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Wayanin and guaijaverin, two active metabolites found in a *Psidium acutangulum* Mart. ex DC (syn. *P. personii* McVaugh) (Myrtaceae) antimalarial decoction from the Wayana Amerindians

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Keywords: *Psidium acutangulum*; Traditional remedy; Cytokines; Antimalarial; Glycosylated flavonols; French Guiana

Abstract

Ethnopharmacological relevance

Psidium acutangulum Mart. ex DC is a small tree used by the Wayana Amerindians from the Upper-Maroni in French Guiana for the treatment of malaria.

Aim of the study

In a previous study, we highlighted the *in vitro* antiplasmodial, antioxidant and anti-inflammatory potential of the traditional decoction of *P. acutangulum* aerial parts. Our goal was then to investigate on the origin of the biological activity of the traditional remedy, and eventually characterize active constituents.

Materials and methods

Liquid-liquid extractions were performed on the decoction, and the antiplasmodial activity evaluated against chloroquine-resistant FcB1 ($[^3\text{H}]$ -hypoxanthine bioassay) and 7G8 (pLDH bioassay) *P. falciparum* strains, and on a chloroquine sensitive NF54 ($[^3\text{H}]$ -hypoxanthine bioassay) *P. falciparum* strain. The ethyl acetate fraction (D) was active and underwent bioguided fractionation. All the isolated compounds were tested on *P. falciparum* FcB1 strain. *In vitro* anti-inflammatory activity (IL-1 β , IL-6, IL-8, TNF α) of the ethyl acetate fraction and of an anti-*Plasmodium* active compound, was concurrently assessed on LPS-stimulated human PBMC and NO secretion inhibition was measured on LPS stimulated RAW murine macrophages. Cytotoxicity of the fractions and pure compounds was measured on VERO cells, L6 mammalian cells, PBMCs, and RAW cells.

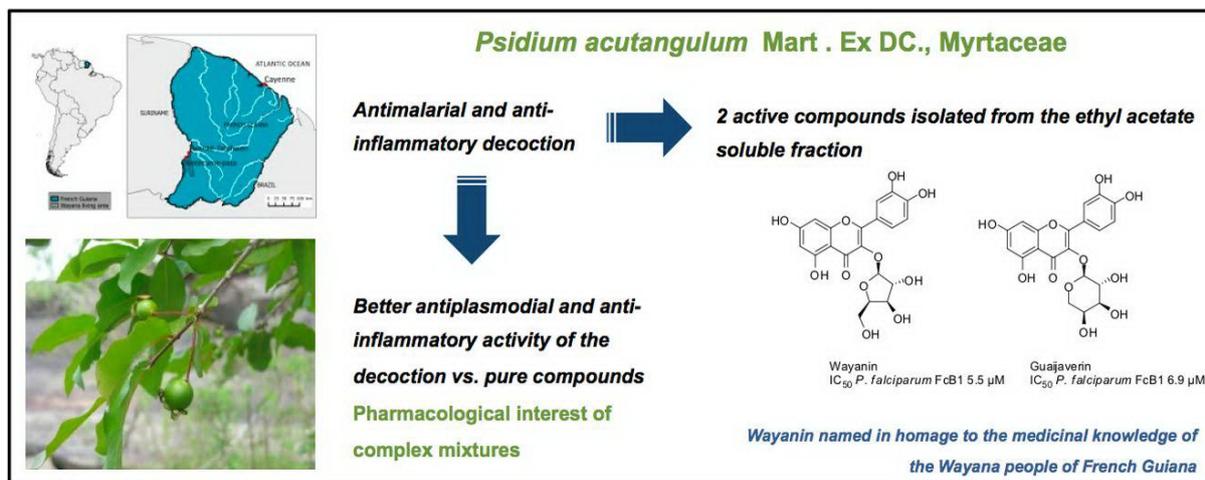
Results

Fractionation of the ethyl acetate soluble fraction (IC_{50} ranging from 3.4 to $< 1 \mu\text{g/mL}$ depending on the parasite strain) led to the isolation of six pure compounds: catechin and five glycosylated quercetin derivatives. These compounds have never been isolated from this plant species. Two of these compounds (wayanin and guaijaverin) were found to be moderately active against *P. falciparum* FcB1 *in vitro* (IC_{50} 5.5 and 6.9 μM respectively). We proposed the name wayanin during public meetings organized in June 2015 in the Upper-Maroni villages, in homage to the medicinal knowledge of the Wayana population. At 50 $\mu\text{g/mL}$, the ethyl acetate fraction (D) significantly inhibited IL-1 β secretion (-46 %) and NO production (-21 %), as previously observed for the decoction. The effects of D and guaijaverin (**4**) on the secretion of other cytokines or NO production were not significant.

Conclusions

The confirmed antiplasmodial activity of the ethyl acetate soluble fraction of the decoction and of the isolated compounds support the previous results obtained on the *P. acutangulum* decoction. The antiplasmodial activity might be due to a mixture of moderately active non-

toxic flavonoids. The anti-inflammatory activities were less marked for ethyl acetate fraction (D) than for the decoction.



1. Introduction

Psidium acutangulum Mart. ex DC. is a shrub used by the local populations against diarrhea and abdominal pain (Fleury, 1991; Grenand et al., 2004). Its antimalarial use was reported during previous studies performed among the Upper-Maroni Wayana community (Fleury, 2003; 2007), where fresh aerial parts are prepared as a decoction. Patients with malaria crisis are washed with the remedy, and take it orally as well.

In Houël et al. (2015), we highlighted *P. acutangulum* decoction's antimalarial properties and demonstrated *in vitro* a significant dose dependent effect of the remedy on selected cytokines secretion and NO production, without cytotoxicity on L6 cell line, RAW cells and PBMC. This species is however poorly described on a phytochemical point of view, aside from the isolation of flavonols and α,β -unsaturated ketones (Miles et al., 1991; Rivero-Maldonado et al., 2013), the latter family of compounds being interestingly known to exhibit antimalarial activity (Singh et al., 2014). Our aim was therefore to gain a better insight into the remedy's antimalarial and anti-inflammatory activities.

2. Materials and methods

2.1. General experimental procedures

^1H NMR spectra were recorded at 400 MHz and ^{13}C NMR spectra at 100.6 MHz on a Varian 400 MR spectrometer equipped with a 5 mm inverse probe (Auto X PGF $^1\text{H}/^{15}\text{N}-^{13}\text{C}$).

Samples were dissolved in deuterated acetone ($\text{C}_3\text{D}_6\text{O}$) or deuterated methanol (CD_3OD) in 5 mm tubes as stated. Chemical shifts are in ppm downfield from tetramethylsilane (TMS), and coupling constants (J) are in Hz (s stands for singlet, d for doublet, t for triplet, q for quartet, m for multiplet, br for broad, *ax* for axial and *eq* for equatorial, and n.d. for not detected). ^{13}C chemical shifts were deduced from proton-carbon correlation experiments.

Water (HPLC grade) was obtained from a Milli-Q system (Milli-Q plus, Millipore Bedford, MA). Analyses were performed on a Discovery C18 column (15 cm x 4.6 mm, 5 μm , Supelco) at 1 mL/min using two HPLC systems (Waters and Varian). The first system was equipped with a W2996 photodiode array absorbance detector and a W2424 light-scattering detector (Waters) and the second system with a photodiode array absorbance detector (Varian). Preparative chromatography was performed at 15 mL/min with a W600 pump and a W2487 double wavelength UV detector (Waters) using a Discovery C18 column (15 cm x 21.2 mm, 5 μm , Supelco). Fractions were automatically collected (Waters Fraction Collector III) and evaporated at 37°C under reduced pressure with a speedvac concentrator (Savant SPD121P, Thermo Scientific).

UV spectra of pure compounds were measured with the W2996 photodiode array absorbance detector (Waters).

Radioactivity was counted using a Wallac® 1450 Betalux scintillation counter (1450-Microbeta Trilux, Wallac Perkin Elmer).

2.2. Plant material

P. acutangulum stems, leaves, and fruits were collected in Twenke-Taluhwen (herbarium voucher MF2176, deposited in Cayenne Herbarium - CAY). Botanical identification was confirmed by Marie-Françoise Prévost. Aerial parts were dried prior to sample preparation to prevent degradation of the vegetal material during transportation between Twenke-Taluhwen and Cayenne.

2.3. Ethical aspects of the study

P. acutangulum was collected in Twenke-Taluhwen in February 2007 prior to the creation of the Amazonian Natural Park of French Guiana (PAG). Therefore, collect authorizations were unnecessary at the time of the collect. Interviews have been undertaken according to French Guiana legislation at the time of the study. The interviewees' participation to the study was voluntary, and as requested by the Wayana population, some of the results were published in a community-oriented booklet (Fleury, 2007). Moreover, public meetings were realized in June 2015 in the Upper-Maroni villages, where we proposed to give the name wayanin to quercetin-3-O- β -D-xylofuranoside, with the agreement of the Wayana traditional authorities and in homage to the Wayana medicinal knowledge.

2.4. Extraction and isolation

300 mL of a decoction (A) were prepared from 50 g of hand-crushed dried aerial parts of *P. acutangulum* (stems, leaves and fruits) placed in a 2-L Erlenmeyer flask with 1 L of cold water, and submitted to decoction during 2 h. Leaves, stems, and fruits were removed by filtration after cooling and the decoction was submitted to successive liquid-liquid extractions. The decoction was first extracted with hexane (3 \times 80 mL). Filtration and evaporation of the solvent yielded 11.1 mg of hexane extract (B). The aqueous layer (C) was then extracted with ethyl acetate (3 \times 80 mL). Filtration and evaporation of the solvent yielded 489.0 mg of ethyl

acetate extract (D). Aliquots of the remaining aqueous phase (E) were also evaporated and submitted to biological evaluation.

The antiplasmodial ethyl acetate extract (D) was diluted in water/acetonitrile (90:10).

Separations were carried out on 1 mL of solution at 10 mg/mL using a linear gradient of water/acetonitrile (90:10 to 50:50 over 20 min) and remaining at 100% acetonitrile during 10 min to give 14 fractions. The eluates from the column were monitored at 214 and 254 nm.

Purification of 10 mg of the crude extract allowed after evaporation for isolation of 0.9 mg of compound **1**, identified as catechin. For the other compounds, successive injections were performed and fractions containing the same compounds were put together, allowing after evaporation for isolation of 5.0 mg of compound **2** from 55.0 mg of crude extract, and for obtaining of 11.4 mg of a mixture of compounds obtained from 70 mg of crude extract. This mixture was then submitted to a second separation. This separation was carried out on using water/acetonitrile 83:17 for 20 min followed by a linear gradient from water/acetonitrile 83:17 to 50:50 over 5 min. The eluates from the column were monitored at 214 and 254 nm.

Fractions containing the same constituents were gathered and evaporation allowed for isolation of compounds **3-6** from 9.1 mg of injected mixture. The isolated amounts of the pure compounds were the following: **3**: 1.0 mg; **4**: 0.4 mg; **5**: 0.8 mg; **6**: 0.4 mg. All compounds were isolated in pure form.

2.5. Analytical data for compounds **1-6**

Catechin (**1**); UV λ_{\max} (H₂O/ACN) 279.6 nm; ¹H NMR (400 MHz, C₃D₆O) δ ppm 6.89 (d, J = 2.0 Hz, 1H, H_{2'}), 6.79 (d, J = 8.2 Hz, 1H, H_{5'}), 6.75 (dd, J = 8.2 ; 2.0 Hz, 1H, H_{6'}), 6.02 (d, J = 2.3 Hz, 1H, H₆), 5.88 (d, J = 2.3 Hz, 1H, H₈), 4.56 (d, J = 7.8 Hz, 1H, H₂), 3.99 (br td, J = 8.2 ; 5.5 Hz, 1H, H₃), 2.91 (dd, J = 16.1 ; 5.5 Hz, 1H, H_{4a}), 2.53 (dd, J = 16.1 ; 8.5 Hz, 1H, H_{4b}). ¹³C RMN (100.6 MHz, C₃D₆O) δ ppm 158.2 (C₇), 157.7 (C₅), 156.9 (C₉), 146.39 (C_{3'}),

146.43 (C_{4'}), 132.9 (C_{1'}), 120.8 (C_{6'}), 116.5 (C_{5'}), 116.0 (C_{2'}), 101.3 (C_{10'}), 96.9 (C_{6'}), 96.2 (C_{8'}), 83.5 (C_{2'}), 69.1 (C_{3'}), 29.6 (C_{4'}).

Quercetin-3-O-β-D-xylofuranoside (wayanin, **2**); UV λ_{max} = 265.3 / 352.3 nm (H₂O/CH₃CN); ¹H NMR (400 MHz, CD₃OD) δ ppm 7.56 (s, 1H, H_{2'}) 7.54 (d, *J* = 8.4 Hz, 1H, H_{6'}), 6.73 (m, 1H, H_{5'}), 6.34 (br s, 1H, H_{6'}), 6.18 (br s, 1H, H_{8'}), 5.20 (d, *J* = 7.3 Hz, 1H, H_{1''}), 4.30 (m, 2H, H_{5''}), 3.49 (m, 3H, H_{2''}, H_{3''}, H_{4''}). ¹³C NMR (100.6 MHz, CD₃OD) δ ppm n.d. (C_{4'}), 165.8 (C_{7'}), 162.6 (C_{5'}), 159.1 (C_{2'}), 158.0 (C_{9'}), 149.4 (C_{4'}), 145.5 (C_{3'}), 135.0 (C_{3'}), 123.3 (C_{6'}), 122.7 (C_{1'}), 117.0 (C_{2'}), 115.8 (C_{5'}), 105.2 (C_{10'}), 104.0 (C_{1''}), 99.7 (C_{8'}), 94.5 (C_{6'}), 77.9 (C_{4''}), 75.6 (C_{2''}), 71.3 (C_{3''}), 64.1 (C_{5''}).

Quercetin-3-O-β-D-xylopyranoside (reynoutrin, **3**); UV λ_{max} (H₂O/ACN) 254.7 / 354.7 nm; ¹H NMR (400 MHz, CD₃OD) δ ppm 7.60 (dd, *J* = 2.1 ; 0.5 Hz, 1H, H_{2'}), 7.59 (dd, *J* = 8.2 ; 2.1 Hz, 1H, H_{6'}), 6.85 (dd, *J* = 8.2 ; 0.5 Hz, 1H, H_{5'}), 6.32 (d, *J* = 2.1 Hz, 1H, H_{6'}), 6.14 (d, *J* = 2.1 Hz, 1H, H_{8'}), 5.13 (d, *J* = 7.2 Hz, 1H, H_{1''}), 3.78 (dd, *J* = 11.7 ; 5.1 Hz, 1H, H_{5''eq}), 3.51 (dd, *J* = 8.8 ; 7.2 Hz, 1H, H_{2''}), 3.51 (m, 1H, H_{4''}), 3.39 (br t, *J* = 8.6 Hz, 1H, H_{3''}), 3.10 (dd, *J* = 11.7; 9.5 Hz, 1H, H_{5''ax}). ¹³C NMR (100.6 MHz, CD₃OD) δ ppm 178.4 (C_{4'}), 166.8 (C_{7'}), 162.5 (C_{5'}), 158.4 (C_{9'}), 158.0 (C_{2'}), 149.6 (C_{4'}), 145.7 (C_{3'}), 134.9 (C_{3'}), 123.0 (C_{6'}), 122.6 (C_{1'}), 116.8 (C_{2'}), 115.7 (C_{5'}), 104.6 (C_{1''}), 104.1 (C_{10'}), 100.7 (C_{8'}), 95.2 (C_{6'}), 77.4 (C_{3''}), 75.0 (C_{2''}), 70.8 (C_{4''}), 66.9 (C_{5''}).

Quercetin-3-O-α-L-arabinopyranoside (guajiverin, **4**); UV λ_{max} (H₂O/ACN) 254.7 / 358.8 nm ; ¹H NMR (400 MHz, CD₃OD) δ ppm 7.74 (d, *J* = 2.2 Hz, 1H, H_{2'}), 7.58 (dd, *J* = 8.5; 2.2 Hz, 1H, H_{6'}), 6.86 (d, *J* = 8.5 Hz, 1H, H_{5'}), 6.33 (d, *J* = 2.0 Hz, 1H, H_{6'}), 6.15 (d, *J* = 2.0 Hz, 1H, H_{8'}), 5.11 (d, *J* = 6.6 Hz, 1H, H_{1''}), 3.90 (dd, *J* = 8.4 ; 6.6 Hz, 1H, H_{2''}), 3.82 (dd, *J* = 12.4 ; 2.9 Hz, 1H, H_{5''b}), 3.81 (m, 1H, H_{4''}), 3.64 (dd, *J* = 8.4 ; 3.2 Hz, 1H, H_{3''}), 3.44 (dd, *J* = 12.4 ; 3.0 Hz, 1H, H_{5''a}). ¹³C NMR (100.6 MHz, CD₃OD) δ ppm 178.0 (C_{4'}), 168.0 (C_{7'}), 160.5 (C_{5'}), 158.6 (C_{9'}), 157.8 (C_{2'}), 149.9 (C_{4'}), 145.8 (C_{3'}), 135.1 (C_{3'}), 122.6 (C_{6'}), 122.5 (C_{1'}), 117.0 (C_{2'}), 115.9 (C_{5'}), 104.7 (C_{1''}), 104.3 (C_{10'}), 100.9 (C_{8'}), 95.4 (C_{6'}), 74.0 (C_{3''}), 72.7 (C_{2''}), 68.9 (C_{4''}), 66.8 (C_{5''}).

Quercetin-3-*O*- α -L-arabinofuranoside (avicularin, **5**); UV λ_{\max} (H₂O/ACN) 255.9 / 351.1

nm. ¹H NMR (400 MHz, CD₃OD) δ ppm 7.52 (d, J = 2.0 Hz, 1H, H₂), 7.48 (dd, J = 8.4 ; 2.0 Hz, 1H, H₆), 6.89 (d, J = 8.4 Hz, 1H, H₅), 6.34 (d, J = 2.1 Hz, 1H, H₆), 6.17 (d, J = 2.1 Hz, 1H, H₈), 5.45 (brs, 1H, H_{1''}), 4.32 (dd, J = 2.9, 1.0 Hz, 1H, H_{2''}), 3.91 (dd, J = 5.2 ; 2.9 Hz, 1H, H_{3''}), 3.87 (brq, J = 4.5 Hz, 1H, H_{4''}), 3.49 (m, 1H, H_{5''a}), 3.48 (dd, J = 12.0 ; 4.6 Hz, 1H, H_{5''b}). ¹³C NMR (100.6 MHz, CD₃OD) δ ppm 176.5 (C₄), 167.9 (C₇), 163.0 (C₅), 158.9 (C₂), 158.7 (C₉), 149.7 (C₄), 146.2 (C₃), 134.6 (C₃), 122.8 (C_{1'}), 122.6 (C₆), 116.5 (C₂), 116.1 (C₅), 109.3 (C_{1''}), 104.6 (C₁₀), 99.7 (C₈), 94.5 (C₆), 87.7 (C_{4''}), 83.0 (C_{2''}), 78.4 (C_{3''}), 62.2 (C_{5''}).

Quercetin-3-*O*- α -L-rhamnopyranoside (quercitrin, **6**); UV λ_{\max} (H₂O/ACN) 254.7 /

345.1 nm. ¹H NMR (400 MHz, CD₃OD) δ ppm 7.33 (d, J = 2.1 Hz, 1H, H₂), 7.30 (dd, J = 8.2 ; 2.1 Hz, 1H, H₆), 6.90 (d, J = 8.2 Hz, 1H, H₅), 6.31 (d, J = 2.1 Hz, 1H, H₆), 6.15 (d, J = 2.1 Hz, 1H, H₈), 5.34 (d, J = 1.7 Hz, 1H, H_{1''}), 4.21 (dd, J = 3.4 ; 1.7 Hz, 1H, H_{2''}), 3.74 (dd, J = 9.4 ; 3.4 Hz, 1H, H_{3''}), 3.42 (m, 1H, H_{5''}), 3.34 (m, 1H, H_{4''}), 0.94 (d, J = 6.1 Hz, 3H, H_{6''}). ¹³C NMR (100.6 MHz, CD₃OD) δ ppm 179.3 (C₄), 168.6 (C₇), 162.7 (C₅), 158.6 (C₂), 159.9 (C₉), 149.4 (C₄), 149.6 (C₃), 133.7 (C₃), 122.8 (C_{1'}), 122.5 (C₆), 116.6 (C₂), 116.1 (C₅), 106.2 (C₁₀), 103.4 (C_{1''}), 100.5 (C₈), 95.1 (C₆), 72.7 (C_{4''}), 71.8 (C_{2''}), 71.8 (C_{3''}), 17.4 (C_{6''}).

Spectra obtained from proton and proton-carbon correlation experiments are given as Supplementary Data.

2.6. *In vitro* antiplasmodial activity

2.6.1. [³H]-hypoxanthine assay

[³H]-hypoxanthine assays were performed on *P. falciparum* FcB1 strain (chloroquine resistant) and NF54 strain (chloroquine sensitive) according to a procedure adapted from Le Bras and Deloron (1983) and Benoit et al. (1996) as reported previously (Houël et al., 2015). Decreasing concentrations of the extracts and pure compounds (100 to 1 μ g/mL, 3 concentrations for FcB1 strain and 100 to 0.002 μ g/mL, 11 concentrations for NF54 strain)

were used. Each experiment was performed 3 times in triplicate, and negative (DMSO) and positive (chloroquine) controls were also performed. Growth/inhibition percentage curves were obtained and the concentration required to inhibit 50% growth (IC_{50}) was determined graphically (FcB1 strain) or calculated by linear regression (NF54 strain). [3H]-hypoxanthine was discarded according to French legislation on elimination of radioactive residue (www.asn.fr/).

2.6.2. pLDH assay

pLDH immunodetection assays were performed against *P. falciparum* 7G8 chloroquine-resistant strain, with a commercially available sandwich enzyme immunosorbent assay (ApDia), as reported previously (Atchade et al., 2013). Parasite culture (1% parasitaemia, 2% hematocrit) was incubated with increasing concentrations of the extract (1 to 100 $\mu\text{g/ml}$, 8 concentrations) for 96 h at 7°C under reduced oxygen conditions (candle jar). Each experiment was performed 3 times in duplicate, and chloroquine was used as a positive control. IC_{50} were calculated with GraphPad Prism 6 software.

2.7. In vitro anti-inflammatory activity

2.7.1. Inhibitory activity on NO production from macrophage-like cell line RAW 264.7

The bioassays were performed as reported previously (Houël et al., 2015). Briefly, RAW 264.7 cells (ATCC TIB-71) were stimulated with lipopolysaccharide (LPS, Sigma, 5 $\mu\text{g/mL}$) and treated with 2 concentrations of ethyl acetate fraction (D) or pure guaijaverin (**4**) (50 $\mu\text{g/mL}$ or 10 $\mu\text{g/mL}$), diluted in DMSO. Quercetin at 50 $\mu\text{g/mL}$ and 10 $\mu\text{g/mL}$ (final DMSO concentration 1% v/v) was used as a positive control. Unstimulated cells (without LPS) are considered as negative control. Plates were then incubated for 24 h and nitrite (NO_2^-) accumulation was determined as an indicator of NO production in culture media as previously described (Kumar-Roiné et al., 2009). Each measurement was performed in triplicate and results were expressed as means for three independent experiments.

Cytotoxicity was measured by the WST-1 assay (Ozyme, according to supplier protocol). All experiments were performed on a fully automated platform (Beckman Coulter).

2.7.2. Peripheral blood mononuclear cells (PBMCs) isolation and culture

PBMCs were prepared from the peripheral blood of healthy donors (Etablissement Français du Sang) as reported previously (Houël et al., 2015). Isolated PBMCs were stimulated with 5 µg/mL LPS (Sigma). Ethyl acetate fraction (D) or pure guaijaverin (**4**) were dissolved in DMSO so that final DMSO concentration is 1% and tested at 10 and 50 µg/mL in triplicate. Positive control dexamethasone dissolved in ethanol so that final EtOH concentration is 1% was simultaneously tested at 10 and 50 µg/mL. Cells without LPS stimulation were considered as negative control (basal level of cytokine). Cells incubation was carried out at 37°C (5% CO₂).

2.7.3. Characterization of cytokine secretion.

After 24 h, PBMCs supernatants were transferred into a 384-well plate. Cytokine detection was performed using HTRF technology (Homogeneous Time Resolved Fluorescence, Cisbio bioassays) for TNFα (62TNFPEC), IL-1β (62IL1PEC), IL-6 (62IL6PEB), IL-8 (62IL8PEB) according to supplier recommendations. Reading was performed after 2.5 h of incubation using Envision multi-labelled reader (Perkin Elmer) with supplier recommended parameters. A standard curve was performed for each cytokine to determine the concentration of released cytokines (in pg/mL) by PBMCs in the supernatant. The cytokine secretion is expressed as a percentage of the cytokine concentration measured in the negative control experiment conducted on the same plate.

2.7.4. Cell viability assay

Cell viability was measured using the WST-1 (Ozyme) assay according to the manufacturers' protocol. Briefly, after supernatant transfer for cytokine determination, WST-1-containing medium was added to cells and cell viability was determined by measuring absorbance at

450 nm using Victor³ reader (Perkin Elmer) after 2 h incubation at 37°C. Each measurement was performed in triplicate and results were expressed as means of three independent experiments.

2.8. *In vitro* cytotoxicity study

2.8.1. *In vitro* cytotoxicity study on VERO cells

Cytotoxicity on VERO cells (African Green Monkey kidney epithelial cells) was evaluated as reported previously (Cachet et al., 2009). Briefly, the cell line was cultured under the same conditions as used for *P. falciparum*, except for the 5% human serum, which was replaced by 10% fetal calf serum (Boehringer). Cells were seeded (2×10^5 cells/well) during their log growth phase in 96-well flat-bottom plates after addition of trypsin. Cells were cultured for 72 h at 37°C and 5% CO₂, extracts/molecules being added after 24 h, at concentrations ranging from 100 µg/ml to 0.01 µg/ml. Cell growth was measured by [³H]-hypoxanthine incorporation after a 48 h incubation. The amount of [³H]-hypoxanthine incorporated in the presence of drugs was compared with that of control cultures (solvent for negative control, doxorubicin for positive control). All experiments were performed in triplicate.

2.8.2. *In vitro* cytotoxicity study on L-6 cells

The cytotoxicity of the fractions was assessed by using the L-6 cell line (rat skeletal myoblasts) and the method of Ahmed et al. (1994). The cytotoxicity assays were performed according to previously published procedures (Houël et al., 2015). After incubation (37°C, 5% CO₂) with the drugs for 70 hours and further 2 hours with resazurin, the plates were read with a Spectramax Gemini XS microplate fluorometer at an excitation wavelength of 536 nm and an emission wavelength of 588 nm. The IC₅₀ values were calculated from the sigmoidal inhibition curves with SoftMax Pro software. Podophyllotoxin (Sigma) was used as a positive control. The given values are the means of two independent assays.

2.9. Statistical analysis

Data were analysed by one-way analysis of variance followed by Tukey's multiple comparison tests using GraphPad Software version 5.04. $P < 0.05$ was considered significant. In figures 2 and 3, whiskers show the maximal and minimal values. The rectangular box represents the 25% quantile to 75% quantile ranges. The dark line shows the median of the distribution.

3. Results and discussion

3.1. Characterization of compounds from the ethyl acetate fraction (D) of the decoction

Antiplasmodial activity of the decoction and of liquid phases obtained during successive extractions was evaluated, and it was found that the ethyl acetate extract (D) was the most active. Naturally, this extract was chosen for bioguided fractionation. The following metabolites were isolated and identified by 1D and 2D NMR spectroscopy, and by comparison with published data: catechin (**1**, 9.1 % w/w) (Davis et al., 1996), wayanin (quercetin-3-O- β -D-xylofuranoside, **2**, 9.1 % w/w) (Moravcova et al., 1997; Saldanha et al., 2013; Saxena et Chaturvedi, 1992; Vvedenskaya et al., 2004; Xue et al., 2006), reynoutrin (quercetin-3-O- β -D-xylopyranoside, **3**, 1.8 % w/w) (Eparvier et al., 2007; Lu and Yeap Foo, 1997), guajaverin (quercetin-3-O- α -L-arabinopyranoside, **4**, 0.7 % w/w) (Stark et al., 2005; Vvedenskaya et al., 2004), avicularin (quercetin-3-O- β -L-arabinofuranoside, **5**, 1.5 % w/w) (Lu and Yeap Foo, 1997; Vvedenskaya et al., 2004) and quercitrin (quercetin-3-O- α -L-rhamnopyranoside, **6**, 0.7 % w/w) (Lu and Yeap Foo, 1997) (Fig. 1). Isolation yields are given in % based on dry ethyl acetate crude extract. Quercetin-3-O- β -D-xylofuranoside, which we named wayanin, was recently isolated from *Myrcia bella* Cambess. (Myrtaceae) leaves hydroalcoholic extract concurrently with avicularin, quercitrin and reynoutrin (Saldanha et al., 2013). Reynoutrin has been isolated recently from the genus *Psidium* (Ho et al., 2012). Avicularin and guajaverin are known since the late 1950s as constituents of *P. guajava*

leaves (El Khadem and Mohammed, 1958). These widely ubiquitous compounds frequently coexist, together with catechin and quercitrin, particularly within the Myrtaceae family (Gallo et al., 2008; Shu et al., 2012; Simirgiotis et al., 2008). These molecules are described for the first time in *Psidium acutangulum*. In fact, only few compounds had been previously described in *P. acutangulum*, among which the α,β -unsaturated ketone 3'-formyl-2',4',6'-trihydroxy-5'-methylidihydrochalcone, isolated from a dichloromethane extract (Miles et al., 1991), and myricetin, identified in *P. acutangulum* leaves (Rivero-Maldonado et al., 2013).

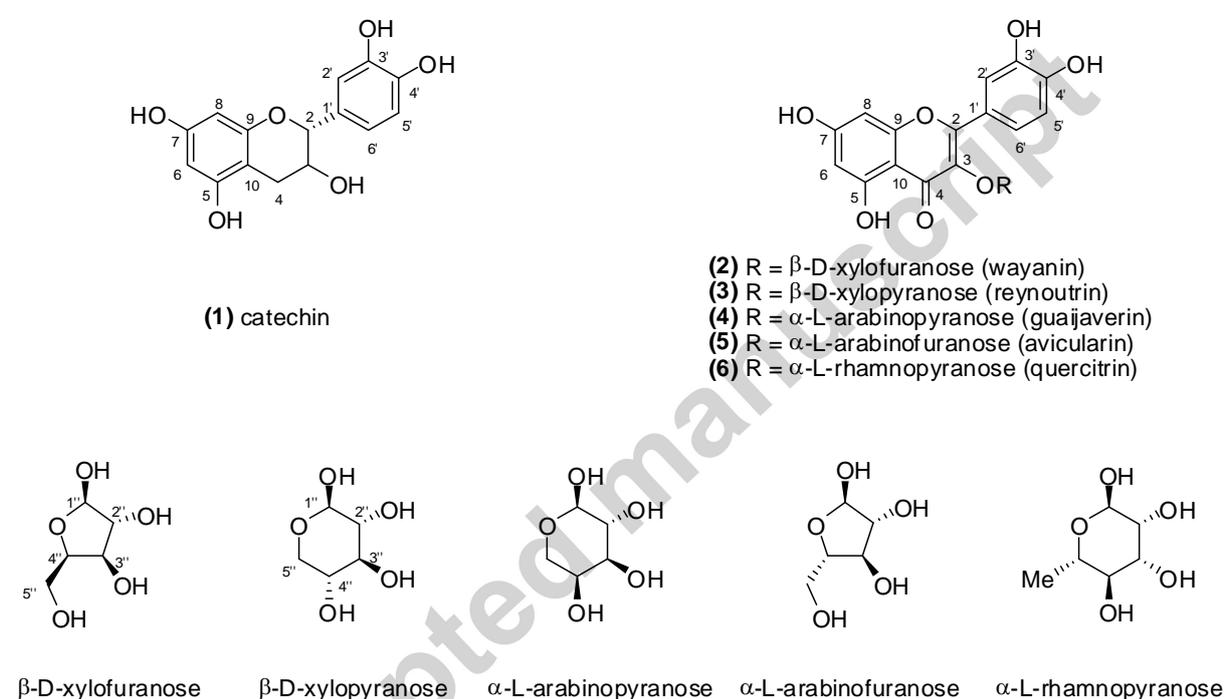


Fig. 1. Catechin and glycosylated quercetin derivatives isolated from *P. acutangulum* decoction

3.2. Biological activities

The decoction (A) had been shown to display a good antiplasmodial activity *in vitro* against various strains, regardless to the bioassay used, with IC_{50} values of 3.3 μ g/mL and 10.3 μ g/mL against *P. falciparum* FcB1 and NF54, respectively and 19.0 μ g/mL against *P. falciparum* 7G8 (Houël et al., 2015), and these results are reminded about in Table 1. The

successive extracts (B-E) obtained through liquid-liquid fractionation were therefore submitted to antimalarial tests *in vitro*, using *P. falciparum* chloroquine-resistant FcB1 strain. The results are presented in Table 1. The antimalarial activity was retained during fractionation, with a good activity observed for the aqueous fraction (C) ($IC_{50} = 2.7 \mu\text{g/mL}$). The most active fraction was the ethyl acetate one (D) which displayed a remarkable IC_{50} value ($< 1 \mu\text{g/mL}$). Cytotoxicity was evaluated on VERO and L6 cells. As previously observed for the decoction, the most active fraction (D) was not cytotoxic against VERO cells ($IC_{50} > 100 \mu\text{g/mL}$), and exhibited a low cytotoxicity against L6 cells ($IC_{50} = 57.4 \mu\text{g/mL}$).

Table 1. *In vitro* antiplasmodial and cytotoxic activities of extracts obtained by successive liquid/liquid fractionation of *P. acutangulum* decoction. Results are in $\mu\text{g/ml}$ for extracts and positive controls.

	$IC_{50} \pm SD$ <i>P. falciparum</i> FcB1	$IC_{50} \pm SD$ <i>P. falciparum</i> NF54	$IC_{50} \pm SD$ <i>P. falciparum</i> 7G8 ^a	IC_{50} VERO cells	$IC_{50} \pm SD$ L6 cells
(A) (decoction)	3.3 ± 0.4	10.3 ± 0.1	19.0 ± 2.7	n.t.	> 100
(B) (hexane)	32.3 ± 4.4	n.t. ^b	n.t.	n.t.	> 100
(C) (water)	2.7 ± 0.7	11.3 ± 1.7	13.6 ± 0.3	n.t.	> 100
(D) (ethyl acetate)	< 1	2.3 ± 0.6	2.6 ± 0.02	> 100	57.4 ± 20.5
(E) (water)	21.2 ± 3.8	9.4 ± 0.2	12.9 ± 0.1	n.t.	> 100
Chloroquine	0.18 ± 0.03	0.002 ± 0.0001	0.13 ± 0.002	n.t.	n.t.
Podophyllotoxin	n.t.	n.t.	n.t.	n.t.	0.007 ± 0.003
Doxorubicin	n.t.	n.t.	n.t.	< 0.02	n.t.

^apLDH (*Plasmodium* lactate dehydrogenase immunodetection) assay

^bNot tested

Pure compounds isolated from the AcOEt fraction were also evaluated for antiplasmodial activity against *P. falciparum* FcB1 and for cytotoxic activity on VERO cells (Table 2).

Table 2. Antiplasmodial activity (*P. falciparum* chloroquine resistant FcB1 strain) and cytotoxicity of the pure compounds isolated from fraction D. Errors for individual

measurements differed by less than 20%. Data represents the average of three independent determinations.

	Antiplasmodial activity <i>P. falciparum</i> FcB1 IC ₅₀ (μM)	Cytotoxic activity VERO cells IC ₅₀ (μM)
Catechin (1)	72.3	n.t. ^a
Wayanin (2)	5.5	> 200
Reynoutrin (3)	26.5	> 200
Guaijaverin (4)	6.9	> 200
Avicularin (5)	64.5	> 200
Quercitrin (6)	71.4	> 200
Chloroquine	0.36	> 200

^a Not tested

Our results thus provide the first evidence for antiplasmodial activity of the well-known guaijaverin and of quercetin-3-O-β-D-xylofuranoside against *P. falciparum*. Although the latter was recently isolated from *Myrcia bella* hydroalcoholic leaves extract (Saldanha et al., 2013), no common name was given to it. We therefore proposed to name it wayanin during public meetings realized in June 2015 in the Upper-Maroni villages, with the agreement of the Wayana traditional authorities, and in homage to the Wayana medicinal knowledge. The two compounds exhibited IC₅₀ of 6.9 and 5.5 μM, respectively, and the other constituents of the fraction D were only moderately active or inactive. To our knowledge, reynoutrin and avicularin had never been evaluated against *P. falciparum*. Quercitrin was already shown to exhibit moderate activity against *P. falciparum* (Ganesh et al., 2012; Liu et al., 2007; Murakami et al., 2001) although it was considered inactive in our study. If flavonoids are generally known for their moderate activity against various strains of *P. falciparum* (Ganesh et al., 2012; Torres-Mendoza et al., 2006), these molecules and their analogs are renowned for their ability to potentialize other molecules' activity such as artemisinin (Bilia et al., 2006; Bourdy et al., 2008; Ganesh et al., 2012; Rasoanaivo et al., 2011). They have also been considered as pharmacophore models in the search for molecules inhibiting fatty acids biosynthesis in the parasite (Gupta et al., 2010). In particular, several flavonoids including quercetin were found to exhibited strong activity (IC₅₀ 0.5-8 μM) towards three important

enzymes (FabG, FabZ and FabI) in a library's screening (Tasdemir et al., 2006) performed further to the discovery of luteolin-7-*O*-glucoside to be the first antimalarial natural product targeting the FabI enzyme of *P. falciparum* (Kirmizibekmez et al., 2004).

Overall, none of the compounds isolated here turned out as active as fraction D (the 5.5 μM for wayanin corresponds to 2.4 $\mu\text{g/ml}$). Hence, the excellent antiplasmodial activity of the fraction might not be due to one single component, but to a mixture of molecules, or to a synergistic action between them, as previously discussed by some authors for other traditional remedies (Deharo and Ginsburg, 2011; Houghton et al., 2007). A striking example is the pharmacodynamic synergy demonstrated *in vitro* between the *Cinchona* alkaloids found in *Cinchona* bark traditionally used for treatment of malaria (Rasoanaivo et al., 2011). Interestingly enough however, none of the isolated molecules was cytotoxic against VERO cells ($\text{IC}_{50} > 200 \mu\text{M}$).

It is worth mentioning that all isolated components except catechin are glycosylated quercetins. Because these compounds seem to be active as a whole, and because our previous work suggested an activity of the decoction per os (300 $\mu\text{L/day}$) on *Plasmodium berghei* NK65 infected Swiss male mice (Houël et al., 2015), it can be postulated that glycosylation plays a role in biological activity. As a matter of fact, it has been demonstrated that some quercetin monoglucosides are transferred more rapidly than the quercetin aglycone across the intestinal epithelium in rats (Gee et al., 2000). Further studies revealed that small intestine absorption in rats is determined by the type of sugar moiety, showing that within quercetin glycosylated derivatives, only Q3- and Q4'-glucosides were hydrolysed by β -glycosidases present in the small intestine, whereas other glycosides (Q3-galactoside, Q3-rhamnoside or Q3-arabinoside) were resistant to intestinal hydrolysis (Arts et al., 2004). Similar results were found in human subjects (Graefe et al., 2001; Hollman et al., 1997), indicating that glycosylated quercetines may really account for the antimalarial potential of the *P. acutangulum* decoction altogether.

Lastly, we investigated the *in vitro* anti-inflammatory activity of the ethyl acetate fraction (D) and the commercially available guaijaverin. For that, we tested their potential to inhibit the release of pro-inflammatory cytokines TNF- α , IL-1 β , IL-6 and IL-8, and the effect of these products on NO production. Indeed, the role of cytokines in malaria pathophysiology has been well documented. For example, it is now widely accepted that cytokines such as TNF and IL-1 are at the origin of systemic manifestations (anorexia, tiredness, aching joints, myalgia, headache, fever...) experienced by malaria patients (Burns et al., 2010; Clark et al., 2006; Lyke et al., 2004). Also, significantly elevated levels of IL-6 and IL-8 in patients' blood samples were associated with *Plasmodium falciparum* malaria severity among children and adults (Berg et al., 2014; Lyke et al., 2004). Diminished nitric oxide bioavailability may also contribute to the pathogenesis of severe malaria. For example, it was observed that children with severe malarial anemia showed evidence of pulmonary vasoconstriction, consistent with low NO bioavailability (Miller et al., 2013). The results of anti-inflammatory assays are presented in Figures 2 and 3. At 50 $\mu\text{g/mL}$, the ethyl acetate fraction (D) significantly inhibited IL-1 β secretion (-46 %) and NO production (-21 %), as previously observed for the decoction (-58% and -13% respectively; Houël et al., 2015). The effect of (D) and guaijaverin (4) on the secretion of other cytokines was not significant. No effect on cell viability was observed during the assays for inhibition of cytokine secretion (data not shown).

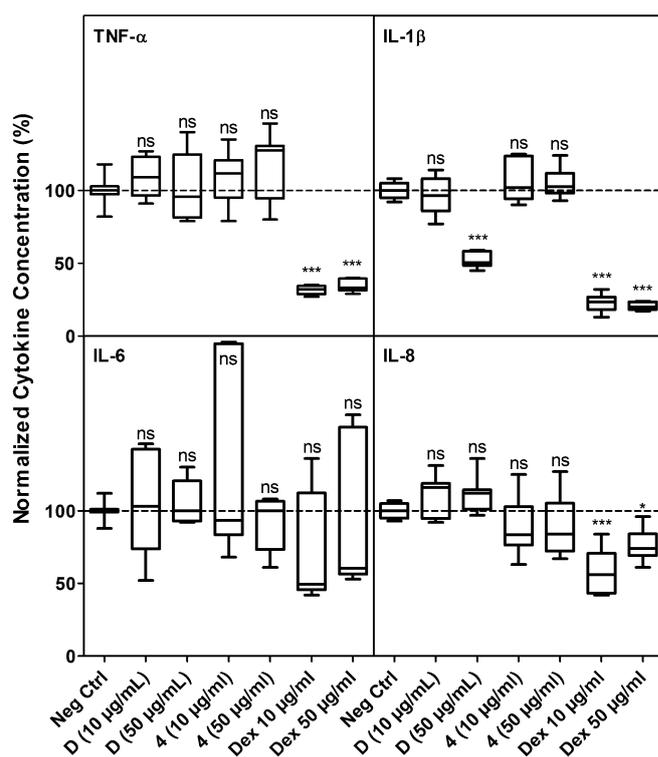


Fig. 2. Effect of *P. acutangulum* decoction ethyl acetate extract (D, 10 & 50 $\mu\text{g/mL}$), guaijaverin (**4**, 10 & 50 $\mu\text{g/mL}$) and dexamethasone (Dex, 10 & 50 $\mu\text{g/mL}$) on the secretion of TNF- α , IL-1 β , IL-6, and IL-8. Concentrations have been normalized relatively to the corresponding negative control of each experiment (Neg Ctrl). Significance levels when compared to negative control as calculated by one-way ANOVA followed by Tukey's multiple comparison test are indicated above the boxes: ns not significant; * $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$.

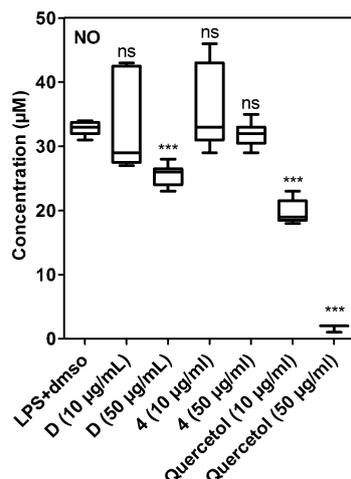


Fig. 3. Effect of *P. acutangulum* decoction ethyl acetate extract (D, 10 & 50 µg/mL), guaijaverin (**4**, 10 & 50 µg/mL) and quercetol (10 & 50 µg/mL) on NO secretion. Significance levels when compared to negative control (LPS+DMSO) as calculated by one-way ANOVA followed by Tukey's multiple comparison test are indicated above the boxes: ns not significant; * $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$.

These results obtained on pro-inflammatory cytokines are of particular interest compared to those obtained with the crude decoction. According to our previous results, the decoction inhibited the secretion of all tested cytokines (TNF α (-18%), IL-1 β (-58%), IL-6 (-32%) and IL-8 (-21%)), suggesting a mild anti-inflammatory activity of the remedy. However, the results obtained on the ethyl acetate fraction (D) and on guaijaverin (**4**) showed a reduced (cytokines spectrum) to in-existent anti-inflammatory activity compared to the decoction, with a significant anti-inflammatory effect on IL-1 β (-46%) for fraction (D) at 50 µg/ml only. Clearly, the crude decoction was more anti-inflammatory than the purified extract or compound. It is reasonable to postulate that other anti-inflammatory compounds were present in the remedy and not isolated during the bioguided fractionation, based on antiplasmodial activity.

4. Conclusion

Taking into account symptoms associated to the first intended pathology is crucial to understand the use and mode of action of traditional preparations. In many cases, interactions between compounds, i.e., pharmacodynamic synergies, complementary biological effects, contribute to its global biological activity (Bourdy et al., 2008; Rasoanaivo et al., 2011). This study adds new elements in this matter to our previous results on the decoction of *P. acutangulum*, a traditional antimalarial remedy of the Wayana Amerindians. Here, two antiplasmodial glycosylated flavonoids, guaijaverin and wayanin, were identified (IC₅₀ 6.9 μ M and 5.5 μ M respectively). However, none of the isolated compounds explained by itself the excellent activity of the ethyl acetate fraction, thus suggesting possible synergistic interactions between them. On the other hand, the ethyl acetate fraction (D) inhibited the secretion of IL-1 β and NO production only, and guaijaverin (**4**) was inactive. Hence, the crude decoction was clearly more anti-inflammatory than (D) or compound (**4**). Therefore, it appears that the interactions between the decoction's constituents clearly contribute to the activity of the remedy. Considering that neither the isolated molecules nor the decoction were cytotoxic, these results speak in favor of the use of *P. acutangulum* as antimalarial phytotherapeutic remedy.

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