Hematopoietic Stem-Cell Transplantation for Hodgkin Lymphoma

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Line(s) of Business: HMO; PPO
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Section: Transplants
Place(s) of Service: Outpatient, Inpatient

Precertification is required for this service.

I. Description

Hematopoietic Stem Cell Transplantation

Hematopoietic stem-cell transplantation (HSCT) refers to a procedure in which hematopoietic stem cells are infused to restore bone marrow function in cancer patients who receive bone-marrow-toxic doses of cytotoxic drugs with or without whole-body radiation therapy. Hematopoietic stem cells may be obtained from the transplant recipient (autologous HSCT) or from a donor (allogeneic HSCT). They can be harvested from bone marrow, peripheral blood, or umbilical cord blood shortly after delivery of neonates. Although cord blood is an allogeneic source, the stem cells in it are antigenically “naïve” and thus are associated with a lower incidence of rejection or graft-versus-host disease (GVHD).

Immunologic compatibility between infused hematopoietic stem cells and the recipient is not an issue in autologous HSCT. However, immunologic compatibility between donor and patient is a critical factor for achieving a good outcome of allogeneic HSCT. Compatibility is established by typing of human leukocyte antigens (HLA) using cellular, serologic, or molecular techniques. HLA refers to the tissue type expressed at the Class I and Class II loci on chromosome 6. Depending on the disease being treated, an acceptable donor will match the patient at all or most of the HLA loci (with the exception of umbilical cord blood).

Conventional Preparative Conditioning for HSCT

The success of autologous HSCT is predicated on the ability of cytotoxic chemotherapy with or without radiation to eradicate cancerous cells from the blood and bone marrow. This permits subsequent engraftment and repopulation of bone marrow space with presumably normal hematopoietic stem cells obtained from the patient prior to undergoing bone marrow ablation. As a consequence, autologous HSCT is typically performed as consolidation therapy when the patient’s
disease is in complete remission. Patients who undergo autologous HSCT are susceptible to chemotherapy-related toxicities and opportunistic infections prior to engraftment, but not GVHD.

The conventional (“classical”) practice of allogeneic HSCT involves administration of cytotoxic agents (eg, cyclophosphamide, busulfan) with or without total-body irradiation at doses sufficient to destroy endogenous hematopoietic capability in the recipient. The beneficial treatment effect in this procedure is due to a combination of initial eradication of malignant cells and subsequent graft-versus-malignancy (GVM) effect mediated by non self immunologic effector cells that develop after engraftment of allogeneic stem cells within the patient’s bone marrow space. While the slower GVM effect is considered to be the potentially curative component, it may be overwhelmed by extant disease without the use of pretransplant conditioning. However, intense conditioning regimens are limited to patients who are sufficiently fit medically to tolerate substantial adverse effects that include pre-engraftment opportunistic infections secondary to loss of endogenous bone marrow function and organ damage and failure caused by the cytotoxic drugs. Furthermore, in any allogeneic HSCT, immunosuppressant drugs are required to minimize graft rejection and GVHD, which also increases susceptibility of the patient to opportunistic infections.

Reduced-Intensity Conditioning for Allogeneic HSCT

Reduced-intensity conditioning (RIC) refers to the pretransplant use of lower doses or less intense regimens of cytotoxic drugs or radiation than are used in traditional full-dose myeloablative conditioning treatments. The goal of RIC is to reduce disease burden but also to minimize as much as possible associated treatment-related morbidity and non-relapse mortality (NRM) in the period during which the beneficial GVM effect of allogeneic transplantation develops. Although the definition of RIC remains arbitrary, with numerous versions employed, all seek to balance the competing effects of NRM and relapse due to residual disease. RIC regimens can be viewed as a continuum in effects, from nearly totally myeloablative to minimally myeloablative with lymphoablation, with intensity tailored to specific diseases and patient condition. Patients who undergo RIC with allogeneic HSCT initially demonstrate donor-cell engraftment and bone-marrow mixed chimerism. Most will subsequently convert to full-donor chimerism, which may be supplemented with donor lymphocyte infusions to eradicate residual malignant cells.

For the purposes of this Policy, the term RIC will refer to all conditioning regimens intended to be non-myeloablative, as opposed to fully myeloablative (traditional) regimens.

Hodgkin Lymphoma

Hodgkin lymphoma (HL) is a relatively uncommon B cell lymphoma. In 2011, the estimated number of cases in the U.S. was approximately 8,830 new diagnoses and 1,300 deaths. (1) The disease has a bimodal distribution, with most patients diagnosed between the ages of 15 and 30 years, with a second peak in adults aged 55 years and older.

The 2008 World Health Organization (WHO) classification divides HL into 2 main types (2):

A. “Classical” HL (CHL)
   o Nodular sclerosis
o Mixed cellularity
o Lymphocyte depleted
o Lymphocyte rich

B. Nodular lymphocyte-Predominant HL (NLPHL)

In Western countries, CHL accounts for 95% of cases of HL and NLPHL only 5%. (3) Classic HL is characterized by the presence of neoplastic Reed-Sternberg cells in a background of numerous non-neoplastic inflammatory cells. NLPHL lacks Reed-Sternberg cells but is characterized by the presence of lymphocytic and histiocytic cells termed “popcorn cells.” (3)

The following staging system for HL recognizes that the disease is thought to typically arise in a single lymph node and spread to contiguous lymph nodes with eventual involvement of extranodal sites. The staging system attempts to distinguish patients with localized HL who can be treated with extended field radiation from those who require systemic chemotherapy.

Staging for Hodgkin Lymphoma

Staging for HL is based on the Ann Arbor staging system. Each stage is subdivided into A and B categories. “A” indicates no systemic symptoms are present, and “B” indicates the presence of systemic symptoms, including unexplained weight loss of more than 10% of body weight, unexplained fevers, or drenching night sweats. (3)

Stage I
- Involvement of a single lymph node region (I) or localized involvement of a single extralymphatic organ or site (IE)

Stage II
- Involvement of 2 or more lymph node regions on the same side of the diaphragm (II) or localized involvement of a single associated extralymphatic organ or site and its regional lymph node(s) with or without involvement of other lymph node regions on the same side of the diaphragm (IIE). The number of lymph node regions involved should be indicated by a subscript (eg, II2).

Stage III
- Involvement of lymph node regions or structures on both sides of the diaphragm. These patients are further subdivided as follows:
  - III-1: disease limited to spleen or upper abdomen
  - III-2: periaortic or pelvic node involvement

Stage IV
- Disseminated (multifocal) involvement of one or more extralymphatic organs, with or without associated lymph node involvement, or isolated extralymphatic organ involvement with distant (nonregional) nodal involvement.
Patients with HL are generally classified into 3 groups: early-stage favorable (stage I–II with no B symptoms or large mediastinal lymphadenopathy), early-stage unfavorable (stage I–II with large mediastinal mass, with or without B symptoms; stage IB–IIB with bulky disease), and advanced-stage disease (stage III–IV). (3)

Patients with nonbulky stage IA or IIA disease are considered to have clinical early-stage disease. These patients are candidates for chemotherapy, combined modality therapy, or radiation therapy alone. (1) Patients with obvious stage III or IV disease, bulky disease (defined as a 10-cm mass or mediastinal disease with a transverse diameter exceeding 33% of the transthoracic diameter), or the presence of B symptoms will require combination chemotherapy with or without additional radiation therapy. (1)

HL is highly responsive to conventional chemotherapy, and up to 80% of newly diagnosed patients can be cured with combination chemotherapy and/or radiation therapy. Patients who prove refractory or who relapse after first-line therapy have a significantly worse prognosis. Primary refractory HL is defined as disease regression of less than 50% after 4 to 6 cycles of anthracycline-containing chemotherapy, disease progression during induction therapy, or progression within 90 days after the completion of first-line treatment. (4)

In patients with relapse, the results of salvage therapy vary depending on a number of prognostic factors, as follows: the length of the initial remission, stage at recurrence, and the severity of anemia at the time of relapse. (5) Early and late relapse are defined as less or more than 12 months from the time of remission, respectively. Approximately 70% of patients with late first relapse can be salvaged by autologous HSCT but not more than 40% with early first relapse. (6)

Only approximately 25 to 35% of patients with primary progressive or poor-risk recurrent HL achieve durable remission after autologous HSCT, with most failures being due to disease progression after transplant. Most relapses after transplant occur within 1 to 2 years, and once relapse occurs posttransplant, median survival is less than 12 months.

II.  Policy

A.  Autologous or myeloablative allogeneic hematopoietic stem-cell transplantation (HSCT) is covered (subject to Limitation/Exclusions and Administrative Guidelines) for patients with primary refractory or relapsed Hodgkin lymphoma (HL).

B.  Tandem autologous HSCT is covered (subject to Limitation/Exclusions and Administrative Guidelines):
   1.  In patients with primary refractory HL; or
   2.  In patients with relapsed disease with poor risk features who do not attain a complete remission to cytoreductive chemotherapy prior to transplantation (see Policy Guidelines).

C.  Reduced-intensity allogeneic HSCT is covered (subject to Limitation/Exclusions and Administrative Guidelines) to treat HL in patients:
   1.  Who have failed a prior autologous HSCT used to treat primary refractory or relapsed disease; or
2. In patients who would otherwise qualify for a myeloablative allogeneic transplant, but would be unable to tolerate a standard myeloablative conditioning regimen (see Policy Guidelines); or

3. When insufficient stem cells are collected for an autologous HSCT.

III. Policy Guidelines

In the Morschhauser et al study of risk-adapted salvage treatment with single or tandem autologous hematopoietic stem-cell transplantation (HSCT) for first relapse or refractory Hodgkin lymphoma (HL), (7) poor-risk relapsed HL was defined as 2 or more of the following risk factors at first relapse: time to relapse less than 12 months, stage III or IV at relapse, and relapse within previously irradiated sites. Primary refractory disease was defined as disease regression less than 50% after 4 to 6 cycles of doxorubicin-containing chemotherapy or disease progression during induction or within 90 days after the end of first-line treatment.

Some patients for whom a conventional myeloablative allotransplant could be curative may be considered candidates for reduced-intensity conditioning (RIC) allogeneic HSCT. These include those with malignancies that are effectively treated with myeloablative allogeneic transplantation, but whose age (typically older than 55 years) or comorbidities (eg, liver or kidney dysfunction, generalized debilitation, prior intensive chemotherapy, low Karnofsky Performance Status) preclude use of a standard myeloablative conditioning regimen.

The ideal allogeneic donors are human leukocyte antigen (HLA)-identical matched siblings. Related donors mismatched at 1 locus are also considered suitable donors. A matched, unrelated donor identified through the National Marrow Donor Registry is typically the next option considered. Recently, there has been interest in haploidentical donors, typically a parent or a child of the patient, with whom usually there is sharing of only 3 of the 6 major histocompatibility antigens. The majority of patients will have such a donor; however, the risk of GVHD and overall morbidity of the procedure may be severe, and experience with these donors is not as extensive as that with matched donors.

In 2003, CPT centralized codes describing allogeneic and autologous hematopoietic stem-cell support services to the hematology section (CPT 38204-38242). Not all codes are applicable for each HDC/stem-cell support procedure. For example, Plans should determine if cryopreservation is performed. A range of codes describes services associated with cryopreservation, storage, and thawing of cells (38208-38215).

CPT 38208 and 38209 describe thawing and washing of cryopreserved cells

CPT 38210-38214 describe certain cell types being depleted

CPT 38215 describes plasma cell concentration
IV. Limitations/Exclusions

A. A second autologous stem-cell transplantation for relapsed lymphoma after a prior autologous HSCT is not covered as it is not known to be effective in improving health outcomes.

B. Other uses of HSCT in patients with HL, including, but not limited to, initial therapy for newly diagnosed disease to consolidate a first complete remission, are not covered as they are not known to be effective in improving health outcomes.

V. Administrative Guidelines

Precertification is required for a transplant evaluation and for the transplant itself and should be submitted by the proposed treating facility. To precertify, please complete HMSA’s Precertification Request and mail or fax the form as indicated along with the required documentation.

<table>
<thead>
<tr>
<th>CPT Codes</th>
<th>Description</th>
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<tbody>
<tr>
<td>38205</td>
<td>Blood-derived hematopoietic progenitor cell harvesting for transplantation, per collection, allogeneic</td>
</tr>
<tr>
<td>38206</td>
<td>Blood-derived hematopoietic progenitor cell harvesting for transplantation, per collection, autologous</td>
</tr>
<tr>
<td>38208</td>
<td>Transplant preparation of hematopoietic progenitor cells; thawing of previously frozen harvest, without washing, per donor</td>
</tr>
<tr>
<td>38209</td>
<td>;thawing of previously frozen harvest, with washing, per donor</td>
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<tr>
<td>38210</td>
<td>;specific cell depletion with harvest, T cell depletion</td>
</tr>
<tr>
<td>38211</td>
<td>;tumor cell depletion</td>
</tr>
<tr>
<td>38212</td>
<td>;red blood cell removal</td>
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<tr>
<td>38213</td>
<td>;platelet depletion</td>
</tr>
<tr>
<td>38214</td>
<td>;plasma (volume) depletion</td>
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<tr>
<td>38215</td>
<td>;cell concentration in plasma, mononuclear, or buffy coat layer</td>
</tr>
<tr>
<td>38220</td>
<td>Bone marrow; aspiration only</td>
</tr>
<tr>
<td>38221</td>
<td>Bone marrow; biopsy, needle or trocar</td>
</tr>
<tr>
<td>38230</td>
<td>Bone marrow harvesting for transplantation; allogeneic</td>
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<tr>
<td>38232</td>
<td>Bone marrow harvesting for transplantation; autologous</td>
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<tr>
<td>38241</td>
<td>;autologous transplantation</td>
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<td>ICD-9 Procedure Codes</td>
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<td>----------------------</td>
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<tr>
<td>41.00</td>
<td>Bone marrow transplant, not otherwise specified</td>
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<td>41.01</td>
<td>Autologous bone marrow transplant, without purging</td>
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<td>41.03</td>
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<td>41.04</td>
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<tr>
<td>410.5</td>
<td>Allogeneic hematopoietic stem-cell transplant, without purging</td>
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<td>41.06</td>
<td>Cord blood stem-cell transplant</td>
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<td>41.07</td>
<td>Autologous hematopoietic stem-cell transplant with purging</td>
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<td>Autologous bone marrow transplant with purging</td>
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<td>41.91</td>
<td>Aspiration of bone marrow from donor for transplant</td>
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<tr>
<td>99.79</td>
<td>Other therapeutic apheresis (includes harvest of stem cells)</td>
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<table>
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<tr>
<th>HCPCS Codes</th>
<th>Description</th>
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<tr>
<td>G0265</td>
<td>Cryopreservation, freezing and storage of cells for therapeutic use, each cell line</td>
</tr>
<tr>
<td>G0266</td>
<td>Thawing and expansion of frozen cells for therapeutic use, each cell line</td>
</tr>
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<td>G0267</td>
<td>Bone marrow or peripheral stem-cell harvest, modification or treatment to eliminate cell type(s) (e.g., T cells, metastatic carcinoma)</td>
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<td>Q0083 - Q0085</td>
<td>Chemotherapy, administration code range</td>
</tr>
<tr>
<td>J9000 - J9999</td>
<td>Chemotherapy drug code range</td>
</tr>
<tr>
<td>S2140</td>
<td>Cord blood harvesting for transplantation, allogeneic</td>
</tr>
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<td>S2142</td>
<td>Cord blood-derived stem-cell transplantation, allogeneic</td>
</tr>
<tr>
<td>S2150</td>
<td>Bone marrow or blood-derived peripheral stem-cell harvesting and transplantation, allogeneic or autologous, including pheresis, high-dose chemotherapy, and the number of days of post-transplant care in the global definition (including drugs; hospitalization; medical surgical, diagnostic and emergency services)</td>
</tr>
</tbody>
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ICD-10 codes are provided for your information. These will not become effective until 10/01/2014.

<table>
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<th>ICD-10-PCS</th>
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<td>30243G0, 30243X0, 30243Y0</td>
<td>Percutaneous transfusion, central vein, bone marrow or stem cells, autologous, code list</td>
</tr>
<tr>
<td>30243G1, 30243X1, 30243Y1</td>
<td>Percutaneous transfusion, central vein, bone marrow or stem cells, nonautologous, code list</td>
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<tr>
<td>07DQ0ZZ, 07DQ3ZZ, 07DR0ZZ, 07DR3ZZ, 07DS0ZZ, 07DS3ZZ</td>
<td>Surgical, lymphatic and hemic systems, extraction, bone marrow, code list</td>
</tr>
</tbody>
</table>

VI. Scientific Background

This policy has been updated regularly with searches of the MEDLINE database. The most recent literature search, including EMBASE, was performed for the period September 1, 2012 through October 15, 2013.

Autologous hematopoietic stem-cell transplantation for front-line therapy of Hodgkin lymphoma

A study published by Federico et al. concluded that high-dose chemotherapy (HDC) with autologous hematopoietic stem-cell transplantation (HSCT) offered no benefit in outcomes over conventional chemotherapy in front-line therapy for advanced Hodgkin lymphoma (HL) patients. (8)

Carella et al. reported the long-term results of 163 patients with unfavorable HL who had received either an autologous HSCT or additional standard chemotherapy for consolidation after initial conventional chemotherapy. (9) Patients were randomly assigned to receive HDC followed by an autologous HSCT (n=83) or 4 additional courses of the same standard chemotherapy used in the induction phase (n=80). After treatment, complete remission (CR) was achieved in 92% of patients in the autologous HSCT arm and 89% in the standard chemotherapy arm (p=0.6). Five-year overall survival (OS) was 88% (95% confidence interval [CI], 80 to 96%) in the autologous HSCT arm and 88% (95% CI, 79 to 96%) in the CT arm (p=0.99). Ten-year OS was 85% (95% CI, 78 to 90%) versus 84% (95% CI, 77 to 89%) for the autologous HSCT versus the standard chemotherapy group, respectively. The authors concluded that, after a median follow-up of 107 months, their data supported that patients who respond to induction therapy with conventional chemotherapy do not achieve superior outcomes with consolidation with HDC and autologous HSCT.

Autologous HSCT for relapsed/refractory HL

Autologous HSCT is widely considered the therapy of choice for relapsed and refractory HL. Two randomized, controlled trials showed benefit in using autologous HSCT in these patients:

The British National Lymphoma Investigation (BNLI) study was the first to show a progression-free survival (PFS) benefit with autologous HSCT over conventional chemotherapy in relapsed or refractory HL patients. (10) Forty patients with relapsed or refractory HL were given chemotherapy without transplant (n=20) or autologous transplant after HDC (n=20). (11) A significantly better
event-free survival (EFS) at 3 years of 53% versus 10% was reported in the patients who underwent transplant versus the group that did not.

Subsequently, these findings were confirmed in a larger trial by the German Hodgkin Study Group (GHSG) and European Group for Blood and Marrow Transplantation (EBMT). (12) Patients relapsing after initial chemotherapy were randomly assigned to chemotherapy without transplant or to autologous HSCT. In the final analysis of 144 patients, freedom from treatment failure at 3 years was 55% in the transplanted group versus 34% in the nontransplanted group. This benefit was maintained in subgroup analysis, regardless of early or late relapse, and the results were confirmed in follow-up data at 7 years. (13)

Several large retrospective studies have reported EFS rates ranging from 25 to 60%, with OS rates from 35 to 66%, showing that disease status before autologous HSCT was the most important prognostic factor for the final outcome. (4)

Limited treatment options exist for patients who relapse following an autologous HSCT and include single-agent palliative chemotherapy or occasionally, localized radiation therapy. (13) When a further remission may be attained with conventional-dose chemotherapy, it is rarely durable, with a median OS of less than 1 year. (14) There is limited experience with second autologous HSCT, and treatment-related mortality is high (25 to 40%). (11) Smith et al. reported the outcomes of 40 patients (21 with HL and 19 with non-Hodgkin lymphoma [NHL]) who underwent a second autologous HSCT for relapsed lymphoma. (15) Results reported were combined for the two populations, but the authors state that the outcomes of patients with HL and NHL were similar. Median age at second HSCT was 38 years (range, 16 to 61). The second HSCT was performed more than 1 year after the first in 82%. Treatment-related mortality at day 100 post-transplant was 11% (95% CI, 3 to 22%). At a median follow-up of 72 months (range, 12 to 124 months) after the second HSCT, 73% of patients had died, 62% of these due to relapsed lymphoma. One-, 3-, and 5-year PFS probabilities were 50% (95% CI, 34 to 66%), 36% (95% CI, 21 to 52%), and 30% (95% CI, 16 to 46%), respectively. Corresponding OS probabilities were 65% (95% CI, 50 to 79%), 36% (95% CI, 22 to 52%), and 30% (95% CI, 17 to 46%), respectively. The authors stated that limitations to their study included the absence of an appropriate comparison group and that it was not known how many patients were considered for a second HSCT but were unable to mobilize sufficient stem cells or were otherwise unable to proceed to the second transplant. Finally, they stated that the heterogeneity of the preparative regimens used in this population precluded comparison of efficacy.

Allogeneic HSCT for HL

The application of allogeneic HSCT to the treatment of patients with HL initially appeared limited due to a procedure-related mortality rate of approximately 50% associated with the myeloablative conditioning regimen.

To date, most of the RIC allogeneic HSCTs have been performed in patients who have failed a previous autologous HSCT for primary relapsed/refractory HL, and most of the studies are
characterized by small numbers of patients, disparate preparative and graft-versus-host disease (GVHD) prophylaxis regimens, and varying lengths of follow-up. Nonetheless, they have demonstrated reduced nonrelapse mortality, and some suggest a graft-versus-HL effect with favorable disease control in these poor-prognosis patients.

Sarina et al. reported a retrospective study of 185 patients with HL who had failed an autologous HSCT. (16) One hundred twenty-two had donors available for a salvage RIC allogeneic HSCT; of these, 104 (85%) were transplanted. Sixty-three patients did not have a suitable donor and were treated with salvage chemotherapy or radiotherapy. Clinical characteristics between the 2 groups did not differ. After a median follow-up of 48 months, PFS and OS were better in the group that underwent the salvage allogeneic HSCT (39.3% vs. 14.2% and 66% vs. 42%, respectively; p<0.001), showing a survival benefit of an RIC allogeneic HSCT versus conventional treatment after a failed autologous HSCT for HL. This study supports one of the policy statements for RIC HSCT.

Peggs et al. investigated outcomes with RIC allogeneic HSCT, and T cell depletion in multiply relapsed patients. (17) Forty-nine patients were enrolled, 90% of whom had failed a previous autologous transplant. Primary study endpoints were engraftment, toxicity, non-relapse-related mortality (NRM), and GVHD incidence. All patients achieved engraftment. Thirty-one patients had an HLA-matched donor and 18, an unrelated donor. The cumulative incidence of NRM was 4.1% at 100 days posttransplant and 16.3% at 730 days posttransplant. Patients with unrelated donors had a significantly higher NRM (34% vs 7%) at 730 days. Projected 4-year OS and PFS were 56% and 39%, respectively.

Alvarez et al. reported the results of a Spanish Cooperative Protocol using RIC allogeneic HSCT in 40 patients with relapsed or refractory HL. (18) Seventy-three percent of patients had failed a previous autologous HSCT. Thirty-eight patients received hematopoietic cells from an HLA-identical sibling. One-year treatment-related mortality was 25%. OS and PFS were 48% and 32%, respectively, at 2 years. For patients who had failed a previous autologous HSCT, 2-year OS and PFS were 75% and 70%, respectively, in the subset that relapsed more than 12 months after autologous HSCT.

Todisco et al. evaluated the efficacy of RIC allogeneic HSCT in 14 patients with refractory or progressive HL after HDC and autologous HSCT. (14) All of the patients had received at least 1 prior course of HDC, and 50% had undergone 2 previous courses. The median time from the first and second courses of HDC and the RIC allogeneic HSCT was 15 and 8 months, respectively (range 2 to 34 and 2 to 31 months, also respectively). With a median follow-up of 21 months post-RIC allogeneic HSCT (range 3 to 74 months), 10 of the 14 patients were alive. Estimated OS at 1 and 2 years was 93% and 73%, respectively, for the entire population; 83% and 44%, respectively, for patients with chemotherapy-resistant disease; and 100% for those with chemotherapy-sensitive disease.

A review of the role of allogeneic HSCT in HL by Laport (19) summarizes the results of the most recent and largest studies of the use of RIC allogeneic HSCT for HL as follows: most patients have failed a prior autologous HSCT and are therefore heavily pretreated going into the RIC allogeneic HSCT; chemotherapy sensitivity is a reliable predictor of outcome; a matched versus an unmatched
related donor did not affect survival in most reports; and approximately one third to one half of these patients may be cured with RIC allogeneic HSCT.

EBMT published the results of the outcomes of 89 HL patients with relapsed or refractory disease who received an RIC allogeneic HSCT and were compared to 79 patients who received myeloablative conditioning. (20) Sixty-two percent of the RIC group had undergone a previous autologous HSCT versus 41% of the patients in the myeloablative group. Although the incidence of relapse was nearly double in the RIC group (57% vs. 30%, respectively), after a median follow-up for surviving patients of 75 months (range, 12-120 months), 24 in the RIC group (26.9%) and 18 in the conventional group (22.8%) were alive. Five-year OS was 22% (95% CI, 13 to 31%) for the conventional group and 28% (95% CI, 18 to 38%) for the RIC group. Independent adverse prognostic factors for OS were a previously failed autologous HSCT (risk ratio [RR], 1.59; 95% CI, 1.07 to 2.35; p=0.02), the use of myeloablative conditioning (RR: 1.62; 95% CI, 1.27 to 3.29; p=0.04), and the presence of refractory disease (RR: 1.51; 95% CI, 1.03 to 2.21; p=.003).

Anderlini et al. published the results of 58 patients from 1 institution with relapsed/refractory HL who received uniform conditioning regimens for RIC allogeneic HSCT. (21) Fifty-seven percent of patients received their allograft from an unrelated donor. Eighty-three percent of patients had failed a prior autologous HSCT. Projected 2-year OS and PFS rates were 64% (range, 49 to 76%) and 32% (range, 20% to 45%), with 2-year disease progression/relapse at 55% (range, 43% to 70%). There were no statistically significant differences in OS, PFS, or disease progression/relapse between matched related and unrelated donor transplants.

Sureda et al. (22) reported the results of a Phase II study of 92 patients with relapsed HL and an human leukocyte antigen (HLA)-identical sibling, a matched unrelated donor, or a 1 antigen mismatched, unrelated donor who were treated with salvage chemotherapy followed by RIC allogeneic transplantation. Fourteen patients had refractory disease and died from progressive lymphoma with a median OS after trial entry of 10 months (range, 6-17 months). Seventy-eight patients proceeded to allograft (unrelated donors, n=23). Fifty were allografted in complete or partial remission and 28 in stable disease. Nonrelapse mortality rate was 8% at 100 days and 15% at 1 year. Relapse was the major cause of failure. The PFS rate was 47% at 1 year and 18% at 4 years from trial entry. For the allografted population, the PFS rate was 48% at 1 year and 24% at 4 years. Chronic GVHD was associated with a lower incidence of relapse. Patients allografted in CR had a significantly better outcome. The OS rate was 71% at 1 year and 43% at 4 years.

**Tandem (autologous-autologous) HSCT**

Fung et al. reported results from a pilot study to evaluate the toxicities and efficacy of tandem autologous HSCT in patients with primary refractory or poor risk recurrent HL. (23) The study involved 28 patients with primary progressive and 18 with recurrent HL who were enrolled in the study between April 1998 and March 2000. Patients had at least one of the following poor prognostic factors: first CR less than 12 months, extranodal disease, or B symptoms at relapse. Forty-one patients (89%) received the second transplant. With a median follow-up of 5.3 years
(range, 1.6-8.1), the 5-year OS and PFS were 54% (95% CI, 40 to 69%) and 49% (95% CI, 34 to 63%), respectively.

Morschhauser et al. reported on the results of a multicenter prospective trial that evaluated a risk-adapted salvage treatment with single or tandem autologous HSCT in 245 patients with relapsed/refractory HL. (7) Median follow-up time was 51 months (range, 20–110 months). Patients who were categorized as poor risk (n=150) had primary refractory disease (n=77) or 2 or more of the following risk factors at first relapse: time to relapse less than 12 months, stage III or IV disease at the time of relapse, or relapse occurring within previously irradiated sites (n=73). In this study, these poor-risk patients were eligible for tandem autologous transplants. Intermediate-risk patients (n=95), defined as 1 risk factor at relapse, were eligible for a single transplant. Overall, 70% of the poor-risk patients received tandem transplants, and 97% of the intermediate-risk patients received a single transplant.

Overall, 94 poor-risk patients responded to cytoreductive chemotherapy (partial response [PR] or CR), whereas 55 patients had chemotherapy-resistant disease. A total of 137 patients (including the 94 patients with chemotherapy-sensitive disease and 43 of 55 with chemotherapy–resistant disease) received the first autologous HSCT. Among 121 patients who were fully restaged, 64 patients had achieved a CR, 37 a PR, and 4 had stable disease. These 105 patients then underwent the second autologous HSCT after a median of 65 days. Among them, 80 patients achieved a CR, including 17 patients who had achieved PR, and 3 patients with stable disease after the first transplant. Among the 55 patients who had cytoreduction failure, 30 responded to the first transplant (9 with CR), and 17 achieved CR after the second transplant.

Outcome analysis based on the intent-to-treat sample showed 5-year freedom from second failure and OS were 73% and 85% for the intermediate-risk group and 46% and 57% for the poor-risk group, all respectively.

In the poor-risk group, patients who underwent tandem transplant and had a complete response to cytoreduction chemotherapy did not have superior outcomes compared to complete responders receiving a single transplant in previous studies. (24) However, in this study, poor-risk patients who were partial responders who underwent tandem transplants did better when compared to partial responders who received a single transplant in previous studies. In this study, 5-year OS rates for poor-risk patients who completed the tandem transplant were 79% and 73% for complete and partial responders, whereas in a previous trial of single autologous HSCT, 5-year OS rates were 86% and 37% for complete and partial responders, all respectively. (24) The authors concluded that a single autologous HSCT is appropriate for intermediate-risk patients and for poor-risk patients who are complete responders to cytoreductive chemotherapy but that tandem autologous HSCT showed a benefit in patients with chemotherapy-resistant disease and in partial responders to cytoreductive conditioning. The authors stated that a trial of random assignment of single versus tandem autologous HSCT was unrealistic, given the low yearly incidence of poor-risk patients, and that the best possible comparisons are with data from previous findings with single transplants.
Clinical Trials and Guidelines

National Cancer Institute (NCI) database of clinical trials (Physician Data Query [PDQ®] database):

An open-label, Phase III trial is actively studying 3 different therapeutic strategies in patients with previously untreated HL, which includes 1 group of poor-prognosis patients who may or may not receive an autologous or allogeneic HSCT based on response to induction chemotherapy (NCT00920153). Estimated enrollment is 810, with an estimated primary completion date of May 2015.

A Phase III randomized study of induction chemotherapy followed by combination chemotherapy and autologous peripheral blood stem cell transplantation with or without high-dose sequential chemotherapy in patients with relapsed Hodgkin’s lymphoma is recruitment status unknown. (NCT00025636) Estimated enrollment is 220.

Summary

- Randomized Phase 3 studies have shown that, in patients with relapsed or refractory Hodgkin lymphoma (HL), autologous hematopoietic stem-cell transplantation (HSCT) leads to improved event- and progression-free survival and freedom from failure, versus conventional chemotherapy.
- The application of allogeneic HSCT to the treatment of patients with HL initially appeared limited due to a high procedure-related mortality rate. To date, most of the reduced-intensity conditioning (RIC) allogeneic HSCTs have been performed in patients who have failed a previous autologous HSCT for primary relapsed/refractory HL. Most of the studies are characterized by small numbers of patients, disparate preparative and graft-versus-host disease prophylaxis regimens, and varying lengths of follow-up. However, they have demonstrated reduced nonrelapse mortality, and some suggest a graft-versus-HL effect with favorable disease control, and possible cure, in these poor-prognosis patients.
- A multicenter, prospective trial showed that tandem autologous HSCT (versus single autologous HSCT) may provide a survival benefit in select patients with poor-risk features.

Practice Guidelines and Position Statements

Comprehensive Cancer Network (NCCN) Guidelines

The NCCN guidelines for HL (v.2.2013) include a recommendation for autologous HSCT in patients with progressive and relapsed HL. The guidelines state that allogeneic transplant is an option in select patients with progressive or relapsed disease as a category 3 recommendation and that allogeneic HSCT with reduced-intensity conditioning remains investigational. The guidelines do not specifically address tandem transplants.

Clinical Input Received through Physician Specialty Society and Academic Medical Center (October 2009)

In response to requests, input was received from 2 academic medical centers while this policy was under review. While the various physician specialty societies and academic medical centers may
collaborate with and make recommendations during this process, through the provision of appropriate reviewers, input received does not represent an endorsement or position statement by the physician specialty societies or academic medical centers, unless otherwise noted.

The 2 reviewers agreed with the policy statements, with the exception of the use of a second autologous HSCT after a prior autologous HSCT, which both reviewers thought would be medically necessary in certain circumstances. The data to support the use of a second autologous HSCT is extremely limited, and the policy statement for this use of HSCT remains investigational.

VII. Important Reminder

The purpose of this Medical Policy is to provide a guide to coverage. This Medical Policy is not intended to dictate to providers how to practice medicine. Nothing in this Medical Policy is intended to discourage or prohibit providing other medical advice or treatment deemed appropriate by the treating physician.

Benefit determinations are subject to applicable member contract language. To the extent there are any conflicts between these guidelines and the contract language, the contract language will control.

This Medical Policy has been developed through consideration of the medical necessity criteria under Hawaii’s Patients’ Bill of Rights and Responsibilities Act (Hawaii Revised Statutes §432E-1.4), generally accepted standards of medical practice and review of medical literature and government approval status. HMSA has determined that services not covered under this Medical Policy will not be medically necessary under Hawaii law in most cases. If a treating physician disagrees with HMSA’s determination as to medical necessity in a given case, the physician may request that HMSA reconsider the application of the medical necessity criteria to the case at issue in light of any supporting documentation.

VIII. References


