Autologous Hematopoietic Stem Cell Transplantation for Patients With Multiple Myeloma: An Overview for Nurses in Community Practice

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Autologous hematopoietic stem cell transplantation (AH SCT) is approved for the treatment of select solid tumors, autoimmune disorders, and most hematologic malignancies. Multiple myeloma (MM) is the most common indication for AH SCT. Despite improvement in response and survival rates in the era of novel agents, AH SCT remains an important treatment option for patients with MM who are eligible. Clinical management of patients with MM requires a multidisciplinary approach that incorporates healthcare professionals in a number of clinical settings as well as caregivers and the patient. Patients about to undergo AH SCT are generally referred to tertiary care centers that specialize in ASCT. Pre- and post-transplantation treatments and long-term follow-up often are managed by a community-based referring oncologist in collaboration with the transplantation team. Oncology nurses play an integral role in the care of patients with MM in each clinical setting. This article aims to provide non-transplantation oncology nurses with guidelines for education, clinical management, and support of patients with MM undergoing AH SCT with a primary focus on the pre- and post-transplantation period.

Hematopoietic stem cell transplantation (HSCT) is an accepted treatment for selected autoimmune and nonmalignant disorders, solid tumors, and hematologic malignancies. High-dose chemotherapy (HDC) is used in these settings to provide intensive cytotoxic therapy with the goal of eliminating malignant cells. However, the toxic effects of treatment are not specific to malignant cells alone, but affect all fast-growing cells. This results in expected side effects, most significantly bone marrow ablation (Antin & Yolin Raley, 2009). As such, reconstitution of the bone marrow and hematopoietic function using either autologous (patient’s own) or allogeneic (related or unrelated donor) stem cells is integral to the treatment process. In both procedures, stem cells are collected prior to receiving HDC, processed, stored, and then infused into the patient following HDC. Without stem cell “rescue” following HDC, patients would not recover bone marrow function, causing significant risk of mortality from life-threatening infection, bleeding, or anemia (Antin & Yolin Raley, 2009; Bensinger, 2009; Kumar, 2009; Rodriguez, 2010b). The general process for HSCT is found in Figures 1 and 2 and further discussed in Faiman, Miceli, Noonan, and Lilleby (2013), published on pages 33–41 of this supplement.
Multiple Myeloma Overview

Multiple myeloma (MM) is a malignant plasma cell disorder. Plasma cells produce immunoglobulin, which are proteins critical to the protective immune response. Immunoglobulins consist of a heavy chain (IgG, IgA, IgM, IgD, IgE) and a light chain (kappa or lambda) (Mangan, 2010). In MM, atypical plasma cells produce excess quantities of one of these proteins, referred to as paraproteins, monoclonal proteins, or M proteins. The patient-specific myeloma subtype is categorized by the involved immunoglobulin (heavy chain and light chain) (e.g., IgG kappa). Several factors are thought to play a role in the malignant transformation of plasma cells, including chromosome changes, molecular characteristics, and elements that affect the bone marrow microenvironment such as cytokine abnormalities. Many of these factors are thought to have prognostic significance (Palumbo & Anderson, 2011). The diagnosis of MM is based on the presence of greater than 10% atypical plasma cells in the bone marrow, presence of a monoclonal protein in the peripheral blood and/or urine, and additional laboratory and clinical findings (Durie et al., 2006; Kyle et al., 2003). The common clinical manifestations of MM are the byproduct of excess paraprotein and its impact on the cellular environment and organs, and include anemia, fatigue, hypercalcemia, bone disease, bone pain, renal dysfunction, and decreased immune function (Kyle et al., 2003; Mangan, 2010).

Autologous Versus Allogeneic Hematopoietic Stem Cell Transplantation

MM is the second most common hematologic malignancy, but is the most common indication for autologous HSCT (AHSCT) (Pasquini & Wang, 2011). Multiple studies demonstrate a survival benefit associated with AHSCT; therefore, AHSCT is considered the standard of care for eligible patients (Attal, 1996; Giralt et al., 2009; Kumar, 2009). Allogeneic HSCT (allo-HSCT) differs from AHSCT in that marrow created by donor cells can promote new immune activity in the recipient, providing a graft-versus-host disease (or antimyeloma) effect. Allo-HSCT is associated with high treatment-related morbidity and mortality and should only be pursued in the setting of a clinical trial (Bensinger, 2009; Lokhorst et al., 2010). Therefore, the focus of these guidelines will be directed toward AHSCT.

Treatment From Induction to Post-Transplantation Recovery

When designing the plan of care for a patient with MM, all treatment options should be considered. Every newly diagnosed patient with MM, even those older than age 70 years, should be considered a candidate for an AHSCT. Eligibility criteria vary by institution. Referral to a transplantation center, most often based on proximity and insurance contracting, should be made early in the treatment process when considering AHSCT (National Comprehensive Cancer Network [NCCN], 2013; Palumbo et al., 2011). More than 150 medical institutions...
exist in the United States that perform AHSC (Blood and Marrow Transplant Information Network, 2013; Center for International Blood and Marrow Transplantation, 2013). Each institution has specific protocols for pretransplantation screening, evaluation, consultation, and treatment planning. 

Transplantation eligibility is largely based on age, performance status, and desire to undergo the procedure (see Figure 3). The screening and approval process may take weeks to months. Ultimately, transplantation eligibility should be determined by a transplantation specialist.

Once a patient is diagnosed with active MM, the patient will initiate a treatment plan that includes chemotherapy. The goal is to effectively suppress the malignant clone and optimally reach a complete response prior to the collection of stem cells. Patients with MM who are eligible for transplantation should not receive regimens containing melphalan prior to stem cell collection because it can interfere with stem cell mobilization (Cavo et al., 2011; Giralt et al., 2009). Novel therapies (thalidomide, lenalidomide, pomalidomide, bortezomib, and carfilzomib), in combination with dexamethasone or standard chemotherapeutic agents, have demonstrated improved response rates (RR) and overall survival (OS) in patients with MM and are considered acceptable regimens prior to HSCT (Cavallio et al., 2011; Kyle & Rajkumar, 2009; Lacy et al., 2012; NCCN, 2013; Sonneveld, Asselbergs, et al., 2012). For those patients deemed transplantation ineligible, melphalan and other alkylating agents in combination with novel agents have shown significant responses and improved OS (Palumbo et al., 2011; Rajkumar et al., 2010; San Miguel et al., 2008). A number of clinical trials that include initial therapy, supportive care, maintenance, or treatment of relapse in the clinical trial setting currently are being conducted. Participation in clinical trials should be considered at all phases of treatment when possible.

AHSC collection and transplantation is a multistep process. The timing for stem cell collection is individualized based on the transplantation plan. For example, a patient may collect stem cells for storage purposes and continue with current therapy, or collect stem cells with the intent to proceed directly to AHSC. Patients who previously collected and stored stem cells will proceed with HDC and AHSC when clinically indicated without repeating the collection process. Certain side effects and clinical implications are common to therapies used at each stage of the process (see Table 1). Preparing for all phases of the transplantation process can be overwhelming for patients. Figure 4 and Appendix A provide considerations and Internet resources that may assist patients and caregivers.

### Post-Transplantation Recovery

The post-transplantation period begins after recovery from acute toxicity of the HDC, including blood count recovery (Kelley, McBride, Randolph, & Leum, 2000; Williams, 2004). This period of time has become obscured because of the many discharge options and individualized practices at each transplantation center. Patient acuity at time of discharge is higher than in years past, and the care of the patient with MM who has...
received AHSCT often is complex (Bevans, 2009). The majority of post-transplantation care becomes the responsibility of non-transplantation practitioners and caregivers. Patients anticipate the return home from the transplantation center, but also may experience anxiety in the transition.

The International Myeloma Foundation Nurse Leadership Board has compiled a summary of guidelines, recommendations, and clinical management strategies intended to optimize the quality of life (QOL) of patients undergoing transplantation and to minimize adverse events during the immediate post-transplantation period. The goal is to assist the community-based healthcare team, including oncology healthcare providers, to ease the transition from transplantation center to community, relieve anxieties, and provide information to guide the recovery of the patient after AHSCT. While reviewing these guidelines, note that QOL may improve over time. In several studies, transplantation-related symptoms and QOL improved or surpassed the pretransplantation level when measured at 6–12 months (Chao et al., 1992; Lyons et al., 2011; McQuellon et al., 1998; Saleh & Brockopp, 2001; Schulmeister, Quiett, & Mayer, 2005).

Considerations for the Nontransplantation Oncology Nurse

Discharge guidelines vary among transplantation centers, but generally include suggested management of psychological and physical needs of the patient. Although patients and their caregivers receive extensive education verbally and in writing prior to their discharge from the transplantation center, the amount of information may be overwhelming, and specific details forgotten. Therefore, ongoing educational reinforcement is essential for both patients and their caregivers. Familiarity with the discharge procedures and post-transplantation policies at the particular transplantation center from which the patient has been discharged will allow for reinforcement of key concepts when healthcare providers meet with patients and their families. If not provided to the patient at discharge, written instructions can be requested from the transplantation center to help guide care. Long-term survivorship issues also should be considered when caring for the patients with MM post-transplantation. Guidelines addressing fertility, sexuality, renal aspects, bone health, health maintenance, and mobility and safety can be found in a previous supplement to the Clinical Journal of Oncology Nursing from the International Myeloma Foundation Nurse Leadership Board (Bilotti et al., 2011; Bilotti, Gleason, & McNeil, 2011; Faiman, Mangan, Spong, & Tariman, 2011; Miceli, Colson, Faiman, Miller, & Tariman, 2011; Richards, Bertolotti, Doss, & McCullagh, 2011; Rome, Jenkins, & Lilleby, 2011).

Post-Transplantation Needs

The psychological impact of AH SCT should not be overlooked. Patients often describe the “let down” feeling after working hard before and during the transplantation, and many reflect on the events leading up to the transplantation and the details of the transplantation after being discharged. Transplantation recovery can be associated with physical setbacks as well as social strain on the caregiver and family. In fact, post-transplantation psychological issues may present greater challenges than the medical needs of the patient for the community-based healthcare team (Cooke, Gemmill, Kravits, & Grant, 2009).

The estimated rate of depression following stem cell transplantation ranges from 25%–50%. Depression affects physical health, can increase symptom-related

FIGURE 3. Treatment Algorithm for Newly Diagnosed Multiple Myeloma

Note. Based on information from Mikhael et al., 2013; National Comprehensive Cancer Network, 2013.
An additional information should be avoided prior to autologous hematopoietic stem cell collection. In some cases, antidepressive medications may be necessary. Caregivers and family members should be made aware of the frequency of post-transplantation distress, decrease survival, and has been associated with a higher incidence of suicide. Early identification of the symptoms of depression will allow the post-transplantation healthcare team to intervene early and refer the patient for more intensive services, such as psychiatric or social services and referral back to the transplantation center. In some cases, antidepressive medications may be necessary. Caregivers and family members should be made aware of the frequency of post-transplantation depression, signs and symptoms they should report, and how best to contact the appropriate healthcare provider (Cooke et al., 2009).

**Post-Transplantation Symptom Management**

Symptom management is vital for patients after stem cell transplantation. Persistent symptoms of HDC-related toxicity are common even after the patient has returned home. Symptom management is vital for patients after stem cell transplantation. Persistent symptoms of HDC-related toxicity are common even after the patient has returned home (see Table 2).

### TABLE 1. Common Multiple Myeloma Therapies, Side Effects, and Clinical Implications

<table>
<thead>
<tr>
<th>Drug, Class, Route</th>
<th>Potential Side Effects and Toxicities</th>
<th>Clinical Implication</th>
<th>Additional Information</th>
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<tbody>
<tr>
<td><strong>Myeloma Therapy Medications</strong></td>
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<tr>
<td><strong>Bortezomib</strong> Proteasome inhibitor IV or SQ administration</td>
<td>MS, PN, diarrhea or constipation, irritation or erythema at injection site; VZV activation</td>
<td>Monitor CBC, monitor PN symptoms, bowel management; use antiviral prophylaxis</td>
<td>Used as combination therapy or single agent; consider SQ administration to reduce PN</td>
</tr>
<tr>
<td><strong>Carfilzomib</strong> Proteasome inhibitor IV administration</td>
<td>Fatigue, anemia, thrombocytopenia, nausea, diarrhea, dyspnea, and fever</td>
<td>Monitor CBC and liver function tests. Prevent tumor lysis syndrome via PO and IV hydration; premedicate with dexamethasone in the first cycle</td>
<td>Approved for patients who have had two or more prior therapies, including bortezomib and an immunomodulatory agent</td>
</tr>
<tr>
<td><strong>Lenalidomide</strong> Immunomodulator Oral administration</td>
<td>MS, thromboembolic event when combined with steroids, and skin rash</td>
<td>Monitor CBC, bowel management, dose adjust for renal impairment; thromboembolic event prophylaxis</td>
<td>Used as combination therapy or as single-agent maintenance; hold for two weeks prior to autologous hematopoietic stem cell collection</td>
</tr>
<tr>
<td><strong>Melphalan</strong> Alkylator IV or oral administration</td>
<td>For conventional doses, MS; for high doses, myelosuppression, GI disturbance, and alopecia</td>
<td>Monitor CBC</td>
<td>Should be avoided prior to autologous hematopoietic stem cell collection; long-term use can cause myelodysplasia</td>
</tr>
<tr>
<td><strong>Pomalidomide</strong> Immunomodulator Oral administration</td>
<td>MS and thromboembolic event</td>
<td>Monitor CBC, bowel management; thromboembolic event prophylaxis</td>
<td>Approved for patients who have had two or more prior therapies, including bortezomib and an immunomodulatory agent</td>
</tr>
<tr>
<td><strong>Thalidomide</strong> Immunomodulator Oral administration</td>
<td>MS, thromboembolic event when combined with steroids, PN, and constipation</td>
<td>Monitor CBC, bowel management; thromboembolic event prophylaxis</td>
<td>Used in combination with dexamethasone</td>
</tr>
<tr>
<td><strong>Supportive Care Medications</strong></td>
<td></td>
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</tr>
<tr>
<td><strong>G-CSF/filgrastim</strong> Cytokine SQ administration</td>
<td>Joint and bone pain; increased white blood cells</td>
<td>Assess and medicate for pain</td>
<td>Management of neutropenia; autologous hematopoietic stem cell mobilization</td>
</tr>
<tr>
<td><strong>Pamidronate</strong> Bisphosphonate IV administration</td>
<td>Initial phase reaction, hypercalcemia, and osteonecrosis of the jaw</td>
<td>Dental evaluation prior to start (if possible), regular dental cleaning; avoid invasive dental procedure while receiving treatment</td>
<td>Inhibition of bone resorption and associated hypercalcemia. See ASCO and IMWG guidelines for duration of use. May be held during transplantation and resumed after.</td>
</tr>
<tr>
<td><strong>Plerixafor</strong> Chemokine inhibitor SQ administration</td>
<td>Diarrhea and erythema at injection site</td>
<td>Bowel management</td>
<td>Used in combination with G-CSF for stem cell mobilization</td>
</tr>
<tr>
<td><strong>Zoledronic acid</strong> Bisphosphonate IV administration</td>
<td>Initial phase reaction, hypercalcemia, and osteonecrosis of the jaw</td>
<td>Dental evaluation prior to start (if possible), regular dental cleaning; avoid invasive dental procedure while receiving treatment</td>
<td>Inhibition of bone resorption and associated hypercalcemia. See ASCO and IMWG guidelines for duration of use. May be held during transplantation and resumed after.</td>
</tr>
</tbody>
</table>

*See package insert for a complete listing of possible side effects. Practical use of medications may differ from U.S. Food and Drug Administration-approved indications and is done at the discretion of a licensed provider.*

ASCO—American Society of Clinical Oncology; CBC—complete blood count; G-CSF—granulocyte–colony-stimulating factor; GI—gastrointestinal; IMWG—International Myeloma Working Group; MS—myelosuppression; PN—peripheral neuropathy; SQ—subcutaneous; VZV—varicella zoster virus

**Note.** Based on information from Amgen Inc., 2013; Bertolotti et al., 2008; Bilotti, Gleason, et al., 2011; Bristol-Myers Squibb, 2005; Celgene Corporation, 2013a, 2013b, 2013c; Genzyme: A Sanofi Company, 2010; GlaxoSmithKline, 2008; Kumar, 2009; Millennium: The Takeda Oncology Company, 2012; Novartis Pharmaceuticals, 2012a, 2012b; Onyx Pharmaceuticals, 2012.
Post-Transplantation Infection Risk and Prevention

Post-transplantation infection is a major cause of morbidity and mortality. Although the patient’s white blood cell count and absolute neutrophil count may be within the normal range, the cells are functionally abnormal, placing the patient at increased risk for infection. In addition, continued physical weakness and malnutrition make recovery from a new infection difficult. Therefore, prophylactic antibiotics to prevent post-transplantation infections, such as invasive pneumococcal infection and pneumocystis pneumonia, are recommended for as long as one year following AHSCST (Tomblyn et al., 2009). Early detection and prompt intervention for infection is essential in caring for patients with MM (Palumbo et al., 2012). Careful assessment of the skin, lungs, gastrointestinal, renal, and skeletal systems are needed in identifying infection. Vital signs should be monitored at each clinic visit and patients should monitor their own vital signs as instructed by their care provider. Potential post-transplantation infections and precautions are listed in Table 3.

Frequent and meticulous hand washing by the patients and those they come in contact with is very important to prevent the transfer of infection. Many transplantation centers recommend that patients wear a mask when coming into a clinic or hospital for appointments. Patients may be advised to avoid public places such as restaurants, movies, or shopping malls. The suspension of these precautions will vary by individual centers and should be discussed in detail with the transplantation center.

Implications for Practice

- Include healthcare professionals from a number of clinical settings to best address the multidisciplinary approach required to manage multiple myeloma.
- Become familiar with the autologous hematopoietic stem cell transplantation process, as it remains an important treatment option for multiple myeloma.
- Incorporate guidelines for post-transplantation management in the community setting to promote quality of life and improve survival for patients.

Recommendations concerning personal hygiene, home maintenance, and cleanliness also may be provided by the transplantation center to further reduce the risk of infection. Guidelines for laundering clothes and housekeeping, particularly facilities used by the patient, are commonly provided. Specific policies regarding personal hygiene also are often recommended. It must be kept in mind that the patient may not be able to perform some of these duties independently in the first months following HSCT, emphasizing the need to include caregivers in the education process (Antin & Yolin Raley, 2009). The role of caregivers in the recovery of patients with HSCT is discussed in greater detail by Kurtin, Lilley, and Spong (2013) on pages 25–32 of this supplement.

Because of the risk of food-borne infection, specific nutritional and dietary guidelines may be mandated by the transplantation center. Nutritional recommendations and restrictions may begin at the start of HDC and continue after discharge. In general, the Advisory Committee on Immune Practices recommends foods that have been refrigerated, pasteurized, or well-cooked for patients during the post-transplantation period (Antin & Yolin Raley, 2009; Tomblyn et al., 2009).

Smoking tobacco is prohibited after an AHSCST for many reasons. People who smoke are at increased risk for developing pneumonia as well as pulmonary and cardiovascular toxicity related to AHSCST. Marijuana use is prohibited because of the heightened risk of fungal infection associated with inhalation. Alcohol consumption also is restricted because of its potential effect on the liver, platelets, and immune function (Sipsas & Kontoyiannis, 2008; Tichelli et al., 2008; Versteeg, Slot, van der Velden, & van der Weijden, 2008).

Post-Transplantation Immunizations

The transplantation process results in a loss of T and B lymphocytes which, in turn, causes loss of immune memory. Immune memory is shaped by the culmination of exposure to infectious agents, environmental antigens, and vaccines during a person’s lifetime (Kroger, Sumaya, Pickering, & Atkinson, 2011; Stadtmauer et al., 2011). Therefore, patients require reimmunization.

Post-transplantation immunizations vary by institution. Based on the Centers for Disease Control and Prevention and Advisory Committee on Immune Practices recommendations, non-live vaccines may be administered as early as three months post-transplantation. Live-attenuated vaccines may be administered two years following transplantation in immune-competent people (Tomblyn et al., 2009). An example of an immunization schedule can be found in Table 4.

American Cancer Society
Information on disease types and available support
www.cancer.org

Be the Match: National Marrow Donor Program
Transplantation-related information for patients and caregivers
www.marrow.org

BMT Information Network
Transplantation-related information for patients and caregivers
www.BMTInfoNet.org

Caring Bridge
A site to create a personal blog or journal that can be shared with family and friends
www.caringbridge.org

International Myeloma Foundation
Information about myeloma, research, and available support
www.myeloma.org

Leukemia and Lymphoma Society
Information on disease types and available support
www.lls.org

Multiple Myeloma Research Foundation
Information about myeloma, research and available support
www.themmrf.org

National Bone Marrow Transplant Link
Transplantation-related information for patients and caregivers
www.nbmtlink.org

National Cancer Institute
Information on disease types and research
www.cancer.gov

Note. Website addresses and content can change; therefore, the information should be reviewed before sharing with patients.
### TABLE 2. Post-Transplantation Symptoms, Clinical Findings, and Management Strategies

<table>
<thead>
<tr>
<th>Symptom</th>
<th>Clinical Findings and Risk Factors</th>
<th>Management Strategies</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anorexia</td>
<td>Weight loss, taste changes, change in performance status, fatigue, nausea and vomiting, and diarrhea</td>
<td>Review medications for possible source. Medical nutritional therapies: oral nutritional supplements, IV hydration. Small frequent meals, calorie counts, weekly weight, nutritional consult. Reinforce improvement with time. Adjust medications as needed. Treat underlying cause (e.g., medication for nausea and vomiting).</td>
</tr>
<tr>
<td>Anxiety and depression</td>
<td>Fatigue, exhaustion, difficulty sleeping, difficulty concentrating, restlessness, irritability and impatience, recurrent thoughts of diagnosis and treatment, and anorexia</td>
<td>Listen to and validate concerns. Referral to social services, psychiatry, and support groups. Pharmacologic: anti-anxiety medication, antidepressants. Complementary and alternative medicine therapy: relaxation therapy, mild exercise such as walking.</td>
</tr>
<tr>
<td>Diarrhea</td>
<td>Increased frequency of bowel movements, abdominal cramps, dehydration, and decrease in weight</td>
<td>Review medications for possible source (i.e., antibiotics, narcotic withdrawal). Electrolyte evaluation. Stool sample for enteric pathogens (i.e. <em>Clostridium difficile</em>). Anti-diarrheal medication. Appropriate fluid and electrolyte replacement. Adjust diet for food sensitivities: milk products, certain spicy foods, nutritional supplements, fatty foods, chocolate. Antibiotics as needed; adjust medications as needed.</td>
</tr>
<tr>
<td>Fatigue</td>
<td>Decrease in energy, inability to complete tasks, insomnia or hypersomnia, not feeling rested after sleeping at night, and generalized weakness</td>
<td>Review medications that may cause fatigue. Assess for anemia. Mild exercise such as walking. Potentially decrease or discontinue medications that cause fatigue. Counsel patient on sleep hygiene, such as minimizing napping or staying in bed throughout the day. Erythropoietin medication if indicated and after obtaining written consent. Red blood cell transfusion, if needed.</td>
</tr>
<tr>
<td>Fever</td>
<td>Diarrhea, muscle weakness, fatigue, confusion, and seizures</td>
<td>Panculture, chest x-ray, and CBC with differential and platelets. Prophylactic antibiotics if neutropenic; therapeutic antibiotics if culture positive. Acetaminophen, IV hydration, symptom management. Monitor for fever greater than 101.3°F (and lower temperatures if patients are not feeling well), blood pressure declining from baseline and tachycardia.</td>
</tr>
<tr>
<td>Nausea and vomiting</td>
<td>Anorexia, nausea and vomiting, weight loss, and diminished skin turgor</td>
<td>Quantify episodes of emesis. Assess fluid and electrolyte status. Review medications for antiemetics and medications that may cause nausea and vomiting. Adjust medications if possible and as needed. IV or oral hydration and replace electrolytes as needed.</td>
</tr>
<tr>
<td>Pain</td>
<td>Assess for new or existing pain symptoms, current pain medication, assess for pain related to infection, and assess for symptoms of depression or anxiety</td>
<td>Appropriate pain medication regimen: long-acting pain medication together with breakthrough pain medication, doses titrated to effectiveness. Consider imaging for source of new or worsening pain. Consult with appropriate specialty, if indicated.</td>
</tr>
<tr>
<td>Thrombosis (DVT or PE)</td>
<td>Painful, swollen and erythematosus extremity (most often lower extremity), shortness of breath, tachycardia, chest pain, and HTN; patients at increased risk: those with obesity, diabetes, cardiovascular disease, HTN, hyperlipidemia, immunomodulatory agents with concurrent high-dose steroids, anthracyclines, ESAs, hospitalizations, and immobility</td>
<td>Prevention: thromboprophylaxis for all patients at risk. Full therapeutic anticoagulation for any patients with more than two risk factors. If DVT or PE is suspected: Doppler ultrasound of suspected extremity. High-resolution chest CT with PE protocol if PE is suspected. Medication to treat thrombosis: low molecular weight heparin, warfarin, and alternative anticoagulants. Consult with coagulation specialist if appropriate.</td>
</tr>
</tbody>
</table>

**Note:** Based on information from Antin & Yolin Raley, 2008; Eaton & Tipton, 2010; Rodriguez, 2010a.

**Table Key:**
- PE—pulmonary embolus
- PN—peripheral neuropathy
- CBC—complete blood count
- CT—computed tomography
- DVT—deep vein thrombosis
- ESA—erythropoietin-stimulating agent
- HTN—hypertension
- PE—pulmonary embolus
**Post-Transplantation Medication Considerations**

Polypharmacy during transplantation is common and can be confusing. Therefore, many transplantation teams will provide patients and their families with a medication chart to help track and maintain the dose and the administration of scheduled and “as needed” medications. Pretransplantation and transplantation-specific medications are added, discontinued, and adjusted frequently according to the patient’s needs during the acute care phase of the transplantation. Hypertension and hyperglycemic management regimens, in particular, often require modifications during the transplantation process. A list of discharge medications should be provided to the patient as well as to the discharge facility or homecare agency and the patient’s referring oncologist involved in the patient’s care. Patients may be restricted from taking certain over-the-counter medications such as nonsteroidal anti-inflammatory drugs and supplements because of drug interactions, organ toxicity, or interference with therapy. Medications must be taken as prescribed and medication changes should be discussed with the staff at the transplantation center.

**Disease Management Following Autologous Hematopoietic Stem Cell Transplantation**

Although AH SCT remains an important treatment strategy for patients with MM, relapse of MM is inevitable for the majority of patients. The timing is unpredictable and relapse can occur at any time following AH SCT, ranging from months to years. Those considered at high risk based on stage of disease and cytogenetics are at greater risk of early relapse (Mikhael et al., 2013; Palumbo & Cavallò, 2012). Providing patients with the clear message is important so that when progression does occur, they understand that it does not necessarily indicate end of life, but, rather, a time for change in therapy. Determining the optimal time to next therapy remains a controversial issue following AH SCT, and several studies are ongoing. Data

<table>
<thead>
<tr>
<th>Infection</th>
<th>Type of Infection</th>
<th>Prevention</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Bordetella pertussis</strong> (whooping cough)</td>
<td>Community-acquired bacterial respiratory infection</td>
<td>If exposed, prevention and treatment available (azithromycin or SMX/TMP) If hospitalized, proper precautions should be taken to avoid transmitting to others. A cellular vaccine is recommended.</td>
</tr>
<tr>
<td>Community respiratory viruses: RSV, influenza, adenovirus, and parainfluenza</td>
<td>Viral infections that can progress to bronchitis or pneumonia RSV is more common in infants, but can be seen in older adults and ICHs. Influenza accounts for about 20% of respiratory viral infection in patients who receive a transplantation. Adenovirus may manifest as a diarrheal illness.</td>
<td>Handwashing; ICHs should wear a mask, clean hard surfaces with anti-infective wash, and avoid crowds and people with respiratory symptoms. If hospitalized, proper precautions should be taken to avoid transmitting to others. Antiviral medication may be available. Inactivated influenza vaccine should be administered to the patient and direct caregivers unless contraindicated. Do not use the live inhaled version. May begin as early as 3–6 months post-transplantation and every year of life.</td>
</tr>
<tr>
<td><strong>Pneumocystis carinii</strong> pneumonia (renamed as PJP)</td>
<td>Protozoal infection that can develop in ICHs. ICHs and patients with AH SCT may develop PJP if prophylaxis is not provided; can occur early after transplantation, particularly if the patient has been heavily treated beforehand.</td>
<td>SMX/TMP, atovaquone, dapsone, or aerosolized pentamidine, depending on allergy profile. Prophylaxis for 3–6 months following AH SCT</td>
</tr>
<tr>
<td>Streptococcus pneumonia or invasive pneumococcal infection</td>
<td>A gram-positive encapsulated organism that can cause sudden and serious systemic infection in patients following AH SCT. Considered a late transplantation complication and is common in patients with multiple myeloma because of decreased humoral immunity</td>
<td>Penicillin or doxycycline, depending on allergy profile. Prophylaxis for 12 months following AH SCT until revaccinated Pneumococcal vaccine should be administered as a 7-valent or 23-valent vaccine as early as 3–6 months post-transplantation.</td>
</tr>
<tr>
<td><strong>Viridans streptococci</strong></td>
<td>Organism found commonly in the oral cavity Greatest concern during times of oral mucositis</td>
<td>Quinolone therapy for neutropenic state longer than seven days.</td>
</tr>
<tr>
<td>VZV (shingles)</td>
<td>Primary infection, commonly known as chicken pox. VZV persists in the sensory nerve ganglia. Reactivation is common in older adults or ICHs.</td>
<td>Acyclovir or valacyclovir therapy for one year or while on active treatment. Also prevents herpes simplex virus, type I and II. The shingles vaccine is a live virus and currently not recommended for patients with multiple myeloma.</td>
</tr>
</tbody>
</table>

AH SCT—autologous hematopoietic stem cell transplantation; ICH—immunocompromised host; PJP—*Pneumocystis jiroveci* pneumonia; RSV—respiratory syncytial virus; SMX/TMP—sulfamethoxazole/trimethoprim; VZV—varicella zoster virus

**Note.** Schedule and use vary between transplantation centers.

**Note.** No live vaccines should be given in the first year following transplantation.

**Note.** Based on information from Antin & Yolín Raley, 2009; Centers for Disease Control and Prevention, 2012; Cordonnier et al., 2010; Stadtmauer et al., 2011; Tomblyn et al., 2009.
regarding the use of maintenance therapy following HSCT for MM continue to be reported. Attal et al. (2012) reported improvement in progression-free survival (PFS) following AHSCT using lenalidomide as maintenance therapy but no increase in OS. McCarthy et al. (2012) also reported an increase in PFS as well as a longer OS. Bortezomib can be used as maintenance post-transplantation as well, and may be associated with improvement in PFS (Sonneveld, Schmidt-Wolf, et al., 2012).

Conclusion

Care of the patient following AH SCT is complex, however, expected side effects of HDC usually are manageable. Although consistent objectives and goals are in place for AH SCT recipients, care must be individualized based on pretransplantation treatment toxicities and transplantation-related side effects. Community oncology professionals play a critical role in the collaborative management of the patient with MM throughout the treatment continuum. Consistent communication among the patient, the referring center, and the transplantation center is vital to ensure all testing, insurance approval, and support services are in place prior to starting the transplantation process. Post-transplantation guidelines are not standardized, and recommendations for post-transplantation care vary between transplantation centers. These factors add to the challenge of caring for the transplantation patient in a community-based setting.

All providers should assist in supporting the patient and family members through the transplantation journey. Discharge is an exciting time for the patient, but also can be physically challenging and emotionally overwhelming. Community providers are instrumental in monitoring and managing post-transplantation concerns. Understanding the AH SCT rationale, process, and needs of the patient post-transplantation will improve the QOL for the patient undergoing transplantation and impact OS. Maintaining a collaborative management approach with consistent communication between the transplantation center and community healthcare provider team will improve overall outcomes for patients undergoing transplantation.

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References


TABLE 4. Post-Transplantation Immunization Schedule

<table>
<thead>
<tr>
<th>Organism</th>
<th>Vaccine</th>
<th>Time Post-HSCT to Initiate Vaccine</th>
<th>Dose and Route</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pneumococcal</td>
<td>PCV7/PPSV23</td>
<td>3–6 months</td>
<td>0.5 ml IM or SQ</td>
<td>Can be given six months post-transplantation</td>
</tr>
<tr>
<td>Pertussis, tetanus, diptheria</td>
<td>DTAP</td>
<td>6–12 months</td>
<td>0.5 m. IM</td>
<td>Can be given six months post-transplantation</td>
</tr>
<tr>
<td>Haemophilus influenza type B</td>
<td>Hib</td>
<td>6–12 months</td>
<td>0.5 ml IM</td>
<td>Can be given six months post-transplantation</td>
</tr>
<tr>
<td>Hepatitis B</td>
<td>–</td>
<td>6–12 months</td>
<td>1 ml IM</td>
<td>Administer to patients who are hepatitis B virus negative.</td>
</tr>
<tr>
<td>Meningococcus</td>
<td>–</td>
<td>6–12 months</td>
<td>0.5 ml SQ</td>
<td>Recommended in areas with an increase in meningococcus</td>
</tr>
<tr>
<td>Influenza</td>
<td>–</td>
<td>4–6 months</td>
<td>0.5 ml IM (the nasal version is live and, therefore, not recommended)</td>
<td>Give annually as available in the autumn months. May administer four months post-transplantation, however, two doses of the vaccine are suggested.</td>
</tr>
<tr>
<td>Measles, mumps, and rubella</td>
<td>MMR</td>
<td>24 months</td>
<td>0.5 ml SQ</td>
<td>MMR should not be given if the patient is immunosuppressed.</td>
</tr>
<tr>
<td>Varicella zoster virus (shingles)</td>
<td>Zoster vaccine</td>
<td>Not currently recommended, Clinical trials are ongoing.</td>
<td>–</td>
<td>Not currently recommended; inactivated version is under investigation. Prevention with antiviral medication is recommended.</td>
</tr>
</tbody>
</table>

HSCT—hematopoietic stem cell transplantation; IM—intramuscular; SQ—subcutaneous

Note. Based on information from Cordonnier et al., 2010; Kroger et al., 2011; Ljungman et al., 2009; Tomblyn et al., 2009.


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### At the Transplantation Center

**Consider bringing these items to the transplantation center consultation**
- Medical records (i.e., radiology on disc, laboratory results, bone marrow reports)
- List of previous chemotherapy and dates received
- Current and recently taken medications (both prescribed and supplements)
- Questions for the physician

**Important paperwork to bring**

Medical Leave of Absence (MLOA), Family Medical Leave Act (FMLA), and insurance benefits. Notify your employer of MLOA and complete FMLA forms. Forms need to be completed for patient and caregivers. Anticipated time for processing is eight weeks.

**Suggested items to bring for your stay**

Comfortable clothing, sedentary activities, multi-unit pill dispenser, personal items of comfort, and cell phone

### Follow-Up Care

**Long-term follow-up plan of action**

Plan to return to the transplantation center near day 100 post-transplantation for a full evaluation. This may be a 2–3 day visit. Post-transplantation immunization may be recommended at 12, 14, and 24 months following transplantation.

**Oral medication management**

Administering scheduled and as needed medications, refills, and renewals

**IV fluids and medications**

In some situations, home infusion of medications for specific conditions (i.e., dehydration, infection, low magnesium) may be ordered by the healthcare provider.

**Symptom monitoring**

Fever, bruising, bleeding, new onset of pain (bone or nerve); changes in energy, appetite, weight (up or down), bowel function, and bladder function. Know your contact information—who to call, where to go.

**Patient advocate**

Communicate with healthcare providers, employers, family and friends; consider creating a blog to keep friends and family informed

**Central line care**

This varies from institution to institution. Instructions will be provided by the transplantation center.

### In the Community

While away from home . . .

- Arrange for caregiver support while at the transplantation center. Caregivers may be rotated. Investigate housing options near the transplantation center.
- Arrange for child and pet care.
- Arrange for care of your home while you are away and assistance when you return after transplantation (i.e., yard maintenance, mail delivery, and utilities).

**Seek help with household chores and activities of daily living**

- Assistance with cleaning, paying bills, grocery shopping, laundry
- Assistance with bathing, dressing, meal preparation, and transportation (medical appointments, shopping, pharmacy)
- Encouragement regarding oral intake, ambulation, and strengthening

**Fundraising**

Out-of-pocket expenses add up quickly. Consider hosting a fundraising event in your community to help cover healthcare costs.

### APPENDIX A. Preparation and Activity Considerations for Patients With Multiple Myeloma Undergoing Hematopoietic Stem Cell Transplantation