T lymphocytes and macrophages in the intestinal tissues of dogs infected with Leishmania infantum

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Abstract

This study was about a semi-quantitative analysis of T lymphocytes (CD4+ and CD8+, FoxP3+ regulatory T cells), and macrophages in the gut wall of dogs naturally infected with Leishmania infantum. Thirteen dogs were divided into three groups: group 1 (G1, n=5), dogs with canine visceral leishmaniasis (CVL) and infected with L. infantum amastigotes in the intestine; group 2 (G2, n=5), dogs with CVL but without intestinal amastigotes; and group 3 (G3, n=3), uninfected dogs (control group). There was no significant difference (p ≥ 0.05) on CD4+ and T reg cell numbers among the groups, whereas the levels of CD8+ T cells and macrophages were significantly higher in dogs from G1 group than in G2 and G3 (p ≤ 0.05), especially in intestinal segments with high parasite burden. Parasite burden correlated positively with levels of CD8+ T cells and macrophages (p ≤ 0.05), but was inversely correlated to levels of CD4+ T lymphocytes and FoxP3+ Treg cells. In conclusion, in the intestine of dogs with CVL, the increase of CD8+ T cells and macrophages population associated with high parasite burdens, but no changes of CD4+ T cells and FoxP3+ T reg cells suggest a possible immunoregulation by the parasite not dependent on T reg cells.

Keywords: T lymphocytes, macrophages, canine visceral leishmaniasis, intestinal tract.

Resumo

Este estudo foi uma análise semi-quantitativa de linfócitos T (CD4+, CD8+ e regulatórios - T reg FoxP3+) e macrófagos na parede intestinal de cães naturalmente infectados com Leishmania infantum. Treze cães foram divididos em três grupos: grupo 1 (G1, n=5) continha cães com leishmaniose visceral canina (LVC) e com amastigotos intestinais; grupo 2 (G2, n=5) continha cães com LVC, mas sem amastigotos intestinais e o grupo 3 (G3, n=3) continha cães não infectados (grupo controle). Verificou-se que não houve diferença significativa (p ≤ 0.05) no número de células CD4+ e de Treg entre os grupos, mas o número de células T CD8+ e macrófagos foi significativamente superior nos cães do grupo G1 em relação ao G2 e ao G3 (p ≤ 0.05), especialmente nos segmentos intestinais com altas cargas parasitárias. As altas cargas parasitárias correlacionaram positivamente com os números de CD8+ e macrófagos (p ≤ 0.05), mas negativamente com as células CD4+ e Treg. Em conclusão, no intestino dos cães com LVC, o aumento das populações de células T CD8+ e de macrófagos associado a altas cargas parasitárias, mas nenhuma alteração de células T CD4+ e células Treg FoxP3+ sugerem uma possível imunorregulação pelo parasita não dependente de células Treg.

Palavras-chave: Linfócito T, macrófagos, leishmaniose visceral canina, trato intestinal.
Introduction

Visceral leishmaniasis (VL) is a chronic parasitic disease caused by the protozoa *Leishmania infantum* (Kinetoplastida: Trypanosomatidae), which infects macrophages in a wide variety of mammals, including humans and dogs (LUCA et al., 1999; NIETO et al., 1999).

VL is transmitted to vertebrates by the bite of infected female phlebotomine sandflies, of which *Lutzomyia longipalpis* (Diptera: Psychodidae) is the main vector of the parasite in the Americas (GALATI et al., 1997; WHO, 2010; WHEELER et al., 2011).

The infection affects the liver, spleen, lymph nodes, and bone marrow and also other organs in the gastrointestinal, central nervous, genital, and urinary systems (BLAVIER et al., 2001). An inflammatory reaction in the gastrointestinal (GI) tract associated with the presence of intramacrophagic *Leishmania* amastigotes has been reported in experimentally infected dogs (KEENAN et al., 1984) and in naturally infected dogs (FERRER et al., 1991; PINTO et al., 2011; TOPLU & AYDOGAN, 2011; SILVA et al., 2016). Although uncommon, occurrences of erosive and ulcerative colitis and hemorrhagic diarrhea have been described in dogs in association with severe clinical signs of disease (GONZÁLEZ et al., 1990; FERRER et al., 1991). However, the prevalence of *Leishmania* parasitism detected in the colonic mucosa through colonoscopy, but without colitis, was surprisingly high in dogs with symptomatic leishmaniasis (ADAMAMA-MORAITO et al., 2007).

*Leishmania* has a strong tropism for macrophages, but they can also infect dendritic cells and other non-phagocytic mammalian cells (NADERER & MCCONVILLE, 2008). It is well established that phagocytic cells can participate in parasite control via processes that depend on free radicals secreted during the respiratory burst pathway activation in neutrophils and macrophages (RODRIGUES et al., 2007).

Macrophages are basically classified as M1 or M2 macrophages and their heterogeneity derives from the distinct effects of cytokines expressed by Th1 (T helper 1) or Th2 (T helper 2) cells during their differentiation. M1 macrophages after stimulation with LPS (lipopolysaccharide) are activated to produce NO (nitric oxide) from arginine, which controls the proliferation of intracellular parasites. M2 macrophages after the same stimulus increase the secretion of cytokines IL-4, IL-5 and IL-10, (DELFS et al., 2001; OHTA et al., 2004). In rats infected with *L. infantum*, CD8+ T cells isolated expressed IFN-γ and TNF, improving the cellular cytotoxic activity against *Leishmania* spp., suggesting that CD8+ T-cells displayed a Tc1 pattern of differentiation, which was important for the control of the parasite (TSAGOZIS et al., 2003). For Ohta et al. (2004) there is increasing evidence that Th1/Th2 and Tc1/Tc2 mediated cytokine imbalance has been of pathogenetic importance in various diseases, such as allergic and autoimmune diseases.

Figueiredo et al. (2014) found no correlation between the clinical signs of canine visceral leishmaniasis (CVL) and pathological, immunological, and parasitological findings in dogs with *L. infantum* amastigotes in GI tissues. However, different segments of the GI tract of infected dogs showed different immunological and parasitological responses. Specifically, increased expression of CD4, FoxP3, IL-10, (DELFS et al., 2001; OHTA et al., 2004). The effector function of CD8+ T-cells alone were unable to induce cure in BALB/c mice. The effector function of CD8+ T-cells probably depends on IFN-γ production (MÜLLER et al., 1991; HERATH et al., 2003).

CD8+ T cells, similar to CD4+ T cells, are classified into two subsets of cytolytic effector cells with distinct cytokine patterns, termed cytotoxic T lymphocytes 1 (Tc1) and cytotoxic T 2 (Tc2). Tc1 cells secrete a Th1-type cytokine pattern, including IL-2 and IFN-γ, whereas Tc2 cells produce Th2-like cytokines, including IL-4, IL-5 and IL-10, (DELFS et al., 2001; OHTA et al., 2004). In rats infected with *L. infantum*, CD8+ T cells isolated expressed IFN-γ and TNF, improving the cellular cytotoxic activity against *Leishmania* spp., suggesting that CD8+ T-cells displayed a Tc1 pattern of differentiation, which was important for the control of the parasite (TSAGOZIS et al., 2003). For Ohta et al. (2004) there is increasing evidence that Th1/Th2 and Tc1/Tc2 mediated cytokine imbalance has been of pathogenetic importance in various diseases, such as allergic and autoimmune diseases.

Materials and Methods

Animals and study location

For this study, 13 dogs were selected from both Ilha Solteira (51°06’35” W and 20°38’44” S) and Andradina cities (20°53’38” S and 51°23’1” W), which are endemic areas for CVL located in the northwestern region of the state of São Paulo (SP), Brazil. Ten dogs were naturally infected with *L. infantum* and three uninfected dogs were selected as controls.
The dogs were divided into three groups: group 1 (G1, n=5), dogs naturally infected with *L. infantum* detected by indirect enzyme immunoassay (ELISA), indirect immunofluorescence (IFAT), and polymerase chain reaction (PCR) from blood, and with intestinal amastigotes detected by immunohistochemistry and PCR of intestinal tissues; group 2 (G2, n=5), dogs naturally infected with *L. infantum* detected by ELISA, IFAT, and PCR from blood, but without intestinal amastigotes on immunohistochemistry and PCR of intestinal tissues; and group 3 (G3, n=3), *L. infantum*-uninfected dogs (control group) on serological, parasitological, and molecular exams.

The study was approved by the Ethics Committee for Animal Use (CEUA) of São Paulo State University, Ilha Solteira Campus (FEIS-UNESP), Department of Biology and Animal Science, under protocol no. 06/2014-CEUA.

**CVL diagnosis**

**CVL-positive dogs (G1 and G2 groups)**

During an epidemiological survey, the dogs were diagnosed with CVL by the Zoonosis Control Center (ZCC) of Ilha Solteira using the following methods: (a) direct parasitological examinations of popliteal or pre-scapular lymph node aspirate stained with Panoptic staining kit (Laborclin®, Pinhais, PR, Brazil) and (b) Dual Path Platform (DPP®) immunochromatographic test (Chembio Diagnostic Systems Inc., Medford, NY, USA) as a screening test and an indirect ELISA test (Biomanguinhos kit; Fiocruz, Rio de Janeiro, RJ, Brazil) for confirmation. Following confirmation, 10 symptomatic infected dogs were spontaneously delivered at ZCC by their guardians to be euthanized and were kindly donated by the ZCC for this study.

The serological tests of these animals were repeated in our laboratory using an indirect ELISA test and IFAT according to Oliveira et al. (2008). The optical density cut-off for ELISA and antibody titration factor for IFAT were 0.3802 and 1:40, respectively. The titer of-anti-*Leishmania* antibodies in IFAT ranged from 1:640 to 1:2560. The cut-off point of the ELISA test was determined multiplying the mean of the negative reference sera in absorbance values by the coefficient 2.5. The negative reference sera were from uninfected dogs from non-endemic areas for canine leishmaniasis.

For DNA extraction from blood and tissue samples of the intestine, we used the QIAamp Blood and Tissue kit (Qiagen, Santa Clarita, CA, USA). The amplification of the 447-base pair (bp) DNA fragments from the promastigote kinetoplast mini circle was performed by PCR using the pair of oligonucleotide primers to *L. infantum* MC1 (5'-GTT CGA TTT TTT TCT AGC TG-3') and MC2 (5'-CCC CCA TTT TTC CGA TTT TG-3') as described by Cortes et al. (2004).

The positive control for the reactions was a DNA sample from *L. infantum* (syn. *Leishmania chagasi*, MCAN/BR/1984/CCC-17,481) provided by the Laboratory of Leishmaniasis of the Leishmanioses Research Center at Fiocruz, Rio de Janeiro, Brazil and the negative control was sterile deionized water. The PCR products were visualized under ultraviolet light (UV) after 1.5% agarose gel electrophoresis containing the SYBR® safe dye (Invitrogen®, Carlsbad, CA, USA) in Tris-borate buffer (pH 8.0). The 100-bp ladder K180 DNA Molecular Weight Marker (Amresco®, Solon, OH, USA) was used to estimate the size of the amplicon.

**Control group – CVL-uninfected dogs (G3)**

The CVL-uninfected dogs (G3) were euthanized after injuries suffered by traffic accidents. Before death, the animals were attended at a veterinary clinic where feces, urine, lymph node, bone marrow, and blood samples were collected for clinical, parasitological, biochemical, serological, and molecular exams. The results of all exams were negative and the dogs had no any clinical signs of leishmaniasis. After euthanasia, samples of intestinal tissues were collected.

The hepatic and renal function of control dogs (G3) was assessed through measurements of alanine aminotransferase (ALT2S kit, ref 04657373), aspartate aminotransferase (ASTL kit, ref 04657543), urea (UREAL kit, ref 04657616), and creatinine (CREJ2 kit, ref 05401755) using the Cobas c111 analyzer (Roche Diagnostics International Ltd, Rotkreuz, Switzerland). The results were compared with reference values for healthy dogs and control dogs had normal enzyme levels.

**Euthanasia and necropsy procedures for tissue sample collection**

Infected dogs were euthanized and necropsied at the ZCC of Ilha Solteira, SP. Euthanasia was performed following the procedure no.1000/2012 from the Brazilian Federal Council of Veterinary Medicine (CFMV) and in accordance with Decree no. 51,838 of Brazilian Federal law published on March 14, 1963, which recommends that dogs infected with *Leishmania* be euthanized. Briefly, the animals received Diazepam (1 mg/Kg) as a preanesthetic drug (MPA) and then sodium thiopental (Thiopentax®, SP, Brazil, 30 mg/Kg) anesthesia. After the anesthesia it was intravenously administered a potassium chloride solution (10%) for animal death.

Following euthanasia, the abdominal cavities of dogs were opened for examination and organs and tissues were removed. First, the small intestine (duodenum, jejunum and ileum) was separated from the large intestine (ascending colon) and 0.5- to 1cm long fragments were collected, washed with phosphate buffered saline (PBS 0.01 M, pH=7.4), and fixed in 10% buffered formalin in 0.01 M PBS for 24 h for histopathological and immunohistochemical procedures.

The necropsy and tissue collection of control dogs followed the same procedures as described above.

Feces samples and intestinal contents of the small and large intestines were collected after death, identified, packed in ice, and sent to the laboratory for parasitological analysis. The results showed that the dogs did not have any gastrointestinal parasites, with the exception of *L. infantum* amastigotes in the intestinal tissues of dogs from the G1 group.
Histological procedures

The intestinal tissue samples collected from G1, G2, and G3 dogs were fixed in 10% formalin in 0.01 M PBS for histochemistry and immunohistochemistry.

Histochemistry: The intestinal tissues were fixed, embedded in paraffin, cut in 5-µm thick sections and stained with hematoxylin and eosin (HE) for histological examination. Histologically stained macrophages were identified in tissues from infected and uninfected dogs. Three sections of each intestinal tissue were evaluated under a light microscope at 40x and/or 100x magnification.

Immunohistochemistry: Intestinal tissue samples were previously hydrated in successive passages in xylen and ethanol. Initially, the endogenous peroxidase enzyme was blocked using 3% hydrogen peroxide solution (H₂O₂) diluted in 0.01 M PBS (pH=7.4). Non-specific secondary antibody reactions were inhibited using the normal serum of the animal of which the secondary antibody was made (goat serum) in 1:50 dilution. Samples were incubated with the primary antibody in a humid chamber at 4 °C overnight. After three rinses with PBS, samples were then incubated with the secondary antibody for 1 h at room temperature, and after successive rinses, an avidin-biotin peroxidase complex (Vectastain ABC Kit; Vector Laboratories Inc., Burlingame, CA, USA) was applied for immune reaction. After appropriated rinses with PBS, 50 to 100 µl of freshly prepared solution of horseradish peroxidase substrate (VECTOR® NovaRED™; Vector Laboratories Inc., SK-4800) was placed on each tissue section and left for 5 min. After rinsing in distilled water, tissues were counterstained with Mayer hematoxylin, dehydrated, and the slides were mounted and coverslipped using Canada balsam.

As reaction control, one slide of each tissue was not incubated with the primary antibody. In addition, tissues from dogs in the control group (CVL-uninfected dogs) were used as negative control for the reaction. Two slides of each tissue from groups G1, G2, and G3 were analyzed.

Leishmania infantum amastigote immunostaining was performed according to the procedures described by Tafuri et al. (2004). The primary antibody used was the immune serum of L. infantum-naturally infected dogs (diagnosed by ELISA and RIFI; titer=1: 2560) and the secondary antibody was the biotinylated rabbit anti-IgG antibody produced in goat (immunoe reaction crossed with dogs; in 1:500 dilution).

Tissue antigen retrieval using TRIS (pH=7.6) + trypsin (1:250) was performed for immunostaining of CD4⁺ and CD8⁺ lymphocytes for 30 min. The primary antibodies used were anti-dog monoclonal antibodies produced in rat anti-CD4 (AbD serotec-MCA1038GA, Bio-Rad Laboratories, Hercules, CA, USA) and anti-CD8 T cells (AbD serotec-MCA1039GA, Bio-Rad Laboratories). Next, we added the biotinylated anti-rat secondary antibody produced in goat (AbD Serotec-STAR131B, Bio-Rad Laboratories) diluted 1:200, which reacted specifically with the respective primary antibodies.

Fluorescence immunolabeling: Treg cells were detected using anti-mouse/rat FoxP3 antibody conjugated with a fluorescein molecule (eFluor® 570; eBioscience, San Diego, CA, USA), which also detects dog FoxP3. The tissue was prepared according to standard techniques for histology and for antigen retrieval using TRIS buffer (pH=7.6) + trypsin (1:250). Next, 5 µl of undiluted primary antibody was added to each tissue section and after 30 min the material was washed with PBS (pH=7.4, 0.01 M) and the slides were covered with a coverslip in glycerin. The material was observed under a fluorescence microscope (Olympus, BX-FLA).

Semi-quantitative analysis

Leishmania infantum amastigotes: The semi-quantitative analysis was performed only in intestinal tissues with L. infantum amastigotes. For this analysis, 100 positive visual microscopic fields of the intestinal tissues were analyzed at 40x magnification and the positive cells containing the parasites were counted. The tissues were scored according to parasite burden (+ to ++++) and positive cell scores (1 to 4), as follows: no positive cells (without amastigotes) or negative tissues; 1-50 positive cells (score 1; +) or low parasite burden; 51-100 positive cells (score 2; ++), or moderate parasite burden; 101-200 positive cells (score 3; +++), or intense parasite burden; > 201 positive cells (score 4; ++++), or very intense parasite burden.

CD4⁺, CD8⁺, and FoxP3⁺ T lymphocytes and macrophages: For a semi-quantitative analysis of T lymphocytes (CD4⁺, CD8⁺, and Treg) and macrophages, approximately 100 visual microscopic fields were visualized under a microscope at 100x magnification. The cellular score was determined by the number of positive microscopic fields where at least one cell from each cell type was detected, as follows: ≤ 25 positive microscopic fields were classified as having low cellular levels (+) and score 1; 26-50 positive fields, moderate cellular levels (+, score 2); 51-75 positive fields, high cellular levels (+++, score 3); and > 76 positive fields were classified as having very high cellular levels (++++, score 4).

All cells evaluated were from the intestinal mucosa, represented by crypt-villus units (CVU) in three segments of the small intestine and one of the large intestine.

Statistical analysis

The residues were submitted to the Shapiro-Wilk normality and outliers test, the homogeneity of the variances was evaluated by the Levene test. The mean scores for the semi-quantitative T lymphocytes (CD4⁺, CD8⁺ and Treg FoxP3⁺) and macrophages from the experimental groups of dogs (G1, G2 and G3) were analyzed by ANOVA using the F test, and the comparison mean of the groups was performed according to the Duncan's test, using the "agricolae" data package.

The correlation between cellular scores and parasite burden was determined using the Spearman correlation test. For both tests, 95% confidence intervals were calculated and p values ≤ 0.05 were considered significant. All analyses were performed using R statistical software version 3.1.1 (R CORE TEAM, 2017).
Results

According to the histological analysis, dogs from the G1 group had *L. infantum* amastigotes in at least one intestinal segment (duodenum, jejunum, ileum, or ascending colon). Two dogs in the G1 group showed a parasite burden ranging from high (score 3; ++++) to very high (score 4; +++++) in all intestinal segments (Table 1, Figure 1A). In addition, the colon was the intestinal segment with the highest parasite burden and also the one more frequently infected; however, there was no significant difference in parasite burden scores between intestinal segments (Table 1).

Macrophages were distributed in the mucosa (CVU) and submucosa of dogs from the G1, G2, and G3 groups (Figure 1B), but macrophage scores between the G1, G2, and G3 groups in the mucosa were significantly different (G1 > G2 > G3, \( p \leq 0.05 \)) (Table 2). Macrophage scores in G1 dogs ranged from 2 (+++) to 4 (++++), and the highest score was detected in animals with highest parasite burden (Tables 1 and 2). When each intestinal segment was analyzed, macrophage scores in the duodenum and colon were significantly higher than in the jejunum and ileum in G1 dogs and significantly different from that of G2 and G3 dogs. Most dogs in the G2 group had moderate macrophage scores (2/++), but their scores were higher than in the G3 group. In addition, macrophage scores of dogs from G1 and G2 groups were significantly different from the control group (G3) only in the duodenum (Table 2).

There was no significant difference in CD4\(^+\) T cell levels among all experimental groups (\( p \geq 0.05 \)). Similarly, no significant difference was observed among all intestinal segments (Table 3, Figures 2A and C). In contrast, the semi-quantitative analysis showed that CD8\(^+\) T lymphocyte scores were significantly higher in the small and large intestines (\( p \leq 0.05 \)) in dogs from G1 and G2 groups in relation to the control dogs (G3) (Table 4, Figures 2B and D). In addition, CD8\(^+\) T lymphocyte scores in the ileum (Peyer’s patches, Figure 2B), duodenum and colon of dog’s numbers 4 and 5 from G1 group were high in these organs, which also had very high parasite burdens (score 3-4; +++, ++++) (Tables 1 and 4).

Table 1. Parasite burden scores (1 [+] to 4 [++++]) in intestinal (duodenum, jejunum, ileum, and colon) mucosal tissues (crypt-villus unit – CVU) from dogs naturally infected with *Leishmania infantum*. Ilha Solteira, SP, Brazil, 2017.

<table>
<thead>
<tr>
<th>Group</th>
<th>Dog</th>
<th>Duodenum</th>
<th>Jejunum</th>
<th>Ileum</th>
<th>Colon</th>
</tr>
</thead>
</table>
| G1    | 01  | N        | N       | N     | 3 (+++)
|       | 02  | 2 (+++)  | N       | 2 (+) | N     |
|       | 03  | N        | N       | N     | 3 (+++) |
|       | 04  | 4 (++++) | 3 (++)  | 4 (++++)| 3 (+++)|
|       | 05  | 3 (+++)  | 2 (+)   | 3 (++)| 4 (++++)|
| Average (scores) | 1.8 | 1.0 | 1.8 | 2.6 |

Note: N = negative. According to the F-test, there is no significant difference to the level of 5% probability (\( p \leq 0.05 \)). R version 3.1.1 (R CORE TEAM, 2017).

Figure 1. Histological sections of the intestine of dogs with CVL. (A) (ileum) Numerous macrophages with amastigotes of *Leishmania infantum* aggregated in the lamina propria at the tips of villi (magnification: 40×, staining: immunohistochemistry); (B) (colon) Basal crypt and submucosa. Red arrows = hypertrophic macrophages with amastigote (magnification: 100×, staining: H&E). Bars = 50 µm. Ilha Solteira, SP, Brazil, 2017.
Table 2. Semi-quantitative analysis of macrophages in intestinal (duodenum, jejunum, ileum, and colon) mucosal tissues (crypt-villus unit – CVU) from dogs naturally infected with *Leishmania infantum* and uninfected dogs. Scores range from 1 to 4. Ilha Solteira, SP, Brazil, 2017.

<table>
<thead>
<tr>
<th>Group</th>
<th>Dogs</th>
<th>Duodenum</th>
<th>Jejunum</th>
<th>Ileum</th>
<th>Colon</th>
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</thead>
<tbody>
<tr>
<td>G1</td>
<td>N=5</td>
<td>3.2&lt;sup&gt;ab&lt;/sup&gt;</td>
<td>2.6&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.0&lt;sup&gt;ab&lt;/sup&gt;</td>
<td>3.8&lt;sup&gt;a&lt;/sup&gt;</td>
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<td>N=3</td>
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<td>2.0&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.0&lt;sup&gt;a&lt;/sup&gt;</td>
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Average of the intestines

<table>
<thead>
<tr>
<th>Group</th>
<th>Dogs</th>
<th>Small intestine</th>
<th>Large intestine</th>
<th>Intestinal tract (all)</th>
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<td>2.0&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Note: Comparison of averages by Duncan’s test. Different lowercase letters (a, b and c) indicate statistical significance between lines and different capital letters (A, B and C) indicate significance statistical between columns at 5% of probability (p ≤ 0.05). Program R version 3.1.1 (R CORE TEAM, 2017).

Table 3. Semi-quantitative analysis of CD4<sup>+</sup> T lymphocytes in intestinal (duodenum, jejunum, ileum, and colon) mucosal tissues (crypt-villus unit – CVU) from dogs naturally infected with *Leishmania infantum* and uninfected dogs. Scores range from 1 to 4. Ilha Solteira, SP, Brazil, 2017.

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<tr>
<td>G1</td>
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Average of the intestines

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<tbody>
<tr>
<td>G1</td>
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<td>N=3</td>
<td>1.0&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.0&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.0&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Note: According to the F-test, there is no significant difference to the level of 5% probability (p ≤ 0.05). For lowercase letters (a, b and c) the comparisons were made in lines and for capital letters (A, B and C) the comparisons were made in columns. R version 3.1.1 (R CORE TEAM, 2017).

The presence of regulatory T cells (FoxP3<sup>+</sup>) was low in all intestinal tissues (small or large intestines) of dogs in the G1, G2, and G3 groups and no significant difference was observed in FoxP3 expression among all experimental groups (p ≥ 0.05; Table 5, Figures 2E and F). Few or none FoxP3<sup>+</sup> T cells were observed in control samples (G3). Thus, dogs with intense parasite burden (Table 1) also had low scores for this type of regulatory cell (Table 5).

The cellular scores of CD4<sup>+</sup>, CD8<sup>+</sup>, Treg (FoxP3<sup>+</sup>), and macrophages were compared in pairs and with parasite burden using correlation analysis. Parasite burden and CD4<sup>+</sup> T lymphocyte scores showed a significant negative correlation (p ≤ 0.05); tissues with higher parasite burdens had low CD4<sup>+</sup> T cell levels (rs=-0.89, p=0.041). Similarly, CD4<sup>+</sup> T cell levels correlated negatively with the levels of CD8<sup>+</sup> T cell (rs=-0.89, p=0.043) and macrophages (rs=-0.92, p=0.028), in the same intestinal region of dogs from G1 group (Table 6).

In contrast, CD8<sup>+</sup> T lymphocyte levels correlated positively with the levels of macrophages in the large intestine of G1 dogs (rs=0.97, p=0.006) and in the small intestine of G2 dogs (rs=0.89, p=0.041; Table 6). The levels of both cell types (CD8<sup>+</sup> and macrophages) also correlated positively with parasite burden in the large intestine (rs=0.91, p=0.03 and rs=0.88, p=0.046, respectively) (Table 6). Conversely, no significant correlation was observed between parasite burden and cell levels in the small intestine of G1 dogs.

There was a negative correlation between FoxP3<sup>+</sup> T cell scores and parasite burden in the small intestine of G1 dogs (rs=-0.89, p=0.041), but no significant correlation was observed in the large intestine of animals from the same group (Table 6).

In Figure 3 it is possible to comparatively evaluate the distribution of T lymphocytes (CD4<sup>+</sup>, CD8<sup>+</sup>, and FoxP3<sup>+</sup>) and macrophages along the small and large intestine from the experimental groups (G1, G2, and G3).

**Discussion**

*Leishmania infantum* is an intracellular parasite that multiplies in macrophages residing in several organs and lymphoid tissues, including intestinal tissues of dogs (PINTO et al., 2011, 2013; SILVA et al., 2016), causing clinical and pathological alterations (FEITOSA et al., 2000). Anderson et al. (1980) observed macrophage infected with *L. infantum* in the lamina propria and submucosa of the stomach, duodenum, jejunum, ileum,
Figure 2. Histological sections of the small intestine. Figures (A, B and E) of dogs infected with *Leishmania infantum* intestinal amastigotes (group G1), (C, D and F) of dogs infected but without intestinal amastigotes (group G2). (A) Payer's patch in the ileum showing that there was no CD4+ T lymphocyte immunolabeling (animal 04); (B) Similar region as in A with immunostained CD8+ T lymphocytes; (C) Villus of the duodenum with immunostained CD4+ T lymphocytes (red arrows); (D) Similar region as in C with immunostained CD8+ T lymphocytes (black arrows); (E and F) villus region of jejunum and ileum, respectively, showing immunofluorescence of FoxP3+ T cells. Magnification = 40x. Bars = 50 µm. Ilha Solteira, SP, Brazil, 2017.
and colon of a dog with CVL associated with a chronic intestinal inflammatory reaction (PINTO et al., 2011, 2013; SILVA et al., 2016). Additional information about histopathological description of intestinal alterations of the dogs (G1 group) of this study can be found in Silva et al. (2016). In the present study, the increase of macrophages in intestinal tissues (small and large intestines) was observed in CVL-positive dogs (G1 and G2 groups), but hypertrophic macrophages with *L. infantum* amastigotes were seen only in G1 dogs (naturally infected dogs with intestinal amastigotes).

### Table 4. Semi-quantitative analysis of CD8+ T lymphocytes in intestinal (duodenum, jejunum, ileum, and colon) mucosal tissues (crypt-villus unit – CVU) from dogs naturally infected with *Leishmania infantum* and uninfected dogs. Scores range from 1 to 4. Ilha Solteira, SP, Brazil, 2017.

<table>
<thead>
<tr>
<th>Group</th>
<th>Dogs</th>
<th>Duodenum</th>
<th>Jejunum</th>
<th>Ileum</th>
<th>Colon</th>
</tr>
</thead>
<tbody>
<tr>
<td>G1</td>
<td>N=5</td>
<td>2.8&lt;sup&gt;A&lt;/sup&gt;</td>
<td>2.2&lt;sup&gt;B&lt;/sup&gt;</td>
<td>2.4&lt;sup&gt;B&lt;/sup&gt;</td>
<td>3.6&lt;sup&gt;A&lt;/sup&gt;</td>
</tr>
<tr>
<td>G2</td>
<td>N=5</td>
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<td>2.0&lt;sup&gt;A&lt;/sup&gt;</td>
<td>2.4&lt;sup&gt;A&lt;/sup&gt;</td>
</tr>
<tr>
<td>G3</td>
<td>N=3</td>
<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

#### Average of the intestines

<table>
<thead>
<tr>
<th>Group</th>
<th>Dogs</th>
<th>Small intestine</th>
<th>Large intestine</th>
<th>Intestinal tract (all)</th>
</tr>
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<tbody>
<tr>
<td>G1</td>
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<td>2.3&lt;sup&gt;A&lt;/sup&gt;</td>
</tr>
<tr>
<td>G3</td>
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<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Note: Comparison of averages by Duncan’s test. Different lowercase letters (a, b and c) indicate statistical significance between lines and different capital letters (A, B and C) indicate significance statistical between columns at 5% of probability (p ≤ 0.05). Program R version 3.1.1 (R CORE TEAM, 2017).

### Table 5. Semi-quantitative analysis of FoxP3+ T reg cells in intestinal (duodenum, jejunum, ileum, and colon) mucosal tissues (crypt-villus unit – CVU) from dogs naturally infected with *Leishmania infantum* and uninfected dogs. Scores range from 1 to 4. Ilha Solteira, SP, Brazil, 2017.

<table>
<thead>
<tr>
<th>Group</th>
<th>Dogs</th>
<th>Duodenum</th>
<th>Jejunum</th>
<th>Ileum</th>
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</thead>
<tbody>
<tr>
<td>G1</td>
<td>N=5</td>
<td>1.4&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.6&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.4&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.2&lt;sup&gt;A&lt;/sup&gt;</td>
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<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
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#### Average of the intestines

<table>
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<th>Group</th>
<th>Dogs</th>
<th>Small intestine</th>
<th>Large intestine</th>
<th>Intestinal tract (all)</th>
</tr>
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<tbody>
<tr>
<td>G1</td>
<td>N=5</td>
<td>1.5&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.2&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.4&lt;sup&gt;A&lt;/sup&gt;</td>
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<tr>
<td>G2</td>
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<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
<td>1.0&lt;sup&gt;A&lt;/sup&gt;</td>
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Note: According to the F-test, there is no significant difference to the level of 5% probability (p ≤ 0.05). For lowercase letters (a, b and c) the comparisons were made in lines and for capital letters (A, B and C) the comparisons were made in columns. R version 3.1.1 (R CORE TEAM, 2017).

### Table 6. Correlation analysis between monocytic cells (CD4+ and CD8+ T cells, FoxP3+ regulatory T cells, and macrophages) and parasite burden of *Leishmania infantum* amastigotes in the intestine of naturally infected dogs (G1 group). Ilha Solteira, SP, Brazil, 2017.

<table>
<thead>
<tr>
<th>Spearman correlation for monocytic cells semi-quantified in the intestines of dogs naturally infected with CVL</th>
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<tbody>
<tr>
<td>Group</td>
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<tr>
<td></td>
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<tr>
<td>G1 (N = 5)</td>
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Note: SIP = parasitic intensity score; SCD4<sup>+</sup> = score of CD4<sup>+</sup> T lymphocytes; SCD8<sup>+</sup> = score of CD8<sup>+</sup> T lymphocytes; SMAC = score of macrophages; SFoxP3 = score of regulatory T lymphocytes (Treg); (*) significant test at 5% probability (p ≤ 0.05). Analysis carried out in accordance with linear correlation coefficient of Spearman (rs); p ≤ .05. Program R, version 3.1.1 (R CORE TEAM, 2017).
In this study, a semi-quantitative analysis showed that G1 dogs had the highest scores for macrophages with amastigotes, particularly in the large intestines (colon) and also showed a positive correlation between the parasite burden and macrophages. These data may suggest the action of M2 macrophages in these animals. According to Moreira et al. (2016), M2 macrophages were the predominant cell type in granulomas and inflammatory infiltrates observed in the skin, lymph nodes and spleens from VL infected dogs, which coincided with the highest parasite burden found in these dogs. These evidences led the authors to conclude that the involvement of M2 macrophages related with amastigotes of *L. infantum* in infected intestinal tissues as in the present work.

Macrophages with *Leishmania* spp. amastigotes can stimulate CD4+ T cells by antigen expression associated with major histocompatibility complex (MHC) class II and also CD8+ T cells via the expression of antigens associated with MHC class I molecules (LANG et al., 1994). As we observed in the intestinal tissues of G1 dogs, the increase in CD8+ T lymphocytes and the non-alteration of CD4+ T lymphocyte population suggest that the infected macrophages of these animals were expressed via MHC class I.

The high parasite burden found in the intestinal tissues of G1 dogs associated with chronic mononuclear inflammatory infiltrate represented by lymphocytes and macrophages, may also suggest that the parasite has the ability to evade the cellular immune response of the host, which may become immunocompromised over time. Previously, Passero et al. (2010) suggested that *L. shawi* parasites regulate different cell populations in infected BALB/c mice to preferentially express immunosuppressive activity or deactivating cytokines, thus avoiding their own destruction. It is well established that the IFN-γ cytokine secreted by Th1 T cells controls the growth of *Leishmania* spp. in macrophages (MURRAY et al., 1992) by inducing nitric oxide synthesis, which activates the microbicidal functions of this cell (REINER & LOCKSLEY, 1995). Moreover, an increase in IL-10, a cytokine secreted by Th2 T cell response associated with disease progression in CVL symptomatic dogs has been observed in the absence of IFN-γ (BOGGIATTO et al., 2010).

Cell-mediated immunity of the host plays an important role in the control of *Leishmania* spp. infection. In the current study, there was no significant difference in CD4+ T cell scores among the animals from G1, G2 or G3 groups (p ≥ 0.05). In addition, CD4+ T cell population had no increase in intestinal tissues even in G1 dogs with high parasite burden in the mucosa (villi and crypts) and submucosa, which cellular scores were inversely correlated with parasite burden. In dogs infected with *L. infantum*, Bourdouiseau et al. (1997) showed a reduction of CD4+ T lymphocytes numbers in the peripheral blood concomitant with the proliferation of the parasite in tissue macrophages, suggesting the absence of an effective immune response to eliminate the parasite. According to Alvar et al. (2004), the reduction of CD4+ T cell levels during the progression of disease in dogs was indicative of the failure of the host to control the parasite, allowing it to migrate to other tissues, increasing the parasite load and the infectivity of the host.

According to Pinelli et al. (1994); Brachelente et al. (2005); Barbieri (2006) and Cruz-Chan et al. (2014) severe and progressive CVL in infected dogs was associated with a predominantly Th2 humoral immune response that was not able to control the infection. Similarly, Stäger and Rafati (2012) showed in rats experimentally infected with *L. major* in subcutaneous tissue that IFN-γ production by Th1 cells was essential to control the infection, but in contrast, Th2 response resulted in susceptibility to the disease. Similarly, Carvalho et al. (2012) also reported that BALB/c mice experimentally infected with *L. (Leishmania) amazonensis* presented a Th2 response that was related to susceptibility to the parasite. Moreover, BALB/c strain mice experimentally infected with *L. (Vianna) brasiliensis* triggered a Th1 cellular immune response in the skin associated with resistance to the parasite. The considerable number of infected macrophages in intestinal tissues of G1 group dogs suggests the intestinal susceptibility of these animals to *L. infantum*. However, studies that relate resistance or susceptibility of dogs to intestinal CVL are still lacking.

Pinelli et al. (1995) demonstrated the involvement of CD8+ lymphocytes in resistance to canine VL. These lymphocytes were detected in asymptomatic dogs experimentally infected with *L. (L.) infantum* and subsequent lysis of *Leishmania*-infected macrophages. In contrast, symptomatic dogs showed a reduced lymphoproliferative response to specific antigenic stimulation, resulting in a lower ability of CD8+ T lymphocytes to lyse infected macrophages. In our study, G1 dogs showed high levels of CD8+ T lymphocytes and low levels of CD4+ T lymphocytes in intestinal tissues. In addition, CD8+ T lymphocyte scores correlated positively with parasite burden in intestinal tissues from dogs of G1 group; significantly higher in the duodenum and colon, where parasite burdens were more intense. Thus, it is likely that CD8+ mediated cellular immune response was not able to control the intestinal infection, possibly due to the
inability of these cells to exert their cytotoxic activity in infected macrophages as suggested by Pinelli et al. (1995). In a study of CD8+ T cells role in Leishmania donovani infection, Joshi et al. (2009) reported that CD8+ T cell responses against the parasite were defective in their cytotoxic activity toward infected cells.

We also detected the presence of parasites in Peyer’s patches and CD8+ T lymphocytes in higher scores than CD4+ T cells indicating that the parasites were invading the lymphoid protective tissue in the intestinal mucosa and that the immune system was not able to control the expansion of the parasite in these organs. For Stäger & Rafati (2012) the protective role of CD8+ T cells during infection by Leishmania spp. is controversial, particularly because of the diversity of different Leishmania species with different tropisms that is reflected in the various clinical manifestations of the disease.

According to Yurdakul (2005), the inability of T cells to activate macrophages to destroy Leishmania spp. amastigotes appears to be the primary defect in this disease, which is associated with parasite multiplication and migration, affecting multiple organs of the host. Our results showed that CD8+ T lymphocytes were unable to destroy parasitized macrophages, suggesting that other anti-inflammatory components such as regulatory T cells could mediate immune response, which may explain the lack of resolution and the maintenance and progression of the disease. The high parasite burden in the colon mucosa with only mild pathological alterations led Pinto et al. (2011) and Pinto et al. (2013) to suggest that L. infantum may benefit of the intestinal immune regulatory response (immunological tolerance). In fact, this type of immunoregulatory response could be mediated by Treg cells. More recently, Figueiredo et al. (2014) suggested that Treg cells could mediate both intestinal leishmaniasis and the homeostasis of the colon under normal physiological conditions. In that study, cellular expression of toll-like receptor 2 (TLR2) and IL-10, and TNF-α were more frequent in the jejunum, which had fewer parasites than the colon. These conflicting findings indicate that further studies are needed to elucidate important immune mechanisms related to parasitic infection in the intestinal wall.

In our study, Treg cell scores in intestinal tissues of dogs in the G1 group (infected dogs with amastigotes in the intestine) were more frequent in the jejunum, which had GITR on CD4+CD25+ T cells in normal skin and inflammatory organs of the host. Our results showed that CD8+ T lymphocytes and macrophages population associated with high parasite loads, but no changes of CD4+ T cells and FoxP3+ Treg cells suggest a possible immunoregulation by the parasite not dependent on Treg cells.

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**References**


