Bronchiolitis Obliterans Organizing Pneumonia: A Late Complication of Stem Cell Transplantation

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Bronchiolitis obliterans organizing pneumonia (BOOP) is a late complication of hematopoietic stem cell transplantation. Many transplant recipients have returned to their community oncologists before BOOP has occurred. The etiology of BOOP in transplant recipients is not understood fully, but it may be associated with chronic graft-versus-host disease. BOOP presents both vague and common symptoms that can progress into respiratory failure but respond to early and appropriate treatment. Early recognition and treatment of BOOP are essential to improving survival of patients who are post-transplantation. Community oncology nurses are in a key position to support initiation of side-effect management. This article presents information about the etiology, presentation, diagnostic testing, and treatment of BOOP. Two case studies are included that illustrate the typical course of BOOP and its treatment.

Key Words: bronchiolitis obliterans organizing pneumonia, peripheral blood stem cell transplantation, steroids

Etiology

BOOP is an inflammatory lung disease that is characterized by the formation of granulation tissue plugs in the alveoli that can extend into the bronchioles. The association with cGVHD suggests that BOOP may represent the rejection of the lungs by transplanted stem cells (Afessa, Litzow, & Tefferi, 2001). Inflammatory cells, such as neutrophils, lymphocytes, and plasma cells, sometimes are seen at the center of intraluminal myxoid polyps (newly formed connective tissue that participates in remodeling and destruction of the interstitium) (Mokhtari, Bach, Tietjen, & Stover, 2002).

Several known causes of BOOP exist, and the most common is idiopathic (Epler, 2001). Other causes include infection, medications, and stem cell transplantation, which is the second most common cause. Many medications used with stem cell transplantation reportedly interact with radiation used as a conditioning regimen and potentially can increase damage to normal lung tissue. Figure 1 lists risk factors for BOOP related to allogeneic HSCT. Although no direct correlation between BOOP and cGVHD exists, blood chemistries performed on patients with respiratory symptoms show decreased immunoglobulin A and G antibody levels. This suggests that immunosuppression plays an important role in this condition (Buchsel, Leum, & Randolph, 1996).

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