Immunodiagnosis of chronic Chagas’ disease using the Tc 46 and Tc 58 antigens

Valéria R.A. Pereira, Mineo Nakazawa, Veridiana C. Furtado, Frederico G.C. Abath and Yara M. Gomes

Abstract The polypeptides of 46 and 58kDa were recognized in different T. cruzi strains (Y, WSL and Colombiana) by serum of all chagasic patients studied. These polypeptides were isolated from T. cruzi Y strain and used in ELISA. The sensitivity and specificity were 97.6% [CI 95%: 86-100%] and 100% [CI 95%: 89.3-100%], respectively when Tc 46 was used. When Tc 58 was used the sensitivity and specificity were 100% [CI 95%: 89.6-100%] and 90.2% [CI 95%: 75.9-96.8%], respectively.


In the acute phase of Chagas’ disease, when parasitemia is high, usually diagnosis can be easily made using conventional parasitological methods (thin blood smear, thick blood smear, fresh blood examination). During the chronic phase, due to the low parasitemia, diagnosis is usually performed by immunological methods in conjunction with clinical epidemiological evidence. Indirect immunofluorescence, indirect hemagglutination and ELISA are widely used in Latin America for individual diagnosis and for screening donated blood, as well as in epidemiological studies.

The antigens commonly used in serodiagnosis of Chagas’ disease are fixed epimastigote forms or complex mixtures of proteins and glycoconjugates extracted from whole parasites which lead to false positive results due to cross-reactions with other parasites. In the last years, several research groups have tried to develop new diagnostic tests employing purified antigens, recombinant proteins or synthetic peptides in order to achieve a more reproducible and specific diagnosis of Chagas’ disease. Thus, the use of chemically defined antigens and the molecular and
immunological characterization of the relevant parasite antigens are of utmost importance. To be useful, these antigens should be common to all known *T. cruzi* strains and should not cross react with sera from patients carrying other diseases.10

In the present communication we report on two polypeptides (46 and 58kDa), present in the insoluble antigenic fraction of different *T. cruzi* strains that were recognized by all chagasic patients studied. In addition, we evaluate their usefulness for the ELISA diagnosis of human chronic Chagas' disease.

Serum samples were collected from 42 patients with chronic Chagas' disease: cardiac form (n = 14), asymptomatic form (n = 21), digestive form (n = 2) and mixed form (n = 5). Diagnosis of chronic Chagas' disease was based upon the collective analysis of a set of elements: clinical manifestations and electrophysiographic alterations, compatible epidemiological history, and a positive reaction by two different serological tests. In addition, serum of patients with other parasitic diseases, such as: visceral (n = 6) and cutaneous (n = 9) leishmaniasis, toxoplasmosis (n = 9), schistosomiasis (7), filariasis (5) and malaria (5), as well as sera from healthy individuals (n = 5) from non endemic areas, were also analysed. Sera were stored at -20°C until use.

Epimastigote forms of three different strains of *T. cruzi* (Y = Type I, WSL = Type II and Colombiana = Type III) were used. These forms were obtained by cultures in RPMI 1640 medium containing 10% calf serum. Parasites were grown at 26°C by serial passages at every 7th, 14th and 21st day for Y, Colombiana and WSL strains, respectively when they were harvested and washed 6 times (2500 x g, 15min) with 150mM PBS pH 7.2 containing proteases inhibitors (1mM PMSF and 1mM EDTA). Parasites were resuspended in 10ml of deionized water containing the same concentration of proteases inhibitors, sonicated 5 times (30 sec pulses) on ice (Ultrasonic Homogenizer, Model CP 501 — Cole Parmer) and centrifuged at 10^5 x g for 1h. Protein concentration of the pellet (insoluble antigens) was determined according to Lowry.

Twelve µg of the insoluble antigen fraction, corresponding to each *T. cruzi* strain were separated in polyacrylamide gels (10%) according to Laemmli using a mini-gel system (Hoefer Scientific Instruments, San Francisco, USA). Proteins were transferred from the gel to a nitrocellulose sheet (pore size = 0.45mm) with a semi-dry blotter at 30mA during 60 min. The sheets containing each of the antigens from the 3 strains were cut into vertical strips, which were treated with a solution of 5% defatted milk (Molico, Nestlé, São Paulo, Brazil), 0.05% Tween 20 in PBS, pH 7.2 (PBS-Tw) for 2h at room temperature with constant shaking. After washing the strips were incubated with serum samples from patients with Chagas' disease, diluted 1:100 in PBS-Tw. Individual sera of normal subjects from non-endemic area as well as of subjects with other parasitic diseases were used as controls. The bound antibodies were detected with peroxidase-conjugated goat anti-human IgG (γ chain specific). The immune complexes were revealed by addition of H_2O_2 and 4-chloro-1-naphthol. Colour development was stopped by washing in distilled water.

The reactivity patterns showed by patients with Chagas' disease are presented in Figure 1. At least 23 polypeptides with molecular weights of 14 to 145kDa could be identified. The intensity of the reaction to different antigens was variable but similar patterns of reactivity were observed with almost all analysed sera (Figure 1). No antigenic pattern characteristic of clinical forms was found. No reaction was observed when the sheet was incubated with normal human sera. The comparative analysis of the reactivity patterns of sera from Chagas' disease and other parasitic diseases indicates that at least 15 polypeptides specifically reacted with sera from chagasic patients. Among them, the 46 and 58kDa polypeptides reacted with sera from 42 Chagas' disease patients. The 16, 20, 29, 32, 33, 34, 35, 40, 44, 48, 74, 88, and 145kDa polypeptides were recognized by sera from chagasic patients with variable frequency (data not shown). The 46 and 58kDa polypeptides were isolated by preparative electrophoresis according to Gomes et al4 and will be referred to as Tc 46 and Tc 58 antigens.

The Tc 46 and Tc 58 antigens isolated from Y strain were diluted in 0.05M Na_2CO_3 buffer, pH 9.6, and coated onto flat-bottomed microtitre plates. The optimum concentration (500ng/well) was determined by checkerboard titration using positive and negative sera. The specific IgG bound to these antigens was determined using a horseradish peroxidase-conjugated goat anti-human IgG. The development of the reaction was
carried out with orthophenylenediamine-OPD and H$_2$O$_2$ and blocked with 2.5M H$_2$SO$_4$. The optical density (OD) was measured at 490nm. Samples were recorded as positive according to the cut-off value, calculated as the mean OD of the negative controls plus two standard deviations (SD). The sensitivity and specificity were calculated according to Camargo\textsuperscript{1}. The confidence interval was calculated at the level of 95\% (CI 95\%).

ELISA results are shown in Table 1. When Tc 46 was used as antigen 41 out of 42 chagasic patients were considered positive for Chagas’ disease. No cross-reaction was observed with other parasitic diseases. Sensitivity and specificity were 97.6\% [CI 95\%: 86-100\%] and 100\% [CI 95\%: 89.3-100\%], respectively. When Tc 58 was used the sensitivity and specificity were 100\% [CI 95\%: 89.6-100\%] and 90.2\% [CI 95\%: 75.9-96.8\%], respectively (Table 1). These results are currently being further evaluated by testing a larger sample and combining both antigens. In addition, experiments are being carried out to clone the encoding genes.

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<thead>
<tr>
<th>Table 1 - Reactivity of Tc 46 and Tc 58 antigens in ELISA with different human sera.</th>
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<tbody>
<tr>
<td>Sera</td>
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<tr>
<td>------</td>
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<tr>
<td>Chagasic (n = 42)</td>
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<tr>
<td>Negative* (n = 5)</td>
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<tr>
<td>Visceral leishmaniasis (n = 6)</td>
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<td>Cutaneous leishmaniasis (n = 9)</td>
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<td>Schistosomiasis (n = 7)</td>
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<td>Filariasis (n = 5)</td>
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* Negative sera were from healthy individuals from non endemic areas.

Figure 1 - Representative Western blot of the antigenic pattern recognized by sera of patients with different clinical forms of chronic Chagas’ disease. A: cardiac form; B: asymptomatic form; C: digestive form and D: mixed form. Y = Ystrain; W = WSL strain; C = Colombiana strain. Molecular weight markers (M) on left indicate the numbers 211, 119, 98, 80.6, 64.4, 44.6 and 38.9 kDa. Arrow on right indicate Tc46 and Tc 58 antigens.

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REFERENCES


