

CARTA AO EDITOR

CUTANEOUS LEISHMANIASIS IN ECUADOR

Sir,

While we welcome the publication of the DIFMA test using *Leishmania* Genus specific monoclonal antibody¹ we are disappointed that the Authors did not include certain fundamental details. In particular, the Authors do not address whether the DIFMA test is equally sensitive for all *Leishmania* spp. reported in Ecuador, for example the DIFMA test may be less suitable for the diagnosis of those *Leishmania* spp. which induce lesions with relatively low amastigote densities. They states in their introductory paragraph that *Leishmania (Viannia) braziliensis*, *L. (V) panamensis*, *L. (V) guyanensis*, *L. (L) mexicana* and *L. (L) amazonensis* are all present in Ecuador. However they do not indicate the species present in the 90 active lesions tested in their patient cohort. Certainly *L. (V) braziliensis*, will not produce amastigotes readily detectable for their DIFMA test. We have no experience of *L. (V) panamensis*. *L. (L) amazonensis* are rare human infections in our experience because of the non human blood meal preference of vector phlebotomine. This leaves *L. (V) guyanensis* which as is the case in Manaus is easily detected in skin scrapings, culture and histology. The last is suggested by the high amastigote rate on histology (74%) found by Chico et al¹. Although more than 40 cultural isolates were made no taxonomy is given.

One can speculate on the value of such precise taxonomy if the situation in a transmission area is defined. For us in Tres Bracos, Bahia, Brazil there is no point in doing further taxonomy since it is almost a pure monotransmission of *L. (V) braziliensis*². Possibly the Authors of this paper are working with isolates from a region where *L. (V) guyanensis* and/or *L. (L) mexicana* are the predominant circulating parasites³.

REFERENCES

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