Tissue Damage in Human Cutaneous Leishmaniasis: Correlations Between Inflammatory Cells and Molecule Expression

Maíra Garcia Saldanha¹, Carla Pagliari², Adriano Queiroz¹, Paulo Roberto Lima Machado³, Lucas Carvalho¹,², Phillip Scott⁴, Edgar M. Carvalho¹,³ and Sérgio Arruda¹,⁵

¹Instituto Gonçalo Moniz, Fundação Oswaldo Cruz (FIOCRUZ), Salvador, Brazil, ²Departamento de Patologia, Faculdade de Medicina, Universidade São Paulo, São Paulo, Brazil, ³Serviço de Imunologia, Complexo Hospitalar Universitário Professor Edgard Santos, Universidade Federal da Bahia, Salvador, Brazil, ⁴Department of Pathobiology, School of Veterinary Medicine, University of Pennsylvania, Philadelphia, PA, United States, ⁵Departamento de Ciências de Vida, Universidade Estadual da Bahia, Salvador, Brazil

Cutaneous leishmaniasis (CL) is caused by the bite of the infected sand fly, which inoculates parasites of Leishmania spp and triggers an immune response. An exacerbated cutaneous inflammatory response is crucial for controlling parasite burden but can also promote tissue damage. This study aimed to characterize the populations of natural killer (NK), CD57⁺, CD4⁺, and CD8⁺ T cells, CD20⁺ B cells, as well as CD68⁺ macrophages, in biopsies of ulcerated CL lesions, and quantify the production of perforin⁺, granzyme B⁺, interleukin 1 beta (IL-1β⁺) and Tumor Necrosis Factor (TNF-α⁺) cells. We then correlated these parameters with necrosis, inflammation and the number of amastigotes. CD4⁺ T cells were positively correlated to the extent of inflammation, B cells and IL-1β⁺ were associated with the extent of necrosis, CD68⁺ macrophages and perforin were correlated with the number of amastigotes, and CD57⁺ NK cells was correlated to CD68⁺ macrophages and amastigotes. In sum, the finding suggests that the production of cytotoxic granules and cytokines by inflammatory cells contributes to tissue damage in CL lesions.

Keywords: cutaneous leishmaniasis, inflammatory cells, necrosis, amastigotes, tissue damage

INTRODUCTION

Cutaneous leishmaniasis (CL) is the most common clinical form presented by individuals infected by protozoa of the genus Leishmania. After inoculation of Leishmania Viannia braziliensis in the skin by sandflies, a nodular lesion and an exuberant satellite lymphadenopathy is documented (Bomfim et al., 2007; Wind et al., 2014). The classical CL ulcers caused by L. braziliensis appear about 2–4 weeks after the presence of nodular lesions and are characterized by well-defined ulcer with raised borders. The evolution of CL is characterized by an exacerbated inflammatory response (Costa et al., 2018). In most infectious diseases, early treatment increases cure rates and decreases healing time; however, the introduction of therapy soon after infection in CL, i.e., prior to the appearance of ulcers, has been associated with a high rate of therapeutic failure (Machado et al., 2002; Unger et al., 2009; Khouri et al., 2014).
The main host defense mechanism against intracellular protozoa is the activation of macrophages by IFN-γ, mainly produced by CD4+ T cells (Santos et al., 2013). However as Leishmania are able to escape this killing mechanism, the persistence of the parasite and leishmanial antigens induce a marked inflammatory response that is associated with tissue damage and the development of skin ulcers (Santos et al., 2013). Several molecules have been associated with the pathology of L. braziliensis infection. Neutrophils are the cells that initially migrate after parasite inoculation, followed by macrophages (Novais et al., 2009; Conceição et al., 2016). The production of IFN-γ by NK cells may contribute to parasite killing, or may be cytotoxic, thereby contributing to this pathology (Muniz et al., 2016; Campos et al., 2017). Subsequently, the activation of CD4+ and CD8+ T cells is observed. T cell activation and the production of cytokines by these cells is determinant in the outcome of infection. An impairment in the host’s Th1 immune response results in diffuse CL, which is characterized by multiple nodular lesions consisting predominantly of macrophages with a high parasite burden (Silveira et al., 2004). Alternatively, a normal Th1 immune response induces an exacerbated inflammatory response, leading to the presence of ulcerating lesions in CL (Bacellar et al., 2002; Antonelli et al., 2005; Castro Gomes et al., 2017).

IL-1β and TNF-α are highly expressed in CL ulcers, and may be involved in the tissue damage arising from L. braziliensis infection (Cardoso et al., 2015; Novais et al., 2017). The production of IL-1β by peripheral blood mononuclear cells is associated with ulcer size (Santos et al., 2013). Treatment with pentoxifylline, a drug that decreases TNF-α production, in combination with meglumine antimoniate, is more effective than antimony alone. Combined therapy not only reduces healing time in patients with mucosal leishmaniasis (ML), but also cures ML patients refractory to antimony therapy alone (Cuba et al., 1984; Lessa et al., 2001; Machado et al., 2002). Further data has indicated the participation of monocytes, CD4+ and CD8+ T cells, in the pathogenesis of CL, due to increased frequencies of intermediate monocytes referred to as an inflammatory monocye subset. In CL and ML, macrophages present increased TLR expression, enhanced respiratory burst and produce higher levels of pro-inflammatory cytokines compared to cells from healthy subjects or individuals with subclinical L. braziliensis infection (Giudice et al., 2012; Carneiro et al., 2016; Muniz et al., 2016). With regard to the adaptive immune response, the size of CL ulcers is directly correlated with the frequencies of CD4+ T cells expressing IFN-γ and CD69, an early marker of T cell activation (Antonelli et al., 2005). More recently, the role of CD8+ T cells has been documented in the pathology of L. braziliensis in both mice and humans (Santos et al., 2013; Cardoso et al., 2015; Novais et al., 2018). Studies have shown that the killing of L. braziliensis-infected cells by CD8+ T cells leads to the increased production of IL-1β through the activation of the inflamasome, which may exacerbate tissue damage (Santos et al., 2013; Cardoso et al., 2015; Novais et al., 2018). However, most of these studies were conducted in mice or in human peripheral blood mononuclear cells, i.e., there is a paucity of data regarding the cells and molecules that participate in the pathogenesis of ulcer formation at the lesion site. Accordingly, the present study evaluated cell frequencies and quantified proinflammatory molecules in CL ulcer biopsies in an attempt to contribute to the knowledge surrounding the immunopathogenesis of CL lesions caused by L. braziliensis.

MATERIALS AND METHODS

Patients

This study included 22 biopsies from 22 CL patients obtained punch biopsy samples (4 mm) after anesthesia from the border of ulcer and immediately fixed in 10% formalin-buffered solution. CL diagnosis was done by to the presence of amastigotes in these biopsies, as well as the detection of L. braziliensis DNA by PCR. Patient demographic characteristics, illness duration and lesion size were documented. All patients were subsequently treated with meglumine antimoniate (Glucantime®, Sanofi Aventis, Gentilly, France) at a dose of 20 mg/Kg.

Immunohistochemistry

Deparaffinization and rehydration of 5-μm thick sections was performed using xylene and alcohol PA, followed by antigen retrieval with buffer pH 9.0 at 96°C for 20 min. Immunohistochemistry was conducted after blocking peroxidase activity with 3% hydrogen peroxide for 10 min, and protein activity with Protein Block Serum-Free (DAKO, California, USA) for 15 min. All slides were incubated at 4°C with the following antibodies: Monoclonal Mouse anti-CD57, clone Ab-1, Monoclonal Mouse anti-Perforin, clone 5B10 (ThermoFisher, Massachusetts, USA); Polyclonal Rabbit anti-TNF-α, clone orb129752 (Biorbyt, Cambridge, UK); Monoclonal Mouse anti-IL-1β, clone 3AC (Cell Signaling Technology, Massachusetts, USA); Polyclonal Rabbit anti-Granzyme B, clone EP230, Monoclonal Mouse anti-CD4, clone EP204, Monoclonal Mouse anti-CD8, clone C8/144B (Cell Marque, California, USA); Monoclonal Mouse anti-CD68, clone PG M1, Monoclonal Mouse anti-CD20, clone L26 (Dako, California, USA); Polyclonal Rabbit anti-Leishmania (in house) (Schubach et al., 2001). A Mouse and Rabbit Peroxidase Kit HorseRadish KP500 (Diagnostic BioSystems, Pleasanton, USA) was used to perform reactions according to manufacturer recommendations. All slides were counterstained with Harris hematoxylin, dehydrated and then mounted using Permount (Thermo Fisher Scientific, Massachusetts, USA) on glass coverslips.

Quantitative Analysis

A Nikon 90i Eclipse microscope (Nikon Corporation, Tokyo, Japan) was coupled to a DS-Fi1 digital camera system (Nikon Corporation, Tokyo, Japan). All slides were photographed (five randomized fields from each section) at 40× and visualized with Nis-Elements software (v. 3.1). In each field, the number of positive cells was counted using the semiautomatic quantification feature of ImageJ software v. 1.48 (National Institutes of Health, Maryland, USA). Positive cells were identified via the amplified molecules that reacted with the chromogenic substrate DAB (eBioscience, California, USA). For all reactions, a preselected pattern was used as a positive control and sections that had
not been incubated with the primary antibody were used as negative controls.

**Morphometry of Inflammation and Necrosis Areas**

The histological sections stained with hematoxylin and eosin were scanned by an optical microscope (Olympus BX51, Tokyo, Japan). The total extension of these sections as well as the areas of inflammatory infiltrate and necrosis was measured by Image J 1.48v (National Institutes of Health). The total length of the biopsy fragment and the sum of the areas of inflammation and necrosis are shown in mm$^2$. The percentage (%) of inflammation and necrosis in the biopsies were calculated by dividing total extension of inflammation and necrosis in mm$^2$ by the total extension of the biopsy fragment multiplied by 100.

**Heatmap and Correlation Matrices**

Language R through the RStudio interface (v1.2.5019) was used for plotting heatmaps and correlation matrices between the parameters and markers observed. The functions corrplot and corrgram was used to track the heatmaps and correlation matrices, respectively, using Pearson’s statistical method. Positive correlations are displayed in blue and negative correlations in red color. Color intensity and the size of the circle are proportional to the correlation coefficients.

**RESULTS**

A total of 22 patients were enrolled with mean age of 40 years and 16 were male, corresponding to 73% of the study sample. Fifteen patients presented lymphadenopathy and lesions were more frequent in the lower limbs (86%) than the upper limbs (14%). The average duration of ulcers was 43 ± 25 days at the time of patient examination, with an average lesion size of 376 ± 336 mm$^2$.

**Figure 1** illustrates the frequencies of cells expressing CD4$^+$, CD8$^+$, CD20$^+$, CD57$^+$, CD68$^+$, IL-1β$^+$, TNF-α$^+$, granzyme B$^+$ and perforin$^+$ in tissue biopsies from CL patients. CD8$^+$ cells were the most frequent, followed by CD68$^+$, CD57$^+$, and CD4$^+$ (Figure 1A). Regarding the molecules secreted by these cells, IL-1β$^+$ and TNF-α$^+$ were found to be the most highly expressed, mainly in individuals with lymphadenopathy, followed by perforin$^+$ and granzyme B$^+$ (Figure 1B).

**Figure 2** depicts correlations between the observed histopathological features and the frequencies of cells and molecules expressed by these cells, in addition to associated R values (Figure 2A). The inflammation seen in CL is characterized...
by the infiltration of mononuclear cells throughout the duration of the illness (Faria et al., 2009; Dantas et al., 2014). Increased infiltration is seen, ranging from mild in the early phase of disease, before the appearance of ulcers, which increases during ulcer development (Saldanha et al., 2017). Inflammation was found to be directly associated with the frequency of CD4+ T cells ($R = 0.40; P = 0.05$; Figure 2B).

Necrosis in CL ulcers are secondary to tissue damage due to the killing of macrophages infected with $L. braziliensis$, or the killing of cells expressing parasite antigens. However, the extent of necrosis in CL ulcers was reduced compared to the area of inflammation. No direct correlations were found between these two variables. However, IL-1$\beta$+ expression was strongly correlated with necrosis ($R = 0.79; P < 0.0001$; Figure 2C). Interestingly, the presence of B cells was also strongly correlated with necrosis ($R = 0.61; P < 0.0001$; Figure 2D).

Figures 2E,F shows the most frequent cells type and molecules associated with the presence of amastigotes in CL ulcers. As expected, the number of parasites was directly correlated with the presence of CD68$^+$ macrophages ($R = 0.63; P < 0.0001$), as well as with the frequency of cells expressing perforin ($R = 0.68; P < 0.0001$).

Furthermore, there was no correlation of CD57$^+$ NK cells with inflammation or necrosis, but these cells were positively correlated to CD68$^+$ macrophages ($R = 0.42; P < 0.05$) and amastigotes numbers ($R = 0.36; P < 0.05$; Figures 2G,H).

**DISCUSSION**

The histological analysis of ulcerated lesions from patients with CL indicated chronic inflammation, with a predominance of mononuclear cellular infiltrate (Faria et al., 2009; Dantas, 2012). In the present study sample, CD4$^+$ and CD8$^+$ lymphocytes were highly expressed in ulcerative lesion biopsies. Both of these lymphocyte populations have been associated with protection and pathology. CD4$^+$ T cells play a protective role through the production of IFN- $\gamma$, the main cytokine responsible for the activation of macrophages, which leads to intracellular parasite killing (Santos et al., 2013). Alternatively, CD8$^+$ T cells have been more associated with tissue damage due to cytotoxic activity (Glennie and Scott, 2016; Castro Gomes et al., 2017). Here, we found cells expressing CD8$^+$, CD4$^+$ and CD68$^+$ were the most frequent in tissue samples, and CD4$^+$ cells were also positively correlated with the extent of inflammation. A previous study in CL patients demonstrated a direct correlation between the frequency of peripheral blood CD4$^+$ T cells expressing IFN-$\gamma$, as well as TNF-$\alpha$, and lesion size (Antonelli et al., 2005). It is well known that the production of IFN-$\gamma$ by CD4$^+$ T cells activates monocytes/macrophages and enhances the production of pro-inflammatory cytokines, which may contribute to the inflammatory response (Bacellar et al., 2002; Faria et al., 2009; Santos et al., 2013; Cardoso et al., 2015). Nonetheless, we found a negative correlation between CD4$^+$ cells and CD68$^+$ macrophages, CD4$^+$ cells and the number of amastigotes and CD4$^+$ cells and necrosis that are against a significant role of these cells in the pathology of CL.

CD8$^+$ T cells were the most frequent cell type found in tissue biopsies from CL patients. Evidence indicates that these cells play an important role in the pathogenesis of $L. braziliensis$ infection (Dantas et al., 2013; Novais et al., 2013). However, despite the high frequency of CD8$^+$ T cells in the present tissue samples, we found no correlations between this cell population and the presence of amastigotes, nor inflammation or necrosis. This apparent discrepancy may be due to several factors. In fact, the production of granzyme and perforin molecules is more important than the frequency of CD8$^+$ T cells, since these are directly involved in the lysing of target. In the tissue, granzyme was positively associated with necrosis and slightly negatively associated with amastigotes and inflammation, while perforin was strongly positively associated with amastigotes and slightly with necrosis, yet slightly negatively associated with inflammation. A large percentage of cells were found to express CD57$^+$, a marker associated with highly cytotoxic CD8$^+$ or NK cells (Kared et al., 2016). Moreover, CD57$^+$ cells were the only cell type whose frequency was directly correlated with macrophages and amastigotes, suggesting the participation of CD57$^+$ cells in killing infected macrophages.

Since *Leishmania* is an intracellular parasite, little emphasis has been placed on the presence of B cells and antibodies in the pathogenesis of the disease. However, B cells and plasma cells are documented in the tissue of CL patients, and anti-leishmanial antibodies are highly produced in severe of *Leishmania* spp. infection, such as the visceral form and diffuse cutaneous lesions (Bomfim et al., 2007; Rodriguez-Pinto et al., 2014). Nonetheless, the role of B cells and antibodies in the pathology of CL has not been well-studied. Opsonization by antibodies may enhance parasite uptake by macrophages. In addition, form immune complex deposition can contribute to tissue pathology. Here we found a strong direct correlation between CD20$^+$ and IL-1$\beta$+ expression, as well as between CD20$^+$ and the extent of necrosis, which highlights the need for future study to evaluate the role of B cells and antibodies in the pathogenesis of CL.

Necrotic cell death may contribute to granuloma formation, inflammation and tissue damage. The role of IL-1$\beta$+ and TNF-$\alpha^+$+ has been well studied in in CL mouse models, and high levels of both cytokines are produced by peripheral blood mononuclear cells in CL patients (Antonelli et al., 2005; Campos et al., 2014; Passos et al., 2015). IL-1$\beta^+$ can be produced by a variety of cells, including inflammatory monocytes, B lymphocytes, and epithelial and NK cells (Carvalho et al., 2012; Novais et al., 2017). Recently, IL-1$\beta^+$, through inflammasome activation, as well as cytotoxic CD8$^+$ T cells have both been associated with tissue damage in $L. braziliensis$ infection (Novais et al., 2017). Here IL-1$\beta^+$ and TNF-$\alpha^+$ were both more frequent in patients with lymphadenopathy, and were positively correlated with granzyme expression, and necrosis was also strongly directly correlated with IL-1$\beta^+$, which suggests that this cytokines plays a key role in the pathology of CL.

Gene expression analysis in biopsied human CL ulcerative lesions revealed high expression of IL-1$\beta^+$, TNF-$\alpha^+$, granzyme and perforin, as well as other molecules (Novais et al., 2017). The perforin and granzymes produced by NK cells and T lymphocytes promote the apoptosis of infected cells (Trapani and Smyth, 2006).
FIGURE 2 | Correlations between histopathological parameters and cellular markers in CL biopsies. (A) Clustered heatmap of Pearson correlation coefficients of the inflammatory cells, inflammation and necrosis; (B–H) Pearson correlation of the cells, areas of inflammation and necrosis, amastigotes and NK cells; (B) CD4⁺ T cells vs. inflammation; (C) IL-1β⁺ cells vs. necrosis; (D) CD20⁺ B cells vs. necrosis; (E) CD68⁺ macrophages vs. amastigotes; (F) perforin⁺ cells vs. amastigotes; (G) CD68⁺ macrophages vs. CD57⁺ NK cells; (H) amastigotes vs. CD57⁺ NK cells.
2002). Perforin forms pores in target cells and also participates in the upregulation of CD8 (Zhou, 2010), while granzymes lyse intracellular pathogens (Arias et al., 2014). Our in situ findings in human CL lesions suggest that the associations between inflammatory cells and the intense production of cytokines and cytotoxic products may contribute to necrosis and potentially participate in the formation of CL ulcers.

**CONCLUSION**

This study attempted to establish correlations between histopathological CL parameters in order to elucidate the pathogenesis of this disease. We suggest that, in an effort to control parasites, infected macrophages are likely to remain in an activated state due to the production of proinflammatory cytokines by CD4+ T cells, in particular IL-1B+, thusly potentially contributing to tissue necrosis. In addition, the exacerbated production of cytotoxic granules containing perforin by CD8+ T cells and NK cells can aggravate tissue destruction, thereby intensifying skin ulcer formation.

**DATA AVAILABILITY STATEMENT**

All datasets generated for this study are included in the article/supplementary material.

**ETHICS STATEMENT**

The studies involving human participants were reviewed and approved by Institutional Review Board (Instituto Gonçalo Moniz—IGM-FIOCRUZ, no. 533.032/2014). The patients/participants provided their written informed consent to participate in this study.

**AUTHOR CONTRIBUTIONS**

MS, SA, EC, and CP planned research and coordinated the study. MS, CP, and LC performed the experiments. PM and AQ provided the clinical samples. MS, EC, PS, and SA analyzed the data. MS and SA wrote the manuscript with contributions from EC, PS, and CP. All authors read and approved the submitted version of the manuscript.

**FUNDING**

This work was supported by National Institutes of Health (grant AI088650) and Conselho Nacional de Pesquisa Instituto Nacional de Ciência e Tecnologia—Doenças Tropicais (project no. 573839/2008-5). The funders had no role in study design, data collection or analysis, the decision to publish, or manuscript preparation.

**ACKNOWLEDGMENTS**

We thank Dr. Mírian Nacagami Sotto and her staff for their assistance and for providing services at the Communicable Diseases Pathology Laboratory of the São Paulo University Medical School. We also thank Cristiano Sampaio Franco for his assistance in the preparation of the manuscript. The authors are grateful to Andris K. Walter for English revision and copyediting services.

**REFERENCES**

Antonelli, L. R. V., Dutra, W. O., Almeida, R. P., Bacellar, O., Carvalho, E. M., and Gollob, K. J. (2005). Activated inflammatory T cells correlate with lesion size in human cutaneous leishmaniasis. *Immunol. Lett.* 101, 226–230. doi: 10.1016/j.imlet.2005.06.004

Arias, M. A., Jiménez de Bagües, M. P., Aguiló, N., Menao, S., Hervás-Stubbs, A., de Martino, A., et al. (2014). Elucidating sources and roles of granzymes A and B during bacterial infection and sepsis. *Cell Rep.* 8, 420–429. doi: 10.1016/j.celrep.2014.06.012

Bacellar, O., Lessa, H., Schriefer, A., Machado, P., Ribeiro de Jesus, A., Dutra, W. O., et al. (2002). Up-regulation of Th1-type responses in mucosal leishmaniasis patients. *Infect. Immun.* 70, 6734–6740. doi: 10.1128/IAI.70.12.6734-67 40.2002

Bromfin, G., Andrade, B. B., Santos, S., Clarência, J., Barral-Netto, M., and Barral, A. (2007). Cellular analysis of cutaneous leishmaniasis lymphenopathy: Insights into the early phases of human disease. *Am. J. Trop. Med. Hyg.* 77, 854–859. doi: 10.4269/ajtmh.2007.77.854

Campos, T. M., Costa, R., Passos, S., and Carvalho, L. P. (2017). Cytotoxic activity in cutaneous leishmaniasis. *Mem. Inst. Oswaldo Cruz* 112, 733–740. doi: 10.1590/0074-02760170109

Campos, T. M., Passos, S. T., Novais, F. O., Beiting, D. P., Costa, R. S., Queiroz, A., et al. (2014). Matrix metalloproteinase 9 production by monocytes is enhanced by TNF and participates in the pathology of human cutaneous leishmaniasis. *PLoS Negl. Trop. Dis.* 8:e3282. doi: 10.1371/journal.pntd.0003282

Cardoso, T. M., Machado, Â., Costa, D. L., Carvalho, L. P., Queiroz, A., Machado, P., et al. (2015). Protective and pathological functions of CD8+ T cells in Leishmania braziliensis infection. *Infect. Immun.* 83, 898–906. doi: 10.1128/IAI.02404-14

Carneiro, P. P., Conceição, J., Macedo, M., Magalhães, V., Carvalho, E. M., and Bacellar, O. (2016). The role of nitric oxide and reactive oxygen species in the killing of Leishmania braziliensis by monocytes from patients with cutaneous leishmaniasis. *PLoS ONE* 11:e0148084. doi: 10.1371/journal.pone.0148084

Carvalho, L. P., Passos, S., Schriefer, A., and Carvalho, E. M. (2012). Protective and pathologic immune responses in human tegumentary leishmaniasis. *Front. Immunol.* 3:301. doi: 10.3389/fimmu.2012.00301

Castro Gomes, C. M., Sousa, M. G. T., Menezes, J. P. B., Batista, M. C., Lima, A. C. S., Belda, W., et al. (2017). Immunopathological characterization of human cutaneous leishmaniasis lesions caused by Leishmania (Viannia) spp. in Amazonian Brazil. *Parasitol. Res.* 116, 1423–1431. doi: 10.1007/s00436-017-5403-4

Conceição, J., Davis, R., Carneiro, P. P., Giudice, A., Muniz, A. C., Wilson, M. E., et al. (2016). Characterization of neutrophil function in human cutaneous leishmaniasis caused by leishmania braziliensis. *PLoS Negl. Trop. Dis.* 10:e0004715. doi: 10.1371/journal.pntd.0004715

Costa, R. S., Carvalho, L. P., Campos, T. M., Magalhães, A. S., Passos, S. T., Schriefer, A., et al. (2018). Early cutaneous leishmaniasis patients infected with leishmania braziliensis express increased inflammatory responses after antimony therapy. *J. Infect. Dis.* 217, 840–850. doi: 10.1093/infdis/jix627

Cuba, C. C., Llanos-Cuentas, E. A., Barreto, A. C., Magalhães, A. V., Lago, E. L., Reed, S. G., et al. (1984). Human mucocutaneous leishmaniasis in Três Braços, Bahia - Brazil: an area of *Leishmania braziliensis braziliensis* transmission. I. Laboratory diagnosis. *Rev. Soc. Bras. Med. Trop.* 17, 161–167. doi: 10.1590/S0037-86821984000040002

---

Frontiers in Cellular and Infection Microbiology | www.frontiersin.org 7 July 2020 | Volume 10 | Article 355
Dantas, M. L. (2012). Aspectos Comparativos da Resposta Inflamatória em Lesões de Leishmaniose Cutânea Localizada e Disseminada (Salvador: Centro de Pesquisas Gonçalo Moniz).

Dantas, M. L., de Oliveira, J. M. G. C., Carvalho, L., Passos, S. T., Queiroz, A., Guimarães, L. H., et al. (2014). Comparative analysis of the tissue inflammatory response in human cutaneous and disseminated leishmaniasis. Mem. Inst. Oswaldo Cruz 109, 202–209. doi: 10.1590/0074-0276130312

Dantas, M. L., Oliveira, J. C., De Carvalho, L., Passos, S. T., Queiroz, A., Machado, P., et al. (2013). Major Article CD8+ T cells in situ in different clinical forms of human cutaneous leishmaniasis. Rev. Soc. Bras. Med. Trop. 46, 728–734. doi: 10.1590/0037-8682-0174-2013

Faria, D. R., Souza, P. E. A., Durães, F. V., Carvalho, E. M., Gollob, K. J., Machado, P. R., et al. (2009). Recruitment of CD8+ T cells expressing granzyme A is associated with lesion progression in human cutaneous leishmaniasis. Parasite Immunol. 31, 432–439. doi: 10.1111/j.1365-3024.2009.01125.x

Giudice, A., Vendrane, C., Bezerra, C., Carvalho, L. P., Delavechia, T., Carvalho, E. M., et al. (2012). Macrophages participate in host protection and the disease pathology associated with Leishmania braziliensis infection. BMC Infect. Dis. 12:75. doi: 10.1186/1471-2334-12-75

Glennie, N. D., and Scott, P. (2016). Memory T cells in cutaneous leishmaniasis. Cell. Immunol. 309, 50–54. doi: 10.1016/j.cellimm.2016.07.010

Kared, H., Martelli, S., Ng, T. P., Pender, S. L. F., and Larbi, A. (2016). CD57 in Memory T cells in situ in different clinical forms of human cutaneous leishmaniasis. J. Immunol. 197, 141:122–123. doi: 10.4049/jimmunol.1601097

Silveira, F. T., Lainson, R., and Corbett, C. E. P. (2004). Clinical and immunopathological spectrum of American cutaneous leishmaniasis with special reference to the disease in Amazonian Brazil - A review. Mem. Inst. Oswaldo Cruz 99, 239–251. doi: 10.1590/S0074-0262200300001

Trapani, J. A., and Smyth, M. J. (2002). Functional significance of the perforin/granzyme cell death pathway. Nat. Rev. Immunol. 2, 735–747. doi: 10.1038/nri911

Wind, B. S., Guimarães, L. H., and Machado, P. R. L. (2014). Erythema exsudativum multiforme after a Leishmania skin test. Am. J. Trop. Med. Hyg. 68, 574–579. doi: 10.4269/ajtmh.2013.13-0166

Zhou, F. (2010). Perforin: more than just a pore-forming protein. J. Immunol. 183, 8088–8098. doi: 10.4049/jimmunol.0803720

Novais, F. O., Wong, A. C., Villareal, D. O., Beiting, D. P., and Scott, P. (2018). CD8+ T Cells Lack local signals to produce IFN-γ in the skin during leishmaniasis infection. J. Immunol. 191, 1006196. doi: 10.4049/jimmunol.1701597

Passos, S., Carvalho, L. P., Costa, R. S., Campos, T. M., Novais, F. O., Magalhães, A., et al. (2015). Intermediate monocytes contribute to pathologic immune response in leishmaniasis braziliensis infections. J. Infect. Dis. 211, 274–282. doi: 10.1093/infdis/jiu439

Rodriguez-Pinto, D., Saravia, N. G., and McMahon-Pratt, D. (2014). CD4 T cell activation by B cells in human Leishmania (Viannia) infection. BMC Infect. Dis. 14:108. doi: 10.1186/1471-2334-14-108

Saldanha, M. G., Queiroz, A., Machado, P. R. L., de Carvalho, L. P., Scott, P., de Carvalho Filho, E. M., et al. (2017). Characterization of the histopathologic features in patients in the early and late phases of cutaneous leishmaniasis. Am. J. Trop. Med. Hyg. 16:0559. doi: 10.4269/ajtmh.16-0559

Santos, C., da S., Boaventura, V., Ribeiro Cardoso, C., Tavares, N., Lordelo, M. J., et al. (2013). CD8+ Granzyme B-mediated tissue injury vs. CD4+ IFN-γ mediated parasite killing in human cutaneous leishmaniasis. J. Invest. Dermatol. 133, 1533–1540. doi: 10.1038/jid.2013.4

Unger, A., Angeles, L., Neal, S. O., Sam, S. W., Park, J., Or, P., et al. (2009). Association of treatment of American cutaneous leishmaniasis prior to ulcer development with high rate of failure in northeastern Brazil. Am. J. Trop. Med. Hyg. 80, 574–579. doi: 10.4269/ajtmh.2009.80.574

Wind, B. S., Guimarães, L. H., and Machado, P. R. L. (2014). Erythema exsudativum multiforme after a Leishmania skin test. Am. J. Trop. Med. Hyg. 90, 587–588. doi: 10.4269/ajtmh.13-0166

Zhou, F. (2010). Perforin: more than just a pore-forming protein. Int. Rev. Immunol. 29, 56–76. doi: 10.1080/088318093349644

Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2020 Saldanha, Pagliari, Queiroz, Machado, Carvalho, Scott, Carvalho and Arruda. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.