

# Childhood leprosy: Clinical and epidemiological study in the Department of Dermatology, Clinicas Hospital, Faculty of Medical Sciences, National University of Asuncion-Paraguay, 2005-2014

**Beatriz Di Martino Ortiz, María Laura Sánchez, Celeste Valiente, Gabriela Martínez, Mirtha Rodriguez Masi, Oilda Knopfmacher, Lourdes Bolla de Lezcano**

*Department of Dermatology, Clinicas Hospital, Faculty of Medical Sciences, National University of Asuncion, Paraguay*

**Corresponding author:** Prof. Dra. Beatriz María Di Martino Ortiz, E-mail: beatrizdimartino@gmail.com

## ABSTRACT

**Introduction:** Leprosy in childhood is not a common finding. The risk of a child to develop the disease is 4 times greater in contact with close people and 9 times higher among household contacts. The maximum risk observed is when the contact is Multibacillary (MB) and intradomiciliary. Leprosy in childhood reflects the clinical characteristics of adult, with some peculiar aspects. Non-contagious forms (IL and TT) are common during childhood. The contagious forms (BB, LB and LL) are less frequent due to higher required incubation period. **Aim:** To describe the clinical and epidemiological characteristics of childhood leprosy in the Department of Dermatology, Clinicas Hospital from January 2005 to July 2014. **Methods:** Retrospective, observational cross-sectional study with an analytical component. **Results:** The total number of leprosy patients was 369, and of these 11 were pediatric patients (2.98%) with a predominance of males (8/11) from 3 to 16 years. The BI ranged from negative to 3+. 6/11 were MB. The evolution was good in all cases and two patients developed leproreactions. The lesions were predominant in facial location. 6/11 patients had family contacts. **Conclusions:** Leprosy in children is more common than is reported, especially in endemic areas. In <5 years, the disease is very rare. More than half of the cases of children with leprosy have a positive contact. It is considered that in <5 years the spread is always intradomiciliary; this shows the importance of monitoring contacts, which will be possible with the determination of all stakeholders in order to banish the undetected cases and prevent damage.

**Key words:** Hansen disease; Leprosy; Childhood hansen disease

## INTRODUCTION

Hansen's disease is a contagious disease of chronic course caused by *Mycobacterium leprae* that has tropism for the skin, mucous membranes and peripheral nerves. Other organs may also be compromised.

Because the consequences, degree of disability and deformity that occurs, are indispensable early diagnosis and treatment, allowing an endemic disease control.

In the field of pediatric dermatology leprosy remains a little described and undervalued in daily practice, so it

becomes a diagnostic challenge because of the diversity of clinical manifestations that may occur, making necessary a thorough skin and neural examination in all children presenting suspicious lesions suggestive and an infectious source [1].

## Aims

### General

To describe the clinical and epidemiological characteristics of childhood leprosy in the Department of Dermatology, Clinicas Hospital, in the period from January 2005 to July 2014.

**How to cite this article:** Di Martino Ortiz B, Sánchez ML, Valiente C, Martínez G, Rodriguez Masi M, Knopfmacher O, de Lezcano LB. Childhood leprosy: Clinical and epidemiological study in the Department of Dermatology, Clinicas Hospital, Faculty of Medical Sciences, National University of Asuncion-Paraguay, 2005-2014. *Our Dermatol Online*. 2016;7(1):17-20.

**Submission:** 17.07.2015; **Acceptance:** 08.09.2015

**DOI:** 10.7241/ourd.20161.4

### Specific

1. To set the epidemiological characteristics of the study population.
2. To determine the clinical characteristics of leprosy in childhood.
3. To describe the type of treatment given and evolution.

## MATERIALS AND METHODS

### Design

Retrospective, observational cross-sectional study with an analytical component. The study was conducted at the Department of Dermatology of CH, FMS-NUA, between January 2005 and July 2014.

### Reference Population

Asunción is the capital of the Republic of Paraguay and is located on the right bank of the Paraguay River (which divides the country into two regions) standing in the Eastern Region. Its metropolitan area called Gran Asunción has a population of 2,870,000 inhabitants. Its area is 118 km<sup>2</sup> and population density 4,411 inhabitants/km<sup>2</sup>.

### Inclusion Criteria

All patients with leprosy in childhood, diagnosed clinically and with pathological confirmation, that have consulted in the Department of Dermatology of CH. FMS-NUA.

### Exclusion Criteria

No leprosy patients in pediatric age or no pathological confirmation of the disease.

### Sources

Medical records of patients with diagnosis of Hansen's disease in childhood.

## RESULTS

- The total number of leprosy patients was 369, and of these 11 were in pediatric age (2.98%).
- The disease predominated in males.
- The age of onset was from 3 years to 16 years.
- Bacillary Index (IB) ranged from negative to 3+.
- Six of the eleven cases presented Multi bacillary forms; the TT and LL forms were 4 cases each.
- The evolution was good in all cases and two developed leproreactions.
- The predominant location of the lesions was on the face.
- Six of the eleven cases had a family contact.

The summary of findings stated in Table 1.

## DISCUSSION

Pediatric patients suffering from leprosy were 11, which corresponds to 2.98% of the total infected patients (369), lower than in other series, as Terencio de las Aguas (7.7%) and Fakhouri et al (17.3%) [1-5].

Among the children affected, males predominated coinciding with a national study of another service of dermatology; other authors did not find any differences in sex [4,5].

Ages of children affected ranged from 3-16 years, with school children (6-12 years) and adolescents the hardest hit, coinciding with the literature reviewed [1].

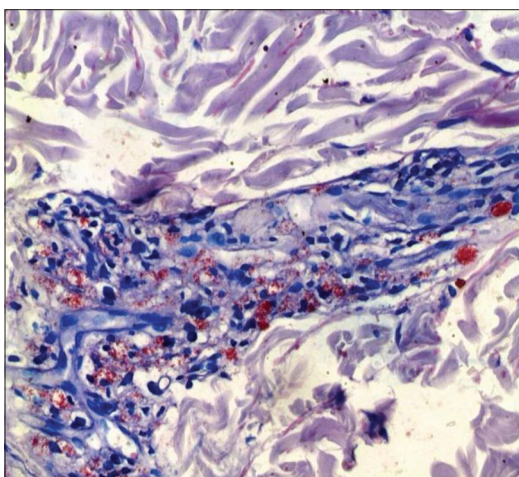
**Table 1:** Summary of findings

Sex/age/orig	FMH	CC	Local	Evol	PD	BI	Treat	Reac
F/3/R	Father LL	Papules	LLL	3 months	TT	Negative	PB	No
F/8/U	Mother LL	Hipocrom macules	LL and UL	1 month	IL	Negative	PB	No
F/10/U	Father LL	Hipocrom macules	RUL	2 years	BB	Negative	MB	No
M/11/U	No	Eritem macules	Face	1 year	TT	Negative	PB	No
M/12/R	No	Eritem macules	UL	3 years	LL	2+	MB	No
M/12/U	Mother LL	Hipocrom macules	Back	1 year	BL	2+	MB	Yes
M/13/U	No	Eritem macules	Face, UL, LL	3 weeks	LL	3+	MB	Yes
M/13/U	No	Hipocrom macules	Face	2 years	TT	Negative	PB	No
M/14/R	Father LL	Eritem macules	RLL	4 months	TT	Negative	PB	No
M/16/U	No	Ulcer	LL	6 years	LL	1+	MB	No
M/16/R	Aunt	Hipocrom macules	Face/back	2 years	LL	3+	MB	No

Sex: M: Male; F: Female, Age: Years, Origen: U: Urban; R: Rural, FMH: Familiar medical history, CC: Cause of consultation, Local: Localization of lesions, Evol: Evolution, PD: Pathological diagnosis, BI: Bacillary index, Treat: Treatment, Reac: Leprosy reactions, LLL: Left lower limb, LL: Lower limb, UL: Uper limb, RUL: Right uper limb, RLL: Right lower limb, PB: Pauci bacillary, MB: Multi bacillary, IL: Indeterminate leprosy, TT: Tuberculoid leprosy, LL: Lepromatous leprosy, BB: Borderline borderline leprosy, BL: Lepromatous borderline leprosy, BT: Tuberculoid borderline leprosy



**Figure 1:** (a) Clinic. Hypochromic macules, between 2 and 6 cm, net limits, in the back, (b) Clinic. Erythematous plates between 0.3 and 0.8 cm, net limits, jagged edges, with the center of colored skin ("inverted dish") in arms. There are nodules, also.



**Figure 2:** Histopathology. Peri adnexal chronic inflammatory infiltrate with multiple bacilli (ZiehlNeelsen 40X).

The BI ranged from negative to 3+, with 5 positive cases (45%), is inconsistent with the literature in which the positivity percentage is lower (2%). This is because the predominant number of MB cases in our study (6/11), opposite as usually PB cases described more frequent in children [5,6].

Evolution was good in all cases and two developed leproreactions.

The elementary dermatological more frequent lesion was a macule, with facial location, and in exposed areas, which is the most frequent site of occurrence in children with leprosy in other countries [7].

Six of eleven cases had a family contact. In most series this is the constant [1,3,8].

In developing countries, where leprosy is a public health problem, it is well accepted that children below five years are more likely to develop it than adults. About 17.13% of all cases of leprosy in India are in children under 15 years. Van Beers et al. show that the risk of a child developing leprosy is 4 times greater in contact with close people and 9 times higher among household contacts. The maximum risk is observed when the contact is Multibacillary and intradomicilliary [3].

Leprosy in childhood presents with a variety of clinical and histological manifestations, which necessitate a thorough skin examination in every child with suggestive or suspicious skin lesions and an infectious source (Figs 1a,b and 2). Many skin lesions are usually asymptomatic and often mimic other dermatologic pictures [1,3].

## CONCLUSIONS

1. Leprosy in children is more common than we tend to think, especially in endemic areas, as is our country [9]. In patients with less than five years, the disease is very rare [10].
2. More than half of the cases of children with leprosy have a positive contact.
3. It is consider that in patients with less than five years the spread is always intradomiciliary; this shows the importance of monitoring contacts, which will be possible with the determination of all stakeholders in order to banish the undetected cases and prevent damage.

## Statement of Human and Animal Rights

All procedures followed were in accordance with the ethical standards of the responsible committee on human experimentation (institutional and national) and with the Helsinki Declaration of 1975, as revised in 2008.

## Statement of Informed Consent

Informed consent was obtained from all patients for being included in the study.

## REFERENCES

1. Táquez A, Cerón C, Chaparro M, Sales A, Nery J, Miranda A, Lupi O. Lepra en la infancia. Desafío diagnóstico. Rev Argent Dermatol. 2011;92:2-10.
2. Fakhouri R, Soto M, Manini M, Margarido L. Nodular leprosy of childhood and tuberculoid leprosy: a comparative, morphologic,

- immunopathologic and quantitative study of skin tissue reaction. *Int J Lepr Other Mycobact Dis.* 2003;71:218-26.
3. Di Martino Ortiz B, Rodriguez Masi M, Knopfmacher O, Bolla de Lezcano L. Lepra infantil. Presentación de un caso. *Childhoodleprosy. Report of a case. Dermatol Online J.* 2011;17:13.
  4. Terencio de las Aguas J. Situación de la lepra en el mundo. *Med Cután Iber Lat Am.* 2005;33:191-2.
  5. Mata Jiménez O, Aguilar Aguilar N, Miranda A, Freitas de Souza M, Azulay R, da Costa Nery J. Lepra en la infancia: caracterización de parámetros clínicos en los estados reaccionales. *Med Cutan Iber Lat Am.* 2006;34:263-9.
  6. Dias Gomes C, Andrade Pontes M, SáGonçalves H, Oliveira G, Penna. *An Bras Dermatol.* 2005;80:283-8.
  7. Rodríguez E, Díaz O, Hernández E. *Boletín Epidemiológico Semanal.* 2013;21:1-13.
  8. Martinez Braga G, Guglielmone C, Di Martino Ortiz B, Bolla de Lezcano L, Aldama A, Mendoza G. Lepra infantil: estudio clínico y epidemiológico en dos servicios de dermatología del Paraguay. Periodo 2005-2011. *Fontilles, RevLeprol.* 2012;28:293-301.
  9. Rongioletti F, Gallo R, Cozzani E, Parodi A. Leprosy: a diagnostic trap for dermatopathologists in non endemic area. *Am J Dermopathol.* 2009;31:607-10.
  10. Dias Gomes CC, de Andrade Pontes MA, de Sá Gonçalves H, Oliveira Penna G. Clinical and epidemiological profile of patients diagnosed with leprosy in a reference center in the northeast of Brazil. *An Bras Dermatol.* 2005;80(supl 3):S283-8.

Copyright by Beatriz Di Martino Ortiz, et al. This is an open access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.  
**Source of Support:** Nil, **Conflict of Interest:** None declared.