Case Report
Peripheral Ulcerative Keratitis with Pyoderma Gangrenosum

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Pyoderma gangrenosum is an unusual necrotizing noninfective and ulcerative skin disease whose cause is unknown. Ophthalmic involvement in pyoderma gangrenosum is an unusual event. Only a few cases have been reported, from which we can highlight scleral, corneal, and orbital cases. Peripheral ulcerative keratitis is a process which destroys the peripheral cornea. Its cause is still unknown although it is often associated with autoimmune conditions. Pyoderma gangrenosum should be included in the differential diagnosis of peripheral ulcerative keratitis. Early recognition of these manifestations can vary the prognosis by applying the appropriate treatment. We introduce a 70-year-old woman who suffered pyoderma gangrenosum associated with peripheral ulcerative keratitis in her left eye. The patient's skin lesions and peripheral keratitis responded successfully to systemic steroids and cyclosporine A.

1. Introduction

Pyoderma gangrenosum (PG) is an unusual necrotizing noninfective and ulcerative skin disease of unknown cause that has been included among the so-called neutrophilic dermatoses. The condition is clinically characterized by necrotic and deep ulcers that are previously preceded by inflammatory pustules [1, 2].

Under the term peripheral ulcerative keratitis (PUK), a group of inflammatory corneal diseases clinically characterized by peripherical corneal thinning, cellular infiltration, ulceration, and variable degree vasoocclusion and injection of the adjacent vascular network are included [3, 4].

Ophthalmic involvement in pyoderma gangrenosum is not a usual event. Only a few cases have been reported, from which we can highlight scleral, corneal, and orbital cases [5–10]. We report a case of PG associated with PUK in a 70-year-old woman. The patient's skin lesions and peripheral keratitis responded successfully to systemic steroids and cyclosporine A.

2. Case Presentation

A 70-year-old woman with a personal history of non-insulin-dependent diabetes mellitus was seen on consultation because of rapid development of an eruption consisting in several ulcerative and painful lesions located on her left leg. Initial lesions were boggy violaceous plaques with pustules that rapidly enlarged for two weeks prior to presentation. The patient was treated with oral and topical antibiotics without results. At the same time, the patient had fever and discomfort and complained of redness and pain and visual acuity decreased in her left eye.

Cutaneous examination revealed scattered shallow ulcers with a necrotic base which were confined to the left leg. The ulcer border was raised, serpiginous, and irregular and it was surrounded by an inflammatory area of erythema (Figure 1). There were no other cutaneous findings.

A wedge-shaped cutaneous biopsy showed neutrophilic abscess formation under areas of ulceration, as well as a dense inflammatory dermal infiltrate composed primarily
Figure 1: Ulcer with a necrotic base, raised border, and halo erythema on the left leg.

Figure 2: Neutrophilic abscess formation under areas of ulceration, as well as a dense inflammatory dermal infiltrate composed primarily of polymorphonuclear leukocytes (H-E ×10).

Figure 3: Ulceration and peripheral stromal infiltrates in the upper and lower limb (slit lamp).

cyclosporine A 3 mg/Kg daily and prednisone 20 mg daily. After three months, the disease was totally resolved with no ocular residual damage and no new active skin lesions were detected.

3. Discussion

PG may appear in healthy patients or in those associated with a variety of systemic diseases. These diseases are present in more than 50% of patients. The most common is the comorbidity inflammatory bowel disease followed by rheumatoid arthritis. Others include immunologic abnormalities, hematologic disorders like monoclonal gammopathy and polycythemia vera, and hematologic malignancies like myeloma, leukemia, lymphoma, and myelodyplasia [1, 2, 5, 7–10]. This relationship supports the hypothesis that the disease may be caused due to underlying defects in the immune system such as abnormalities of cellular or humoral immunity, reduced production of macrophage inhibitory factor, disorder of chemotaxis, and phagocytosis by neutrophils and monocytes. However, a specific immune defect has not been demonstrated [1, 2, 5, 8, 10].

The skin lesions have classic appearance and evolution, starting as a papule or pustules that rapidly progress to a well defined and very painful ulcer with necrotic or mucopurulent debris at the base. The ulcer is surrounded by violaceous undermined borders and an inflammatory halo of erythema. Cutaneous lesions may be present at any site of the skin surface but mucosal membranes are usually spread. It has a propensity to appear on the lower limbs or the trunk and sometimes occurs at areas of the skin previously damaged by trauma or surgical wounds (pathergic phenomenon). The diagnosis of PG is essentially clinical and PG is considered a diagnosis of exclusion. The histopathologic findings are not specific and there are no diagnostic laboratory test markers of the disease [1, 2, 10].

PUK is a destructive process of the peripheral cornea that is often associated with autoimmune conditions including rheumatoid arthritis, Sweet syndrome, systemic lupus erythematosus, Wegener’s granulomatosis, and polyarteritis nodosa. Patients may present decreased visual acuity,
Table 1: Description of cases reported of pyoderma gangrenosum with peripheral ulcerative keratitis (PUK). Literature review.

<table>
<thead>
<tr>
<th>Authors</th>
<th>Gender</th>
<th>Age</th>
<th>PUK</th>
<th>Association</th>
<th>Therapy</th>
<th>Response to therapy</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bouchard et al. [5]</td>
<td>Male</td>
<td>37</td>
<td>Bilateral</td>
<td>Chronic myelogenous leukaemia</td>
<td>Systemic corticosteroids</td>
<td>Complete response</td>
</tr>
<tr>
<td>Bishop and Tullo [6]</td>
<td>Female</td>
<td>59</td>
<td>Left eye</td>
<td>Monoarticular arthritis</td>
<td>Cyclophosphamide and systemic corticosteroids</td>
<td>Improved but with intermittent flares of ocular diseases</td>
</tr>
<tr>
<td>Bishop and Tullo [6]</td>
<td>Male</td>
<td>56</td>
<td>Right eye</td>
<td>Leukocytoclastic vasculitis</td>
<td>Systemic corticosteroids</td>
<td>Improved but with recurrence of ocular disease</td>
</tr>
<tr>
<td>Wilson et al. [7]</td>
<td>Female</td>
<td>60</td>
<td>Left eye</td>
<td>Rheumatoid arthritis and leukocytoclastic vasculitis</td>
<td>Cyclosporine A and systemic corticosteroids</td>
<td>Improved but with intermittent flares of ocular and skin diseases</td>
</tr>
<tr>
<td>Brown et al. [8]</td>
<td>Male</td>
<td>54</td>
<td>Right eye</td>
<td>Chronic obstructive pulmonary disease and diabetes mellitus</td>
<td>Systemic corticosteroids and azathioprine</td>
<td>Complete response</td>
</tr>
<tr>
<td>Teasley et al. [9]</td>
<td>Female</td>
<td>30</td>
<td>Left eye</td>
<td>Graves’ disease</td>
<td>Dapsone</td>
<td>Complete response</td>
</tr>
<tr>
<td>Fournié et al. [10]</td>
<td>Male</td>
<td>78</td>
<td>Left eye</td>
<td>Multiple myeloma</td>
<td>Cyclophosphamide, systemic corticosteroids, and human intravenous immunoglobulins</td>
<td>Complete response</td>
</tr>
</tbody>
</table>

blindness, eye pain, redness, or irritation. The diagnosis is confirmed by slit lamp examination [3, 4, 7–10].

As with PG the aetiology of PUK is poorly understood, the postulated reasons include autoimmune reactions to corneal antigens, circulating immunocomplex deposition, vasculitis, and hypersensitivity reactions to exogenous antigens. PUK may result from humoral or cell-mediated immune mechanisms or both, causing obliterator microangiitis at the level of the limbal vascular arcades. Subsequent leakage of inflammatory cells with destructive collagenases and proteases leads to scleral inflammation and destruction [3–5, 8, 10].

There have been only seven reported cases of PUK associated with PG [5–10] (Table 1). There have been four reported cases in males and three cases in females. The age of clinical appearance varies between 30 and 78 years. PUK is usually unilateral and the left eye is the most frequently affected. There is only one case of bilateral ocular involvement [5]. Other autoimmune disorders were associated like monoarticular arthritis [6], rheumatoid arthritis [7], leukocytoclastic vasculitis [6, 7], and Graves’ disease [9], as well as diseases producing immunosuppression like chronic obstructive pulmonary disease [8], diabetes mellitus [8], multiple myeloma [10], and chronic myelogenous leukemia [5]. All the patients were prescribed systemic corticosteroids. All but 1 were treated with immunosuppressive agents. These include cyclophosphamide, cyclosporine A, azathioprine, dapsone, and human intravenous immunoglobulins. All the patients had an initial adequate response but the course of the disease varied between cases with some relapsing cases. PG and PUK do not follow a parallel course.

Our patient had no other autoimmune disease associated and she had a complete response of PG and PUK with cyclosporine A and systemic corticosteroids. PG and other autoimmune disorders or diseases producing immunosuppression should be considered in the differential diagnosis of PUK. Early recognition of these manifestations can lead to the application of appropriate treatment improving the prognosis [3–10].

Acronyms

PG: Pyoderma gangrenosum
PUK: Peripheral ulcerative keratitis.

Conflict of Interests

The authors declare no conflict of interests.

References


