#### **PROTOZOOLOGY - ORIGINAL PAPER**



# Antileishmanial activity evaluation of thiazolidine-2,4-dione against *Leishmania infantum* and *Leishmania braziliensis*

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#### Abstract

Leishmaniasis is responsible for approximately 65,000 annual deaths. Despite the mortality data, drugs available for the treatment of patients are insufficient and have moderate therapeutic efficacy in addition to serious adverse effects, which makes the development of new drugs urgent. To achieve this goal, the integration of kinetic and DSF assays against parasitic validated targets, along with phenotypic assays, can help the identification and optimization of bioactive compounds. Pteridine reductase 1 (PTR1), a validated target in *Leishmania* sp., is responsible for the reduction of folate and biopterin to tetrahydrofolate and tetrahydrobiopterin, respectively, both of which are essential for cell growth. In addition to the in vitro evaluation of 16 thiazolidine-2,4-dione derivatives against *Leishmania major* PTR1 (*Lm*PTR1), using the differential scanning fluorimetry (ThermoFluor®), phenotypic assays were employed to evaluate the compound effect over *Leishmania braziliensis* (MHOM/BR/75/M2903) and *Leishmania infantum* (MHOM/BR/74/PP75) promastigotes viability. The ThermoFluor® results show that thiazolidine-2,4-dione derivatives have micromolar affinity to the target and equivalent activity on *Leishmania* cells. 2b is the most potent compound against *L. infantum* (EC<sub>50</sub> = 23.45 ± 4.54  $\mu$ M), whereas 2a is the most potent against *L. braziliensis* (EC<sub>50</sub> = 44.16 ± 5.77  $\mu$ M). This result suggests that lipophilic substituents on either—*meta* and/or—*para* positions of the benzylidene ring increase the potency against *L. infantum*. On the other hand, compound 2c (CE<sub>50</sub> = 49.22 ± 7.71  $\mu$ M) presented the highest selectivity index.

**Keywords** Dissociation constant  $\cdot$  Phenotypic assay  $\cdot$  *Leishmania braziliensis*  $\cdot$  *Leishmania infantum*  $\cdot$  *Leishmania major*  $\cdot$  Pteridine reductase 1  $\cdot$  ThermoFluor®  $\cdot$  Thiazolidine-2,4-dione

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# Introduction

Leishmaniasis is an infectious disease caused by protozoa of the genus *Leishmania* sp., which is endemic in more than 98 countries, distributed throughout Africa, Asia, and Latin America (WHO 2017; 2019). According to World Health Organization (WHO 2019), the incidence of new cases of leishmaniasis per year is estimated at 700,000 to 1 million, and up to 65 thousand deaths are caused by the different clinical forms (WHO 2019).

Despite the public health impact, the therapeutic arsenal to fight leishmaniasis is limited, what underscores the need for new alternatives to treat the patients infected with different species of *Leishmania* (Gontijo and Carvalho 2003; Swinney and Anthony 2011; Alvar and Arana 2018). One way to achieve this goal is through the inhibition of targets that are unique for the parasite (Eggert 2013). *Leishmania* genus is auxotrophic for free or conjugated pteridines, such as folic acid and biopterin, which

are cofactors in the process of DNA synthesis only in their fully reduced forms (tetrahydrofolate and tetrahydrobiopterin, respectively) (Nare et al. 2009; Leprohon et al. 2015). As a consequence, these parasites developed a complex and versatile biochemical pathway capable of using conjugated and unconjugated pteridines from the host (Nare et al. 1997; Gourley et al. 1999).

In Leishmania sp., the preferred route to obtain tetrahydrofolate  $(H_4F)$  is through the action of the enzyme dihydrofolate reductase-thymidylate synthase (DHFR-TS; E. C 1.5.1.3/ 2.1.1.45), which is considered a validated target for the discovery of new drugs. However, the use of classic antifolates such as methotrexate and trimethoprim is ineffective for the treatment of patients with leishmaniasis. The therapeutic failure is associated with the presence of pteridine reductase 1 (PTR1; EC 1.5.1.33) (Nare et al. 1997). Gene-silencing assays showed that PTR1 is essential for the growth and development of the parasite (Cavazzuti et al. 2008). The selective inhibition of PTR1 is sufficient to kill the parasite, what suggests this is a promising target for the development of new antileishmanial drugs (Vickers and Beverley 2011). Leite et al. (2016) showed that thiazolidine-2,4dione derivatives are micromolar inhibitors of Leishmania major PTR1 that display competitive behavior with the enzyme cofactor (NADPH) (Fig. 1). In order to improve the compound's biological profile, 16 novel thiazolidine-2,4-dione derivatives were designed, synthesized, and evaluated against L. major PTR1. This most active compound (4 Kd = 18.56  $\mu$ M) has a 1.3-fold-higher affinity for LmPTR1 than our previous hit (2d  $Kd = 25.35 \mu$ M). In addition, cell-based assays suggest that high/ moderate selectivity (Leishmania braziliensis (MHOM/BR/75/ M2903) and Leishmania infantum (MHOM/BR/74/PP75) versus WI-26VA4 fibroblast cells) can be achieved, depending on the substituents decorating the benzylidene ring.

### Materials and methods

#### Chemistry

#### Experiment

All melting points were measured in a capillary tube on a Buchi apparatus. Infrared spectra (1% KBr, cm<sup>-1</sup>) pellets were recorded on a Bruker IFS66 spectrophotometer and are uncorrected. The <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra were recorded on a VARIAN VNMRS 400-MR, using 400 MHz for <sup>1</sup>H and 75.4 MHz in DMSO-d<sub> $\delta$ </sub>, using tetramethylsilane (TMS) as the internal standard. Chemical shift values are reported in parts per million units. The <sup>13</sup>C NMR in CDCl<sub>3</sub> and DMSO was maintained at 25 °C using Me<sub>4</sub>Si (TMS) as an internal standard. The following abbreviations were used to indicate the peak multiplicity: s (singlet), d (doublet), dd (double doublet, doublet), t (triplet), and m (multiplet). The chemical shifts were reported in  $\delta$  units, and the

coupling constants (J) were reported in Hertz. Mass spectra were recorded on a Varian MAT 711 spectrometer 70-eV electron impact. Thin-layer chromatography (TLC) was performed on pre-coated silica plates (Merck Kiesegel 60  $F_{254}$ ), and the spots were visualized under ultravioleta light (254 nm/long (365 nm) UV wavelength). The chemical reagents were supplied by Sigma-Aldrich (USA) and were used without further purification. Purity of the compounds was confirmed by using TLC (Kiesel gel 60 G F 254) using the appropriated system for each compound.

#### Procedure for preparation synthesis of thiazolidine-2,4-dione

The thiazolidine-2,4-dione was obtained by using the method described by Libermann in 1948 and modified by Albuquerque in 1995 (Albuquerque et al. 1995; Libermann et al. 1948; Gouveia et al. 2009). This reaction occurs by condensation of monochloroacetic acid and thiourea in an aqueous medium under reflux for 24 h (Bozdag-Dündar et al. 2007; Bruno et al. 2002) (Fig. 2). Molecular formula  $C_3H_3O_2NS$ ; yield 78%; mp 118–120 °C; Rf 0.48 (0.9:0.1 CHCl<sub>3</sub>/MeOH). Recrystallization: water.

#### General method for the synthesis of 5-arylidene-thiazolidine-2,4-diones (1–16)

The compounds of 1–16, in turn, have the description of the synthesis below.

#### General method for the synthesis of thiazolidine-2,4-dione

The compounds (1-16) were synthesized in two steps. In step 1, the thiazolidine 2,4-dione compound was synthesized using monochloric acetic acid and thiourea. The solvent used was water. The reagents were heated at 100 °C for 16 h. The compounds 2a-s was prepared by using a simple method, which involve the Knoevenagel condensation with different aromatic aldehydes. The structures of the desired compounds were determined by <sup>1</sup>H NMR, <sup>13</sup>C NMR, IR, and mass spectrometry.

### General method for the synthesis of 5-arylidene-thiazolidine-2,4-dione (1–16)

The compounds 1–16 and compounds 2a–2e were synthesized from a mixture of thiazolidine-2,4-dione (1—Fig. 2) (2.5 g, 21.36 mmol), aldehyde (21.36 mmol), piperidine (14.11 mmol), and ethanol (150 mL). The reaction mixture was heated under reflux and continuously stirred for a period of 16 h. The course of the reaction was monitored by using TLC. The reaction mixture was poured into water and acidified with acetic acid. The resulting precipitate was filtered off and recrystallized from acetic acid to give (2—Fig. 2).



(Z)-5-(3-methoxy-4-hydroxy-benzilydene)-thiazolidine-2,4dione (1) Yield 70%; mp 260 °C; Rf 0.55(CH<sub>2</sub>Cl<sub>2</sub>/MeOH 9:1) Recrystallization: ethanol; IR (KBr 1%,  $v_{max}$  cm<sup>-1</sup>) 1566 (C=C); 1730–1670 (C=O). <sup>1</sup>H NMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$ ppm): 8.48 (s 1H, NH); 7.71 (s 1H, CH=); 3.77 (s 3H, OCH<sub>3</sub>); 6.02 (s 1H, OH). 6.92 (s 1H<sub>(2)</sub>); 6.86 (d 1H<sub>(5)</sub> J= 8.33); 6.91 (d 1H<sub>(6)</sub> J = 8.34); <sup>13</sup>C NMR (Acetone-d<sub>6</sub>, 75.4 MHz,  $\delta$  ppm): 169.81 (C=O<sub>2</sub>); 166.89 (C=O<sub>4</sub>); 119.55 (C<sub>5</sub>, heterocycle); 143.28 (CH=); 56.50 (OCH<sub>3</sub>); 129.20 (C<sub>1</sub>); 112.58 (C<sub>2</sub>), 148.06 (C<sub>3</sub>); 144.76 (C<sub>4</sub>); 115.56 (C<sub>5</sub>); 122.30 (C<sub>6</sub>). Anal. Calcd. for C<sub>11</sub>H<sub>9</sub>NO<sub>4</sub>S: C, (52.58%); H, (3.61%); N, (5.57%); Found: C, 53.01%; H, 3.83%; N, 5.25%. HRMS<sup>+</sup>: calcd 251.0252; found 251.0253.

(Z) 5-(2,4-dimethoxy-arylidene)-thiazolidine-2,4-dione (2) Yield 67%; MP 250 °C; MF  $C_{12}H_{11}NO_4S$ ; MW 265.2850; Rf 0.50 (CHCl<sub>3</sub>/MeOH 9:1 Recrystallization: ethanol; IR (KBr 1%,  $\nu_{max}$  cm<sup>-1</sup>) 1571 (C=C); 1733–1691 (C=O). <sup>1</sup>H NMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$  ppm): 11.30 (s 1H, NH); 8.25 (s 1H, CH=); 3.86 (s 3H, OCH<sub>3</sub> (<sub>orto</sub>); 6.57 (s 1H, H3); 3.87 (s 3H, OCH<sub>3 para</sub>); 6.0 (d 1H<sub>(6)</sub> J = 8.34); 11.40 (s 1H, NH, (imide). HRMS<sup>+</sup>, calcd 250.0048; found 265.0409.

(Z) 5-(2-nitro-benzylidene)-thiazolidine-2,4-dione (3) Yield 83%; MP 205–206 °C; MF  $C_{10}H_6N_2O_4S$ ; MW 250.2306; Rf 0.50 (Hex: Ethyl acetate 9:1) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>) 1569 (C=C); 1736–1660 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$  ppm): 11.97 (s 1H, NH

(imide)); 8. 57 (d 1H C-H), 7.81 (dd 2H, H<sub>4</sub>, J = 7.3), 7.77 (dd 1H, H<sub>5</sub>) J = 7.6; J = 1.5, 8.13 d, 1H J = 7.6 (H<sub>6</sub>). HRMS<sup>+</sup>, calcd. 250.0048; found 250.0049.

(Z) 5-(2,3,5-Trichloro-benzylidene)-thiazolidine-2,4-dione (4) Yield 73%; MP 170.5–171.0 °C; MF  $C_{10}H_4Cl_3NO_2S$ ; MW 308.5683. Rf 0.50 (CHCl<sub>3</sub>/MeOH 9:1) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>) 1569 (C=C); 1738–1667 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$  ppm): 11.97 (s 1H, NH (imide)); 8. 35 (s 1H C=H), 7.55 (s 1H, (H<sub>4</sub>). 7.23 (s 1H, H<sub>6</sub>) HRMS<sup>+</sup>, calcd. 306.9028; found 306.9028.

(Z)-5-(3-methoxybenzylidene) thiazolidine-2,4-dione (5) Yield 83%; MP 170–171 °C; MF C<sub>11</sub>H<sub>9</sub>NO<sub>3</sub>S; MW 235.2591; Rf 0.50 ((Hex: Ethyl acetate 5:5) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>) 1564 (C=C); 1733–1665 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$  ppm): 11.87 (s 1H, NH (imide)); 7.87 (s 1H C=H), 7.19 (d 1H, H<sub>2–4</sub>, J=7.5), 3.83 (s 3H OCH<sub>3</sub>) 7.56 (t 1H, H<sub>5</sub>) J=7.6; J=1.5), 6.90.(d,1H J=7.6 (H<sub>6</sub>). HRMS<sup>+</sup>, calcd. 235.0303; found 235.0303.

(Z) 5-(3-hydroxy-benzylidene)-thiazolidine-2,4-dione (6) Yield 69%; MP 265–266 °C; MF  $C_{10}H_7NO_3S$ ; MW 221.2025; Rf 0.50 ((Hex: Ethyl acetate 5:5) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>) 1568 (C=C); 1738–1669 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$ ppm): 12.31 (s 1H, NH (imide)); 7.97 (s 1H C=H), 6.72 (d 1H, H<sub>2</sub>, J = 7.48), 5.37 (s 1H OH) 6.86 (d 1H, H<sub>4</sub>) 7.56.(s 1 H)

**Fig. 2** Synthetic route of thiazolidine-2,4-dione compounds



R=a) R=4-OH; 3-OCH3; b) 2,4-OCH3; c) 2-NO2; d) 2,3,5-Cl; e) 3-OCH3, f) 3-OH); g ( 3,4-OCH3 ; h) 4-S-CH3; i) 2-Cl; j) 1,4-Biphenyl; k) 3- indol; l) 3-Indol-N-COCH3; m) 4-NO2,3-indol; n) 2-Methyl-phenylallyl

7.56; J = 1.5), 7.19 (d,1H, J = 7.16; (H<sub>6</sub>). (d,1H) HRMS<sup>+</sup>, calcd. 235.0303; found 235.0303.

(Z) 5-(3,4-dimethoxy-benzylidene)-thiazolidine-2,4-dione (7) Yield 73%; MP 213–214 °C; MF  $C_{12}H_{11}NO_4S$ ; MW 265. 0409; Rf 0.46 ((Hex: Ethyl acetate 7:3) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>) 1568 (C=C); 1735–1668 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$  ppm): 12.31 (s 1H, NH (imide)); 7.95 (s 1H C=H), 7.24 (s 1H, H<sub>2</sub>), 3.84 (s 3H OCH<sub>3</sub>) 3.86 (s 3H, OCH<sub>3</sub>) 6.96 (d,1H, J = 7.7; (H<sub>5</sub>). 7.21 (d,1H, J = 7.6; (H<sub>6</sub>). HRMS<sup>+</sup>, calcd. 265.0409; found 265.0409.

(Z) 5-(4-methylthio)-benzylidene)-thiazolidine-2,4-dione (8) Yield 63%; MP 209–210 °C; MF  $C_{11}H_9NO_2S_2$ ; MW 251.3247–251.3251. Rf 0.65 (Hex: Ethyl acetate 6:4) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>) 1571 (C=C); 1735–1668 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$ ppm): 12.31 (s 1H, NH (imide)); 7.95 (s 1H C=H), 7.24 (s 1H, H<sub>2</sub>), 3.84 (s 3H OCH<sub>3</sub>) 3.86 (s 3H, OCH<sub>3</sub>) 6.96 (d,1H, J = 7.7; (H<sub>5</sub>). 7.21 (d,1H, J = 7.6; (H<sub>6</sub>). HRMS<sup>+</sup>, calcd. 265.0409; found 265.0409.

(Z) 5-(2-Chloro benzylidene)-thiazolidine-2,4-dione (9) Yield 70%; MP 165.9–166.5 °C; MF  $C_{10}H_6CINO_2S$ ; MW 239.6781. Rf 0.50 (Hex: Ethyl acetate 6:4) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>) 1568 (C=C); 1733–1667 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$  ppm): 12.29 (s 1H, NH (imide)); 8.23 (s 1H C=H); 7.54 (d 1H, H<sub>3</sub> J = 7.5; 1.4), 7.05 (t 3H (H<sub>4</sub>) J = 7.4; 1.5) H<sub>5</sub> 7.25 (t 2H, (H<sub>5</sub>) J = 7.4; 1.5) 7.28 (d,1H, J = 7.7) (H<sub>5</sub>). 7.28 (d,1H, J = 7.6; (H<sub>6</sub>). HRMS<sup>+</sup>, calcd. 238.9808.; found 238.9808.

(Z) 5-([1,1'-biphenyl]-4-ylmethylene) thiazolidine-2,4-dione (10) Yield 78%; MP 238.5–239.8 °C; MF  $C_{16}H_{11}NO_2S$ ; MW 281.0510. Rf 0.53 (CH<sub>2</sub>Cl<sub>2</sub>/Acetone (0.53) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>) 1567 (C=C); 1731–1665 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$ ppm): 12.29 (s 1H, NH (imide); 7.99 (s 1H C=H); 7.56 (d 1H, H<sub>2</sub>-H<sub>2'</sub>,J=7.5; 1.4), 7.74. (d 2H (H<sub>3</sub>-H<sub>3'</sub>) J=7.4; 1.5) 7.40 (t 2H (H<sub>6</sub>-H<sub>6'</sub>) J=7.5–1.50; 1.50. (t 2H, (H<sub>7</sub>-H<sub>7'</sub>) J= 7.6; 1.5; 1.5). 7.40 (t 2H, (H<sub>6</sub>-H<sub>6'</sub>) J=7.6; 1.5; 1.5). HRMS<sup>+</sup>, calcd. 281.0510.; found 281.0510.

(Z) 5-((1H-indol-3-yl) methylene) thiazolidine-2,4-dione (11) Yield 78%; MP 165.9–166.5 °C; MF  $C_{12}H_8N_2O_2S$ ; MW 239.6781. Rf 0.50 (Hex: Ethyl acetate 6:4) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>) 1567 (C=C); 1733–1667 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$  ppm): 12.29 (s 1H, NH (imide)); 8.11 (s 1H C=H); 7.76 (d 1H, H<sub>2</sub> indol J = 7.5; 1.4); 12.0 (d 1H (NH)) 7.05 (d 1H (H<sub>1</sub>, Ar) J = 7.4; 1.5); 7.11 (t 2H, (H<sub>2</sub>) J = 7.4; 1.5; 1.5) 7.18 (t, 2H, (H<sub>3</sub>). J = 7.7; 1.5, 1.5)) 7.57 (d, 1H, J = 7.6, H<sub>6</sub>). HRMS<sup>+</sup>, calcd. 244.0306; found 244.0306. (Z)5-((6-methylene-4-oxo-4H-chromen-3-yl) methylene)) thiazolidine-2,4-dione (12) Yield 72%; MP 287.5-  $^{0}$ C; MF C<sub>14</sub>H<sub>9</sub>NO<sub>4</sub>S; MW 287.2906. Rf 0.5 ((Hex: Ethyl acetate 6:4) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>) 1559 (C=C); 1730–1666 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$  ppm): 12.29 (s 1H, NH (imide)); 7.43 (s 1H C=H); 6.71 (s 1H (H<sub>1</sub>)) 7.63 (d 1H (H<sub>2</sub>)) 7.67 (d 1H (H<sub>3</sub>), Ar) J = 7.47.2.44 (s 3H, (CH<sub>3</sub>) 7.41 (s 1H), HRMS<sup>+</sup>, calcd. 289.0409; found 289. 0409.

(Z)-5-((4-nitro-1H-indol-3-yl)methylene)thiazolidine-2,4-dione (13) Yield 57%; MP 129.5–131.0 0C; MF C12H7N3O4S; MW 289.2667; Rf 0.54 (Dichloro metane: Methanol 9.9:0.1) Recrystallization: ethanol, IR (KBr 1%,  $\nu$ max cm-1) 1560 (C=C); 1729–1657 (C=O). 1HNMR (CDCl3, 300 MHz,  $\delta$  ppm): 12.29 (s 1H, NH (imide)); 7.21 (s 1H C-H); 7.95 (t 2H (H2) J = 7.63, J = 1.4 J = 1.6). 7.23 (d 1H (H3) J = 7.47); 9.9 (d 1H J = 7.80 (NH), 8.81 (d 1H, J = 7.79). HRMS+, calcd. 289.0157; found 289.0156.

(Z)-5-(quinolin-4-ylmethylene)-thiazolidine-2,4-dione (14) Yield 83%; MP 273–274 °C; MF  $C_{14}H_{10}N_2O_2S$ ; MW 270.3036. Rf 0.50 ((Hexane Ethyl acetate 9:1) Recrystallization: ethanol, IR (KBr 1%,  $\nu$ max cm<sup>-1</sup>) 1559 (C=C); 1733–1661 (C=O). 1HNMR (CDCl3, 300 MHz,  $\delta$  ppm): 12.26 (s 1H, NH (imide)); 7.82 (s 1H C-H); 6.73 (t 2H (H<sub>2</sub>) J = 7.60, J = 1.5 J = 1.5). 7.76 (t 1H (H<sub>3</sub>) J = 7.45; 1.5, 1.5) 8.12 (d 1H (H<sub>4</sub>), 8.75 (d 1H, J = 7.47, J = 1.5) 7.12 (d 1H, J = 7.3, J = 1.5 HRMS<sup>+</sup>, calcd. 256.0306; found 256.0306.

(Z)-5-((5-Bromothiophen-2-ylmethylene)-thiazolidine-2,4dione (15) Yield 83%; MP 180.3–180.7 °C; MF  $C_8H_4BrNO_2S_2$ ; MW 290.1490. Rf 0.50 (CHCl<sub>3</sub>/MeOH 9:1) Recrystallization: ethanol, IR (KBr 1%,  $\nu$ max cm<sup>-1</sup>) 1558 (C=C); 1735–1662 (C=O). 1HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$  ppm): 12.25 (s 1H, NH (imide)); 7.55 (s 1H C-H); 7.55 (d 1H (H<sub>1</sub>, C=H J=7.50); 7.24 (d 2H (H<sub>2</sub>,-H<sub>2</sub>' J=7.46; J=1.45; J=1.45) 7.45 (d d H<sub>3</sub>-H<sub>5</sub> CH) J=7.24. HRMS<sup>+</sup>, calcd. 288.01878; found 288.01878.

(Z)-5-((E)-2-methyl-phenylallylidene (thiazolidine-2,4-dione) (16) Yield 79%; MP 204.6–205.2 °C; MF  $C_{13}H_{11}NO_2S$ ; MW 245–246. Rf 0.49 ((Hexane: Ethyl acetate 9.9:0.1) Recrystallization: ethanol, IR (KBr 1%, vmax cm<sup>-1</sup>); 1555 (C=C); 1733–1665 (C=O). <sup>1</sup>HNMR (CDCl<sub>3</sub>, 300 MHz,  $\delta$ ppm): 12.27 (s 1H, NH (imide)); 7.41 (C=C; 6.37 (s 1H C-H); 2.21 (s 3H CH<sub>3</sub>); 6.34 (s 1H CH) 7.61 (d 2H (H<sub>2</sub>,-H<sub>2</sub>·) J=7.46; J=1.45; J=1.45) 7.45 (d 2H (H<sub>3</sub>-H<sub>5</sub>·) CH) J=1.45; J=1.45) 7.33 (t 2H. HRMS<sup>+</sup>, calcd. 245.0510 found 245.0510.

The 5-arylidene thiazolidine-2,4-diones compounds (2a–e) were synthesized as described in Leite et al. 2016.

#### In vitro assays

#### ThermoFluor® assays

Thermal shift assays were carried out using a CFX Touch RT-PCR Detection System (Bio-Rad CFX Manager®, Bio-Rad Laboratories, Inc.) fitted with custom filter sets. The data was recorded in the CFX Manager<sup>™</sup> v 3.1 Software. Initially, conditions of the assay (choice of buffer, protein concentration, and DMSO concentration) were optimized for LmPTR1. Each parameter was tested in triplicate on a 96-well PCR plate (PCR plates 96-well Bio-Rad®), manually sealed with transparent capping strips (Flatcap strips Bio-Rad®). The final protein concentration, for optimization trials, was set to 5 µM, except for the determination of optimum PTR1 concentration, which varied from 1 to 5 µM. Fluorescence changes were monitored with SYPRO Orange® dye, using 492- and 610-nm wavelengths for excitation and emission respectively. Thermal shift assays were carried out from 25 to 85 °C in increments of 1 °C per minute. Fluorescence data (raw data) was recorded using the CFX Manager<sup>™</sup> v 3.1 Software, and then exported to Excel 2007 worksheet (ftp://ftp.sgc.ox.ac.uk/ pub/biophysics) for processing and analysis. Tm values employed for compound comparison were those calculated from the non-linear fitted melting curves (Boltzmann sigmoidal function) implemented in GraphPad Prism version 5.0 for Windows (GraphPad® Software, San Diego, CA, USA, www.graphpad.com).

#### Single-concentration assays

The PTR1 expression and purification steps were carried out as described previously (Leite et al. 2016). All compounds were assayed at the final concentration of 50  $\mu$ M. In each well, a solution containing 4  $\mu$ L of LmPTR1 (to obtain the final concentration, 5  $\mu$ M), 14  $\mu$ L of 20-mM sodium acetate buffer pH 4.7, 1- $\mu$ L SYPRO Orange (Invitrogen<sup>TM</sup>) (5X), and 1- $\mu$ L of each compound was added. The values of T*m* obtained in the presence of each compound were compared with the value of the reference solution, which contains 5% DMSO P.A. (v/v), instead of the inhibitor. The assays were performed in triplicate.

#### Concentration-response assays

Compounds with  $\Delta Tm > 1.0$  °C, in single-concentration assays, also had their effect evaluated at the following concentrations: 6.25, 12.5, 25, 50, and 100 µM. The obtained Tm values were compared with the value of the reference solution, which contains 5% DMSO P.A. (v/v), instead of the inhibitor. All assays were carried out in triplicate.

# Determination of the dissociation constant by ThermoFluor $\ensuremath{^{\scriptscriptstyle \odot}}$

Compounds that show a concentration-response behavior, in the previous step, had their dissociation constants (K*d*) determined according to the equation established by Vivoli et al. (2014), using data from ThermoFluor assays carried out in the presence of the following concentrations of each compound: 6.25, 12.5, 25, 50, 100.0, 200.0, and 400.0  $\mu$ M. The five compounds that presented the best  $\Delta$ T*m* values in the study by Leite et al. (2016) also had their dissociation constants determined, following the same concentrations and previous parameters of ThermoFluor® assays. The assays were performed in triplicate.

#### **Cellular assays**

#### **Cultivation of cell lines**

The human cell line WI-26VA4 (pulmonary fibroblast ATCC CCL-75) was used for toxicity analysis. This lineage is part of the animal cell bank of the Cell Biology Service (SBC) of the Ezequiel Dias Foundation (FUNED) in Belo Horizonte - Minas Gerais.

Cells were cultured from cryopreserved ampoule; they were thawed at 37 °C; their contents were transferred to a 15-mL tube containing 10-mL RPMI 1640 medium. Cells were centrifuged at  $1200 \times g$  for 5 min; the supernatant was discarded, and the pellet was resuspended in RPMI 1640 medium supplemented with 10% heat-inactivated fetal bovine serum (SBF) (complete medium). The cells were transferred to plastic bottles of treated cell culture to promote cell adhesion, T75 (75 cm<sup>2</sup>) 10 mL (Corning Costar Inc., USA), and maintained as monolayers at 37 °C in a greenhouse (Thermo electron co.) humid atmosphere of 5% CO2. The culture medium was replaced every 48 h of incubation. Cellular morphology and monolayer formation were observed with the aid of an inverted microscope at ×100 magnification (Olympus model, CKX 41). After confluence of 80% of the culture in the T75 bottle, the cells were replicated or used in the cytotoxicity assays (Patra et al. 2011; Pereira et al. 2012).

#### Cytotoxicity assays

Cells with 80% confluency were trypsinized with 1.5 mL of trypsin (1:250 Sigma), incubated at 37 °C for 5 min, resuspended in 10 mL of complete RPMI medium, and centrifuged at  $1200 \times g$  for 5 min. The supernatant was discarded, and the pellet was resuspended with complete RPMI medium. Cells were distributed into 96-well microplates at a density of 4 ×  $10^5$  cells/100 µL per well and incubated in a CO<sub>2</sub> oven at 37 °C for 24 h for adhesion to the plate. After this period of adhesion, the medium was removed, a new RPMI medium

supplemented with 1% FBS was added, and then the diluted compounds were added in RPMI medium containing different concentrations (0.01, 0.10, 1.0, 10, and 100  $\mu$ g/mL). The plates were incubated for an additional 48 h.

At the end of the treatment period, 100  $\mu$ L/well of a solution of 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazole (MTT) (Sigma-Aldrich) bromide, tetrazole salt, concentration of 0.5 mg/mL in RPMI 1640 without phenol red (Denizot and Lang 1986) were add in the plate. After 3 h of incubation in the CO<sub>2</sub> incubator at 37 °C in the dark with MTT, the supernatant was aspirated and the formazan crystals were dissolved in 50  $\mu$ L/well of DMSO P.A. and held at 37 °C for 10 min. The absorbance per well was measured at a wavelength of 550 nm using the Gen5 software (Data-Analysis Software-Bio-Tek). The assays were performed in triplicate.

#### Cultivation of Leishmania sp.

The parasite cultures were carried out with reference strains of *Leishmania* (V.) *braziliensis* (MHOM/BR/75/M2903) and *Leishmania* (L.) *infantum* (MHOM/BR/74/PP75) belonging to the Laboratory of Parasitic Infectious Diseases of the UFSJ/CCO (LAB-DIP/CCO).

The parasites were cultured in biphasic medium, the liquid phase being composed of Schneider's Insect Medium with Lglutamine and sodium bicarbonate (Sigma®), plus 10% Fetal Bovine Serum (FBS) and 1% antibiotic Pen Strep-GIBCO (10,000 units/mL penicillin and 10,000-µg/mL streptomycin), and maintained in solid phase with NNN culture medium (Novy, McNel, and Nicolle) plus 1% antibiotic Pen Strep for storage. The parasites were kept in bottles for cell culture in polystyrene, growth volume of 25 cm<sup>3</sup>, sterile, and in BOD greenhouse at 25 °C. For the maintenance of the parasites, peaks were performed every 4 days. All procedures involving the maintenance and cultivation of the strains were carried out in a laminar flow hood, avoiding any type of contamination by microorganisms. Follow-up of the cultures was carried out by observation in a ZEISS optical microscope, AXIOSTAR PLUS model, by the preparation of crop slides, in order to observe flagellar mobility and absence of contamination.

#### Antileishmania activity assays

Antileishmania activity assays were performed by the same cytotoxicity test, i.e., the MTT chemosensitivity test. After 48 h of boiling, to allow the parasites to grow, that is, in the log phase (ideal phase for testing the compounds), the viability of the parasites was evaluated by microscopy. Aliquots of the viable parasites were removed from the growth bottles, and the ideal volume was calculated for a final concentration of  $1.0 \times 10^6$  parasites/well. The reaction volume per well was 200 µL, with 20 µL corresponding to the compounds to be tested (at the concentrations of 1.5625, 3.123, 6.25, 12.5, 25,

50, and 100  $\mu$ g/mL in triplicate for the compounds and 50  $\mu$ g/mL for amphotericin B deoxycholate) and 180  $\mu$ L of parasite at the concentration of  $1.11 \times 10^6$  parasites/mL for each well in 96-well microplates. From this, the incubation was incubated at 26 °C for 48 h.

At the end of the incubation period, the MTT was diluted in incomplete Schneider's medium (without fetal bovine serum), so that it was at a concentration of 5 mg/mL. From this moment, the experiments were performed with the laminar flow hood light off, since the MTT is photosensitive. A volume of 20- $\mu$ L MTT was added to the wells. The plates were then wrapped in aluminum foil and incubated for 4 h at 37 °C. The plates are maintained at this temperature, as the compounds have already acted, the temperature no longer affects the viability of *Leishmania* protozoan, and some proliferative capacity is diminished. It is also worth noting that in low temperatures, the production of formazan salts is lower.

After 4 h of incubation with MTT, the plates were centrifuged at  $3500 \times g$  for 10 min at 4 °C in the plate centrifuge. At the end of centrifugation, the supernatant was carefully removed to preserve the formazan crystals and parasites, 50 µL/well of DMSO P.A. was added to dissolve the formazan, and the plates were homogenized and allowed to stand for 15 min for reading in ELISA reader at 570 nm using the Gen5 software (Data-Analysis Software-Bio-Tek). The assays were performed in triplicate.

#### Calculation of the selectivity index

After determination of the effective concentration to kill 50% of the parasites ( $CE_{50}$ ) and determination of the lethal dose capable of killing 50% of human cells ( $LD_{50}$ ), the selectivity index (SI) was calculated according to the equation:

 $\mathrm{SI}=\mathrm{LD}_{50}/\mathrm{CE}_{50}.$ 

# **Results and discussion**

## Synthesis and characterization of thiazolidine-2,4dione

Leite et al. (2016) showed that thiazolidine-2,4-dione derivatives inhibit LmPTR1 at micromolar range concentration. Compound 2d, the most promising compound identified in that study, was used as a prototype to design novel compounds with optimized effect against *Leishmania* cells in vitro. A total of 16 compounds have been synthetized based on features of the prototype and their structural variations: compounds 1 to 10 (except for 2, 3, and 6) were synthetized based on chemical viability, while 11–15 were designed to explore additional LmPTR1 pockets nearby the active site. Compounds 2, 3, and 9 were designed to the effect of bulk substituents (steric clashes) on binding affinity, whereas compound 16 aimed at investigating the effect of a spacer moiety, between the two rings on the binding profile.

#### ThermoFluor<sup>®</sup> assays

# Single-concentration assays for the identification of LmPTR1 inhibitors

Sixteen structural analogs of compound 2d were evaluated for their ability to interact with LmPTR1. ThermoFluor® is a high/medium throughput screening that has enabled the prioritization of potential molecules due to the ability to evaluate the (de)stabilizing effect of hundreds of compounds over the tertiary structure of the protein and, therefore, their ability to bind to the target and generate a biological response. Molecules with  $\Delta Tm > 1.0$  °C are described as promising and tend to confirm the biological potential in secondary assays (Pantoliano et al. 2001; Ericsson et al. 2006; Cimmperman and Matulis 2011; Kopec and Schneider 2011). This technique has already been employed to identify thiazolidine-2,4-dione that inhibit LmPTR1 (Leite et al. 2016) or that inhibit both DHFR-TS and PTR1 (Teixeira et al. 2019). Compounds with higher degree of affinity to the target tend to have higher  $\Delta Tm$  when all other variables are kept equal (Zaia et al. 1998; Pantoliano et al. 2001; Ericsson et al. 2006; Niesen et al. 2007; Zhang and Monsma 2010). Among the 16 novel thiazolidine-2,4-dione derivatives evaluated by a singleconcentration ThermoFluor® assay against LmPTR1 (Graph 1), only two compounds, 4 ( $\Delta Tm = 12.12 \pm 0.34$  °C) and 9 ( $\Delta Tm = 4.69 \pm 0.21$  °C), may be considered as potential inhibitors against pteridine reductase 1 of L. major.

However, both compounds have lower  $\Delta Tm$  values than compound 2d, (compound 4  $\Delta Tm = 12.12 \pm 0.34$  °C and compound 9  $\Delta Tm = 4.69 \pm 0.21$  °C versus compound 2d  $\Delta Tm = 44.67 \pm 1.74$  µM). Single-concentration assays also show that compound 12 shows negative thermal displacement. Some studies have shown that compounds with this profile can stabilize the target with whom they interact in a conformation other the native one. However, a direct relationship between negative values of thermal shift and biological profile has not been described (Waldron and Murphy 2003; Dupeux et al. 2011; Garbett and Chaires 2012; Sancho 2013; Contessoto et al. 2018).

Thus, only molecules 4 and 9 were evaluated at different concentrations (6.25, 12.5, 25, 50, and 100  $\mu$ M) to probe if the thermal stabilization follows a dose-dependent behavior. Both compounds were found to increase LmPTR1 thermal stability at higher concentrations, with  $\Delta Tm$  values for compound 4 higher than those seen for compound 6 (Graph 2). For compound 4, a significant increase (p < 0.01) in  $\Delta Tm$  is observed between 6.25-µM and 12.5-µM and between 12.5-µM and 25-µM concentrations. Higher concentrations afford negligible increases in LmPTR1 Tm. Noteworthy is an eight-fold increase in thermal shift between the lowest and highest ligand concentration. In contrast, compound 9 showed a significant increase (p < 0.01) in  $\Delta Tm$  only at low concentrations (between 6.25  $\mu$ M and 12.5  $\mu$ M), and despite the overall tendency to increase LmPTR1 TM at higher concentrations, only a 6fold increase in thermal shift is observed if one compares the lowest and the highest concentration of compound 9.

# Determination of apparent dissociation constant by ThermoFluor®

The inhibition constant (K*d*) can also be determined by ThermoFluor® assays (Lo et al. 2004). This assay does not require a high degree of sample purity and does not have to include solution enthalpy in the K*d* calculation, which sometimes makes it difficult to characterize (Cimmperman and Matulis 2011; Craig and Newton 1991; Vivoli et al. 2014). Alternatively, Vivoli et al. (2014) developed a method to calculate the apparent dissociation constant (K $d_{app}$ ) from





**Graph 1** Thermal stability of LmPTR1 in the presence of the thiazolidine-2,4-dione derivatives in the single-concentration assay. The values represent the median and interquartile range of the variation of Tm

( $\Delta$ Tm). Statistical differences were calculated between each compound (n = 3) and compound 2d, \*\*\*p < 0.001 (Kruskal-Wallis test with Dunn's post-test), previously reported by Leite et al. (2016)



**Graph 2** Effect of different concentrations of compound 4 (a) or 9 (b) over the *Lm*PTR1 thermal stability. The values represent the median and interquartile range of the variation of  $Tm(\Delta Tm)$  by multiple comparisons (n = 3), \*\*p < 0.01. (Kruskal-Wallis test with Dunn post-test)

information, provided by differential scanning fluorimetry assays, which in most cases is very close to the real Kd value (Petrauskas et al. 2019).

In order to make a fair comparison of the current compounds to those already published (Leite et al. (2016)), not only compounds 4 and 9 had their  $Kd_{app}$  calculated but also compounds 2a–2e (Table 1).

According to this method, compound 4 (18.56  $\mu$ M) has the lowest K $d_{app}$  value, followed by compound 2b (23.12  $\mu$ M), 2d (25.35  $\mu$ M), 2e (89.98  $\mu$ M), and 9 (98.11  $\mu$ M). Hence, compound 4 seems to have higher affinity to LmPTR1 than our previous lead compound (2d). Moreover, compounds with  $Kd_{app} \leq 100 \mu$ M are expected to have moderate to high affinity for the target (Vivoli et al. 2014). As compound 2d is among the most promising compounds, the assumptions made by Cimmperman and Matulis (2011) and Vivoli et al. (2014) are in good agreement with our previous results (Leite et al. 2016), since the IC<sub>50</sub> is related to the ability of a compound to inhibit the catalytic activity of the target at a fixed substrate concentration and Kd accounts for the ability of a ligand to bind with the target.

#### **Cellular assays**

It is known that the use of enzymatic assays for the characterization of specific target inhibitors has been commonly used

**Table 1** Apparent dissociation constant values (Kd) of the seven compounds screened with  $\Delta Tm > 1$  °C. The assays were performed in triplicate, and these were analyzed by non-linear regression with p < 0.05 by using the program GraphPad Prima version 5.0

Compounds	K <i>d</i> (μM)	$R^2$			
2a	43.42	0.97			
2b	23.12	0.94			
2c	196.40	0.97			
2d	25.35	0.99			
2e	89.98	0.99			
4	18.56	0.92			
9	98.11	0.99			

since the advent of molecular cloning (Pikkemaat et al. 2002; Gasteiger et al. 2005; Sievers et al. 2011; Priest and Erdemli 2014). However, this approach is considered a simplistic view of the effect of the compound against the pathology (Grasberger et al. 2005; Lesyk et al. 2006; Vicini et al. 2006; Silva et al. 2014; Wagner and Schreiber 2016). An example to be highlighted is the research carried out by Sienkiewicz et al. (2010), which, after genetic validation of Trypanosoma brucei PTR1 as a potential target, developed a potent and selective inhibitor of TbPTR1 (IC<sub>50</sub> for TbPTR1 =7.0 nM; *Tb*DHFR  $\geq$  30.0 µM) that is unable to inhibit parasite growth. Thus, enzymatic assays only account for the compound's ability to interact with the target, whereas its permeability through the cell membranes, efflux process of the compounds, the retention capacity in lipophilic compartments, the high rate non-specific binding to proteins, among other factors remain largely ignored by such assays (Gilbert 2013). Although the in vitro assays were performed on pteridine reductase 1 from Leishmania major, in Brazil, the endemic species responsible for the clinical manifestations of CL, CML, and VL are, mainly, L. braziliensis and L. infantum, respectively. Nevertheless, the NADPH-dependent oxidoreductase enzymes present conservation of their supersecondary structure (Vidal et al. 2018) and highly similar active sites. In fact, a sequential identity analysis of PTR1 (Fig. 3) among the three species reveals 77.8% sequence identity between LmPTR1 and LbPTR1 and 88.9% sequence identity between LmPTR1 and LiPTR1 or LiPTR1 and LbPTR1. Despite overall similarity, the cofactor-binding site has two substitutions, His-241 for a Tyr-241 and a Tyr-194 for a Phe-194, that might impact the binding of ligands. In fact, Tyr-194 is considered an essential amino acid in the mechanism proposed by Gourley et al. (2001). However, previous results from our group (Leite et al. 2016) suggest that this residue does not interact with 2d, which only interact with residues that are conserved in LbPTR1 and LiPTR1. Thus, based on the sequential alignment data as well as the interaction profile described for the 2d, it is reasonable to assume that the L. major PTR1 inhibitors described in this work also bind to PTR1 from other Leishmania species. In order to support this hypothesis, compounds 2a, 2b, 2c, 2d, 2e, 4, and 9 were evaluated against the

LmPTR1 LiPTR1 LbPTR1	-MTAPTVPVALVTGAAK <mark>R</mark> LGRSIAEGLHAEGYAVCLHYHRSAAEANALSATLNARRPNSA -MAAPTVPVALVTGAAKRLGSGIAEGLHAEGYAVCLHYHRSAAEANTLAATLNARRPNSA MTSVATVPVALVTAAAKRLGCGIAEILHAQGYAVCLHYHRSAEDANTLTATLNSRRPNSA :. *************	59 59 60
LmPTR1 LiPTR1 LbPTR1	ITVQADLSNVATAPVSGADGSAPVTLFTRCAELVAACYTHWGRCDVLVNNASSFYPTPLL ITVQADLSNVAKAPAGGADGAAPVTLFKRCADLVAACYTHWGRCDVLVNNASSFYPTPLL IAVQADLSAVATASVSSAHDAAPISLVQRCAGLVDACYNHWGRCDVLVNNASAYYPTPLL *:****** **.***.:**::*. *** *** *	119 119 120
LmPTR1 LiPTR1 LbPTR1	RNDEDGHEPCVGDREAMETATADLFGSNAIAPYFLIKAFAHRFAGTPAKHRGTNYSIINM RKDEDGHVPCVGDREAMEAAAADLFGSNAMAPYFLIKAFAHRVADTPAEQRGTNYSIVNM NKVEEGHEPSMNEET-EAAAAADLFGTNALAPFFLIKAFAQRVADIPAEQRSDNYSIVNI .: *:** *: :::::::::::::::::::::::::	179 179 179
LmPTR1 LiPTR1 LbPTR1	VDAMTNQPLLGYTIYTMAKGALEGLTRSAALELAPLQIRVNGVGPGLSVLVIDMPPAVWE VDAMTSQPLLGYTIYTMAKGALEGLTRSAALELAPLQIRVNGVGPGLSVLAIDMPPAVRE IDAMTNQPLLGFTMYTMAKEALEGLTRSAALELAPLQVRVNGVSPGLSLFPADMPSAVQA :****	239 239 239
LmPTR1 LiPTR1 LbPTR1	GHRSKVPLYQRDSSAAEVSDVVIFLCSSKAKYITGTCVKVDGGYSLTRA 288 TYRSKVPLYQRDSSAAEVSDVVIFLCSSKAKYVTGTCVKVDGGYSLTRA 288 TYRSKVPLYQRESTAEEVAAAVFFLCSSSAKYITGTCVKVDGGYSLTRA 288	

**Fig. 3** Multiple sequential alignment of PTR1 of *L. major* (*Lm*PTR1), *L. infantum* (*Li*PTR1), and *L. braziliensis* (*Lb*PTR1). Multiple alignment was generated on the online Clustal Omega server. The rectangles delimit

the residues present at cofactor binding site. ("\*" = Identical, ":" = similar, "." = not conserved, "-" = absent)

strains of *L. braziliensis* and *L. infantum*. In addition, their cytotoxicity profile in human cells was also evaluated.

The evaluation of promising compounds against *Leishmania* sp. consists in determining the efficacy in inhibiting parasite growth by 50% ( $CE_{50}$ ). However, cytotoxicity assays in human cell lines are essential for evaluating the possible damages caused by the compounds on the human organism and to understand these compound selectivity profile.

Initially the seven compounds were analyzed for cytotoxicity against the WI-26VA4 cell line (ATCC CCL-75 lung fibroblast) in order to determine the dose capable of killing 50% of cells (LD<sub>50</sub>) (Table 2). The seven compounds evaluated had an LD<sub>50</sub> value higher than the drug used as reference (amphotericin B deoxycholate). Compounds 2b and 2c are the least cytotoxic in the series. The high standard deviation of error presented by compound 9 prevents a correct classification of its toxicity. Perhaps the structure of the compound enables it to emit fluorescence at the same wavelength as MTT, causing interference. The action of the compounds on human cells is unknown, since humans do not possess the enzyme PTR1. Although there are NADPH-dependent oxidoreductase enzymes in human cells, this does not guarantee the action of the compounds on them. Teles et al. (2018) have shown that analogous compounds, which bind to DHFR of Schistosoma mansoni (SmDHFR), have a competitive mechanism of inhibition towards the substrate. However, compounds 2a, 2b, and 2c were inactive against the SmDHFR. Thus, it is not possible infer these compounds' mechanism of action from those studies, since interference with normal physiology can be triggered by the impairment of basic cellular functions, such as energy metabolism, cytoskeleton organization, membrane integrity, glycogen storage, electrical conductivity, glutathione depletion, radical formation, or

 Table 2
 Evaluation of the biological activities and selectivity index for the compounds evaluated against Leishmania braziliensis, Leishmania infantum, and the WI-26VA4 cell line

Compounds	Cells line WI-26VA4 LD <sub>50</sub> (μM)	L. braziliensis EC <sub>50</sub> (µM)	L. infantum EC <sub>50</sub> (µM)	SI WI-26VA4/ L. braziliensis	SI WI-26VA4/ L. infantum
2a	$33.41 \pm 8.90$	44.16 ± 5.77	40.89 ± 5.65	0.76	0.82
2b	$43.29 \pm 10.05$	$65.69 \pm 13.85$	$23.45\pm4.54$	0.66	1.85
2c	>100	$49.22 \pm 7.71$	$66.09 \pm 14.17$	> 2.03	>1.51
2d	$26.67\pm5.27$	$48.23 \pm 13.05$	$35.90\pm 6.93$	0.55	0.74
2e	$31.24\pm7.13$	$47.60 \pm 12.51$	$51.24\pm3.39$	0.66	0.61
4	$29.22 \pm 6.37$	$46.44 \pm 9.58$	$30.36\pm5.11$	0.63	0.96
9	$61.29 \pm 26.85$	$70.98 \pm 11.25$	$68.77 \pm 12.62$	0.86	0.89
Amphotericin B deoxycholate	$3.89 \pm 1.64$	$4.23 \pm 0.44$	$3.12 \pm 0.74$	0.92	1.24



Fig. 4 Structure-activity relationship of the 7 compounds tested against *L. infantum* (EC<sub>50</sub>). Shaded atoms in blue promote improved potency, and those shaded in red promote potency loss

alteration of uptake or excretion (Schoonen et al. 2005). It should be noted that high standard deviations are characteristic for MTT phenotypic assays (Stockert et al. 2012).

Next, the effective concentration to inhibit 50% growth (EC<sub>50</sub>) of *Leishmania* (V.) *braziliensis* (MHOM/BR/75/M2903) and *Leishmania* (L.) *infantum* (MHOM/BR/74/PP75) was determined (Table 2).

For the L. *braziliensis* species, all compounds show similar EC50 values; however, compound 2c shows higher selectivity for *L. braziliensis* than the other compounds. In contrast, the biological activity of the compounds against the species of *L. infantum* exhibited a wider distribution of EC50 values, compound 2b was the most potent (EC50 =  $23.5 \pm 4.5$ ) and the most selective (SI 1.85). On the other hand, compound 2c is among the least potent ((EC50 =  $66.1 \pm 14.2$ ), despite its selectivity profile remains high. Thus, compound 2b is the best lead compound. No compound showed superior biological potency than amphotericin B deoxycholate, the control drug for both species.

Given that compound 2b presented the best biological activity (EC<sub>50</sub> =  $23.45 \pm 4.54 \mu$ M) and one of the best K*d* values (23.12  $\mu$ M), it seems that the presence of small lipophilic substituents in the *meta* and/or *para* position is essential for effective compounds against *Leishmania* species. Although these

compounds have lower biological activity than amphotericin B deoxycholate, they are less cytotoxic. Lipophilic substituents at the orto position affect the affinity of the compounds for the L. infantum species and, consequently, the biological activity. The relationship between the biological affinity of compounds 2a, 2b, 2d, 2e, and 4 against LmPTR1 and the biological potency of these compounds compared with the L. infantum species may be associated with greater conservation of residues at the binding site of LiPTR1 and LmPTR1, since the same behavior is assumed in the mechanism of action of these compounds with the prototype compound 2d. Hence, the conservation of the Tyr-194 residue, essential for the catalytic mechanism of PTR1, in the species LiPTR1 and LmPTR1, may be responsible for the improved activity profile of the compounds against L. infantum (Fig. 4). This line of thought does not hold for the L. braziliensis species, since there is no relationship between structure and activity. This difference, therefore, is expected, since L. major and L. infantum are closer, from the phylogenetic point of view, than L. braziliensis, a neotropical species. Considering the structure-activity relationship data, it can be seen that both Kd and  $EC_{50}$  values follow the same structural relationship profile.

Following the determination of the  $LD_{50}$  of the compounds against the human cell line and the  $EC_{50}$  against the species of

*Leishmania*, the selectivity index was calculated (Table 2) in order to assess to which organism the compounds are most harmful.

Only compounds 2b for *L. infantum* and 2c for both *L. braziliensis* and *L. infantum* had a selectivity index greater than 1 and were more effective against the parasite than cytotoxic to human cells. Although the literature adopts a value greater than 10 for the selectivity index, in order to consider that a compound is promising in relation to phenotypic assays (Di Giorgio et al. 2004; De Mesquita et al. 2009), this does not mean that compounds with the values below are insignificant. Thus, compound 2b, due to both enzymatic and cellular characteristics, is considered the most promising lead compound.

# Conclusions

The knowledge about compounds' biophysical characteristics is critical to set a suitable screening process. The information provided by LmPTR1 thermal shift assays suggest that compound 2c has low affinity to the target, whereas compounds 2a, 2b, 2d, and 4 have micromolar affinity to LmPTR1. As might be expected, these compounds are also active against other Leishmania species, as they share greater phylogenetic proximity. The most potent compound for the L. infantum is compound 2b (CE<sub>50</sub> =  $23.45 \pm 4.54 \mu$ M), while the most potent compound for the L. braziliensis species is compound 2a  $(CE_{50} = 44.16 \pm 5.77 \mu M)$ . These results highlight that kinetic characterization is the first step to identify a novel inhibitor; however, subsequent phenotypic assays make it possible to probe the compound behavior in complex media that allow pharmacokinetic parameters to be extrapolated for in vivo assays.

Results from phenotypic assays have been routinely employed, by research groups and the pharmaceutical industry alike, to decide which compound to move forward, or not, based on their pharmacokinetic and toxicological profile. Accordingly, it is found that only compounds 2b and 2c for L. infantum and 2c for L. braziliensis showed an antileishmania activity superior to their cytotoxic effect. Although, cell permeability assays are required to make a final decision on these compounds, some knowledge on their structure-activity relationships is crucial to unravel structural modifications that maintain the activity, but reduce toxicity to human cells, or increase their lipophilicity. Considering the molecular diversity of the compounds presented in this work (modification of the substituents of the -meta and -para, addition of other lateral lipophilic residues of different sizes), compound 2b shows the best selectivity index and the highest potency against L. infantum. This results are in good agreement with the Kd value for LmPTR1, which might be related to their high sequential identity within the active site. It should also be considered that a future analysis of the PTR1 structure for the *Leishmania* species circulating in Brazil is something to be explored, given the diversity of species in this country and their respective epidemiological impacts.

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