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merican visceral leishmaniasis (AVL) is a serious parasitic disease distributed in Argentina, Bolivia, Brazil, Colombia, Costa Rica, El Salvador, Guatemala, Honduras, Mexico, Nicaragua and Venezuela (Arias et al., 1996). In Venezuela its magnitude is not well known, mainly because clinical features are shared with other endemic diseases and diagnosis is difficult. It is accepted that some fatal AVL cases might be erroneously attributed to malaria, schistosomiasis, Chagas disease or toxoplasmosis. Finding Leishmania spp. parasites in bone marrow, which is the unequivocal evidence for etiologic diagnosis, is not practiced in Venezuelan rural areas. On the other hand, this method has a low sensitivity because of a low parasitic burden. Serological methods (e.g. IFAT), which are only available in specialized diagnostic centers in Venezuela, show high sensitivity but not always high specificity (Camargo & Rebonato 1969; Badaró et al., 1983). The use of a dipstick based on the recombinant rK39 antigen of a sequence of 298 aminoacids and an improved serological procedure has showed to be practical and reliable for the diagnosis of visceral leishmaniasis in the New World (Badaró et al., 1996) and in the Old World (Sundar et al., 1998). Here we present results of the first application of the rK39 dipstick for differential diagnosis between AVL and other sympatric parasitic diseases in Venezuela.

**MATERIALS AND METHODS**

Serology of patients kept at the Instituto de Medicina Tropical of the Universidad Central de Venezuela with diagnosis of AVL (Badaró et al., 1998) were tested using a recombinant rK39 antigen (InBios International, Inc). The diagnosis of AVL was based on the positivity of at least two of the following tests: bone marrow examination, immunofluorescence antibody test (IFAT), counterimmuno-electrophoresis (CIEP) and Western blotting, as previously described (Delgado et al., 1998). Plasmodium vivax and P. falciparum had been detected by microscopical observation. Schistosomiasis had been demonstrated by Schistosoma mansoni eggs in feces and by the circumovular test (Oliver-Gonzalez, 1954). Complement fixation test (Machado Guerreiro’s test) and IFAT were used for Chagas Disease diagnosis (Almeida & Fife, 1976). Toxoplasmosis was determined by ELISA (Voller et al., 1976) and indirect agglutination test (Jacobs & Lundle, 1957).

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Sera from 10 patients with localized cutaneous leishmaniasis (LCL) and 10 with mucocutaneous leishmaniasis (MCL) were also included in the study to discard cross reaction to species-specific Leishmania spp. antibodies. The rK39 dipstick test was assayed three times for each sample. Simultaneous IFAT using leishmanial antigen was always done, allowing us to determine the comparative specificity and sensitivity of both tests. The methodology used for the dipstick test followed standard recommendations: one drop of serum was applied to the absorbent pad at the bottom of the strip. After air drying, three drops of test strip buffer (protein A-colloidal gold conjugate) were added. All dipstick results were recorded at five minutes after applying the buffer solution.

RESULTS

The appearance of a red upper band (control) indicating the presence of IgG, demonstrated proper test functioning. The appearance of a lower red band revealed the presence of IgG anti-rK39, indicating a positive test for AVL. When comparing results of proved AVL positive sera against sera positive to other diseases, a 100% of specificity was demonstrated. Conversely, IFAT showed cross-reaction for Chagas disease, LCL and MCL (titer = 1:256). Dipstick sensitivity in sera positive for AVL was 87.8% (36/41) while IFAT sensitivity was 100%. When results were analyzed as function of time of sera storage it was detected that sera that were negative to dipstick test (but positive to IFAT) had been kept in the laboratory at –70°C for more than 10 years (1974-1983). In fact, no significant difference was detected when comparing results obtained with sera stored during 1974-1978 (24-20 years before), 1979-1983 (19-15 years) and 1984-1988 (10-15 years of storage) (Fisher exact test), but significant difference was obtained comparing results of each of these groups of samples with those from sera collected between 1989-1993 and 1994-1998 (Fisher exact test: p > 0.01).

DISCUSSION

A 60-second dipstick for a rapid diagnosis of visceral leishmaniasis has being recently used (Sundar et al., 1998; Jelinek et al., 1999). Results on the sensitivity of tests using this antigen have led, to a certain extent, to different interpretations. Some authors report a very high sensitivity in active AVL, being positive in acute infections and subclinical progressing to VL infections, while asymptomatic and subclinical self-healing patients would be negative (Badaró et al., 1996). Sundar et al. (1998) report 100% sensitivity in all patients with positive spleen aspirate, that would indicate active infection, while they obtained four positive results in patients with negative spleen aspirate direct smears that were also interpreted as subclinical no-self healing infection. On the other hand, false negatives (28.6%) were detected among 14 samples by Jelinek et al. (1999). These authors state that the reason for the false negative reactions remains unclear, so they do not consider that the rK39 dipstick as a reliable and conclusive test for VL diagnosis. We observed loss of reactivity in sera kept more than nine year, which may be due to protein degradation in long-term stored samples (Margulies, 1996).

The presence of specific antibodies is detected by the recombinant product rK39 with a repetitive epitope closely conserved between Leishmania chagasi and L. donovani, which is part of a large kinesin related protein expressed predominantly by amastigotes (Reed, 1990; Burns et al., 1993). A 100% specificity of this protein had been previously reported in Brazil using sera from patients with tropical diseases other than AVL (Badaró et al., 1996). A good specificity has been also reported in the Old World (Singh et al., 1995; Sundar et al., 1998; Jelinek et al., 1999). In Venezuela AVL, Chagas Disease, malaria, schistosomiasis and toxoplasmosis overlap in some areas. These diseases share clinical symptoms and signs, therefore the clinical-epidemiological diagnosis is of scarce value, even in endemic areas. The major need for a rural medical doctor is to handle a reliable test for an early diagnosis of AVL in order to apply the opportune specific treatment. The high specificity of the rK39 antigen to differentiate from sympatric parasitic diseases other than AVL in Venezuela, allows us to recommend the routine use of this simple and low cost dipstick test for a rapid diagnosis of suspected AVL cases in rural area primary care centers, since it may indeed avoid deaths. Suspected AVL cases which would result negative, should be referred to the closest hospital to be confirmed by other serological or parasitological techniques.

ACKNOWLEDGEMENTS

This work received financial support from CDCH-UCV, Project 09-10-3682-99 and PCEE-VEN96-002-006 (Commitment Venezuelan Government-World Bank).

REFERENCES


