Case Letter

Leukemia Cutis Presenting as Scrotal Ulcerations in a Patient With Acute Myelogenous Leukemia

To the Editor:

Cutaneous manifestations of leukemia can be defined as specific or nonspecific skin lesions. Specific skin lesions include papules, nodules, tumors, or ulcerating plaques caused by the infiltration of leukemic cells (eg, leukemia cutis), while nonspecific skin lesions include petechiae, purpura, ecchymosis, or infection. Leukemia cutis is a rare entity that denotes a poor prognosis. We present a case of leukemia cutis with scrotal ulcerations, which heralded the onset of acute myelogenous leukemia (AML). Other causes of painful cutaneous ulcers to consider in a patient with a hematologic malignancy include a variety of infectious and inflammatory etiologies.

A 63-year-old man with long-standing essential thrombocytosis was admitted to the hospital with increasing fatigue and fever. A complete blood cell count revealed anemia, thrombocytopenia, neutropenia, and leukocytosis with 49% blasts. He was immediately placed on broad-spectrum antifungal, antiviral, and antibiotic agents after blood and urine cultures were obtained and while subsequent diagnostic tests were being performed. The patient developed antibiotic-associated *Clostridium difficile* diarrhea after several days in the hospital. The nursing staff noted "genital irritation from feces" while cleaning the patient and reported the findings to the primary team. Dermatology was consulted for treatment recommendations and wound care.

A full-body skin examination revealed discrete and coalescing painful scrotal and perineal ulcers with fecal contamination (Figure 1), along with numerous petechial macules on the anteromedial thighs. The patient was unsure of the onset of the ulcers, but he claimed to have scrotal and perineal pain prior to admission, which suggested their long-standing presence. No

Correspondence: Darshan C. Vaidya, MD, Department of Dermatology, The Warren Alpert Medical School of Brown University, 593 Eddy St, APC 10, Providence, RI 02903 (dvaidya83@gmail.com). crepitus or rapid contiguous spread was noted on serial physical examination over the next 12 to 24 hours. Punch biopsies for tissue culture and histopathologic examination were obtained from the scrotum.

Subsequent diagnostic studies, including a bone marrow biopsy touch preparation that showed 44.7% blasts, along with cytogenetics analysis yielded the diagnosis of AML with an unfavorable karyotype: 5q deletion, 17p deletion, and monosomy 16. Induction chemotherapy with multiple cycles of a combination of cytarabine and daunorubicin was initiated, albeit with an unfavorable response. Blood, urine, bone marrow, and stool cultures remained negative. Serologic testing for herpes simplex virus (HSV), cytomegalovirus, and Epstein-Barr virus was negative.

Histopathology showed an ulcerated epidermis along with a dense dermal perivascular and interstitial infiltrate of myeloid blasts (Figure 2). Lesional cells were diffusely CD68⁺ (Figure 3), myeloperoxidase positive (Figure 4), and CD34⁻, correlating with findings on the bone marrow biopsy. Gomori methenamine-silver, periodic acid–Schiff, and Gram stains highlighted several bacterial cocci, yeast forms, and pseudohyphae on the ulcer surface with no organisms seen in the dermis. Tissue culture did not yield any organisms. There was no overt evidence of HSV infection. The histopathologic features were consistent with a diagnosis of leukemia cutis in the setting of AML. After failed chemotherapy and a complicated hospital course with a poor long-term



Figure 1. Discrete and coalescing painful scrotal and perineal ulcers with fecal contamination.

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Figure 2. Dense dermal infiltrate of myeloid blasts (H&E, original magnification ×10).



Figure 3. CD68 stain highlighted lesional cells (original magnification ×20).

prognosis, the patient and his family opted for home hospice care.

Cases of leukemia cutis manifesting as scrotal ulcerations are rare.¹ Typically, most dermatologic lesions seen in leukemia patients are of a nonleukemic origin.² Leukemia cutis is a rare entity that is associated with a poor prognosis.³ It was accordingly an ominous finding in our patient, as his AML had an unfavorable karyotype and was unresponsive to chemotherapy.

In hematologic malignancies, the morphology and distribution of specific cutaneous lesions tends to vary by the type of leukemia. For example, chronic lymphocytic leukemia presents with papules, nodules, and ulcerating plaques more often than acute lymphocytic leukemia.⁴ Acute myelocytic papules, nodules, or tumors tend to appear on the face, while chronic lesions commonly are seen on the face and extremities.⁵ Generally, for all forms of leukemia, lesions on the genitalia are uncommon.

This case of ulcerative scrotal AML lesions is unique, as myelogenous leukemia cutis typically appears as a tumor that infrequently ulcerates and rarely is located on the genitalia.⁶ Tissue culture, blood culture, viral serology, and biopsy failed to reveal an infectious etiology of the genital ulcerations in our patient. Therefore, the leukemic infiltrate likely preceded the ulceration and an antecedent infection did not elicit an inflammatory response, resulting in the attraction of a leukemic infiltrate to the area. This latter phenomenon also has been demonstrated in the setting of recent surgical scars, trauma, burns, herpes zoster, HSV, and intramuscular injection sites.^{1,7}



Figure 4. Myeloperoxidase stain highlighted lesional cells (original magnification ×10).

The differential diagnosis of painful cutaneous ulcers in the setting of hematologic malignancy includes a variety of infectious and inflammatory etiologies such as ulcerative HSV, cytomegalovirus, Epstein-Barr virus, amoebic skin ulcerations, opportunistic fungal infections, bacterial infections, pyoderma gangrenosum, erythema nodosum, acute febrile neutrophilic dermatosis (Sweet syndrome), histoplasmosis,¹ Fournier gangrene, and vasculitis, all of which were ruled out in our case.

We report a case of a patient with a history of essential thrombocytosis who developed leukemia cutis presenting as scrotal ulcerations, which preceded the diagnosis of AML. Ulcerative leukemic lesions on the genitalia are atypical and extremely rare. However, the lesions are notable because they may be suggestive of a poor prognostic

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outcome, as in our patient whose induction therapy with cytarabine and daunorubicin failed. We propose that leukemia cutis be included in the differential diagnosis of chronic skin ulcerations of the genitalia.

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