



One step Rapid Immuno-chromatographic test for the detection of antibodies specific to Leishmaniasis in human serum, plasma or whole blood.

Leishmaniasis is a vector-borne disease caused by protozoan parasites that belong to the genus *Leishmania* and is transmitted by the bite of certain species of sand fly. Leishmaniasis is one of the most important diseases of humans. This parasitic disease can be caused by many species of *Leishmania*, most of which are zoonotic. In humans, different species of the parasite are associated with different forms of the disease. Many *Leishmania* spp. cause skin ulcers and nodules. A few of these organisms can also affect the mucous membranes, and may cause disfiguring lesions of the nose. Other species damage the internal organs and cause human visceral leishmaniasis, a life-threatening condition. Leishmaniasis affected 88 countries, of which 72 are classed as developing countries, including 13 of the least developed countries.

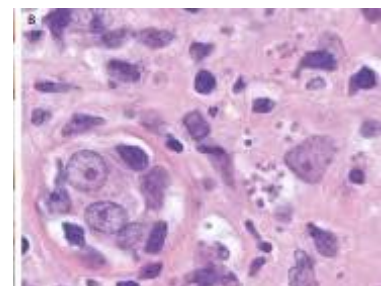
Sensit Leishmaniasis antibody Rapid Test is a lateral flow immunoassay for the qualitative detection of antibodies (IgG) against *Leishmania donovani*, (*Leishmania* causative Protozoan) in Whole blood, plasma or serum. This test is intended to be used as a screening test and as an aid in the diagnosis of *Leishmania* infection. Any reactive specimen with the Sensit Leishmaniasis antibody Rapid Test must be confirmed using supplemental assay.

Assay Overview and Usage

Organism detected	IgG to <i>L. donovani</i>
Sample type	Wholeblood, Serum, plasma
Shelf Life	23 months
Storage	2-30 °C
Capture Antigens	rK39 antigen
Sensitivity	90% vs ELISA
Specificity	98% vs Elisa
Packing	10T, 30T, Bulk, Uncut sheets
Ref. No	S009-04

Rapid Testing

- Quick results available in just a few minutes.
- Require no instrumentation.
- Easy to use.
- No capital expenditure.
- Do not require refrigerated storage.



Why Leishmaniasis Rapid Test

- Early identification of infected individuals, to allow intervention strategies in a clinically relevant time frame.
- Its analytical sensitivity is comparable to FDA-approved laboratory methods



Positive



Negative



I nvalid

Test Interpretation



Blood Collection
test device

Serum Separation

Add sample to

Manufactured &Marketed By

ubio Biotechnology Systems Pvt. Ltd.

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