Pemphigus foliaceus as a differential diagnosis in vesicobullous lesions
Pênfigo foliáceo como diagnóstico diferencial em lesões vesicobolhosas

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INTRODUCTION
The differential diagnosis of vesicobullous diseases can be considered a challenge in the daily clinic of internal medicine and dermatology. The lesions present an acute onset, and several mechanisms may be responsible for their formation.

Blisters occur at different skin levels. Initially, in the histological, immuno-histological and electron microscopy evaluations, the goal is to find the plane in which the loss of cell adhesion occurs. Histological diagnosis becomes mandatory to complement the investigation, although, only the histologic assessment and pathogenic mechanisms do not necessarily assure a correct diagnosis. Therefore, immunofluorescence techniques are a valuable supplement.

Pemphigus foliaceus (PF) is a chronic autoimmune bullous skin disease, which is histopathologically characterized by the formation of intra-epidermal bulla with acantholysis, and immunologically by circulating autoantibodies to the epidermis, which are responsible for the cutaneous lesions.

CASE REPORT
A 78-years old Caucasian male, born and raised in the urban areas of Jundiaí, State of São Paulo, Brazil, presented “burst wounds in the body” for 20 days. He referred that approximately during 2 months, expressive pruritic lesions appeared on the scalp associated with local burning (burning sensation) with craniocaudal progression. He also reported that the lesions progressed to cervical, trunk, abdomen and underarms areas involvement for approximately 20 days. He had used zinc oxide ointment and bath with Stryphnodendron barbatiman, which are popular generic treatments, with worsening of the lesions. He had no previous skin lesions, and denied any mucosal involvement or other clinical complaints.

Previously he had hypertension, type 2 diabetes mellitus, and obesity. He denied any family history of skin lesions. His current prescription consists in omeprazole, losartan, simvastatin, aspirin and subcutaneous NPH insulin.
Dermatological examination revealed erythematous and bullous lesions, with thick crusts on the surface. Lesions were distributed in scalp, trunk, back, abdomen and underarms. A few bullous lesions on the back presented with flaccid bulla. Nikolsky sign (slight rubbing of the skin results in exfoliation of the outermost layer) was clearly positive. After this finding, dermatologists made the clinical diagnosis of PF. Treatment was started with prednisone 1mg/kg per day associated with antibiotic therapy (initially with cephalaxin and later teicoplanin to control a secondary nosocomial infection), topical corticosteroid therapy with hydrocortisone and an immnosuppressant corticosteroid-sparing agent (azathioprine). After hospital discharge, histologically examination of biopsied lesions confirmed PF.

This patient is currently followed as an outpatient and still presents partial remission, with dependence on corticosteroid therapy at lower doses, after several months.

Histological analysis showed a subcorneal lesion with inflammation (Figure 1) and apoptotic keratinocytes (Figure 2), characteristic of PF.

**DISCUSSION**

The patient had a recent history of itching with burning sensation, first in the scalp region, which had a fast craniocaudal progression to the neck, back, abdomen, and underarms. On examination, he presented with flaccid vesicular and bullous lesions with areas of ulceration and thick crusts. The positive Nikolsky sign suggests an intra-epithelial cleavage.

Due to the frequently heterogeneous clinical features, it is often not possible to make an exact diagnosis of bullous dermatoses solely on the basis of the clinical findings. Initially, in the histological and immuno-histological evaluations, and even in electron microscopy, the goal is to find the plane in which the loss of cell adhesion occurs. From the identification of this region, we have some possible pathologies: sub-stratum corneum (represented by PF, impetigo, pustular psoriasis and neonatal toxic erythema), suprabasal (spongiotic dermatitis, Darier’s disease, Hailey-Hailey’s disease, multiform erythema and pemphigus vulgaris), subepidermal (porphyria cutanea, epidermolysis bullosa acquisita, vesicular lichen, bullous pemphigoid, dermatitis herpetiformis, linear IgA bullous dermatosis) (Figure 3).

Once the cleavage site is identified, the mechanism responsible for the lesion is investigated. A histologic finding of intraepidermal bullae with the presence of acantholysis is the main histopathological feature of autoimmune intraepidermal diseases from the pemphigus group. The autoimmune intraepidermal bullous diseases include defined entities from the pemphigus family: pemphigus vulgaris, PF, pemphigus herpetiformis, IgA pemphigus, paraneoplastic pemphigus and drug-induced pemphigus. In pemphigus vulgaris, there are intraepidermal, suprabasal acantholysis, loss of keratinocytes adhesion, neutrophilic and eosinophilic granulocytes, and a scant perivascular round-cell
infiltrate is found in the upper dermal vascular plexus. Pemphigus foliaceus presents a superficial subcorneal acantholysis in the upper stratum spinosum or granulosum and a slight inflammatory infiltration of the upper dermis. In paraneoplastic pemphigus, interface dermatitis with vacuolar degeneration of basal keratinocytes and keratinocytes necrosis with slight acantholysis are seen histologically in association of a lichenoid infiltration in the vicinity of the dermo-epidermal junction zone. IgA pemphigus exhibits an intraepidermal or subcorneal infiltration of neutrophils and classical signs of acantholysis are usually missing. In pemphigus herpetiformis, acantholytic cells are usually located subcorneally, and eosinophilic spongiosis is most typical for this variant (Chart 1).

| Pemphigus diagnosis                  | Histology                                    | Autoantigen |
|--------------------------------------|----------------------------------------------|-------------|
| Pemphigus vulgaris                   | Intraepidermal suprabasal acantholysis       | Dsg 3       |
|                                      | Tomb stone appearance                        | Dsg 1       |
| Pemphigus foliaceus                  | Superficial subcorneal acantholysis          | Dsg 1       |
|                                      | Dyskeratosis                                 | Dsc 1       |
| Pemphigus herpetiformis              | Subcorneal acantholysis                      | Dsg 1       |
|                                      |                                              | Dsg 3       |
|                                      |                                              | Dsc 1       |
|                                      |                                              | Dsc 3       |
|                                      |                                              | 178kDa protein |
| IgA pemphigus                        | Intraepidermal infiltration by neutrophils   | Dsg 1 (IgA) |
|                                      | or subcorneal infiltration by neutrophils    | Dsg 3 (IgA) |
|                                      | usually without classical acantholysis       | Dsc (IgA)   |
| Drug-induced pemphigus               | Suprabasal acantholysis                      | Dsg 1       |
|                                      |                                              | Dsg 3       |
| Paraneoplastic pemphigus             | Interface dermatis, lichenoid infiltration   | Enoplakin    |
|                                      | of the dermo-epidermal junction, keratinocyte necrosis | Pariplakin (IgA, IgG) |

The major pemphigus sub-types are pemphigus vulgaris and PF. In this disease, IgG autoantibodies (desmoglein-1 and/or 3) break the connection between desmosomes, causing acantolisis. The pathophysiological process is controversial, but it is certain that pemphigus vulgaris causes mucosal and skin lesions, reaching the suprabasal layer by producing desmoglein-3. Pemphigus foliaceus only causes skin lesions, reaching the subcorneal layer by production of desmoglein-1.

Fogo selvagem or “wild fire”, is an endemic form of PF in rural regions of Brazil, and appears to be triggered by an environmental agent, not yet identified.

Direct immuno-fluorescence is used to determine the type of immunoglobulin (IgG, IgA and IgM), and complement fraction (C3, C5b-9) or fibrinogen may be necessary to identify the underlying pathophysiology and its distribution pattern on the basis of the vesicle or bubble in the epidermis, basal linear, or granular membrane. In the case of PF, an intercellular deposition of IgG is observed. Direct immuno-fluorescence is similar between pemphigus vulgaris and PF, but once they have been differentiated by histopathology, direct immuno-fluorescence findings are suggestive of PF up to 100%. It is important to emphasize this is a severe illness and potentially fatal. The diagnosis needs to be quick and the treatment is based on immunosuppression with steroids (topical and systemic), corticosteroid-sparing immunosuppressants, immunoglobulins, and anti CD-20.

**REFERENCES**

1. Kumar V, Abbas AK, Aster JC, Fausto N, Robbins & conran pathologic basis of diseases. 8th ed. United States: Saunders; 2009. p. 1165. [Series Robbins Pathology].
2. Levitt J, Nazarian R. Vesiculobullous diseases. Dermatol Educ. 2011;1-22. DOII:10.2310/7900.1105.
3. Abesman J, Grover R, Helm TN, Beutner EH. Can direct immunofluorescence testing still be accurate if performed on biopsy specimen after brief inadvertent immersion in formalin? J Am Acad Dermatol. 2011;65(1):106-11.
4. Kneisel A, Hertl M. Autoimmune bullous skin diseases. Part 2: diagnosis and therapy. J Dtsch Dermatol Ges. 2011;9(11):927-47. Review.
5. Radó J. Autoimmune blistering diseases: histologic meaning. Clin Dermatol. 2011;29(4):377-88.
6. Hertl M, Eming R. Pemphigus vulgaris. CME Dermatol [Internet]. 2009 [cited 2009 July 31];4(2):94-115. Available from: www.akademos.de/derma
7. Stanley JR. Pemphigus. In: Freedberg IM, Eisen AZ, Wolff K, Austen KF, Goldsmith LA, Katz SI, editors. Fitzpatrick’s dermatology in general medicine. 6th ed. vol. 1. New York: McGraw-Hill; 2003. p. 558-67.
8. Cunha PR, Baraviera SR. Autoimmune bullous dermatoses. An Bras Dermatol. 2009;84(2):111-24. Review.
9. Shams-Davatchi C, Valikhani M, Daneshpazhooh M, Esmaeil N, Balighi K, Hallaji Z, et al. Pemphigus: analyses of 1209 cases. Int J Dermatol. 2005;44(6):470-6.
10. Inchara YK, Rajalakshmi T. Direct immunofluorescence in cutaneous vesiculobullous lesions. Indian J Pathol Microbiol. 2007;50(4):730-2.