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Triphenylmethane Derivatives Have High In Vitro and In Vivo Activity against the Main Causative Agents of Cutaneous Leishmaniasis

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Abstract

The current standard of care for cutaneous leishmaniasis (CL) is organic antimonial compounds, but the administration of these compounds is complicated by a low therapeutic-toxic index, as well as parenteral administration. Thus, there is an urgent need for the development of new and inexpensive therapies for the treatment of CL. In this study, we evaluate the activity of the triphenylmethane (TPM) class of compounds against three species of Leishmania which are pathogenic in humans. The TPM have a history of safe use in humans, dating back to the use of the original member of this class, gentian violet (GV), from the early 20th century. Initially, the in vitro efficacy against Leishmania (Viannia) braziliensis, L. (Leishmania) amazonensis and L. (L.) major of 9 newly synthesized TPM, in addition to GV, was tested. Inhibitory concentrations (IC) IC₅₀ of 0.025 to 0.84 μM had been found in promastigotes in vitro assays. The four most effective compounds were then tested in amastigote intracellular assays, resulting in IC₅₀ of 0.10 to 1.59 μM. A high degree of selectivity of antiparasitic activity over toxicity to mammalian cells was observed. Afterwards, GV and TPM 6 were tested in a topical formulation in mice infected with L. (L.) amazonensis leading to elimination of parasite burdens at the site of lesion/infection. These results demonstrated that TPM present significant anti-leishmanial activities and provide a rationale for human clinical trials of GV and other TPM. TPM are inexpensive and safe, thus using them for treatment of CL may have a major impact on public health.

Introduction

The genus Leishmania spp. protozoa are pathogenic to a wide variety of hosts, including humans, and are most prevalent in tropical climates of developing countries. The major forms of leishmaniasis include cutaneous, mucosal and visceral leishmaniasis [1]. Leishmania (Leishmania) major is one of the main etiological agents of CL in the Old World, while Leishmania (Viannia) braziliensis, Leishmania (L.) amazonensis and Leishmania (V.) guyanensis are the main causative species of CL in the Americas. Lesions caused by these species frequently appear as ulcers at the site of infection and are commonly located in poorly-protected areas of the body, such as the face, arms and legs [2,3]. In addition, L. (L.) amazonensis may also lead to development of diffuse cutaneous leishmaniasis in a few patients, which is characterized by nodular lesions, refractory to chemotherapy [4].

Current therapeutic alternatives for CL treatment are unsatisfactory. The conventional first-line therapy consists of pentavalent antimonials (sodium stibogluconate - Pentostan™ and meglumine antimoniate - Glucantime™). However, these drugs present inconvenient aspects that limit their use, such as the necessity of parenteral administration and a high incidence of toxic and adverse reactions [2]. Pentavalent antimonials have long been considered highly effective [3,5], however, there is a growing body of evidence of variable efficacy, depending on species, geographic region, presence of resistant strains, and therapeutic schemes [2,6–8]. Among the alternative therapeutic schemes, intraleisonal administration of pentavalent antimonials has been used to treat old world cutaneous leishmaniasis [9].

The second line therapies for leishmaniasis include amphotericin B (AmB), liposomal AmB, and pentamidine. AmB is a very powerful polyenic antibiotic against Leishmania but also presents significant adverse effects, including nephrotoxicity and infusion reactions. Liposomal AmB was developed to improve the tolerability profile of AmB deoxycholate [10]. In Brazil, liposomal AmB is recommended for CL treatment only upon failure of first line therapies. In addition, another limitation of liposomal AmB is
its high cost [11]. Pentamidine is complicated by hypoglycemia and the requirement of intravenous administration.

Finally paromomycin, an aminoglycoside antibiotic, is an antileishmanial drug that has been on the market since the 1960’s and has been used in several formulations for the topical treatment of CL, with inconclusive results [12–17]. Therefore, further research and studies based on new technologies aimed at improving the delivery and efficacies of topical treatments are still required, especially in regards to safety, efficacy, and cost [18].

Compounds with the triphenylmethane pharmacophore (TPM), such as gentian violet (GV), have a long history of human use as anti-bacterial and antymycotic agents. In addition, GV has been shown to have antiparasitic activity against various human parasites and have been used in blood banks to circumvent Chagas’ disease transmission [19–24]. However, TPM have not been previously evaluated in vivo against CL.

We synthesized 9 novel TPM derivatives, as part of a structure-function study of TPM compounds and tested, in addition to GV, against 3 species of pathogenic Leishmania, including both Old World and New World Leishmania. These derivatives were tested against both promastigotes of L. (L.) amazonensis, L. (L.) major and L. (V.) braziliensis and intracellular amastigotes of L. (L.) amazonensis and L. (V.) braziliensis. Finally, we demonstrate that topical treatment with either GV or one of the novel TPM is highly effective in treating L. (L.) amazonensis infected mice.

**Materials and Methods**

**Ethics statement**

This study has been approved by Ethics Committee for Animal Experimentation from University Federal of Minas Gerais (CETEA/UFMG: 12/2009). The University Federal of Minas Gerais adheres to the standards as outlined by relevant national (CONCEA - Brazilian Government Council for Control of Animal Experimentation) and international guidelines for care and use of laboratory animals.

**Parasites**

Promastigotes of L. (L.) amazonensis (IFLA/BR/1967/PB-8), L. (V.) braziliensis (MHOM/BR/75/M2903), and L. (L.) major (MHOM/IL/80/Friedlin) were maintained at 23°C in Schneider’s Drosophila medium (Merck, Germany) supplemented with 20% heat-inactivated fetal calf serum (FCS) (Gibco, Eggenstein, Germany), pH 7.2. The same strain of L. (L.) amazonensis was used for both in vitro and in vivo experiments.

**Triphenylmethane compounds (TPM)**

Novel TPM were synthesized by reacting aromatic substrates with 4,4’diis (diethylaminobenzophenone) in the presence of phosphorus oxychloride in a calorimeter bomb at 140°C under pressure. TPM formula and molecular weight are present in table 1 and structures in figure 1. All 124 reagents were obtained from Sigma-Aldrich, and were purified on silica column 125 chromatography. The molecular weight of TPM compounds were obtained by mass spectrometry. For in vitro assays, a stock solution was prepared in ethanol (EtOH) and maintained at −20°C. All subsequent dilutions were prepared in the respective fresh culture Schnecker’s medium on the day of the assay, and the final maximum concentration of EtOH was 0.1%.

**Promastigotes assay**

Promastigotes of L. (L.) amazonensis, L. (V.) braziliensis or L. (L.) major with 3 or 4 days of growth were plated in 24 well plates at a plating density of 1×10^6 parasites/mL in Schneider’s medium supplemented with 20% FCS, pH 7.2. Drugs were diluted at the same medium and added to a parasite suspension in different concentrations, in triplicate. After 48 h of incubation, the parasites were counted and compared to the controls containing parasites in the absence of drugs. The drug concentration corresponding to 50% of the parasite growth inhibition was expressed as IC50. Three independent experiments were performed to confirm the results. Data are presented as mean and 95% CI.

**Intracellular amastigote assay**

L. (L.) amazonensis and L. (V.) braziliensis promastigotes with 7–8 days and 5–6 days of growth, respectively, were harvested from cultures and added to a fresh Schneider’s medium supplemented with 5% FCS, pH 6.0, and incubated at 32°C for 7 days for L. (L.) amazonensis and 4–5 days for L. (V.) braziliensis until a complete transformation to amastigote-like.

Peritoneal macrophages from BALB/c mice were harvested by washing with ice-cold RPMI 1640 medium, 4 days after induction with a 3% thiglycollate solution. Macrophages were diluted in RPMI 1640 medium (Sigma, Poole, United Kingdom) plus 10% FCS and plated in 24 well plates with a circular cover glass at a plating density of 2×10⁴ macrophages/well. Macrophages were allowed to adhere for 2 h at 37°C and 5% CO₂, when the medium was replaced by a fresh one and incubated overnight.

Macrophages were infected with amastigote-like parasites from L. (L.) amazonensis or L. (V.) braziliensis. The parasites were counted in a Neubauer’s chamber and adjusted to a macrophage-amastigote ratio of 1:3. Infected cultures were maintained at 37°C and 5% CO₂. After 4 h, extracellular parasites were removed by washing, and a fresh medium containing either TPM compounds, no drug, or the reference drug. Amb (Fungizone®-Bristol- Meyers Squibb Pharmaceuticals Ltda, Bedford, USA) was the chosen reference drug and was used at 0.2 μg/mL. After 72 h, the cover glass was removed from the well, washed in RPMI 1640 medium, set in microscope slides, fixed with methanol, and stained with Giemsa for evaluation. Under the immersion microscope, infection indexes (number of amastigotes/100× percentage of infected macrophages) were determined by counting the numbers of intracellular amastigotes in 100 macrophages. The experiments were considered valid only when the control group (without drugs) displayed at least 80% of infection. Each point was tested in triplicate and three independent experiments were then performed. Results are presented as mean and 95% CI [25].

**Cytotoxicity**

To evaluate the toxicity of selected compounds, an in vitro cytotoxicity assay on macrophages from BALB/c mice was performed through 3-(4,5-dimethylthiazol-2-yl)-2,5- diphenyltetrazolium bromide (MTT) (Sigma, Poole, United Kingdom) assay. Briefly, peritoneal macrophages from BALB/c mice were harvested, as described previously and plated in 96-well flat-bottom microplates at a plating density of 1×10⁵ macrophages/well. Macrophages were allowed to adhere for 2 h at 37°C and 5% CO₂, at which time the medium was replaced by a fresh one and incubated over night. Then, the cells were exposed to ten points of serial dilution of TPM 1, 2, 6, 9 or GV (0.195 and 20 μM), which were used to obtain a curve to determine the IC50. After 68 h of incubation, 10 μL of MTT (10 mg/mL) was added to each well and the plates were further incubated for 4 h. The enzymatic reaction was then stopped by addition of 100 μL of 50% isopropanol-10% sodium dodecyl sulfate solution. The optical density at 570 nm was quantified using an ELISA plate reader (BioSource, Inc., EUA). Three independent experiments, in
Figure 1. TPM structures.
doi:10.1371/journal.pone.0051864.g001
triplicate, were performed to determine the cytotoxicity and data were expressed as mean and 95% CI. An experiment was also done to control the possibility of dye color interference on MTT assay. TPM 1, 2, 6, 9 and GV were plated diluted in RPMI for 68 h and then the MTT was added. The optical density at 570 nm was measured using an ELISA plate reader (BioSource, Inc., EUA).

In vivo assay

Gel formulation. The formulation of the GV and TPM 6 gel was prepared by mixing equal amounts of a 2% hydroxyethylcellulose gel (HEC; Natrosol 250 HR, Aqualon) and a 2% GV or TPM 6 hydroethanolic solution (mixture ethanol/water 1/5), until a homogeneous preparation had been attained. Therefore, GV and TPM 6 concentration in these formulations was 1%. The gel lower concentrations for dose-response experiments was obtained by diluting the 1% GV gel with 1% HEC gel.

Treatment of infected animals. BALB/c mice (females, 5–6 weeks old) were inoculated with 1 x 10^7 stationary growth phase promastigotes of L. (L.) amazonensis through subcutaneous injections at the base of the tail, after trichotomy.

To evaluate the in vivo efficacy of GV and TPM 6, after development of ulcerated lesions (average diameter of 7 to 9 mm), BALB/c mice were divided into three groups. For treatment with TPM 6 and GV, lesions were covered with 50 μl of a gel formulation containing either 1% GV or 1% TPM 6, twice a day, for 20 days, using an Eppendorf pipettor. Control group: animals from control group were treated with the gel formulation without the compounds had a low activity against L. (L.) amazonensis,

TPM 7 and TPM 10) the IC50 could not be precisely calculated as for 5 out of 10 compounds evaluated (TPM 3, TPM 4, TPM 5, TPM 4, TPM 5, TPM 6. A linear relationship between the drug concentration and parasitic growth inhibition was obtained for TPM 1, TPM 2, TPM 6, TPM 9 and GV. Table 2 summarizes the data of IC50 obtained. The highest activity was observed for GV (IC50 0.025 μM), followed by TPM 6, TPM 1, TPM 2 and TPM 9. For 5 out of 10 compounds evaluated (TPM 3, TPM 4, TPM 5, TPM 2 and TPM 10) the IC50 could not be precisely calculated as the compounds had a low activity against L. (L.) amazonensis, requiring higher concentrations, which exceeded the maximum EtOH concentration of 0.1% (data not shown). The compounds presenting the highest activity were then selected and tested on L. (L.) major and L. (V.) braziliensis promastigote assays. As observed in TPM 1, TPM 2, TPM 6 and TPM 9, which showed a lower activity against L. (V.) braziliensis than that observed for L. (L.) amazonensis.

Intracellular amastigote assay

The compounds selected from the promastigote assays were subsequently tested on intracellular amastigote assays. Table 3 summarizes the results obtained. TPM 6 was the most effective compound against intracellular amastigotes of L. (L.) amazonensis, followed by GV, TPM 9, TPM 1 and TPM 2. Similar findings were observed for L. (V.) braziliensis, except for GV, which it was not tested against this species. The mean value of parasite growth inhibition observed with the control drug (0.2 μg/ml AmB) was 98% for L. (V.) braziliensis and 99.5% for L. (L.) amazonensis.

Cytotoxicity

The MTT assay was performed to determine the cytotoxicity of TPM 1, TPM 2, TPM 6, TPM 9 and GV. Table 3 summarizes the results of cytotoxicity assays against peritoneal macrophages from

Table 1. Molecular weight and chemical formula for all TPM compounds tested.

<table>
<thead>
<tr>
<th>TPM</th>
<th>Molecular Weight</th>
<th>Formula</th>
</tr>
</thead>
<tbody>
<tr>
<td>TPM1</td>
<td>502.71</td>
<td>C_{23}H_{32}N_{3}</td>
</tr>
<tr>
<td>TPM2</td>
<td>588.87</td>
<td>C_{23}H_{32}N_{4}</td>
</tr>
<tr>
<td>TPM3</td>
<td>455.46</td>
<td>C_{23}H_{32}O_{3}</td>
</tr>
<tr>
<td>TPM4</td>
<td>432.60</td>
<td>C_{23}H_{32}N_{2}</td>
</tr>
<tr>
<td>TPM5</td>
<td>512.69</td>
<td>C_{23}H_{32}O_{2}</td>
</tr>
<tr>
<td>TPM6</td>
<td>400.60</td>
<td>C_{23}H_{32}N_{4}</td>
</tr>
<tr>
<td>TPM7</td>
<td>672.86</td>
<td>C_{23}H_{32}N_{4}</td>
</tr>
<tr>
<td>TPM9</td>
<td>521.76</td>
<td>C_{23}H_{32}N_{4}</td>
</tr>
<tr>
<td>TPM10</td>
<td>662.95</td>
<td>C_{23}H_{32}N_{4}</td>
</tr>
<tr>
<td>VG</td>
<td>407.98</td>
<td>C_{23}H_{32}C_{N}_{3}</td>
</tr>
</tbody>
</table>

doi:10.1371/journal.pone.0051864.t001

Triphenylmethane Activity against Leishmaniasis

10% bovine fetal serum and 100 U/mL penicillin and 100 μg/mL streptomycin. Next, the tissue was centrifuged at 50 g for two minutes for sedimentation (Hitachi, Himac). The supernatant was separated and centrifuged again at 1700 g for 15 minutes (Express, Jouan). The pellet formed was resuspended in 1 mL of Schneider’s modified medium supplemented with 10% FCS and 1% of a 100 U/mL penicillin and 100 μg/mL streptomycin solution. The homogenate was submitted to serial dilutions in duplicates in sterile 96 well culture plates and incubated at 23°C. Each well was examined for the presence of parasites, and the number of parasites was quantified by the highest dilution at which parasites could grow over a 7-day period. The lowest dilution that parasites were detected was 10^{-3}, which was considered the limit of quantification.

Statistical Analysis

The data were processed using MiniTab 15.1 and Sigma Stat 3.5 software. For in vitro assay, IC50 values were calculated by linear regression analysis. The statistical significance of differences among groups was evaluated using the one-way analysis of variance (ANOVA) test followed by Tukey’s test. The Kruskal–Wallis non-parametric test followed by Dunn’s Method for multiple comparisons was used to compare parasite quantification among groups. The difference was considered significant when the p value was less than 0.05.

Results

Promastigote assay

All ten TPM compounds were initially tested against L. (L.) amazonensis promastigotes. Figure 2 shows the results obtained for TPM 6. A linear relationship between the drug concentration and the parasite growth inhibition was obtained for TPM 1, TPM 2, TPM 6, TPM 9 and GV. Table 2 summarizes the data of IC50 obtained. The highest activity was observed for GV (IC50 0.025 μM), followed by TPM 6, TPM 1, TPM 2 and TPM 9. For 5 out of 10 compounds evaluated (TPM 3, TPM 4, TPM 5, TPM 2 and TPM 10) the IC50 could not be precisely calculated as the compounds had a low activity against L. (L.) amazonensis, requiring higher concentrations, which exceeded the maximum EtOH concentration of 0.1% (data not shown). The compounds presenting the highest activity were then selected and tested on L. (L.) major and L. (V.) braziliensis promastigote assays. As observed in table 2, the IC50 determined for these compounds were similar for the three species tested, except for TPM 6, which showed a lower activity against L. (V.) braziliensis than that observed for L. (L.) amazonensis.

Intracellular amastigote assay

The compounds selected from the promastigote assays were subsequently tested on intracellular amastigote assays. Table 3 summarizes the results obtained. TPM 6 was the most effective compound against intracellular amastigotes of L. (L.) amazonensis, followed by GV, TPM 9, TPM 1 and TPM 2. Similar findings were observed for L. (V.) braziliensis, except for GV, which it was not tested against this species. The mean value of parasite growth inhibition observed with the control drug (0.2 μg/ml AmB) was 98% for L. (V.) braziliensis and 99.5% for L. (L.) amazonensis.

Cytotoxicity

The MTT assay was performed to determine the cytotoxicity of TPM 1, TPM 2, TPM 6, TPM 9 and GV. Table 3 summarizes the results of cytotoxicity assays against peritoneal macrophages from
The number of parasites in the control group (2.2 ± 10^6) increased, although a linear dose-response has not been observed. Parastomes within the lesion decreased when gel concentration were 1%. Five animals per group were treated twice a day for 20 days, group. whereas, no parasites were found at lesion site in the GV treated group.

In vivo assay

The quantification of parasites within lesions was used to evaluate the efficacy of different treatments in BALB/c mice. The ratio of cytotoxicity to biological activity was used to determine the selectivity index (SI) of the compounds (Table 3). It is generally considered that biological efficacy is not due to in vitro cytotoxicity when this index is ≥10 [23]. The IC_{50} values observed in macrophage assays for TPM 1, TPM 2, and TPM 9 were higher than that observed for TPM 6 and GV, indicating that those compounds provide lower toxicity to macrophages. However, TPM 6 and GV presented higher selectivity indexes as compared to TPM 1, TPM 2 and TPM 9 for L. (L.) amazonensis infected macrophages. TPM 6 also presented the highest SI for L. (V.) braziliensis infected macrophages (Table 3).

In a dose-effect assay, GV was tested in a gel either at 0.1, 0.5 or 1%. Five animals per group were treated twice a day for 20 days, as above described. As shown in Figure 3B, the number of parasites within the lesion decreased when gel concentration were increased, although a linear dose-response has been not observed. The number of parasites in the control group (2.2 ± 10^6) was higher than that observed in the groups treated with GV at 0.1% (2.2 ± 10^5), 0.5% (2.62 ± 10^5), or 1% (parasites were not detected).

In a dose-effect assay, GV was tested in a gel either at 0.1, 0.5 or 1%. Five animals per group were treated twice a day for 20 days, as above described. As shown in Figure 3B, the number of parasites within the lesion decreased when gel concentration were increased, although a linear dose-response has been not observed. The number of parasites in the control group (2.2 ± 10^6) was higher than that observed in the groups treated with GV at 0.1% (2.2 ± 10^5), 0.5% (2.62 ± 10^5), or 1% (parasites were not detected). Statistical analysis showed a significant reduction in parasite numbers only in 1% GV treated group when compared with the control group (p<0.05).

Discussion

Given the worldwide prevalence of Leishmania infection in countries that have low budgets for health care, finding a safe and inexpensive treatment for leishmaniasis is still an unmet need. In this study, 10 novel TPM were evaluated against promastigotes and amastigotes from 3 species of Leishmania, recognized worldwide as major etiological agents of CL. The most effective compounds proved to be GV and TPM 6 for all the Leishmania species tested. Overall, there was no significant difference in the efficacy of the same compound against the promastigotes of three different species of Leishmania.

Table 2. In vitro anti-leishmanial activity of TPM compounds expressed as IC_{50} (µM) on promastigotes assay.

<table>
<thead>
<tr>
<th>TPM</th>
<th>IC_{50} (µM)</th>
<th>L. (L.) amazonensis</th>
<th>L. (L.) major</th>
<th>L. (V.) braziliensis</th>
</tr>
</thead>
<tbody>
<tr>
<td>TPM 1</td>
<td>0.436</td>
<td>0.567</td>
<td>0.492</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(0.375; 0.497)</td>
<td>(0.521; 0.613)</td>
<td>(0.445; 0.539)</td>
<td></td>
</tr>
<tr>
<td>TPM 2</td>
<td>0.546</td>
<td>0.764</td>
<td>0.551</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(0.462; 0.630)</td>
<td>(0.677; 0.851)</td>
<td>(0.476; 0.626)</td>
<td></td>
</tr>
<tr>
<td>TPM 3</td>
<td>&gt;1.0</td>
<td>n.d.</td>
<td>n.d.</td>
<td></td>
</tr>
<tr>
<td>TPM 4</td>
<td>&gt;2.0</td>
<td>n.d.</td>
<td>n.d.</td>
<td></td>
</tr>
<tr>
<td>TPM 5</td>
<td>&gt;1.0</td>
<td>n.d.</td>
<td>n.d.</td>
<td></td>
</tr>
<tr>
<td>TPM 6</td>
<td>0.031*</td>
<td>0.045^a,b</td>
<td>0.063^b</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(0.026; 0.036)</td>
<td>(0.041; 0.049)</td>
<td>(0.054; 0.072)</td>
<td></td>
</tr>
<tr>
<td>TPM 7</td>
<td>&gt;5.0</td>
<td>n.d.</td>
<td>n.d.</td>
<td></td>
</tr>
<tr>
<td>TPM 9</td>
<td>0.769</td>
<td>0.734</td>
<td>0.839</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(0.614; 0.924)</td>
<td>(0.658; 0.810)</td>
<td>(0.745; 0.933)</td>
<td></td>
</tr>
<tr>
<td>TPM 10</td>
<td>&gt;4.0</td>
<td>n.d.</td>
<td>n.d.</td>
<td></td>
</tr>
<tr>
<td>GV</td>
<td>0.025</td>
<td>0.034</td>
<td>n.d.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(0.016; 0.034)</td>
<td>(0.029; 0.039)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

IC_{50} values correspond to mean and 95% CI of results obtained from triplicates; n.d., not determined; data obtained for linear regression on MiniTab® 15.1 software; a,b p<0.05 compared IC_{50} determined for L.(L.) amazonensis, L. (L.) major and L.(V.) braziliensis.

doi:10.1371/journal.pone.0051864.t002

Table 3. Cytotoxicity, anti-leishmanial in vitro activity and selectivity index (SI) of TPM 1, TPM 2, TPM 6, TPM 9 and GV against L. (L.) amazonensis and L. (V.) braziliensis on intracellular amastigotes assay.

<table>
<thead>
<tr>
<th>TPM Cytotoxicity</th>
<th>L. (L.) amazonensis</th>
<th>L. (V.) braziliensis</th>
</tr>
</thead>
<tbody>
<tr>
<td>IC_{50} (µM)</td>
<td>IC_{50} (µM)</td>
<td>SI</td>
</tr>
<tr>
<td>TPM 1</td>
<td>8.21</td>
<td>0.76</td>
</tr>
<tr>
<td></td>
<td>(7.46; 8.96)</td>
<td>(0.53; 0.99)</td>
</tr>
<tr>
<td>TPM 2</td>
<td>9.49</td>
<td>1.59</td>
</tr>
<tr>
<td></td>
<td>(6.68; 10.30)</td>
<td>(1.25; 1.93)</td>
</tr>
<tr>
<td>TPM 6</td>
<td>4.16</td>
<td>0.10</td>
</tr>
<tr>
<td></td>
<td>(3.18; 5.14)</td>
<td>(0.08; 0.11)</td>
</tr>
<tr>
<td>TPM 9</td>
<td>7.03</td>
<td>0.34</td>
</tr>
<tr>
<td></td>
<td>(6.07; 7.99)</td>
<td>(0.29; 0.39)</td>
</tr>
<tr>
<td>GV</td>
<td>4.03</td>
<td>0.17</td>
</tr>
<tr>
<td></td>
<td>(3.36; 4.70)</td>
<td>(0.16; 0.18)</td>
</tr>
</tbody>
</table>

IC_{50} values correspond to mean and 95% CI of results obtained from triplicates; n.d., not determined; data obtained for linear regression on MiniTab® 15.1 software; mean value of parasite growth inhibition observed for control drug (0.2 µg/ml AmB) was 98% for L. (V.) braziliensis and 99.5% for L. (L.) amazonensis.

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Leishmania (V.) braziliensis, as it was observed for TPM 1, TPM 6, TPM 9 and GV efficacy is not due to Moreover, based on the cytotoxicity data, the selectivity index was demonstrated the highest selectivity indexes (Table 3). In addition, humans have tolerated systemic exposure of GV as well, given that GV is routinely added to transfused blood in areas where Chagas disease is prevalent [19–24].

Limitations of this study may include the use of an extreme susceptible experimental model and the lack of an extended period to follow up lesions healing. It is noteworthy that previous data have indicated that intramuscular administration of Glucantime, at doses equivalent to those used in human chemotherapy, to either hamsters infected L. (V.) braziliensis or BALB/c mice infected with L. (L.) amazonensis did not lead to significant reductions in lesions size [15]. Similar findings were observed when testing topical administration of PA to BALB/c mice infected with either L. (L.) amazonensis or L. (V.) braziliensis [15,32,33].

Thus, for the first time, we have demonstrated that GV and other members of the TPM class are effective against Leishmania species in vitro and in vivo. Humans have a long history of topical GV use as an antibacterial and antifungal agent, and we extend the potential use of GV as a potent antileishmanial agent. In addition, humans have tolerated systemic exposure of GV as well, given that GV is routinely added to transfused blood in areas where Chagas disease is prevalent [19–24].

This study indicates that some TPM derivatives have in vitro anti-leishmanial activity and that this activity is not limited to a single species. In addition, we have demonstrated that topical GV is highly effective against the usually refractory species L. (L.) amazonensis in vivo, perhaps making it an alternative treatment agent where species diagnosis is not possible. The findings described herein are of public health relevance for the following reasons. First, these drugs are inexpensive and stable at room temperature,
making them ideal for use in areas where *Leishmania* is endemic. Second, GV is readily accessible and has an established safety record, making clinical trials rapidly feasible. Finally, gentian violet has anti-angiogenic properties, which might lead to an enhanced host response, in addition to direct anti-parasitic activity.

**References**


**Author Contributions**

Conceived and designed the experiments: RCCdSP LAMF APF. Performed the experiments: RCCdSP LFR ET LF BL. Analyzed the data: RCCdSP JA LAMF APF. Contributed reagents/materials/analysis tools: LF BL AR JA LAMF APF. Wrote the paper: RCCdSP JA APF.