Mucocutaneous Granulomatous Disease in a Patient With Hermansky-Pudlak Syndrome

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Report of a Case

An albino woman in her mid-40s with a 30-year history of Crohn disease (CD) was referred for evaluation and treatment of recurrent, exquisitely tender, persistent ulcerations involving the vulvar, perianal, inguinal, and left axillary skin. She noted that her ulcers had partially responded to prednisone, infliximab, and azathioprine administered for her CD several years ago. Her gastrointestinal CD was quiescent, but she continued to have skin manifestations that had been diagnosed as hidradenitis suppurativa. On multiple occasions, exudate cultures from the ulcers grew mixed flora. She had been treated with long-term antibiotics, including cephalexin monohydrate, trimethoprim-sulfamethoxazole, doxycycline hyclate, metronidazole hydrochloride, and rifampin without improvement. A left axillary ulcer had been excised 1 year prior, but it recurred soon after surgery.

In addition to CD, the patient’s medical history included oculocutaneous albinism with horizontal nystagmus, episodic iritis, emphysema, and interstitial lung disease. Her surgical history included a proctocolectomy and end-ileostomy and perianal fistula repair for CD. She had required a blood transfusion following her gastrointestinal surgery but had had no other problems with excessive bleeding.

Her family history was significant for 2 sisters with albinism; her only child, a son, does not have albinism. There was no family history of gastrointestinal diseases.

Physical examination revealed an afebrile and mildly dyspneic woman with pale skin, white hair, light blue eyes, and horizontal nystagmus (Figure 1). Examination of her groin revealed well-demarcated, deep, tender, linear, inguinal and vulvar ulcers with surrounding erythematous, indurated, pink plaques and scaly patches extending to the medial thighs (Figure 2A). The vulva was diffusely swollen, and indurated firm nodules were present on the labia majora and labia minora. Erythematous indurated plaques with central ulceration extended along the bilateral gluteal folds. Examination of her left axilla revealed a deep, crusted ulcer overlying a surgical scar. All ulcers exuded serosanguineous or thick yellow discharge. There were no sinus tracts, comedones, or fistulas in any of the affected sites.

Biopsy samples from the vulvar ulcer and axilla demonstrated deep dermal and subcutaneous lymphoplasmacytic infiltrates with noncaseating granulomas including numerous multinucleated giant cells (Figure 3). Tissue cultures from the vulvar ulcers were negative for atypical mycobacteria and fungi. Mixed flora were cultured from ulcer exudate, including *Staphylococcus aureus*, *Escherichia coli*, and *Pseudomonas aeruginosa*. Tuberculin skin test and serologic analysis for lymphogranulo...
loma venereum had negative results. Routine laboratory studies, including complete blood cell count with differential and a comprehensive metabolic panel, had unremarkable results.

Because of her constellation of cutaneous and systemic findings, the patient was evaluated for Hermansky-Pudlak syndrome (HPS). Electron microscopic examination of her platelets revealed an absence of dense bodies, a finding considered pathognomonic for HPS. Genetic testing identified a homozygous mutation of the \textit{HPS1} gene (c.1189 del C), thus confirming the diagnosis of HPS.

The patient was treated with prednisone, azathioprine, and nabumetone and initially experienced improvement of pain and swelling in the vulvar area, although the ulcers remained unchanged in size. Several months later, however, she developed increased vulvar pain and edema. A combined medical and surgical approach was commenced, including infliximab infusions at a dose of 10 mg/kg monthly, prednisone 30 mg daily, and surgical debridement of granulation tissue in the vulvar ulcers followed by the use of a negative-pressure wound therapy system. No bleeding complications arose as a result of the surgical debridement. She continued to improve 12 months after starting this regimen, with nearly complete healing of the vulvar ulcers, pain relief, and reduction in edema and erythema (Figure 2B). The prednisone dosage was decreased to 10 mg daily, and she continues to receive infliximab 10 mg/kg infusions every 4 weeks; attempts to reduce the dosages of these medications have been unsuccessful because of cutaneous disease flares.

Discussion

Hermansky-Pudlak syndrome is a rare autosomal-recessive genodermatosis. All patients have oculocutaneous albinism and...
variably severe platelet dysfunction. Some patients have pulmonary fibrosis and granulomatous colitis. It is unclear whether our patient’s skin findings were due to cutaneous CD or rather were a manifestation of HPS. Her skin findings were clinically and histologically indistinguishable from those of metastatic cutaneous CD, just as the colitis of HPS is indistinguishable from that of CD. There has been only 1 other report of a patient with HPS with cutaneous granulomatous disease; a 9-year-old Puerto Rican girl with HPS developed granulomatous colitis and subsequently an erosive genital and peristomal eruption, which on biopsy showed nonnecrotizing granulomas. An infectious workup was unrevealing. These 2 cases support the possibility of metastatic cutaneous involvement of granulomatous colitis in HPS. Herein we review the features of cutaneous CD and HPS, genetically distinct diseases that share a common granulomatous inflammatory reaction pattern.

Hermansky-Pudlak syndrome (OMIM 203300) is a rare, autosomally recessive, multisystem disorder resulting from mutations in genes that encode proteins involved in the biogenesis and function of intracellular organelles found in melanocytes, platelets, T cells, neutrophils, and lung epithelial cells. To date, 8 HPS genes have been identified; these genes encode subunits of the adaptor protein complex 3 and biogenesis of lysosome-related organelles complexes. Defects in these distinct genes result in 1 of 8 subtypes of HPS (Table). An inability to form melanosomes in melanocytes and dense bodies in platelets causes the oculocutaneous albino- nism and bleeding diathesis that occur to variable extents in all affected patients. The mechanisms through which mutations in the HPS genes cause the pulmonary, intestinal, immunologic, or, rarely, renal and cardiac manifestations of HPS have yet to be fully elucidated. Defects in the production or transport of lysosomes appear to be central to the pathogenesis.

Patients with mutations in the HPS1 gene tend to have more severe disease and a greater likelihood of pulmonary fibrosis and granulomatous colitis. A granulomatous reaction to intracellular ceroid lipofuscin, a compound believed to be derived from lipid peroxidation of polyunsaturated lipids of subcellular membranes in the lungs and gastrointestinal tract of patients with HPS, may lead to the pulmonary and gastrointestinal manifestations of HPS. Patients with HPS may be unable to eliminate this compound as a result of the aberrant genesis or function of lysosomes. On histopathological examination, however, granulomas are not always formed in proximity to ceroid deposits. In some cases, however, no ceroid deposits are observed, and the clinical and histopathologic features of the enterocolitis of HPS can be identical to those of CD. This has prompted speculation that there may be a genetic linkage between HPS and CD.

The pulmonary fibrosis and the enterocolitis associated with HPS are often recalcitrant to medical therapy. Several recent reports have documented successful treatment of some patients with agents used to treat intestinal CD including the tumor necrosis factor inhibitor infliximab.

**Table. Features of Hermansky-Pudlak Syndrome (HPS) Subtypes**

<table>
<thead>
<tr>
<th>HPS Subtype</th>
<th>Gene</th>
<th>Protein</th>
<th>Clinical Features</th>
</tr>
</thead>
<tbody>
<tr>
<td>HPS1</td>
<td>HPS1</td>
<td>BLOC-3</td>
<td>Restrictive lung disease, granulomatous colitis</td>
</tr>
<tr>
<td>HPS2</td>
<td>AP3B1</td>
<td>APC-3</td>
<td>Neutropenia, recurrent infections</td>
</tr>
<tr>
<td>HPS3</td>
<td>HPS3</td>
<td>BLOC-2</td>
<td>Mild hypopigmentation, no lung or GI disease</td>
</tr>
<tr>
<td>HPS4</td>
<td>HPS4</td>
<td>BLOC-3</td>
<td>Restrictive lung disease, granulomatous colitis</td>
</tr>
<tr>
<td>HPS5</td>
<td>HPS5</td>
<td>BLOC-2</td>
<td>Mild hypopigmentation, no lung or GI disease</td>
</tr>
<tr>
<td>HPS6</td>
<td>HPS6</td>
<td>BLOC-2</td>
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<tr>
<td>HPS8</td>
<td>BLOC1S</td>
<td>BLOC-1</td>
<td>Mild hypopigmentation, no lung or GI disease</td>
</tr>
</tbody>
</table>

Abbreviations: APC, adaptor protein complex; BLOC, biogenesis of lysosome-related organelles complex; GI, gastrointestinal.

Albinism and bleeding diathesis are common to all subtypes.
Crohn disease is a multisystem granulomatous inflammatory disease that typically begins in the gastrointestinal tract. The disease may involve any portion of the gastrointestinal tract. Dermatologic findings are the most common extraintestinal manifestations of intestinal CD, occurring in 10% to 75% of patients. Other extraintestinal manifestations of CD include inflammatory arthropathies, hepatobiliary disease, ocular disease (episcleritis, scleritis, and uveitis), osteoporosis, anemia, nephrolithiasis, urinary tract fistulas, and renal amyloidosis.

The cutaneous manifestations of CD can be generally classified into nutritional deficiency–associated dermatoses (acrodermatitis enteropathica and pellagra resulting from zinc and niacin deficiency, respectively, among patients with CD), reactive dermatoses (pyoderma gangrenosum, erythema nodosum, palmar plantar pustulosis, Sweet syndrome, oral aphthae, epidermolysis bullosa acquisita, necrotic or granulomatous vasculitis, hidradenitis suppurativa, finger clubbing, palmar erythema, pyoderma faciale, and pathergy reactions), and cutaneous granulomatous disease.

Cutaneous granulomatous disease is the most common cutaneous manifestation of CD. It may be contiguous or noncontiguous with the gastrointestinal tract. Contiguous disease affects 36% to 80% of patients with CD and is the most common form of all CD-related skin diseases. Noncontiguous or “metastatic” granulomatous disease, conversely, is the rarest skin manifestation of CD and is defined as the presence of noncasing granulomas arising in the skin at a site distant to and separate from the intestine. Patients with granulomatous skin disease are more likely to have extensive gastrointestinal disease.

Clinical features of metastatic cutaneous CD vary by patient age and anatomic site. The genital skin is the most common site of metastatic cutaneous CD in both adults (59%) and children (75%). In children, genital swelling with or without erythema is the most common presentation, but ulceration is uncommon. In contrast, adult metastatic cutaneous CD is ulcerative in the majority of cases. Deep “knifelike,” linear ulcerations are characteristic, and these preferentially affect the vulva and intertriginous areas. Vegetative, papillomatous nodules mimicking condyloma have been described. Genital infiltration and edema with or without erythema have also been reported in adults. Nongenital metastatic CD more commonly presents with erythematous or violaceous nodules and plaques. Up to 20% of adults and 86% of children present with cutaneous disease prior to the diagnosis of intestinal disease, which makes diagnosis more challenging.

Emanuel and Phelps recently reviewed the histopathologic characteristics of 12 cases of metastatic cutaneous CD. All cases were characterized by a superficial and/or deep granulomatous dermatitis. Granulomas consisting of epithelioid histiocytes were present in 8 cases. Granulomas with Langhans-type giant cells were also common. Five of 12 cases showed ulceration. Granulomatous vasculitis was present in 2 cases, and in 2 others, massive papillary dermal edema was noted.

The clinical differential diagnosis of metastatic cutaneous CD may include infectious etiologies such as cellulitis, erysipelas, tuberculosis, deep fungal infection, tertiary syphilis, filariasis, actinomycosis, schistosomiasis, leishmaniasis, lymphogranuloma venereum, granuloma inguinale, and chronic herpetic simplex virus infection. Appropriate cultures and serologic studies should be used to exclude infectious etiologies. Metastatic cutaneous CD may also be mistaken for dermatitis, foreign-body reactions, intertrigo, Behçet disease, Wegener granulomatosis, lichenoid eruptions, hidradenitis suppurativa, pyoderma gangrenosum, or sarcoidosis. Hidradenitis suppurativa can be ruled out clinically by the absence of interconnecting sinuses and comedones. Deep, linear ulcerations in flexural skin and vegetative papillomatous nodules in the vulva are characteristic findings of genital metastatic cutaneous CD.

The pathogenesis of metastatic cutaneous CD is unclear, but it has been postulated that T lymphocytes react with unknown antigens in the skin and lead to the granulomatous infiltrates of metastatic cutaneous CD.

Metastatic cutaneous CD, particularly genital disease, often responds poorly to treatment. There are no randomized, controlled studies of therapy for this entity. Antibiotics, immunosuppressive medications, and anti-inflammatory agents have been used with limited success.

Conclusions

Our patient’s history of albinism, pulmonary fibrosis, and colitis were suggestive of HPS, which was then confirmed ultrastructurally and genetically. Her gastrointestinal biopsies revealed the histopathologic features typical of both metastatic CD and HPS. It is unclear whether the cutaneous findings in this case are manifestations of CD or, in fact, sequelae of HPS. Cutaneous granulomatous disease has been reported in only 1 other patient with HPS; this is likely due to the rarity of HPS in general. Clinicians should be aware of the possibility of skin manifestations in patients with HPS. Further observation of patients with this rare disease is required to determine whether such cutaneous findings are due to HPS or whether skin lesions originate from a separate disease process, such as CD, that may be in some way genetically linked with HPS.


