

Psoriasis with verrucous appearance: A case report

Sarra Ben Rejeb¹, Amen Dhaoui¹, Dorra Ben Ghachem¹, Asmahane Souissi², Khadija Bellil¹

¹Department of Pathology, FSI Hospital, La Marsa, Tunisia, ²Department of Dermatology, FSI Hospital, La Marsa, Tunisia

Corresponding author: Dr. Sarra Ben Rejeb, E-mail: sarrabenrejeb88@yahoo.fr

ABSTRACT

Psoriasis verrucosa is a rare clinical variant of psoriasis with peculiar histologic features. Only few cases have been reported in the literature. We herein report a rare case of psoriasis with verrucous appearance occurring in a 63 year-old woman who presented with verrucous and scaly erythematous plaque of the legs which was developed thirty years ago. The biopsy specimen showed regular psoriasiform epidermal hyperplasia with acanthosis, hyperkeratosis, and focal spongiosis with a superficial perivascular infiltrate. The patient was diagnosed with verrucous psoriasis. Recognition of this entity should preempt confusion with verruca vulgaris or other entities capable of producing wart-like epidermal changes.

Key words: Psoriasis; Verrucous; Histology

INTRODUCTION

Psoriasis is an inflammatory disorder characterized by peculiar skin and joint manifestations. The most common clinical presentation is a scaly erythematous plaque with thick silvery white scale. Psoriasis with verrucous appearance is a rare variant of the disease with characteristic clinical and pathological features; it might be confused with many other lesions including warts, epidermal nevus, contact dermatitis, eczema and fungal infection [1,2].

We herein describe another case of this rare entity. The clinicopathological features of this disease are discussed.

CASE REPORT

A 63 year-old woman with no past medical history presented with erythematous, scaly lesions of the lower extremities developed thirty years ago. They were painless but were progressively growing, causing significant physical disfigurement and discomfort on wearing shoes. Physical examination revealed verrucous plaques of the legs (Fig. 1). A skin biopsy of these lesions has been performed.

Pathologic examination revealed psoriasiform epidermal hyperplasia of rete ridges with prominent papillomatosis, marked parakeratosis and hyperkeratosis. It also showed epidermal hypogranulosis, thin suprapapillary epidermal plates, focal spongiosis of the Malpighi mucous body and a superficial perivascular infiltrate. No inflammatory collection has been noted within the epidermal layers (Figs. 2 and 3). Koilocytic changes were not observed.

Clinical and pathological findings were consistent with verrucous psoriasis (VP). The patient was treated with local dermocorticoid. The patient has not been seen since the beginning of the treatment.

Prior to the study, patient gave written consent to the examination and biopsy after having been informed about the procedure.

DISCUSSION

Verrucous psoriasis is a rare variant of the disease with less than thirty cases reported in the literature [3]. It is mostly occurring on men with a sex ratio of 1.6 and a mean age of 53 year old [1,4]. Clinically, VP is characterized by a scaly erythematous plaque with

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Figure 1: Warty plaques of the foot.

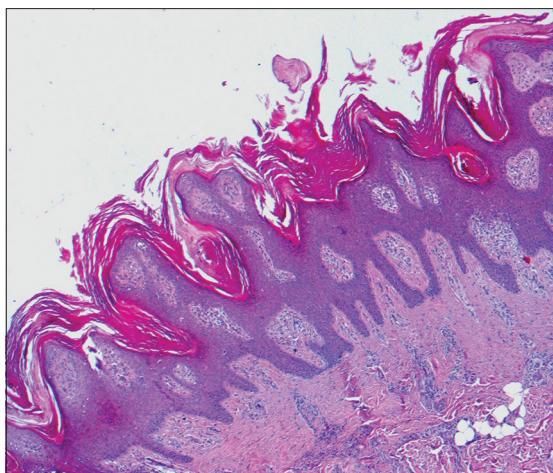


Figure 2: Wart-like appearance of the epidermis (H&E x400).

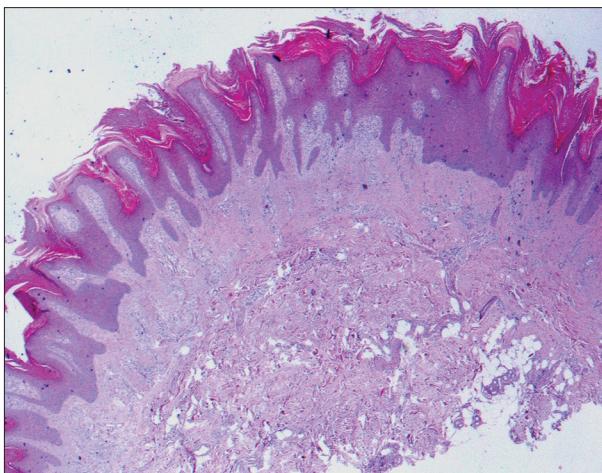


Figure 3: Epidermal hyperplasia with parakeratosis and prominent papillomatosis (HE x200).

wart-like changes. It is commonly occurring on friction areas such as elbows, hands, knees and feet. VP may be confused with many other benign lesions, including

verruca vulgaris, epidermal nevus, contact dermatitis, eczema and fungal infection [1]. However, a verrucous carcinoma must be ruled out on biopsy. Microscopically, the lesion has a characteristic feature of psoriasis with wart-changes, it is typically showing parakeratosis, epidermal psoriasiform hyperplasia with elongation of rete ridges, thin suprapapillary epidermal plates, thinning of the granular layer, and dilated, tortuous capillaries with an inflammatory infiltrate, which may contain admixed neutrophils in the papillary dermis. Munro abscesses and spongiform pustules are frequently noted. In our case, despite the absence of characteristic neutrophils collection's of psoriasis within the epidermal layer, the correlation of clinical and other histologic features including papillomatosis and epithelial buttressing suggested the diagnosis. Moreover, Munro abscesses might be absent in 35% of authentic psoriasis. Classically, absence of koilicytic changes and Human papilloma virus (HPV) immunostaining support the diagnosis of verrucous psoriasis.

The etiology of verrucous psoriasis remains unclear. It has been associated in some cases to lymphatic obstruction, microangiopathy, diabetes and obesity [5,6]. Other authors consider it as a progressive form of vulgaris psoriasis [1,5]. The verrucous changes in this case have been related to repeated trauma.

However, although it is considered to be a variant from psoriasis, little is known about its adequate treatment because of poor response to classic local therapy (dermocorticosteroid, vitamine D, puvatherapy). Some authors reported the efficacy of etretinate, adalimumab and methotrexate. However, further investigations are required to determine an optimal treatment for this rare entity.

CONCLUSION

Verrucous psoriasis is a rare variant of psoriasis that might prompt consideration of verruca vulgaris. Combination of clinical and pathological findings is required for precise diagnosis. Because of its rarity, no codified therapy has been established yet.

Consent

The examination of the patient was conducted according to the Declaration of Helsinki principles. Written informed consent was obtained from the patient for publication of this article.

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