PUBLICATION REPORT

1693

8/92

ISOLATION AND CHARACTERIZATION OF LEISHMANIA MAJOR
FROM PHILETOMUS PAPATASI AND MILITARY PERSONNEL
IN NORTH SINAI, EGYPT

BY

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92-10-2-057

92-26392
Short Report

Isolation and characterization of Leishmania major from Phlebotomus papatasi and military personnel in north Sinai, Egypt

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Since October 1982, cutaneous leishmaniasis has been repeatedly diagnosed among military members of the multinational force and observers (MFO), an international peace keeping force based in the north Sinai desert of Egypt (Dunn & Smertz, 1983; Mansour et al., 1987). In an effort to assist in the recognition, prevention, and treatment of this disease among MFO personnel, the US Naval Medical Research Unit no. 3 initiated epidemiological studies in July 1989 at locations in north-eastern Sinai. As part of these epidemiological investigations, an entomological survey sought to determine the seasonal dynamics of local sandflies and the prevalence of Leishmania infection among these potential vectors.

Sandflies were obtained from human landing and resting collections during the months of July, August and September, 1989. Flies were cryopreserved following a modification of the procedure developed by Young et al. (1987). Sandfly pools in cryopreservant were slowly frozen on dry ice and transferred to liquid nitrogen for storage until examined. Cryopreserved sandflies were thawed and individually dissected on Whatman No. 1 filter paper using sterile techniques. The midgut and surrounding media from these specimens containing promastigotes were removed from the slide, triturated in approximately 1.0 ml of sterile PBS with antibiotics (as above), and inoculated into NN medium and into the hind footpads of BALB/c mice. These procedures were repeated monthly for the first three months and then monthly thereafter for the duration of the study. Culture supernatants from infected cultures were examined for the presence of promastigotes by phase microscopy throughout the culture period. Inoculated Balb/c mice were monitored for development of lesions at the inoculation site. Lesions were scored on a scale of 0-7, with 0 representing no visible pathology. Lesions were scored at 1-2 week intervals.

The promastigotes were centrifuged, washed, resuspended in PBS, and stored at -80°C until isoenzyme electrophoresis was performed. Promastigotes were serotyped according to the method of Schnur & Zuckerman (1977), using specific parasite excretory factors (EF) present in the culture medium. Comparative cellulose acetate electrophoresis against World Health Organization reference strains of L. major MHOM/IL/87/Jericho-II (= LRC-L137), L. tropica MHOM/SU/58/OD (= LRC-L39), and L. donovani MHOM/IN/80 DD8 was used to identify 2 sandfly isolates, IPAP/EG/89-Si-177 and IPAP/EG/90-Si-1614, and 3 human isolates, MHOM/EG/89/Si-15, MHOM/EG/89/Si-17 and MHOM/EG/90/Si-18, from MFO personnel, all from the Sinai study area. The electrophoretic methods of Kreutzer & Christensen (1980) were used for the separation of glucose-6-phosphate dehydrogenase (G6PD; EC.1.1.1.49), glucose phosphate isomerase (GPI; EC.5.3.1.9), mannose phosphate isomerase (= phosphomannose isomerase; MPI; EC.5.3.1.8), phosphoglucomutase (PGM; EC.2.7.5.1) and 6-phosphogluconic dehydrogenase (6PGD; EC.1.1.1.44), and that of Harris & Hopkinson (1976) for the visualization of malate dehydrogenase (MDH; EC.1.1.1.37).

Phlebotomus papatasi was the only man-biting sandfly species identified in the 1594 specimens dissected. The promastigote infection rate for this sample, representing the period July–September 1989, was approximately 0.7% (11/1594). Promastigotes from 2 sandfly infections were successfully cultured in NNN medium, and produced lesions after inoculation into the footpads of BALB/c mice. Serotyping of both the sandfly isolates revealed that they were of the same serotype as the 2 recent MFO isolates (EF serotype A/B) and also several other human Leishmania isolates from the same area in the Sinai (Mansour et al., 1987). In addition, each isolate yielded isoenzyme profiles identical to those of the L. major reference strain used for comparison (Figure).
It is reasonable to consider the 10 unidentified promastigote infections in the wild-caught sandflies as *L. major*. This consideration is based upon their position in the sandfly midgut and the tendency of *P. papatasi* to feed preferentially upon warm-blooded hosts. Their position in the midgut, and not in the hind-gut, excludes the possibility of their being trypanosomatids of reptiles, upon which *P. papatasi* only occasionally feeds (KILICK-KENDRICK, 1979). Because *Leishmania* isolated from the naturally-infected sandflies in this study was indistinguishable from the parasite isolated from cutaneous lesions in man, *P. papatasi* is considered to be a grade 3 vector (KILICK-KENDRICK & WARD, 1981) in Egypt. WAHBA et al. (1990) had previously characterized a single isolate of *L. major* from Egyptian collections of *P. papatasi*. However, this is the first report of repeated, concurrent isolations of this parasite from both sandflies and humans in Egypt.

**Acknowledgements**

We acknowledge, with appreciation, confirmatory isoenzyme electrophoresis of human and sandfly isolates by Dr. Richard Kreutzer of the Youngstown State University, Youngstown, Ohio, USA. Grateful thanks are extended to members of the multinational force and observers for their participation.

**References**


Received 20 September 1990; revised 7 February 1991; accepted for publication 8 February 1991.
Cutaneous leishmaniasis (CL) was repeatedly diagnosed among members of the Multinational Forces and Observers (MFO) in the north Sinai. Leishmania parasites were isolated from both sandflies and suspected human cases of (CL). This study revealed that Phlebotomus papatasi is the only man-biting species in this area. Leishmania parasites produced lesions in the footpads of BALB/c mice, their excretory factor (EF) gave the serotype A,B, and each isolate yielded isoenzyme profiles identical to those of the L. major reference strain. This is the first report of repeated, concurrent isolation of L. major from both sandflies and humans in Egypt.