Acrokeratoelastoidosis as an example of marginal papular acrokeratoderma with prominent elastorrhexis

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Acrokeratoelastoidosis (AKE) is a rare condition described by Oswaldo Costa in 1953 [1]. Clinically, it is characterized by the presence of hyperkeratotic or umbilicated papules located on the margins of palms and/or soles. The first lesions usually develop in childhood or early adulthood and gradually increase in size and number, become grouped or form plaques [1–3]. The prominent histological feature of the condition is the diminished density and fragmentation (elastorrhexis) of elastic fibres, which can be detected using Verhoeff-van Gieson or orcein staining [1–3].

A 37-year-old Caucasian woman was admitted to the Department of Dermatology for evaluation of asymptomatic papular lesions located on the margins of hands and feet. The first papules had developed when the patient was 20 years old and had gradually increased in number. The lesions had been initially diagnosed as viral warts and had been treated with keratolytics and several sessions of cryosurgery with no improvement. Family history for similar skin changes was unremarkable. Several years prior to hospitalization, the patient had been diagnosed with ectopic gastric mucosa in the oesophagus, but was otherwise healthy. On physical examination, flesh-coloured papules of 2–4 mm in diameter were present along the margins of both hands accompanied by slight hyperkeratosis of the palms (Figures 1 A, B). Additionally, several hyperkeratotic papules were present at the line of transgression between the dorsal and plantar surface of feet. Routine laboratory work-up was within normal limits. A 5-mm punch biopsy was performed from one of the papules, which revealed prominent irregular acanthosis, papillomatosis, hypergranulosis, focal hyperkeratosis and scarce perivascular inflammatory infiltration in the dermis with slight epidermotropism (Figure 2 A). Verhoeff-van Gieson staining revealed reduced density of elastic fibres with elastorrhexis (Figure 2 B). On the basis of clinical presentation and histological image the diagnosis of AKE was made. The patient was informed about the benign nature of the condition and she refused to try other treatment options.

AKE is considered to be a very rare disease, but its actual incidence may be underestimated. Over 50 cases have been reported so far. The number of articles has risen significantly in the last decade, with 18 cases reported in 2008–2018. Interestingly, recent reports include predominantly females (a female to male ratio of 7 : 2) with late-onset AKE (mainly at 20–30 years of age). The time between the occurrence of first papules and the diagnosis is usually several years, but may be extended up to 20 years. Unilateral lesions have been reported mainly in
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Children and have been present since birth. Both sporadic and familial cases of autosomal dominant inheritance have been reported in the literature [4–9]. A potential linkage to chromosome 2 was suggested by Greiner et al. on the basis of genetic mapping of 21 family members affected by AKE [10]. No particular race predominance has been found so far. The electron microscope (EM) has recently proved to be a valuable tool in making the diagnosis [5]. Fragmented elastic fibres resembling rooster crests can be observed on scanning EM, while transmission EM shows decreased thickness and fragmentation of the elastic fibres, surface indentations, irregular black spots in the fibre core and normal collagen bundles. In single case reports, AKE was associated with nail dystrophy [11] or aquagenic palmoplantar keratoderma [9].

The pathogenesis of the disease remains unknown. One of the most common hypotheses implies a reduced number and impaired function of fibroblasts in the dermis, which secrete abnormal elastic material [12]. An interesting finding was reported by Fiallo et al. [3], who detected fragmentation of the elastic fibres not only in the lesional skin, but also in normal-appearing, non-sun-exposed skin, and suggested that AKE may be in fact a primary elastic tissue disease. A confirmation of this hypothesis might be delayed wound healing, which Fiallo et al. observed in one of their patients with AKE [3]. A potential association between late-onset AKE and immunosuppression was suggested by Hussain et al. [6], who reported a patient with AKE developing during therapy with vedolizumab for Crohn’s disease.

The nosological position of AKE is not well established, either. The condition was included by Rongioletti et al. [13] in a group of marginal papular acrokeratodermas, which consisted of two acquired entities (degenerative collagenous plaques of the hands and digital papular calcinosis) and five hereditary conditions (AKE, focal acral hyperkeratosis, palmoplantar keratoderma of the punctate type, hereditary papulotranslucent keratoderma and mosaic acral keratosis). All these entities as well as acrokeratosis verruciformis of Hopf and viral warts, should be taken into consideration in differential diagnosis. In case of doubt, histological examination should be performed. The presence of cell vacuolization should raise the suspicion of viral warts, dyskeratosis with “churh spire” configuration points to acrokeratosis verruciformis of Hopf, while elastorrhexis is the distinctive histological feature of AKE. Focal acral hyperkeratosis is considered to have the same clinical presentation as AKE, but lacks signs of elastorrhexis on histological examination [14]. Degenerative collagenous plaques of the hands typically develop in middle-aged or older individuals with a history of chronic sun exposure. They are present predominantly on the sides of hands and/or fingers, rarely feet, and are characterized by the presence of degenerative collagen and elastin fibers with extensive actinic damage on histology. Palmoplantar keratoderma of the punctate type presents with dome-shaped papules on the palms and soles, which display hyperkeratosis, parakeratosis, vacuolated epidermis, basal layer spongiosis and dilated sweat ducts on histological examination [2]. On the other hand, Abulafia and Vignale [15] suggested that AKE, focal acral hyperkeratosis and other entities classified by Rongioletti et al. [13] as hereditary marginal papular acrokeratodermas are in fact a result of a variable genetic expression of the same condition and should be considered as clinicopathological variants rather than separate entities.

The prognosis in AKE is very good and for the majority of patients the lesions are only a cosmetic problem. No cases associated with malignancies have been published so far. A few authors reported a potential association of AKE with hyperhidrosis. Nevertheless, in most of the cases,
the affected patients, as well as the present patient, did not complain of hyperhidrosis of the palms or soles.

The therapy of AKE remains challenging. Cryosurgery or standard keratolytics are usually ineffective. Other attempts to use systemic steroids, methotrexate, dapsone or antibiotics were not successful either [16]. Due to the rarity of the condition, treatment recommendations are mainly based on single cases. Several reports on the efficacy of acitretin or other systemic retinoids in marginal papular acrokeratoderma have been published recently [4, 17, 18]. However, the potential teratogenicity of this therapy should always be taken into consideration in young patients with AKE.

Conflict of interest

The authors declare no conflict of interest.

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