Pemphigus herpetiformis: A rare clinical variant of pemphigus

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Abstract

Pemphigus herpetiformis is an autoimmune blistering disease. We report a case of pemphigus herpetiformis, a generalized form, manifested as pruritic grouped vesicles at extremities and trunk, in a seventy years old man. Neutrophil dominant inflammatory substrate was seen in histopathology. Direct immunofluorescence revealed IgG reactivity in net like pattern at upper epidermis, confirming pemphigus. The patient responded well to prednisolone and colchicine combination initially and maintained remission later on colchicine alone. Colchicine has anti-mitotic, anti-inflammatory and immunosuppressive mechanism of actions and proven benefits in neutrophilic dermatoses. Therefore we recommend colchicine as mono therapy or in combination with immunosuppressive, for the treatment of pemphigus herpetiformis where neutrophil is predominant inflammatory infiltrate histologically.

Keywords: Pemphigus, Herpetiform, Pruritic, Vesicles

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Introduction

Pemphigus is an autoimmune blistering disease. Many variants have been described but pemphigus herpetiformis is relatively uncommon. Clinically it presents as dermatitis herpetiformis and immunopathologically as pemphigus. Incidence is equal in both genders, with mean age of onset around 60 years. These patients usually present with intensely pruritic vesicles and blisters with variable duration. Two forms have been described; generalized and localized types. There is a paucity of studies on pemphigus herpetiformis, so whether it is an atypical variant of pemphigus foliaceus or it is separate entity is so far not known.
Case Report

atorvastatin 10mg, amlodipine 5mg and diltiazem 60mg. He had unexplained iron deficiency anaemia (hemoglobin-8.4gm/dl, ferritin-4.73ng/ml, total iron binding capacity-386ug/dl, transferring saturation % -10%), vitamin B12 deficiency (155.4pg/ml) and blunting of duodenal villi with presence of helicobacter pylori in the endoscopic biopsy. Occult blood test was negative. After taking iron and vitamin B12 supplements, his hemoglobin increased to 11gm/dl. Apart from occasional epigastric pain, he does not give history of any gastrointestinal and neurological symptoms. He doesn’t recall worsening of the skin conditions with any particular diet. Clinically his iron deficiency anemia and blunting of duodenal villi couldn’t be correlated with his skin condition.

![Figure 1: Intensely pruritic vesicles and urticated papules arising on an erythematous base at right arm](image1)

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![Figure 2: Excoriated scabbed lesions resolving with post inflammatory hypopigmentation symmetrically at bilateral lower limbs](image2)

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![Figure 3: Few intact grouped vesicles with superficial de-epithelization at the sacral region](image3)

Figure 3: Few intact grouped vesicles with superficial de-epithelization at the sacral region

On physical examination, he had few intact itchy grouped vesicles arising on normal to urticated skin and crusted scabbed lesions over the upper chest, sacral region, extensor of both upper and lower limbs bilaterally (Figure 1, 2 & 3). These lesions healed with faint hypopigmentation and no scarring. Nikolsky’s sign was negative. Clinically no secondary infection was seen. Mucous membrane, nail, scalp, palms and soles were spared. Rest of the physical examination was normal. So a presumptive diagnosis of dermatitis herpetiformis with differential diagnoses of bullous pemphigoid- vesicular type, pemphigus foliaceus- dermatitis herpetiformis and bullous lupus was made.

Skin biopsy was taken from left arm. Histologically epidermis showed mild acanthosis, collection of neutrophils at malphigian layer of the epidermis with few areas of microabcess (Figure 4). Upper dermis showed mild papillary dermal edema with few collection of neutrophils. No clear separation or cleft was seen. Hence direct immunofluorescence (DIF) was suggested for confirmation.

![Figure 4: Histopathology showing neutrophilic spongiosis with papillary dermal edema](image4)

Figure 4: Histopathology showing neutrophilic spongiosis with papillary dermal edema
Hence a diagnosis of pemphigus herpetiformis (PHF) was made on the basis of clinical features where dermatitis herpetiformis like intensely itchy grouped vesicles in generalized distribution were present along with immunopathologic features of pemphigus. Treatment was started with prednisone 40 mg/day and topical clobetasol. His itching and occurrence of new lesion substantially reduced, so after two weeks steroid was tapered to 20mg/day and further reduced and colchicine 0.5mg twice daily was added for next two weeks. No recurrence was noted at subsequent follow-up, therefore systemic steroid was finally stopped and was continued only on colchicine for next two months only. There was no relapse at six months after all the drugs were stopped. He continued his previous cardiac medicines and vitamin supplements during the treatment and remission.

Discussion
Pemphigus herpetiformis (PHF) was first introduced by Jablonska and colleagues as a variant of pemphigus. PHF is atypical variant of pemphigus that combines the clinical features of dermatitis herpetiformis with the immunopathologic features of pemphigus. Mean age of onset around is 60 years. However childhood cases has also been reported. Patients affected with pemphigus herpetiformis usually have a subacute onset of disease. Approximately half the patients experience severe pruritus. The clinical distribution of PHF is almost invariably generalized, involving both the trunk and limbs, characteristic skin manifestations are pruritic erythematous vesicular, bullous or papular lesions in herpetiform pattern, similar to those of dermatitis herpetiformis. However localized form of PHF has been described. PHF occurrence in association with malignancies such as lung cancer, prostate cancer, and cutaneous angiosarcoma has also been reported. Histologically demonstrated eosinophil and/or neutrophil infiltration into the epidermis may be relevant pathogenically in the disease process of pemphigus herpetiformis. Huhn et al found that the inflammatory infiltrate in patients with PHF was 68% eosinophil-dominant, 16% neutrophil-dominant, and 16% mixed eosinophil/neutrophil. In the neutrophil-dominant subset, epidermal cells secrete a neutrophil chemokine interleukin 8 (IL-8), which apparently is induced by IgG autoantibodies to desmoglein and may be responsible for the recruitment of neutrophils to the epidermis, resulting in the subsequent blistering process. Pemphigus herpetiformis appears to be mediated by the immunoglobulin G (IgG) class of autoantibodies that target the skin epidermis desmoglein components, commonly Desmoglein 1 and 2.

DIF helps detect molecules such as immunoglobulins and complement components within biopsy specimens. For bullous diseases, DIF is performed using perilesional skin, that is, normal-appearing skin immediately adjacent to a lesion (vesicle, bulla, urticarial plaque, or erythematous patch). Indirect immunofluorescence (IIF) confirms the presence in the patient's serum of IgG-circulating autoantibodies that bind to epidermal cell surfaces. The titer of autoantibodies often parallels the clinical level of activity of pemphigus herpetiformis and often is helpful in following the patient's disease activity after the diagnosis has been established. Other tests such as immunoblotting, ELISA, immunoprecipitation,
chest xray, prostate specific antigen are recommended if symptoms suggest.

Pemphigus herpetiformis is responsive to anti-inflammatory and immunosuppressive medications. Dapsone is the drug of choice if the patient tolerates the treatment, systemic corticosteroid and immunosuppressives, monoclonal antibody anti-CD20 (rituximab). We treated our patient with prednisolone and colchicine combination initially and colchicine alone later as a steroid sparing agent. Colchicine has anti-mitotic, anti-inflammatory and immunosuppressive mechanism of actions and proven benefits in neutrophilic dermatoses. In our patient, neutrophil dominant infiltrate was seen histologically and he maintained remission on colchicine alone and 6 months after it was stopped.

**Conclusion**

We would like to stress the importance of performing direct immunofluorescence to test for the presence of an autoimmune blistering disorder when histology reveals neutrophilic and/or eosinophilic spongiosis for correct diagnosis. Also, colchicine can be safe steroid sparing agent in pemphigus herpetiformis with neutrophil dominant infiltrate.

**References**


