
Vellus hair cysts presenting as an atypical acneiform eruption

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Abstract: A 32-year-old male patient presented for 8 months an asymptomatic therapy-resistant acneiform eruption on his back and buttocks. Skin examination showed several inflammatory papules, which evolved to hyperpigmentation. At the same distribution non inflammatory papules, which resembled rice grains, were also observed. Light microscopy showed small keratin-filled cysts, with an epithelial multilayered wall, without granular layer. Keratin and some vellus hairs were identified inside the cyst, confirming the diagnosis of vellus hair cysts. Diagnosis of vellus hair cysts should be suspected in cases of multiple papules or therapy-resistant cases of acneiform eruptions.

Keywords: Acne vulgaris; Acneiform eruptions; Follicular cyst; Isotretinoin; Young adult; Keratins

INTRODUCTION

Vellus hair cysts is a rare condition which has no racial or gender predisposition. It normally affects adolescents or young adults and could be sporadic or autosomal dominant. 1,2,3 This term was used for the first time by Esterly and cols in 1977, who described two pediatric patients with multiple monomorphic asymptomatic hyperchromic papules on the trunk and extremities. 2,4

CASE REPORT

A 32-year-old male patient presented an asymptomatic acneiform eruption on his back and buttocks for 8 months. He was treated with benzoyl peroxide, topical antibiotics and oral isotretinoin with no response. Skin examination showed inflammatory papules, which resolved with hyperpigmentation (Figure 1). At the same distribution non inflammatory papules, which resembled rice grains, were also observed. One of these papules was removed and light microscopy showed a small keratin-filled cyst, with an epithelial multilayered wall, without granular layer (Figures 2A and 2B). Some vellus hairs were identified inside the cyst, confirming the diagnosis of vellus hair cyst (Figure 2C).

DISCUSSION

The pathogenesis of vellus hair cysts is not fully understood, probably development abnormalities of this peculiar small follicle, which is typically found outside from scalp and face, lead to occlusion and dilatation of the follicular unit. Moreover mutations in the gene that codifies keratin 17 were described.5
This condition is characterized by multiple normochromic or hyperpigmented papules, ranging from 1 to 5 mm, that appear mainly on the chest and extremities; rarely abdomen, neck, axillae and face may be involved.  6,7 They may undergo inflammatory changes and be misdiagnosed as acne, as in the case here described.

Diagnosis should be suspected in cases of multiple papules or therapy-resistant cases of acne. It can be confirmed by light microscopy, which shows the dermal keratinized cysts with hairs in their interior.  

Differential diagnoses include steatocystoma multiplex, milium, moluscum contagiosum, other cysts (infundibular, trichilemmal, epidermal), folliculitis and acne.  4,8,9 Spontaneous remission occurs in 25% of the cases, due to transepidermal elimination or inflammatory destruction. Treatment is a challenge, with no response to oral isotretinoin.  3,8 There are some reports of improvement with dermabrasion, CO2 Laser and needle incision, but the large number of lesions may preclude complete resolution.  3,10 There is only one report of non-inflammatory vellus hair cysts in the Brazilian literature, different from our case, which had an inflammatory onset.  11

REFERENCES


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