Soulamarin Isolated from *Calophyllum brasiliense* (Clusiaceae) Induces Plasma Membrane Permeabilization of *Trypanosoma cruzi* and Mytochondrial Dysfunction

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**Abstract**

Chagas disease is caused by the parasitic protozoan *Trypanosoma cruzi*. It has high mortality as well as morbidity rates and usually affects the poorer sections of the population. The development of new, less harmful and more effective drugs is a promising research target, since current standard treatments are highly toxic and administered for long periods. Fractioning of methanol (MeOH) extract of the stem bark of *Calophyllum brasiliense* (Clusiaceae) resulted in the isolation of the coumarin soulamarin, which was characterized by one- and two-dimensional ¹H- and ¹³C NMR spectroscopy as well as ESI mass spectrometry. All data obtained were consistent with a structure of 6-hydroxy-4-propyl-5-(3-hydroxy-2-methyl-1-oxobutyl)-6,6'-dimethylpyrane-[2',3',8,7]-benzopyran-2-one for soulamarin. Colorimetric MTT assays showed that soulamarin induces trypanocidal effects, and is also active against trypomastigotes. Hemolytic activity tests showed that soulamarin is unable to induce any observable damage to erythrocytes (cₘₐₓ = 1,300 μM). The lethal action of soulamarin against *T. cruzi* was investigated by using amino[4-(6-(amino(iminio)methyl)-1H-indol-2-yl)phenyl)methaniminium chloride (SYTOX Green and 1H,5H,11H,15H-Xantheno[2,3,4-ij:5,6,7-i']diquinoliniz-18-iium, 9-[4-(chloromethyl)phenyl]-2,3,6,7,12,13,16,17-octahydrochloride (MitoTracker Red) as fluorimetric probes. With the former, soulamarin showed dose-dependent permeability of the plasma membrane, relative to fully permeable Triton X-100-treated parasites. Spectrofluorimetric and fluorescence microscopy with the latter revealed that soulamarin also induced a strong depolarization (ca. 97%) of the mitochondrial membrane potential. These data demonstrate that the lethal action of soulamarin towards *T. cruzi* involves damages to the plasma membrane of the parasite and mitochondrial dysfunction without the additional generation of reactive oxygen species, which may have also contributed to the death of the parasites. Considering the unique mitochondrion of *T. cruzi*, secondary metabolites of plants affecting the bioenergetic system as soulamarin may contribute as scaffolds for the design of novel and selective drug candidates for neglected diseases, mainly Chagas disease.

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Introduction

The tree *Calophyllum brasiliense* is known in Brazil as “Guanandi” or “Jacareúba”. It can reach up to 40 meters in high, 1–3 meters in diameter and is usually found in Brazil in the rain forest regions of the Amazon. Its stem bark is used in folk medicine to treat rheumatism, varicose veins, haemorrhoids and ulcers, whereas the leaves have anti-inflammatory properties [1]. Previous chemical studies on *C. brasiliense* resulted in the isolation of several interesting secondary metabolites of plants affecting the bioenergetic system as soulamarin may contribute as scaffolds for the design of novel and selective drug candidates for neglected diseases, mainly Chagas disease.

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Author Summary

Chagas disease is a parasitic protozoan that affects the poorest population in the world, causing a high mortality and morbidity. As a result of highly toxic and long-term treatments, the discovery of novel, safe and more efficacious drugs is essential. Natural products isolated from plants are commonly used as drug prototypes or precursors to treat parasitic diseases. As part of our investigation of bioactive compounds from Brazilian flora, the present study was undertaken in order to determine the antitrypanosomal effects of the soulamarin, a coumarin isolated from the stem bark of Callophyllum brasiliense (Clusiaceae), against Trypanosoma cruzi. This study moreover investigated the lethal action of soulamarin towards the parasite. Considering the obtained results, secondary metabolites of plants affecting the bioenergetic system as soulamarin may contribute as scaffolds for the design of novel and selective drug candidates for neglected diseases, mainly Chagas disease.

General experimental procedures

NMR spectra were recorded on a Bruker DRX-500 (1H: 500 MHz, 13C:125 MHz) spectrometer at ambient temperature. Chemical shifts (δ) are reported in ppm and coupling constants (J) in Hz. All resonances were referenced to residual NMR solvent resonance. Low-resolution electrospray ionization mass spectra (LR-ESI-MS) were measured in positive mode on a Platform II-Micromass (quadrupole) mass spectrometer.

Materials and Methods

Chemical reagents and drugs

The compounds 3-[4,3-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT; Thiazol blue), mesoxalondinitrile 4-trifluoro-omethoxyphenylhydrazide (FCCP), 4’,6-diamidino-2-phenylindole dihydrochloride (DAPI), sodium dodecyl sulfate (SDS), M-199 and RPMI-1640 medium (without phenol red) as well as the NMR solvents CDCl3 and CD3OD were purchased from Sigma-Aldrich (USA). Dimethylsulfoxide (DMSO) was purchased from Merck (Brazil). 1H,5H,11H,15H-Xantheno[2,3,4-ij:5,6,7-tr']diquinoliniz-18-i um, 9-[4-(chloromethylphenyl]-2,3,6,7,12,13,16,17-octahydro-chloride (MitoTracker Red CM-H2XROS), amino(-6-aminooiminio)methyl]-1H-indol-2-ylphenylmethanaminium chloride (SYTOX Green) and 2’,7’-dichlorodihydrofluorescein diacetate (H2-DCF-DA) were purchased from Molecular Probes (Invitrogen, Carlsbad, CA, USA). Silica gel (230–400 mesh) and Sephadex LH-20, used for column chromatography and analytical TLC (60 PF254), were either purchased from Merck (USA) or Sigma-Aldrich (USA). Benznidazole (2-nitroimidazole) was obtained from the Laboratorio Farmaceutico do Estado de Pernambuco – LAFEPE (Recife, Brazil).

Plant material

Samples of the stem bark of C. brasiliense were collected in the Amazonian rain forest of Brazil during September 2011. The authenticity of the plant material was verified by Dr. Eliana Rodrigues from ICAQF-UNIFESP. Sample specimens were deposited at the herbarium of the Instituto de Botânica - SEMA of São Paulo (SP, Brazil).

Extraction and isolation of 6-Hydroxy-4-propyl-5-(3-hydroxy-2-methyl-1-oxobutyl)-6’,6”-dimethylpyrane-[2’,3’;8,7]-benzopyran-2-one (soulamarin)

Dried and powdered stem bark samples of C. brasiliense (72 g) were washed exhaustively with hexane (10×500 mL) at room temperature in order to remove any residual fats. Subsequently, the plant material was extracted with MeOH (10×1 L) at room temperature. The combined organic fractions afforded, after removal of all solvents under reduced pressure, 4.7 g of crude residue. This crude extract was dissolved in MeOH:H2O (1:2) and extracted with EtOAc. The removal of the solvent under reduced pressure resulted in the deposition of a residue (3.0 g), which was subsequently subjected to column chromatography (Sephadex LH-20) with MeOH as the eluent. Nine fractions (I–IX) were separated like this. Fraction III (1.31 g) was further purified by column chromatography over silica gel with a crude solvent mixture of EtOAc:H2O (1:1), giving 544 mg of soulamarin (see Figure 1).

General experimental procedures

NMR spectra were recorded on a Bruker DRX-500 (1H: 500 MHz, 13C:125 MHz) spectrometer at ambient temperature. Chemical shifts (δ) are reported in ppm and coupling constants (J) in Hz. All resonances were referenced to residual NMR solvent resonance. Low-resolution electrospray ionization mass spectra (LR-ESI-MS) were measured in positive mode on a Platform II-Micromass (quadrupole) mass spectrometer.

Materials and Methods

Chemical reagents and drugs

The compounds 3-[4,3-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT; Thiazol blue), mesoxalondinitrile 4-trifluoro-omethoxyphenylhydrazide (FCCP), 4’,6-diamidino-2-phenylindole dihydrochloride (DAPI), sodium dodecyl sulfate (SDS), M-199 and RPMI-1640 medium (without phenol red) as well as the NMR solvents CDCl3 and CD3OD were purchased from Sigma-Aldrich (USA). Dimethylsulfoxide (DMSO) was purchased from Merck (Brazil). 1H,5H,11H,15H-Xantheno[2,3,4-ij:5,6,7-tr’]diquinoliniz-18-i um, 9-[4-(chloromethylphenyl]-2,3,6,7,12,13,16,17-octahydro-chloride (MitoTracker Red CM-H2XROS), amino(-6-aminooiminio)methyl]-1H-indol-2-ylphenylmethanaminium chloride (SYTOX Green) and 2’,7’-dichlorodihydrofluorescein diacetate (H2-DCF-DA) were purchased from Molecular Probes (Invitrogen, Carlsbad, CA, USA). Silica gel (230–400 mesh) and Sephadex LH-20, used for column chromatography and analytical TLC (60 PF254), were either purchased from Merck (USA) or Sigma-Aldrich (USA). Benznidazole (2-nitroimidazole) was obtained from the Laboratorio Farmaceutico do Estado de Pernambuco – LAFEPE (Recife, Brazil).
mice and deposited on a 16-well chamber slide (16 co-workers [23] was used with minor modifications. Peritoneal intracellular amastigotes, the method described by De Souza and (10% v/v; 100 μM carried out for 18 hours at 24°C with sodium dodecylsulfate (SDS) [22]. A suspension (5%) of Trypanosoma cruzi trypomastigotes was incubated with soulamarin at 25°C for 1 hour in a U-shaped microplate (96 wells). The absorption of the supernatant at 550 nm was recorded (FilterMax F5 Multi-Mode Microplate Reader-Molecular Devices).

Cytotoxicity against mammalian cells

Peritoneal macrophages were collected from female BALB/C mice, seeded at 1×10^6 cells/well in 96-well microplates and incubated with soulamarin for 72 h at 37°C in an incubator with 5% CO2. The viability of the cells was determined using MTT [16]. The data represent the mean of two independent assays (triplicates).

Hemolytic activity

The hemolytic activity of soulamarin in concentrations up to 1,300 μM was evaluated from the erythrocytes of BALB/c mice [24]. A suspension (5%) of erythrocytes in PBS (phosphate buffered saline) was incubated with soulamarin at 37°C for 1 hour in a U-shaped microplate (96 wells). The absorption of the supernatant at 550 nm was recorded (FilterMax F5 Multi-Mode Microplate Reader-Molecular Devices).

Spectrofluorimetric detection of the permeability of the cell membrane

Trypomastigotes were washed with PBS (phosphate buffered saline), deposited on a microplate (1×10^6 cells/well) and incubated with SYTOX Green (1 μM) for 15 minutes at 24°C [25]. Soulamarin was added in three concentrations (IC_{50} = 386 μM, IC_{25} = 219 μM and IC_{25} = 103 μM) and the fluorescence was measured after 20, 40 and 60 minutes. The maximum permeability was observed with 0.1% Triton X-100 (positive control). The fluorescence intensity was determined using a spectrofluorimeter. Untreated trypomastigotes and DMSO-treated parasites were used as negative controls.

Effect of soulamarin on the mitochondrial membrane potential

Trypomastigotes were washed with PBS, deposited on a microplate (2×10^6 cells/well) and incubated with soulamarin (IC_{50} = 219 μM) for 60 minutes at 37°C. Then MitoTracker Red CM-H2XROS (500 nM) was added and the incubation was continued for 40 minutes in the dark. Cells were washed twice with BSS (Hank’s buffered salt solution) and the fluorescence was measured using a spectrofluorimeter. Untreated trypomastigotes and DMSO-treated parasites were used as negative controls. Mesoxalonitrile 4-trifluoromethoxyphenylnitrazene (FCCP; 10 μM) was used as a positive control [27]. For the fluorescence microscopy analysis, trypomastigotes were co-stained with 4’,6-diamidino-2-phenylindole (DAPI; 10 μM) and examined at 1000× magnification. Merged images of blue (DAPI) and red (MitoTracker Red) images were obtained using the Nikon NIS - Elements AR software. A Nikon D-FL Epimicroscope equipped with a DS-U3 digital camera was used for the experiment.

Parasites and mammalian cells

In all in vitro assays, Y strains of T. cruzi trypomastigotes were used, which were kept at 37°C in LLC-MK2 (ATCC CCL 7) cells using RPMI-1640 medium with calf serum (2%) [20]. To keep the Y strains infective, trypomastigotes were also kept in swiss mice and regularly harvested from the bloodstream by heart puncture of infected animals at the peak of the parasitemia [21]. LLC-MK2 cells were maintained at 37°C in RPMI-1640 medium with fetal calf serum (10%) in an incubator (5% CO2 atmosphere).

Antitrypanosomal activity

Trypomastigotes were counted in a hemocytometer (Neubauer) and deposited on a microplate (96 wells; 1×10^6 cells/well). Subsequently, soulamarin was added to the cells in concentrations up to 386 μM and the cells were allowed to incubate for 24 hours at 37°C (5% CO2 atmosphere). Benzimidazole was used as standard. Trypomastigote activity was based on the conversion of the soluble tetrazolium salt 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT) into the insoluble formazan by mitochondrial enzymes. The extraction of formazan was carried out for 18 hours at 24°C with sodium dodecylsulfate (10% v/v; 100 μL/well) [22].

In order to determine the IC_{50} value for soulamarin against intracellular amastigotes, the method described by De Souza and co-workers [23] was used with minor modifications. Peritoneal macrophages were collected from the peritoneal cavity of BALB/c mice and deposited on a 16-well chamber slide (1×10^7 cells/well) before being incubated for 24 hours at 37°C (5% CO2 atmosphere). Trypomastigotes from LLC-MK2-infected cultures were washed twice in RPMI-1640 medium, counted in a hemocytometer and added to the macrophages (parasite:macrophage ratio = 10:1). After an incubation period of 18 hours at 37°C (5% CO2), residual free parasites were removed with two washings with medium. Soulamarin was subsequently incubated with infected macrophages (60 h, 37°C, 5% CO2) in a non-toxic concentration range between 3.01 and 306 μM. Benzimidazole was used as a standard. At the end of the assay, slides were fixed with methanol and stained with Giemsa prior to counting under a light microscope. IC_{50} concentrations were obtained by counting 300 amastigotes per well (in duplicate) and determining the number of amastigotes infected per macrophage.

Soulamarin Induce Death in T. cruzi

Soulamarin was added in three concentrations (IC_{100} = 386 μM, IC_{50} = 219 μM and IC_{25} = 103 μM) and the fluorescence was measured after 20, 40 and 60 minutes. The maximum permeability was observed with 0.1% Triton X-100 (positive control). The fluorescence intensity was determined using a spectrofluorimeter. Untreated trypomastigotes and DMSO-treated parasites were used as negative controls.

Effect of soulamarin on the mitochondrial membrane potential

Trypomastigotes were washed with PBS, deposited on a microplate (2×10^6 cells/well) and incubated with soulamarin (IC_{50} = 219 μM) for 60 minutes at 37°C. Then MitoTracker Red CM-H2XROS (500 nM) was added and the incubation was continued for 40 minutes in the dark. Cells were washed twice with BSS (Hank’s buffered salt solution) and the fluorescence was measured using a spectrofluorimeter. Untreated trypomastigotes and DMSO-treated parasites were used as negative controls. Mesoxalonitrile 4-trifluoromethoxyphenylnitrazene (FCCP; 10 μM) was used as a positive control [27]. For the fluorescence microscopy analysis, trypomastigotes were co-stained with 4’,6-diamidino-2-phenylindole (DAPI; 10 μM) and examined at 1000× magnification. Merged images of blue (DAPI) and red (MitoTracker Red) images were obtained using the Nikon NIS - Elements AR software. A Nikon D-FL Epimicroscope equipped with a DS-U3 digital camera was used for the experiment.

Approved by the local ethics committee for animal use (CEUA-IAL/Pasteur 002/2011).

Figure 1. Chemical structure of soulamarin.

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Figure 1. Chemical structure of soulamarin.
Analysis of reactive oxygen species (ROS)

Trypomastigotes (2×10^6 cells/well) were washed in HBSS (Hanks Balanced Salt Solution) medium and incubated with soulamarin (IC_{50} = 219 μM) for 60 minutes at 37°C. To these cells 2',7'-dichlorodihydrofluorescein diacetate (H₂DCF-DA) was added (5 μM) and incubation was prolonged for 15 minutes. Then the fluorescence was measured using a fluorimetric microplate reader (FilterMax F5 Multi-Mode Microplate Reader-Molecular Devices) with excitation and emission wavelengths of 485 and 520 nm, respectively. Oligomycin (20 μM) was used as positive control [26]. Untreated trypomastigotes and parasites treated with DMSO were included as negative controls.

Statistical analysis

Results are displayed as mean values ± standard deviations, which were obtained from at least two independent assays (n=2). IC_{50} values were calculated from sigmoidal dose-response curves using the Graph Pad Prism 3.0 software. Confidence intervals of 95% are included in parentheses. The Student’s t-test was used for significance testing (p<0.05) for all assays.

Results

Chemical characterization of soulamarin

The structure of soulamarin is shown in Figure 1. The assigned structure is consistent with the results obtained from NMR data and LR-ESI mass spectrum. The individual assignment of proton and carbon atoms was accomplished by 1D (1H, 13C) and 2D (HMQC, HMBC and NOESY) NMR measurements. The 1H NMR spectrum of soulamarin in CDCl₃/CD3OD displayed two doublets at δ 6.35 and 5.26, with coupling constants of 10.0 Hz, which were assigned to H-9 and H-10. Together with the presence of a singlet at δ 6.64 (H-12d), three peaks suggested the presence of a chromene moiety [29]. The presence of a dihydrocoumarin segment was based on the multiplets at δ 2.4–2.5 (2H) and δ 3.48 (1H), which were assigned to H-3a/H-3b and H-4, respectively. The multiplets at δ 1.30 (H-14), 1.32 (H-15) and the triplet at δ 80.64 (7 = 7.5 Hz, H-16) were assigned to an ˛-propyl chain linked to C-4 [30]. The doublets at δ 0.97 and 1.28 (7 = 6.4 Hz) were attributed to the methyl groups H-5β and H-5α, while the multiplets at δ 2.93 and 3.93 were assigned to H-2α and H-2β of the isopropyl moiety at C-5. The 13C NMR spectra showed carbonyl carbons at δ 199.1 (C-1), 195.4 (C-2), 171.6 (C-3), 171.0 (C-4), as well as the sp² carbon atoms of the chromene unit at δ 125.5 (C-10) and 115.2 (C-9). Resonances for aromatic carbon atoms were observed between δ 160 and 101, while the carbonil carbon atoms C-3’ and C-11 were observed at δ 78.4 and 77.4, respectively. Additional peaks, corresponding to an n-propyl group were observed at δ 35.1 (C-15), 20.4 (C-14) and 13.7 (C-16). Resonances corresponding to the isopropyl unit were observed at δ 45.5 (C-2’), 19.2 (C-4’) and 10.0 (C-5’). The relative configurations of C-2’ and C-3’ were assigned by comparison of the NMR data with those reported for (2R*,3R*)-2'-hydroxy-4-propyl-5-(3-hydroxy-2-methyl-1-oxobutyl)-6β-(dimethylamino)pyrane-2,3,5,7-tetrahydrobenzopyran-2-one (see Figure 1). The assigned structure was furthermore supported by comparison of our spectroscopic data with those reported in the literature [32].

Antitrypanosomal, cytotoxicity and hemolytic activity of soulamarin

Soulamarin was incubated with trypomastigotes and the activity of cells was determined after 24 hours via MTT assay. Soulamarin thereby demonstrated activity against parasites, killing all the cells at the highest tested concentration. An IC_{50} value of 219.5 μM (95% confidence interval for 186.9–250.5 μM) was established (see Table 1). Benznidazole was used as standard against and resulted in an IC_{50} value of 440.7 μM (95% confidence interval for 272.4–478.4 μM). Soulamarin was also effective against intracellular amastigotes; IC_{50} = 210.1 μM; 95% confidence interval for 174.5–252.6 μM, while benznidazole showed an IC_{50} of 319.7 μM (95% confidence interval for 283.8–360.1 μM). The cytotoxicity of soulamarin was determined with peritoneal macrophages by the MTT assay. Soulamarin showed an IC_{50} value of 908.95 μM and IC_{50} value of 278.3 μM (95% confidence interval for 224.9–342.8 μM). The hemolytic activity was also examined, but soulamarin did not induce any observable hemolysis up to concentrations of 1,300 μM (Table 1).

Modified permeability of the plasma membrane induced by soulamarin

Three different concentrations of soulamarin were incubated for up to 60 minutes with trypomastigotes and the permeability of the plasma membrane was examined by SYTOX Green assay. Soulamarin induced significant increased (p<0.05) fluorescence for all tested concentrations. Highest fluorescence intensities were observed after 60 minutes of incubation (Figure 2). Relative to fully permeabilized parasites (Triton X-100, 60 min), soulamarin induced the following percentages of permeability: i) 81% for IC_{50} = 306 μM (standard error of the mean SEM 6.2) (p<0.05); ii) 60% for IC_{50} = 219 μM (SEM 8.5); iii) 28% for IC_{50} = 103 μM (SEM 1.02). DMSO was used as internal control and resulted in lack of alteration.

Soulamarin–induced depolarization of the mitochondrial membrane potential of T. cruzi

Soulamarin was incubated with trypomastigotes (60 min) and the mitochondrial membrane potential was examined using MitoTracker Red. Spectrofluorimetric measurements indicated that soulamarin induced a significant (97%, p<0.05) decrease in fluorescence levels compared to untreated trypomastigotes (Figure 3A). The control group showed a typical mitochondrial membrane potential. FCCP was used as positive control, which reduced the fluorescence levels by 70% (p<0.05) relative to untreated parasites. Additional fluorescence microscopy experiments corroborated the spectrofluorimetric analysis, demonstrating a substantial reduction of fluorescence levels in soulamarin-treated parasites (Figure 3B), as well as in FCCP (Figure 3D). Untreated parasites showed intense fluorescence levels of mitochondria after labeling with MitoTracker Red, which is consistent with a normal mitochondrial membrane potential (Figure 3C). Panels I represent images with blue fluorescence channel labeled with the fluorescent probe DAPI, panels II represent images with red fluorescence channel labeled with the fluorescent probe MitoTracker Red and panels III, represent the merged images.

Analysis of reactive oxygen species (ROS)

Soulamarin was incubated with trypomastigotes and the up/down-regulation of ROS was examined using 2’,7’-dichlorodihydrofluorescein diacetate (H₂DCF-DA). No changes in the production of ROS could be observed after 60 minutes. Oligomycin was used as positive control (100% ROS up-regulation). Untreated parasites were used as a negative control, showed a normal level of ROS production and were used for normalization (data not shown here).
Discussion

Antitrypanosomal drugs targeting the bioenergetic metabolism as well as the plasma membrane have been considered as potentially chemotherapeutics for Chagas disease [33]. As part of an ongoing search aiming at the isolation of antiparasitic compounds from Brazilian plants [34–36], the coumarin 6-hydroxy-4-propyl-5-(3-hydroxy-2-methyl-1-oxobutyl)-6,6'-dimethylpyrano[2',3':8,7]-benzopyran-2-one (soulamarin) was isolated for the first time from the stem bark of *C. brasiliense* and characterized by $^1$H- and $^{13}$C-NMR analysis as well as by mass spectrometry. Ee and co-workers have recently isolated soulamarin from *C. soulattri*, but no biological activity was described [32]. To the best of our knowledge, this is the first time that an antiparasitic activity against *Trypanosoma cruzi* is reported for soulamarin. The comparable IC$_{50}$ values of soulamarin and benznidazole, which is the currently drug in clinical use, suggests a similar effectiveness against trypomastigotes and intracellular amastigotes. In our assays, benznidazole showed an IC$_{50}$ value of 440.7 μM (95%CI 272.4–478.4 μM) and against intracellular amastigotes - 319.7 μM (95%CI 283.8–360.1 μM).

**Table 1. Evaluation of the 50% Inhibitory Concentration of soulamarin against trypomastigotes and intracellular amastigotes.**

| compound | IC$_{50}$ (μM) trypomastigotes (95%CI) | IC$_{50}$ (μM) amastigotes (95%CI) | Hemolytic (%) activity at 1,300 μM | IC$_{50}$ (μM) Cytotoxicity (95%CI) |
|----------|--------------------------------------|----------------------------------|-----------------------------------|-------------------------------------|
| Soulamarin | 219.8 (186.9–258.5) | 210.1 (174.5–252.6) | 0 | 278.3 (229.4–342.8) |

The viability of trypomastigotes was determined with MTT at 550 nm and the hemolytic activity was determined at 550 nm. The efficacy of soulamarin against intracellular amastigotes was determined using light microscopy counting.

95%CI – 95% confidence interval; IC$_{50}$ – 50% inhibitory concentration. IC$_{50}$ of benznidazole against trypomastigotes - 440.7 μM (95%CI 272.4–478.4 μM) and against intracellular amastigotes - 319.7 μM (95%CI 283.8–360.1 μM).

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Figure 2. Fluorescence measurements (SYTOX Green) of *T. cruzi* after incubation with soulamarin, reflecting the modified permeability of the plasma membrane. Parasites were treated with soulamarin (IC$_{100}$ = 386 μM, IC$_{50}$ = 219 μM and IC$_{25}$ = 103 μM) and compared to Triton X-100 (100% permeability = positive control) as well as an untreated negative control* (p < 0.05).

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evaluation of T. cruzi survival, resulting in different IC_{50} values. Such variations include: i) evaluation of cell lysis by light microscopy counting [37]; ii) spectrophotometric evaluation of MTT oxidation by mitochondrial dehydrogenases [38]; iii) culture-derived trypomastigotes and bloodstream trypomastigotes; iv) parasite strain; v) number parasites per well; vi) time of incubation with benzimidazole, and vii) the source of the drug, which has been produced by different laboratories. Then, comparisons to other IC_{50} values of benzimidazole should be carefully analyzed. Natural products and synthetic compounds affecting the biosynthesis [33] or the permeability of the plasma membrane [39] of T. cruzi have been identified as interesting targets for drug discovery studies. By way of using different fluorimetric probes (SYTOX Green, MitoTracker Red), we targeted in this study the mechanistic aspects on how soulamin rapidly altered the permeability of the plasma membrane, resulting in a dose- and time-dependent influx of the vital dye SYTOX Green into the cell. Fluorescence levels were constant during the observation period (60 min) for all tested concentrations of soulamin. At 386 μM [IC_{100}], soulamin induced the highest fluorescence intensity, which was close to the positive control Triton X-100, suggesting long-term effects on the membrane of the parasite. A similar effect was also observed at the lowest tested concentration [IC_{25} = 103 μM]. Edelfosine, a synthetic lyosphosphilipid drug has also been shown to induce alterations in the plasma membrane and mitochondria of T. cruzi, suggesting that these organelles could be potential targets [40]. However, it has also been proposed that a drug-induced change of the permeability of the plasma membrane is not necessarily accompanied by changes of the mitochondrial function. Digoxin for example, a natural glycoside obtained from Digitalis sp., has been shown to increase plasma membrane permeability in T. cruzi, without affecting the mitochondria [41].

The respiratory apparatus of protozoans typically displays a greater diversity in electron pathways compared to their host cells. The mitochondrion of the protozoan can be considered as a valuable drug target, because of its unique structure and function compared to mammalian cells [42]. The predominant physiological function of the mitochondrion is the generation of ATP by oxidative phosphorylation. Additional functions include the generation and detoxification of ROS, the involvement in some forms of apoptosis and the regulation of cytoplasmic and mitochondrial calcium [43]. Furthermore, a proper mitochondrial membrane potential is essential for the survival of cells and changes can result in a variety of consequence, such as the inhibition of the electron transport chain, the inhibition of ATP synthase, the stimulation of uncoupling proteins or the permeability of the inner membrane [44]. In our assays, soulamin rapidly induced a depolarization of the mitochondrial membrane potential in trypomastigotes, resulting in a reduction of the fluorescence intensity by 97%, relative to the untreated group. This effect may have contributed to deleterious cellular damages associated with bioenergetic system. A similar, potent dose-dependent collapse of the mitochondrial membrane potential, resulting in the killing of T. cruzi parasites, has been reported for synthetic naphthofuranquinones [45].

Under physiological conditions, the oxidative phosphorylation involving an electron transport to pump hydrogen ions across the inner membrane, releases ROS, amounting to 3–5% of the total amount of oxygen consumed [46]. Under pathologic conditions, several pathways result in excessive ROS production, which causes - if not efficiently scavenged by the antioxidant system - oxidative stress. Proteins, lipids, and DNA are readily oxidized by ROS, resulting in dysfunction of vital physiological processes, oxidative damage, and cell death [47]. In our assays, despite the substantial depolarization of the mitochondrial membrane potential of T. cruzi, soulamin induced no up-regulation of ROS compared to untreated trypomastigotes. Within the mitochondria, the primary site of ROS production is the electron transport chain, which involves four protein-associated complexes [48]. Several cellular enzyme systems are potential sources of ROS: NAD(P)H oxidase, xanthine oxidase, uncoupled endothelial nitric oxide synthase (eNOS), arachidonic acid metabolizing enzymes such as cytochrome P450 enzymes, lipooxygenase and cyclooxygenase, as well as the mitochondrial respiratory chain. Considering that a large number of drugs, which affect mitochondria also contribute to an up-regulation of ROS [49–50], we propose that soulamin could target T. cruzi mitochondria without affecting the enzymes mentioned above.
Conclusion

Soulamin was isolated for the first time from the stem bark of *C. brasiliensis* and showed desirable anti-trypanosomal activity. Our results furthermore indicated that soulamin-induced death in *T. cruzi* is associated with mitochondrial dysfunction and a modified permeability of the plasma membrane. Therefore, the natural product soulamin could serve as a scaffold for the development of selective new drugs against neglected diseases, in particular Chagas disease.

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Author Contributions

Conceived and designed the experiments: AGT JHGL. Performed the experiments: AR AGT EGP JTM LGMS JHGL. Analyzed the data: AR AGT JHGL. Contributed reagents/materials/analysis tools: AGT PS ER JHGL. Wrote the paper: AGT JHGL.

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