Abstract
The mastocytosis is characterized by local or systemic accumulation of mast cells of unknown etiologic, for which is included in groups of infrequent disease. It generally appears in the childhood or in adults between the third and fourth decade of life and it may lead to different clinical pictures. The case of a six month-old female is presented with lesion cutaneous and histopathological characteristics from this illness.

Keywords: Mastocytosis; Children; Progenitor cells; Urticaria Pigmentosa; VDRL

Introduction
Mastocytosis refers to a heterogeneous group of diseases characterized by the hyperplastic proliferation and accumulation of mast cells in one or more organs. The skin is the most organ frequently affected [1,2]. The incidence of 2 cases per 300,000 people per year is estimated and may be observed in all ethnic group. The disease is higher predominant in children and in adults between the third and fourth decade of life [3,4]. Children are more affected before the age of six months in 50% of the cases [5].

The etiology of the disease remains unknown. The current hypothesis is that occurs an initial mutation involving KIT D816V. It triggers an increase in differentiation from progenitor cells and cell clustering, through adhesion molecules CD2, CD58, CD29, CD47 and others [1].

A new classification was made by the World Health Organization in 2008 which divides in cutaneous mastocytosis and systemic mastocytosis. The systemic mastocytosis characterized with involvement of the organs including liver, spleen, lymph nodes and/or bone marrow. Among the cutaneous forms, there are three ways: maculopapular cutaneous mastocytosis, cutaneous mastocytoma and diffuse cutaneous mastocytosis [6]. The patient may present more than one clinical variant. Some studies indicate a slight predominance in male [7,8].

There is a great heterogeneity of clinical presentation and prognosis even within the same clinical form and it may be related to patient’s age. Mastocytosis may be associated with hypersensitivity, chronic diseases and neoplasms. This disease has an indolent initial course which may delay the diagnosis of this uncommon pathology in medical practice.

Case Presentation
A 6-month-old female, caucasian, born and resident in João Pessoa, Brazil, presents a yellow-brownish solitary nodule with 2 cm on the face, which appeared at one month of age and remained of stable size (Figure 1). No other symptoms or previous pathologies are reported. The patient didn’t use any medication preceding the cutaneous lesion. He does not have any similar cases in the family. As possible diagnosis, were suggested the hypotheses of pilomatrixoma (Malherbe calcified epithelioma), tricoepitelioma, juvenile xanthogranuloma, X histiocytosis, cutaneous sarcoïdosis, Spitz nevus, dermatofibroma and eruptive xanthoma. We chose to perform the skin biopsy whose histopathology was compatible with the diagnosis of solitary mastocytoma (mastocytosis, urticaria pigmentosa). Long bone radiography, VDRL, blood count, abdominal ultrasonography, liver and renal function were requested and all showed no alterations.

Discussion
The Cutaneous mastocytoma is the second most common type of mastocytosis in childhood and presents as a single or multiple nodule of firm consistency and yellowish, pinkish or brownish color, usually located at the extremities of the limbs (Figure 2) [4]. If these nodules are exposed to any trauma including friction, they may lead to flushing and systemic symptom, which characterizes the Darier’s sign, considered pathognomonic of this illness [9].

The diagnosis is based on a typical cutaneous condition associated with the presence of focal infiltrate composed of mast cells in the dermis in cutaneous biopsy, after exclusion of criteria that establish the diagnosis of systemic mastocytosis, through laboratory tests (Figure 3). The lesions are symptomatic in 60 to
70% of the cases and the main complaints are pruritus, erythema and edema and the presentation of Darier’s sign [Figure 4] [8].

Conclusion

In this case report the patient had a single brownish nodule on the face, an unusual local of appearance of this dermatosis. He was asymptomatic and did not have the Darier’s sign. The most reliable marker for the severity and extent of disease is the serum level of tryptase [10]. In our reality, this test is not widely available and not performed by the patient, as well as the analysis of the mutation in receptor KIT, since histopathology presented characteristics that allowed the conclusion of the case.

There is no consensus on the laboratory tests that should be performed or the frequency with which they should be ordered in children with cutaneous mastocytosis, since most cases of children are restricted to the skin. Therefore, the definitive diagnosis of systemic involvement by the disease would require bone marrow biopsy, which is not routinely performed. In the case presentation, we chose to request long bone radiography, VDRL, blood count, abdominal ultrasonography, liver and renal function. It was planned an annual periodic evaluation of the patient.

There is no curative treatment for cutaneous mastocytosis, only symptomatic, which does not change the course of the disease. As the patient was asymptomatic at the time of the evaluation, no specific therapy was required. It was recommend to the mother to avoid physical, biological and stimulating factors for the release of histamine by mast cells. The mastocytoma was excised at the time of skin biopsy. The prognosis in childhood is good and usually the mastocytoma regresses throughout the first years of life [10].

Acknowledgement

None.

Conflicts of Interest

The authors declared that there no conflicts of interest.

References