Bell’s palsy in a case of Darier’s disease - a rare disease association or coincidental finding?

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ABSTRACT
Darier’s disease (DD) is a rare acantholytic dyskeratotic autosomal dominant genodermatosis characterized by the presence of warty, brown papules and plaques affecting the seborrhoeic areas. Frequent bacterial, fungal and viral particularly herpes simplex virus (HSV) infections complicate DD. Bell’s palsy is an acute onset, idiopathic facial paralysis resulting from a dysfunction anywhere along the peripheral part of the facial nerve. Reactivation of HSV is considered to be the main cause of Bell’s palsy. This case represents, to the best of our knowledge, the first case of DD presenting with Bell’s palsy. This case underlines the importance of recognizing HSV infection in DD.

Key words: Darier disease; Bell’s palsy; Skin disease

INTRODUCTION
Darier’s disease (DD) is an autosomal dominant condition characterized by a persistent eruption of hyperkeratotic papules, histological examination of which shows suprabasal acantholysis with dyskeratosis. Patients with DD have an increased susceptibility to herpes simplex infection. A number of clinical studies have described co-occurrence of various neurological and psychiatric symptoms with DD, but occurrence of Bell’s palsy with DD has never been reported [1]. We report here a case of DD with Bell’s palsy.

CASE REPORT
A 20 year old man presented with a 3-year history of itchy greasy yellow brown papules and plaques over face, neck, shoulders, upper back, axillae and groins (Fig. 1). On examination patient also had minute Palmar pits (Fig. 1) and V- shaped nicking of the free edge of the nails, characteristic of DD (Fig. 2). A skin biopsy was done from back to support clinical diagnosis, came consistent with diagnosis of DD (Fig. 3). Five days later, the patient presented with pain and weakness over right side of the face, was diagnosed as Bell’s palsy (Fig. 4) and treated with oral steroids and valacyclovir. Serological test for both HSV-1 and HSV-2 were non-reactive. A week later, there was improvement in both facial palsy and skin lesions (Fig. 5).

Prior to the study, patient gave written consent to the examination and biopsy after having been informed about the procedure.

DISCUSSION
Darier’s disease was described independently by White and Darier in 1889. It is an autosomal disorder with variable penetrance, related to mutations in ATP2A2 gene at chromosome 12q24.1, which encodes the sarco-endoplasmic reticulum calcium ATPase type 2 (SERCA2). This defect results in impaired intercellular adhesions [2,3].

It has world-wide distribution, with the prevalence estimated to vary from one in 36,000 to one in 100,000 and an incidence of new cases of four per million per 10 years [2,4].
The diagnosis can be easily established in patients with unexplained unilateral isolated facial weakness. The onset is sudden and symptoms typically peak within a few days. Additional symptoms may include pain in or behind the ear, numbness or tingling in the affected side of the face usually without any neural deficit, hyperacusis and disturbed taste on the ipsilateral anterior part of the tongue [7].

Autoimmune process, viral infections, and even ischemia are the cause of initiation of inflammation. Different viruses from herpes virus family, herpes simplex virus -1 (HSV-1), HSV-2, human herpes virus -6 (HHV-6), and varicella zoster virus (VZV) have been considered to play role in bell’s palsy. HSV has been considered particularly as the etiological agent in recent studies. Despite studies in favour of seropositivity of HSV in Bell’s palsy, most studies could not find any definite association between antibody titres and Bell’s palsy [8,9].
The main aim of treatment in the acute phase of Bell’s palsy is to speed up the recovery and to prevent corneal complications. Strategies to speed recovery include physical therapy, corticosteroids and antiviral agents. Inflammation and edema of the facial nerve are implicated in causing Bell’s palsy. The rationale for the use of corticosteroids in acute phase of Bell’s palsy is that they have a potent anti-inflammatory action which should minimize nerve damage and thereby improve the outcome. The rationale for the use of antiviral agents is the evidence that the inflammation of the facial nerve in Bell’s palsy might be related to the HSV [10].

CONCLUSION

This case represents, to the best of our knowledge, the first case of DD presenting with Bell’s palsy. The cause of Bell’s palsy in this case, whether it is the reactivation of HSV due to disease per se or due to stress after skin biopsy remains eluded. This case underlines the importance of recognizing HSV infection in DD and also highlights the possible therapeutic effect of antiviral agents in DD.

Consent

The examination of the patient was conducted according to the Declaration of Helsinki principles.

REFERENCES