Revista da Sociedade Brasileira de Medicina Tropical

Journal of the Brazilian Society of Tropical Medicine Vol.:52:e20190354: 2019

doi: 10.1590/0037-8682-0354-2019



Authors Reply

Authors' response to "De novo histoid leprosy—further points to be discussed"

Claudia J. Díaz^[1], Kevin Escandón-Vargas^[2] and Lina M. Piedrahíta-Rojas^[1]

[1]. Universidad del Valle, Hospital Universitario del Valle, School of Dermatology and Dermatological Surgery, Cali, Colombia.
[2]. Universidad del Valle, School of Medicine, Cali, Colombia.

Dear Editor:

We thank Dr. Pugazhenthan and colleagues for their interest in our recently published article "De novo histoid leprosy in a Colombian patient with multiple skin nodules on the ears and extremities" http://dx.doi.org/10.1590/0037-8682-0502-2016¹, which describes an unusual case of histoid leprosy in an adult male from Cali, Colombia, who did not exhibit previous clinical signs or treatment of the disease. While we appreciate and agree with their observations, we would like to provide some information about the management and diagnosis of leprosy in general and specifically in Colombia.

The diagnosis of histoid leprosy is challenging because of the clinical variations in the disease. Histoid leprosy may even go unnoticed by experts, thereby requiring a high clinical suspicion, especially in patients without apparent health concerns. A complete physical examination for epistaxis, skin nodules and thickening of nerves; histopathological examination of skin biopsy specimens from papules, nodules, plaques or tumorlike lesions; and skin smear examination (bacilloscopy) of the earlobes, nose, and elbows are all critical for early and accurate diagnosis of histoid leprosy^{1,2}.

Histoid leprosy can occur as a relapsing or recurrent disease on lepromatous leprosy and other forms of multibacillary leprosy, even when there is no history of inadequate or irregular therapy^{3,4}. Although irregular treatment and inadequate dapsone monotherapy are arguably the most common scenarios, cases of histoid leprosy have also been reported in patients treated with adequate doses of multibacillary-multidrug therapy (dapsone,

Corresponding author: Dr. Claudia J. Díaz.

e-mail: clajudiaz@yahoo.com Orcid: 0000-0003-3582-7748 Received 18 July 2019 Accepted 24 July 2019 clofazimine, rifampin)⁵ as well as in *de novo* (untreated) patients^{1,3,6}, as reported in our article. Patients with lepromatous leprosy who have received adequate treatment may develop leprosy as early as within 5 years following the cure, but also up to 20 years later⁷, prompting strict monitoring and epidemiological surveillance. Therefore, in Colombia, we recommend biannual follow-up of paucibacillary patients for 5 years and that of multibacillary patients for 10 years. Concerning leprosy patients' cohabitants, those with suspected disease are subjected to clinical assessment and laboratory diagnostic testing.

Finally, histoid leprosy poses a threat to campaigns aiming to eliminate leprosy because of the delay in diagnosis, frequent high bacillary index of infected patients, data inaccuracies, obstacles during follow-up of patients and their household contacts, and the potential for emerging resistance in *Mycobacterium leprae* to the treatment. Consequently, the Colombian Ministry of Health and Social Protection has established the healthcare route (*Ruta Integral de Atención en Salud* in Spanish) for leprosy management and the "National Strategic Plan for Prevention and Control of Hansen Disease 2016–2025" in line with the "2016–2020 Global Leprosy Strategy", which has the target of achieving <1 newly diagnosed leprosy patient with visible deformities per million people and no child cases diagnosed with leprosy and visible deformities.

REFERENCES

- Piedrahíta-Rojas LM, Díaz CJ, Escandón-Vargas K. De novo histoid leprosy in a Colombian patient with multiple skin nodules on the ears and extremities. Rev Soc Bras Med Trop. 2019;52:e20160502.
- Gupta SK. Histoid leprosy: review of the literature. Int J Dermatol. 2015;54(11):1283–8.
- Nair SP, Kumar GN. Tropical medicine rounds A clinical and histopathological study of histoid leprosy. Int J Dermatol. 2013;52:580-6.
- Kaur I, Dogra S, De D, Saikia UN. Histoid leprosy: A retrospective study of 40 cases from India. Br J Dermatol. 2009;160(2):305–10.

- Shaw IN, Rao GS, Natrajan MM, Balasundaram B, Ebenezer G. Relapse as histoid leprosy after receiving multidrug therapy (MDT); a report of three cases. Int J Lepr Other Mycobact Dis. 2000;68(3):272-6.
- Mendiratta V, Jain A, Chander R, Khan A, Barara M. A nine-year clinico-epidemiological study of Histoid Hansen in India. J Infect Dev Ctries. 2011;5(2):128–31.
- 7. Mansfield RE. Histoid leprosy. Arch Pathol. 1969;87(6):580–5.
- Ministerio de Salud y Protección Social. Plan Estratégico Nacional de Prevención y Control de la Enfermedad de Hansen 2016-2025
- "Compromiso de todos hacia un país libre de enfermedad de Hansen"; MinSalud, version 3.0, 2018. Available at: https://www.minsalud.gov.co/sites/rid/Lists/BibliotecaDigital/RIDE/VS/PP/ET/Plan-strategico-enfermedad-hansen-2016-2025.pdf
- World Health Organization. Global Leprosy Strategy 2016-2020: Accelerating towards a leprosy-free world. WHO Regional Office for South-East Asia; 2016. Available at: http://www.searo.who.int/ entity/global_leprosy_programme/documents/global_leprosy_ strategy_2020/en/.

