Immunoregulatory mechanisms in Chagas disease: modulation of apoptosis in T-cell mediated immune responses

Ana Thereza Chaves1, Juliana de Assis Silva Gomes Estanislau1,2,3, Jacqueline Araújo Fiuza1, Andréa Teixeira Carvalho4, Karine Silvestre Ferreira2, Rafaelle Christine Gomes Fares1, Pedro Henrique Gazzinelli Guimarães5, Elaine Maria de Souza Fagundes6, Maria José Morato1, Ricardo Toshio Fujiwara5, Manoel Otávio da Costa Rocha3 and Rodrigo Correa-Oliveira1,7,8

Abstract

Background: Chronic Chagas disease presents different clinical manifestations ranging from asymptomatic (namely indeterminate) to severe cardiac and/or digestive. Previous results have shown that the immune response plays an important role, although not all mechanisms are understood. Immunoregulatory mechanisms such as apoptosis are important for the control of Chagas disease, possibly affecting the morbidity in chronic clinical forms. Apoptosis has been suggested to be an important mechanism of cellular response during T. cruzi infection. We aimed to further understand the putative role of apoptosis in Chagas disease and its relation to the clinical forms of the disease.

Methods: Apoptosis of lymphocytes, under antigenic stimuli (soluble T. cruzi antigens – TcAg) where compared to that of non-stimulated cells. Apoptosis was evaluated using the expression of annexin and caspase 3* by T cells and the percentage of cells positive evaluated by flow cytometry. In addition activation and T cell markers were used for the identification of CD4+ and CD8+ subpopulations. The presence of intracellular and plasma cytokines were also evaluated. Analysis of the activation status of the peripheral blood cells showed that patients with Chagas disease presented higher levels of activation determined by the expression of activation markers, after TcAg stimulation. PCR array were used to evaluate the contribution of this mechanism in specific cell populations from patients with different clinical forms of human Chagas disease.

Results: Our results showed a reduced proliferative response associated a high expression of T CD4+CD62L- cells in CARD patients when compared with IND group and NI individuals. We also observed that both groups of patients presented a significant increase of CD4+ and CD8+ T cell subsets undergoing apoptosis after in vitro stimulation with T. cruzi antigens. In CARD patients, both CD4+ and CD8+ T cells expressing TNF-a were highly susceptible to undergo apoptosis after in vitro stimulation. Interestingly, the in vitro TcAg stimulation increased considerably the expression of cell death TNF/TNFR superfamily and Caspase family receptors genes in CARD patients.

Conclusions: Taken together, our results suggest that apoptosis may be an important mechanism for the control of morbidity in T. cruzi infection by modulating the expression of apoptosis genes, the cytokine environment and/or killing of effector cells.

Keywords: Chagas disease, Immunoregulation, Apoptosis, TNF/TNFR superfamily, Caspase family, T lymphocytes, Trypanosoma cruzi

*Correspondence: correaa@cpqrr.fiocruz.br
1Laboratório de Imunologia Celular e Molecular, Centro de Pesquisas René Rachou, Fiocruz, Belo Horizonte, Brazil
2Instituto Nacional de Ciência e Tecnologia em Doenças Tropicais – INCyT, Minas Gerais, Brazil
© 2016 Chaves et al. Open Access This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The Creative Commons Public Domain Dedication waiver (http://creativecommons.org/publicdomain/zero/1.0/) applies to the data made available in this article, unless otherwise stated.
Background
Chagas disease, a neglected disease caused by *Trypanosoma cruzi*, remains a serious public health problem and affects about 10 million people in Latin America [1]. Chronic cardiomypathy represents the most important and severe manifestation of human Chagas disease, eventually affecting approximately 20–30% of individuals. The majority of the chronically affected individuals present the indeterminate (IND) form of the disease, with an apparent absence of morbidity [2, 3]. Epidemiological studies in endemic areas have shown that 2–5% of patients will evolve each year from the indeterminate to the cardiac clinical form of the disease [4].

Although the pathophysiology of Chagas disease is not completely understood, it is widely accepted that the involvement of the immune response is critical in determining the disease outcome [5–8]. While the balance between inflammatory and anti-inflammatory cytokines produced by circulating cells in patients with IND form leans towards an anti-inflammatory profile, patients with chagasic cardiomyopathy seem to display a predominately inflammatory pattern [6, 9, 10]. The type of immune response induced in these individuals seems to be critical for the maintenance of a “healthy” balance between the parasite and the host [11]. In fact, several studies have demonstrated that immunoregulatory mechanisms are important for the control of infection, possibly affecting disease morbidity in chronic clinical forms [11, 12] such as T-cell suppression, polyclonal lymphocyte activation, and regulatory cytokines [13, 14]. Infection with *T. cruzi* leads to polyclonal lymphocyte activation [15], which, by itself, promotes T-cell apoptosis [16, 17]. In addition, antigens released by *T. cruzi*, such as trans-sialidase and HSP70, induce lymphocyte apoptosis [18, 19]. Therefore, it is possible that the parasite exploits host cell apoptosis to evade the immune response. Evidences also indicate that apoptosis plays a role in the resolution of inflammation [20].

In the present work, we evaluated the contribution of apoptosis in specific T-cell populations on the development/maintenance of different clinical manifestations of human Chagas disease. Our findings demonstrated that in vitro stimulation with *T. cruzi* antigens induce lymphocyte apoptosis by continued cell activation, modulation of the expression of apoptosis genes and cytokine secretion profile. These findings may contribute to the regulation of immune response during human Chagas disease.

Methods
Study population
The patients that agreed to participate in this study were identified and selected from those being attended at the Referral Outpatient Center for Chagas Disease, which is located at the Clinics Hospital of the Federal University of Minas Gerais (UFMG), Brazil, under the medical care of one of us (MOCR). These patients were enrolled in a prospective cohort study initiated 20 years ago as previously described [13]. Patients infected with *T. cruzi* were grouped as indeterminate (IND) or with cardiomypathy (CARD). The IND group included 15 asymptomatic individuals with age ranging from 24 to 66 years (mean of 39.6 ± 10.3), with no significant alterations in electrocardiography, chest X-ray and echocardiogram. The CARD group included 15 patients with age ranging from 23 to 69 years (mean of 48 ± 12.52) presenting dilated cardiomypathy, characterized by the echocardiographic finding of a dilated left ventricle with impaired ventricular systolic function. Left ventricular ejection fraction (LVEF) and left ventricular diastolic diameter (LVDD) were used as clinical parameters of the ventricular function for Chagas disease patients, where LVEF <55% and LVDD/ body surface area ≥31 mm were used to define Chagas disease dilated cardiomypathy [3]. None of the patients had undergone chemotherapeutic treatment, nor been previously treated for *T. cruzi* infection. Healthy individuals with age ranging from 29 to 55 years [mean of 42.6 ± 8.8], from a non-endemic area for Chagas disease with negative serological tests for the infection were included in the control group (non-infected NI).

Ethics statement
This study was carried out in full accordance with all international and Brazilian accepted guidelines and was approved by the Ethics Committee of the René Rachou Research Center – FIOCRUZ (14/2006 CEPSH-IRR) and UFMG protocol COEP-ETHIC 001/79). All enrolled patients gave written informed consent prior to the inclusion in the study.

*Trypanosoma cruzi* soluble antigen preparations
The CL strain of *T. cruzi* was used for antigenic preparation as described elsewhere [21]. After preparation, the protein concentration was determined, aliquoted and stored at -70 °C prior use.

Short-term in vitro whole blood cultures with *T. cruzi* antigens
Whole blood samples (final concentration of 1 x 10^6 cells/mL) were treated with staurosporine (Sigma, St. Louis, MO, USA) (4 µM), soluble *T. cruzi* antigens (TcAg) (25 µg/mL) or untreated (stimulated with medium alone – RPMI 1640 supplemented with 1.6% L-glutamine, 3% antibiotic-antimycotic, 5% of AB Rh-positive heat-inactivated normal human serum), and incubated for approximately 24 h at 37 °C in 5% CO₂. Following incubation, the cultures were treated with 220 µL of EDTA at 20 mM and maintained at room temperature for 15 min
prior immunophenotypic staining for apoptosis assay, cell surface markers, and intracellular cytokine analysis.

Cell preparation and proliferation assay
PBMC from Chagas patients and healthy individuals were isolated by FicollPaque density gradient centrifugation (LSM; Organon Teknika, Charleston, S.C.) as previously described (Gomes, 2003). The cells were washed in RPMI 1640 medium and cultured in flat-bottom 96-well plates (Nunc Brand Products). Proliferative responses were evaluated by incubating 2.5 × 10^5 cells/well for TcAg (25 μg/mL) or 1.5 × 10^5 cells/well for mitogen stimulation (PHA, 10 μg/mL), respectively, in a final volume of 200 μL of complete RPMI-1640. Incubation was carried out in a humidified 5% CO2 incubator at 37 °C for 3 days for PHA-stimulated cultures and 6 days for antigen-stimulated cultures. Cells were pulsed for the last 6 h of incubation with 1 μCi of [3H] methyl thymidine (PerkinElmer LAS, Shelton, CT, USA), and harvested onto glass fiber filters (Printed Filtermat A, Walenc, Finland). Radioactive incorporation was determined by liquid scintillation spectrometry (MicroBeta JET, PerkinElmer Inc., USA).

Analysis of apoptosis profile
Short-term in vitro whole blood cultures were washed with 6 mL of FACS buffer (PBS supplemented with 0.5% bovine serum albumin-BSA and 0.1% sodium azide) by centrifugation at 600 xg for 7 min at room temperature and resuspended in 5 mL of FACS buffer.

For the annexin V analysis, aliquots of 150 μL were transferred to polystyrene tubes and incubated for 30 min at room temperature (RT) with 2 μL of allophycocyanin (APC) – labeled anti-CD4 (RPA-T4) (BD Pharmingen) or anti-CD8 (RPA-T8) (BD Pharmingen) monoclonal antibodies. Following incubation, the red blood cells were lysed by the addition of 3 mL of FACS lysing solution (Becton Dickinson, CA, USA) for 10 min, and cells washed with 2 mL of PBS by centrifugation at 600 xg for 7 min at room temperature. The cells were resuspended in annexin V binding buffer (0.1 M Hepes/NaOH (pH 7.4) 1.4 M NaCl, 25 mM CaCl2 - Biosciences, San Jose, CA), for working solution (1X), then incubated for 15 min at RT (25 °C) in the dark with 5 μL annexin V-PE and 5 μL 7AAD. The reaction was stopped by the addition of 100 μL of 1X binding buffer for each tube.

For caspase-3 analysis, aliquots of 150 μL of whole blood cultures were transferred to polystyrene tubes and incubated for 30 min at room temperature with 2 μL of fluorescein isothiocyanate (FITC)-labeled anti-CD45 (2D1) (BD Biosciences) and 2 μL of peridinin chlorophyll-a protein (PerCP)-labeled anti-CD14 (M5E2) (BD Biosciences) monoclonal antibodies. The tubes were incubated in the dark for 10 min at RT. The cells were permeabilized in saponin buffer (0.5%) (Sigma) for 15 min at RT in the dark. Finally, the cells were incubated with PE-conjugated rabbit anti-active caspase-3 mAb (C92-605) (BD Pharmingen) using 20 μL/1×10^6 cells for 60 min at RT in the dark. Phenotypic analyses were performed by flow cytometry using a Becton Dickinson FACSCalibur flow cytometer. Analysis was performed on 7 × 10^4 lymphocytes (gated according to their forward and side scatter properties. The sample acquisition and data analysis were performed using CellQuest software (BD Biosciences, USA).

Analysis of cell surface markers and intracytoplasmic cytokines
Cultured cells were washed twice in PBS containing 1% BSA and stained with monoclonal antibodies specific for cell-surface markers. Antibodies to CD4 (RPA-T4), CD8 (RPA-T8) and CD62L (DREG-56) (all from BD Pharmingen) were used. The cells were then fixed in formaldehyde (4%) and permeabilized in saponin buffer (0.5%) (Sigma, USA) for 15 min. Finally, the cells were incubated with anti-TNF-α (PE) (L293) (BD Biosciences) washed and resuspended in FACS buffer prior acquisition in flow cytometer.

Phenotypic analyses were performed by flow cytometry using a Becton Dickinson FACSCalibur flow cytometer, collecting data on 7 × 10^4 lymphocytes gated according to their forward and side scatter properties. The sample acquisition and data analysis were performed using CellQuest software (BD Biosciences, USA).

Detection of plasmatic cytokine levels by Cytometric Bead Array (CBA)
A cytometric beads array (CBA) immunoassay kit (BD Biosciences, USA) was used to measure cytokine levels (IFN-γ, TNF-α, IL-2 and IL-10) in plasma as described in previous studies [6]. The data were acquired in a Becton Dickinson FACSCalibur flow cytometer and analyzed using BD CBA software (BD Biosciences, USA). The results were expressed by mean intensity of fluorescence (MIF).

Apoptotic pathways triggered by T. cruzi infection in different clinical forms of Chagas disease
In order to determine putative apoptotic pathways triggered by T. cruzi infection, PBMC from infected patients IND (n = 2) and CARD (n = 2) were incubated only with culture medium- culture non-stimulated; or in the presence of antigen- culture stimulated with TcAg at a final concentration of 25 μg/mL. After 18 h of incubation at 37 °C and 5% CO2 atmosphere, cells were recovered and washed with PBS. Subsequently, cells were submitted to a total RNA extraction protocol using NucleoSpin* RNA II kit (Macherey-Nagel, Germany). The total RNA was quantified according to standard procedures using spectrophotometer
(Thermo Scientific, USA) and evaluated in agarose gel to confirm its integrity [22]. The cDNAs were obtained with Superscript II kit (Invitrogen, USA) using 120 ng of total RNA, according to manufacturer’s instructions. Afterwards, a RT-PCR reaction was performed using previously established constitutive human primers to confirm the cDNA synthesis.

The apoptotic transcripts were evaluated using a Human Apoptosis RT² Profiler PCR Array kit (SABiosciences, USA) in a qPCR machine (7500, Applied Biosystem, USA), according to manufacturer’s instructions. Thirty one transcripts involved with pro-apoptotic activity were evaluated. Out of thirty one genes, twenty belong to TNF and TNF receptor superfamily, cell death domains and inductors of apoptosis; and eleven to the caspasases family, which are also involved with pro-apoptotic activity. The qPCR data were analyzed by PCR Array Data Analysis Web Portal (SABiosciences, USA), and the results were expressed using the method of $2^{-\Delta\DeltaCT}$.

**Statistical analyses**

Statistical analyses were conducted using the R 2.15.0 software. Initially, the Anderson-Darling test was applied to verify whether the obtained data represent a normal distribution. Statistical comparative analyses were performed using the non-parametric: Mann–Whitney test to compare two groups (NI x IND or NI x CARD or IND x CARD); Kruskal-Wallis test to compare three groups (NI x IND x CARD) and, together with the Bonferroni correction (significance level, 0.05/3 = 0.0167). All tests were performed considering a significance level of 5% ($\alpha = 0.05$).

**Results**

**Cellular proliferative response is decreased in CARD patients**

The proliferative immune response was analyzed in PBMC from IND and CARD patients as well as NI individuals stimulated with soluble T. cruzi antigens (TcAg) by the quantification of $\delta$-thymidine incorporation. We observed that PBMC proliferative response of the IND group had a significantly higher ($p < 0.05$) level of cellular proliferation after stimulation with TcAg when compared with CARD and NI groups (Fig. 1). No significant difference was observed in the proliferative response induced by PHA (data not shown).

**Increased downregulation of CD62L in CD4+ T cells is related to CARD patients**

We also evaluated the loss of expression ($\Delta$CD62L) in circulating CD4+ and CD8+ T cells, before and after in vitro stimulation with TcAg. The analysis showed significant increase on the percentage of CD4+CD62L$^{-}$ cells in CARD patients ($p < 0.05$) in ex vivo analysis as well as non-stimulated cultures when compared with NI and IND groups (Fig. 2a). Moreover, after TcAg in vitro stimulation, a higher percentage of CD4+CD62L$^{-}$ cells in CARD patients ($p < 0.05$) it was also observed when compared to NI individuals (Fig. 2a). Ex vivo expression of CD62L$^{-}$ by CD8$^{+}$ cells and after in vitro culture in the presence or not of TcAg did not show any statistically significant differences between the study groups (Fig. 2b).

**Lymphocytes from IND and CARD patient presented elevated frequency of apoptosis after in vitro TcAg stimulation**

The low proliferative response and high percentage of CD4+CD62L$^{-}$ T cells observed in CARD patients might be related to regulatory mechanisms involved in the control of the immune response induced by T. cruzi. In this context, apoptosis has been previously demonstrated as play a role in experimental model of acute T. cruzi infection [23–26] and also in human heart tissues [27]. In order to investigate if apoptosis might be a factor related to the lower proliferative response and if it is caused by high T cell activation in the CARD patients, the frequency of annexin V$^{+}$ and active caspase 3$^{+}$ was analyzed as markers of apoptosis.

Our data show that after in vitro stimulation with TcAg the lymphocytes from CARD displayed high frequency of annexin V$^{+}$ when compared with IND and NI groups (Fig. 3a). The higher frequency of annexin V$^{+}$ in CD4+ T cells was observed in IND and CARD patients as compared to NI (Fig. 3b). On the other hand, lower frequency of annexin$^{+}$ in CD8$^{+}$ T cells were observed in
IND and NI groups as compared with CARD group (Fig. 3c). A similar profile was observed for the cultures in the presence of Staurosporin-STP, a positive control for apoptosis.

The frequency of total lymphocytes and CD4+ or CD8+ T cells subsets from IND and CARD patients exhibited similar apoptosis profile as compared with NI individuals when active caspase 3" was analyzed (Fig. 3a and b, respectively).

Activated Lymphocytes' TNF-α form CARD patients were susceptible to apoptosis

A second hypothesis to explain the induction of programmed cell death in lymphocytes from chagasic patients involves the cytokine environment. It is known that TNF-α is a major cytokine implicated in the apoptosis pathway. Thus, we have further evaluated serum levels and intracytoplasmic expression of this cytokine ex vivo and following incubation with TcAg. We observed that plasma from
CARD patients presented a higher expression of TNF-α than IND and NI groups ($p < 0.05$). Also, after ToAg stimulation, chagasic patients (IND and CARD) presented a higher frequency of TNF-α by total lymphocytes (Fig. 4b), and CD4$^+$ T or CD8$^+$ T cells when compared with NI groups (Fig. 4c and d, respectively). Analysis of serum levels of IFN-γ and IL-10 were also performed. IFN-γ was elevated in sera from cardiac patients and IL-10 in indeterminate patients as previously shown [28]. In this study we observed that IL-2 was also elevated in sera from cardiac patients (Additional file 1). These results did not show any association with apoptosis.

To determine whether TNF-α expression by total lymphocytes and CD4$^+$ or CD8$^+$ T cells subsets is associated with the frequency of apoptosis, we performed a correlative analysis between the frequency of apoptosis and TNF-α expression. We observed a positive and significant correlation between high frequency of annexin$^+$ in CD4$^+$ T cells and high frequency of TNF-α in CD4$^+$ T cells both in IND and CARD groups. Moreover, a significant positive correlation was found, between frequency of annexin$^+$CD8$^+$ T cells and TNF-α$^+$CD8$^+$ T cells in CARD group (Fig. 4d). Together, these results suggested that both CD4$^+$ and CD8$^+$ T cells from CARD patients

---

**Fig. 4** Analyses of cytokine levels and their association with cardiac morbidity expressed by the clinical classification. **(a)** The analysis of plasma cytokine levels was performed as described in material and methods. The groups evaluated were: NI ($n = 15$, white box), IND ($n = 15$, light gray box), and CARD ($n = 15$, dark gray box). The results were expressed by mean intensity of fluorescence (MF). Plasma TNF-α levels in NI, IND, and CARD groups and their association with cardiac morbidity. **(b, c)** Analysis of percentage of CD4$^+$TNFα$^+$ and CD4$^+$TNFα$^+$ T cells in the peripheral blood from chagasic patients, in ex vivo context and after in vitro stimulation with T. cruzi antigens, from patients with distinct clinical forms of Chagas disease. Significant differences ($P$-value $< 0.05$) in the charts are identified by connecting lines and the symbol ‘*’ for comparisons between the groups. **(d)** Correlation analysis between percentage of annexin$^+$ CD4$^+$ and annexin$^+$ CD8$^+$ T cells and TNF-α intracytoplasmic in the IND ($n = 15$) and CARD ($n = 15$) groups. Mann–Whitney test was used for comparison and the results were expressed as the median. Correlation analysis were done using the Spearman correlation coefficient, and the results were considered significant with a $p$-value $< 0.05$. Significant differences ($P$-value) are indicated in each graph together with the $r$ values.
expressing TNF-α were highly susceptible to undergo apoptosis upon in vitro stimulation.

Cell apoptosis in CARD patients might be induced by TNF receptors superfamily and/or caspase family pathway

Once we showed that CARD patients presented higher frequency of apoptotic cells, the next step was to identify the possible apoptotic pathway used to activate T apoptosis in T cells from patients with this clinical form, before and after in vitro stimulation with TcAg antigen. Initially, 31 apoptosis-related genes from TNF/TNFFR superfamily and Caspase family were grouped and categorized according to their respective families and function. Analysis of the others 53 genes of human apoptosis pathway were also performed and these results did not show any significant differences in all groups (data not shown).

In the absence of in vitro TcAg stimulation, when we evaluated the TNF/TNFFR superfamily genes from CARD group was determined. The results showed that cell death receptors (FADD, TRADD, TNFRSF10A (TNF receptor superfamily member 10A), TNFRSF10B, TNFRSF11B, TNFRSF21 and TNFRSF25) were up-regulated when compared with IND group (Fig. 5a). Moreover, other pro-apoptotic genes (FAS, FASL, TNFSF8, TNFRSF9, TRAIL, TRAF2) were also up-regulated in CARD patients when compared with individuals from the IND group (Fig. 5a). When also evaluated the caspase family genes, the results showed that caspase 1 (CASP1), caspase 2 (CASP2), caspase 3 (CASP3), caspase 4 (CASP4), caspase 5 (CASP5), caspase 7 (CASP7), caspase 9 (CASP9), caspase 10 (CASP10) and caspase 14 (CASP14) genes, were up-regulated in CARD group in comparison with IND patients (Fig. 5b).

Interestingly, the in vitro TcAg stimulation increased considerably the expression of cell death TNF/TNFFR superfamily as well as caspase family receptors genes (FADD, TRADD, TNFRSF10A, TNFRSF10B, TNFRSF11B, TNFRSF21, TNFRSF25, TNFRSF10, FAS, FASLG, TNFRSF9, TNFSF8, TRAF2, TRAF4) (Fig. 5c and d, respectively). Furthermore, high expression of genes such as TNFRSF1A, CD27, LTA, TRF3, TRF4 and CASP3 were observed after PBMCs cultures from CARD when compared to IND patients (Fig. 5c and d, respectively). Moreover, the CD70 gene was downregulated in CARD patients when compared to IND patients (Fig. 5c).

Discussion

In the present work, we have shown that apoptosis is associated with reduced proliferative response, a high expression of CD4+CD62L- cells, an increase TNF-α intracellular production and expression of genes of cell death TNF/TNFFR superfamily and caspase family in CARD patients.

In this context, some studies have been shown that CARD patients presented low proliferation of T cells when compared with healthy and non-chagasic cardiomyopathy donors [29–32]. Although, the mechanisms of proliferative dysfunction in Chagas disease need further investigation, studies suggest that they could be related to the decrease of co-stimulatory molecules expression, receptor cytokine starvation or expression of inhibitory receptors as PD-1 [32, 33].

Some studies suggest that activated cells are susceptible to apoptosis, which may represent a mechanism of immunoregulation [34, 35]. To assess the activation status of T cell subsets, we evaluated whether CD62L is downregulated in these cell subpopulations. Our data showed that CARD patients presented a significantly higher percentage of TCD4+CD62L- cells suggesting that a putative involvement of this cell type in the exacerbation of the immune response to the parasite and, consequently, on the development of myocarditis by cell death induced by activation. Dos Santos et al. [36] showed that the majority of TCD4+ and TCD8 lymphocytes in the inflammatory foci from the heart of chagasic patients did not express or slightly expressed CD62L and T CD8+ cells are the majority of the activated cells in the tissue when compared with CD4+ T cells. The inflammatory process occurring in Chagas' disease mainly consists of CD8 T lymphocytes, CD4 T lymphocytes and macrophages. The extent of inflammatory reaction and the tissue damage caused may contribute to loss of myocardial cells, and to the heart failure that is observed on more severe cases of chronic Chagas' disease. Tostes Jr et al. [37] showed that myocardial cell loss by apoptosis and fibrosis contributes to heart failure in the chronic phase of Chagas' disease. In fact, apoptosis has been considered a cause of heart failure in other diseases such as myocardial infarction and heart hypertrophy [38, 39], as well as a mechanism involved in the control of the immune response in experimental models [26]. On the other hand, heart failure by itself may induce apoptosis [40].

In this work, high expression of annexin by lymphocytes from CARD group was observed when compared with IND and NI groups, as well as an increase on the expression of annexin by CD4+ T cells and caspase by both CD4+ and CD8+ T cells by chagasic patients when compared with NI group after in vitro stimulation with T. cruzi antigens. In Chagas' disease, the occurrence of apoptosis in T lymphocytes was observed after antigen stimulation in experimental models [26] although its significance in terms of clinical form or outcome of the disease was not clear. Apoptosis-like death has been reported in amastigote nests and trypomastigotes forms from T. cruzi, and this mechanism has been associated with control of parasite burden regulated by the parasite itself or by the host, parasite evasion of the host's immune
response and clonal selection [41-44]. Together, these findings suggest that although the lymphocytes from CARD patients presented lower proliferative response upon antigenic recall, lymphocytes from IND and CARD patients presented a singular ability to undergo apoptosis that may reflect different regulatory mechanisms.

Rodrigues et al. [45] demonstrated a high percentage of lymphocyte apoptosis in patients with a severe cardiomyopathy, associating this event to activation of programmed death pathways, by Fas/Fas-L, or TNF-α receptors, leading to parasite escape, and consequently, to a continuous stimulation of the immune system. In this context, T cells apoptosis in experimental model leads to an increase of parasite growth [46]. Indeed, apoptosis has been suggested to be an important mechanism to control the immune response and heart damages [47].

In order to determine whether TNF-α contributes to T cell apoptosis, the plasmatic and intracellular production by T cell subsets was evaluated. Our data demonstrated that CARD group showed increased levels of circulating and intracytoplasmic TNF-α, and up-regulation of the TNF receptor gene superfamily. Lula et al. [48] have shown correlation among soluble ligands of TNF superfamily (TNF-α, TRAIL and FasL/CD95L) and functional disorders of the left ventricle in chronic chagasic patients with cardiomyopathy. These results are associated with ligand receptors associated with programmed cell death, suggesting that apoptotic mechanisms are involved with the development of miocardioopathy in Chagas disease.

Ferreira et al. [49] showed a correlation between high serum levels of TNF-α and the occurrence of severe Chagas cardiomyopathy. Also, it has been shown that
there is an inverted correlation between high levels of TNF-α with the lowest left ventricular ejection fraction seen in patients with chronic Chagas cardiomypathy [50]. Moreover, patients with heart failure are shown to have a significantly lower PBMC proliferative response but higher levels of apoptosis and Fas and Fas-L expression [45]. TNF-α may play a role on the high levels apoptosis and in the low proliferative response observed in patients with heart failure. TNF-α may contribute to the induction of apoptosis by the interaction with its receptor or by induction of Fas and Fas-L expression [51]. Our results suggest that the high levels of TNF-α detected in plasma from CARD group may contribute to the heart condition. Additionally, our data support the hypothesis that high levels of TNF-α and up-regulation of the TNF receptor gene superfamily lead to an increase in apoptosis, and consequently the exacerbation of the pathology.

Conclusions
Here, we showed that apoptosis is associated with low proliferative response, intense T cell activation, high TNF-α production and up-regulation of genes associated with TNF receptors superfamily and caspase family in CARD patients. These results suggest that apoptosis could interfere on the development and/or maintenance of the different clinical forms of Chagas disease. Assuming that the immunological regulation in the IND group, may control the development of Chagas cardiomypathy, the absence of this mechanism in the CARD group, may be one of the factors associated with sustained inflammation which would, consequently, lead to a higher morbidity in the latter group. The association of lymphocyte apoptosis, induced by the constant activation of the immunological system, with high levels of inflammatory cytokines and associated pathological events (fibrosis and apoptosis) may contribute with the development and progression of heart injuries in the chronic phase of the human Chagas disease.

Additional file

Additional file 1: Figure S1. Analyses of plasma cytokine levels expressed by the clinical classification. The analysis of plasma levels was performed as described in material and methods. The groups evaluated were: NI (n = 15, white box), IND (n = 15, light grey box), and CARD (n = 15, dark grey box). The results were expressed by mean intensity of fluorescence (MF). (A) Plasma TNF-α levels in NI, IND, and CARD groups. (B) Plasma IL-2 levels in NI, IND, and CARD groups. (C) Plasma IFN-γ levels in NI, IND, and CARD groups. (D) Plasma IL-10 levels in NI, IND, and CARD groups. Significant differences (P value < 0.05). Statistical comparative analysis were performed, in groups of two, between the NI, IND, and CARD groups, using the non-parametric Kruskal-Wallis test and Mann-Whitney U test, together with the Bonferroni correction (significance level 0.05/3 = 0.0167). The letters represent statistically significant differences (p < 0.05) between the groups: a = difference when compared to NI group; b = difference when compared to IND group; c = difference when compared to CARD group.

References


