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Incidence of agranulocytosis in allogeneic bone marrow transplant recipients. The occurrence of agranulocytosis in 64 patients receiving allogeneic stem cell transplants was evaluated prospectively. Twenty-five of these patients (39%) developed agranulocytosis during the course of their grafting following high-dose immunosuppressive therapy. Severe granulocytopenia and/or monocytopenia was observed in all these patients but only nine of these 25 developed bacterial, fungal, or viral infections. Seven of these nine developed life-threatening septicemia and in two of these, pulmonary abscesses were suspected as the infection. Bone marrow biopsy performed in five of these nine patients revealed leukemic infiltration (four patients), graft versus host disease (two patients), pure granulocytic hyperplasia (one patient), and fibrosis (one patient). Immunosuppression was stopped in eight of these nine patients, while antibiotics and/or antifungal therapy was initiated in two patients with suspicion of infection. Seven of these nine developed complete recovery without additional therapy within 3 months of the onset of agranulocytosis and the other two had complete recovery after the administration of broad-spectrum antibiotics. Immunosuppression was restarted after 2-4 months in the latter two patients and they recovered. One patient died from fungal pneumonia during the recovery phase. No association of the type of graft donor or conditioning regimen with the occurrence of agranulocytosis could be established. These findings indicate that allogeneic bone marrow transplantation is associated with a significantly increased incidence of agranulocytosis after myeloablative immunosuppressive therapy.", "clear-table-header": "□□□□◆"