Atrophying Pityriasis Versicolor

Pityriasis versicolor atrófica

To the Editor:

Pityriasis versicolor (PV) is a common superficial fungal infection caused by yeasts of the genus *Malassezia*. In clinical terms, it generally presents on the trunk as round or oval macules that vary in color from hypopigmentation to blackish.

Its symptoms are easily recognizable, with several atypical forms reported, including the variant with atrophic lesions.

A 28-year-old man with plaque psoriasis treated with topical corticosteroids consulted for the appearance of cutaneous abnormalities that he perceived as being different from his psoriasis lesions. The physical examination revealed several circular areas on the back and the extensor surface of the upper right limb. The areas measured 0.5 cm to 4 cm and were depressed and atrophic in appearance (Fig. 1). Some were hypopigmented, whereas others were slightly pink; some had mild desquamation, whereas others were similar to typical psoriasis lesions (Fig. 2). The Wood lamp examination did not reveal fluorescence. The skin biopsy revealed laminated hyperkeratosis in the epidermis, with flattening of the rete ridges and an edematous appearance with vascular dilations in the dermis that was more evident in the papillary dermis. The vessels were surrounded by a discreet lymphocytic infiltrate. Periodic acid-Schiff staining revealed abundant short, septate hyphae and spores between the layers of keratin. Orcein staining revealed diminished elastic fibers in the papillary dermis and fragmentation thereof in the superficial reticular dermis (Fig. 3).

Direct examination of the lesions revealed the presence of yeasts and pseudomycelium. The patient was prescribed itraconazole 200 mg/d for 7 days combined with topical flutrimazole at night for 1 month. The topical corticosteroid was replaced by...
calcitriol ointment for treatment of his psoriasis. Although improvement was slow, the patient had fully recovered at 6 months, with no new lesions after 1 year of follow-up.

Atrophying or atrophic PV is an uncommon presentation of this condition.\textsuperscript{2-10} It manifests clinically as depressed erythematous or hypopigmented areas with discreet desquamation located mainly on the trunk, generally the back and shoulders. The lesions are usually numerous, tend to group together, and measure a few millimeters to several centimeters, although they are generally uniform on the same patient. The diagnostic suspicion can be confirmed by direct examination, which reveals accumulations of yeasts and short hyphae; if these occur together, they have the typical "spaghetti and meatballs" appearance. Skin biopsy may help in atypical cases such as the present one. The most characteristic and differentiating histopathological findings of the atrophic variant include an epidermis with flattened rete ridges and, especially, reduction and fragmentation of the dermal elastic fibers.\textsuperscript{5,9} The differential diagnosis includes processes that induce skin atrophy, that is, anetoderma, morphea/atrophoderma, mycosis fungoides, sarcoidosis, and corticosteroid-induced atrophy.\textsuperscript{6,9,10}

The mechanism by which species of Malassezia can induce atrophy in the skin remains unclear. Since many of the patients reported were being treated with topical corticosteroids\textsuperscript{3-7} or systemic corticosteroids,\textsuperscript{9} atrophy was associated with the atrophogenic effect of the drugs.\textsuperscript{3-7} The fungal infection is thought to alter the epidermal barrier, thus increasing absorption of corticosteroids in the PV lesions and inducing local cutaneous atrophy. However, in corticosteroid-induced atrophy, in addition to epidermal atrophy and vasodilation, we can see atrophy of skin adnexa and reduction and fragmentation of dermal collagen, whereas in atrophic PV, we do not see alteration of collagen but alteration of the elastic fibers.\textsuperscript{5,6,9,10} We did not find collagen abnormalities in the case we report, suggesting the possibility of another pathogenic mechanism, given that cases not associated with corticosteroids have been reported.\textsuperscript{5,10} Crowson and Magro\textsuperscript{7} proposed that this form of PV appeared through a type 1 helper T-cell immune response to Malassezia antigens, in which histocytes recruited and activated by interferon-\textgamma would generate the elastases responsible for the dermal elastolysis observed. Flattening of the rete ridges, on the other hand, is mediated by cytokines, such as tumor necrosis factor \textalpha and interleukin 1\beta, which can inhibit the nuclear factor-kB pathway of keratinocytes to induce apoptosis and hinder proliferation thereof.\textsuperscript{7} In parallel, the reduced thickness and loss of hair in areas of PV has also been related to this mechanism.\textsuperscript{11} Furthermore, a type 2 helper T cell response, which could

**Figure 2** Psoriatic plaques in the center of depressed areas.

**Figure 3** Epidermis with abundant hyphae and spores in the stratum corneum and flattening of the rete ridges (periodic acid–Schiff, original magnification \times 200). The inset shows diminished fine elastic fibers in the papillary dermis (arrow) and fragmented elastic fibers in the superficial reticular dermis, as well as vascular ectasia (orcein stain, original magnification \times 100).
favor fungal overgrowth, has been implicated, together with the type 1 response, in the pathogenesis of atrophic PV.\(^\text{10}\)

Treatment of this variant of PV is standard,\(^\text{10}\) although it has been suggested that longer regimens may be necessary and even that oral and topical approaches should be combined.\(^\text{11}\) Obviously, corticosteroids should be withdrawn.\(^\text{11}\) As a rule, the atrophy disappears; therefore, compared with other atrophying diseases, atrophying PV has a good prognosis.

**Conflicts of Interest**

The authors declare that they have no conflicts of interest.

**References**


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**Human Papilloma Virus and the Risk of Squamous Cell Carcinoma Arising in Hidradenitis Suppurativa**

**Virus del papiloma humano y riesgo de desarrollo de carcinoma escamocelular en la hidradenitis suppurativa**

To the Editor:

We report the case of a 51-year-old man who was referred to the dermatology department with a painful, suppurative nodule in his left groin that had appeared 1 year previously and was refractory to various antibiotic regimens prescribed by his primary care physician. The patient had a personal history of type 2 diabetes mellitus and was a smoker. Physical examination revealed an erythematous subcutaneous lesion with areas of suppuration in the lower left groin. On palpation, the lesion extended to the perineal region (Fig. 1). The examination also revealed 3 lesions that were clinically compatible with condylomata acuminata in the perianal area. He was diagnosed clinically for hidradenitis suppurativa (HS). Ultrasonography with an 18-MHz linear probe revealed the presence of a fistulous tract, with areas of fluid collection, together with intense vascularization on Doppler mode (Fig. 2). Given the location and depth of the lesion, the patient was sent to the clinic of the surgery department for complete removal of the lesion. Histopathology showed changes that were compatible with HS and well-differentiated squamous cell carcinoma (SCC) over the fistulous tract, which was located close to the deep resection margin. No vascular invasion or perineural infiltration was observed (Fig. 3 A and B). Extension of the surgical margins revealed no evidence of residual tumor, although foci suggestive of viral infection were observed (Fig. 3 C). Immunohistochemical staining with p16 was intensely positive (Fig. 3 D). Polymerase chain reaction and genotyping based on in situ hybridization with a microarray of paraffin-embedded tissue were performed to investigate the presence of human papillomavirus (HPV). The result was positive for genotype 6 (considered low risk); the same finding was recorded for the perianal condylomata acuminata. Laboratory analysis yielded negative results.

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\(^\text{\#}\) Please cite this article as: Segura Palacios JM, García Montero P, Fúnez Liébana R, Repiso Jiménez JB. Human Papilloma Virus and the Risk of Squamous Cell Carcinoma Arising in Hidradenitis Suppurativa. Actas Dermosifiliogr. 2018;109:457–459.