
- Primary induction failure (M3 marrow) after achieving MRD negative status
- Presence of MRD after initial therapy; MRD ≥ 0.01% following consolidation (9–12 weeks from diagnosis)
- High/very high-risk CR1, including:
  - Philadelphia chromosome positive slow-TKI responders or with *IKZF1* deletions; Philadelphia-like if MRD+ at end of consolidation, or persistently detectable low level of molecular disease
  - *iAMP21* if MRD+ at end of consolidation
  - 11q23 rearrangement if MRD+ at end of consolidation
- First relapse with aim to transplant in CR2
- CR2 and beyond, if not previously evaluated, including:
  - all young adults in CR2
  - early relapse (≤36 months from diagnosis for medullary disease, ≤18 months from diagnosis for EMD)
  - MRD+ (>0.1% for medullary disease or equivalent for EMD) after re-induction (4–8 weeks from relapse)
  - T cell ALL
  - CR3 and beyond
- Chimeric Antigen Receptor Therapy (CAR-T), including:
  - patients who receive CD19 4-1BB and achieve MRD negative CR if they have not already received HCT
  - patients who receive CD22 or other investigational therapies

### Myelodysplastic Syndromes (MDS)

- At diagnosis for all subtypes

### Juvenile Myelomonocytic Leukemia (JMML)

- At diagnosis

## Lymphomas

### Non-Hodgkin Lymphoma

#### Follicular (FL)

- Poor response to initial treatment
- Initial remission duration < 24 months
- At relapse (CAR or allo HCT can be offered to patients with multiple relapsed FL)
- Transformation to diffuse large B-cell lymphoma

#### Diffuse Large B-cell

- Primary induction failure, including residual PET avid disease
- First relapse
- CR2 or subsequent remission
- Double or triple hit (MYC and BCL-2 and/or BCL-6) - at diagnosis
- Primary CNS lymphoma at diagnosis PIF or first relapse

#### High Grade B-cell

- MYC and BCL-2 and/or BCL-6 rearrangements
- Primary induction failure
- CR1
- First relapse
- CR2 or subsequent remission

#### Mantle Cell

- At diagnosis
- First relapse
- Bruton's tyrosine kinase (BYK) intolerant or resistant disease

## Mature T-cell and NK Cell Lymphoma

- At diagnosis or CR1
- First relapse

## Other High-Risk Lymphomas

- At diagnosis

## Hodgkin Lymphoma

- Primary refractory disease
- First or subsequent relapse
- Brentuximab vedotin and check point inhibitor refractory and/or intolerant disease (for allo HCT)

## Other Malignant Diseases

### Germ Cell Tumors

High-dose therapy with autologous stem cell support may be considered for patients with:

- Non-germinomatous germ cell tumors (NGGCTs) for refractory disease post induction chemotherapy as long as the patient has chemo-responsive disease and does not have bulky residual tumor
- Germinoma or NGGCT if the patient has chemo-responsive disease to reinduction chemotherapy and does not have bulky residual tumor

### Neuroblastoma

- INRGSS L2 at diagnosis
  - *MYCN* amplification
  - age > 18 months with unfavorable histology and segmental chromosome aberration
- INRGSS stage M at diagnosis
  - *MYCN* amplification
  - age > 18 months at diagnosis
  - age 12-18 months with unfavorable histology, segmental chromosome aberrations, or diploid DNA content

### Ewing family of tumors

- Metastatic disease at diagnosis
- First relapse or CR 2

### Medulloblastoma

High-dose therapy with autologous stem cell support may be considered standard therapy for several infant (younger than 3 years of age) embryonal tumors as first line, including:

- Medulloblastoma
- Atypical teratoid rhabdoid tumor (AT/RT)
- Embryonal tumor with multilayered rosettes (ETMR)
- Primitive neuroectodermal/embryonal tumors, not otherwise specified (NOS)

High-dose therapy with autologous stem cell support may be considered at relapse for patients with:

- Embryonal tumors as long as the patient has chemoresponsive disease to re-induction chemotherapy and does not have bulky residual disease

## Plasma Cell Disorders

### Multiple Myeloma

- At diagnosis
- At progression and/or relapse

### Light Chain Amyloidosis

- At diagnosis
- At progression and/or relapse

### POEMS Syndrome (Osteosclerotic Myeloma)

- At diagnosis

## Non-Malignant Disorders

### Immune Deficiency Diseases

Including severe combined immunodeficiency syndromes, Wiskott-Aldrich syndrome, Omenn syndrome, X-linked lymphoproliferative syndrome, severe congenital neutropenia, and others.

- At diagnosis or if detected on newborn screening

### Inherited Metabolic Disorders

Including Hurler syndrome, adrenoleukodystrophy, and others.

- At diagnosis
- Adreno leukodystrophy (ALD): following the diagnosis of the cerebral form of ALD

## Hemoglobinopathies

### Sickle Cell Disease

- Children (especially under age 13) with available matched sibling donors
- All patients with aggressive course (stroke, end-organ complications, frequent pain crises)
- All patients with an alternative donor option and any of the following:
  - Stroke or silent cerebral infarct or cognitive impairment > 24 hours
  - Abnormal transcranial Doppler (TCD) velocity of  $\geq 200$  cm/sec or  $> 185$  cm/sec with intracranial vasculopathy
  - 
  - Frequent episodes of acute chest syndrome or severe vaso-occlusive pain crises or combination of both in the preceding 2–3 years
  - Regular red blood cell transfusions to prevent sickle cell disease complications
  - Tricuspid valve regurgitant jet (TRJ) velocity  $\geq 2.7$  m/sec (mainly in adults)
  - Chronic pain  $\geq 6$  months (leg ulcers, avascular necrosis)

### Transfusion-dependent Thalassemias

- At diagnosis

### Hemophagocytic Lymphohistiocytosis (HLH)

- At diagnosis

### Other Marrow Failure Syndromes

Including Diamond-Blackfan anemia, Shwachman-Diamond syndrome, Fanconi anemia, Dyskeratosis Congenita and others

- 
- Diamond-Blackfan Anemia: continued transfusion dependent anemia following a course of steroid therapy, development of significant infections, MDS/AML



- Shwachman-Diamond syndrome, Fanconi anemia, Dyskeratosis Congentia and others: development of cytopenias, transfusion dependence, or significant infections; high-risk cytopenic clones, high-risk somatic mutation patterns; MDS/AML

### **Systemic Sclerosis**

- At diagnosis or with diffuse disease, with increasing skin tightness score (modified Rodnan skin score, [mRSS]) and evidence of decrease (< 80%) in % predicted pulmonary function tests: forced vital capacity (FVC) and/or diffusion capacity (DLCO)

### **Multiple Sclerosis (MS)**

After MS relapse, with  $\geq 2$  relapse episodes in past 3 years, while on disease modifying therapy. Refer patient prior to progression of severe disability: patient must be able to walk 100 meters (with unilateral assistance: cane, crutch or brace)

# References

Alaggio, R., Amador, C., Anagnostopoulos, I., et al. The 5th edition of the World Health Organization Classification of Haematolymphoid Tumours: Lymphoid Neoplasms. *Leukemia* 2022; 36(7), 1720–1748. <https://doi.org/10.1038/s41375-022-01620-2>.

American Cancer Society. (2024). Key Statistics for Hodgkin Lymphoma. Hodgkin Lymphoma Statistics | How Common Is Hodgkin Disease? | American Cancer Society

Bashey A, Pérez WAS, Zhang M-J, et al. Comparison of twin and autologous transplants for multiple myeloma. *Biol Blood Marrow Transplant*. 2008;14(10):1118-24.

Bensinger WI. Reduced intensity allogeneic stem cell transplantation in multiple myeloma. *Front Biosci*. 2007 May;12:4384-4392.

Blade J, Rosin, L, Cibeira MT, et al. Hematopoietic stem cell transplantation for multiple myeloma beyond 2010. *Blood*. 2010;115(18):3655-3663.

Blaes AH et al. Positron emission tomography scanning in the setting of post-transplant lymphoproliferative disorders. *Clin Transplant*. 2009 Nov-Dec;23(6):794-9.

Brown E, Hebra A, Jenrette J, Hudspeth MP. Successful treatment of late, recurrent Wilms' tumor with high dose chemotherapy and autologous stem cell rescue in third complete response. *J Pediatr Hematol Oncol*. 2010;32(6):e241-e243. doi:10.1097/MPH.0b013e3181e5e25b.

Bruno B, et al. A comparison of allografting with autografting for newly diagnosed myeloma. *N Engl J Med*. 2007;356(11):1110-1120.

Bruno B, et al. Unrelated donor hematopoietic cell transplantation after non-cytoreductive conditioning for patients with high-risk myeloma. *Eur J Haematol*. 2007 Apr;78(4):330-337.

Bruno B, et al. Nonmyeloablative allografting for newly diagnosed multiple myeloma: the experience of the Gruppo Italiano Trapianti di Midollo. *Blood*. 2009;113:3375-3382.

Bruno B, Giaccone L, Sorasio R, Boccadero M. Role of allogeneic stem cell transplant in multiple myeloma. *Semin Hematol*. 2009 Apr;46(2):158-165.

Brunstein CG, Barker JN, Weisdorf DJ, et al. Umbilical cord blood transplantation after nonmyeloablative conditioning: impact on transplantation outcomes in 110 adults with hematologic disease. *Blood*. 2007 Oct;110(8):3064-3070.

Brunstein CG, Cantero S, Coa Q, et al. Promising progression-free survival for patients low and intermediate grade lymphoid malignancies after nonmyeloablative umbilical cord blood transplantation. *Biol Blood Marrow Transplant*. 2009;15:214-222.

Bunnapradist S, Danovitch G. Evaluation of Adult Kidney Transplant Candidates. *Am J Kidney Dis*. 2007 Nov; 50(5):890-898.

Campbell AD, Cohn SL, Reynolds M, et al. Treatment of relapsed Wilms' tumor with high-dose therapy and autologous hematopoietic stem-cell rescue: the experience at Children's Memorial Hospital. *J Clin Oncol*. 2004;22(14):2885-2890.

Campo, E., Jaffe, E. S., Cook, J. R., et al. The International Consensus Classification of Mature Lymphoid Neoplasms: a report from the Clinical Advisory Committee. *Blood* 2022; 140(11), 1229–1253. <https://doi.org/10.1182/blood.2022015851>

Castagna L, Sarina B, Bramanti S, et al. Donor lymphocyte infusion after allogeneic stem cell transplantation. *Transfus Apher Sci*. 2016 Jun;54(3):345-55. doi: 10.1016/j.transci.2016.05.011. Epub 2016 May 13. PMID: 27216544.

Cervantes F, Dupriez B, Pereira A, et al. New prognostic scoring system for primary myelofibrosis based on study of the International Working Group for Myelofibrosis Research and Treatment. *Blood*. 2009;113:2895-2901.

Charrow J, Andersson HC, Kaplan P, et al. Enzyme replacement therapy and monitoring for children with type 1 Gaucher disease: Consensus recommendations. *J Pediatr*. 2004;144:112-120.

Cheson BD, Pfistner B, Juweid ME, et al. Revised response criteria for malignant lymphoma. *J Clin Oncol*. 2007;25:579-86.

Cohen JA, Baldassari LE, Atkins HL, et al. Autologous Hematopoietic Cell Transplantation for Treatment-Refractory Relapsing Multiple Sclerosis: Position Statement from the American Society for Blood and Marrow Transplantation. *Biol Blood Marrow Transplant*. 2019;25(5):845-854. doi:10.1016/j.bbmt.2019.02.014

DeFilipp Z, Ciurea SO, Cutler C, et al. Hematopoietic Cell Transplantation in the Management of Myelodysplastic Syndrome: An Evidence-Based Review from the American Society for Transplantation and Cellular Therapy Committee on Practice Guidelines. *Transplantation and Cellular Therapy* 2023, 29(2), 71–81. <https://doi.org/10.1016/j.jtct.2022.11.014>

D'Souza A, Lacy M, Gertz M, et al. Long-term outcomes after autologous stem cell transplantation for patients with POEMS syndrome (osteosclerotic myeloma): a single-center experience. *Blood*. 2012 Jul;120(1):56-62.

de Lima M, Shpall E. Strategies for widening the use of cord blood in hematopoietic stem cell transplantation. *Haematologica*. 2006;91(5):584-7.

Devine SM, Flomenberg N, Vesole, et al. (1) Rapid mobilization of CD34+ cells following administration of the CXCR4 antagonist AMD3100 to patients with multiple myeloma and Non-Hodgkin's Lymphoma. *J Clin Oncol*. 2004;22:1095-1102.

Devine SM, Vij R, Rettig M, et al. (2) Rapid mobilization of functional donor hematopoietic cells without G-CSF using AMD3100, an antagonist of the CXCR4/SDF-1 interaction. *Blood*. 2008;112(4):990-998.

Dietrich S, Weidle J, Rieger J, et al. Rituximab Maintenance Therapy After Autologous Stem Cell Transplantation Prolongs Progression-Free Survival in Patients with Mantle Cell Lymphoma. *Leukemia*. 2014;28:708-709. doi:10.1038/leu.2013.332

DiPersio J, Stadtmauer EA, Nademanee AP, et al. (1) A phase III, multicenter, randomized, double-blind, placebo-controlled, comparative trial of AMD3100 (Plerixafor)+G-CSF vs. G-CSF+placebo for mobilization in multiple myeloma (MM) patients for autologous hematopoietic stem cell (aHSC) transplantation. *Blood (ASH Annual Meeting Abstracts)*. 2007;110: Abstract 445. © 2007 American Society of Hematology.

DiPersio J, Micallef I, Stiff PJ, et al. (2) A phase III, multicenter, randomized, double-blind, placebo controlled, comparative trial of AMD3100 (Plerixafor)+G-CSF vs. placebo+G-CSF in Non-Hodgkin's lymphoma (NHL) patients for autologous hematopoietic stem cell (aHSC) transplantation. *Blood (ASH Annual Meeting Abstracts)*. 2007;110: Abstract 601. © 2007 American Society of Hematology.

Döhner H, Estey E, Grimwade D, et al. Diagnosis and management of AML in adults: 2017 ELN recommendations from an international expert panel. *Blood*. 2017;129(4):424-447. doi:10.1182/blood-2016-08-733196

Epperla N, et al. Fludarabine and Busulfan versus Fludarabine, Cyclophosphamide, and Rituximab as Reduced-Intensity Conditioning for Allogeneic Transplantation in Follicular Lymphoma. *Biol Blood Marrow Transplant* 2018; 24:78–85.

Fernand JP, Bridoux F, Kyle RA, et al., International Kidney and Monoclonal Gammopathy Research Group. How I treat monoclonal gammopathy of renal significance (MGRS). *Blood*. 2013 Nov 21;122(22):3583-90. doi: 10.1182/blood-2013-05-495929. Epub 2013 Oct 9. PMID: 24108460.

Ferri F. Clinical Overview: Primary Immunodeficiency Diseases. ClinicalKey®. Elsevier 2024.

Fisher SA, Doree C, Mathur A, et al. Stem cell therapy for chronic ischemic heart disease and congestive heart failure. *Cochrane Database Syst Rev.* 2016 Dec 24;12(12):CD007888. doi: 10.1002/14651858.CD007888.pub3.

Filosto M, Scarpelli M, Tonin P, et al. Course and management of allogeneic stem cell transplantation in patients with mitochondrial neurogastrointestinal encephalomyopathy. *J Neurol.* 2012;259(12):2699-706.

Flomenberg N, Devine SM, Dpersio JF, et al. The use of AMD3100 plus G-CSF for autologous hematopoietic progenitor cell mobilization is superior to G-CSF alone. *Blood.* 2005;106(5):1867-1874.

Freed J, Talano J, Small T, Ricci A, Cairo MS. Allogeneic cellular and autologous stem cell therapy for sickle cell disease. *Bone Marrow Transplant.* 2012 Dec;47(12):1489-98. doi: 10.1038/bmt.2011.245. Epub 2011 Dec 19.

Gagelmann N, Ditschkowski M, Bogdanov R, et al. Comprehensive clinical-molecular transplant scoring system for myelofibrosis undergoing stem cell transplantation. *Blood.* 2019 May 16;133(20):2233-2242. doi: 10.1182/blood-2018-12-890889. Epub 2019 Feb 13. PMID: 30760453.

Giralt S, Garderet L, Durie B, et al. American Society of Blood and Marrow Transplantation, European Society of Blood and Marrow Transplantation, Blood and Marrow Transplant Clinical Trials Network, and International Myeloma Working Group Consensus Conference on Salvage Hematopoietic Cell Transplantation in Patients with Relapsed Multiple Myeloma. *Biol Blood Marrow Transplant.* 2015 Dec;21(12):2039-51. doi:10.1016/j.bbmt.

Halter J, Schüpbach WM, Casali C, et al. Allogeneic hematopoietic SCT as treatment option for patients with mitochondrial neurogastrointestinal encephalomyopathy (MNGIE): a consensus conference proposal for a standardized approach. *Bone Marrow Transplant.* 2011 Mar;46(3):330-7.

Harousseau JL, Moreau P. Autologous hematopoietic stem-cell transplantation for multiple myeloma. *N Engl J Med.* 2009;360(25):2645-2654.

Heese BA. Current strategies in the management of lysosomal storage diseases. *Semin Pediatr Neurol.* 2008;15:119-126.

Herzog CE & Huh WW (2025). Gonadal and Germ Cell Neoplasms. In *Nelson Textbook of Pediatrics* (pp3150-3153.e1. Elsevier.

Horwitz ME, Stiff PJ, Cutler C, et al. Omidubicel vs standard myeloablative umbilical cord blood transplantation: results of a phase 3 randomized study. *Blood.* 2021 Oct 21;138(16):1429-1440. doi: 10.1182/blood.2021011719. PMID: 34157093; PMCID: PMC9710469.

Host L, Nikpour M, Calderone A, et al. Autologous stem cell transplantation in systemic sclerosis: a systematic review. *Clin Exp Rheumatol.* 2017;35 Suppl 106(4):198-207.

Ji ZF, Zhang DY, Weng SQ, et al. POEMS Syndrome: A report of 14 cases and review of the literature. *ISRN Gastroenterol.* 2012;2012:584287.

Jmoudiak M, Futerman AH. Gaucher disease: Pathological mechanisms and modern management. *Br J Haematol.* 2005;129:178-188.

Kalaycio ME, Kukreja M, Woolfrey AE, et al. Allogeneic hematopoietic cell transplant for prolymphocytic leukemia. *Biol Blood Marrow Transplant.* 2010;16:543.

Kanter J, Liem RI, Bernaudin F, et al. American Society of Hematology 2021 guidelines for sickle cell disease: stem cell transplantation. *Blood Adv.* 2021 Sep 28;5(18):3668-3689. doi: 10.1182/bloodadvances.2021004394C. PMID: 34581773; PMCID: PMC8945587.

Khedmat H. Early onset post transplantation lymphoproliferative disorders: analysis of international data from 5 studies. *Ann Transplant.* 2009;14(3):74-7.

Klingebiel T, Cornish J, Labopin M, et al. on behalf of the Pediatric Diseases and Acute Leukemia Working Parties of the European Group for Blood and Marrow Transplantation (EBMT). Results and factors influencing outcome after

fully haploidentical hematopoietic stem cell transplantation in children with very high-risk acute lymphoblastic leukemia: impact of center size: an analysis on behalf of the Acute Leukemia and Pediatric Disease Working Parties of the European Blood and Marrow Transplant group. *Blood*. 2010;115(17):3437-3446.

Krishnan B, Else M, Tjonnfjord GE, et al. Stem cell transplantation after alemtuzumab in T-cell prolymphocytic leukaemia results in longer survival than after alemtuzumab alone: a multicentre retrospective study. *Br J Haematol*. 2010;149:907.

Kumar A, Kharfan-Dabaja MA, Glasmacher A, et al. Tandem versus single autologous hematopoietic cell transplantation for the treatment of multiple myeloma: a systematic review and meta-analysis. *J Natl Cancer Inst*. 2009;101:100-106.

LeMaistre CF et al. Standardization for terminology of episodes of hematopoietic stem cell patient transplant care. *Biol Blood Marrow Transplant* 2013; 19(6):851-57.

Leukemia Foundation. (2024). Primary myelofibrosis. [www.leukaemia.org](http://www.leukaemia.org)

Leung N, Bridoux F, Batuman V, et al. Publisher Correction: The evaluation of monoclonal gammopathy of renal significance: a consensus report of the International Kidney and Monoclonal Gammopathy Research Group. *Nat Rev Nephrol*. 2019 Feb;15(2):121. doi: 10.1038/s41581-018-0102-7. Erratum for: *Nat Rev Nephrol*. 2019 Jan;15(1):45-59. PMID: 30568288; PMCID: PMC7608309.

Li J, Zhang W, Duan MH, et al. PBSC mobilization in newly diagnosed patients with POEMS syndrome: outcomes and prognostic factors. *Bone Marrow Transplant*. 2013 Feb;48(2):233-7.

Lin C, Horwitz ME. Corrigendum to 'Multicenter Long-Term Follow-Up of Allogeneic Hematopoietic Cell Transplantation with Omidubicel: A Pooled Analysis of Five Prospective Clinical Trials'. *Transplant Cell Ther*. 2023 Jul 5:S2666-6367(23)01356-8. doi: 10.1016/j.jtct.2023.06.009. Epub ahead of print. Erratum for: *Transplant Cell Ther*. 2023 May;29(5):338.e1-338.e6. PMID: 37421972.

Lin C, Schwarzbach A, Sanz J, et al. Multicenter Long-Term Follow-Up of Allogeneic Hematopoietic Cell Transplantation with Omidubicel: A Pooled Analysis of Five Prospective Clinical Trials. *Transplant Cell Ther*. 2023 May;29(5):338.e1-338.e6. doi: 10.1016/j.jtct.2023.01.031. Epub 2023 Feb 10. Erratum in: *Transplant Cell Ther*. 2023 Jul 5; PMID: 36775201; PMCID: PMC10149622.

Manning, E. (2022). Myelofibrosis. Salem Press Encyclopedia of Health.

McNeer JL, Devidas M, Dai Y, et al. Hematopoietic Stem-Cell Transplantation Does Not Improve the Poor Outcome of Children with Hypodiploid Acute Lymphoblastic Leukemia: A Report From Children's Oncology Group. *J Clin Oncol*. 2019;37(10):780-789. doi:10.1200/JCO.18.00884

Miano M, Lanino E, Gatti R, et al. Four-year follow-up of a case of fucosidosis treated with unrelated donor bone marrow transplantation. *Bone Marrow Transplant*. 2001;27:747-751.

National Cancer Institute (2024). <https://www.cancer.gov/types/leukemia>

National Marrow Donor Program (NMDP) and the American Society for Transplantation and Cellular Therapy (ASTCT). 2024 Recommended Timing for Transplant Consultation. Accessed 9/6/2024. Available at: <https://bethematchclinical.org/transplant-indications-and-outcomes/referral-timing-guidelines/>

Navarro WH, Loberiza, Jr. FR, Bajorunaite R, et al. Effect of body mass index on mortality of patients with lymphoma undergoing autologous hematopoietic cell transplantation. *Biol Blood Marrow Transplant*. 2006;12:541-551.

Navarro WH, Agovi M-A, Logan BR, et al. Obesity does not preclude safe and effective myeloablative hematopoietic cell transplantation (HCT) for acute myelogenous leukemia (AML) in adults. *Biol Blood Marrow Transplant*. 2010;16:1-9.

Nicolaides T, Tihan T, et al. High-dose chemotherapy and autologous stem cell rescue for atypical teratoid/rhabdoid tumor of the central nervous system. *J Neurooncol*. 2010 May; 98(1): 117-123. doi: 10.1007/s11060-009-007.

Oliansky DM, Rizzo JD, Aplan PD, et al. The role of cytotoxic therapy with hematopoietic stem cell transplantation in the therapy of acute myeloid leukemia in children: an evidence-based review. *Biol Blood Marrow Transplant.* 2007;13:1-25.

Oliansky DM, Appelbaum F, Cassileth PA, et al. The role of cytotoxic therapy with hematopoietic stem cell transplantation in the therapy of acute myelogenous leukemia in adults: an evidence-based review. *Biol Blood Marrow Transplant.* 2008;14:137-180.

Oliansky DM, Antin JH, Bennett JM, et al. The role of cytotoxic therapy with hematopoietic stem cell transplantation in the therapy of myelodysplastic syndromes: an evidence-based review. *Biol Blood Marrow Transplant.* 2009;15:137-172.

Oliansky DM, Gordon LI, King J, et al. The role of cytotoxic therapy with hematopoietic stem cell transplantation in the treatment of follicular lymphoma: an evidence-based review. *Biol Blood Marrow Transplant.* 2010;16:443-468.

Oliansky DM, Czuczman M, Fisher RI, et al. The role of cytotoxic therapy with hematopoietic stem cell transplantation in the treatment of diffuse large B cell lymphoma: update of the 2001 evidence-based review. *Biol Blood Marrow Transplant.* 2011;17:20-47.

Optum Stem Cell Transplant Expert Panel Meeting. June 17, 2015.

Panagiotidis E, Quigley AM, Pencharz D, et al. (18)F-fluorodeoxyglucose positron emission tomography/computed tomography in diagnosis of post-transplant lymphoproliferative disorder. *Leuk Lymphoma.* 2014;55(3):515-519. doi:10.3109/10428194.2013.813501

Passamonti F, Cervantes F, Vannucchi AM, et al. A dynamic prognostic model to predict survival in primary myelofibrosis: a study by the IWG-MRT (International Working Group for Myeloproliferative Neoplasms Research and Treatment). *Blood.* 2010;115:1703-1708.

Pastores GM, Weinreb NJ, Aerts H, et al. Therapeutic goals in the treatment of Gaucher disease. *Semin Hematol.* 2004;41:4-14.

Patel JP, Levine RL. How do novel molecular genetic markers influence treatment decisions in acute myeloid leukemia? *Hematology Am Soc Hematol Educ Program.* 2012;2012:28-34.

Peters C, Steward CG. Hematopoietic cell transplantation for inherited metabolic diseases: An overview of outcomes and practice guidelines. *Bone Marrow Transplant.* 2003;31:229-239.

Popat U, Mehta RS, Rezvani K, et al. Enforced fucosylation of cord blood hematopoietic cells accelerates neutrophil and platelet engraftment after transplantation. *Blood.* 2015 May 7;125(19):2885-92. doi: 10.1182/blood-2015-01-607366. Epub 2015 Mar 16. PMID: 25778529; PMCID: PMC4424412

Rajkumar SV. Multiple myeloma: 2022 update on diagnosis, risk stratification, and management. *Am J Hematol.* 2022 Aug;97(8):1086-1107. doi: 10.1002/ajh.26590. Epub 2022 May 23. PMID: 35560063; PMCID: PMC9387011. Rotta et al. Long-term outcome of patients with multiple myeloma after autologous hematopoietic cell transplantation and nonmyeloablative conditioning. *Blood.* 2009;113:3383-3391.

Schiller GJ. High-risk acute myelogenous leukemia: treatment today...and tomorrow. *Hematology Am Soc Hematol Educ Program.* 2013;2013:201-8.

Schuchman EH. The pathogenesis and treatment of acid sphingomyelinase-deficient Niemann-Pick disease. *Int J Clin Pharmacol Ther.* 2009;47 Suppl 1:S48-57.

Short NJ, Jabbour E. Minimal Residual Disease in Acute Lymphoblastic Leukemia: How to Recognize and Treat It. *Curr Oncol Rep.* 2017;19(1):6. doi:10.1007/s11912-017-0565-x

Stiff PJ, Agovi M-A, Antman KH, et al. High-dose chemotherapy with blood or bone marrow transplants for rhabdomyosarcoma. *Biol Blood Marrow Transplant.* 2010;16:525-532.

Sullivan KM, Majhail NS, Bredeson C. et al. Systemic sclerosis as an indication for autologous hematopoietic cell transplantation: position statement from the American Society for Blood and Marrow Transplantation. *Biol Blood Marrow Transplant.* 2018;24:1961-64.

Sureda A. et al. Allogeneic Hematopoietic Stem Cell Transplantation for Relapsed Follicular Lymphoma: A Combined Analysis on Behalf of the Lymphoma Working Party of the EBMT and the Lymphoma Committee of the CIBMTR. *Cancer* 2018;124:1733-42.

Swerdlow SH, Campo E, Harris, NL (Eds.). WHO classification of tumours of haematopoietic and lymphoid tissues. World Health Organization Classification of Tumours. Revised 4th edition. Volume 2. 2017.

Vellodi A, Cragg H, Winchester B, et al. Allogeneic bone marrow transplantation for fucosidosis. *Bone Marrow Transplant.* 1995;15:153-158.

Venkatramani R. et al. Outcome of infants and young children with newly diagnosed ependymoma treated on the "Head Start" III prospective clinical trial. *J Neurooncol.* 2013;113(2):285-91.

# Appendix A: Clinical, Cytogenetic and Mutational Risk Stratification for AML

## Favorable risk:

- Cytogenetics
  - t(8;21)
  - inv(16) or t(16;16)
- Mutations
  - Kit

## Intermediate risk (one or more of the following):

- Cytogenetics
  - Normal
  - +8
- Mutations
  - Flt3 ITD-positive
  - Mutant TET2, MLL-PTD, DNMT3A, ASXL1, PHF6

## Unfavorable high risk (one or more of the following):

- Cytogenetics
  - -5/-7
  - 11q23, 20q
  - 3 or more
- Clinical features:
  - CR2 and beyond
  - Age > 70
  - Refractory to induction chemotherapy
  - Persistence of minimal residual disease following induction

Patel JP, Levine RL. How do novel molecular genetic markers influence treatment decisions in acute myeloid leukemia? Hematology Am Soc Hematol Educ Program. 2012; 2012:28-34.



# **Appendix B: Prognostic Risk Scores for Myelodysplastic Syndrome**

Medicare coverage for allogeneic hematopoietic stem cell transplant using bone marrow, peripheral blood or umbilical cord blood stem cell products for Medicare patients with MDS are dependent on prognostic risk scores of:

- $\geq 1.5$  (Intermediate-2 or high) using the International Prognostic Scoring System (IPSS), or
- $\geq 4.5$  (high or very high) using the International Prognostic Scoring System - Revised (IPSS-R), or
- $\geq 0.5$  (high or very high) using the Molecular International Prognostic Scoring System (IPSS-M)

<https://www.cms.gov/medicare-coverage-database/view/ncd.aspx?ncdid=366>

## **Appendix C: The Dynamic International Prognostic Scoring System (DIPSS) and Dynamic International Prognostic Scoring System-Plus (DIPSS-Plus) for Primary Myelofibrosis (PMF)**

<b>DIPSS Factors</b>	<b>Point Value</b>
Age > 65	1
Hemoglobin level < 10 g/dl	2
White blood cell count (WBC) > 25 x 10 <sup>9</sup> /L	1
Peripheral blood blasts ≥ 1%	1
Peripheral blood blasts ≥ 1%	1

DIPSS Risk Categories: Low (0 points), Intermediate 1 (1 point), Intermediate 2 (2-3 points), High (≥ 4 points).

<b>DIPSS-Plus Factors</b>	<b>Point Value</b>
Adverse karyotypes*	1
Platelets < 100 x 10 <sup>9</sup> /L	1
RBC transfusion need	1

\*Adverse karyotypes include +8, -5/del5q, -7/del7qi(17q), inv(3), 11q23 rearrangements.

DIPSS-Plus Risk Categories: Low (0 points), Intermediate 1 (1 point), Intermediate 2 (2-3 points), High (4-6 points).

Gagelmann N, Ditschkowski M, Bogdanov R, et al. Comprehensive clinical-molecular transplant scoring system for myelofibrosis undergoing stem cell transplantation. *Blood*. 2019 May 16;133(20):2233-2242. doi: 10.1182/blood-2018-12-890889. Epub 2019 Feb 13. PMID: 30760453.

# **Appendix D: Complete Remission and Partial Remission Highlights from Revised Response Criteria for Malignant Lymphoma**

Complete Remission (CR): Disappearance of all evidence of disease.

Nodal masses

- FDG-avid or PET positive prior to therapy: mass of any size permitted if PET negative
- Variably FDG-avid or PET negative: regression to normal size on CT

Spleen, Liver

- Not palpable, nodules disappeared

Bone marrow

- Infiltrate cleared on repeat biopsy; if indeterminate by morphology, immunohistochemistry should be negative

Partial Remission (PR): Regression of measurable disease and no new sites.

Nodal masses

- Greater than 50% decrease in sum of the products of diameters (SPD) of up to 6 largest dominant masses, no increase in size of other nodes
  - FDG-avid or PET positive prior to therapy; one or more PET positive at previously involved site
  - Variably FDG-avid or PET negative; regression on CT
  - NOTE: In the absence of adequate size measurements one can use a greater than 50% decrease in the Standardized Uptake Value (SUV) to document PR.

Spleen, Liver

- Greater than 50% decrease in SPD of nodules (for single nodule in greatest transverse diameter); no increase in size of liver or spleen.

Bone marrow

- Irrelevant if positive prior to therapy; cell type should be specified.

Cheson BD, Pfistner B, Juweid ME, et al. Revised response criteria for malignant lymphoma. J Clin Oncol. 2007;25:579–86. Available at: <http://jco.ascopubs.org/content/25/5/579.full.pdf+html>.

# Appendix E: Multiple Sclerosis Definitions

## Relapsing-Remitting MS (RRMS)

A pattern of symptoms of multiple sclerosis in which symptomatic attacks occur that last 24 hours or more., followed by complete or almost complete improvement. This is the most common form of multiple sclerosis. About 85% of people with MS are initially diagnosed with RRMS. People with RRMS have temporary periods called relapses, flare-up or exacerbations, when new symptoms appear (Hooper, 2011).

## Secondary-Progressive MS (SPMS)

A pattern of symptoms of multiple sclerosis in which there are relapses and remissions, followed by more steady progression of symptoms. In SPMS, symptoms worsen more steadily over time, with or without the occurrence of relapses and remissions. Most people who are diagnosed with RRMS will transition to SPMS at some point (National Multiple Sclerosis Society, 2011).

## Relapse

A relapse of MS (also known as also known as an exacerbation attack or flare-up) is the occurrence new symptoms or the worsening of old symptoms. It can be very mild, or severe enough to interfere with a person's ability to function. No two exacerbations are alike. Symptoms vary from person to person and from one exacerbation to another. For example, the exacerbation might be an episode of optic neuritis (caused by inflammation of the optic nerve that impairs vision), or problems with balance or severe fatigue. Some relapses produce only one symptom (related to inflammation in a single area of the central nervous system). Other relapses cause two or more symptoms at the same time (related to inflammation in more than one area of the central nervous system).

To be a true exacerbation, the attack must last at least 24 hours and be separated from the previous attack by at least 30 days. It must also occur in the absence of infection, or other cause. Most exacerbations last from a few days to several weeks or even months (National Multiple Sclerosis Society, 2011).

Hooper K. Managing Progressive MS. New York, NY: National Multiple Sclerosis Society; 2011.  
Gale Encyclopedia of Medicine. Copyright 2008 The Gale Group, Inc. All rights reserved.

Kurtzke JF. Rating neurologic impairment in multiple sclerosis: an expanded disability status scale (EDSS). *Neurology* 1983;33(11):1444-1452.

Multiple Sclerosis: Just the Facts New York, NY; National Multiple Sclerosis Society;2011  
Gale Encyclopedia of Medicine. Copyright 2008 The Gale Group, Inc. All rights reserved.  
<https://www.nationalmssociety.org/Treating-MS/Managing-Relapses>

Gangat N, Caramazza D, Vaidya R. et al. DIPSS Plus: A refined dynamic international prognostic scoring system for primary myelofibrosis that incorporates prognostic information from karyotype, platelet count, and transfusion status. *J Clin Oncol.* 2011;29(4):392-97.

Salit, RB & Deeg HJ. Transplant decisions in patients with myelofibrosis: should mutations be the judge? *Biol Blood Marrow Transplant.* 2018; 24: 649-58.

# Appendix F: Hematopoietic Stem Cell Transplant Reference Sheet

The following is a list of rare and unusual conditions where allogeneic transplant may be indicated. The list was reviewed and accepted by the 2018 Optum Hematopoietic Stem Cell Transplant Expert Panel. **Note:** If there is a condition found on this list that is not included in the “Indications” section above, refer to Medical Director.

## 1. Lymphocyte Immunodeficiencies (many fall under ‘severe combined immunodeficiency’ classification)

Adenosine deaminase deficiency

Artemis deficiency

Calcium channel deficiency

Cernunnos-XLF immunodeficiency

CHARGE syndrome with immune deficiency Common gamma chain deficiency

Deficiencies in CD 45, CD3, CD8

DiGeorge syndrome

DNA ligase IV

DOCK8 immunodeficiency syndrome

GATA2 deficiency

Interleukin-7 receptor alpha deficiency

Janus-associated kinase 3 (JAK3) deficiency

Major histocompatibility class II deficiency

Purine nucleoside phosphorylase deficiency

Recombinase-activating gene (RAG) 1/2 deficiency

Reticular dysgenesis

Winged helix deficiency

Zeta-chain-associated protein-70 (ZAP-70) deficiency

## 2. Phagocytic Deficiencies

Chediak-Higashi syndrome

Griscelli syndrome, type 2

Interferon-gamma receptor deficiencies

Leukocyte adhesion deficiency

Shwachman-Diamond syndrome\*

\*may be considered as marrow failure syndrome rather than immunodeficiency

### 3. Other Immunodeficiencies

Autoimmune lymphoproliferative syndrome  
Cartilage hair hypoplasia  
CD25 deficiency  
Familial hemophagocytic lymphohistiocytosis  
Hyper IgD and IgE syndromes  
ICF syndrome IPEX syndrome NEMO deficiency  
NF- $\kappa$ B inhibitor, alpha (I $\kappa$ B-alpha)

Antoine C, Muller S, Cant A, et al. Long term survival and transplantation of hematopoietic stem cells for immunodeficiencies: report of the European experience 1968-99. *Lancet*. 2003 Feb;361(9357):553-60. PMID: 12598139254.

Burroughs L, Woolfrey A, Shimamura A. Shwachman-Diamond syndrome: a review of the clinical presentation, molecular pathogenesis, diagnosis, and treatment. *Hematol Oncol Clin North Am*. 2009 Apr;23(2):233-48. PMID: 19327581.

Coppa GV, Gabrielli O, Zampini L, et al. Bone marrow transplantation in Hunter syndrome (mucopolysaccharidosis type II): two-year follow-up of the first Italian patient and review of the literature. *Pediatr Med Chir*. 1995 May-Jun;17(3):227-35. PMID:7567644.

Ehlert K, Roth J, Frosch M, et al. Farber's disease without central nervous system involvement: bone-marrow transplantation provides a promising new approach. *Ann Rheum Dis*. 2006;65(12):1665-6.

Filipovich A. Hematopoietic cell transplantation for correction of primary immunodeficiencies. *Bone Marrow Transplant*. 2008 Aug;42 Suppl 1:S49-S52. PMID: 18724301.

Guffon N, Bertrand Y, Forest I, et al. Bone marrow transplantation in children with Hunter syndrome: outcome after 7 to 17 years. *J Pediatr*. 2009 May;154(5):733-7.

Heese BA. Current strategies in the management of lysosomal storage diseases. *Semin Pediatr Neurol*. 2008 Sep;15(3):119-26. PMID: 18708002.

Myers KC, Davies SM. Hematopoietic stem cell transplantation for bone marrow failure syndromes in children. *Biol Blood Marrow Transplant*. 2009 Mar;15(3):279-92. PMID:19203719.

Orange JS, Hossny EM, Weiler CR, et al. Use of intravenous immunoglobulin in human disease: a review of evidence by members of the Primary Immunodeficiency Committee of the American Academy of Allergy, Asthma and Immunology. *J Allergy Clin Immunol*. 2006 Apr;117(4 Suppl):S525-53. PMID: 16580469.

Tolar J, Blazar BR, Wagner JE. Concise review: Transplantation of human hematopoietic cells for extracellular matrix protein deficiency in epidermolysis bullosa. *Stem Cells*. 2011 Jun;29(6):900-6. doi: 10.1002/stem.647. Review. PubMed PMID: 21557391.

Vellodi A, Young E, Cooper A, et al. Long-term follow-up following bone marrow transplantation for Hunter disease. *J Inher Metab Dis*. 1999 Jun;22(5):638-48.

Vormoor J, Ehlert K, Groll AH, et al. Successful hematopoietic stem cell transplantation in Farber disease. *J Pediatr*. 2004 Jan;144(1):132-4.

# **Appendix G: Updated Criteria for Diagnosis of Multiple Myeloma**

## **Multiple myeloma**

DIAGNOSTIC CRITERIA: ALL 3 REQUIRED

- Monoclonal plasma cells in the bone marrow > 10% and/or presence of a biopsy-proven plasmacytoma
- Monoclonal protein present in the serum and/or urine \*
- Myeloma-related organ dysfunction (1 or more) \*\*

Traditional CRAB Criteria:

[C] Calcium elevation in the blood S. Calcium >10.5 mg/l or upper limit of normal

[R] Renal insufficiency S. Creatinine > 2 mg/dl

[A] Anemia Hemoglobin < 10 g/dl or 2 g < normal

[B] Lytic bone lesions or osteoporosis \*

NOTE: These criteria identify stage IB and stages II and IIIA/B myeloma by Durie Salmon stage. Stage IA becomes smoldering or indolent myeloma.

\* If no monoclonal protein is detected (non-secretory disease), then > 30 % monoclonal bone marrow plasma cells and/or a biopsy-proven plasmacytoma required.

\*\* The revised International Myeloma Working Group (IMWG) criteria will allow, in addition to the classic CRAB features, the following three markers as “myeloma defining events” (MDEs):

- Sixty percent or greater clonal plasma cells on bone marrow examination
- Serum involved/uninvolved free light chain ratio of 100 or greater, provided the absolute level of the involved free light chain is at least 100 mg/l (a patient’s “involved” free light chain – either kappa or lambda – is the one that is above the normal reference range; the uninvolved light chain is the one that typically is in, or below, the normal range)
- More than one focal lesion on MRI that is at least 5 mm or greater in size

The presence of at least one of these markers will be considered sufficient for a diagnosis of multiple myeloma, regardless of the presence or absence of symptoms or CRAB features. Each of these markers has been shown in two or more independent studies to be associated with an approximately 80 % or higher risk of developing myeloma-related organ damage within two years.

In addition, the IMWG criteria allow the use of CT and PET-CT for detecting osteolytic bone lesions in order to make the diagnosis of myeloma. In patients with equivocal findings on MRI, CT, and/or PET-CT, the IMWG recommends follow-up imaging. The use of modern imaging methods at diagnosis and follow-up will enable the diagnosis of myeloma to be made before serious bone damage, such as pathologic fractures, can develop.

## Monoclonal gammopathy of undetermined significance (MGUS)

DIAGNOSTIC CRITERIA: ALL 3 REQUIRED

- Serum monoclonal protein and/or urine monoclonal protein level low\*
- Monoclonal bone marrow plasma cells < 10 %
- Normal serum calcium, hemoglobin level and serum creatinine

\* Low is defined as:

- Serum IgG < 3.5 g/dl
- Serum IgA < 2.0 g/dl

No bone lesions on full skeletal x-ray survey and/or other imaging if performed.

No clinical or laboratory features of amyloidosis or light chain deposition disease.

Urine monoclonal kappa or lambda < 1.0 g/24 hours.

The definition of MGUS has not changed. However, a new entity termed light chain MGUS has been defined.

## Smoldering or indolent myeloma

DIAGNOSTIC CRITERIA: ALL 3 REQUIRED

- Monoclonal protein present in the serum and/or urine
- Monoclonal plasma cells present in the bone marrow and/or a tissue biopsy
- Not meeting criteria for MGUS, multiple myeloma, or solitary plasmacytoma of bone

NOTE: These criteria identify stage IA myeloma by Durie Salmon stage.

The diagnosis of smoldering myeloma will now have an upper limit of 60% for the percentage of clonal plasma cells in the marrow. Patients considered to have smoldering myeloma should not have any myeloma defining events or amyloidosis.

A new kind of smoldering multiple myeloma, termed light chain smoldering multiple myeloma, has been recently described in a study conducted at the Mayo Clinic, and the specific monoclonal protein level required for this diagnosis has also been added.

Rajkumar SV. Multiple myeloma: 2022 update on diagnosis, risk stratification, and management. Am J Hematol. 2022 Aug;97(8):1086-1107. doi: 10.1002/ajh.26590. Epub 2022 May 23. PMID: 35560063; PMCID: PMC9387011



# Appendix H: Hematopoietic Stem Cell Transplant Quick Reference Guide

Appendix H is to be used solely as a quick reference guide to identify standard of care. This guide does not reflect potential exceptions such as enrollment in a clinical trial. It is required that the user read the guideline for important criteria and directions concerning medical director escalation.

Disease/Indication	Autologous	Allogeneic
<b>Leukemias</b>		
Acute Lymphoblastic Leukemia (ALL)	Yes	Yes
Acute myeloid leukemia (AML)	Yes	Yes
Chronic lymphocytic leukemia (CLL)	No	Yes
Chronic myeloid leukemia (CML)	No	Yes
Prolymphocytic leukemia	Yes	Yes
<b>Myelodysplastic Syndromes &amp; Mixed Myelodysplastic/Myeloproliferative Neoplasms</b>		
Myelodysplastic syndromes (MDS)	No	Yes
Juvenile myelomonocytic leukemia (JMML/JCML)	No	Yes
Chronic myelomonocytic leukemia (CMML)	No	Yes
<b>Myeloproliferative Disorders</b>		
Primary myelofibrosis and related conditions	No	Yes
Secondary myelofibrosis	No	Yes
<b>Brain Tumors</b>		
Anaplastic astrocytoma	No	No
Brain stem glioma	No	No
Ependymoma	No	No
Germinoma	No	No
Glioblastoma Multiforme (GBM)	No	No
Medulloblastoma	Yes	No

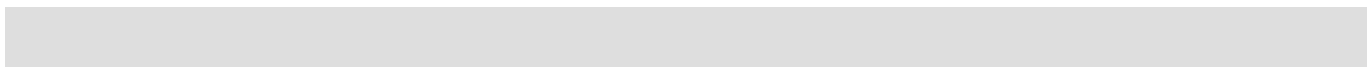
Oligodendroglioma	Yes	No
Pineoblastoma	Yes	No
Embryonal Tumors with Multi-layered Rosettes (ETMR). Formerly known as Primitive Neuroectodermal Tumor (PNET)	Yes	No

<b>Disease/Indication</b>	<b>Autologous</b>	<b>Allogeneic</b>
<b>Germ Cell Tumors</b>		
Testicular germ cell tumor	Yes	No
Extragenadal germ cell tumor	Yes	No
Seminoma	Yes	No
Choriocarcinoma	Yes	No
Embryonal carcinoma	Yes	No
Mixed germ cell tumors	Yes	No
Teratoma	Yes	No
Yolk-sac tumor (endodermal sinus tumor)	Yes	No
Germ cell tumor of the ovary	Yes	No

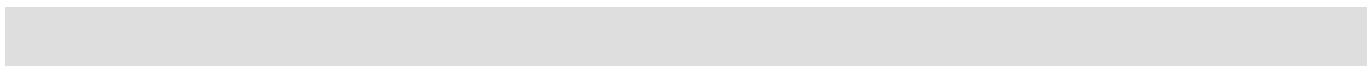
### **Multiple Myeloma/ Plasma Cell Disorders**

<b>Multiple Myeloma</b>		
Single Auto	Yes	No
Tandem (auto followed by auto)	Yes	No
Tandem (auto followed by allo)	Yes	No
Allogenic	No	Yes
AL-Amyloidosis	Yes	No
Waldenstrom macroglobulinemia	Yes	Yes
Monoclonal gammopathy of renal significance (MGRS)	Yes	No
Monoclonal gammopathy of uncertain significance (MGUS)	No	No
Polyneuropathy organomegaly endocrinopathy, monoclonal gammopathy skin defects syndrome (POEMS)	Yes	No
Solitary Plasmacytoma	No	No

<b>Hodgkin Lymphoma</b>		
Hodgkin Lymphoma	Yes	Yes
<b>Disease/Indication</b>		
	<b>Autologous</b>	<b>Allogeneic</b>
<b>Non-Hodgkin Lymphoma (NHL)</b>		
Small B-cell lymphocytic lymphoma	No	Yes
Follicular lymphoma	Yes	Yes
Lymphoplasmacytoid lymphoma/immunocytoma	Yes	Yes
Marginal zone lymphoma (mucosa-associated lymphoid tissue, splenic, nodal)	Yes	Yes
Burkitt lymphoma	Yes	Yes
Diffuse, large cell lymphoma (mediastinal large cell, primary effusion)	Yes	Yes
Mantle cell lymphoma	Yes	Yes
Precursor B-cell leukemia/lymphoma	Yes	Yes
T-cell Lymphoma	Yes	Yes
<b>Other Malignancies</b>		
Atypical teratoid rhabdoid tumors	Yes	No
Blastic plasmacytoid dendritic cell neoplasm	No	Yes
Epithelial ovarian cancer	No	No
Ewing tumor (Ewing sarcoma)	Yes	No
Neuroblastoma	Yes	No
Osteogenic sarcoma	No	No
Renal cell carcinoma	No	No
Retinoblastoma	Yes	No
Rhabdomyosarcoma/soft tissue sarcoma	No	No
Supratentorial ependymoma	Yes	No
Wilms tumor	Yes	No



<b>Hematological Disorders</b>	<b>Autologous</b>	<b>Allogeneic</b>
Aplastic Anemia	No	Yes
Blackfan-Diamond Syndrome	No	Yes
Chronic Granulomatous Disease	No	Yes
Congenital Agranulocytosis (Kostmann Syndrome)	No	Yes
Congenital Amegakaryocytic Thrombocytopenia	No	Yes
Dyskeratosis Congenita	No	Yes
Fanconi Anemia	No	Yes
Paroxysmal Nocturnal Hemoglobinuria (PNH)	No	Yes
Shwachman-Diamond Syndrome	No	Yes
Sickle Cell Disease (SCD)	No	Yes
Thalassemia Major	No	Yes



<b>Immunodeficiency Syndromes</b>		
CD40 ligand deficiency	No	Yes
Chediak-Higashi syndrome	No	Yes
Hemophagocytic Lymphohistiocytosis (HLH) (same as familial Erythrophagocytic lymphohistiocytosis - FEL)	No	Yes
Leukocyte adhesion deficiency	No	Yes
Omenn syndrome	No	Yes
Severe Combined Immunodeficiency Disease (SCID)	No	Yes
Wiskott-Aldrich syndrome	No	Yes
X-linked lymphoproliferative syndrome	No	Yes
Gaucher disease type 1	No	Yes
Niemann-Pick type B	No	Yes
Fucosidosis	No	Yes
Lysosomal storage diseases	No	Yes



### Autoimmune Diseases

Crohn's disease	No	No
Multiple sclerosis	Yes	No
Rheumatoid arthritis	No	No
Systemic lupus erythematosus (SLE)	No	No
Systemic sclerosis (Scleroderma)	Yes	No

### Inherited Metabolic Disorders

Adrenoleukodystrophy	No	Yes
Epidermolysis bullosa	No	Yes
Globoid cell leukodystrophy (Krabbe Disease)	No	Yes
Hurler syndrome (MPS I)	No	Yes
Hunter syndrome (MPS II)	No	Yes
Mannosidosis	No	Yes
Maroteaux-Lamy Syndrome (MPS VI)	No	Yes
Metachromatic leukodystrophy	No	Yes
Mitochondrial neurogastrointestinal encephalopathy (MNGIE)	No	Yes
Osteopetrosis	No	Yes
Rett syndrome	No	Yes

### Cardiac Conditions

Heart disease	No	No
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### Additional condition/disease indications

Refer to section titled: Hematopoietic Stem Cell Transplant Reference Sheet Error! Reference source not found.	No	Yes
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# Review and Approval History

Version	Date and Description
1.0	07/19/2012: New guideline. Approved by Medical Technology Assessment Committee
1.0	08/14/2012: Approved by National Medical Care Management Committee
2.0	10/10/13: Revised and updated. Approved by Medical Technology Assessment Committee
2.0	10/16/2013: Approved by Complex Medical Conditions Policy Committee
2.0	11/12/2013: Approved by the National Medical Care Management Committee
3.0	08/07/2014: Approved by Medical Technology Assessment Committee
3.0	09/09/2014: Approved by National Medical Care Management Committee
4.0	8/25/2015: Annual review; revised and updated
4.0	09/03/2015: Approved by Medical Technology Assessment Committee
4.0	10/13/2015: Approved by National Medical Care Management Committee
5.0	08/15/2016: Annual review. Revised and updated. Transplant Review Guidelines separated into two documents: Hematopoietic Stem Cell Transplantation and Solid Organ Transplantation.
5.0	09/01/2016: Approved by Medical Technology Assessment Committee
5.0	09/13/2016: Approved by National Medical Care Management Committee
6.0	6/22/2017: Approved by Optum Policy and Guideline Committee
6.0	07/06/2017: Approved by Medical Technology Assessment Committee
6.0	07/11/2017: Approved by National Medical Care Management Committee
7.0	09/07/2017: New content relevant to CAR-T Therapy approved by Medical Technology Assessment Committee.
7.0	09/12/2017: New content relevant to CAR-T Therapy approved by National Medical Care Management Committee.
7.0	11/1/2017: Updated to reflect FDA-approval of new CAR-T Therapy agent axicabtagene ciloleucel (Yescarta™)
7.0	11/13/2017: Corrected CAR-T prior authorization statement on page 7.
8.0	08/02/2018: Approved by Medical Technology Assessment Committee
8.0	09/11/2018: Approved by National Medical Care Management Committee
9.0	4/17/2019: Annual review with Optum Stem Cell Expert Panel. Minor revisions including addition of CMML to approved indications for allogeneic stem cell transplant; revised the preferred scoring system for primary myelofibrosis; revised systemic sclerosis indication to approve autologous transplant; added allogeneic transplant evaluation for secondary myelofibrosis in patients with polycythemia vera and essential thrombocytopenia; and added DIPSS-Plus factors table and scoring directions. Updated references.
9.0	06/06/2019: Approved by Medical Technology Assessment Committee

9.0	06/11/2019: Approved by National Medical Care Management Committee
9.0	12/2/2019: Corrected follicular lymphoma indication on page 10. Updated supporting references.
10.0	6/10/2020: Annual review with Optum Stem Cell Expert Panel. Revisions to the MRD statement, Relative Contraindications and Special Considerations sections, and NMDP recommendations for timing of transplant consultation. References updated throughout.
10.0	8/6/2020: Approved by Medical Technology Assessment Committee
10.0	8/11/2020: Presented to National Medical Care Management Committee
10.0	11/11/2020: Updated minimal/measurable disease terminology
11.0	6/15/2021: Annual Review with Optum Stem Cell Transplantation Expert Panel. No revisions.
11.0	9/9/2021: Approved by Medical Technology Assessment Committee
11.0	9/14/2021: Presented to National Medical Care Management Committee
12.0	7/29/2022: Annual Review with Optum Stem Cell Transplantation Expert Panel. Added link to American Society of Hematology Stem Cell Transplantation in Sickle Cell Disease Guideline. Updated references.
12.0	9/1/2022: Approved by Medical Technology Assessment Committee
12.0	9/8/2022: Presented to National Medical Care Management Committee
12.0	1/5/2023: Interim update. Added criteria for autologous HSCT in patients with monoclonal gammopathy of renal significance (MGRS). Approved by Medical Technology Assessment Committee.
12.0	1/10/2023: Presented to National Medical Care Management Committee
13.0	7/12/2023: Annual Review with Optum Stem Cell Transplantation, Chimeric Antigen Receptor T-cell Therapy, and Gene Therapy Expert Panel. Added literature review and medical necessity criteria for Omidubicel only; updated NMDP/ASBMT Recommended Timing for Stem Cell Transplantation Consultation.
13.0	9/11/2023: Approved by Optum Clinical Guideline Advisory Committee
13.0	10/05/2023: Approved by Medical Technology Assessment Committee
13.0	11/17/2023: Approved by the Pharmacy and Therapeutics (P&T) Committee
14.0	8/28/2024: Annual review with Optum Hematopoietic Stem Cell Transplant and Chimeric Antigen Receptor T-Cell Therapy Expert Panel . Guideline reformatted. Updated NMDP/ASTCT Recommended Timing for Transplant Consultation.
14.0	10/09/2024: Approved by Optum Clinical Guideline Advisory Committee
14.0	11/07/2024: Approved by Medical Technology Assessment Committee