Preputial Ectopic Sebaceous Glands Mimicking Molluscum Contagiosum

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The authors describe a 12-year-old boy with a peculiar presentation of preputial papular lesions similar to molluscum contagiosum. Histopathologic investigation revealed the presence of a sebaceous gland opening directly onto the surface and hyperplasia of the epithelium. Key words: Prepuce; Puberty.

(Accepted December 30, 1989.)

Acta Derm Venereol (Stockh) 1990; 70: 344–348.

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The presence of ectopic sebaceous glands on the mucocutaneous surfaces of the oral cavity and of male and female genital organs is well known (1, 2). Clinically, such glands appear as small roundish, yellowish papules about 2–3 mm in size, easily detected because they are superficially located and coated by a thin epidermal layer (3).

We report here the case of a young male with ectopic sebaceous glands of the preputium, characterized by unusual clinical features.

CLINICAL CASE

A 12-year-old boy was brought to our notice because of asymptomatic preputial papular lesions present for about 2 months. The boy was in good general health. Psychosexual development was normal for his age.

The dermatological examination revealed many papular lesions of the mucosal aspect of the prepuce, some discrete and some arranged in small groups. Such lesions, of the same colour as the surrounding skin, were hemispherical, smooth and shiny, about 2–6 mm in diameter, hard and elastic at compression, with central umbilication (Fig. 1). Upon squeezing, no discharge from the umbilication could be seen.

Microscopic examination of one element showed a protrusion of the chorion coated by hyperplastic epithelium, with a sebaceous gland opening directly onto the surface, corresponding to an invagination of the epidermis (Fig. 2).

Laboratory findings were within normal limits. Plasma levels of hypophyseal gonadotropins (LH and FSH) and of testosterone were also normal. The urinary 17-ketosteroids were at the upper limits of normal for the patient’s age (10.6 mg/24 h; n.v. 0.7–10.0 mg/24 h).

The patient was checked again after about one year, the preputial lesions being unchanged with regard to both number and size.

DISCUSSION

In the case reported, the morphology of the preputial lesions was not immediately suggestive of the clinical picture of ectopic sebaceous glands as described in the literature (3). Rather, their size and the presence of a central umbilication suggested a clinical diagnosis of molluscum contagiosum. Histological examination showed a small sebaceous gland in the superficial dermis, not connected with a hair follicle and opening directly onto the skin surface, as an ectopic sebaceous gland.

According to the literature, visible ectopic sebaceous glands seem to be present on the shaft of the penis in about one-third of adult males (4). Clinically they appear as papules the size of a pinhead, being only rarely more prominent. If this is the case, histologically a marked hypertrophy of the sebaceous glands, a broken glandular duct or a cystic formations enclosing hair fragments is usually observed (4). In the case described by us, the clinical lesions seem to be histologically related to epidermal hyperplasia rather than to an increase in size of the sebaceous glands, which appear in the centre of the sections.

In the past literature there are confusing data about the presence of sebaceous glands on the prepuce and the corona (5). They were actually based on the incorrect definition of glands Tyson had given to the papillae of the corona glandis. It is better to withdraw the term of ‘Tyson’s glands’ (4) and to define as Papillomatosis corona penis the papular structures of the corona and as ectopic sebaceous glands those tiny papules microscopically related to free sebaceous glands of the prepuce or (rarely) of the glans.

In conclusion, on the basis of the site and of some histological features of the lesions we have described, we believe we are dealing with an unusual clinical presentation of ectopic sebaceous glands of the prepuce.

REFERENCES

Urticarial Vasculitis Occurring in Association with Visceral Malignancy

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This is a report of a solitary patient who had urticarial vasculitis and an adenocarcinoma of the colon. Urticarial vasculitis has not been described in association with malignancy. It is considered that tumor-associated immune complexes might have been involved in the pathogenesis of the vasculitis. Key words: Circulating immune complexes; Leukocytoclastic vasculitis; Paraneoplastic.

(Accepted December 11, 1989.)

Acta Derm Venereol (Stockh) 1990; 70: 345–347.

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The syndrome of urticarial vasculitis has found its niche in the spectrum of cutaneous vasculitis as attested by reports and reviews which have been numerous in the recent literature (1–5). A case is reported herein of urticarial vasculitis occurring in a patient with a visceral malignancy. The vasculitis was kept well under control with a low-dose regimen of indomethacin. The presence of circulating immune complexes was documented.

CASE REPORT

A 69-year-old white woman presented May 16, 1984 with a