Squamous Cell Carcinoma Developing within Lesions of Disseminated Superficial Actinic Porokeratosis

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Disseminated superficial actinic porokeratosis (DSAP) consists of multiple annular, hyperkeratotic lesions that have a bilateral distribution on sun-exposed areas, particularly the extremities. DSAPs have a wider distribution than porokeratosis of Mibelli and usually develop during the 3rd or 4th decade of life. Squamous cell carcinoma that arises in the classical type of porokeratosis of Mibelli is well-documented, but there are only a few reports of squamous cell carcinoma in DSAP. Here, we describe a 62-year-old man with DSAP who developed squamous cell carcinoma on his right forearm. (Ann Dermatol 23(4) 536-538, 2011)

Keywords- Disseminated superficial actinic porokeratosis, Squamous cell carcinoma

INTRODUCTION

Disseminated superficial actinic porokeratosis (DSAP) is the most common form of porokeratosis, and is inherited as an autosomal dominant condition, with reduced penetrance at a younger age. The characteristic lesions of DSAP represent multiple annular, hyperkeratotic, brownish macules measuring 2~5 mm in diameter. The center of a macule is minimally atrophic or depressed and the border spreads centrifugally in raised ridges. The distribution is symmetric and usually affects sun-exposed areas.

Development of squamous cell carcinoma within the classic type of porokeratosis of Mibelli is well-documented, but there are only a few reports of squamous cell carcinoma in DSAP. We describe a case of squamous cell carcinoma developing within lesions of DSAP.

CASE REPORT

A 62-year-old male presented with pruritic eruptions on sun-exposed portions of both forearms that had gradually increased in number over a period of 5 years (Fig. 1). The lesions were exacerbated during the summer months. Along with these lesions, an erythematous, irregular, marginated, scaly, crusted plaque had developed on the right forearm 3 years earlier. There was no significant medical or family history.

On physical examination, a 2×3 cm erythematous, irregular, marginated, scaly, crusted plaque was noted on the right forearm. The lesions were exacerbated during the summer months. Along with these lesions, an erythematous, irregular, marginated, scaly, crusted plaque had developed on the right forearm 3 years earlier. There was no significant medical or family history.

On physical examination, a 2×3 cm erythematous, irregular, marginated, scaly, crusted plaque was noted on the right forearm. In addition, the patient had numerous annular, brown, atrophic, and symmetric macules surrounded by well-demarcated, raised ridges on extensor aspects of both forearms, which are characteristics of DSAP. Complete blood count, as well as liver and kidney function tests were all within normal limits.

A skin biopsy specimen of the multiple, brown, annular lesions showed histologic changes of typical DSAP. There was a cornoid lamella composed of a column of parakeratosis with underlying hypogranulosis and perivascular lymphocytic infiltrations in the dermis localized beneath the cornoid lamella. A skin biopsy obtained from the erythematous plaque on the right forearm showed dysregulated keratinocytes with hyperchromatic, atypical nuclei, consistent with squamous cell carcinoma. A cornoid lamella was observed in the lesion of the
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Fig. 1. An irregular, margined, erythematous plaque and multiple, brown, atrophic macules surrounded by well-demarcated, raised ridges on the right forearm.

Fig. 2. Biopsy specimen obtained from the erythematous plaque on the right arm. In epidermis, acanthosis and dysregulated keratinocytes with hyperchromatic, atypical nuclei are observed. A cornoid lamella composed of a column of parakeratosis is seen in the lesion of the squamous cell carcinoma (H&E, ×100).

Fig. 3. Overexpression of p53 in the epidermis of a disseminated superficial actinic porokeratosis lesion. A column of parakeratosis with underlying hypogranulosis is observed. Perivascular lymphocytic infiltrations are localized beneath the cornoid lamella (p53 immunohistochemical stain, ×200).

DISCUSSION

Porokeratosis has several clinical varieties including porokeratosis of Mibelli, giant porokeratosis, DSAP, porokeratosis palmaris et plantaris, punctuate porokeratosis, and linear porokeratosis. The most common form of porokeratosis is DSAP, which was first described as a clinical entity by Chernosky in 1966. The distribution of typical lesions is symmetric and usually affects sun-exposed areas. The lesions generally spare the face, palms, soles, and mucosal surfaces. DSAP can affect people of all ages, but manifests during the 3rd or 4th decade of life.

The pathogenesis of DSAP is not clearly understood, but frequent p53 overexpression in the epidermis of porokeratotic lesions have been detected. Overexpression of p53 can be induced by p53 gene mutations and other DNA damaging agents, such as ultraviolet (UV) light and ionizing radiation. In a previous study, mutations of the p53 gene were not detected in porokeratotic lesions. This finding suggests that other causative mechanisms exist for overexpression of p53 in the epidermis of porokeratotic lesions. The patient in this case showed overexpression of p53 in the epidermis of DSAP lesions (Fig. 3).

The occurrence of malignancies in porokeratotic lesions is clinical evidence of the pre-cancerous nature of this disease. Malignancies have been reported for porokeratosis of Mibelli, linear porokeratosis, porokeratosis palmaris et plantaris and DSAP. Associated malignancies are squamous cell carcinoma, Bowen’s disease and basal cell carcinoma. Squamous cell carcinoma arising in the classic type of porokeratosis of Mibelli is well-documented, but there are only a few reports of squamous cell carcinoma in DSAP. All of the reported squamous cell carcinoma cases arising from DSAP lesions have originated in the distal extremities. These findings indicate that there is a significant role of UV light on the evolution of squamous cell carcinoma from DSAP. In addition, results from previous reports show that p53 gene mutations are
Porokeratosis does not usually need treatment, but in some cases, treatment is necessary due to potential for progression to a malignancy and for cosmetic purposes. In contrast, treatment of DSAP lesions is unsatisfactory. As in this case, the potential for progression to cancer induced by UV light exists, thus sun avoidance must be emphasized when treating DSAP.

The prognosis of squamous cell carcinoma in DSAP lesions has not been reported yet. However, the reported cases have had no recurrences after excision of the cancerous lesions.

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