The Role of Inflammatory, Anti-Inflammatory, and Regulatory Cytokines in Patients Infected with Cutaneous Leishmaniasis in Amazonas State, Brazil

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The authors discuss in this paper the role of inflammatory, anti-inflammatory, and regulatory cytokines in patients infected with different species of Leishmania in Amazonas State, Brazil. A comparative analysis was made of serum concentrations of these cytokines in the peripheral blood of 33 patients infected with cutaneous leishmaniasis. The isolates were identified as Leishmania guyanensis, L. naiffi, and L. amazonensis. Most (64%) of the patients were male ranging in age from 18 to 58 years. Protein expression profiles of IL-2, IL-4, IL-6, IL-10, IFN-γ, TNF-α, and IL-17 cytokines were shown to vary significantly between infected and noninfected (control group) individuals and according to the Leishmania species. Infection caused by L. guyanensis accounted for 73% of the cases and patients with this parasite also showed higher concentrations of IL-2, IFN-γ, IL-4, and IL-17 when compared to infection by L. amazonensis. Patients with infection caused by L. naiffi showed higher concentration of the cytokines analyzed when compared to uninfected patients; however, there was no statistically significant difference with the other species analyzed.

1. Introduction

Leishmaniasis is endemic in several parts of the world, with a global prevalence of over 12 million cases. Divided in two main groups, leishmaniasis can affect the skin (cutaneous leishmaniasis/CL) or internal organs (visceral leishmaniasis/VL). There are 1.5 million new cases of cutaneous leishmaniasis emerging every year [1]. The infection is caused by protozoan parasites of the genus Leishmania, transmitted by the bite of the female sand fly vector. Several Leishmania species are able to cause a wide spectrum of clinical manifestations of CL, ranging from the mild cutaneous form (localized cutaneous leishmaniasis, LCL) to multiple non ulcerative nodules (diffuse cutaneous leishmaniasis, DCL) and the disfiguring mucosal form (mucocutaneous leishmaniasis, MCL) [2]. In Brazil, American cutaneous leishmaniasis (ACL) is notable for its wide distribution, occurring in all states. In the State of Amazonas, in 2011 alone, 2,230 new cases were notified and the transmission occurred mainly in the cities of Manaus (752 cases), Presidente Figueiredo (213), and Rio Preto da Eva (203) [3]. In the Amazon Basin, seven species of
Leishmania have been identified as causative agents of ACL. Six such species belong to the subgenus Viannia, Leishmania (V.) braziliensis, L. (V.) guyanensis, L. (V.) lainsoni, L. (V.) naiffi, L. (V.) shawi, and L. (V.) lindenbergi, and one to the subgenus Leishmania, Leishmania (Leishmania) amazonensis [2, 4]. However, thus far, the State of Amazonas, only L. guyanensis, L. naiffi, L. braziliensis, and L. amazonensis have been isolated from humans, with L. guyanensis being the predominant species [5, 6].

Clinical manifestations of ACL vary widely, from asymptomatic infections to small lesions limited to or contained within a small area of the skin, to disseminated ulcers and the mutilating mucocutaneous forms [2, 7, 8]. The pathogenesis of ACL is strongly influenced by factors inherent to the host (such as genetic and immune response), the parasite (such as virulence of Leishmania species infecting), and vector (as its vectorial capacity). As a result of this interaction between different species of the parasite, the vector, and the mechanisms of the immune response of the vertebrate host, a spectrum of clinical, histopathological, and immunopathological manifestations is observed in humans [2, 9, 10]. Cutaneous leishmaniasis may develop into spontaneous healing or progress to the formation of papules, nodules, plaques, and especially ulcers. Invasion of peripheral lymph nodes and progression to mucosal lesions may also occur. The types of immune response mounted by the individual, crucial for eliminating the parasites, drive such outcomes. Such response, however, is dependent upon host genetic factors and the species of Leishmania responsible for infection [2, 8]. Factors associated with the immune response include tumor necrosis factor (TNF); an effective response of natural killers (NK) cells against interleukin-12 (IL-12); and an appropriate production of interferon gamma (IFN-γ) among other cytokines that might be regulated by the Leishmania responsible for infection [11, 12].

Generally, treatment for CL in Brazil is based on antimicrobial chemotherapy using N-methylglucamine antimoniate (Glucantime, Sanofi-Aventis) [13] and may depend on the results of controlled studies of different dosages employed [9]. Moreover, the response to antimicrobial therapy is likely influenced by the infecting Leishmania species [14, 15].

According to the diversity and biological complexity of the parasite, the vector, and the host cellular immune response in ACL, different clinical forms may develop, with Leishmania able to direct the production of cytokines and chemokines, cell differentiation, and immune response profile that can lead to cure or persistent disease. Thus, an understanding of the immune response by patients to the infecting Leishmania and how effective the current therapy is to the different species of the parasite circulating in the Amazon are of paramount importance for disease control. Hence, the main purpose of this study was to compare levels of inflammatory, anti-inflammatory, and regulatory cytokines in the sera of patients infected by distinct leishmanial parasites and to evaluate the success of the antimicrobial therapy in 33 patients with ACL treated at a Primary Health Care and the Hospital of Rio Preto da Eva municipality, Amazonas State.

2. Materials and Methods

2.1. Study Area. The city of Rio Preto da Eva, State of Amazonas, is located near the 80 Km marker on AM-010 Highway (3° 07'06.5" S, 59° W), 575 Km from the state capital, Manaus, and it is part of the Metropolitan Area of Manaus. The vegetation is tropical moist forest upland, with great diversity of species.

2.2. Clinical Cases. All cases of suspected ACL were examined at the Hospital Thomé de Medeiros Raposos and at the Basic Health Unit Manoel Ruminão in Rio Preto da Eva, Amazonas. Positive diagnosis in each case was made by direct examination of skin fragments obtained by scraping the edge of skin lesions, followed by staining using a Panoptic kit (LB Laborclin) and observation of Leishmania amastigotes under optical microscope. Treatment was performed as recommended by the Brazilian Health Ministry: 10–20 mg/Kg/day (15 mg/Kg/day) for 20 days [16]. A skin fragment obtained from the lesion of each patient also was seeded into biphasic NNN blood agar medium [17, 18] and later expanded in liquid Schneider’s Drosophila complete medium pH 7.2 (GIBCO-BRL, Gaithersburg, MD) supplemented with 10% heat-inactivated Fetal Calf Serum (iFCS). Following expansion, parasites were prepared for biochemical identification and cryopreservation. Stocks from each parasite species isolated were preserved in liquid nitrogen using 8% glycerol in Schneider’s medium containing 30% iFCS. The study was conducted in accordance with the requirements of the National Health Council (resolution 196/96) and was approved by the INPA’s ethics committee on research involving human under protocol 193/2008. Patients included in the study were those diagnosed with ACL caused by different species of Leishmania and a control group without any symptoms of Leishmania infection and no history of previous infection (negative control), matched for sex and age. All study participants signed the informed consent form (ICF) agreeing to participate in the study.

2.3. Identification of Leishmania Isolates. The strains isolated were typed by multilocus enzyme electrophoresis using eight enzymatic systems: malate dehydrogenase (MDH, EC1.1.1.37), isocitrate dehydrogenase (IDH, EC1.1.1.42) with substrate NAD and NADP, malic enzyme (ME, EC1.1.1.40), glucose-6-phosphate dehydrogenase (G6PDH, EC1.1.1.49), 6-phosphogluconate dehydrogenase (6GPDH, EC1.1.1.44), Aconitate hydratase (ACON, EC4.2.1.3), and hexokinase (HK, EC2.7.1.1) [19]. Species identification was performed by comparing enzyme profiles with reference strains of Leishmania (Viannia) braziliensis (MHOM/BR/1975/M2903), L. (V.) guyanensis (MHOM/BR/1975/ M4147), L. (V.) naiffi (MDAS/BR/1979/M5333), and Leishmania (Leishmania) amazonensis (IFLA/BR/1967/PH8).

2.4. Peripheral Blood Samples. A total of 5 mL of peripheral blood was collected from 33 confirmed cases of ACL as
Table 1: Clinical and epidemiological data from patients* infected with Leishmania spp. from Rio Preto da Eva, State of Amazonas, BR.

| Leishmania isolates | Patients data | Infection*  |
|---------------------|---------------|-------------|
| Designations1       | Species2      | Sex | Age (years) | Length (days) | Status | Number of lesions | Lesion size (mm) |
| MHOM/BR/09/IM5553   | L. naiffi     | M   | 34          | 120           | P      | 1               | ND              |
| MHOM/BR/09/IM5562   | L. naiffi     | M   | 26          | 160           | P      | 2               | 2 × 2           |
| MHOM/BR/09/IM5584   | L. amazonensis| M   | 58          | 45            | P      | 1               | 8 × 8           |
| MHOM/BR/10/IM5637   | L. guyanensis | F   | 31          | 60            | P      | 1               | 17 × 18         |
| MHOM/BR/10/IM5641   | L. guyanensis | F   | 41          | 150           | S      | 3               | 21 × 15         |
| MHOM/BR/10/IM5653   | L. guyanensis | M   | 18          | 90            | P      | 1               | 30 × 20         |
| MHOM/BR/10/IM5657   | L. guyanensis | M   | 56          | 37            | P      | 1               | 19 × 17         |
| MHOM/BR/10/IM5665   | L. naiffi     | M   | 23          | 90            | P      | 3               | 8 × 6           |
| MHOM/BR/10/IM5668   | Leishmania sp.| M   | 41          | ND            | S      | 1               | ND              |
| MHOM/BR/10/IM5690   | L. guyanensis | F   | 58          | 60            | P      | 1               | 17 × 18         |
| MHOM/BR/10/IM5692   | L. guyanensis | F   | 30          | 15            | P      | 2               | 60 × 60         |
| MHOM/BR/10/IM5694   | L. guyanensis | M   | 29          | 14            | P      | 1               | 20 × 23         |
| MHOM/BR/11/IM5697   | L. guyanensis | M   | 19          | 14            | P      | 1               | 13 × 16         |
| MHOM/BR/11/IM5749   | L. guyanensis | M   | 38          | 30            | P      | 1               | 17 × 14         |
| MHOM/BR/11/IM5752   | L. guyanensis | M   | 34          | 21            | P      | 4               | 11 × 12         |
| MHOM/BR/11/IM5773   | L. guyanensis | M   | 30          | 10            | P      | 1               | 32 × 21         |
| MHOM/BR/11/IM5775   | L. guyanensis | M   | 24          | 21            | P      | 4               | 8 × 5           |
| MHOM/BR/11/IM5828   | L. guyanensis | M   | 43          | 150           | P      | 1               | 11 × 8          |
| MHOM/BR/11/IM5833   | L. guyanensis | F   | 21          | 30            | P      | 1               | 70 × 70         |
| MHOM/BR/11/IM5869   | L. guyanensis | M   | 24          | 60            | P      | 8               | 60 × 60         |
| MHOM/BR/11/IM5875   | L. guyanensis | M   | 49          | 30            | P      | 1               | 11 × 7          |
| MHOM/BR/11/IM5894   | L. guyanensis | M   | 27          | 30            | S      | 1               | 20 × 12         |
| MHOM/BR/11/IM5950   | L. amazonensis| F   | 30          | 30            | P      | 1               | 25 × 26         |
| MHOM/BR/11/IM5955   | L. amazonensis| F   | 29          | 15            | P      | 1               | 60 × 90         |
| MHOM/BR/11/IM5962   | Leishmania sp.| M   | 30          | 60            | S      | 2               | 6 × 5           |
| MHOM/BR/11/IM5969   | Leishmania sp.| F   | 30          | 15            | S      | 1               | 10 × 7          |
| MHOM/BR/11/IM5976   | L. guyanensis | F   | 18          | 21            | P      | 1               | 17 × 13         |
| MHOM/BR/11/IM5985   | L. guyanensis | F   | 25          | 15            | P      | 1               | 12 × 12         |
| MHOM/BR/12/IM6006   | Leishmania sp.| M   | 33          | 180           | S      | 1               | ND              |
| MHOM/BR/12/IM6016   | Leishmania sp.| F   | 30          | 120           | S      | 04              | 11 × 11         |
| MHOM/BR/12/IM6025   | Leishmania sp.| M   | 31          | 7             | S      | 1               | 1 × 1           |
| MHOM/BR/12/IM6034   | L. amazonensis| M   | 42          | 21            | P      | 1               | 29 × 15         |
| MHOM/BR/12/IM6038   | Leishmania sp.| F   | 48          | 30            | S      | 1               | 17 × 17         |

*All biological material was collected by healthcare professionals at the Joint Health Unit Thomé de Medeiros Raposo and the Basic Health Unit Manoel Rumião, Municipality of Rio Preto da Eva.

1Designations: host [M = Mammalia; HOM = Homo sapiens]/country of origin/year of isolation/original code used by INPA; 2stock identification was established by enzyme electrophoresis; 3length of infection indicates the approximate time in days when patients recalled appearance of symptoms or lesion; number of lesion = number of ulcerated skin lesion in a patient; infection status refers to either primary (P) or secondary (S) infection; lesion size = measurements (L × W) of lesions. ND: not determined.

well as from 19 noninfected controls using BD vacutainer tubes (BD Biosciences). Sera were separated by centrifugation at 1,500 x g for 15 minutes at 4°C and stored at −80°C. All serum samples were tested for infections with HIV (ELISA HIV Tetra, Biotest/Diasorin), HTLV (Murex HTLV-I+II kit, Abbott), and Trypanosoma cruzi (ELISA Chagas III, BIOSChile).

2.5. Measurement of Cytokine Levels in Sera. Levels of IL-2, TNF-03B1, IFN-03B3, IL-4, IL-6, IL-10, and IL-17 cytokines were compared among the 33 serum samples obtained from patients with confirmed ACL caused either by L. amazonensis, L. naiffi, or L. guyanensis and from unidentified Leishmania sp. Cytokine levels were measured using a FACSCanto II flow cytometer (BD Biosciences, San Jose, CA, USA) and the BD Cytometric Bead Array (CBA) Human Inflammatory Cytokine Kit for TH1/TH2/TH17 (BD Biosciences, San Diego, CA, USA). For each cytokine assessed, the concentration (in pg/mL) and the mean fluorescence intensity (MFI) were calculated using the Array-FCAP software (v3.0.1). The
results were analyzed using GraphPad Prism (v6.0) and nonparametric tests (Kruskal-Wallis and Mann-Whitney). $P$ values <0.05 were considered statistically significant.

3. Results

Twenty-four out of the 33 peripheral blood samples were of cases representing primary *Leishmania* infections and nine were collectively referred to as secondary. This latter group was comprised of patients who had either relapse or were infected a second time. In all patients, skin lesions varied in terms of size, appearance, and number of lesions depending on the infecting *Leishmania* species, as shown in Table 1. Nineteen peripheral blood samples were collected from noninfected individuals (control group). All patients also tested negative for HIV, HTLV, and *T. cruzi*.

In the group of patients referred to as secondary, seven (approximately 78%) displayed a single skin lesion, and one patient (approximately 11%) displayed two lesions in the lower limbs. Most (66%) were unable to or did not indicate the type of treatment received at the time of their first infection. Three patients (33%) confirmed previous antimonial treatment received. All nine patients did recall the presence of typical skin lesion characteristic of *Leishmania* infection, which was confirmed by the attending physician at the time.

Infections with *L. guyanensis* accounted for 19 (73%) of the cases and were followed by infections with *L. amazonensis* with 12% (four individuals) and *L. naiffi* with 9% (three cases). In seven patients (representing 21% of the total number of cases) the infecting *Leishmania* species was not identified. Information of clinical and epidemiological details of the 33 ACL cases is summarized in Table 1. Twenty-one (64%) individuals were males and 12 (36%) were females and ranging from 18 to 58 years in age. Although single skin lesions were observed in the majority of cases (24 individuals or 69%), eight skin lesions were observed in a single individual. The estimated time for the evolution of disease (i.e., appearance of skin lesion) varied widely from seven to 180 days (Table 1) and was based on individual patient account. In terms of their occupation, 100% of the patients were indicated to be farmers. Patient information was obtained via a survey administered individually.

### 3.1. Measured Levels of Cytokine in Patient Sera

The levels of inflammatory and Th1 profile cytokines IL-2, TNF-$\gamma$, and IFN-$\gamma$ observed in the serum samples from the *Leishmania*-infected patients differed significantly from the noninfected controls, as shown in Figure 1. Similar comparisons between patients infected with different *Leishmania* indicated that infection with *L. guyanensis* led to higher serum levels of both IL-2 ($P = 0.0002$) and IFN-$\gamma$ ($P = 0.0412$) than those from patients infected with *L. amazonensis*. For infections with *L. naiffi*, higher levels of TNF-$\gamma$ and IFN-$\gamma$ were consistently detected in comparison to infections with *L. guyanensis* and *L. amazonensis*, though no statistical differences were observed. Higher IFN-$\gamma$ levels also were observed on infection by *L. guyanensis* compared to other species.

For the regulatory cytokine IL-10, and for the Th12 and Th17 cytokines such as IL-4, IL-6, and IL-17, our results indicate significant differences when comparing all primary infections with *Leishmania* spp. with noninfected controls. Assessed IL-10 levels were significantly higher in the sera of infected individuals as compared to controls ($P = 0.0002$). Similar observations were made with regard to the levels of IL-4 ($P = 0.0016$), IL-6 ($P < 0.0001$), and of IL-17 ($P = 0.0002$). A significant difference also was observed between the subgroup (i.e., distinct *Leishmania* species) analyzed: higher levels of IL-4 ($P = 0.0132$), IL-10 ($P = 0.0085$), and IL-17 ($P = 0.0042$) were detected in patients with infections caused by *L. guyanensis* than those infected with *L. amazonensis* (Figure 2). Meanwhile, IL-6 levels showed no significant difference in the analysis between the species studied, in spite of being higher in samples from patients infected with *L. guyanensis*.

Seric cytokine levels were generally higher in samples of primary infections as compared with sera obtained from patients with secondary *Leishmania* infection and the control samples. The analysis of Th1 cytokine profile revealed significant differences between infected (primary or secondary infections) and noninfected controls for for IL-2 and IFN-$\gamma$ ($P < 0.0001$ for both cytokines) and TNF-$\gamma$ ($P = 0.0009$). When comparing the concentrations of these cytokines in the patients with secondary *Leishmania* infection no significant differences between the two groups were observed, despite higher serum levels of IL-2, IFN-$\gamma$, and TNF-$\gamma$ detected in primary infections. Levels of TNF-$\gamma$ were lower in secondary than in primary infections but again this difference was not statistically significant ($P = 0.2100$, Figure 3).

The comparative analysis between the primary and secondary infections and the noninfected controls revealed significant differences for IL-6, IL-17, and IL-10 (Figure 4). For Th12 and Th17 regulatory cytokines, significant differences in the serum concentrations of IL-4 ($P = 0.0086$) and IL-10 ($P = 0.0297$) were detected between the primary and the secondary infections. For IL6 (0.1200) and IL17 (0.3718) such differences were not statistically significant.

4. Discussion

The ACL is characterized by a spectrum of cutaneous manifestations. The classical clinical symptom of the disease is the appearance of an erythematous nodule, which may be single or multiple, usually located at the exposed region in the tegument. In this study we assessed the levels of selected Th1, Th12, and Th17 cytokines in the sera of patients infected with different species of *Leishmania*. Patients had been positively diagnosed with cutaneous leishmaniasis and had not yet started treatment with the antimonial drug Glucantime (intravenous or intramuscular applications of 10–20 mg/kg/day for 20 consecutive days) according to guidelines from the Brazilian Ministry of Health [16].

Cytokines have a central role in the immune response, presenting both local and systemic effects [20]. The balance in the production of inflammatory and regulatory cytokines determines the profile of immune response and influence on disease severity [21–23]. In mouse models, resistance to infection or healing is associated with a Th1 response that
is dependent upon expression of IL-12, IFN-γ, and TNF-α with production of nitric oxide (NO) [22, 24–26]. In contrast, susceptibility to disease or the inability to control lesion development is associated with a regulatory response with the concomitant production of IL-10 as well as expression of T\(_{H2}\) cytokines such as IL-4 and TGF-β [27–29].

Cytokines and precursors of T\(_{H1}\) profile such as IL-2, IFN-γ, and TNF-α were present at higher concentrations during the primary *Leishmania* infections when compared to controls. These cytokines stimulate macrophage activity that plays a crucial role in parasite killing [30, 31]. In the comparative analysis between the *Leishmania* species identified in this study, higher concentration of IL-2 and IFN-γ in the sera of patients with primary infection, compared to sera from noninfected or those with secondary *Leishmania* infection, was observed. Moreover, levels of IL-2 and IFN-γ also were higher in the primary infection group when infected with *L. guyanensis*.

Many cell types, including T cells, dendritic cells, macrophages, and NK cells, produce IFN-γ. IFN-γ limits the growth of *Leishmania* in human and murine macrophages by stimulating nitric oxide synthesis in macrophages [32]. In murine models, the modulation of local or systemic levels of IFN-γ is a critical determinant in the resolution of infection. TNF-α produced mainly by macrophages and in concert with IFN-γ also leads to induction of nitric oxide synthesis and consequently the death of amastigotes inside macrophages [33]. In 2007, Pinheiro and Rossi-Bergmann [34] indicated that IFN-γ is necessary for the control of *infection by L. major* reducing the expansion of T\(_{H2}\) cells in experimental infections caused by this species. These authors also suggested that IFN-γ role remains distinct in infections with *L. major* or *L. amazonensis* and that in *L. amazonensis* IFN-γ is effective in the first week of infection. In addition, IL-10 has an important role with regard to IFN-γ, both in terms of expression control as well as induction
of IFN-γ effectors’ function, thus determining the course of *Leishmania* infection [35].

Regarding TNF-α, it has been shown that, in the sera of patients with active *L. major* infection or in the sera of patients who received treatment and relapsed, levels of this cytokine do change significantly when compared to a control group [36]. Although that study did not indicate whether these were primary or secondary infections, low levels of TNF-α appear to be associated with disease severity during *Leishmania* infection [36]. Here, significantly higher levels of TNF-α were detected in primary infections but did not significantly differ between secondary and controls, though were generally lower. Low levels of TNF-α are associated with persistence of infection and greater difficulty in resolution of skin lesion [36]. Our results also support this view.

For IL-10, IL-4, IL-6, and IL-17, significant increases in the measurable levels of these cytokines were observed in the infected group compared to controls. In addition, patients with ACL caused by *L. amazonensis* displayed lower levels of sera IL-4 and IL-17 when compared to infection by *L. guyanensis*. IL-17 is known to play a role in inflammation, in autoimmune diseases, and in cancer [37–39]. IL-17 has multiple effects, acting both in the induction of proinflammatory cytokines and in the recruitment and activation of leukocytes. Although its role in CL is not fully understood, this cytokine seems to play an important role in the pathogenesis of the disease [40, 41]. Bacellar et al. [40] observed that IL-17 is produced during infection with *L. braziliensis* and lymphocytes obtained from patients with MCL and CL yielded higher concentrations of IL-17 compared to lymphocytes obtained from uninfected individuals. The presence of T<sub>H</sub>17 cells in lesions of CL/MCL was associated with neutrophils and tissue destruction [40, 42]. Our data point to higher levels of T<sub>H</sub>17 in the sera of patients infected with *L. guyanensis* when compared to the other two species identified (*L. amazonensis* and *L. naiffi*). Nevertheless, no significant differences were

Figure 2: Levels of IL-4, IL-6, IL-10, and IL-17 in serum samples from patients infected with different *Leishmania* species and in noninfected control. Cytokine levels (pg/mL) were calculated for each serum sample. Means (horizontal lines) are shown for each group. Ctrl: noninfected control; L.g: *L. guyanensis*; L.n: *L. naiffi*; L.a: *L. amazonensis*. 
observed between the sera of primary infections and sera of patients reinfected (secondary infection) with *L. guyanensis*.

IL-6 is produced by many cell types and is involved in acute-phase response in B cell maturation and differentiation of macrophages. IL-6 promotes Th2 differentiation and simultaneously inhibits Th1 via independent molecular mechanisms. In this work the primary infected patients had high serum concentrations of IL-6 when compared to controls (*P* < 0.0001); the same was not observed in the analysis of patients with secondary infection. In the group of patients with a secondary infection, IL-6 levels were slightly lower than in the primary infection group, but no statistical difference was observed between the two groups. There was no difference in the concentration of IL-6 in the comparison among the three species.

In the murine model, it was demonstrated that the modulation of the immune response by IL-4 produced mainly by T lymphocytes led to the regulation of specific Th1 cytokines, such as IFN-γ and IL-12 [43]. As suggested by Hurdayal and Brombacher [44], depending on the model utilized, IL-4 appears to be a “double-edged sword” driving a susceptible Th2 response and mediating a Th1 response during disease. In our study, patients with primary infection caused by parasites of the genus *Leishmania* (Figure 4) displayed higher levels of IL-4 (*P* = 0.0086) when compared to individuals with secondary infection, suggesting an activation of IL-4 during the first contact with the parasite and its presence in the clinical course of infection caused by various species of *Leishmania*. It was also observed that, in primary infections with *L. amazonensis*, the levels of IL-4 were not as elevated as in infections with *L. guyanensis* or *L. naiffi* when compared with the control group (Figure 2).

The IL-10 plays regulatory role in the immune response, by inhibiting inflammatory mediators and activation of monocytes [45, 46]. It has been demonstrated that the production of this cytokine by CD4+ CD25 T cells (Tregs)
is associated with persistence of the parasite and reactivation of the disease [47–49]. IL-10 secretion by T_{H}1 cells likely is a mechanism that minimizes self-regulatory immunopathology of ACL [50, 51]. We observed lower levels of IL-10 in the secondary infection group of patients in comparison with primary infections. Furthermore, patients infected with *L. guyanensis* had higher IL-10 levels than patients infected with *L. amazonensis* (P = 0.0085). These results suggest a profile of mixed T_{H}1/T_{H}2 response during infection with *L. guyanensis*. Interestingly, we also observed that three individuals infected with *L. guyanensis* (MHOM/BR/11/IM5749; MHOM/BR/11/IM5752; MHOM/BR/11/IM5775) displayed higher IL-10 levels in comparison with other individuals infected with the same *Leishmania* species. Two of these patients with primary infections (MHOM/BR/11/IM5752; MHOM/BR/11/IM5775) had a total of four skin lesions and progression of the disease in about 21 days. It has been shown that, prior to treatment, patients infected with *L. braziliensis* have higher levels of IFN-γ and decreased production of IL-10 [52].

Currently, there is no mechanism available to predict whether or in whom the most serious forms of CL may develop [53]. The results presented here suggest that certain cytokines, such as IL-2, IL-4, IL-6, IL-10, IFN-γ, TNF-α, and IL-17, are involved in the clinical course of human CL in the Amazon and are likely associated with disease outcome, being responsible for either complete healing or remission or on the other hand the development of relapses and more severe forms of the disease.

**Conflict of Interests**

The authors declare that there is no conflict of interests regarding the publication of this paper.

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