Case Report

Dyschromatosis Symmetrica Hereditaria of Late Onset?

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Dyschromatosis symmetrica hereditaria (DSH), also known as reticulated acropigmentation of Dohi, is an autosomal dominant disease with high penetrance, characterized by hypo- and hyperpigmented macules of varying sizes on the dorsal of the extremities with reticulated pattern. This paper presents a female patient with typical dermatological lesions, but only diagnosed in adulthood. It is necessary to perform differential diagnosis with other pigmentary disorders. This entity is not very common in South America, and the vast majority of cases were described in Japanese population. Since it is a benign disease, it is important to be aware of this diagnosis in order to establish the correct conduct for these patients.

1. Introduction

Dyschromatoses are characterized by the presence of hyper- and hypopigmented macules arranged in a reticular pattern. Dyschromatosis symmetrica hereditaria (DSH) is a rare genodermatosis, autosomal dominant with high penetrance, with some sporadic reported cases. It is often reported in Japanese patients, but there have been also cases in India, Europe, and South America [1, 2]. After the literature review in PubMed database, this is the fourth case reported in Brazil [3, 4].

2. Case Report

A 40-year-old female patient presented with multiple small hyperchromic and hypochromic macules distributed symmetrically on the dorsum of the hands and feet (Figures 1 and 2). She noted these lesions when she was 26 years old, and they were asymptomatic. Moreover multiple freckle-like pigmented macules were also noted on her face (Figure 3). She denies other comorbidities. Investigation for autoimmune disease and other laboratory findings were negative. Family history was positive with a nephew with similar lesions.

Two different biopsies were performed from hypochromic and hyperchromic areas. From hypochromic lesion, histology demonstrated a compact stratum corneum and discrete acanthosis. Using Fontana Masson staining, hypochromic lesion demonstrated marked decrease, with almost absence, of melanin. On the other hand, the hyperchromic lesion demonstrated an intense pigmentation (Figures 4(a) and 4(b)). Those findings were compatible with the clinical hypothesis of DSH.

3. Discussion

DSH is a genodermatosis characterized by multiple small hypo- and hyperpigmented macules, with irregular size and shape, symmetrically distributed on the backs of the hands and feet. Some patients also show freckle-like macules on the face. Palms, plants, and mucous membranes are free from the disorder [5]. Lesions appear in childhood, usually before the age of six, remaining during life without any color or distribution changes after stabilization in adolescence. In general, the involvement is limited to skin, without systemic involvement or evidence of photosensitivity. However, there are isolated reports of association with neurofibromatosis type I, thalassemia, polydactyly, and torsion dystonia [6].

Since 2003, a genetic mutation has been identified on chromosome 1q11-1q21 as responsible for the production and
distribution changes of melanin. Miyamura et al. were the first to identify heterozygous mutations of the gene DSRAD or ADAR1, responsible for codifying the double-stranded adenosine deaminase specific RNA, as the cause of DSH [1, 6]. Despite more than 90 different mutations in the DSRAD gene having been described in the literature, it is still uncertain how these changes can cause the same phenotype [7].

Histologically, in hypopigmented areas, the deposit of melanin is scarce, in contrast to the areas of hyperpigmentation that contain melanocytes with an increased metabolic activity and high concentration of melanosomes [6]. The number of melanocytes in the areas of hypopigmentation is diminished compared to individuals without the disease [1].

This disease must be differentiated from other pigmented disorders such as reticulate acropigmentation of Kita-mura, which is marked by presence of atrophy and absence of hypopigmented lesions; Dowling-Degos disease by reticulate hyperpigmentation in the body's folds, accompanied by comedogenic lesions on back and neck, and depressed or pitted scars; initial cases of xeroderma pigmentosum distinguished by the development of more serious symptoms of xerosis, atrophy, telangiectasia, and tumors in photo-exposed areas; dyschromatosis universalis hereditaria with predominating lesions in trunk starting in childhood; vitiligo with repigmentation areas which can simulate hyperchromic macules of DSH but have perifollicular distribution [1, 2, 6, 8].

Interestingly the age of onset of DSH is 4.4 years old in average, which differs from our case (26 years old). Dowling-Degos disease and Galli-Galli disease are characterized by a later onset (24.5 and 45.6, resp.) and may be considered as a differential diagnosis [2]. Nevertheless, we favor the diagnosis of DSH once Dowling-Degos disease presents with reticulate hyperpigmented macules in the flexures and large body folds. Galli-Galli disease is characterized by reticulated pigmented and hyperkeratotic erythematosus macules and papules, located in the same areas of Dowling-Degos disease, but has a very peculiar histology demonstrating moderate-to-severe acantholysis of the suprabasal epidermis, which was
not observed in our case. Adult-onset dyschromatosis often is secondary to chemicals, drugs, physical agents, cutaneous lupus erythematosus, or infection [9]. We excluded those causes by history and laboratory investigation. We believe that this is a true case of DSH since the patient has very typical clinical findings and positive family history of similar lesions. It is possible that lesions have been unnoticed until adulthood.

No treatment is effective for this genodermatosis [6]. The exact frequency of the DSH is unknown since the main changes are confined to the skin with no systemic involvement, and many cases remain unreported. Despite the several reports and genetic studies from East Asian countries, the DSH is a very rare diagnosis in Brazil. We report this case due to its rare occurrence in South America and in view of the need of proper diagnosis and guidance for these patients.

Conflict of Interests
The authors declare that there is no conflict of interests regarding the publication of this paper.

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