

Research Article

Cite this article: Santos CC, Zhang H, Batista MM, de Oliveira GM, Demarque KC, da Silva NL, Moreira OC, Ogungbe IV, Soeiro MdeNC (2022). Phenotypic investigation of 4-nitrophenylacetyl- and 4-nitro-1*H*-imidazolyl-based compounds as antileishmanial agents. *Parasitology* **149**, 490–495. <https://doi.org/10.1017/S0031182021002079>

Received: 25 February 2021
Revised: 22 September 2021
Accepted: 2 December 2021
First published online: 3 February 2022


Key words:

Cutaneous leishmaniasis; *in vitro* experimental chemotherapy; *in vivo*; *Leishmania amazonensis*; shape-based virtual screening

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Phenotypic investigation of 4-nitrophenylacetyl- and 4-nitro-1*H*-imidazolyl-based compounds as antileishmanial agents

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Abstract

Cutaneous leishmaniasis (CL) is a spectrum of clinical manifestations characterized by severe skin ulcerations that leads to social stigma. There are limited treatment options for CL, and the available drugs are becoming less efficacious due to drug resistance. More efficacious and safer antileishmanial drugs are needed. In this study, the biological effect of seven synthetically accessible nitroaromatic compounds was evaluated *in vitro* against amastigotes of *Leishmania amazonensis*, followed by *in vivo* evaluation using mouse models of CL. Two compounds (**6** and **7**) were active against amastigotes *in vitro* [half-maximal effective concentration (EC₅₀): 4.57 ± 0.08 and 9.19 ± 0.68 μM, respectively], with selectivity indexes >50, and the other compounds were not selective. *In vivo*, compounds **6** and **7** (10 mg kg⁻¹, twice a day for 14 days) failed to reduce skin lesion sizes and parasite loads determined by light microscopy of lesion *imprints* and quantitative polymerase chain reaction. Nevertheless, the *in vitro* leishmanicidal efficacy sustained their use as templates for nitroimidazole-based antileishmanial drug discovery programmes focusing on analogues with more suitable properties.

Introduction

Cutaneous leishmaniasis (CL) comprises a broad spectrum of clinical manifestations caused by over 20 different kinetoplastid *Leishmania* species transmitted by female sandflies (WHO, 2020). In Asia and Africa, most CL cases are caused by *Leishmania major* and *Leishmania tropica*, leading to self-limiting ulcers. In the Americas, the disease is caused by several species, including *Leishmania braziliensis*, *Leishmania mexicana*, *Leishmania guyanensis*, *Leishmania naiffi* and *Leishmania amazonensis*. The clinical presentations range from self-healing localized skin lesions to disfiguring mucocutaneous ulcerations (Martins *et al.*, 2014; de Vries *et al.*, 2015). Annually, about 1.2 million new CL cases are reported, and a significant number of patients develop severely disfiguring permanent scars, which results in social stigma and loss of economic productivity (Alvar *et al.*, 2012; Bailey *et al.*, 2019). CL is one of the most mistreated illnesses among neglected tropical diseases (NTDs). The large variation of drug susceptibility among the diverse parasite species makes the drug discovery process even more difficult (de Vries *et al.*, 2015; Van Bocxlaer *et al.*, 2019).

The first-line treatment for CL includes pentavalent antimonials: sodium stibogluconate and meglumine antimoniate. Antimonials present several drawbacks in terms of efficacy, safety (cardiotoxicity and hepatotoxicity) and protracted periods of drug administration (Weisz, 2006). Alternative systemic agents include miltefosine, amphotericin B, pentamidine isethionate, paromomycin and granulocyte-macrophage colony-stimulating factor, limited by severe adverse effects (de Vries *et al.*, 2015). There is a need for safer, more selective and easily accessible therapies for CL.

A few new small molecules are currently in preclinical and clinical development against *Leishmania*. One example is the first-in-class kinetoplastid-selective proteasome inhibitor compound LXE408. LXE408 was discovered by Novartis with financial support from the Wellcome Trust. LXE408 is efficacious in the murine model of CL, and it is under clinical trial for the treatment of visceral leishmaniasis (Nagle *et al.*, 2020). Partnerships between not-for-profit research and private pharmaceutical companies for the research and development of new treatments for NTDs such as leishmaniasis are increasing (Katsuno *et al.*, 2015).

Some of the compounds in development include nitroaromatic compounds, which are historically used to treat several diseases caused by protozoans, for example, DNDI-8219, DNI-VL2098 and DNDI-0690 (Van Bocxlaer *et al.*, 2019). In this study, we envision that scaffold-driven and easily synthesizable nitroaromatic compounds based on prior art can facilitate rapid phenotypic screenings using *in vitro* and *in vivo* experimental models. The scaffolds were selected based on our ongoing research on the use of natural product-derived motifs in the design and development of antiprotozoal agents (Zhang *et al.*, 2018). The compounds were evaluated against the free amastigote forms (AFs) of *L. amazonensis* (strain LTB0016),

purified from paw lesion, followed by evaluation against the intracellular forms as well as on peritoneal macrophages (PMMs). The most selective were evaluated in *L. amazonensis*-infected BALB/c mice. The methods and results are presented below.

Materials and methods

Compounds: Dimethyl sulphoxide (DMSO, Merck) solutions (30 mM) of the compounds (Fig. 1) and miltefosine $\geq 98\%$ (Sigma-Aldrich, St. Louis, USA) were diluted just before the *in vitro* experiments into RPMI-1640 medium (pH: 7.2–7.4) without phenol red (Sigma-Aldrich (St. Louis, USA), R7509) but supplemented with 1% L-alanyl-L-glutamine (GLUTAMAX I, Gibco™), 1% penicillin–streptomycin 5000 U mL⁻¹ (Gibco™, PEN-STR) and 10% heat-inactivated sterile-filtered fetal bovine serum (FBS), from Cultilab, Brazil. The concentration of DMSO was <0.6% for all *in vitro* experiments to prevent non-specific toxicity to host cells (Romanha *et al.*, 2010). For *in vivo* assays, compounds 6 and 7 were prepared daily in a solution composed of 5% Gum Arabic from acacia tree (Sigma-Aldrich, St. Louis, USA), 6% DMSO and 3% Tween™ 80 Surfact-Amps™ detergent solution (Thermo Scientific™). The reference drug, Milteforan™ (Mt™), was dissolved in sterile deionized water.

Parasite strain and mammalian host cell cultures

Leishmania amazonensis (strain LTB0016) was used throughout the study. AFs were purified from male BALB/c mice. Briefly, the foot paws were inoculated (subcutaneously) with 20 μ L containing 10⁶ amastigotes, and the animal skin lesions were aseptically removed 30 days post-infection (dpi). The parasites in the lesions were mechanically dispersed by pipetting and used for the assays. The compounds were tested on (1) extracellular AFs; (2) intracellular amastigote forms (IA) in PMMs and (3) BALB/c mouse models infected with amastigotes as recently described (Cardoso Santos *et al.*, 2021). PMMs were obtained from Swiss male mice (18–20 g) inoculated with 3% Brewer thioglycollate medium (Merck) previously diluted in water and autoclaved. After 4 days of stimulation, the cells were collected by rinsing the animals' peritoneum with RPMI 1640. The cells were subsequently seeded in 24- (3 \times 10⁵ cells well⁻¹) and 96-well (5 \times 10⁴ cells well⁻¹) plates for analysis of infection and cytotoxicity assays, respectively. Cell cultures were maintained at 37°C, 5% CO₂ atmosphere in RPMI 1640 medium (pH: 7.2–7.4) without phenol red, but supplemented with 1% L-glutamine, 1% (PEN-STR) and 10% FBS. Assays using AF were maintained at 32°C using RPMI culture medium containing 5% FBS.

Cytotoxicity and in vitro leishmanicidal analysis

The toxicity of the compounds on host cells was evaluated 24 h after PMMs were seeded into culture plates. The compounds (0–500 μ M) were added to the wells and incubated as described above for 48 h, and cellular viability was evaluated using the AlamarBlue test (Invitrogen) according to the manufacturer's recommendations. The effect of the compounds (0–20 μ M) on AFs was evaluated using 10⁶ parasites per well (0.2 mL) for 48 h, and parasite viability assessed with AlamarBlue tests as reported (Mikus and Steverding, 2000; Santos *et al.*, 2020). Phenotypic screenings against IAs were performed by infecting 3 \times 10⁵ PMMs with 9 \times 10⁵ amastigotes, multiplicity of infection being 3. After 48 h of drug treatment, cultures were rinsed with saline, fixed 5 min with Bouin and stained for 15 min with Giemsa solution (Sigma-Aldrich (St. Louis, USA), 32884), filtered and diluted five times in distilled water (Santos *et al.*, 2018; Feitosa *et al.*, 2019). Samples were subsequently evaluated by

light microscopy and photographs were obtained by using a Zeiss AxioObserver MI microscope (Oberkochen, Germany). The percentage of infected host cells, the number of parasites per infected cell and the corresponding infection index were obtained. Parasites with well-defined nuclei and kinetoplasts were counted as surviving since irregular structures could mean parasites undergoing death. The results were expressed as % reduction in parasite burden, and the half-maximal effective concentration (EC₅₀) and 90% maximal effective concentration (EC₉₀) were calculated by non-linear regression analysis using GraphPad Prism v.9.1.2. All assays were run at least twice with three independent replicates each time.

Hep G2 viability assay

Hep G2 (CRL-11997™) cells were grown in complete growth medium (Dulbecco's modified Eagle medium: F12 containing L-glutamine and sodium bicarbonate, 10% FBS and 1% penicillin/streptomycin) incubated at 37°C in a 5% CO₂ environment. Cells were seeded into 96-well plates at 5 \times 10⁵ cells mL⁻¹ and incubated for about 24 h. Cells were treated with the compounds prepared in DMSO for 72 h at a final concentration range of 160–1.2 μ M in triplicates. Cell viability was assessed using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-2H-tetrazolium bromide assay as previously described (Zhang *et al.*, 2018). Podophyllotoxin was used as cytotoxicity control, and sodium dodecyl sulphate (10%) was used as assay control.

In vivo efficacy analysis

Male BALB/c mice (18–20 g) obtained from the animal facilities of ICTB (Institute of Science and Biomodels Technology, Fiocruz, RJ, Brazil) were housed five per cage, kept in a room at 20–24°C under a 12-light and 12-h dark cycle and provided with sterile water and chow ad libitum. Animals were subcutaneously inoculated with 5 \times 10⁵ amastigotes on the left foot paw (Godinho *et al.*, 2012). The treatment started 15 dpi, corresponding to the lesion onset (size diameter of 3–4 mm) (Van Bocxlaer *et al.*, 2019). The compounds were given for 14 days by oral gavage (100 μ L) at 10 mg kg⁻¹, twice a day (q12h). Mt™ was administered at a standardized dose (40 mg kg⁻¹) once a day (q24h). The lesions were measured regularly in three dimensions (height, width and depth), and animals were euthanized at 31 dpi. Skin lesions aseptically removed were used for molecular analysis of parasite load by quantitative polymerase chain reaction (qPCR) and light microscopy inspections of *imprints* (Ribeiro-Romão *et al.*, 2016). All procedures were carried out in accordance with the guidelines established by the FIOCRUZ Committee of Ethics for the Use of Animals (CEUA L038/2017). Statistical analysis was conducted in GraphPad Prism v.9.1.2 by analysis of variance (ANOVA): (1) one-way Fisher's LSD test or (2) two-way unpaired *t*-test with Welch's correction, significance $P < 0.05$ [95% confidence interval (CI)].

Results

All experimental details are fully described in the Supplementary material. The compounds (Fig. 1) were synthesized as shown in Fig. S1 and diluted just before the *in vitro* experiments. First, the compounds were screened at 10 μ M against extracellular amastigotes of *L. amazonensis* purified from male BALB/c paw lesions (Feitosa *et al.*, 2019) using colorimetric assays (AlamarBlue test, Invitrogen) (Mikus and Steverding, 2000; Godinho *et al.*, 2012). Two (6 and 7) out of the seven compounds caused a significant decline in the total number of live parasites ($\geq 50\%$ decrease compared to untreated parasites) with only 48 h of drug exposition.

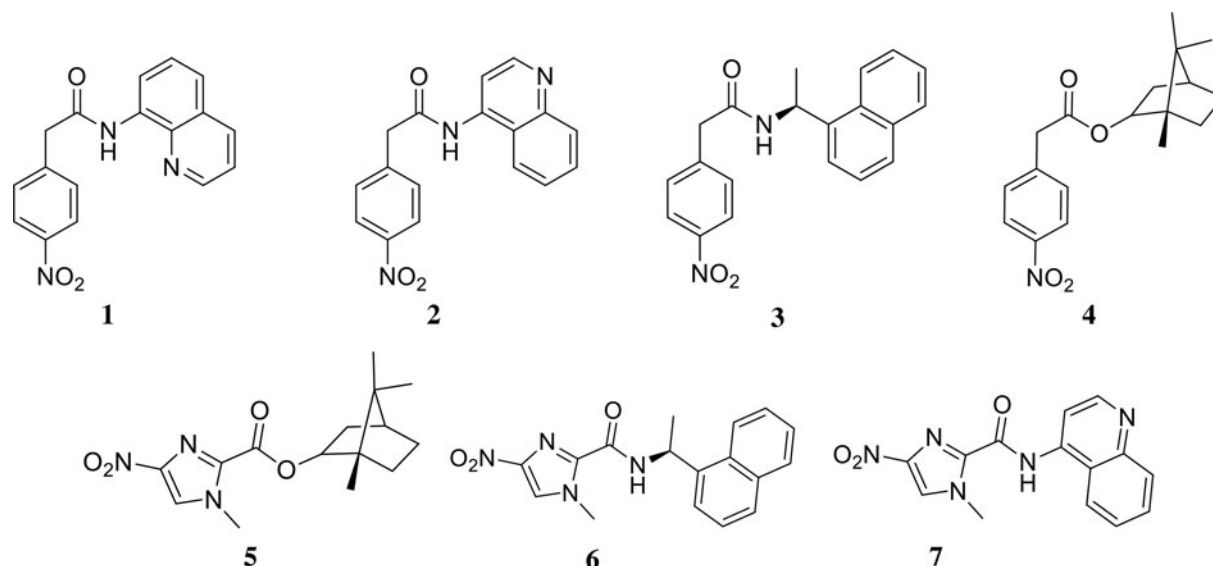


Fig. 1. Structures of the tested compounds.

Table 1. Phenotypic studies of compounds 6 and 7 against *Leishmania amazonensis* and their toxicity profile

Compound	EC ₅₀ (μM) _{AF}	EC ₅₀ (μM) _{IA}	CC ₅₀ (μM) _{PMM}	SI _{AF}	SI _{IA}	CC ₅₀ (μM) Hep G2
6	10.78 ± 0.80	4.57 ± 0.08 (<i>P</i> = 0.027) ^a	>500	>46	>109	>160
7	13.12 ± 1.70	9.19 ± 0.68	>500	>38	>54	>160
Miltefosine	4.37 ± 0.11	13.30 ± 3.03 ^a	169.69 ± 5.07	>38	>12	–

AF, amastigote forms of *L. amazonensis* (LTB0016 strain) purified from animal lesions; IA, intracellular forms in peritoneal macrophages; SI, selective index. EC₅₀ is reported as mean and s.d. values.

^aStatistical analysis was performed with GraphPad prism v.9.1.2 by ordinary ANOVA test.

Next, 6 and 7 were further analysed in dose–response assays. The EC₅₀ values obtained were 10.78 ± 0.80 and 13.12 ± 1.70 μM, respectively (Table 1). Host cell viability showed that compounds did not exhibit any toxicity up to the highest tested concentration, leading to CC₅₀ PMM values >500 μM. Also, when cellular viability was assessed using Hep G2 cultures, no toxicity was observed (CC₅₀ Hep values >160 μM, Table 1). The selective antiparasitic activity was encouraging, and we decided to evaluate the compounds on intracellular amastigotes (IA) in PMMs (9 × 10⁵ IA: 3 × 10⁵ PMMs well⁻¹), which mimics typical infection in mammalian cells. Light microscopy analysis showed that 6 was about 3-fold more active compared to the reference drug (miltefosine, *P* = 0.0273). Its EC₅₀ was 4.57 μM, and it is quite selective [selective index (SI) > 100]. Compound 6 also displayed low EC₉₀ (9.14 μM) and caused a 92% decrease in PMM infection at 10 μM. Compound 7 is slightly less potent (EC₅₀ = 9.19 μM and EC₉₀ = 17.2 μM) with an SI of >50 (Table 1). Both compounds displayed an impressive leishmanicidal effect due to a drastic decrease in the number of parasites per cell and the ratio of infected cells at 20 μM (Fig. S2). These favourable results prompted us to conduct preliminary *in vivo* proof of concepts using both hit compounds.

In the *in vivo* experiment, we observed a gradual increase in skin lesion size of mice treated with the vehicle alone, attaining 438.8 ± 42.05 mm³ at the endpoint (Fig. 2). Mice treated with compound 6 displayed similar lesions size (443.7 ± 47.76 mm³, *P* = 0.4798), whereas a slight increase was observed using 7 (550.9 ± 30.23 mm³, *P* = 0.2398), as compared to the vehicle-treated group. MtTM achieved a 72% decrease in the size of skin lesions and significantly improved the clinical condition (*P* < 0.0001 (Fig. 2A and B). The qPCR standardization curves achieved all the desired efficiency and linearity coefficient criteria,

ranging between 93.2–91.4% and 0.98–0.99% for the parasite kDNA amplification (Fig. S3A) and mice GAPDH target (Fig. S3B), respectively. At 31 dpi, qPCR readout expressed as DNA equivalents per mg of tissue (14) showed a 5–10-fold increase in the parasite load of mice treated with 6 (276 526 ± 40 519 eq. par mg⁻¹ tissue, *P* = 0.0208) and 7 (153 765 ± 54 932 eq. par mg⁻¹ tissue, *P* = 0.0829) as compared to the vehicle-treated group (27 753 ± 9539 eq. par mg⁻¹ tissue) (Fig. 2B). MtTM completely suppressed (99.99%) parasite load (0.256 ± 0.1022 eq. par mg⁻¹ tissue, *P* = 0.0437). The qPCR showed that the hit compounds lacked *in vivo* efficacy, and it was corroborated by a qualitative light microscopy analysis of lesions imprints (Fig. 2C). The findings clearly showed that both 6- and 7-treated animal groups have high parasitaemia coupled with the presence of lymphomononuclear diffuse infiltrates and several parasite niches inside mononuclear host cells (Fig. 2).

The lack of *in vivo* efficacy suggests low bioavailability of the compounds in mice. Low bioavailability is generally caused by impaired absorption or high clearance of compounds. Therefore, an *in silico* tool, pkCSM, was used to predict some critical absorption, distribution and metabolism parameters of 6 and 7, comparing to miltefosine (Gleeson, 2008; Pires et al., 2015). Compounds 6 and 7 do not violate Lipinski's rule. The cLog *P*s of 6 and 7 are 2.97 and 2.13, respectively, while that of miltefosine is 5.68 (Table 2). Since miltefosine is zwitterionic, its relatively high cLog *P* is ideal for high intestinal absorption. Similarly, since compounds 6 and 7 have low molecular weights (324.34 and 297.27 g mol⁻¹, respectively) and are neutral molecules at physiological pH with relatively low cLog *P*, sufficient intestinal adsorption was expected from oral gavage. The predicted high intestinal absorption further supports this in addition to the

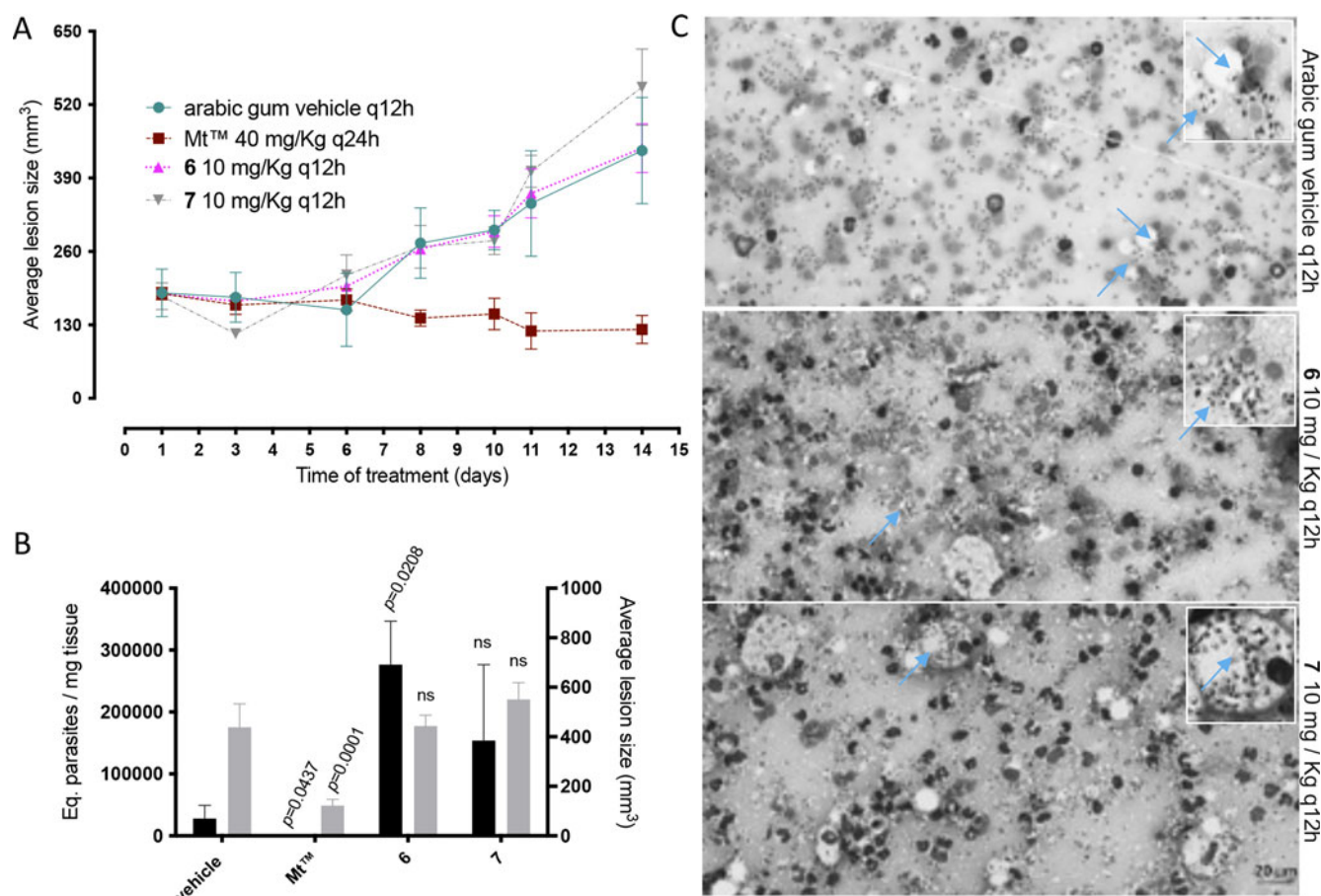


Fig. 2. Effect of **6** and **7** in CL experimental mouse models using BALB/c mice infected with *Leishmania amazonensis*. The graphics show: average lesion size during treatment (A), the correlation between parasite load by qPCR (black bars) and average lesion size measurements (grey bars) at 31 dpi (B), according to each experimental group. Light microscopy of lesion imprints of infected mice treated with vehicle and after administration of compounds **6** and **7** at 10 mg kg⁻¹ q12h po (C). Arrow: intracellular parasites. Statistical analysis was performed with GraphPad prism v.9.1.2 by ANOVA test (95% CI). When *P* value ≥ 0.05 = not significant (ns).

Table 2. Predicted ADME properties of **6**, **7** and miltefosine

Compound	cLog <i>P</i>	Caco2-permeability	Intestinal absorption ^a	VD _{ss} ^a	Fraction unbound ^a	CYP3A4 substrate
6	2.97	0.83	90.23	1.28	0.05	Yes
7	2.13	1.043	90.03	0.81	0.21	Yes
Miltefosine	5.68	1.049	92.02	0.36	0.16	Yes

^aThe predicted values are for humans.

predicted moderate to high Caco-2 permeability (0.83 and 1.04, respectively) from *in silico* Absorption, Distribution, Metabolism, and Excretion (ADME) calculations. Oral gavage was selected for this study because oral administration is the preferred route for the clinical use of new antileishmanial drugs. The predicted volume of distribution (VD_{ss}) is high and ideal for tissue distribution for both compounds. However, the predicted unbound fraction for **6** is relatively low (5%) and ideal for **7** (21%). Both compounds are predicted as CYP3A4 substrates, which suggest significant first-pass hepatic clearance and low overall plasma and tissue concentration. Therefore, a comparative pharmacokinetics study (oral and intravenous) can provide insight into plasma concentrations and bioavailability of both compounds in addition to determining the appropriate dosage and structural changes necessary for *in vivo* efficacy. Also, *in silico* toxicological analysis of both compounds suggests probable mutagenicity (positive for Ames tests) and potential to inhibit hERG II and that **6** is potentially hepatotoxic. However, animals treated

with compounds **6** and **7** did not show any observable side-effects. A cell proliferation assay using Hep G2 cells indicates that the compounds are not hepatotoxic at the tested concentrations (160–1.2 μM).

Discussion

Leishmania amazonensis is one of the most common *Leishmania* species in Brazil, known to cause a wide spectrum of pathologies, including highly severe and diffuse CL (Lainson *et al.*, 1994). Most CL patients live in impoverished communities with limited access to primary healthcare facilities (Okwor and Uzonna, 2016). The lack of access to primary healthcare typically means that the disease is usually at an advanced stage before medical intervention is sought, if available (Ruoti *et al.*, 2013). In this study, we explored the *in vitro* and *in vivo* leishmanicidal effects of compounds derived from two nitroaromatics scaffolds (4-nitrophenylacetyl and 4-nitro-1*H*-imidazolyl). The *in vitro* data demonstrated that

two 4-nitro-1*H*-imidazolyl compounds have acceptable selectivity indices (>50 for intracellular forms of *L. amazonensis*) as previously discussed by Caridha *et al.* (2019) as well as by Alcântara *et al.* (2018). Therefore, their antileishmanial activities were investigated further. The compounds were administered to BALB/c mouse models of CL for 14 days at a relatively low dose, starting at the onset of lesions (Godinho *et al.*, 2012). The results showed that the *in vitro* potency of the hit compounds 6 and 7 did not translate to desired *in vivo* efficacy. The measurements of the mouse lesions (using a paquimeter) and the molecular analysis of animal parasitism (qPCR) demonstrated a lack of *in vivo* activity of both compounds, whereas MtTM suppressed both parameters, as reported (Van Bocxlaer *et al.*, 2019). The lack of *in vivo* efficacy of the tested compounds could be, at least in part, due to low bioavailability and/or high metabolic clearance. Nevertheless, as nitro-drugs are activated by *Leishmania* nitroreductases (NTR2) (Wyllie *et al.*, 2016), it is possible that 6 and 7 may be also activated in a similar way, and thus, the *in vitro* leishmanicidal efficacy supports their use as templates for nitroimidazole-based antileishmanial drug discovery programmes focusing on analogues that have the appropriate Target Product Profile (TPP) for new CL drugs, as recommended (Drugs for Neglected Diseases Initiative, 2020).

Supplementary material. The supplementary material for this article can be found at <https://doi.org/10.1017/S0031182021002079>

Acknowledgements. The authors acknowledge the Fortalecimento dos Programas de Gestão Estratégica de Pesquisa da Fiocruz Rede de Plataformas Fiocruz (VPPLR-001-Fio 14) and Programa de Excelência Acadêmica (PROEX) from CAPES. They appreciate the excellent technical contribution of Roberson Donola Girão, Dr Ana Lia Mazzetti and Dr Cristiane França da Silva (Laboratory of Cellular Biology/IOC/Fiocruz) and Kingsley Bimpeh (Jackson State University).

Financial support. The current study was supported by grants from Fundação Carlos Chagas Filho de Amparo à Pesquisa do Estado do Rio de Janeiro (FAPERJ), Conselho Nacional Desenvolvimento científico e Tecnológico (CNPq), Coordenação de Aperfeiçoamento de Pessoal de Nível Superior (CAPES) and Fundação Oswaldo Cruz PAEF/CNPq/Fiocruz. MNCS and OCM are research fellows of CNPq and CNE and JCNE researchers. The work at Jackson State University was supported, in part, by the US National Institutes of Health (SC3GM122629 and G12MD007581).

Conflict of interest. None.

Ethical standards. All procedures were carried out in accordance with the guidelines established by the FIOCRUZ Committee of Ethics for the Use of Animals (CEUA L038/2017).

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