# THE CLINICAL PICTURE

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# The Clinical Picture Ulcerative leg nodules in a transplant recipient



FIGURE 1. Purpuric nodules with surrounding erythema.

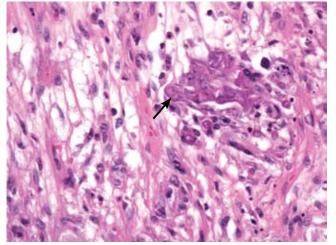
A 66-YEAR-OLD WHITE MAN presents with a 1-month history of two nontender, nonpruritic skin lesions on his leg 3 weeks after undergoing bilateral lung transplantation for idiopathic pulmonary fibrosis (FIGURE 1). Except for transient lymphopenia, the postoperative course has been uneventful, with no episodes of rejection. Immunosuppressive drugs include tacrolimus (Prograf) and steroids. He has a history of insulin-dependent diabetes mellitus, and no history of significant trauma to his legs. Other than the skin lesions, he feels well and has no systemic symptoms. He has no history of iron overload or deferoxamine (Desferal) therapy.

**Q:** What is the probable diagnosis?

- □ Kaposi sarcoma
- 🗌 Majocchi (trichophytic) granuloma
- □ Staphylococcus aureus furunculosis
- □ Primary cutaneous mucormycosis
- Disseminated zoster

A: This patient has primary cutaneous mucormycosis. Microscopic examination of the skin biopsy showed suppurative granulomatous inflammation

\*Dr. Billings has disclosed that he has received consulting fees from Allos Therapeutics. doi:10.3949/ccjm.76a.09006



**FIGURE 2.** Irregular nonseptated, branching, fungal hyphae (arrow) throughout the dermis are surrounded by a histiocytic granulomatous infiltrate (hematoxylin-eosin, original mangification × 400).

with numerous nonseptated, branching fungal hyphae (FIGURE 2). A fungal culture grew *Absidia* species. No evidence of disseminated disease was found on computed tomography of the brain, paranasal sinuses, chest, abdomen, and pelvis. Histologic studies and cultures taken from both native and transplanted lungs were negative for fungi.

# CAUSES AND RISK FACTORS

Mucormycosis is an invasive infection caused by fungi of a variety of genera (*Rhizopus*, *Rhizomucor*, *Mucor*, *Absidia*, *Apophysomyces*, *Cunninghamella*, *Saksenaea*, *Conidiobolus*, and *Basidiobolus* species) belonging to the class Zygomycetes in the order Mucorales. These ubiquitous environmental fungi have been found widely distributed in hospital sources, having been cultured from plaster casts,<sup>1</sup> tongue depressors,<sup>2</sup> cloth tapes for securing endotracheal tubes,<sup>3</sup> peritoneal dialysis catheters,<sup>4</sup> surgical wounds,<sup>5</sup> contaminated dressings, needles, and intravenous catheters.<sup>6</sup>

These fungi rarely cause disease in healthy people, but patients with certain risk factors may develop disseminated disease, in which the death rate is high.<sup>6</sup> Risk factors include diabetes mellitus, renal failure, trauma, burns, malnutrition, solid organ transplantation, hematologic malignancies, use of immunosuppressive drugs (eg, chemotherapeutic agents, corticosteroids, cyclosporine, methotrexate, infliximab [Remicade]<sup>7</sup>), and iron overload and iron-chelating therapy with deferoxamine.<sup>8</sup> Patients such as ours—an organ transplant recipient with concomitant diabetes mellitus—are highly susceptible to this infection.

## DIAGNOSIS

The clinical presentation of mucormycosis can be rhino-orbitocerebral (most common),<sup>9</sup> pulmonary, primary cutaneous, gastrointestinal, cardiac, or disseminated infection.<sup>6</sup> Primary cutaneous mucormycosis occurs from fungal inoculation of the dermis. Small areas of trauma may be all that is needed for this inoculation to occur.

The initial lesion may be an erythematous patch, plaque, or nodule that may subsequently ulcerate and become gangrenous or necrotic.<sup>10</sup>

The differential diagnosis of new cutaneous nodules in an immunocompromised patient includes a wide variety of infections, such as ecthyma gangrenosum caused by *Pseudomonas* or *Candida* species, herpes simplex virus infection, cryptococcal infection, phaeohyphomycosis, and cutaneous aspergillosis.

Because the morphology of cutaneous mucormy-

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cosis is not distinctive, and because we need to cast our net wide for the numerous, potentially lethal diagnostic possibilities in an immunosuppressed patient, it is crucial that we have a low threshold for prompt biopsy to establish the diagnosis and initiate definitive treatment. While skin biopsy for microscopy can suggest certain fungi, the exact diagnosis is confirmed by a fungal culture of a biopsy specimen.

### TREATMENT

Localized soft tissue infection is more amenable to therapy and therefore carries a better prognosis than visceral or disseminated disease.

The best treatment outcomes of primary cutaneous mucormycosis are achieved with both complete excision and debridement of necrotic tissue and systemic antifungal therapy. Amphotericin B in conventional form (Fungizone) and liposomal form (AmBisome) and posaconazole (Noxafil)<sup>11–12</sup> are effective.

Paradoxically, in contrast to deferoxamine, other iron chelators such as deferiprone (Ferriprox, not available in the United States) and deferasirox (Exjade), which do not supply iron to the fungus, had been shown to be effective against Zygomycetes in in vitro and animal models. The role of iron chelators as adjunctive therapy for mucormycosis needs further investigation.<sup>8</sup>

Primary cutaneous mucormycosis may become a disseminated infection. Therefore, one should have a high index of suspicion. When a transplant recipient develops a cutaneous plaque or nodule, biopsy should be performed promptly. Failure to do so can increase the risk of morbidity and death.

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**CME ANSWERS** Answers to the credit test on page 616 of this issue 1D 2A 3B 4C 5D 6C 7A 8E