

Evaluation of antibacterial activity of “Mangabarana” *Austroplenckia populnea* Reissek (Celastraceae)

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RESUMO: “Avaliação da atividade antibacteriana de “Mangabarana” *Austroplenckia populnea* Reissek (Celastraceae)”. *Austroplenckia populnea* (Mangabarana) é popularmente utilizada em Minas Gerais, Brasil, para o tratamento de disenterias. A ela também são atribuídas propriedades antitumoral e antiúlcera. Extratos de partes desta planta obtidos com solventes de diferentes polaridades e triterpenos pentacíclicos (TTPCs) isolados destes, por métodos fitoquímicos foram submetidos a testes de atividade antibacteriana. Os resultados mostraram a existência desta atividade e abriram perspectivas para a continuidade dos estudos com outros compostos orgânicos isolados desta planta.

Unitermos: *Austroplenckia populnea*, Celastraceae, atividade antibacteriana, triterpenos pentacíclicos.

ABSTRACT: *Austroplenckia populnea* (Mangabarana) is popularly used by people from Minas Gerais, Brazil for dysenteries diseases treatment. Antitumor and antiulcer activities were also attributed to this plant. Extracts obtained using solvents of different polarities and pentacyclic triterpenes (PCTTs) isolated from these extracts through phytochemical methods were submitted to antibacterial assays. The results showed the existence of this activity and open perspectives for news studies with other organic compounds isolated from this plant.

Keywords: *Austroplenckia populnea*, Celastraceae, antibacterial activity, pentacyclic triterpenes.

INTRODUCTION

The indiscriminate use of antibiotics has been caused bacterial resistance, allergic sensibility, blocking of natural fermentation processes and other adverse biological effects. This fact induces the development of new antibiotics with a good efficacy and with low incidence of adverse effects.

Ethnobotanical and ethnopharmacological studies have shown that products obtained from plant can present healing action related to specific pathologies. Consequently, a great variety of medicinal plants have been investigated to evaluate and validate their real biological activity (Cardoso and Santos, 1948; Kinghorn and Balandrin, 1993; Rao, 1996; Gnan and Demello, 1999; Hernandez et al., 2000; Deena et al., 2000; Cardoso-Lopes et al., 2008). The use of plants and their derivatives for different diseases occurs since the beginning of the humanity (Antunes et al.,

2006). With the advances of pharmaceutical industries, mainly after World War II, the use of herbal medicines diminishes in function of the dissemination idea of its inefficiency (Antunes et al., 2006). However, the efficacy and trustworthiness in medicinal plant products have demonstrated the real value of these materials. Brazil have a great diversity of native plants (Mors, et al., 2000, Barbosa-Filho, et al., 2005, Brandão, et al., 2006; Quintans-Júnior et al., 2008), but its medicinal properties had had not been adequately studied yet. The scientific study of plants used in folk medicine is essential to lower the adverse effect risk for populations (Agra et al., 2008).

Frequent deforestation and fires are threatening a substantial number of native plant species with extinction. The *Austroplenckia populnea* Reissek, specie of the Celastraceae family is among of them. This plant, commonly known as “Mangabarana”, “Marmelo-do-campo” or “Mangabeira-brava”, can be found in a

vast tropical ecoregion known as "Cerrado" mainly in the State of Minas Gerais, Brazil, and has been of great interest to researchers because of its chemical constituents, larvicidal effect, and molluscicidal effect (Vichniewski et al., 1984).

Other biological activities were also attributed to *Austroplenckia populnea* including antidysenteric action (Correa, 1969), antitumoral action (Monache, 1972) and antirheumatic action (Gonzalez et al., 1982). In relation to the dysenteries, it is important to point out that epidemiological studies indicate high *E. coli* prevalence, mainly in people who live in Brazilian cities peripheries (Tavares-Dias & Grandini, 1999), and, are exactly these people that use *A. populnea* and other medicinal plants more frequently. Previous studies demonstrate antibacterial activity of the following pentacyclic triterpenes (PCTTs) isolated from *Austroplenckia populnea*: abruslactone-A, 3-epiabraslactone-A, populnicic acid, polpuninic acid and katononic acid (Vieira Filho et al., 1999). Treatment of adult Wistar rats with hexane extract obtained from dried *Austroplenckia populnea* leaves showed an antispermatogetic action characterized by a significant decrease of the epididymis spermatozoa number (Mazaro et al., 2000). Anti-*Trypanosoma cruzi* effect presented by the PCTTs epikatonic acid and 20-hydroxy-tingenone isolated from *A. populnea* was also related (Duarte et al., 2002).

The isolation and identification of other PCTTs, sesquiterpenes and other chemical constituents from this specie have been reported by several authors (Monache, 1972; Vichniewski et al., 1984; Souza et al., 1988; Souza et al., 1990; Silva et al., 2002; Vieira Filho et al., 1999; Vieira Filho et al., 2000; Vieira Filho et al., 2001; Cotta et al., 1990 a, b).

The objectives of this work involve the antibacterial evaluation of the *A. Populnea* extracts obtained using solvents of different polarities and the PCTTs isolated from them, followed by determination of the minimum inhibitory concentration (MIC), and by the minimum bactericidal concentration (MBC). Because of the previous antibacterial effect identified for some PCTTs (Vieira Filho et al., 1999) and the popular use of the decoct from *A. Populnea* branches as an antidysenteric (Correa, 1969), the PCTTs friedelin (1), 3 β -friedelinol (2), 28-hydroxyfriedelin (canofilol) (3), populnicic acid (4), katononic acid (6), epikatonic acid (8), pristimerin (10), abruslactone A (11) and α -amirin (12) isolated from *Austroplenckia populnea* and the transformation products: methyl populnonate (5), methyl katotonate (7) and methyl epikatotate (9) (Figure 1) were tested to determine the *in vitro* bacterial activity.

MATERIAL AND METHODS

Plant

A. Populnea was collected in the proximities of Miguelão Lake, at Nova Lima city region, Minas Gerais, Brazil. This plant was identified by Prof. Wagner Pedersoli (*in memoriam*). A sample of the collected material was compared and identified with a voucher specimen (N^o 10473) deposited at the Herbarium of the Natural History Museum of Universidade Federal de Minas Gerais, Belo Horizonte, Minas Gerais, Brazil.

Extracts obtainment

Leaves, branches, (bark and heartwood) and root were separated and dried over kraft paper at room temperature. After fragmentation on a mill, each part was submitted to continuous extraction process in a Soxhlet apparatus using hexane, chloroform, ethyl acetate, methanol and *n*-butanol as solvent extractor.

Afterwards, each extract was submitted column chromatography (CC), flash-CC, high-pressure liquid chromatography (HPLC) and other phytochemical methods. By these processes it was possible to isolate the PCTTs: friedelin (1), 3 β -friedelinol (2), 28-hydroxyfriedelin (canofilol) (3), populnicic acid (4), katononic acid (6), epikatonic acid (8), pristimerin (10), abruslactone A (11) and α -amirin (12). The PCTTs (4), (6) and (8) were esterified to obtain methyl populnonate (5), methyl katotonate (7) and methyl epikatotate (9), respectively. Infrared (IR), mass spectrometry (MS) and nuclear magnetic resonance (¹H and ¹³C NMR including 2D experiments) were used to elucidate the chemical structure of the compounds (Souza et al., 1990; Vieira Filho et al., 1999; Vieira Filho et al., 2000; Vieira Filho et al., 2001; Cotta et al., 1990; Silva et al., 2002; Zanon et al., 2008).

Antibacterial activity preliminary assays

Chloroform extract obtained from the bark (ChE) and from the heartwood of the branches (HChE) were submitted to qualitative microbiological tests against wild-type strains of *Staphylococcus aureus* and *Escherichia coli*, both sensitive to vancomycin through Bawer & Kirby disc diffusion method (Bawer and Kirby, 1966) (Table 2).

Minimum inhibitory concentration (MIC) of the extracts

Through microdilution technique, four samples (2 mg/mL initial concentration) of *A. populnea* extracts were submitted to microbiological assays to determine the growth minimum inhibitory concentration (MIC) of *Staphylococcus aureus* (ATCC 21027), *Proteus*

vulgaris (ATCC 13315), *Pseudomonas aeruginosa* (ATCC 15442), *Escherichia coli* (ATCC 10536) and *Staphylococcus aureus* (MARSA), microorganisms commonly found in human infectious diseases. All experiments were carried out in triplicate and the average rate found was the rate taken in account (Table 3).

Evaluation of minimum bactericidal concentration of the extracts

After the MIC have been established, aliquots of the extracts in which growth inhibition was observed were inoculated onto Muller-Hinton agar and incubated at 37 °C, during 24 hours. Each experiment was performed in triplicate and a non-growth in the cultivation conditions showed the minimum bactericidal concentration of the extract (Table 4).

Evaluation of the antibacterial activity of the constituents isolated from *A. populnea*

The PCTTs 1 to 12 (Figure 1) were respectively

dissolved in chloroform. Each chloroform solution was impregnated in sterile Whatmann paper discs (6.0 mm diameter) to obtain five different PCTT concentrations, after solvent evaporation. Dried discs containing PCTT were deposited on the surface of agar Müller-Hinton previously inoculated with *Salmonella typhimurium*, *Staphylococcus aureus* ATCC 25923, *Escherichia coli* ATCC 25922 or *Shigella sonnei* ATCC 11060, in according to Bawer & Kirby method (Bawer and Kirby, 1966). Oxacillin, tetracycline, amoxicillin, clindamycin and chloramphenicol were used as internal standards.

The analysis of the results was done after 24 hours of incubation at 37 °C. The samples that produced inhibition halos equal or superior to 7 mm were considered active. Each experiment was developed using five Petri plates. Discs were soaked individually with each one of the solvents that were used for the incorporation of the compounds to be tested. After the process of drying, these discs also were submitted to growth inhibition tests to verify possible interferences caused by the presence of residual nonvolatile compound. For these discs, any growth inhibition halo was detected, which demonstrated the purity of the solvents used and the efficiency of the evaporation process.

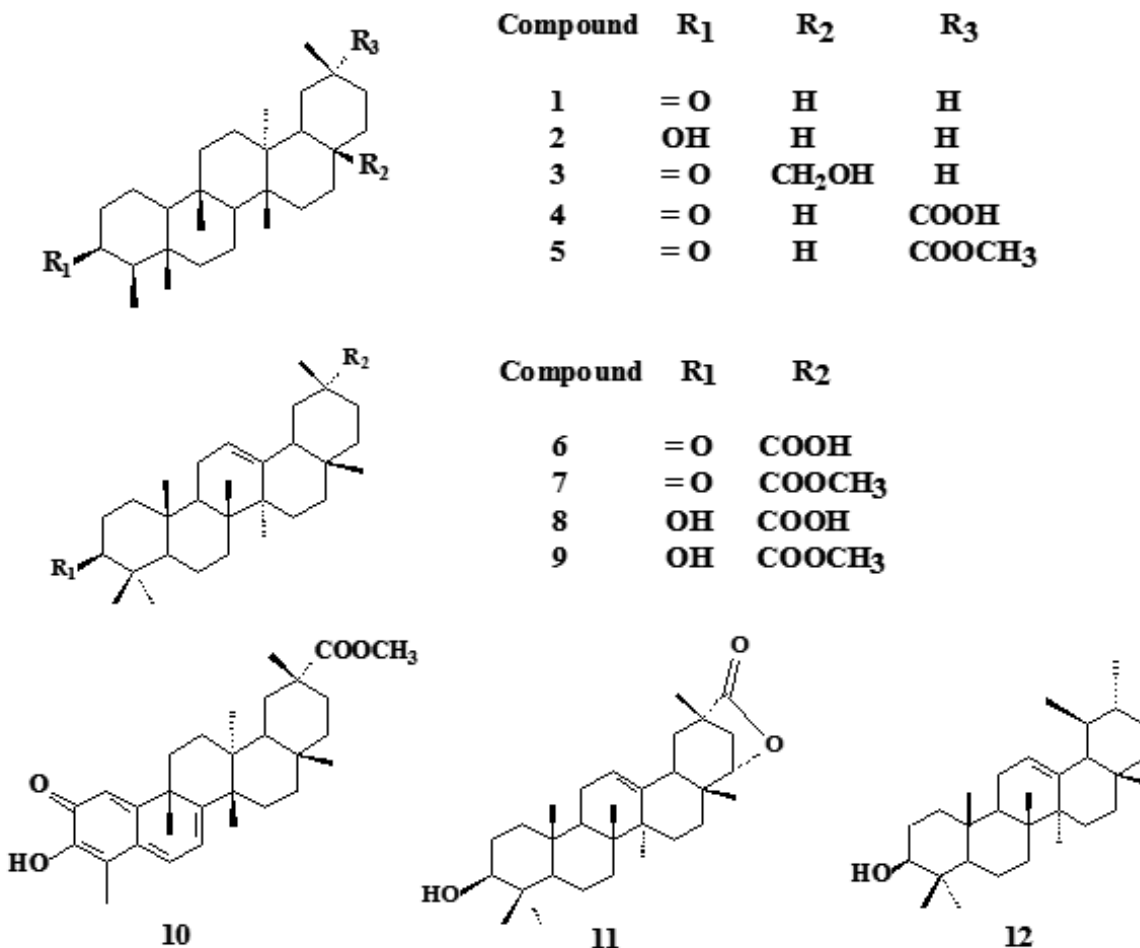


Figure 1. Pentacyclic triterpenes from *A. populnea* submitted to antibacterial activity.

Table 1. Parts of *Austroplenckia populnea* and respective extracts used in the antibacterial experiments.

Parts of <i>Austroplenckia populnea</i>	Extracts submitted to antibacterial assays
Leaves	Hexanic (LHE) Ethanol (LEE)
Branches	Chloroformic of bark (ChE) Chloroformic of heartwood (HChE) Ethylacetic (BEaE) Butanolic (BBE) Methanolic (BME)
Roots	Methanolic (RME)

Table 2. Results of the preliminary antibacterial activity tests realized with chloroform extract obtained from bark and heartwood of *A. populnea* branches.

Material tested	Microorganism	
	<i>S. aureus</i>	<i>E. coli</i>
ChE	+	-
HChE	+	-
Vancomycin	+	+

+ = Showed growth inhibition halo.

- = Don't showed growth inhibition halo.

Table 3. Minimum inhibitory concentration (MIC) of the extracts (mg/mL) obtained from *A. populnea*.

Bacteria submitted to tests	Parts of <i>A. populnea</i>			
	Branches bark			Roots
	BBE	BEaE	BME	RME
<i>Escherichia coli</i> (ATCC 10536)	nd	nd	nd	nd
<i>Pseudomonas aeruginosa</i> (ATCC 15442)	nd	nd	2	1
<i>Staphylococcus aureus</i> (MARSA)	nd	nd	2	nd
<i>Staphylococcus aureus</i> (ATCC 21027)	0,5	0,5	0,5	nd
<i>Proteus vulgaris</i> (ATCC 13315)	nd	nd	nd	nd

E = extract; B = branches and butanolic; Ea = ethylacetic; M = methanolic

nd = inhibitory effect not detected in the experimental conditions.

Table 4. Minimum bactericidal concentration (MBC) of the extracts (mg/mL) obtained from *A. populnea*.

Bacteria submitted to tests	Extracts of <i>A. populnea</i>			
	Branches bark			Roots
	BBE	BEaE	BME	RME
<i>Escherichia coli</i> (ATCC 10536)	nd	nd	nd	nd
<i>Pseudomonas aeruginosa</i> (ATCC 15442)	nd	nd	nd	nd
<i>Staphylococcus aureus</i> (MARSA)	nd	nd	nd	nd
<i>Staphylococcus aureus</i> (ATCC 21027)	nd	1	nd	nd
<i>Proteus vulgaris</i> (ATCC 13315)	nd	nd	nd	nd

E = extract; B = butanolic; Ac = ethylacetic; M = methanolic

nd = inhibitory effect not detected in the experimental conditions.

RESULTS AND DISCUSSION

The antibacterial activity of five pentacyclic triterpenes isolated from *Austroplenckia populnea* extracts was previously reported in relation to *Escherichia coli*, *Streptococcus faecalis*, *Staphylococcus aureus*, *Pseudomonas aeruginosa*, *Staphylococcus epidermidis*

and *Salmonella* sp. (Vieira Filho et al., 1999). The results obtained in this work demonstrated that among the studied extracts, four of them, with polar character, also showed antibacterial activity. The butanol (BBE) and ethyl acetate (BEaE) extracts obtained from *Austroplenckia populnea* branches bark showed activity against *S. aureus* (ATCC 21027). The methanol extract from branches bark (BME) also showed activity against

Table 5. Antibacterial activity of pentacyclic triterpenes isolated from *A. populnea*.

Compound	<i>S. typhimurium</i>		<i>S. aureus</i> (ATCC 25923)		<i>E. coli</i> (ATCC 259222)		<i>S. sonnei</i> (ATCC 11060)	
	C	H	C	H	C	H	C	H
1	650	nd	970	nd	1080	nd	170	7
2	700	nd	930	nd	1030	nd	230	8
3	930	nd	520	nd	2250	8	180	9
4	860	nd	2080	nd	810	nd	190	7
5	600	nd	1620	nd	250	9	270	8
6	-	-	1050	nd	930	nd	1250	8
7	-	-	570	nd	1430	nd	240	8
8	630	nd	430	nd	160	7	500	nd
9	950	nd	1360	nd	950	nd	250	7
10	1010	nd	490	nd	200	10	170	8
11	650	nd	440	nd	150	10	140	7
12	-	-	810	nd	860	8	230	9

C = µg of compound;

H = inhibition halo (mm) of bacteria growth;

nd = not detected and

- = not submitted to the experiments.

None inhibition halo was observed for the solvents used during dilution process.

Inhibition halo observed for standards: Amoxicillin: 30 µg (10 mm); Clindamycin: 28 µg (2 mm); Cloramphenicol: 30 µg (mm); Oxacyclin: 29 µg (1 mm); Tetracyclin: 24 µg (30 mm).

S. aureus (MARSA) and *S. aureus* (ATCC 21027) and inhibited the growth of *P. aeruginosa* (ATCC 15442). The growth of this microorganism also was inhibited by root methanol extract (RME) (Table 3).

The corroboration of antibacterial activity of the vegetable extracts evaluated in this work suggests more accurate phytochemical studies to quantify and identify other biologically active chemical substances, which can be pointed as sources of new antibiotics.

None of the PCTTs (Table 5) showed activity against *S. typhimurium* and *S. aureus*. It can be observed that despite the fact that the chloroform extract obtained from the bark and from the heartwood of the branches showed activity against *S. aureus* (Table 3). When abruslactone A (**11**) a PCTT isolated from this extract (200 µg/disc) was submitted to a test by plate cavity method, it showed activity against *S. aureus* ATCC 25923 (Vieira Filho et al., 1999).

Salaspermic acid was also isolated from the chloroform extract of the *Autroplenckia populnea* heartwood of the branches. In according to the literature, this PCTT showed anti-HIV activity (Chen et al., 1992). It allows the premise that the activity of this extract should be due to the presence of this compound, that

was not tested in this work, or of other substance found in the plant. The data presented in Table 5, show that all the PCTTs showed activity against *Shigella sonnei*.

For *E. coli* it was observed that compounds **3**, **5**, **8**, **10**, **11** and **12** showed low activity (Table 5). Compound **10** showed significant activity against *E. coli*. On the other hand, this compound was inactive when tested against *E. coli* ATCC st330/84. It was observed that the strains tested are different, which can justify this difference in antibacterial action answer (Vieira Filho et al., 1999).

The results stimulate the accomplishment of new experiments where new tests must be performed, using other pathogenic bacteria, others PCTTs and respective derivatives in order to establish chemical structure *versus* antibacterial activity correlations.

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