PL can occur at any age, but it is particularly common in children, with incidence peaking at 2, 5, 10, and 12 years of age. It is very rare in the first year of life. In our review of the literature, we found only 9 cases of onset before the age of 12 months. These included 1 congenital case, 3 cases with onset at 8 months of age, 5-7 and 1 case with onset at 11 months of age. 8 We also found mention of an additional 4 cases in children aged under 1 year in a retrospective study of 124 children with PL. 9 In the 9 cases described in the literature, all the patients (8 boys and 1 girl) were otherwise healthy. There was just 1 case with a history of previous infection, in which pneumococci and Haemophilus influenzae were isolated. 3 Six of the patients had lesions clinically consistent with PLEVA, with ulceronecrotic lesions that resolved with topical corticosteroid therapy in a maximum of 2 months. 5-7, 8 The remaining 3 patients had lesions that were clinically and histologically consistent with PL chronica. 4-6 The disease followed an indolent course in 2 patients, 1 of whom developed an indurated plaque in the right iliac fossa after a year and a half. The plaque was initially compatible with lymphomatoid papulosus but 2 years later, it was histologically diagnosed as cutaneous T-cell lymphoma. 7 The second patient developed parapsoriasis at 10 years of follow-up. 5

Given the scarcity of cases in the literature, no clear conclusions can be drawn, but based on our review, PL would appear to be very rare in children under 12 months of age, with just 10 cases, including ours, reported to date. Within this age group, PL is more common in boys and tends to occur in an acute form and to resolve spontaneously without complications. Nevertheless, given the possible association with lymphoproliferative disorders, long-term monitoring is recommended.

References


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Periungual Acral Fibrokeratoma: Surgical Excision Using a Banner Flap

Fibroqueratoma acral periungual: exéresis quirúrgica en bandera

To the Editor

The surgical treatment of periungual tumors is complicated by the risk of causing irreversible nail dystrophy. However, incomplete resection leads to a high risk of recurrence, and surgery must therefore eliminate the whole tumor while preserving the germinal cells of the nail matrix.

We present the case of a 31-year-old man who was seen in outpatients for a firm, pink, digitiform tumor of about 5 × 3 mm that had been present on the fifth toe of the left foot for about 5 months (Fig. 1). The base of the lesion was beneath the eponychium but above the nail plate. The patient did not recall any history of trauma, and there was no family history or presence of other lesions suggestive of tuberous sclerosis. The clinical diagnosis was periungual fibroma and, as the patient reported discomfort due to friction against his footwear, it was decided to excise the lesion as an elective procedure.

The operation was performed under local anesthesia with 2% mepivacaine, and a glove was tied around the proximal part of the digit as a tourniquet to provide a bloodless
subungual and periungual fibromas characteristic of tuberous sclerosis. These tumors should be excised when they cause symptoms or significant cosmetic problems.

Various surgical techniques can be used to resect these tumors, depending on whether they arise from above or below the nail plate. Lesions beneath the nail plate require partial or total removal of the nail. Lesions above the nail plate, on the other hand, can be removed by shaving followed by phenolization, carbon-dioxide laser vaporization, or, more usually, surgical resection after lifting the proximal nail fold as a banner flap.

Surgical resection is usually performed under nerve block anesthesia. Two oblique incisions are made in the proximal nail fold, one at each end of the eponychium; the whole proximal nail fold can then be lifted to expose the lesion. This allows the base of the lesion to be evaluated and complete excision performed, preventing future recurrence, without damaging the nail matrix. We used a simpler variant of this technique in which the 2 incisions were made along the borders of the tumor instead of making them at the medial and lateral ends of the eponychium.

It is important to note that this type of lesion has a tendency to bleed despite suturing, and correct hemostasis is therefore essential. The use of vasoconstrictors is not recommended in acral regions due to the risk of distal ischemia; a simple digital tourniquet will adequately control bleeding. In addition, Monsel solution—a ferric sulphate solution that has a high local hemostatic potential—can be used to ensure correct hemostasis. Monsel solution is applied directly with a cotton bud or gauze swab after suturing.

We thus propose this technique as a useful alternative to wedge or punch excision. These other procedures usually provide poorer visualization of the lesion, with a higher risk of recurrence due to incomplete excision of the lesion or nail dystrophy if the resection is too wide. In addition, the technique described preserves the suprallesional skin of the proximal nail fold, which reduces tension in the scar and achieves excellent cosmetic results. The banner flap is therefore a simple and very practical technique for small periungual lesions.

References

Fox-Fordyce Disease With an Atypical Clinical Presentation

Enfermedad de Fox-Fordyce con presentación clínica atípica

To the Editor:

Fox-Fordyce disease (FFD), or apocrine miliaria, is a rare inflammatory disease of the apocrine glands characterized by the presence of pruritic follicular papules in apocrine-rich areas of the skin. We present a case of histologically confirmed FFD with extensive involvement of 2 unusual sites, the chest and abdomen.

A 39-year-old woman was referred for evaluation of an asymptomatic rash that had appeared 2 years earlier. There was no significant personal or family medical history. The rash had first appeared in the presternal area and had then spread over the abdomen to the pubic region. The rash consisted of multiple minute nonconfluent red-to-yellow papules on the breasts and especially the lower chest and abdomen, as far as the pubic region (Fig. 1). The axillae, areolae, and genitals were uninvolved.

Histologic examination of a papule from the lower presternal region revealed dilatation and hyperkeratotic plugging of the follicular infundibulum, which was surrounded by abundant foamy histiocytes (xanthomized macrophages) (Fig. 2); a mild-to-moderate periadnexal chronic lymphocytic inflammatory response; and clusters of apocrine cells, with a tendency to cystic dilatation, between the reticular dermis and hypodermis (Fig. 3).

There were no relevant findings in the blood panel and the basic hormone panel, which included testosterone, dehydroepiandrosterone sulfate, progesterone, estradiol, luteinizing hormone, and follicle-stimulating hormone.

No improvement was noted after successive attempts at topical treatment with tretinoin, corticosteroids, and clindamycin.

FFD was first described by Fox and Fordyce in 1902. This rare skin condition is most common in women aged 13 to 35 years. The etiology and pathogenesis of FFD are unknown.

Figure 1 A, Multiple minute nonconfluent red-to-yellow papules on the chest and abdomen. B, The axillae were not involved.

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