Inflammation and structural changes of splenic lymphoid tissue in visceral leishmaniasis: A study on naturally infected dogs

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SUMMARY

The aim of this study was to identify splenic immuno-inflammatory patterns associated with natural infection by Leishmania chagasi. Spleen samples were obtained from 72 stray dogs from an endemic area of visceral leishmaniasis. The animals were grouped into four categories as follows: (i) potentially resistant to visceral leishmaniasis, with a positive leishmanin skin test result, and negative splenic culture for Leishmania parasites (ii) potentially susceptible to visceral leishmaniasis, with a negative leishmanin skin test and positive splenic culture for Leishmania (iii) infected with undefined susceptibility status, with a positive leishmanin skin test and positive splenic culture for Leishmania, and (iv) noninfected, with a negative leishmanin skin test, negative splenic culture for Leishmania, and negative serology for anti-Leishmania antibodies. Histopathological analyses showed that there was a higher frequency of perisplenitis (18/25, \( P < 0.0001 \)), granuloma (7/25, \( P = 0.0102 \)), structural disorganization (14/25, \( P < 0.0001 \)), and atrophy of the lymphoid follicles (20/25, \( P = 0.0036 \) and of the marginal zone (15/25, \( P = 0.0025 \)) in the potentially susceptible group than in the other groups. The data presented here show changes in the white pulp of the spleen that are associated with naturally acquired visceral leishmaniasis.

Keywords: canine visceral leishmaniasis, Leishmania chagasi, Leishmania infantum, lymphoid tissue, Spleen

INTRODUCTION

Visceral leishmaniasis is used to describe two clinically, parasitologically and epidemiologically different diseases: antroponotic visceral leishmaniasis, caused by Leishmania donovani, and zoonotic visceral leishmaniasis (VL), caused by L. chagasi/L. infantum. The latter disease is endemic in Mediterranean countries and the American continent, and its causal agents are transmitted to mammalian hosts by sand flies of the Phlebotomus and Lutzomyia genera (1). Leishmania chagasi/L. infantum infection does not always lead to VL (2). Most infected humans (2,3) and at least a proportion of infected dogs (4,5) remain asymptomatic or develop a mild disease, which is spontaneously cured. Occasionally, dogs and humans develop a severe form of VL (2,6) that is usually lethal if left untreated. In this report, we use the term VL to refer to this severe form of the disease. In both humans and dogs, the disease proceeds with emaciation, enlargement of the liver and spleen, fever, anaemia, and an increased predisposition to bacterial infection (2,7–10). In dogs, such signs of disease are further accompanied by a variety of skin and ocular lesions (9). Much that is known about the immune response to Leishmania parasites, and even the immune response in general has come from studies in murine models of leishmaniasis (11–13). Most of these models, however, are intrinsically compromised by the artificial nature of the infective inocula. Instead of consisting only of naturally infective, metacyclic promastigotes, these inocula usually contain (i) either relatively large number of complement-fixing, noninfective procyclic promastigotes (usually constituting more than 70% of the inoculum) (14) or (ii) amastigotes, which are not the usual infectious form of the parasite. Canine VL, arising from natural infection by L. chagasi, is a disease worthy not only of study in its own right for veterinary and epidemiological reasons (15), but also for consideration as an attractive animal model for the human disease, because...
the infective inoculum is very likely identical to that causing human infection in endemic areas. Observations made in canine VL may therefore confirm and refine conclusions drawn from studies using murine experimental models.

Recently, we observed that a positive leishmanin skin test (LST) was negatively associated with emaciation and specific antibody activity in Leishmania-infected dogs (16). In the present study, we examined the spleen histology of these animals to identify alterations potentially associated with disturbances in the immune response to Leishmania and other pathogens that occur at late stages of the disease.

We decided to examine the spleen because it plays a central role in VL. The spleen is infected in all cases of the disease. In contrast to other organs like the liver, the spleen maintains the infection during the entire course of VL (11,12). In fact, a splenic index of parasitism is used as a clinical criterion of therapeutic response in human VL (17). The spleen is also a lymphoid organ with cellular diversity and compartmentalized microenvironments, suitable for the definition of immunologic processes that may influence the outcome of infections (18,19). During the course of VL, the spleen becomes an evident site of interaction between the immune system and the Leishmania, because all of the obligatory participants in the immune response against the parasite are present in large quantities. These include the antigen (live parasites or their debris), antigen-presenting cells, and lymphocytes capable of responding to these antigens. Changes in the splenic microenvironment have been identified following the experimental infection of mice by a variety of pathogens, including Leishmania (20,21).

Our hypothesis is that such alterations are also present in the course of natural infection by L. chagasi in dogs and that they reflect a status of susceptibility to the disease.

MATERIALS AND METHODS

Animals

Spleen samples from 72 dogs were used in this study. The specimens were obtained from the archive of histological samples of the Gonçalo Moniz Research Center, FIOCRUZ, Salvador, Brazil. These animals were stray dogs of different breeds and estimated ages collected from the streets of Jequié (Bahia State, Brazil, an endemic area for visceral leishmaniasis) at different periods between 1998 and 2001 in collaboration with the Endemic Diseases Surveillance Program of the State Health Service. The nutritional status of the animals was recorded into two categories: showing evidence of emaciation (when prominences of bones from the vertebral column, iliac crest and ribs were present), or normal (when bone prominences were absent). The presence of anti-Leishmania antibodies in the serum were investigated by ELISA, and a cellular immune response against L. chagasi antigens was detected by LST. The details of these tests are reported elsewhere (16,22,23). The animals were kept for at least 48 h in the municipality kennel. When not claimed by their owners, the animals were culled, as recommended by the Brazilian program for the control of zoonotic diseases. Spleen specimens were then collected for both Leishmania, detection by in vitro culture, and histology. All procedures were conducted in accordance with the Oswaldo Cruz Foundation guidelines for animal experimentation. All of the animals that had detectable Leishmania in their spleen cultures or a positive LST, with available spleen samples, were included in the study. The animals were distributed into the following categories: (i) Infected and potentially resistant to VL – 22 animals with a positive LST and negative spleen culture for Leishmania; (ii) Infected and potentially susceptible to VL – 25 animals with a negative LST and positive spleen culture for Leishmania; (c) Infected with undefined susceptibility status – 11 animals with both a positive LST and positive spleen culture for Leishmania. As a control group, specimens from 14 animals with negative spleen cultures, a negative LST, and the absence of anti-Leishmania antibody activity in the serum as measured by ELISA were randomly selected; we referred to this group as noninfected. Table 1 summarizes the characteristics of the groups used in the study.

Spleen samples and histological analysis

Three to 4 mm-thick slices of spleen tissue were cut transversally to the capsule and fixed in 10% formalin. After fixation, tissue slices were embedded in paraffin. Four to 5 μm-thick sections were cut and stained with haematoxylin and eosin (H&E). These sections were examined by at least two of the authors who were blind to previous knowledge of the identities of the animals. The authors observed the following parameters:

1. Presence of perisplenitis, as evidenced by a focal thickening of the splenic capsule with an inflammatory infiltrate frequently associated with incipient fibrosis (Figure 1a).
2. Presence of granuloma, with focal aggregates of four or more epithelioid macrophages (Figure 1b,c).
3. Lymphoid follicle frequency, in which the density of lymphoid follicles in the splenic tissue sections was classified as rare (follicles were infrequently found), low (follicles were easily found but sparse in the red pulp), average (follicles were uniformly distributed and easily found in the section) and high (follicles were frequent and large).
4. Size of lymphoid follicles, germinal centre and marginal zone, in which the lymphoid follicles and the marginal zone were classified as tiny, small, average and large.
5. Degree of structural organization of the white pulp, in which the white pulp was classified as: well organized, with distinct...
peri-arteriolar lymphocyte sheath, germinal centre, mantle zone and marginal zone (24); slightly disorganized, with either hyperplastic or hypoplastic changes leading to a loss in definition of any of the regions of the white pulp; moderately disorganized, when the white pulp was evident, but its regions were poorly individualized or indistinct; extensively disorganized, when the follicular structure was barely distinct from the red pulp and T-cell areas (Figure 1d–i). The last two categories were frequently associated with lymphoid atrophy.

6. Frequencies of cell populations in the red pulp, with the frequencies of lymphoblasts, plasma cells, macrophages, granulocytes, and megakaryocytes semiquantified as absent, rare (seen as isolated cells or small aggregates in a small proportion of the examined ×400-magnified microscopic fields), few (seen as isolated cells or small aggregates in approximately half of the examined ×400-magnified microscopic fields), average (seen as single cells or aggregates observed in most of the examined ×400-magnified microscopic fields), or frequent (consisting of single cells or aggregates seen in roughly all of the examined ×400-magnified microscopic fields). The cells were characterized morphologically according to the following criteria:

a. Lymphoblasts: Cells with round nuclei containing condensed chromatin, inconspicuous nucleoli, and scant cytoplasm.

b. Plasma cells: Cells with eccentric nuclei, heterochromatin dispersed around the edge in a pattern similar to the numerals of an analogue clock face, and basophilic cytoplasm with clear, perinuclear vacuoles.

c. Macrophages: Large cells with oval or reniform nuclei containing loosely packed chromatin surrounded by a rim of eosinophilic cytoplasm with indistinct edges.

d. Granulocytes: Cells with highly lobulated nuclei with densely packed chromatin.

e. Megakaryocytes: Very large cells with wide and clearly defined cytoplasm and large, lobulated nuclei containing condensed chromatin.

7. Parasite burden, where the amount of amastigotes in the spleen sections was estimated by conventional light microscopy examination of a minimum of 40 and a maximum of 100 nonoverlapping high-power (×1000-magnified) fields per section. The examined fields were equally distributed between the subcapsular (Figure 1k) and internal red pulp (Figure 1l). The results are expressed as the proportion of independent fields with amastigotes.

Expression and analysis of the results

The numerical data shown in the text, table and graphs represent absolute or relative numbers of animals with the corresponding condition, unless otherwise stated. The significance of the differences between the proportions of the studied variables among the groups was tested using the $\chi^2$-test with a Yates’ correction, or a Fishers’ exact probability test when recommended. When appropriate, a paired $t$-test or the Kruskal–Wallis statistic were used for testing differences involving the means of numerical data obtained from different groups (25). The critical level of significance was set at $P < 0.05$.

RESULTS

Perisplenitis

Perisplenitis was present in 20 (28%) out of 72 animals (Table 2 and Figure 1a). It was more frequent in animals in the infected and potentially susceptible to VL group (18 out of 25 animals, 72%) than other groups ($\chi^2$-test, $P < 0.0001$).

Table 1 General characteristics of a sample of dogs, from an endemic area of visceral leishmaniasis, used in this study

| Parameters                  | Infected | Potential susceptible to VL | Undefined susceptibility status | Noninfected |
|-----------------------------|----------|-------------------------------|---------------------------------|-------------|
| Number of animals           | 22       | 25                            | 11                              | 14          |
| Spleen culture              | –        | +                             | +                               | –           |
| Leishmanin skin test        | +        | –                             | +                               | –           |
| Serology                    | 0.55 ± 0.39 | 1.54 ± 0.91$^b$               | 0.87 ± 0.77                     | 0.28 ± 0.13 |
| With emaciation             | 6/20 (30%) | 15/25 (60%)$^d$               | 2/11 (18%)                      | 4/13 (31%)  |

$^a$ Expressed as mean ± SD of the OD. 490 nm obtained in indirect ELISA for detection of anti-Leishmania antibodies in the serum; $^b$Statistically significant from the control group, Kruskal–Wallis test, $P < 0.001$; $^c$Number of emaciated animals/total number of animals in which emaciation was investigated (%); $^d$Statistically significant from the other groups, $\chi^2$-test $P = 0.02$. 

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Granuloma

Poorly structured aggregates, consisting of at least four large epithelioid macrophages (Figure 1b), to slightly organized granulomas (Figure 1c) were observed in 8 (11%) out of 72 animals (Table 2 and Figure 1b). They were present in 1 (9%) out of 11 animals in the undefined susceptibility state group and 7 (28%) out of 25 animals in the infected and potentially susceptible to VL group. The difference in the presence of spleen granulomas between the infected and potentially susceptible to VL and other groups was statistically significant ($\chi^2$-test, $P = 0.0102$).

Cell populations in the red pulp

The frequency of plasma cells was higher in the groups with evidence of *Leishmania* infection than it was in the noninfected group ($\chi^2$-test, $P = 0.0366$, Table 2 and Figure 1j). The distribution of lymphoblasts, granulocytes, resident macrophages and megakaryocytes did not differ significantly among the groups (Table 2).

Parasite burden

Amastigotes were visualized in the spleens of 10 (14%) out of the 72 animals (Table 2); all of these animals belonged to...
Table 2 Perisplenitis and red pulp changes in dogs from an endemic area of *L. chagasi* infection, with different patterns of response to the infection

| Parameters                        | Animal groups                             | Infected |
|-----------------------------------|-------------------------------------------|----------|
|                                   | Potentially resistant to VL (%) | Potentially susceptible to VL (%) | Undefined susceptibility status (%) | Noninfected (%) | Total (%) |
|-----------------------------------|-------------------------------------------|----------|----------------------------------|-----------------|-----------|
| Number of animals                 | 22                                        | 25       | 11                               | 14              | 72        |
| Perisplenitis                     | 2 (10)²                                   | 18 (72)³  | 0 (0)                            | 0 (0)           | 20 (28)   |
| Red Pulp:                         |                                           |          |                                  |                 |           |
| Granuloma                         | 0 (0)                                     | 7 (28)³  | 1 (9)                            | 0 (0)           | 8 (11)    |
| Plasma cells:                     |                                           |          |                                  |                 |           |
| Absent–few                        | 5 (23)                                    | 1 (4)    | 2 (18)                           | 6 (43)³         | 14 (19)   |
| Average–frequent                  | 17 (77)                                   | 24 (96)  | 9 (82)                           | 8 (57)          | 58 (81)   |
| Macrophage:                       |                                           |          |                                  |                 |           |
| Absent–few                        | 13 (59)                                   | 15 (60)  | 0 (0)                            | 4 (29)          | 32 (44)   |
| Average–frequent                  | 9 (41)                                    | 10 (40)  | 11 (100)                         | 10 (71)         | 40 (55)   |
| Lymphoblasts:                     |                                           |          |                                  |                 |           |
| Absent–few                        | 19 (86)                                   | 17 (68)  | 9 (82)                           | 9 (64)          | 54 (75)   |
| Average–frequent                  | 3 (14)                                    | 8 (32)   | 2 (18)                           | 5 (36)          | 18 (25)   |
| Megakaryocyte:                    |                                           |          |                                  |                 |           |
| Absent–few                        | 14 (64)                                   | 19 (76)  | 8 (72)                           | 12 (86)         | 53 (74)   |
| Average–frequent                  | 8 (36)                                    | 6 (24)   | 3 (27)                           | 2 (14)          | 19 (26)   |
| *Leishmania* amastigotes:*        |                                           |          |                                  |                 |           |
| Positive histology:               |                                           |          |                                  |                 |           |
| Sub-capsular                      | 0 (0)³                                    | 10 (40)³ | 0 (0)                            | 0 (0)           | 10 (14)   |
| Red pulp                          | 0 (0)³                                    | 10 (40)³ | 0 (0)                            | 0 (0)           | 10 (14)   |
| Parasite density:                 |                                           |          |                                  |                 |           |
| Sub-capsular                      | 0                                         | 45 ± 40  | 0                                | 0               | 0         |
| Internal red pulp                 | 0                                         | 37 ± 44  | 0                                | 0               | 0         |

²Capsular tissue was absent or poorly represented in one animal. ³Statistically significant differences (*χ²*-test with Yates' correction): ⁴*P* = 0.02; ⁵different from the other groups, *P* < 0.0001; ⁶different from the other groups, *P* = 0.0102; ⁷different from the other groups, *P* < 0.0366; ⁸different from the other groups, *P* < 0.0001; ⁹Statistically different from the parasite density in the internal red pulp (paired *t*-test, *P* = 0.0425). ¹⁰Expressed as the mean of the relative (%) number of high power fields positive for amastigotes among 20–50 examined.

the infected and potentially susceptible to VL group. In all of these animals, amastigotes were seen in both the subcapsular area and the internal red pulp (Figure 1j–k). The parasite burden, measured as the proportion of positive fields among the examined high power fields, was higher in the subcapsular area (45 ± 40%) than the internal red pulp (37 ± 44%; paired *t*-test, *P* = 0.0425). Although parasite distribution in the spleen was usually uneven, three animals with high parasite burden showed evidence of parasites in all of the examined fields (i.e. in the subcapsular area and in the internal red pulp).

**White pulp structural organization**

Distinction of the different regions inside the white pulp of the spleen was absent or poor in 17 (24%) out of 72 animals (Table 3 and Figure 1d–i). The frequency of animals with moderate or extensive structural disorganization of the white pulp areas was higher in the infected and potentially susceptible to VL group (14 out of 25 animals, 56%) than it was in the other groups (χ²-test, *P* < 0.0001).

**Lymphoid follicle frequency**

Lymphoid follicles were present in specimens from all of the animals (Table 3). The frequency of these follicles was rare or low in 36 (50%) out of the 72 animals. The frequency of animals with a low proportion of lymphoid follicles was higher in the infected and potentially susceptible to VL group (20 out of 25, 80%) than it was in the other groups (χ²-test, *P* = 0.001).

**Lymphoid follicle size**

Lymphoid follicles were tiny or small in 38 (53%) out of the 72 animals (Table 3). A higher proportion of animals with small or tiny lymphoid follicles was observed in the infected
and potentially susceptible to VL group (20 out of 25 animals, 80%) than other groups ($\chi^2$-test, $P = 0.0036$). In contrast, the frequency of animals with large follicles in the infected and potentially resistant to VL group (16 out of 22 animals, 73%) was higher that in the other groups ($\chi^2$-test, $P = 0.044$).

### Germinal centre size

Germinal centres were absent or small in 39 (54%) out of the 72 animals (Table 3). The frequency of animals with lymphoid follicles containing small, tiny, or the absence of germinal centres was higher in the infected and potentially susceptible to VL group (21 out of 25 animals, 84%) than other animal groups ($\chi^2$-test, $P = 0.0015$). Conversely, the frequency of animals with average or large germinal centres was higher in the infected and potentially resistant to VL group (16 out of 22 animals, 73%) than the other groups ($\chi^2$-test, $P = 0.0162$).

### Marginal zone size

The marginal zone was absent, tiny, or small in 25 (35%) out of the 72 animals (Table 3). The proportion of animals with marginal zones that were small, tiny, or absent was higher in the infected and potentially susceptible to VL (15 out of 25 animals, 60%) than the other animal groups ($\chi^2$-test, $P = 0.0025$). The marginal zone size was moderately correlated with lymphoid follicle size (Spearman $r = 0.5247$, $P < 0.0001$, Figure 2a) and germinal centre size (Spearman $r = 0.5354$, $P < 0.0001$, Figure 2b).

### Associations between morphological changes in the spleen and emaciation

Evidence of weight loss was present in 27 (39%) out of the 69 animals from which data were recorded. Weight loss was more frequent in the infected and potentially susceptible to VL group (15 out of 25 animals, 60%) than the other animal groups ($\chi^2$-test, $P = 0.0025$). Being underweight was associated with the following alterations in the splenic tissue. First, it was associated with disorganization of the white pulp structure. Among the 27 animals with emaciation, 19 (70%) had some degree of white pulp disorganization; only 18 (43%) out of 42 animals with normal nutritional status had loss of white pulp architecture definition ($\chi^2$-test, $P = 0.0467$). This association between

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### Table 3 White pulp changes in dogs from an endemic area of *L. chagasi* infection, with different patterns of response to the infection

| Parameters                        | Animal groups                        | Infected | Potentially resistant to VL (%) | Potentially susceptible to VL (%) | Undefined susceptibility status (%) | Noninfected (%) | Total (%) |
|-----------------------------------|--------------------------------------|----------|--------------------------------|----------------------------------|------------------------------------|----------------|-----------|
| Number of animals                 |                                      | 22       | 25                             | 11                               | 14                                 | 72             |           |
| Structural disorganization:       | Absent – Slight                      | 21 (95)  | 11 (44)                        | 10 (91)                          | 13 (93)                            | 55             | 76        |
|                                  | Moderate – Extensive                  | 1 (5)    | 14 (56)$^b$                    | 1 (9)                            | 1 (7)                              | 17             | 24        |
| Lymphoid follicle:                | Frequency:                           |          |                                |                                  |                                    |                |           |
|                                  | Rare – Low                           | 6 (27)   | 20 (80)$^b$                    | 2 (18)                           | 8 (57)                             | 36             | 50        |
|                                  | Average – High                       | 16 (73)  | 5 (20)                         | 9 (82)                           | 6 (43)                             | 36             | 50        |
|                                  | Size:                                |          |                                |                                  |                                    |                |           |
|                                  | Tiny – Small                         | 6 (27)   | 20 (80)$^b$                    | 4 (36)                           | 8 (57)                             | 38             | 53        |
|                                  | Average – Large                      | 16 (73)$^b$| 5 (20)                         | 7 (64)                           | 6 (43)                             | 34             | 47        |
| Germinal centre:                 | Size:                                |          |                                |                                  |                                    |                |           |
|                                  | Absent – Small                       | 6 (27)   | 21 (84)$^b$                    | 5 (45)                           | 7 (50)                             | 39             | 54        |
|                                  | Average – Large                      | 16 (73)$^b$| 4 (16)                         | 6 (55)                           | 7 (40)                             | 33             | 46        |
| Marginal zone:                   | Size:                                |          |                                |                                  |                                    |                |           |
|                                  | Absent – Small                       | 5 (23)   | 15 (60)$^b$                    | 2 (18)                           | 3 (21)                             | 25             | 35        |
|                                  | Average – Large                      | 17 (77)  | 10 (40)                        | 9 (82)                           | 10 (79)                            | 47             | 65        |

$^a$–$^g$Statistically different from the other groups ($\chi^2$-test with Yates’ correction): $^aP < 0.0001$; $^bP = 0.0010$; $^cP=0.0036$; $^dP = 0.044$; $^eP = 0.0015$; $^fP = 0.0162$; $^gP = 0.0025$. 
emaciation and disorganization of the lymphoid tissue was observed even in animals with negative splenic culture. Out of 10, 8 (80%) animals who were undernourished in this group had loss of white pulp architecture definition, whereas only 9 (36%) out of 25 without emaciation exhibited this alteration. In addition, being underweight was associated with perisplenitis. This condition was present in 12 (46%) out of 26 animals with emaciation, but in only 6 (15%) out of 41 animals with normal nutritional status ($\chi^2$-test, $P = 0.0107$).

DISCUSSION

Changes in lymphoid organs associated with VL have been previously reported in various studies. However, the current study presents a systematic view of changes in different compartments of the spleen in naturally infected dogs from an endemic area of VL, and describes different patterns of immunological responses to the parasite. In a previous study, we have shown that negative LST results were associated with high levels of serum antibody activity and emaciation in L. chagasi-infected dogs. We have now shown that infected animals with negative LST also have (i) increased frequencies of perisplenitis, granuloma, and parasites in the red pulp (ii) a variety of white pulp alterations, such as atrophy and loss of structural definition (iii) lymphoid follicles that are reduced in number and size, and (iv) germinal centres and marginal zones that are frequently small or absent. These animals, with loss of lymphoid follicle definition, perisplenitis, and a high parasite density in the spleen (in which parasites were seen by conventional histology), were also more frequently underweight; this association likely reflects the status of the clinical disease. Furthermore, we observed that the distribution of parasites in the spleen was not homogeneous. In heavily infected animals, parasites were distributed almost continuously in the red pulp. In animals with a less intense parasite burden, a discontinuous distribution of infected cells was observed, predominating in the subcapsular area.

Perisplenitis is a common finding in canine VL (26,27). Although the increase in spleen size may predispose that organ to trauma or mechanical stress, the actual genesis of perisplenitis in leishmaniasis remains unknown. It is interesting, however, that the perisplenitis was associated with the presence of parasite-containing macrophages in subcapsular areas. The presence of such macrophages in capsular and subcapsular inflammatory infiltrates has also been observed in other studies (26,27). These findings may indicate a role for inflammation in parasite dissemination (28,29), or that the parasite elicits an inflammatory reaction.

In light of previously published data regarding immunity to Leishmania, it seems contradictory that splenic granulomas were more frequently found in the group of infected animals with negative LST than in infected animals with positive LST. In effect both, granuloma and positive LST, may result from DTH reactions to Leishmania antigens. The granulomas observed in the spleen of the animals in this study varied from poorly structured aggregates, consisting of at least four large epithelioid macrophages (Figure 1b), to the barely organized structure shown in Figure 1(c). Although parasites were not usually seen in the interiors of epithelioid cells, amastigotes were more frequently observed in the spleen of animals with granulomas. Hence, as observed in humans with long duration VL (30), these granulomas appear to reflect a persistent chronic inflammation in response to uncontrolled infection, rather than protection against the disease. Although granulomas may constitute a step towards infection control, the conditions in their interior may be
subverted by a parasite to favour its survival and growth (31). In fact, interleukin-10 and other cytokines produced by granuloma cells may provide conditions for the survival of *Leishmania* (12) and other microorganisms (32,33). The fact that granulomas were only present in animals with positive spleen culture in our study supports the view that these granulomas may reflect a state of inefficient control of *Leishmania* infection by the host immune system.

The increase in the plasma cell population in lymphoid organs is a common finding in leishmaniasis (7,27,34), and it may be present at relatively early stages of the disease (35). In all of the infected groups examined in this study, a significant increase in the plasma cell population was observed to replace other cell populations in areas of the red pulp. A progressive replacement of T-cell areas in the white pulp was also observed in some animals (data not shown). The determinants of the changes in number or distribution of plasma cells in VL are poorly understood. However, in visceral leishmaniasis, several factors like the well-characterized polyclonal B-cell activation (36,37) and production of cytokines [e.g. IFN-γ, IL10 (38–41), and IL6 (42,43)] and chemokines (e.g. CXCL12) may contribute to plasma cell differentiation and retention in the red pulp (44). Further studies are necessary to define the mechanisms involved in the plasma cell replacement of normal cell populations in the spleen, and future work should also examine the potential contribution of cell population replacement to both the increase of polyclonal immunoglobulin levels observed in VL (45) and disease development.

Lymphoid hyperplasia, atrophy, and disorganization were all present in the infected dogs examined in this study. Hyperplasia was common in animals from the infected and potentially resistant to VL group, and atrophy and lymphoid tissue disorganization were most frequently present in the animals from the infected and potentially susceptible to VL group. Hyperplasia followed by atrophy of the lymphoid follicles may occur in the spleen during the course of VL (7,34,35). However, such atrophy and hyperplasia are not uniform in all compartments of the white pulp. As shown in Figure 2, marginal zone hyperplasia may occur with a variable degree of follicle atrophy. This finding, together with the effacement of the boundaries between usually distinct areas of the lymphoid tissue, indicates that structural disorganization is present in the white pulp, as suggested by Veress *et al.* (1977). The loss of the architectural structure of splenic tissue has been the subject of recent studies performed in mice infected with *Leishmania* (20) or other pathogens (21). A variety of changes, including the loss of cell populations and the impairment of cell migration to the different splenic compartments, have been observed in murine models of visceral leishmaniasis induced by TNF and IL-10 (20,46,47). We also observed that disorganization of the splenic lymphoid tissue was more frequent in animals with emaciation, even for those animals without evidence of *Leishmania* infection, than it was in animals without evidence of undernourishment. In these emaciated animals, weight loss may result from insufficient access to food, which, in its turn, may be associated with other infections (9,48,49). For instance, studies on experimental models have shown that viral infections may also cause splenic lymphoid tissue disorganization (21). A factor common to many infections, such as high levels of TNF (unfortunately not tested in this study), may account both for emaciation and for lymphoid tissue disorganization (20,50). Nevertheless, the fact that the intensity of white pulp disorganization was higher in animals with splenic parasitism than in those without indicates that an active *Leishmania* infection may play a causal role in a spleen disorganization process. Further studies are necessary to clarify the mechanisms involved in the structural disorganization of the splenic tissue in canine leishmaniasis and assess the potential role of such splenic changes in mediating increased susceptibility to bacterial infections in the late stages of the disease.

In conclusion, our study shows that natural infection by *L. chagasi* is associated with splenic architecture disruption, which is characterized by disorganization of normal lymphoid tissue, loss of normal spleen leucocyte diversity via replacement of leucocytes by plasma cells, and eventual atrophy of the lymphoid tissue. Studies examining the redistribution of the main cell populations, the profile of cytokine expression in the spleen, and the potential association of splenic alterations with increased susceptibility to bacterial infection in these animals are currently underway.

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