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# Cutaneous Leishmaniasis

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**CITATION:**

Cundiff L, Byrd R, Short C. Cutaneous leishmaniasis [published online December 6, 2019]. *Consi*

During a medical mission to the indigenous Cabécar people of Costa Rica, a 9-year-old girl presented with a long-standing concern of fleabites of several months' duration. On physical examination, the girl was noted to have multiple painless ulcerations of her left forearm (**Figure 1**), right upper arm (**Figure 2**), and chin (**Figure 3**).



**Figure 1.** Ulceration on the patient's left forearm (photo by Richard L. Byrd, MD).



**Figure 2.** Ulceration on the patient's right upper arm (photo by Richard L. Byrd, MD).



**Figure 3.** Ulceration on the patient's chin (photo by Richard L. Byrd, MD).

A clinical diagnosis of cutaneous leishmaniasis was made. The girl was given a 10-day course of suspected secondary infection of the lesions and was referred to the state health department for diagnosis and further treatment.

## **EPIDEMIOLOGY**

Leishmaniasis is a vector-borne parasitic disease endemic to 98 countries worldwide. It is primarily tropical regions of Southern Europe, Northern Africa, the Middle East, Central and South America subcontinent.<sup>1,2</sup> With an estimated global annual incidence rate of between 0.9 million and 1.7 million, leishmaniasis remains one of the most common neglected tropical diseases.<sup>2</sup> The Centers for Disease Prevention reports isolated cases of leishmaniasis originating in Texas and Oklahoma, although the

not typically arise in the United States. However, cases may be diagnosed in travelers, immigrant personnel upon their return to the United States and therefore remains an important clinical consi

The causative organism is the protozoan *Leishmania*. At least 20 species of *Leishmania* have been causing a range of clinical manifestations in different geographical areas.<sup>1</sup> The parasite is transmitted by infected *Phlebotomus* sand flies, with more than 98 species of this vector having been identified.<sup>2</sup> Canines, and humans serve as the primary reservoirs of *Leishmania*, and thus this disease undergoes zoonotic (animal-to-human) and anthroponotic (human-to-animal) transmission when female sand flies feed on meals from their mammalian hosts.<sup>2</sup>

Three clinical syndromes are possible, depending on the *Leishmania* species. Once inside the host, the promastigote develops into an amastigote within macrophages of the skin, causing the cutaneous disease.<sup>2</sup> Mucosal leishmaniasis can evolve from dissemination of the parasite from the skin to the naso-oropharyngeal mucosa, which can erode the nasal cartilage and produce a disfiguring scar. Visceral leishmaniasis (kala-azar) occurs.<sup>2</sup> (This review focuses primarily on the diagnosis and treatment of cutaneous manifestations, as seen in our patient.)

## DIAGNOSIS

Cutaneous leishmaniasis most commonly presents with a painless papule developing at the site of the bite. It increases in size and eventually crusts and ulcerates.<sup>1</sup> Lesions may develop between 2 weeks and several months after the initial bite; some people never develop clinical manifestations.<sup>1</sup> Laboratory evaluation is necessary to rule out other etiologies (**Table**).<sup>3,4</sup>

## References

1. Piscopo TV, Mallia Azzopardi C. Leishmaniasis. *Postgrad Med J*. 2007;83(976):649-657. doi:10.1136/pgmj.2006.047340corr1
2. Steverding D. The history of leishmaniasis. *Parasit Vectors*. 2017;10(1):8 doi:10.1186/s13071-017-2028-5
3. Tirelli F, Vernal S, Roselino AM. Final diagnosis of 86 cases included in differential diagnosis of American tegumentary leishmaniasis in a Brazilian sample: a retrospective cross-sectional study. *An Bras Dermatol*. 2017;92(5):642-648. doi:10.1590/abd1806-4841.20175794
4. Aronson N. Cutaneous leishmaniasis: clinical manifestations and diagnosis. UpToDate. <https://www.uptodate.com/contents/cutaneous-leishmaniasis-clinical-manifestations-and-diagnosis>. Updated June 18, 2019. Accessed December 6, 2019.
5. Braz LMA. Tegumentary leishmaniasis diagnosis: what happened with MST (Montenegro skin test) in Brazil? *Rev Inst Med Trop Sao Paulo*. 2019;61:e17. doi:10.1590/S1678-9946201961017
6. Zanetti ADS, Sato CM, Longhi FG, Ferreira SMB, Espinosa OA. Diagnostic accuracy of enzyme-linked immunosorbent assays to detect anti-*Leishmania* antibodies in patients with American tegumentary leishmaniasis: a systematic review. *Rev Inst Med Trop Sao Paulo*. 2019;61:e42. doi:10.1590/S1678-9946201961042

7. Wyllie S, Cunningham ML, Fairlamb AH. Dual action of antimonial drugs on thiol redox metabolism in the human pathogen *Leishmania donovani*. *J Biol Chem*. 2004;279(38):39925-39932. doi:10.1074/jbc.M405635200
8. Rugani JN, Gontijo CMF, Frézard F, Soares RP, Monte-Neto RLD. Antimony resistance in *Leishmania (Viannia) braziliensis* clinical isolates from atypical lesions associates with increased ARM56/ARM58 transcripts and reduced drug uptake. *Mem Inst Oswaldo Cruz*. 2019;114:e190111. doi:10.1590/0074-02760190111
9. Soyer TG, Mendonca DVC, Tavares GSV, et al. Evaluation of the *in vitro* and *in vivo* antileishmanial activity of a chloroquinolin derivative against *Leishmania* species capable of causing tegumentary and visceral leishmaniasis. *Exp Parasitol*. 2019;199:30-37. doi:10.1016/j.exppara.20102.019
10. Darcis G, Van der Auwera G, Giot JB, et al. Recurrence of visceral and muco-cutaneous leishmaniasis in a patient under immunosuppressive therapy. *BMC Infect Dis*. 2017;17(1):47-x. doi:1186/s12879-017-2571-x
11. Ribeiro RR, Michalick MSM, da Silva ME, Dos Santos CCP, Frézard FJG, da Silva SM. Canine leishmaniasis: an overview of the current status and strategies for control. *Biomed Res Int*. 2018;2018:3296893. doi:10.1155/2018/3296893
12. Ghorbani M, Farhoudi R. Leishmaniasis in humans: drug or vaccine therapy? *Drug Des Devel Ther*. 2017;12:25-40. doi:10.2147/DDDT.S146521