

Splenic mucormycosis

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Mucormycosis is a rare opportunistic infection caused by fungi of the order *Mucorales*. It has usually been associated with underlying conditions such as diabetic ketoacidosis, neutropenia, organ transplantation, and immunosuppressive therapy.¹ The clinical presentation is rhinocerebral or pulmonary disease in most cases.^{2,3} Cutaneous, gastrointestinal, splenic and disseminated infections are much less common.^{4,5}

A 64-year-old man was diagnosed as having multiple myeloma IgA- κ in June 1992. The patient was treated with melphalan and prednisone and an objective response was documented. He received treatment with melphalan and prednisone in July 1995, but only a partial response was observed. The patient remained asymptomatic without therapy during the next year and follow-up studies showed moderate pancytopenia



Figure 1. Abdominal CT scan showing a hypodense mass in the spleen (arrow).

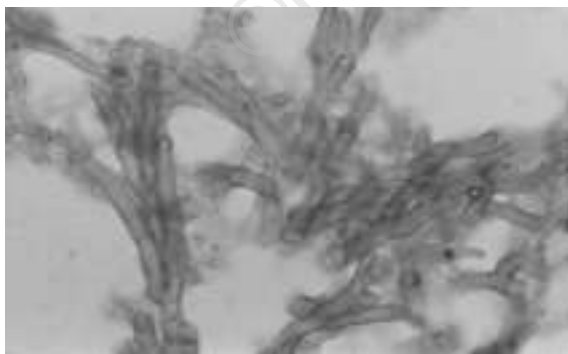


Figure 2. Spleen section stained with hematoxylin-eosin (x1000). *Mucorales* appears as broad, non- or sparsely-septate hyphae with irregular branching.

(neutrophil count $> 0.5 \times 10^9/L$) and a stable M-component. In November 1997 the patient was admitted because of rectal hemorrhage. A colonoscopy revealed an adenocarcinoma in the left colon. At that time, chest radiography showed a round mass in the left lower lung lobe. Abdominal CT scan showed a hypodense mass of 4 cm of diameter in the spleen (Figure 1). Segmentary colectomy and splenectomy were performed. Histology of a cutaneous nodule and the spleen revealed necrotic abscesses containing hyphal elements compatible with a *Mucorales* infection (Figure 2). Therapy with liposomal amphotericin B (LAB) therapy (1.5-3.5 mg/kg/d) and G-CSF (300 $\mu\text{g/d}$) was started. After seven days of LAB therapy the patient died.

Mucormycosis has a high mortality rate and diagnosis is usually made at autopsy.^{6,7} The standard therapy of mucormycosis consists in high doses of amphotericin B and surgical resection of localized infection.^{1,3} Some authors have suggested that the use of novel therapies such as LAB and G-CSF might improve the prognosis.^{6,8} Unfortunately, our patient died despite surgical resection of an infectious focus, and treatment with LAB and G-CSF.

References

1. Sugar AM. Mucormycosis. *Clin Infect Dis* 1992; 14 (Suppl 1):S126-129.
2. Rangel-Guerra RA, Martínez HR, Sáenz C, Bosques-Padilla F, Estrada-Bellmann I. Rhinocerebral and systemic mucormycosis. Clinical experience with 36 cases. *J Neurol Sci* 1996; 143:19-30.
3. Tedder M, Spratt JA, Anstadt MP, Hegde SS, Tedder SD, Lowe JE. Pulmonary mucormycosis: results of medical and surgical therapy. *Ann Thorac Surg* 1994; 57:1044-50.
4. Adam RD, Hunter G, DiTomasso J, Comerchi G Jr. Mucormycosis: emerging prominence of cutaneous infections. *Clin Infect Dis* 1994; 19:67-76.
5. Singh N, Gayowski T, Singh J, Yu VL. Invasive gastrointestinal zygomycosis in a liver transplant recipient: case report and review of zygomycosis in solid-organ transplant recipients. *Clin Infect Dis* 1995; 20:617-20.
6. Gonzalez CE, Couriel DR, Walsh TJ. Disseminated zygomycosis in a neutropenic patient: successful treatment with amphotericin B lipid complex and granulocyte colony-stimulating factor. *Clin Infect Dis* 1997; 24:192-6.
7. Pagano L, Ricci P, Tonso A, et al. Mucormycosis in patients with haematological malignancies: a retrospective clinical study of 37 cases. *Br J Haematol* 1997; 99:331-6.
8. Sahin B, Paydas S, Cosar E, Bicakci K, Hazar B. Role of granulocyte colony-stimulating factor in the treatment of mucormycosis. *Eur J Clin Microbiol Infect Dis* 1996; 15:866-9.