Net Case

Epidermolytic hyperkeratosis with a rare digital contracture

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ABSTRACT

A sixteen year-old male patient with no history of consanguinity in the family, reported with patchy, thickened lichenified plaques over the whole body. Some areas had normal skin while some were Blaschkoid lesions. The child had delayed milestones along with hypogonadism. Digital contracture with palmoplantar keratoderma was present. Histopathology showed characteristic vacuolar degeneration of the upper epidermis and suprabasilar keratinocytes with hyperkeratosis.

Key Words: Epidermolytic hyperkeratosis, Digital contracture

INTRODUCTION

Epidermolytic hyperkeratosis (EHK) is transmitted as an autosomal dominant trait with a prevalence of approximately 1 in 200,000 to 300,000. However, there is a high frequency of spontaneous mutation and as many as one-half of the cases have no family history and have new mutational events.[1]

CASE REPORT

We report a 16 year-old male patient from Murshidabad with patchy hyperkeratosis all over the body and face, present since a year of age. The lichenified plaques were extensive on the knees, elbows and all flexors. There were also linear streaks of similar lichenified rugose plaques. The patient complained of generalized itching, palmoplantar keratoderma and digital contractures of both hands involving middle, ring and little fingers. Both palms and soles showed keratoderma, but there was no uniform thickening in palms and soles and a transgrediens pattern was seen. There was no history of erythroderma or colloidion baby at any point of time during the evolution of the disease. The patient had reported a retarded growth and delayed developmental milestones. The size of the testis was 3 cm x 3 cm (smaller). Routine semen analysis revealed oligospermia with very sluggish motility. There was no history of any similar disease or consanguinity in the family. Histopathology revealed vacuolar degeneration of the upper epidermis and suprabasilar keratinocytes with hyperkeratosis compatible with the clinical diagnosis.

DISCUSSION

Sporadic EHK due to postzygotic spontaneous mutation during embryogenesis can present in a mosaic pattern of skin involvement. Areas of hyperkeratosis alternating with normal skin are often distributed in streaks along Blaschko’s lines.[3] The characteristic histopathology of vacuolar degeneration of epidermis and associated hyperkeratosis are the hallmarks of the disease.

A useful differentiating characteristic feature is the presence or absence of severe palmar / plantar hyperkeratosis. Three subtypes have palm / sole involvement (PS-type), while the other three have no palm / sole involvement (NPS-type) [Table 1].

We therefore report this case of EHK (PS-2) type with
occasional linear Blaschkoid strips of hyperkeratotic lesions, possibly suggesting a sporadic mosaic mutation of hystrix pattern of EHK. Digital contracture and palmoplantar hyperkeratosis in association with EHK is reported in PS variants. Only PS-2 types have shown digital contracture, as seen in our patient. Retarded growth and delayed developmental milestones are not very common in EHK. We report this case because of these atypical features.

Table 1: Clinical subtypes of epidermolytic hyperkeratosis

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>NPS-1</th>
<th>NPS-2</th>
<th>NPS-3</th>
<th>PS-1</th>
<th>PS-2</th>
<th>PS-3</th>
</tr>
</thead>
<tbody>
<tr>
<td>Palm/sole hyperkeratosis</td>
<td>_</td>
<td>_</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Palm/sole surface</td>
<td>Normal</td>
<td>Normal</td>
<td>Hyperlinear, minimal scale</td>
<td>Smooth</td>
<td>Smooth</td>
<td>Cerebriform</td>
</tr>
<tr>
<td>Digital contractures</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>+</td>
</tr>
<tr>
<td>Scale</td>
<td>Hystrix</td>
<td>Brown</td>
<td>Fine, white</td>
<td>Mild</td>
<td>White scale to peel</td>
<td>Tan</td>
</tr>
<tr>
<td>Distribution</td>
<td>Generalized</td>
<td>Generalized</td>
<td>Generalized</td>
<td>Localized</td>
<td>Generalized</td>
<td>Generalized</td>
</tr>
<tr>
<td>Erythroderma</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Blistering</td>
<td>Localized</td>
<td>+</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>-</td>
</tr>
</tbody>
</table>

NPS - No palm / sole, PS - Palm / sole

REFERENCES