

Rare Angioinvasive Fungal Infection in Association With Leukemia Cutis

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Practice Points

- Immunosuppressed patients are at risk for atypical presentations of common infections as well as infection with rare pathogens.
- Skin biopsy and tissue culture play an important role in identifying infectious agents in immunosuppressed patients.
- Leukemic infiltrates may mask pathogens, and pathologists should strongly consider additional stains when indicated.

*Leukemia cutis (LC) is characterized by the infiltration of malignant neoplastic leukocytes or their precursors into the skin and is most often seen in conjunction with systemic leukemia. Patients with LC frequently are in a relative or absolute immunocompromised state. We report the case of a 52-year-old man with primary refractory acute myelogenous leukemia (AML) following allogeneic stem cell transplant (SCT) who presented with a progressive reddish purple nodule with surrounding erythema and central necrosis in the setting of leukocytosis and possible fungal pneumonia. Histopathologic examination revealed an ulcerated dense diffuse dermal infiltrate of large atypical lymphocytes consistent with LC and septate hyphae with acute-angle branching in the dermal blood vessels. Cultures from a biopsied lesion grew *Paecilomyces species*, a rare but emerging*

opportunistic infection, despite the patient being on antifungal prophylaxis. This novel report of a rare angioinvasive infection occurring within a lesion of LC supports the need to maintain a high index of suspicion for invasive infection in patients with hematologic malignancy, even those on antifungal prophylaxis.

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Leukemia cutis (LC) is characterized by the infiltration of malignant neoplastic leukocytes or their precursors into the skin, most often in conjunction with systemic leukemia.¹ Acute myelogenous leukemia (AML) is the second most common cause of LC and the most common form of leukemia among adults.¹ Patients with leukemia often are in a relative or absolute immunocompromised state, which may be secondary to neutropenia, chemotherapy regimens, or immunosuppressive regimens following stem cell transplant (SCT). Thus, when evaluating cutaneous lesions consistent with LC in immunocompromised patients, there must be a high index of suspicion for concomitant opportunistic infections.

We report the case of a 52-year-old man with primary refractory AML following allogeneic SCT with relapse who presented with an LC lesion below the knee with concomitant invasive fungal infection despite being on prophylactic oral antifungal therapy.

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Case Report

A 52-year-old man with primary refractory AML (M1) of 1 year's duration presented for evaluation of a slowly progressing reddish purple nodule on the right knee of 2 to 4 months' duration. The patient had undergone a matched unrelated donor allogeneic SCT 6 months following diagnosis of AML with subsequent disease progression despite reduction of posttransplant graft-versus-host disease prophylactic immune suppression and a cycle of clofarabine. The patient was hospitalized 2 months after the SCT for neutropenic fever and was found to have vancomycin-resistant enterococcal bacteremia, *Clostridium difficile* colitis, and possible fungal pneumonia. He was treated with voriconazole 200 mg twice daily, which he continued following discharge for antifungal prophylaxis. At the time of discharge, the patient reported that he noticed an asymptomatic "purple papule" on the right knee but did not seek further workup.

Two months later, the patient presented with a fever (temperature, 38.6°C) and leukocytosis (white blood cell count, 130 cells/mL [increased from 53 cells/mL 1 week prior to admission]). Due to his history of immunosuppression and neutropenia, the patient was placed on a broad-spectrum antibiotic regimen of cefepime, daptomycin, and linezolid on admission. Later, vancomycin and gentamicin were added and voriconazole was switched to caspofungin. The patient also received granulocyte-macrophage colony-stimulating factor for neutropenia. During the current hospitalization, blood cultures demonstrated vancomycin-resistant enterococemia, and computed tomography of the chest revealed findings consistent with multilobar pneumonia.

Dermatology was consulted to evaluate the purple nodule on the right knee, which had slowly progressed since his last admission. Physical examination revealed a violaceous, 1.5×1.5-cm nodule with central necrosis covered by black eschar with surrounding erythema (Figure 1). Biopsy specimens for routine histology and a tissue culture were obtained. Histopathologic examination revealed a dense diffuse infiltrate of large hyperchromatic mononuclear cells extending through the dermis, which was consistent with the patient's known AML (Figure 2). Acid-fast bacillus staining was negative for mycobacterial organisms. Grocott-Gomori methenamine-silver stain demonstrated an overwhelming number of septate fungal hyphae with acute-angle branching, concerning for *Aspergillus* species (Figure 3). Of note, an *Aspergillus* serum antigen test was performed at this time and was negative. On repeat review of the routine histologic sections, angioinvasion by hyphae was detected amidst the dense lymphocytic infiltrate (Figure 4).



Figure 1. A large round, reddish purple, centrally ulcerated plaque with dark hemorrhagic crust and a surrounding patch of erythema inferior to the knee on the right lateral lower leg.

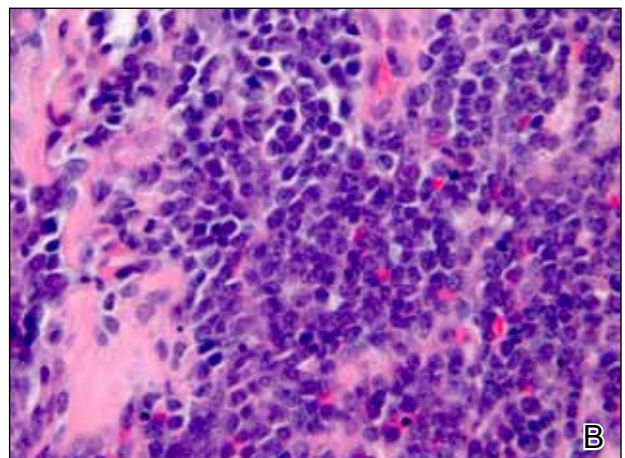
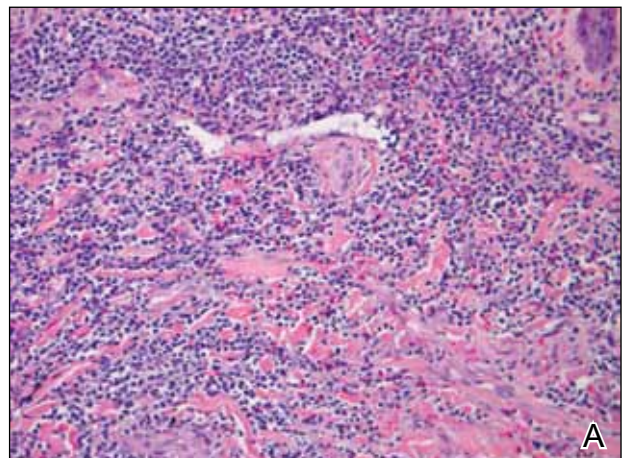


Figure 2. A punch biopsy specimen demonstrated a dense interstitial infiltrate of hyperchromatic cells extending through the dermis (A) (H&E, original magnification ×200). Higher-power view of monomorphic cells exhibited elevated nuclear-cytoplasmic ratios (B) (H&E, original magnification ×600).

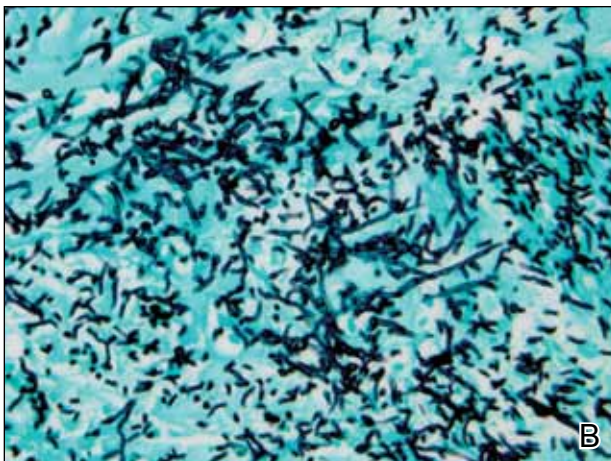
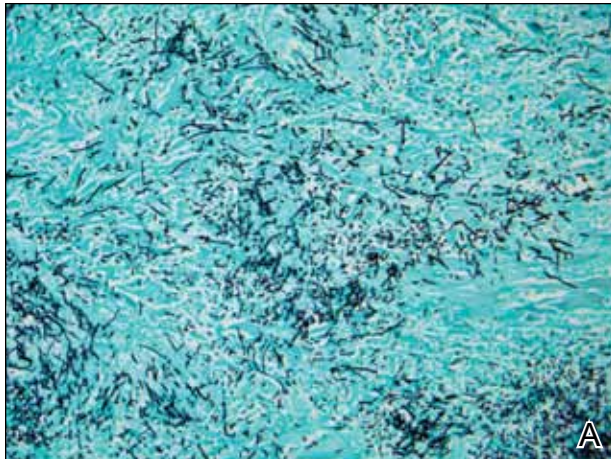


Figure 3. Grocott-Gomori methenamine-silver stain revealed numerous fungal hyphae (A)(original magnification $\times 200$). Higher-power magnification showed septate fungal hyphae (B)(original magnification $\times 600$).

Given the patient's immunocompromised state, the presence of angioinvasive fungi on the skin biopsy, and unresolved pneumonia, the patient was restarted on voriconazole for treatment of likely *Aspergillus* infection. He was continued on chemotherapy for the primary refractory AML and received a donor lymphocyte infusion prior to discharge. After the patient was discharged, the tissue culture grew *Paecilomyces*, a rare fungal species. The patient died 1 week after discharge.

Comment

Leukemia cutis is an extramedullary manifestation of leukemia that appears in 10% to 15% of patients with AML.² The frequency of LC differs widely for the various types of AML, with the majority of cases occurring in the acute myelomonocytic

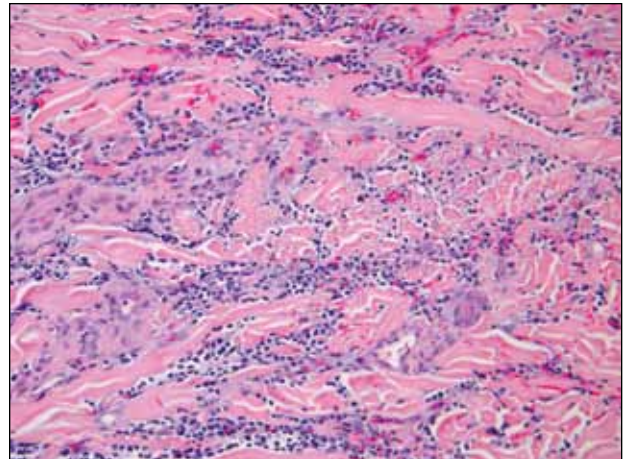


Figure 4. Leukemia cutis and numerous fungal hyphae demonstrated angioinvasion (H&E, original magnification $\times 200$).

leukemia (M4) or acute monocytic leukemia (M5) subtypes.^{3,4} One large study of AML patients (N=381) demonstrated an incidence of LC in 28.6% of patients with the M4 subtype and 42.9% of those with the M5 subtype, with an incidence of only 7.1% of patients with the M1 subtype.³ It occurs less frequently in chronic myeloproliferative diseases.^{2,4}

Leukemia cutis has a wide range of cutaneous manifestations and may present with solitary or multiple papular, nodular, or plaque-like lesions that are red-brown, blue, violaceous, or hemorrhagic.⁴ Leukemia cutis occurs most commonly on the legs, followed by the arms, back, chest, scalp, and face.^{2,4} Leukemia cutis may be hard to distinguish clinically from other conditions such as cutaneous metastases of visceral malignancies, lymphoma, drug eruptions, and opportunistic infections. Leukemia cutis ulcers often measure only a few centimeters in diameter with a firmly adherent purulent or hemorrhagic crust and may occur in unusual locations. These lesions usually are treatment resistant and their persistence may help to lead to diagnosis.⁴

Microscopically, most LC lesions show a perivascular or periadnexal pattern of involvement or a dense diffuse, interstitial, or nodular atypical lymphocytic infiltrate involving the dermis and subcutis with sparing of the upper papillary dermis.² The cytologic appearance of M1 and M2 subtypes of AML are characterized by medium-sized to large mononuclear cells with a light cytoplasm and large basophilic cell nuclei. The M4 and M5 subtypes of AML generally are dominated by medium-sized, round or oval-shaped mononuclear cells that may have eosinophilic cytoplasm and segmented or kidney-shaped basophilic

nuclei.⁴ Immunophenotyping is crucial for diagnosis. In myeloid disorders, there is positive staining with markers of myeloid lineage such as myeloperoxidase, lysozyme, CD34, CD15, CD68, CD43, and CD117.⁵

In our patient, there was an ulcerated dense diffuse dermal infiltrate of large atypical lymphocytes consistent with LC and positive immunostaining consistent with LC associated with AML. Additionally, septate hyphae with acute-angle branching also were noted in the dermal blood vessels on hematoxylin and eosin and fungal staining, demonstrating concomitant fungal infection. Angioinvasion of organisms demonstrated on skin biopsy and persistent pneumonia noted on chest imaging suggested a disseminated infectious process.

Invasive fungal infections are an increasing cause of morbidity and mortality in immunocompromised individuals, including those with hematologic malignancies and hematologic SCTs. Despite an increasing number of antifungal therapies, outcomes are frequently suboptimal with mortality rates often greater than 50% depending on the pathogen and disease.⁶ Thus, there must be a high index of suspicion of infection even when a separate histopathologic diagnosis is available, such as the finding of leukemic infiltrates in this patient's biopsy specimen. A similar case of LC has been reported with concomitant fungal infection involving *Fusarium* and *Enterococcus*.⁷ Patients with leukemic cells may develop leukemic infiltrates in response to cutaneous infection, and a high index of suspicion for 2 related but distinct processes is necessary.

Paecilomyces species are an emerging cause of opportunistic and usually severe human infections.⁸⁻¹¹ The *Paecilomyces* species are saprophytic filamentous fungi that are found worldwide in soil as well as contaminants in the air and water.¹² *Paecilomyces* infection is generally associated with the use of immunosuppressive therapies, implants, or ocular surgery. Among species in this genus, *Paecilomyces lilacinus* and *Paecilomyces variotii* are of clinical importance. Most species have high susceptibility to the newer azoles such as voriconazole.⁶

Conclusion

Despite continued treatment with voriconazole, our patient still developed a rare fungal infection arising in a lesion of LC. He had signs of infection,

including an elevated white blood cell count, fever, and malaise, which are nonspecific clinical findings that could have been attributed to known relapse of systemic leukemia or to known enterococemia. Even in patients on antifungal prophylaxis and with other possible causes of leukocytosis, this case illustrates that there must be a high index of suspicion for angioinvasive fungal infection.

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