AN EPIDEMIOLOGICAL ANALYSIS OF CHILDREN AND ADOLESCENTS PSORIASIS IN A TERTIARY REFERRAL DERMATOLOGY INSTITUTE IN THE DOMINICAN REPUBLIC

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Introduction
Psoriasis is one of the most common chronic inflammatory diseases, affecting about 3.5% of the population. Despite psoriasis carries a high risk of morbidity, few epidemiological studies provide estimates on the prevalence of psoriasis in children.

Aim: The objective of this study is to report the frequency of children and adolescents diagnosed with psoriasis at Instituto Dermatologico y Cirugía de Piel “Dr. Huberto Bogaert Díaz” (IDCP-DHBD) between March 2007 and March 2012.

Material and Methods: Examination were done on 76 children and adolescents with psoriasis confirmed by histopathological examinations at the IDCP DHBD in the Dominican Republic between March 2007 and March 2012. The data was retrospectively reviewed to assess age, gender, clinical localization, treatment modalities and delay in diagnosis.

Results: The frequency of children and adolescents with psoriasis among dermatological patients was 0.96 cases for every 10,000 patients seen in the Institute. There were 43 (56.6%) girls and 37 (43.4%) boys. The mean age of onset was 14 years. Children from 0-2 years, were the least affected with 3% of the cases. The most frequent site of onset were the trunk (28.8%) and the scalp (27.4%). 67% of the patients had 2 or more sites involved. The mean delay in diagnosis was 6 months. Topical therapy was the treatment of choice in all the patients except one.

Conclusions: Even though psoriasis may cause a profound impact on the quality of life of children and adolescents the epidemiological data in the countries of Central America and the Caribbean is scarce.

Key words: psoriasis; children; Dominican Republic

Etiology
Psoriasis is a chronic inflammatory condition probably mediated by T lymphocytes, endothelial cells, dendritic cells, monocytes, neutrophils, keratinocytes and cytokines and chemokines characterized by hyperproliferation of keratinocytes, endothelial vascular proliferation and an infiltrate of inflammatory cells. The IL-23/Th17 seems to be crucial in the pathogenesis of psoriasis, mediating the host’s inflammatory response [14].
It has been demonstrated that the PSORS1 gene is determinant in the early onset of non-pustular type 1 psoriasis [5,15]. The HLA-Cw6 is the most important allele in the PSORS1 locus which confers susceptibility for the early onset of the disease [2,5,16,17].

Contrary to what has been described in adults, infections such as pharyngitis or skin trauma may trigger the disease [5,18]. Additionally, the onset of new lesions in periods of emotional stress is seen more frequently in children [18].

**Aim**

To report the number of diagnosed cases of psoriasis in children and adolescents at Instituto Dermatológico y Cirugía de Piel “Dr. Huberto Bogaert Díaz” (IDCP-DHBD) between March 2007 and March 2012.

**Material and Methods**

This is a retrospective epidemiologic study from 76 children and adolescents with psoriasis confirmed by histopathological examinations at the IDCP DHBD in the Dominican Republic between March 2007 and March 2012. Information data was obtained from the clinical and histopathology records of the patients seen at the Institute. The data was retrospectively reviewed to assess age, gender, clinical localization, treatment modalities and delay in diagnosis.

**Results**

In total 76 children and adolescents with psoriasis were included in the study. The center registered a total of 789,558 patients seen in the clinical dermatology division in the period of time of the study. The frequency of children and adolescents with psoriasis among dermatological patients was 0.96 cases for every 10,000 patients seen in the Institute (Fig. 2). There were 43 (56.6%) girls and 37 (43.4%) boys, with a female to male ratio of 1.3:1 (Fig. 3). The mean age of onset was 14 years, with a peak age of onset seen in the group of adolescents from 12-18 years with 40.2% of the cases. Children from 0-2 years, were the least affected with 3% of the cases (Fig. 4). The most frequent site of onset were the trunk (28.8%) and the scalp (27.4%) (Fig. 5). It was found that 67% of the patients had 2 or more sites involved, while 32% of patients were affected in one body site.

The period between the onset of the first skin lesions and the final diagnosis was designated as the delay in diagnosis. The mean delay in diagnosis was 6 months. Topical therapy was the treatment of choice in all the patients except one. Antralin was prescribed for 42% of the patients followed by betamethasone combined with calcipotriol, cade emulsion, topical corticosteroids and salicylic acid ointments. Systemic therapy with methotrexate was used in only one patient.
Discussion

In terms of age of onset our data shows that 3.4% of patients were infants of 0-2 years, 12.6% less than 5 years and 54% less than 15 years. This results differs to the data of other regions of the world, for example in Australia, where 27% of the cases of psoriasis in children were reported in the group of patients of 0 - 2 years. But they are similar to areas like California which reports 2% of affected patients less than 2 years, 7% in 5 years or less, and 45% in the group of 12 years or less [12].

There have been reported racial differences on the groups affected by psoriasis; White and Asians seem to be the most affected while Hispanics and blacks are the least affected [12]. Our data reflects a low frequency of children and adolescents psoriasis. There is a mixture of races in the Caribbean area where there is a significant African American population.

In the present series, girls were slightly more affected than boys in a 1.3:1 ratio. It was interesting to see that 67% of the patients presented with 2 or more body segments involved, which could correlate with a delay in the diagnosis found of 6 months. There is a need to educate the population about what is psoriasis, how the symptoms may be found in children and adolescents and how they may differ from the typical presentation of the adults.

Treatment

It is important to consider certain facts that in children could alter the course of the disease, for instance, infections may trigger the onset of psoriasis or may perpetuate it. [19] In this young age it is essential to verify the weight and height of the children and adolescents, especially to detect metabolic syndrome risk factors [20,21], and also search for sign and symptoms of psoriatic arthritis. The objective of the treatment is to improve the physical symptoms minimizing the effect of the disease in the psychosocial development of the child. When choosing a therapy it is imperative to think in the future health of the child as well as in his growth and development [7].

Based on the European consensus on the management of juvenile psoriasis [22] we propose that the first line of treatment for induction to remission on mild psoriasis defined as PASI ≤ 10, or BSA ≤ 10 or DLQI ≤ 10 should be topical steroids class II- IV; for transitional therapy topical steroids class IV- VI or calcipotriol plus steroids; and for maintenance therapy it should be considered the use of mild potency steroids, calcipotriol, tacrolimus and finally tar, antraline and salicylic acid compounds.

In cases of moderate-to-severe psoriasis, defined as PASI ≥ 10 or BSA ≥ 10 or DLQI ≥ 10, it is indicative of the use of systemic therapy such as methotrexate, cyclosporine, oral retinoids or biologic agents. Alternatively narrow band ultraviolet B phototherapy could be considered for older children and adolescents. It is important to take into consideration the rotation, combination and sequential strategies when using these agents in order to improve the effectiveness of the treatment, its tolerance and to minimize the long term side effects of these agents (Fig. 6).

Conclusions

Even though psoriasis may cause a profound impact in the quality of life of children and adolescents the epidemiological data in the countries of Central America and the Caribbean is scarce. Psoriasis may impact the quality of life of children and adolescents and their parents in different ways than in adults, thus requiring separate studies. Many times the initial clinical presentation, the different trigger factors, and the different associations may be unrecognized by the parents and thus contribute to the delay in the diagnosis therefore more education should be given to the general population regarding this entity and specific therapeutic guidelines should be proposed.

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Figure 4. Age distribution of patients.

Figure 5. Most frequent site of onset.
REFERENCES