Detection of Human Papillomavirus Type 16 in Bowen's Disease of the Web-space of the Foot

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Accepted July 19, 2004.

Sir,
Recent studies have identified various types of human papillomavirus (HPV) in Bowen’s disease (BD) on the hands and feet, especially at periungual and subungual sites (1, 2). BD represents a squamous cell carcinoma in situ. HPV 16 and other ‘high risk’ types have frequently been identified as in genital BD. Only two cases of BD on the web-spaces of the foot have been reported (3, 4). We describe a case of BD arising in the web-space of the foot associated with HPV 16.

CASE REPORT
A 33-year-old Japanese man had observed a brownish macule on his left foot for 1 year. He had previously been treated for tinea pedis with topical antifungal agents. Physical examination revealed a brownish, relatively well demarcated macule, 7×3 mm in diameter, in the web-space between the third and fourth toes of his left foot (Fig. 1a). A skin-coloured keratotic papule was observed adjacent to the brownish macule. He had no history of genital disorders such as Bowenoid papulosis or condyloma acuminatum. Histopathological examination of a skin biopsy revealed proliferation of irregularly arranged atypical keratinocytes throughout the epidermis, hyperkeratosis with dyskeratotic, clumping cells and mitotic figures (Fig. 1b). A diagnosis of BD was made. Fontana-Masson staining showed an increased amount of melanin in the basal layer and numerous pigment-filled dendritic melanocytes in the lesion. Melanophages were also prominent in the upper dermis. The HPV capsid antigen was detected by immunohistochemical analysis (K1H8, Dako, Kyoto, Japan). Southern blot hybridization was employed to detect HPV DNA and determine the HPV genotype. The cleavage patterns of the lesion were similar to those of prototype HPV 16 (Fig. 1c). The lesion was surgically removed and no relapse was observed after 1 year.

DISCUSSION
Our case is clinically characterized by a very small, brownish and keratotic macule on the web-space of the patient’s left foot. Histopathological findings were compatible with those of BD. Pigmentation was caused by the presence of pigment-filled melanocytes, a large amount of melanin in the basal layer and prominent melanophages in the upper dermis. HPV 16 DNA was identified in the lesion.

Over 90 HPV genotypes have been characterized. HPV 16 was first found in extragenital BD in 1983 (5), and subsequently also in BD of the hands and feet (2).
HPV 16 is the type most often found in cervical intraepithelial neoplasia grade III (CIN III) or invasive cervical carcinoma (6). Since BD histopathologically corresponds to CIN III, it is assumed that specific genital HPVs may induce the same histological lesions both on the skin and in the genital region. In some reported cases, the same HPV type has been found in BD both on the skin and in the genital lesions of the same patient (7, 8). Thus, it is assumed that the viral transmission occurs via a genital-digital route in the development of extragenital BD (8). Our patient was not aware of genital lesions such as Bowenoid papulosis or condyloma acuminatum, but had a history of tinea pedis. The scratching may result in autoinoculation of HPV into the skin. His lesion may thus be related to HPV inoculation or environmental trauma.

The occurrence of BD in the web-spaces of the feet has been rarely reported (3, 4). Burns (3) described a patient with multiple pigmented, warty macules in the web-spaces of both feet; however, a virological investigation was not conducted. Stones et al. (4) described a patient with hyperkeratotic plaques on the dorsal surface and interdigital spaces over the second, third and fourth toes of both feet and detected HPV 16 DNA in the tissue using Southern blot hybridization. These three cases including our own had similar clinical features with pigmented, hyperkeratotic macules. However, the previously described lesions have been multiple, well circumscribed and verrucous plaques (3, 4), a type also described by Kettler et al. (9).

REFERENCES


DOI: 10.1080/00015550410023491