



miR-548d-3p Is Up-Regulated in Human Visceral Leishmaniasis and Suppresses Parasite Growth in Macrophages

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Visceral leishmaniasis caused by *Leishmania (Leishmania) infantum* in Latin America progress with hepatosplenomegaly, pancytopenia, hypergammaglobulinemia, and weight loss and maybe lethal mainly in untreated cases. miRNAs are important regulators of immune and inflammatory gene expression, but their mechanisms of action and their relationship to pathogenesis in leishmaniasis are not well understood. In the present study, we sought to quantify changes in miRNAs associated with immune and inflammatory pathways using the *L. (L.) infantum* promastigote infected- human monocytic THP-1 cell model and plasma from patients with visceral leishmaniasis. We identified differentially expressed miRNAs in infected THP-1 cells compared with non-infected cells using qPCR arrays. These miRNAs were submitted to *in silico* analysis, revealing targets within functional pathways associated with TGF- β , chemokines, glucose metabolism, inflammation, apoptosis, and cell signaling. In parallel, we identified differentially expressed miRNAs in active visceral leishmaniasis patient plasma compared with endemic healthy controls. *In silico* analysis of these data indicated different predicted targets within the TGF- β , TLR4, IGF-I, chemokine, and HIF1 α pathways. Only a small number of miRNAs were commonly identified in these two datasets, notably with miR-548d-3p being up-regulated in both conditions. To evaluate the potential biological role of miR-548d-3p, we transiently transfected a miR-548d-3p inhibitor into *L. (L.) infantum* infected-THP-1 cells, finding that inhibition of miR-548d-3p enhanced parasite growth, likely mediated through reduced levels of MCP-1/CCL2 and nitric oxide production. Further work will be required to determine how miR-548d-3p plays a role *in vivo* and whether it serves as a potential biomarker of progressive leishmaniasis.

Keywords: *Leishmania (Leishmania) infantum*, microRNA, visceral leishmaniasis, THP-1 cells, pathogenesis, inflammation

INTRODUCTION

The leishmaniasis are vector-borne diseases caused by protozoan parasites of the order Kinetoplastida, family Trypanosomatidae, and genus *Leishmania*. The leishmaniasis are endemic in 98 countries, affecting 0.9 to 1.6 million people globally each year, and are considered the second most important of the neglected diseases caused by protozoa (Bi et al., 2018; Burza et al., 2018; Sasidharan and Saudagar, 2021; WHO, 2021). Transmission occurs when sand flies inject metacyclic promastigotes into the skin. These parasite forms enter phagocytic myeloid cells, transform into amastigotes, and then proliferate, establishing the infection. The diversity of *Leishmania* parasites and vector species and host genetic and immunological conditions lead to different clinical presentations. The three main clinical forms of the disease are cutaneous, mucosal, and visceral leishmaniasis (VL). Human active VL is a severe disease caused by either *Leishmania (Leishmania) donovani* or *L. (L.) infantum* and affects organs rich in mononuclear phagocytes. Active VL manifests with fever, hepatosplenomegaly, pancytopenia, hypergammaglobulinaemia, and significant weight loss and may be lethal mainly in untreated cases (Burza et al., 2018; Ibarra-Meneses et al., 2020; Serafim et al., 2020). Most cases occur in Brazil, East Africa, and India. An estimated 50 000 to 90 000 new cases of VL occur worldwide annually (WHO, 2021).

Data from endemic areas for VL in Brazil indicate that *L. infantum* efficiently induces lifelong protective immunity in most infected individuals that remain asymptomatic (Badaro et al., 1986; Silveira et al., 2009), highlighting the still need to search for factors and mechanisms leading to disease development. Among factors leading to the development of active disease, an immunosuppressive state resulting from either protein-energy malnutrition (Malafaia, 2009) or HIV infection (Lindoso et al., 2014) rank highest. However, participation of other factors gains evidence, including lipoprotein and lipoprotein fraction levels (Carvalho et al., 2014) and growth factors (Reis et al., 2021), and inhibition of secretion of pro-inflammatory cytokines and chemokines (Tasew et al., 2021), along with alterations in the adaptive immune response (Saporito et al., 2013; Palacios et al., 2021).

In active human VL, studies using peripheral blood mononuclear cells (PBMCs) have identified *Leishmania* antigen-specific T cell suppression associated with increased abundance of Transforming growth factor β (TGF- β), interleukin (IL-)10, IL-35 and apoptosis of CD4⁺ T cells. In contrast, studies directly analyzing lymphoid tissues where parasites proliferate, such as bone marrow and spleen, suggest intense immune activation in active VL, including abundant mRNA for tumor necrosis factor α (TNF- α) and interferon γ (IFN- γ) (Goto and Prianti, 2009) and consequently high serum IFN- γ levels (de Medeiros et al., 1998; Samant et al., 2021). Data in the experimental hamster model of VL supports this view, with a broad inflammatory environment found in the spleen that favors parasite proliferation (Kong et al., 2017). This view is consistent with studies showing increased T cell expression of Cytotoxic T-lymphocyte-associated protein 4 (CTLA-4) and

Programmed cell death protein 1 (PD-1), indicative of exhausted or anergic T cells (Gautam et al., 2014). The pathogenesis of VL is complex and the factors that cause the disease are not well understood. In this context, many changes in the gene expression profile may occur during infection development, including microRNAs (miRNAs).

miRNAs are a group of small non-coding RNAs (23-25 nucleotides), acting to post-transcriptionally fine-tune gene expression by regulating mRNA levels (Guo et al., 2012; Meng et al., 2012; Liu and Abraham, 2013). miRNAs exert a critical regulatory role in various biological processes, such as cell proliferation, death, reprogramming, signaling, participating in homeostasis, hematopoiesis, cell migration, and regulation of the immune response (Bartel, 2004; Tufekci et al., 2014). In recent years, miRNAs have been shown to play a critical role in developing functional immune responses (Liu and Abraham, 2013). miRNAs have been implicated in various aspects of immunity to viruses, bacteria, and protozoan parasites, including *Trypanosoma*, *Toxoplasma*, *Plasmodium*, and *Leishmania* (Chandan et al., 2019; Acuna et al., 2020; Rashidi et al., 2021). miRNAs have been considered as targets for therapeutic strategies, and since some miRNAs are considered stable in plasma and other biological fluids and are resistant to environmental conditions, they are also attractive as biomarkers of disease or therapeutic response (Sohel, 2016; Takahashi et al., 2019; Souza et al., 2021).

In *Leishmania* infection, modulation of miRNAs expression has been identified in experimental models of visceral and cutaneous leishmaniasis (Acuna et al., 2020; Paul et al., 2020). In VL caused by *L. donovani*, decreased miR-122 expression in the liver of infected mice was associated with an increase in the parasite load (Ghosh et al., 2013). In symptomatic dogs naturally infected with *L. infantum*, expression of miRNAs in PBMCs has been associated with parasite load, with miR-194 and miR-371 positively and miR-150 negatively correlating with the parasitism (Bragato et al., 2018).

In human leishmaniasis, we recently identified a role for miR-548d-3p in cutaneous leishmaniasis caused by *L. (Viannia) braziliensis*. miR-548d-3p was differentially expressed in plasma from patients with active compared to self-healed cutaneous leishmaniasis and up-regulated in *L. braziliensis*-infected THP-1 cells. A role for miR-548d-3p in directly regulating parasite growth was demonstrated *in vitro*, besides modulation of chemokines MCP1/CCL2, RANTES/CCL5, and IP10/CXCL10 production through miR-548d-3p inhibition (Souza et al., 2021). In another *in vitro* study, more than 300 differentially expressed miRNAs were identified in THP-1 cells infected with an *L. donovani* strain isolated from a patient with post-kala-azar dermal leishmaniasis (Kumar et al., 2020).

In the present study, we analyzed the expression of miRNA related to immune-inflammatory processes during *L. infantum* infection. We studied miRNA abundance in plasma of patients with active VL caused by *L. infantum* compared to healthy non-endemic controls and in THP-1 cells infected *in vitro* with *L. infantum* compared to uninfected THP-1 cells. We observed that differentially expressed miRNAs in these two settings targeted

different cellular pathways, including TCR and chemokine signaling and JAK-STAT signaling, as well as cellular processes like apoptosis and peroxisome and phagosome function. miR-548d-3p, previously implicated in *L. braziliensis* infection, was one of the few miRNAs commonly regulated in these two experimental conditions. Transfection of THP-1 cells with a miR-548d-3p inhibitor revealed an effect on inhibiting *Leishmania* growth, likely through induction of MCP-1/CCL2 and nitric oxide production.

MATERIAL AND METHODS

Ethics Statement

All individuals agreed to participate by signing the Informed Consent Form. The experimental protocols were approved by the Ethics Committee of the Faculdade de Medicina, Universidade de São Paulo (CAAE 71602217.4.0000.0065) and are in accordance with the World Medical Association Declaration of Helsinki on Ethical Principles for Medical Research Involving Human Subjects of 1964, with the latest amendment of 2013. All animal procedures were approved by the Faculdade de Medicina, Universidade de São Paulo, to adhere to institutional guidelines for animal care and use (Protocol n° 336).

Patients

Individuals were selected from endemic areas for VL in Pernambuco state, Northeastern Brazil. The sample was collected from 8 active VL patients, five males and three females, mean 14 and median 9 years old (1- 48 years old) with confirmed diagnosis by PCR (using the primers RV1 5'-CTTTTCTGGTCCCGCGGGTAGG-3' and RV2 5'-CCACCTGGCCTATTTTACACCA-3' targeting the *L. infantum* kinetoplast minicircle DNA) (Ravel et al., 1995). They had no comorbidities such as HIV/AIDS. The control group was six healthy adults recruited from non-endemic areas without previous leishmaniasis or coinfections. Four milliliters of whole blood were collected in EDTA from each individual, and plasma was stored at -80°C until use.

Parasites

L. (L.) infantum (MHOM/BR/72/LD) was maintained through regular passages in the hamster (*Mesocricetus auratus*) to preserve infectivity. Promastigotes were derived from amastigotes purified from the spleen of a hamster and expanded and maintained in 199 medium (Cultilab, Brazil) containing penicillin (100 IU/ml), gentamicin (10 µg/ml), L-glutamine (2mM), HEPES (10 mM) and Hemin (20 mg/ml) and supplemented with 10% heat-inactivated fetal calf serum (FCS) (Cultilab, Brazil) at 26°C. The promastigotes used in the experiments were in the stationary phase of growth and had undergone no more than four passages in culture.

Infection of Macrophages With *L. infantum*

The THP-1 human monocytic cell line (ATCC) was grown in RPMI 1640 medium (Sigma-Aldrich, USA) supplemented with

2 mM L-glutamine, 1 mM sodium pyruvate, 0.2% sodium bicarbonate, and 5% FCS (complete medium). Cells (5×10^5) were dispensed onto round 13-mm² glass coverslips in 24-well plates for infectivity analysis, or 2×10^6 cells in the wells of 24-well plates (Corning Costar, USA) and incubated in the presence of 20 ng/mL phorbol myristate acetate (PMA; Sigma-Aldrich, USA) for 48 hours at 37°C in a humid atmosphere with 5% CO₂ to allow differentiation into macrophages (Tsuchiya et al., 1982). Then, the wells were washed twice with a culture medium to remove non-adherent cells. Promastigote suspensions (at a multiplicity of infection of 8 parasites per cell) were dispensed into the wells, and infection was allowed to occur for 4 hours at 37°C in a humid atmosphere with 5% CO₂. After incubation, the non-internalized parasites were washed away, and a complete RPMI medium was added to the wells, beginning the experimental period (0h). The plates were then maintained for 6h or 24h at 37°C in a humid atmosphere with 5% CO₂. The cells into coverslips were stained for evaluation of parasitism. Cells in 24-well plates were prepared in triplicates per plate, and a pool of cells was considered one biological replicate, and experiment was developed three times to generate three biological replicates for RNA extraction. Uninfected cells were considered as controls.

Parasite Load in Macrophages

Coverslips were removed from plates, stained with panoptic dyes (Instant Newprov, Brazil), mounted, and processed to evaluate parasitism under a light microscope (Carl Zeiss, Germany); 900 cells per group were counted. The data are presented as the number of parasites per 100 cells [(number of parasites/number of infected cells) × (number of infected cells/total number of cells) × 100]. Two independent observers who were blinded to the experimental conditions performed the analysis.

RNA Extraction, Reverse Transcription and Pre-Amplification

Total RNA extraction from THP-1 cells was performed using the miRVana PARIS isolation kit (Thermo Fisher, USA), and from plasma samples was performed using the miRNeasy Serum/Plasma kit (Qiagen, USA), with the addition of a spike-in control (*Caenorhabditis elegans* cel-miR-39) to ensure the quality of the procedure and to allow qPCR normalization, according to the manufacturer's instructions. RNA integrity was determined in a spectrophotometer as an OD260/280 absorption ratio between 1.8 and 2.1.

Complementary DNA (cDNA) to template RNA purified from THP-1 cells and plasma samples was synthesized with miScript II RT kit (Qiagen, USA). Briefly, 250 ng of total RNA from THP-1 cells were added to 2 µL of 5X miScript HiSpec Buffer, 1 µL of 10X Nucleics Mix, and 1 µL of miScript Reverse Transcriptase Mix. RNase-free water was added to a final volume of 10 µL. The RNA was incubated for 60 min at 37°C to insert poly-A tail downstream of the miRNA sequence and anneal a T-tail tag for the cDNA elongation. The enzyme was inactivated at 95°C for 5 min. The reaction was performed in the Mastercycler Gradient thermal cycler (Eppendorf, Germany), and the product was stored at -20°C until use. The reverse transcription reaction

for plasma samples followed the same protocol, with the manufacturer's instructions to add 4.5 μL of the purified total RNA. Then, 40 μL of DEPC water was added into each 10 μL RT-PCR product and submitted to a pre-amplification reaction (preAmp), using the miScript PreAmp PCR Kit (Qiagen, USA) according to the manufacturer's instructions. Then the samples were diluted 10x and stored at -20°C .

Quantitative Real-Time PCR for miRNA

miRNA expression was evaluated in three biological replicates with the miScript microRNA PCR array (Qiagen, USA), focusing on inflammation and auto-immunity pathway-related molecules (MIHS-105Z). Ready-to-use qPCR plates containing a set of eighty-four specific primers for miRNAs and twelve internal controls were filled in with the previously prepared master mix containing PCR Buffer, SYBR Green, and the 10-fold diluted cDNA for *in vitro* infected THP-1 macrophages or preAmp samples of plasma samples. Quantitative PCR conditions were 40 cycles of 94°C for 15 seconds, 55°C for 30 seconds, and 70°C for 30 seconds (Souza et al., 2021). Normalization of miRNA expression in THP-1-derived macrophages was performed using SNORD95 and RNU6-6p as reference genes amplified in the qPCR plate. The relative expression levels were calculated using the Comparative Ct method, with non-infected cells considered the calibrator group.

For plasma samples, miRNA expression was also evaluated by relative quantification after previous normalization using the miScript miRNAPCR Array Data Analysis software (Qiagen, USA). The cel-miR-39 spike-in control was considered as a technical reference. Simultaneously, a geometric mean of all expressed miRNAs was used as a normalization factor to calculate relative expression to a calibrator group, which varied depending on the analysis.

In Silico miRNA Target Prediction

Target prediction strategy was performed using DIANA-miRpath 3.0 server in the reverse search module (Vlachos et al., 2015), with Targetscan (Agarwal et al., 2015) as the chosen algorithm. To discover potential interactions with other biological pathways related to human leishmaniasis pathogenesis, we performed a second analysis using MiEEA (MiRNA Enrichment Analysis and Annotation), which integrates data from different databases such as miRBase, miRWalk, and miRTarBase (Backes et al., 2016).

In Vitro miRNA Inhibition

The inhibition of miR-548d-3p in THP-1-derived macrophages was performed through a transient transfection protocol. Assays with three different concentrations (3 nM, 10 nM, and 30 nM) of the mirVana[®] miR-548d3p inhibitor (Ambion, USA) or scrambled Negative Control (Ambion, USA) were performed, and 10 nM concentration was chosen for further use. Before the addition of *L. infantum* promastigotes, a solution containing the miR-548d-3p inhibitor or the negative control together with 3 μL of FUGENE transfection reagent (Promega, USA) diluted in 500 μL of RPMI medium previously incubated for 20 min at room

temperature was added into each well containing the cells and maintained for 24h (Souza et al., 2021). Simultaneously, non-transfected cells received only a complete RPMI medium. The experiment continued with promastigote infection to evaluate parasitism, chemokine, and nitrite levels in supernatants collected and stored at -80°C until use.

Evaluation of Chemokine Production

Chemokine quantification was performed using CBA – Human Chemokine Kit (BD Biosciences, USA), following the manufacturer's instructions in culture supernatants. Briefly, 50 μL of capture beads for MCP1/CCL2, RANTES/CCL5, IL-8/CXCL8, MIG/CXCL9, and IP10/CXCL10, 50 μL of Detection Reagent and 50 μL of the studied sample or standard were added consecutively to each sample tube and incubated for 3 h at room temperature, in the dark. Next, the samples were washed with 1 ml of wash buffer and centrifuged. After discarding the supernatant, the pellet was resuspended in 300 μL buffer and analyzed in a FACS LSR Fortessa flow cytometer (BD Biosciences, USA) (Souza et al., 2021). Raw data were then analyzed using FCAP Array software (BD Biosciences, USA). The detection limits of each chemokine were as follows: 2.7 pg/mL for MCP1/CCL2, 1.0 pg/mL for RANTES/CCL5, 0.2 pg/mL for IL-8/CXCL8, 2.5 pg/mL for MIG/CXCL9 and 2.8 pg/mL for IP10/CXCL10.

Nitric Oxide (NO) Production

Nitrite (NO_2) accumulation in the cell culture supernatants was used as an indicator of NO production and was determined using the standard Griess reaction (Green et al., 1982). Fifty microliters of the culture supernatant were reacted with 50 μL of Griess reagent (1% sulfanilamide, 0.1% N-(1-Naphthyl)ethylenediamine dihydrochloride and 2.5% phosphoric acid in bidistilled water) for 10 min at room temperature. The absorbance was measured at 540 nm using a Multiskan MCC/340 P version 2.20 plate reader (Labsystems, Brazil), and the nitrite concentration was calculated using a standard curve for sodium nitrite (NaNO_2). The tests were run in triplicate.

Statistical Analysis

Statistical analyses of *in vitro* miRNA expression were performed with Qiagen miScript miRNA PCR Array Data Analysis online software, where data from three independent experiments were submitted to an integrated Student's t-test under the manufacturer's recommendation as done in the previous similar work (Muxel et al., 2017). The fold regulation (FR) was considered to be the negative inverse of the fold change [function = $-1^*(1/\text{fold change value})$]. $\text{FR} \geq 2$ were considered to indicate upregulation, and levels ≤ -2 were considered to indicate downregulation, as previously described (Muxel et al., 2017). Ex vivo data were also submitted to Student's t-test, with Bonferroni's correction, using Microsoft Excel 365. Parasite load data were analyzed by ANOVA with Tukey's post-test and data from chemokine quantification by Kruskal-Wallis test with Bonferroni's correction. The differences were considered significant when $P < 0.05$.

RESULTS

miRNA Expression in *L. infantum*-Infected THP-1 Cells

miRNA expression was evaluated at 6h and 24h post infection (p.i.), with non-infected THP-1 cells used as a calibrator group. Twenty two (26%) out of 84 miRNAs assayed were significantly altered in expression ($P < 0.05$) across one or more condition. At 6h p.i., nine miRNAs were upregulated (miR-302a-3p, miR-302b-3p, miR-372-3p, miR-373-3p, miR-381-3p, miR-524-5p, miR-543, miR-548c-3p, miR-607), while seven were down-regulated (let-7b-5p, let-7d-5p, miR-144-3p, miR-19b-3p, miR-21-5p, miR-29c-3p, miR-548d-3p) (Figure 1A). At 24h p.i., eight miRNAs were upregulated (let-7b-5p, let-7d-5p, miR-302b-3p, miR-340-5p, miR-372-3p, miR-373-3p, miR-548d-3p, miR-607) and five were downregulated (miR-144-3p, miR-29c-3p, miR-302a-3p, miR-524-5p, miR-548c-3p) (Figure 1B). Through the time course of infection, we observed that four miRNAs (miR-302b-3p, miR-372-3p, miR-373-3p, miR-607) were upregulated at both 6h and 24h, while miR-144-3p was down-regulated at both times. let-7b-5p, let-7d-5p, miR-340-5p and miR-548d-3p were down-regulated at 6h and upregulated at 24h. In contrast, miR-302a-3p and miR-548c-3p were up-regulated at 6h and

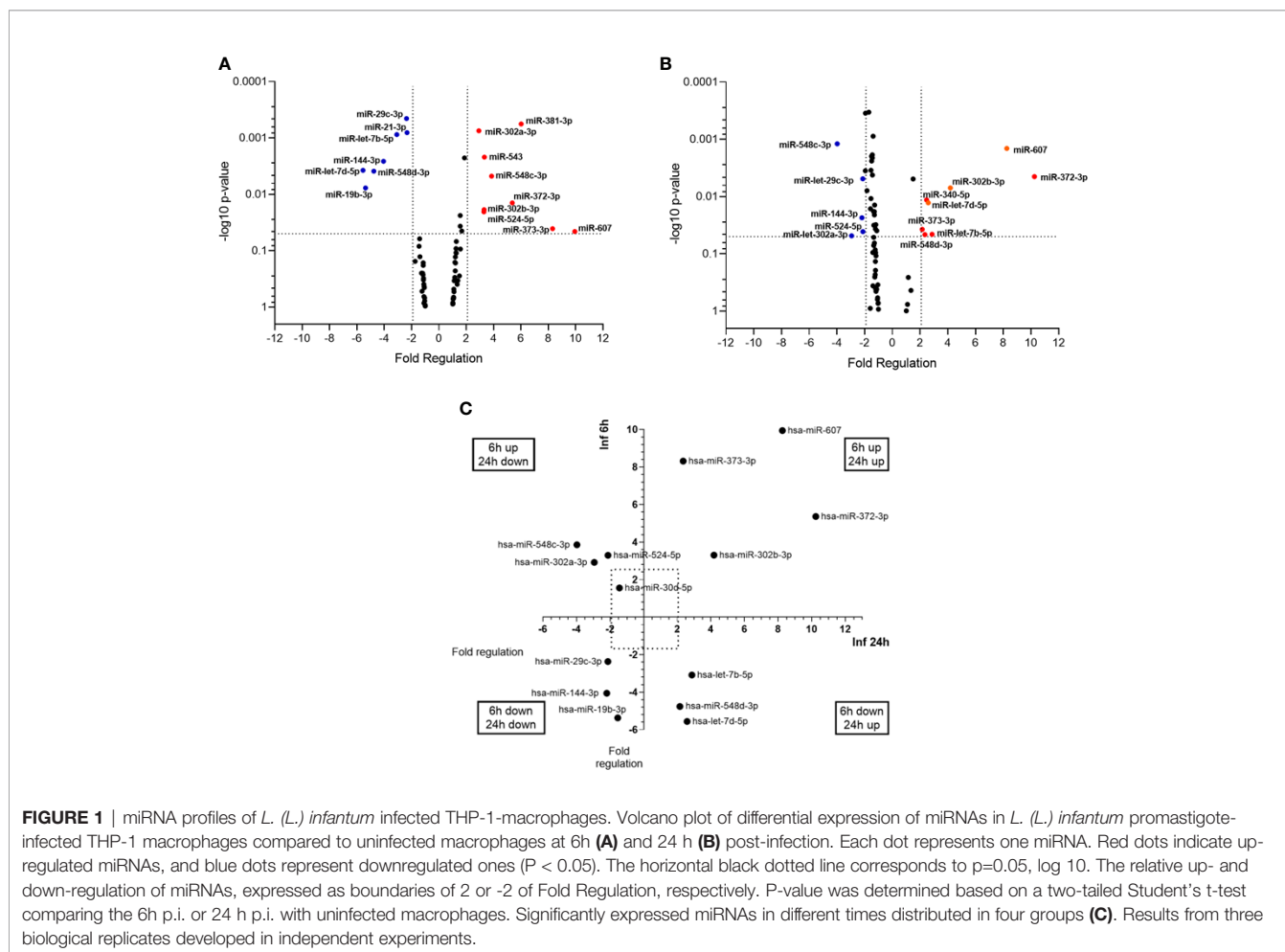
down-regulated at 24h (Figure 1C). These data indicate that *L. infantum* infection selectively alters miRNA abundance in human macrophage in a time-dependent manner.

miRNA Expression in Plasma From Active VL Patients

miRNA expression was next evaluated in plasma samples from patients with active disease compared to a healthy endemic control group. Forty-two (50%) out of 84 miRNAs assayed were significantly altered in abundance ($P < 0.05$) (Figure 2). 32 miRNAs were up-regulated in patients with active VL, whereas 10 were downregulated (Table 1). Of note, 6 miRNAs were commonly up-regulated both in these plasma samples and in infected THP-1 cells (miR-302a-3p, miR-302b-3p, miR-372-3p, miR-373-3p, miR-340-5p, and miR-548d-3p) (Table 1).

miRNA Predicted Targets and Their Interactions With Biological Pathways Related to VL

Considering all differentially expressed miRNAs, we searched for target genes and metabolic pathways using Diana MiRPath 3.0 with TargetScan2 as the chosen algorithm to predict miRNA/mRNA interactions. Analyzing modulated miRNAs in infected



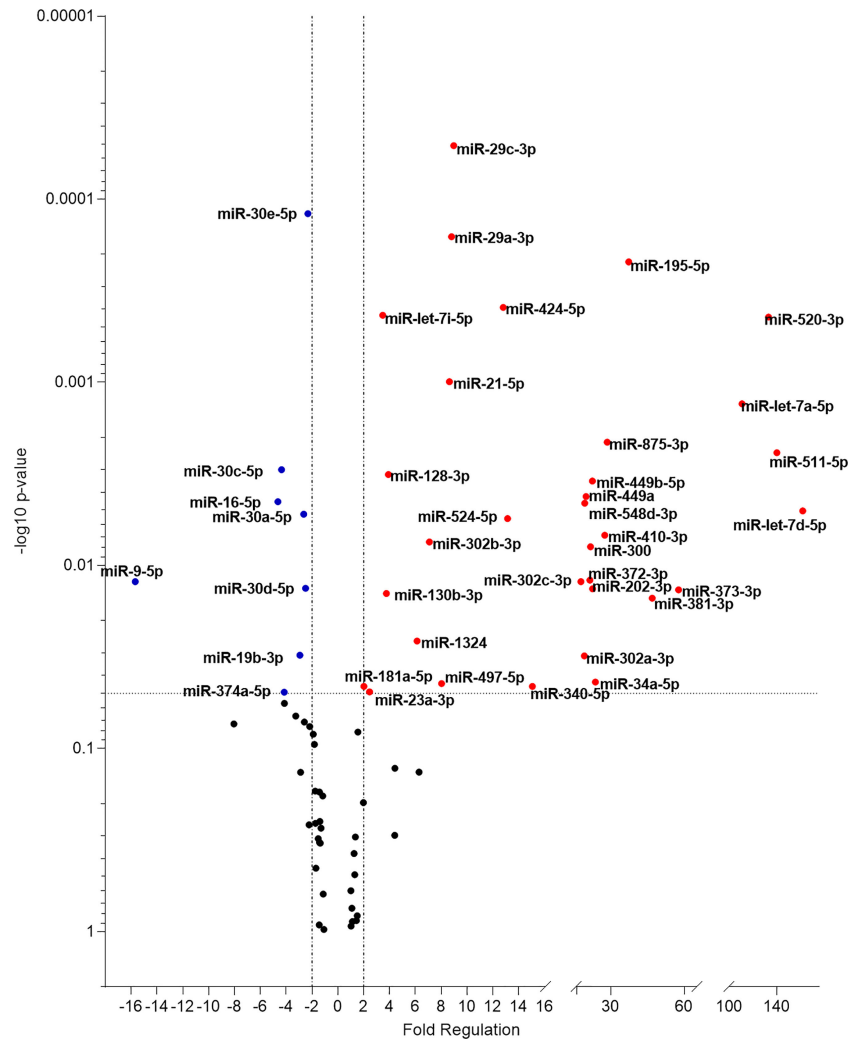


FIGURE 2 | Volcano plot of differential expression of miRNA in plasma from VL patients. Volcano plot showing differential expression of miRNA in plasma samples of active VL patients compared to healthy individuals. Each dot represents one miRNA. Red dots indicate up-regulated miRNAs, and blue dots represent down-regulated ones ($P < 0.05$). The horizontal black dotted line corresponds to $p=0.05$, log 10. The relative up- and down-regulation of miRNAs, are expressed as boundaries of 2 or -2, of Fold Regulation respectively. P-value was determined based on a two-tailed Student's *t*-test. $P < 0.05$ (Student *t*-test and Bonferroni correction).

TABLE 1 | miRNAs significantly expressed in plasma samples of patients with active disease.

Up-Regulated miRNAs		Down-Regulated miRNAs	
hsa-let-7a-5p	hsa-miR-29a-3p	hsa-miR-410-3p	hsa-let-7c-5p
hsa-let-7d-5p	hsa-miR-29c-3p	hsa-miR-424-5p	hsa-miR-101-3p
hsa-let-7i-5p	hsa-miR-300	hsa-miR-449a	hsa-miR-16-5p
hsa-miR-128-3p	hsa-miR-302a-3p	hsa-miR-449b-5p	hsa-miR-19b-3p
hsa-miR130b-3p	hsa-miR-302b-3p	hsa-miR-497-5p	hsa-miR-30a-5p
hsa-miR-1324	hsa-miR-302c-3p	hsa-miR-511-5p	hsa-miR-30c-5p
hsa-miR-181a-5p	hsa-miR-340-5p	hsa-miR-520d-3p	hsa-miR-30d-5p
hsa-miR-195-5p	hsa-miR-34a-5p	hsa-miR-524-5p	hsa-miR-30e-5p
hsa-miR202-3p	hsa-miR-372-3p	hsa-miR-548d-3p	hsa-miR-374a-5p
hsa-miR-21-5p	hsa-miR-373-3p	hsa-miR-875-3p	hsa-miR-9-5p
hsa-miR-23a-3p	hsa-miR-381-3p		

miRNAs in bold, miRNAs up-regulated in infected THP-1 cells and plasma samples of VL patients.

THP-1 cells, four important metabolic pathways were predicted: the TGF- β pathway with 56 possible target genes; the FOXO transcription factor pathway (Forkhead box) with 74 putative target genes, associated with the control of apoptosis and glucose metabolism; Wnt signaling pathway with 64 putative targets related to foxo, PI3/AKT and the HIF-1 factor pathway with 55 putative target genes associated with anaerobic metabolism in M1 macrophages (Table S1).

New targets were predicted using the MiEEA platform to obtain coverage from more than one classification system (PANTHERDB, WikiPathways, and KEGG). This analysis is worth highlighting those likely targeted by more than one miRNA. There were some pathways known to be important in the *Leishmania*-host interaction like T cell receptor, chemokine signaling pathway, peroxisome, JAK-STAT signaling pathway, apoptosis, PPAR signaling pathway, phagosome, JAK1, MAPK1, PTPN6, PTGS2, of which MAPK1, JAK are involved in the TGF β , FOXO, and HIF1 pathways (Figure 3). Some of these pathways have also been observed in our previous study of *L. braziliensis* infection (Souza et al., 2021). Hsa-miR-548d-3p also targets common genes in these pathways. A similar analysis was then conducted with the miRNAs that were found to be

differentially regulated in plasma from VL patients (Figure 4), revealing a similarly broad and diverse set of potential targets influenced by infection. Among various differentially expressed miRNAs identified, we selected miR-548d-3p, up-regulated in both plasma samples from active VL patients and *L. infantum*-infected THP-1 cells for further validation.

Effect of miR-548d-3p Inhibition on the Parasite Load in *L. infantum*-Infected THP-1 Cells

Having identified miR-548d-3p as one of 6 miRNAs affected both in infected THP-1 cells and in plasma from patients with VL, we next sought to identify how this miRNA affected the outcome of infection using the THP-1 model by conducting targeted knockdown experiments. We inhibited miR-548d-3p during *L. infantum* infection using different concentrations (3, 10, and 30 nM) of specific inhibitor or scrambled miRNA. THP-1 cell viability was not affected by the inhibitors at the concentrations used. All concentrations of miR-548d-3p inhibitor increased parasite load in THP-1 cells with the maximal response (1.5 fold increase in parasite numbers) observed at 10 nM and above (Figure S1). At both 6h and 24h

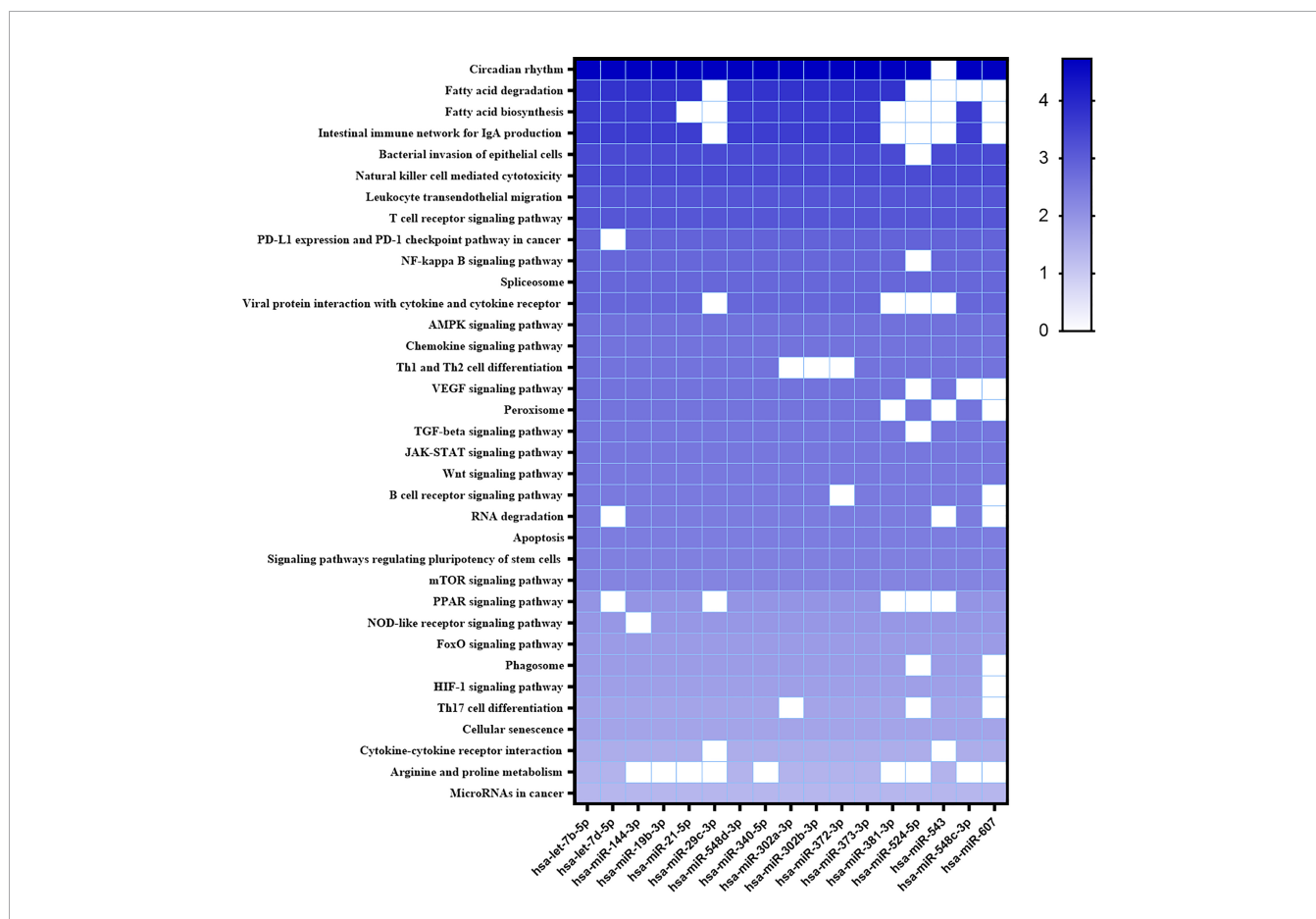


FIGURE 3 | Heatmap of predicted interactions between differentially expressed miRNAs in infected THP-1 cells. The set of differentially expressed microRNAs in *L. infantum*-infected THP-1 at 6 and 24 h and the biological pathways on which they are suggested to act, according to MiEEA algorithms, are shown in heat map format. Results are shown as $-\log_{10}$ of prediction p-value, and the degree of color saturation corresponds to the $-\log_{10}$ of the p-value.

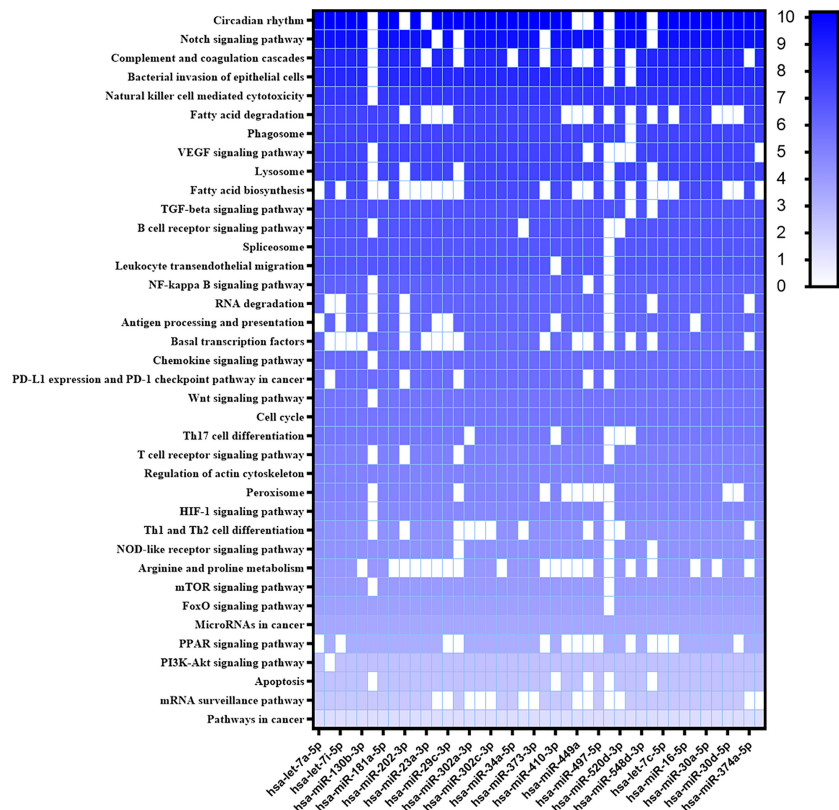


FIGURE 4 | Heatmap of predicted interactions between differentially expressed miRNAs in VL patient plasma. The set of differentially expressed microRNAs in plasma samples from active VL patients caused by *L. infantum* and the biological pathways on which they are suggested to act according to MiEAA algorithms are shown in heat map format. Results are shown as $-\log_{10}$ of prediction p-value, and the degree of color saturation corresponds to the $-\log_{10}$ of the p-value.

post-infection, we observed a significant increase in the parasite load when miR-548d-3p was inhibited ($P < 0.05$), compared with transfection with scrambled RNA (negative control) and non-transfected cells (Figure 5A).

Effect of miR-548d-3p Inhibition on the Nitric Oxide and Chemokine Production in *L. infantum*-Infected THP-1 Cells

As we observed a change in parasite load, we next evaluated NO production, given its role as a major leishmanicidal effector mechanism in macrophages. The effect of miR-548d-3p on nitric oxide (NO) production was assessed at 6 and 24 hours post-infection. We observed a tendency towards decreased NO production at 6h p.i. and a significant decrease at 24h when miR-548d-3p was inhibited ($P < 0.05$) compared with transfection with scrambled RNA (negative control) and non-transfected cells (Figure 5B).

Given that chemokines can regulate and be regulated by NO production and are also stimulated by infection with *Leishmania*, we analyzed the production of chemokines in the supernatant of infected THP-1 cells with or without miR-548d-3p inhibition (Figures 5C–F). Since the approach was on THP-1 cells, we selected the chemokines related to monocyte/macrophages

biology. Production of MCP-1/CCL2, RANTES/CCL5, and IP-10/CXCL10, but not IL-8/CXCL8, was readily observed at 6h p.i. MIG/CXCL9 was undetectable in all samples. Although MCP-1/CCL2 and RANTES/CCL5 expression was maintained at 24h p.i., MIG/CXCL10 expression was reduced at this later time, possibly reflecting limited induction of interferon in this system. Amongst these four chemokines, only the production of MCP-1/CCL2 (Figure 5D) was significantly decreased by miRNA inhibition and only at 24 h p.i. Although NO has been shown to be positively regulated by MCP-1/CCL2 (Biswas et al., 2001), the more immediate impact of miR-548d-3p inhibition on NO production (Figure 5A) suggests that this may occur independently of its effect on MCP-1/CCL2 production.

Chemokine Levels in the Plasma of VL Patients

Finally, we examined the plasma of VL patients and controls for expression of the same set of chemokines. VL patients showed significantly increased IP-10/CXCL10, MCP-1/CCL2, MIG/CXCL9, and IL-8/CXCL8 chemokines compared to the control group (healthy individuals). No difference was observed in RANTES/CXCL5 production (Figure 6).

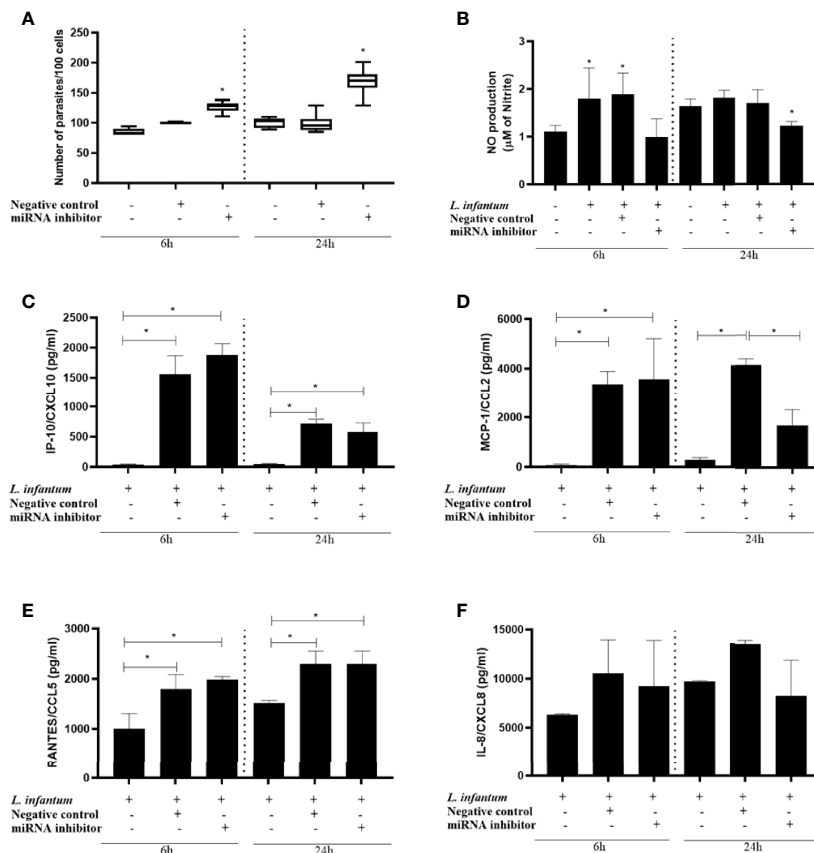


FIGURE 5 | Parasite load, nitric oxide production, and chemokine levels in *L. (L.) infantum* promastigote-infected THP-1 cells transiently transfected with miR-548d-3p inhibitor. THP-1 cells were transiently transfected with miR-548d-3p inhibitor (10 nM) or negative control (scrambled miRNA; 10 nM) as described in Methods. After 24h, macrophages were infected with *L. (L.) infantum* promastigotes and incubated for 6 and 24h post-infection for different evaluations. **(A)** Parasite load represented as the number of parasites/100 cells. **(B)** Nitric oxide (NO) production was determined by the accumulation of nitrite in the cell culture supernatants. IP-10/CXCL10 **(C)**, MCP-1/CCL2 **(D)**, RANTES/CXCL5 **(E)**, and IL-8/CXCL8 **(F)** concentrations (pg/mL) were measured by flow cytometry using the CBA kit. * = $P < 0.05$ (one way ANOVA and Tukey's test). One representative experiment from three independent assays is shown.

DISCUSSION

In the present study, we examined miRNA expression in *L. infantum* infection to identify signatures likely associated with the pathogenic process. Few studies have evaluated the participation of miRNAs during *Leishmania* infection (Lemaire et al., 2013; Muxel et al., 2017; Muxel et al., 2018; Fernandes et al., 2019; Paul et al., 2020; Rashidi et al., 2021; Souza et al., 2021). We compared miRNAs in plasma of patients with VL caused by *L. infantum* and in human monocyte-derived THP-1 cells infected *in vitro* with *L. infantum*.

Our data are the first to report on the profile of miRNAs in plasma from patients with active VL caused by *L. infantum*. We identified 32 miRNAs up-regulated and 10 down-regulated in plasma during VL infection compared to plasma-derived from healthy endemic controls. Amongst up-regulated miRNAs, six were also up-regulated *in vitro* in THP-1 cells, namely miR-302a-3p, miR-302b-3p, miR-340-5p, miR-372-3p, miR-373-3p, and miR-548d-3p, suggesting that these miRNAs can be directly regulated in macrophages as a consequence of intracellular

parasitism. One of these, miR-548d-3p, had not previously been described with *L. infantum* infection. Therefore, we determined whether it played a functional role during *in vitro* infection.

The miR-548 family, only described in primates, comprises over 69 identified miRNAs present on almost all human chromosomes and is one of the larger yet more poorly conserved microRNA families. The miR-548 family is implicated in different biological processes, including regulating important signaling pathways: MAPK, Phosphatidylinositol (PI3K), p53, B cell receptor, T cell receptor, TGF-beta, PPAR, Calcium, and Insulin signaling pathways. It also has a role in human tumorigenesis (Liang et al., 2012).

Members of this miRNA family have been shown to target immune system-associated genes and have been described as differentially expressed in other diseases. miR-548ah-5p is up-regulated at the immune activation phase of chronic hepatitis B patients, and *in vitro* studies using THP-1 cells treated with miR-548ah-5p mimics suggest that it may involve the reduced expression of IFN- γ R1, which may, in turn, facilitate viral

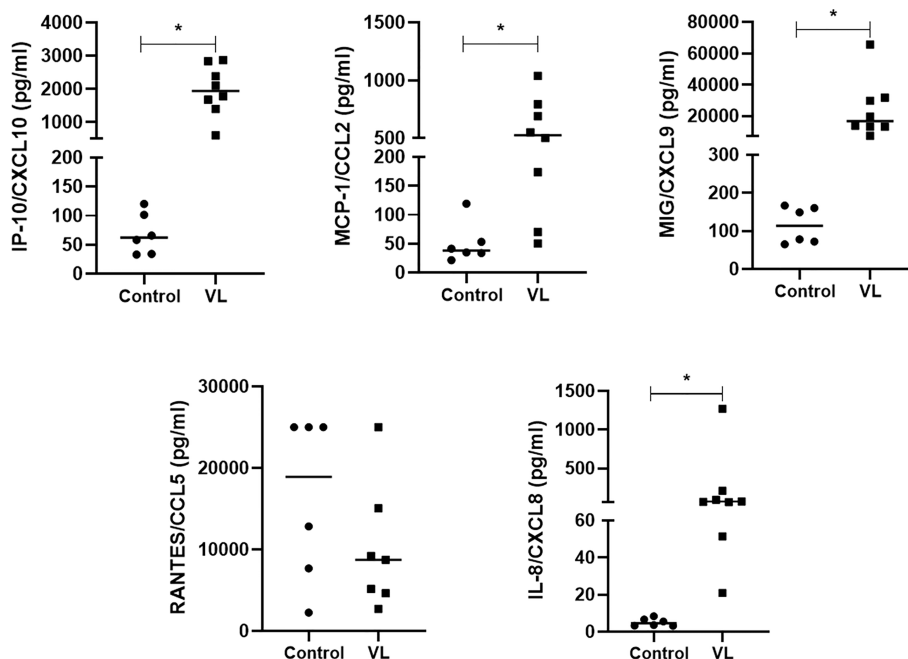


FIGURE 6 | Chemokines in plasma of active VL patients and controls. IP-10/CXCL10, MCP1/CCL2, MIG/CXCL9, RANTES/CXCL5, and IL-8/CXCL8 concentrations were measured in the CBA kit using flow-cytometry. Data are shown as pg/mL, and each symbol corresponds to one subject * = $p < 0.05$ (Mann-Whitney test). Patients (N=8) and controls (N=6).

pathogenesis (Xing et al., 2014). Other members of the miR-548 family down-regulate host antiviral response *via* direct targeting interferon- λ 1 (IFN- λ 1). For example, inhibition of miR-548b-5p, miR-548c-5p, miR-548i, and miR-548j increased IFN- λ 1 secretion, and the targeted IFN- λ 1 gene can induce IP-10, MCP1, IL-8 (Fu et al., 2021). The chemokine MCP-1 was seen affected by miR548d-3p in the present study. Due to the diversity of signaling pathways regulated by the miR-548 family and its participation in *Leishmania* infection in our previous results in *L. braziliensis* infection affecting parasite growth and chemokine production (Souza et al., 2021), we proceeded to evaluate the role of miR-548d-3p in *L. infantum* infection.

miR-548d-3p was up-regulated in plasma from VL patients and in *L. infantum* infected- THP-1 cells *in vitro*. Our results also show that *L. infantum* infected-THP-1 cells treated with miR-548d-3p inhibitor had increased parasite load and decreased expression of MCP1 and NO. MCP-1 is known to induce the production and release of NO, a leishmanicidal molecule, explaining in part the increase in parasitism in the presence of the miRNA inhibitor. However, the MCP-1 level in serum samples was elevated in active VL that presents with high parasitism. However, in another study, it was even higher after 6 months of treatment and in asymptomatic patients (Ibarra-Meneses et al., 2017). These data suggest that MCP-1 affects parasite growth inhibition but not as a single anti-leishmanial element. In addition, full concordance across *in vitro* and *in vivo* data would not be expected as multiple cells, and tissue sources

may contribute to MCP1 plasma levels *in vivo*, many of which would not be directly subject to intracellular parasitism. In our previous study, miR-548d-3p was up-regulated in plasma from self-healed cutaneous leishmaniasis patients and *in vitro* *L. braziliensis*-infected THP-1 cells, and when miR-548d-3p was inhibited in infected THP-1 cells, a decrease in parasite load was observed, and an increase in the production of MCP1/CCL2, RANTES/CCL5, and IP-10/CXCL10 (Souza et al., 2021). These differences may be due to species-specific mechanisms for immune system activation and evasion and the differences (cutaneous or systemic) in clinical presentation. Other members of the 548-miRNA family, miR-548at-5p and miR-548t-3p, were up-regulated in THP-1 cells infected with promastigotes isolated from VL and post kala-azar dermal leishmaniasis patients caused by *L. donovani*. These were seen as related to inflammation and apoptosis (Kumar et al., 2020).

It was observed that MCP-1 cytokine mRNA is more abundant in lesions of *L. mexicana*-infected patients who developed self-healing localized lesions when compared to lesions from chronic diffuse cutaneous leishmaniasis. This finding suggests that the cytokine MCP-1 may contribute to the spontaneous healing process (Ritter et al., 1996). Using human monocytes infected with *L. major*, it was demonstrated that MCP1 chemokine activates monocytes and decreases parasitism, possibly by producing intermediate reactive oxygen species (Ritter and Moll, 2000). In the same line, human peripheral macrophages infected with *L. infantum* showed that MCP-1 enhances macrophage's nitric oxide production and

release at levels induced by INF- γ stimulation. This NO production results in decreasing parasitism (Brandonisio et al., 2002).

Another study demonstrated that *L. infantum*-infected whole blood cells from asymptomatic patients, when stimulated with soluble *Leishmania* antigen, produce significantly higher levels of MCP1 than patients with active or cured VL. Whole blood cells from cured patients have significantly higher levels of MCP-1 after 6 months of the treatment (Ibarra-Meneses et al., 2017). These data reinforce that MCP-1 is essential for a protective immune response and can be used as a biomarker for a protective immune response against *L. infantum*. The increase in parasitism observed in the present study may suggest inhibition of NO production *via* MCP1. However, other studies showed the activation of PI3K/AKT signaling pathway activation upon knockdown of miR-548b-3p with further suppression of cell apoptosis and enhancement of cell proliferation (Xie et al., 2017; Wang et al., 2020). The increase the parasitism in the presence of the miRNA inhibition *in vitro* may also be related to the activation of the PI3K/AKT signaling pathway.

Our ex-vivo results show that patients with active VL have significantly higher IP10/CXCL10, MIG/CXCL9, IL-8/CXCL8, and MCP1/CCL2 compared with the control group, probably an attempt of the immune system to control *Leishmania* infection. IP-10 seems to be related to the control of *L. amazonensis* infection and also protects against VL (Vasquez and Soong, 2006; Fallahi et al., 2016). In the active VL, if miR-548d-3p is acting to control the parasite growth, it should have other elements counteracting to maintain infection progression seen in these cases.

In addition to miR-548d-3p, other miRNAs identified in this study have also been observed to change expression in studies of *Leishmania* infection. These miRNAs presented below in sequence are related to activation or modulation of the immune system and may play a crucial role in *Leishmania* infection either in the control or progression of the infection. The following miRNAs may contribute to the progression of the infection: a) miR-21 seen up-regulated in dendritic cells, macrophages, inflammatory monocytes, polymorphonuclear neutrophils, and in the spleen and liver tissues after *L. donovani* infection and in splenic leukocytes from *L. infantum*-naturally infected dogs and related to the inhibition of IL-12 expression and impairment of the Th1 response implied in the control of the infection (Melo et al., 2019; Varikuti et al., 2021); b) miR-302a-3p that regulates receptor activator of nuclear factor kappa-B ligand (RANKL) (a member of the tumor necrosis factor TNF superfamily) that regulates T-cell-dependent immune response and apoptosis (Irwandi et al., 2018) and also up-regulated in *in vitro* *L. braziliensis* infection (Souza et al., 2021); c) miR-372-3p that inhibits cell proliferation and apoptosis and seen up-regulated in *L. braziliensis* infection (Chen et al., 2015; Souza et al., 2021). Otherwise, distinct miRNAs may act on the control of the infection: a) miR-340-5p that targets the IL-4, major Th2 cytokines secreted by CD4⁺ T cells, and this miRNA, different from observed with *L. infantum*,

was down-regulated in *in vitro* *L. donovani* infection (Kumar et al., 2020); b) miR-302b-3p that inhibits cell proliferation through the AKT pathway by targeting insulin-like growth factor 1 receptor (IGF-1R) (Guo et al., 2017) and was also up-regulated in *in vitro* *L. amazonensis* and *L. braziliensis* infection (Muxel et al., 2017; Souza et al., 2021); c) miR-373-3p that is implicated in the regulation of cell proliferation, apoptosis, senescence, migration, and cell invasion and was up-regulated in *L. braziliensis* infection (Wei et al., 2015; Souza et al., 2021).

Our *in silico* results show that up-regulated microRNAs in plasma and THP-1 cells can target processes or pathways of TGF- β signaling, T cell receptor signaling chemokine signaling, cytokine-cytokine receptor interaction, NOD-like receptor signaling, arginine a proline metabolism. From the total up-regulated microRNAs seen in the present study, six miRNAs are also up-regulated in plasma and THP-1 samples and target important pathways. When inhibited, phagosome and peroxisome processes may result in a favorable microenvironment for amastigote parasites, including the inhibition of apoptosis by infection. Other pathways like the NOD pathway associated with inflammasome, TGF- β signaling pathway, INHBA gene that may modulate MCP-1, IP-10, NO production, arginine, proline metabolism, chemokine signaling were targeted. These pathways are involved directly or indirectly in chemokine, cytokine signaling, or gene expression, including MCP-1 and NO production altered or modulated in the present experiments upon miR-548d-3p inhibition.

In conclusion, this study of VL patients and using *L. infantum*-infected THP-1 cells extend previous observations on the role of miR-548d-3p in cutaneous leishmaniasis caused by *L. braziliensis* (Souza et al., 2021). Our data open new possibilities to study the interactions of the 548 family and the respective targeted genes, using the miR-548d-3p as a tool directed to the regulation of immune or pathological processes in humans/primates leishmaniasis. In addition, miR-548d-3p may be explored as a biomarker of prognostic value, and targeting miR-548d-3p may provide new therapeutic opportunities in leishmaniasis and other inflammatory diseases.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/**Supplementary Material**. Further inquiries can be directed to the corresponding author.

ETHICS STATEMENT

The studies involving human participants were reviewed and approved by Ethics Committee of the Faculdade de Medicina, Universidade de São Paulo (CAAE 71602217.4.0000.0065). The patients/participants provided their written informed consent to participate in this study. All animal procedures were approved by the Faculdade de Medicina, Universidade de São Paulo to adhere to institutional guidelines for animal care and use (Protocol n° 336).

AUTHOR CONTRIBUTIONS

HG: Conceptualization, study design, project and researcher supervision, data analysis, and manuscript preparation. ER-S, LR, and MS: Conceptualization, study design, experimental work, data analysis, manuscript preparation. SM: study design, experimental work, data analysis, manuscript preparation. KS: experimental work; VP and MB: coordination of sample and data collection in the endemic area; data interpretation. LF-W: study design, researcher supervision, data analysis, and manuscript preparation. DL and PK: data analysis, manuscript preparation. All authors contributed to the article and approved the submitted version.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fcimb.2022.826039/full#supplementary-material>

Supplementary Figure 1 | Parasite load (number of parasites/100 cells) in *L. (L.) infantum* promastigote-infected THP-1 cells transiently transfected with miR-548d-3p inhibitor. THP-1 cells were transiently transfected with miR-548d-3p inhibitor (miR = 3, 10 and 30 nM) or negative control (NC - scrambled miRNA = 3, 10 and 30 nM) as described in Methods. After 24h, cells were infected with *L. (L.) infantum* promastigotes, and parasite load in macrophages was assessed at 6 hours post-infection (**A**) and 24 hours post-infection (**B**). * = $P < 0.05$ (one-way ANOVA and Tukey's test) compared to the control groups. One representative experiment from three independent assays is shown.

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