

Images in Infectious Diseases

Chronic paronychia as a manifestation of skin leishmaniasis

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FIGURE 1: Erythematous swelling with scales and eroded surface in the nail fold of the right ring finger.

A 42-year-old male, who was a rural worker in the municipality in the interior of Amazonas, Brazilian state, presented with swelling, erythema, and infiltration of ungual folds with eroded surface and scales on the right ring finger along with fever and pain for 45 days (**Figure 1**). Direct smear of the lesion, stained with Giemsa showed the presence of amastigote forms of *Leishmania* parasite upon microscopic examination, thereby diagnosing it as a case of cutaneous leishmaniasis (LC) (**Figure 2**). The patient was administered with miltefosine for 28 days but showed therapeutic failure 60 days after the treatment ended. Thereafter, pentavalent

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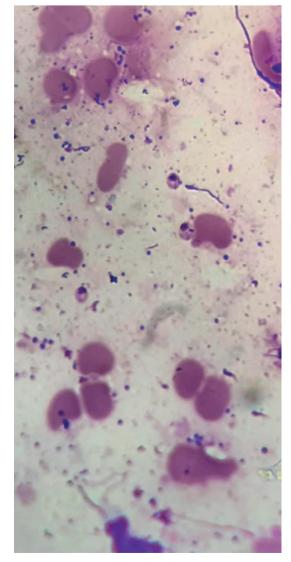


FIGURE 2: Slit-skin smear showing the presence of amastigote forms of the *Leishmania* parasite. Giemsa staining, original magnification 100x.

antimony (Sb^v) was administered at 20mg/kg daily total dose for 20 days, which resulted in complete cure (**Figure 3**). Initial therapy was conducted using miltefosine because the patient was part of a multicenter clinical trial that evaluated the effectiveness of this drug in combination with a topical immunomodulator.



FIGURE 3: Clinical inspection of the lesion after six months of treatment, showing atrophic scarring and improvement of the ungual plate.

CL is endemic in the Amazonas State and in most cases, it is caused by *Leishmania (Viannia) guyanensis*¹, exhibiting low response to Sb^v treatment with only 53% of cure rate². The classical evolution of this disease is characterized by the emergence of ulcers with infiltrated regular and elevated edges³, but atypical forms have also been described³. Despite the diversity of clinical presentations in leishmaniasis, paronychia, known as inflammation of the ungual folds, is poorly described, with only nine reports on PubMed. It is important to diagnose this disease, mainly in endemic regions, given the possibility of occurrence of permanent ungual dystrophy if the diagnosis and treatment are delayed.

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AUTHORS' CONTRIBUTION

LMS, JAOG: conception and design of the study, protocol writing, critical review and final approval of the manuscript; JLDS: contributed to the project administration, prepared the manuscript and figures.

CONFLICT OF INTEREST

The authors declare that there is no conflict of interest.

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