Introduction

Nevus comedonicus is a rare hamartoma of the pilosebaceous unit, first described by Kofmann in 1895 [1]. In most cases it is present at birth but may also develop later, usually before the age of 10 years, without racial or sexual preponderance. When appearing in adults, although rare, it is frequently associated with irritation or trauma [2].

The lesions typically present a linear or zosteriform distribution, but extensive areas have already been described, including variant with bilateral involvement. The most frequently affected sites are the face, trunk, neck, and upper limbs. It may affect areas without hair follicles, such as the palms, soles, and the glans penis. Scalp involvement is rare [3]. There may be associations with other skin lesions or abnormalities of the central nervous, musculoskeletal, and ocular system [4].

We present a case of nevus comedonicus in a woman suffering from obesity and acne vulgaris.

Case Presentation

A 40-year-old woman with a history of obesity and acne presented with an asymptomatic lesion that had appeared 20 years previously on the right thigh. The clinical aspect and extent of the lesion caused relevant psychosocial repercussions, interfering with her personal relationships. At dermatological examination, bundles of dilated hair follicles were observed, filled with dark-colored plugs, presenting a linear distribution in the right thigh with alteration of the skin pigmentation at the site of the lesion; at dermoscopy, well-defined homogeneous areas in dark brown shades were noted (Figure 1, A and B). Some cysts had been treated in the past with antibiotics, leaving some hyperchromic signs on the thigh. The patient also suffered from facial acne (Figure 2) that was resistant to numerous treatments performed in previous years: topical and systemic antibiotics and topical and systemic retinoids. The histological examination performed with a 5-mm diameter punch showed a large group of dilated follicular ostia devoid of hair shafts but filled with keratin layers (Figure 3).

On the basis of the history and typical features of the lesion and histology, a nevus comedonicus in a patient with resistant acne and obesity was diagnosed. Consultations with other specialists excluded the presence of extracutaneous signs or association with genetic syndromes and other diseases or nevus comedonicus syndrome. The patient refused genetic analysis. Because of the extent of the lesion and the possibility of its turning into an unaesthetic scar, we treated...
In each case of nevus comedonicus, it is obligatory to rule out the comedonicus syndrome, which may include ocular lesions (cataract, corneal erosion), skeletal abnormalities (syndactyly, clinodactyly, the absence of hand bones on x-ray, scoliosis, vertebral defects), and neurological disturbances (microcephaly, mental deficiency, the dysgenesis of the corpus callosum) [7].

In some cases of nevus comedonicus, a somatic mutation in fibroblast growth factor-receptor gene 2 (FGFR2) has been identified, namely the Ser252Trp missense mutation. FGFR2 is expressed in keratinocytes, hair follicles, and sebaceous glands and has been implicated in induction of hypercornification and comedogenesis. Germline FGFR2 mutations are also associated with acne, as seen in dominant Apert syndrome. When occurring as a mosaic, the acne lesions follow the lines of Blaschko, the pattern of embryological cell development and proliferation [8]. In a recent case, a postzygotic mosaicism was found in exon 4 of FGFR2 (c.758C>G, p.Pro253Arg) in low copy number. A similar mosaicism was also found in 2 other patients in p.Ser252Trp of exon 4 [9].

FGF signaling pathways lead to the interplay of the mTORC1 and FoxO1 pathways; the same pathway is regulated by increased caloric intake, leading to the onset of acne and metabolic syndrome [10]. The 2 pathways could be strictly dependent, explaining our clinical case.

Discussion

The prevalence of nevus comedonicus has been estimated from 1 in 45,000 to 1 in 100,000. Nevus comedonicus lesions might present with various patterns of distribution: unilateral, bilateral, linear, interrupted, segmental, or blaschkoid [5]. The major histopathological features are large-grouped, dilated follicular ostia devoid of hair shafts but filled with keratin layers. At some locations in the bases of the follicular invaginations one may observe singular rudimentary glands, which are not, however, obligatory as they may be absent. Small cysts, cystic invaginations, and occasionally large cysts may be seen in histopathological investigation; the variable cystic structures are lined by keratinizing squamous epithelium. Hyperkeratosis and acanthosis of the epidermis may be present but not para- or dyskeratosis [6].

Nevus comedonicus syndrome (ORPHA:64754) belongs to the group of epidermal nevus syndromes including Schimelpenning syndrome, phacomatosis pigmentokeratotica, angora hair nevus syndrome, and Becker nevus syndrome among others in which a genetic basis has not yet been identified [7].

Figure 1. (A) Clinical presentation: plaque-like lesion with wide-open follicles with blaschkoid distribution and alteration of the skin pigmentation. (B) Dermoscopy (polarized ×10): well-defined homogeneous area in dark brown shade in the area of whitish skin of the right thigh.

Figure 2. Papulopustular acne lesions of the face and erythematous acne scars due to past treatments.

Figure 3. Histology presentation: dilated follicular ostia devoid of hair filled with keratin layers.

the patient with 33% salicylic acid peeling and home-based topical retinoid therapy, obtaining only partial benefit.
Conclusions

We suggest that the coexistence of comedonic nevus, acne, and obesity may not be a random association. Future studies should investigate a screening of phenotypically similar patients for FGFR2 mutations and a possible therapeutic modulation of FGFR signaling.

References