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Ichthyosis follicularis with alopecia and photophobia syndrome (IFAP): A Case Report

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Abstract

IFAP syndrome is a rare autosomal recessive X-linked disease characterized by the triad of alopecia universalis, severe photophobia, and follicular ichthyosis. It is caused by loss of function of the gene MBTPS2. Its severity varies and there are only a few reports in the literature. We present a patient with characteristic clinical features and a mutation not previously reported.

Keywords: Icthyosis, alopecia; photophobia

Introduction

IFAP syndrome is an unusual autosomal recessive X-linked disease characterized by the triad of alopecia universalis, severe photophobia, and follicular ichthyosis [1-4]. Other findings may determine variations in its clinical presentation [1,4,5]. It is caused by loss of function of the gene MBTPS2, essential for cholesterol homeostasis. Twenty different mutations of this gene have been reported [2]. Treatment is based on emollients and retinoids [1,5].

Case Synopsis

An 18-month-old boy with a history of collodion baby and congenital hypothyroidism was referred for cutaneous signs present since birth. These were characterized by spiny hyperkeratotic papules localized on scalp, ears, cheeks, elbows, and knees and periungual erythema on his hands associated with generalized alopecia and photophobia (Figure 1).

He also showed facial dysmorphism, atopic dermatitis-like lesions, and plantar keratoderma (Figure 2).

The patient’s height was below normal for his age. Dysgeusia was also observed. Ophthalmic examination revealed keratitis and corneal neovascularization. The pathology of follicular papules showed follicular keratosis. The genetic study found a mutation in the gene MBTPS2 E11 homozygous c. 1433 C> A, p. 478 A>D. This mutation had not been reported previously. These data confirmed the presumptive diagnosis of IFAP syndrome. The patient was treated with levothyroxine, general humectant measures, and eye lubrication.

Figure 1. A, B) Spiny hyperkeratotic papules localized on the scalp and left arm. C) Dystrophic nail with periungual erythema.
Case Discussion

IFAP syndrome is a rare autosomal recessive X-linked disease [1-4]. It is characterized by the triad of alopecia universalis, severe photophobia, and spiny keratotic follicular papules distributed symmetrically on scalp and along the extensor surfaces of the extremities [5]. At birth, patients may present as collodion babies. Alopecia is non scarring, congenital [5], non-progressive, and widespread. Up to 40% of patients also have psoriasiform plaques, cheilitis, hypohydrosis, nail dystrophy, atopic dermatitis, and rarely, keratoderma [4,5]. Photophobia may be congenital or develop later in life. It is owing to corneal defects: erosion, ulcers, scars, and neovascularization [1]. Blindness and myopia are common [1,5]. Mental retardation, seizures, and hypotonia are frequent [1,4,5]. Nevertheless, our patient did not show neurologic involvement. Other findings include asthma, allergic reactions, recurrent infections, and enteropathy [4,5]. Histopathology of skin lesions is nonspecific [4,5].

Loss of function of the MBTPS2 gene is the cause of this disease. This gene encodes a zinc metalloproteinase intramembrane essential for cholesterol homeostasis and response of the endoplasmic reticulum to stress [3]. Clinical severity ranges from mild to severe cutaneous forms [2]. At present, 20 different mutations of this gene [2] have been reported. In our patient, a previously unreported mutation was found. Treatment for follicular ichthyosis is based on urea emollients, local corticosteroids, and retinoids [1,5]. Photophobia and seizures may improve spontaneously [1], although sometimes antiseizure drugs are required [5]. Life expectancy varies between death in the neonatal period and a normal survival [4,5].

References