Ectodermal dysplasia: familial report of six cases

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ABSTRACT

Ectodermal dysplasia is a hereditary disease characterized by a congenital dysplasia of one or more ectodermal structures and their accessory appendages. There are two main types, Hypohidrotic/Christ-Seimens- Tourain syndrome and Hidrotic/Clouston syndrome. Common manifestations include defective hair follicles and eyebrows, frontal bossing with prominent supraorbital ridges, nasal bridge depression and protuberant lips. Intraorally common findings are anodontia or hypodontia and conical shaped teeth. The patient may suffer from dry skin, hyperthermia and unexplained high fever because of the deficiency of sweat glands. The present article reports unique case series of ectodermal dysplasia cases in two families, where three generations in the both the families were affected.

Key-words: Christ-Siemens-Touraine syndrome, ectodermal dysplasia, hypohidrosis, hypotrichosis, oligodontia

Introduction

Ectodermal dysplasia (ED) is a rare heterogeneous group of inherited disorder that share primary defects in the development of two or more tissues derived from ectoderm like skin, hair, nails, eccrine glands and teeth. The disorders are congenital, diffuse and non-progressive; more than 192 distinct disorders have been described. There are 2 major types, depending on the number and functionality of the sweat glands: X-linked anhidrotic or hypohidrotic, where sweat glands are either absent or significantly reduced in number (Christ-Siemens-Touraine syndrome) second one is hidrotic, where sweat glands are normal and the condition is inherited as autosomal dominant (Clouston’s
syndrome). The Hypohidrotic form exhibits the classic triad of hypohidrosis, hypotrichosis and hypodontia. Hypohydrotic form is most common type accounting for 80% of EDs, which is X-linked recessive, affects males and is inherited through a female carrier. In the hidrotic form teeth, hair and nails are affected; the sweat glands are usually not affected.

An estimated incidence of ED is about 7 in 10,000 births and all Mendelian modes of inheritance have been reported. Typical facial features include frontal bossing, sunken cheeks, saddle nose and protuberant lips with wrinkled, hyperpigmented skin around the eyes. In some cases, mucous glands are absent in the upper respiratory tract and in the bronchus, esophagus, and duodenum. Scalp hair may be sparse and short sometimes, there may be complete absence of hairs. Dental manifestations include conical shaped teeth, hypodontia or complete anodontia, and delayed eruption of teeth. This article reports six cases of ectodermal dysplasia in two families where all the siblings are affected.

Case Report

Family I: All the three children were affected and presented with scanty hairs on scalp, eyebrows, eyelashes, dry skin and oligodontia. A five-year-old boy reported with a complaint of missing teeth in upper and lower jaws. His family history revealed similar complaint of missing teeth in his mother, maternal grandfather, younger and elder sisters. On extraoral examination showed frontal bossing, prominent supraorbital ridges with midfacial hypoplasia with no nail dystrophies. Examination of hairs showed a fine, sparse scalp hair with thin eyebrows and eyelashes.
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On radiographic investigation, OrthoPantomogram showed oligodontia. (Figure 1d) Extraoral examination of patients elder and younger sisters showed similar features. Intraoral examination and OPG view of elder and younger sisters showed oligodontia, with tongue-tie in the elder sister. (Figure 2)

Family II: Both the children and their mother were affected, past family history revealed multiple missing teeth in their maternal grandmother. Twelve-year boy reported with a complaint of deposits on his teeth surface and missing teeth in lower front region of jaw. Extraoral examination (Figure 3a) showed prominent supraorbital ridge with scanty hairs on head and eyebrows. Intraoral examination (Figure 3b & 3c) and OPG showed oligodontia.
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Fig. 3c Intraoral picture of mandible showing absence of multiple teeth

Discussion
Ectodermal dysplasia was first described by Thurnam [1, 5] in 1848. Freire-Maia and Pinheiro proposed the first classification system of the EDs in 1982 and classified it into different subgroups according to the presence or absence of hair anomalies or trichodysplasias, dental abnormalities, nail abnormalities or onychodysplasias, [1] and dyshidrosis. In 2003, Lamartine [6] reclassified it into the following four functional groups based on the underlying pathophysiologic defect: cell-to-cell communication and signaling, adhesion, development and others.

Thurman first reported two male cousins and their maternal grandmother with a hereditary syndrome associated with sparse hair, missing teeth and dry skin. In our cases, in both the families all siblings were affected with history of similar complaints in their parents and grandparents. ED is an X linked recessive disorder affecting males and is inherited through female carriers. Clinical findings in carrier females are same as those in affected males. One third of the carriers appears healthy, another third of them show mild symptoms, and the last third exhibits significant symptoms, but often milder than the affected males. In our cases, both males and females were affected but features were more prominent in males.

The presentation of facial deformity, dry skin, and sparse hair in this report is similar to previous reports. These features are due to anomalies of the skin appendages, which include the hair follicles, sweat glands and sebaceous glands. [7-10]

Pure ED is characterized by defects in ectodermal structures alone, while ED syndromes are combined with other anomalies like cleft lip and palate. In autosomal recessive condition, there is a total absence of permanent teeth with or without taurodontism of primary molars, which was not observed in our patients.

Cases have been reported where both primary and permanent teeth were congenitally missing, [3] which was seen in all our cases. Few cases of tongue-tie have also been reported which was noted in one of the female patient.

The family history of similar clinical features helps in diagnosis of ED, as seen in our cases. ED is a rare genodermatosis, it is invariably characterized by its clinical features. The dermatologist, pediatrician, and dentist are usually the medical personnel that these patients first visit, and therefore these individuals should be acquainted with this disease in order to provide appropriate care. A multidisciplinary approach of treatment involving pediatric dentist, orthodontist, prosthodontist, and oral-maxillofacial surgeon are required to help these patients to lead a normal life.

References
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