

QUANTIFICATION OF BERGENIN, ANTIOXIDANT ACTIVITY AND NITRIC OXIDE INHIBITION FROM BARK, LEAF AND TWIG OF *Endopleura uchi*Magno P. Muniz^{a,*}, Sergio M. Nunomura^c, Emerson S. Lima^b, Arleilson S. Lima^b, Patrícia D. O. de Almeida^b and Rita C. S. Nunomura^a^aDepartamento de Química, Instituto de Ciências Exatas, Universidade Federal do Amazonas, 69077-000 Manaus – AM, Brasil^bFaculdade de Ciências Farmacêuticas, Universidade Federal do Amazonas, 69080-900 Manaus – AM, Brasil^cCoordenação de Pesquisa de Tecnologia e Inovação, Instituto Nacional de Pesquisas da Amazônia, 69067-375 Manaus – AM, Brasil

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Phytochemical investigation of *Endopleura uchi* led to the isolation of a gallic acid derivate, known as bergenin (2) and friedelin (1), a pentacyclic triterpene. The present work also reports the bergenin quantification in different *Endopleura uchi* bark, twig and leaves extracts. Our findings showed the highest bergenin concentration in bark methanol extracts (4.75%) and the lowest concentration in twig aqueous extracts (1.89%). Phenolics quantification by Folin-Ciocalteu, revealed phenolic compounds level values from 16.69 to 43.02 mg GAE g⁻¹ dry extract. The ferric reducing antioxidant activity ranged from 230.43 to 567.89 mmol Fe⁺² g⁻¹ dry extract. DPPH IC₅₀ free radicals showed to range from 12.04 to 24.20 µg mL⁻¹. Leaves' chloroform fraction exhibited the highest nitric oxide inhibiting activity bearing IC₅₀ = 3.2 µg mL⁻¹. Results show this species to hold significant bergenin concentrations as well as phenolic and anti-inflammatory compounds in all extracts suggesting it bear therapeutic potential.

Keywords: Humiriaceae; liquid chromatography; anti-inflammatory agents; natural products; DPPH.

INTRODUCTION

Species *Endopleura uchi* (Huber) Cuatrec. [sin. *Sacoglottis uchi* Huber] belongs to the Humiriaceae family and is typically wild in primary forests growing on non-flooded land, widely dispersed throughout the Amazon Basin. It is mainly found in the state of Amazonas, and popularly known as “uxi-amarelo”, “uxi-liso”, “uxi-pucu” and “pururu”.^{1,2} *Endopleura uchi* barks have been sold in open fairs, markets, nationwide online shops and commercial natural products outlets and used in the form of tea as a potent anti-inflammatory medicine against tumors, myomas, uterine infections, fibroids and polycystic ovaries.³ Friedelin, a pentacyclic triterpene, and a 4-O-methyl gallic acid C-glycoside derivate known as bergenin, both bearing anti-inflammatory properties (Figure 1).^{4,5} Bergenin shows activity against the growth of yeast for different *Candida* species,⁶ inhibitory activity against COX-2,⁷ which is an enzyme mediating inflammatory processes in the body, antioxidant activity,⁸ hepatoprotective,⁹ neuroprotective,¹⁰ anti-HIV¹¹ and antifungal activities against complete inhibition of germination of spores of *Fusarium udum* and *Erysiphe pisi*.¹² Earlier studies have reported the isolation of bergenin from the aqueous extract of the bark of *Endopleura uchi* of about 3.19%,⁷ in fruits and in other species of the family Humiriaceae as *Humiria balsamifera* Aubl. and *Sacoglottis gabonensis*.^{13,14} The several bergenin biological activities have led this work toward the bergenin quantification in different *Endopleura uchi* leaves, twigs and bark extracts through RP-HPLC to find out the active principle concentration in different parts of the species.

Considering the sp., *Endopleura uchi* popular use as anti-inflammatory medicine and on account of this inflammatory process being associated with oxidative stress, extracts and fractions antioxidant activity and free radicals scavenging ability, were evaluated through the FRAP (ferric reducing antioxidant power) and DPPH (2,2-diphenyl-1-picrylhydrazyl) methodologies, respectively. Furthermore, the content of phenolic compounds, which are known for their ability to prevent

oxidative stress that yields reactive oxygen (ROS) and nitrogen (RNS) species, causing physiological alterations and even cell death, have been determined. ROS and RNS can be generated by external factors or, within the organism, by neutrophils and macrophages during the inflammatory action.¹⁵ A method was employed to estimate *Endopleura uchi* extracts ability to suppress nitric oxide (NO), an inflammation mediator, production in LPS (lipopolysaccharide)-induced J774 murine macrophages cultures, for assessing their anti-inflammatory aspect.¹⁶

Earlier studies in the literature on the species *Endopleura uchi* have just been targeted on its fruits and barks. This paper also discusses unpublished studies addressing *Endopleura uchi* twigs and leaves, which have presented relevant aspects that may contribute to the development of new drugs or herbal medicines. The scientific research of this species through understanding their chemical constituents and confirmed biological activity can direct the popular use to another part of the plant since the inadequate extraction of tree bark can result in the death of the species.¹⁷

EXPERIMENTAL SECTION**Solvents, reagents and chemicals**

All solvents used were of HPLC grade by J. T. Baker (Xalostoc, Edo. Mexico). Water was purified using a Milli-Q system (Millipore®, Bedford, USA). Folin-Ciocalteu, 2,2-diphenyl-1-picrylhydrazyl (DPPH), quercetin; gallic acid; (Sigma-Aldrich, Inc, St. Louis, Missouri, USA); sodium bicarbonate, ferrous sulphate heptahydrate and ferric chloride III (Synth, Diadema, SP, Brazil); 2,4,6-tri(2-pyridyl)-s-triazine (TPTZ) (Fluka Chemie GmbH, Buchs, CH); hydrochloric acid and trifluoroacetic acid (Tedia, Fairfield, OH, USA); ion (II) chloride tetrahydrate (Merck, Darmstadt, GmbH).

Plant material

Leaves, twigs and barks of the species *Endopleura uchi* identified by botanist G. T. Prance and deposited in the herbarium of the INPA

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under the number 190992 were collected at Adolpho Ducke Reserve (INPA) located at Km 23 from Manaus (AM) during the dry period on October 11, 2010 and the rainy season on April 16, 2011. This project has been registered with the National Genetic Resource Management and Associated Traditional Knowledge Management System (SisGen – A04356A).

General experimental procedures

The NMR spectra were obtained on a Unity Inova spectrometer model 500 MHz (^1H NMR) and 125 MHz (^{13}C NMR) in CDCl_3 or CD_3OD (Cambridge Isotope Laboratories Inc.). HR-ESI-MS and HR-APCI-MS analyses were measured in the positive ion mode (Bruker Daltonics, model II - TOF, APCI). Column chromatography was performed on silica LC60A (70 - 200 μm , Grace) and silica 60 (40 - 63 μm , Merck). Melting points (uncorrected) were recorded on a FISATOM 430 D Melting Point Apparatus.

Extraction

Leaves (F), twigs (G) and barks (C) were ground and dried. Aqueous extracts (EA) were prepared by decoction of 30 g of dried plant material in 500 mL boiling distilled water for 5 min as the popular use. Methanol extracts (EM) were prepared by soxhlet extraction, the solvent replaced every 6 hours, totaling 18 h of extraction. Ethanol extract (EE) prepared by using 40 g of dried plant material in 400 mL of ethanol three times in ultrasound for 15 min each, followed by maceration for one day.

Friedelin and bergenin isolation

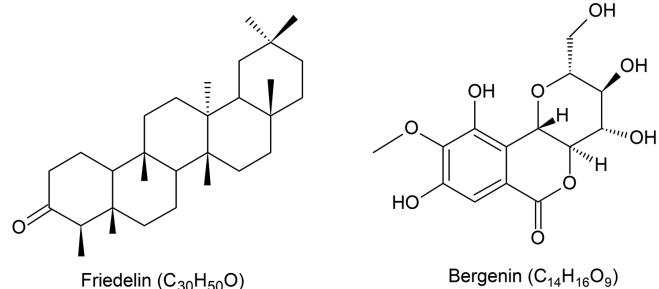


Figure 1. Structures of friedelin and bergenin

Friedelin was isolated from twigs and leaves hexane fractions. Bergenin was isolated from barks, twigs and leaves ethyl acetate fractions. Compounds were identified by ^1H , ^{13}C NMR, HOMOCOSY, HSQC, HMBC and MS spectrometry. Twigs (GEM-1, 9.6 g), leaves (FEM-1, 10.0 g) and barks (CEM-1, 13.8 g) methanol extracts prepared in liquid-liquid extraction solubilized in methanol/water (9:1) and partitioned with hexane, chloroform and ethyl acetate, successively. The hexane phase of twigs (GEM-1-Hex, 260.0 mg) was concentrated and then recrystallized with acetone that resulted in impure crystals. These crystals were purified by chromatographic column in silica gel 60 (40-63 μm), using hexane/chloroform gradient (7:3, 1:1, 3:7) and chloroform (100%) as mobile phase to yield friedelin (**1**) (5.0 mg). The same procedure was performed with leaves hexane fraction (FEM-1-Hex, 500.0 mg) to yield friedelin (**1**) (17.5 mg). The ethyl acetate phase (GEM-1-EtOAc, 1.87 g) concentrated and washed with acetone that resulted in a crystalline solid impure. This material was separated in a chromatographic column using silica gel LC60A (70-200 μm) and chloroform/ethanol (7:3) isocratic as the mobile phase to yield bergenin (**2**) (550.0 mg). The same procedure was both carried out with leaves ethyl acetate

fraction (FEM-1-EtOAc, 1.7 g) to yield bergenin (**2**) (540.0 mg) and barks ethyl acetate fraction (CEM-1-EtOAc, 2.5 g) to yield bergenin (**2**) (840.0 mg).

Friedelin¹⁸ (**1**): White crystalline solid; m.p: 252-255 °C; ^1H δ_{H} (500 MHz, CDCl_3): 1.97 (m, H-1a), 1.69 (1H, dd, $J = 5.0$ and 13.0 Hz, H-1b), 2.39 (1H, ddd, $J = 2.0, 5.0, 14.0$ Hz, H-2a), 2.30 (1H, dd, $J = 7.5, 13.5$ Hz, H-2b), 2.25 (1H, q, $J = 6.5$ Hz, H-4), 1.76 (1H, dd, $J = 3.0, 16.0$ Hz, H-6a), 1.28 (m, H-6b), 1.60 (m, H-7a), 1.48 (m, H-7b), 1.45 (m, H-8a), 1.40 (m, H-8b), 1.56 (m, H-10), 1.39 (m, H-11), 1.36 (m, H-12), 1.48 (m, H-15a), 1.28 (m, H-15b), 1.55 (m, H-16a), 1.37 (m, H-16b), 1.57 (m, H-18), 1.38 (m, H-19), 1.50 (m, H-21), 1.49 (m, H-22), 0.88 (3H, d, $J = 6.5$ Hz, H-23), 0.73 (3H, s, H-24), 0.89 (3H, s, H-25), 1.01 (3H, s, H-26), 1.05 (3H, s, H-27), 1.18 (3H, s, H-28), 1.00 (3H, s, H-29) and 0.96 (3H, s, H-30); ^{13}C δ_{C} (125 MHz, CDCl_3): 22.6 (C-1), 41.8 (C-2), 213.5 (C-3), 58.5 (C-4), 42.4 (C-5), 41.6 (C-6), 18.5 (C-7), 53.6 (C-8), 37.8 (C-9), 59.8 (C-10), 35.9 (C-11), 30.8 (C-12), 40.0 (C-13), 38.6 (C-14), 33.1 (C-15), 36.3 (C-16), 30.3 (C-17), 43.1 (C-18), 35.7 (C-19), 28.5 (C-20), 32.7 (C-21), 39.6 (C-22), 7.1 (C-23), 15.0 (C-24), 18.2 (C-25), 20.6 (C-26), 19.0 (C-27), 32.4 (C-28), 32.1 (C-29) and 35.3 (C-30); HR-APCI-TOF-MS: 427.3942 $[\text{M} + \text{H}]^+$, $\text{C}_{30}\text{H}_{50}\text{O}^+$; calc. 427.3934.

Bergenin^{5,7} (**2**): White crystalline solid; m.p: 150-152 °C¹⁹; ^1H δ_{H} (500 MHz, CD_3OD): 7.08 (1H, s, H-4), 4.95 (1H, d, $J = 10$ Hz, H-9), 3.69 (1H, m, H-11), 3.43 (1H, dd, $J = 8.5, 8.5$ Hz, H-12), 3.81 (1H, dd, $J = 8.8, 9.5$ Hz, H-13), 4.06 (1H, dd, $J = 9.5, 10.0$ Hz, H-14), 3.91 (3H, s, H-15), 3.68 (1H, m, H-16), 4.02 (1H, dd, $J = 3.0, 13.0$ Hz, H-16). ^{13}C δ_{C} (125 MHz, CD_3OD): 164.6 (C-2), 118.2 (C-3), 109.9 (C-4), 151.1 (C-5), 141.1 (C-6), 148.2 (C-7), 116.1 (C-8), 73.1 (C-9), 81.9 (C-11), 70.7 (C-12), 74.4 (C-13), 80.2 (C-14), 59.8 (C-15), 61.5 (C-16). HR-ESI-TOF-MS: 329.0866 $[\text{M} + \text{H}]^+$, $\text{C}_{14}\text{H}_{16}\text{O}_9$; calc. 329.0867.

Total phenolics determination

The Folin-Ciocalteu²⁰ method was used to quantify the total phenolic content in extracts and fractions. Two-hundred (200) μL of the sample (1 mg mL^{-1} in methanol) were mixed with 1.5 mL of aqueous solution of Folin-Ciocalteu (1:10). Following 5 min, 1.5 mL of sodium bicarbonate (Na_2CO_3 , 6%) solution added. Then, after it incubated at room temperature for 90 min, the absorbance was measured at 725 nm in a UV/Vis spectrophotometer (Femto 800XI) and Milli-Q water used as blank. The analyses were performed in triplicate and the total phenolic content mean was expressed in milligrams of gallic acid equivalents (GAE) per gram of dry extract. Gallic acid analytical curve was constructed in the range of 15.6-250 $\mu\text{g mL}^{-1}$ ($y = 4.9604x - 0.0428$, $R^2 = 0.9986$).

DPPH free radical scavenging through antioxidant activity

DPPH free radical scavenging capacity was measured by the photometric method described by Mensor.²¹ For each fraction, extract or standard (quercetin) six different dilutions were prepared (4-57 mg mL^{-1} in methanol, 2.5 mL) and 1 mL of DPPH (0.2 mg mL^{-1}) added. After 30 min, in the dark at room temperature, a reading was taken at 518 nm in a spectrophotometer (Femto 800XI). Blank was obtained by the higher concentration samples and 2.5 mL methanol in 1 mL of DPPH as a negative control. The antioxidant activity was calculated as follows:

$$\text{AA}\% = 100 - [(\text{Abs sample} - \text{Abs blank}) / \text{Abs control}] \times 100 \quad (1)$$

where, Abs sample = average absorbance of the sample, Abs blank = absorbance of the higher concentration sample, Abs control = average absorbance of the negative control. The 50% inhibitory capacity (IC_{50}) was calculated from the slope with antioxidant activity versus sample concentration, which analyses were performed in triplicate.

Antioxidant activity by FRAP method (ferric reducing antioxidant power)

The FRAP assay was performed as described by Luximon-Ramman.²² The FRAP reagent was prepared fresh by mixing in 10 mM TPTZ (2,4,6-tripiridyl-triazine), 20 mM of iron (III) chloride solution and sodium acetate buffer (0.25 M, pH 3.5) at a ratio of 1:1:10. Then, 100 μ L of the sample in methanol (1 mg mL⁻¹) were added to 300 μ L of Milli-Q water and 3.0 mL of FRAP reagent. The absorbance was measured at 593 nm after being incubated for 4 minutes. Findings were expressed as Fe⁺² μ mol g⁻¹ dry weight of plant material and the analytical curve was constructed using FeSO₄ (62.5 – 2000 μ mol L⁻¹; $y = 0.0007x - 0.0428$, $R^2 = 0.9999$).

Cell culture and treatments

Murine macrophages J774 cells (donated from the cell bank of the Rio de Janeiro) were grown at 37 °C in an incubator containing 5% CO₂ in RPMI-1640 medium included fetal bovine serum (10%, FBS), streptomycin (50 μ g mL⁻¹, Invitrogen) and penicillin (50 U mL⁻¹). The used LPS stock solution (1 mg mL⁻¹) was prepared in sterile PBS [137.0 mM NaCl, 2.7 mM KCl, 4.3 mM Na₂HPO₄·7H₂O, 1.4 mM KH₂PO₄, pH 7.4]. LPS is used to stimulate iNO in cells. Samples were added to the culture medium before LPS treatment.²³

NO production assay

After pre-incubation of cells (1x10⁶ cells mL⁻¹) with differently concentrated samples (2.5 to 25.0 μ g mL⁻¹) for 2 h, cells were incubated for 24 hours either with or without LPS (1 mg mL⁻¹) at 37 °C in an incubator containing 5% CO₂. The same procedure was carried out for negative control and blank. Then, nitric oxide production (NO₂) in the culture supernatant was determined using Griess reagent²³ in triplicate. Samples absorbance was measured at 560 nm using a microplate reader (DTX 800, Beckman). Sodium nitrite was used as a standard to calculate nitrite concentrations.

Bergenin quantification

Endopleura uchi extracts bergenin quantification was performed using a method previously described by Nunomura *et al.*,⁷ with some modifications. Analyses were performed by Liquid Chromatography using a SHIMADZU Prominence LC-20A, equipped by LC-10AT Vp quaternary pump, automatic injector SIL-20A and DAD detector (DAD) SPD-M20A and data process using a LC-Solution software. The separation was performed in a Merck LiChrospher 100 RP-18e (250 mm x 4.0 mm, 5 μ m and 120 Å) column using, methanol

as solvent B (20%) and THF solution in water (pH 2.0) as solvent A (80%), as mobile phase during 30 minutes. The mobile phase was degassed before analyses. The flow rate was 1.0 mL min⁻¹ and detection on 215, 254 and 272 nm. The column temperature was 25 °C and the injection volume 50 μ L. The quantification was performed by integrating peaks using bergenin, as an external standard, diluted in methanol at 6.25, 12.5, 25.0, 50.0, 100.0 and 200.0 μ g mL⁻¹ concentrations, analyzed intra-day and inter-day ($R^2 > 0.995$), in duplicate. Extracts were submitted to SPE using C18 cartridge (Sep-Pak) and analyzed at 250 μ g mL⁻¹ in duplicate. Extract sample peaks identification was performed by comparison with bergenin by retention time and λ_{max} in nm. Extract and standard samples were filtered on 0.2 μ m Millipore membrane.

RESULTS AND DISCUSSION

Bergenin quantification

Bergenin (2) was quantified in leaves, twigs and barks from *Endopleura uchi* by High Performance Liquid Chromatography-Diode Array Detector (HPLC-DAD) on aqueous and methanol extracts using Nunomura *et al.*⁷ method and some modifications. Bergenin standard solutions (6.25-200 μ g mL⁻¹) were analyzed repeatedly intraday and interday and resulted in an excellent linear regression slope ($R^2 > 0.995$, CV < 5%) (Supplementary Material Table 1S and Figure 12S). Detection and quantification limits (DL) and (QL) showed a high detection level in addition to sensitivity for both bergenin quantification and trace analysis (Table 1).

Methanol and aqueous extracts from *Endopleura uchi* collected in the dry period (1st collect) and rainy period (2nd collect) were analyzed at 250 μ g mL⁻¹. The quantification of bergenin in *Endopleura uchi* extracts were performed using the calibration slope with detection at 272 nm (Supplementary Material Table 2S). Bergenin was identified in chromatogram extracts by comparison with the chromatogram of the standard using UV parameters with λ_{max} 215, 254 and 272 nm and retention time at 11 min. The concentration of bergenin in dried bark (%) was determined using the yield of extracts (%) at Table 2.

Both dry and wet season extracts exhibit similar bergenin concentrations. Each extract showed similar extractive and bergenin levels when dry and rainy seasons have to be compared. When bergenin concentration is compared between methanol and aqueous extracts, one finds the extraction using methanol to yield a higher bergenin concentration, though in aqueous extracts it showed to be about 10% lower than in leaves methanol extracts and 30% lower than in bark and twigs methanol extracts. Therefore, these findings indicate aqueous extracts to bear high bergenin concentration, especially the leaves aqueous extracts that show similar to methanol extracts values. The mean barks aqueous extracts bergenin concentration in dry season and rain season showed to be similar to that reported in Nunomura *et al.*,⁷ that showed a concentration of 3.19%. The lower concentration of bergenin is observed in aqueous extract of twigs with an average concentration of 1.9%. These findings indicate bergenin to be the *Endopleura uchi* main component being found in all parts of the plant and possibly its active principle that is used as a medicinal

Table 1. Curve data of standard solutions of bergenin (6.25-200 μ g mL⁻¹)

λ (nm)	Equation curve ^a	Linear coefficient (R^2)	DL ^b	QL ^c
272	$y = 66.349,9399 x - 17.824,6393$	1.000	9.85	29.84
254	$y = 36.770,1945 x - 4.814,1791$	1.000	8.88	26.92
215	$y = 242.018,3217 x - 956.260,4163$	0.996	0.78	2.37

^aSix points (n= 2, interday and intraday). ^bDetection limits (μ g mL⁻¹). ^cQuantification limits (μ g mL⁻¹).

Table 2. Quantification of bergenin in different *Endopleura uchi* extracts with detection at 272 nm

Barks		Twigs		Leaves	
Extract ^a	Bergenin (%±CV) ^b	Extract ^a	Bergenin (%±CV) ^b	Extract ^a	Bergenin (%±CV) ^b
CEA-1	3.31 ± 0.4	GEA-1	1.95 ± 1.0	FEA-1	3.24 ± 2.0
CEA-2	2.87 ± 3.0	GEA-2	1.89 ± 2.4	FEA-2	2.21 ± 2.1
CEM-1	4.17 ± 0.5	GEM-1	3.44 ± 5.0	FEM-1	2.23 ± 0.7
CEM-2	4.75 ± 1.0	GEM-2	2.66 ± 0.4	FEM-2	3.87 ± 2.0

^aCEA = aqueous barks extract, CEM = methanol barks extract, GEA = aqueous twigs extract, GEM = methanol twigs extract, FEA = aqueous leaves extract, FEM = methanol leaves extract, 1 = rain season, 2 = dry season. ^bCV = Coefficient of Variation, from triplicate analyses (n=3).

plant in Amazonia. Bergenin can be used as a standard reference to analyze *Endopleura uchi* commercial products quality control.

Total phenolic analysis

Plant phenolic compounds, such as phenolic acids, flavonoids, coumarins, lignins and lignans, tannins and stilbenes are well known. These compounds have presented antioxidant activities by free radical scavenging or neutralization. Besides, they can act by chelating of transition metals.²⁴ Therefore, the total phenolic analysis was performed by Folin-Ciocalteu method in *Endopleura uchi* barks, twigs and leaves aqueous, methanol and ethanol extracts. The results are shown in Table 3 confirm the presence of significantly higher-valued phenolic compounds than those found on other plant species, as reported on all extracts in other studies.²⁵ The best results were observed in methanol extracts (38.24-43.02 mg GAE g⁻¹ of dry extract), aqueous extracts (23.69-38.94 mg GAE g⁻¹ of dry extract) and ethanol extracts (18.43-28.30 mg GAE g⁻¹ of dry extract).

Antioxidant activities

The antioxidant capacity was accomplished through two different methods as a way of attaining better results and avoiding unreal conclusions due to each method's limitations.^{26,27} The aqueous, ethanol and methanol extracts were tested by FRAP (Ferric Reduction Power Assay) and by DPPH free radical scavenging methods according to Table 3.

Twigs, leaves and barks methanol extracts, respectively, resulted in the highest antioxidant activities by FRAP method. These extracts also presented the best phenolic compounds concentration and, these findings suggest these compounds be responsible for the antioxidant activity achieved by FRAP.²⁸ In aqueous extracts the antioxidant activity was higher in leaves (469.99-567.89 mmol Fe⁺² g⁻¹), twigs (316.97-359.27 mmol Fe⁺² g⁻¹) and barks (230.43-239.30 mmol Fe⁺² g⁻¹), respectively. Leaves ethanol extracts presented better results (365.38-395.96 mmol Fe⁺² g⁻¹) than barks (241.93-326.53 mmol Fe⁺² g⁻¹) and twigs (211.47-290.24 mmol Fe⁺² g⁻¹) ones. Bergenin presented lower values (176.1 mmol Fe⁺² g⁻¹) than *Endopleura uchi* extracts by FRAP method.

DPPH free radical scavenging capacity by the method is a rapid method to assess the compounds and extracts antioxidant activity.²⁹ Our findings were compared with quercetin as a positive control (IC₅₀ = 4.50 µg mL⁻¹).

Extracts from twigs resulted in the best activities (IC₅₀ = 12.08-13.89 µg mL⁻¹), followed by leaves (IC₅₀ = 14.00-20.60 µg mL⁻¹) and barks (IC₅₀ = 12.04 to 24.20 µg mL⁻¹). Although bergenin is the main extracts component, it presented a low antioxidant activity by DPPH method. The concentration of 1000 µg mL⁻¹ inhibits only 33% of DPPH radical scavenging activity, therefore the concentration to inhibit 50% should be higher than

Table 3. Total phenolics and antioxidant activities of extracts of *Endopleura uchi*

Extracts ^a	FT (mg GAE ^b g ⁻¹ dry extract)	FRAP (mmol Fe ⁺² g ⁻¹ dry extract)	DPPH (IC ₅₀ µg mL ⁻¹)
Barks (value ± CV ^c)			
CEA-1	23.69 ± 3.7	230.43 ± 0.4	24.20 ± 1.6
CEA-2	23.94 ± 1.5	239.30 ± 0.8	22.13 ± 0.4
CEM-1	42.49 ± 0.7	449.47 ± 0.3	15.63 ± 2.6
CEM-2	42.17 ± 0.1	495.55 ± 0.9	12.04 ± 0.7
CEE-1	18.43 ± 2.4	241.93 ± 0.6	13.89 ± 0.3
CEE-2	26.49 ± 1.0	326.53 ± 0.7	12.51 ± 2.0
Twigs (value ± CV ^c)			
GEA-1	27.94 ± 1.4	359.27 ± 0.2	13.89 ± 4.2
GEA-2	26.04 ± 4.3	316.97 ± 1.4	12.36 ± 1.4
GEM-1	42.16 ± 2.8	527.85 ± 0.8	13.62 ± 0.9
GEM-2	40.03 ± 1.3	512.58 ± 1.2	12.08 ± 2.3
GEE-1	16.69 ± 1.3	211.47 ± 1.6	13.51 ± 1.7
GEE-2	21.68 ± 0.5	290.24 ± 1.0	12.38 ± 1.7
Leaves (value ± CV ^c)			
FEA-1	38.94 ± 2.0	567.89 ± 2.7	14.76 ± 1.1
FEA-2	35.86 ± 1.6	469.99 ± 0.6	18.40 ± 0.9
FEM-1	43.02 ± 2.6	512.39 ± 0.9	14.00 ± 1.7
FEM-2	38.24 ± 3.1	476.35 ± 1.8	20.36 ± 0.3
FEE-1	32.46 ± 2.9	395.96 ± 0.4	17.50 ± 0.3
FEE-2	28.30 ± 1.4	365.38 ± 0.5	20.60 ± 1.5
Standard (value ± CV ^c)			
Quercetin	-	-	4.50 ± 0.4
Bergenin	-	-	>1000

^aCEA = aqueous barks extract, CEM = methanol barks extract, CEE = ethanol barks extract, GEA = aqueous twigs extract, GEM = methanol twigs extract, GEE = ethanol twigs extract, FEA = aqueous leaves extract, FEM = methanol leaves extract, FEE = ethanol extract of leaves, 1 = rain season, 2 = dry season. ^bGAE = gallic acid equivalent. ^cCoefficient of Variation (CV), from triplicate analyses (n=3).

1000 µg mL⁻¹ that is considered no significant antioxidant activity. According to Subramanian *et al.*,³⁰ bergenin showed mild activity by the methods of DPPH, FRAP and nitric oxide inhibition. The antioxidant activity of extracts by DPPH method suggests *Endopleura uchi* extracts' potential antioxidant activity when compared with the leaves' commercial extracts from *Ginkgo biloba* (IC₅₀ = 40 µg

mL⁻¹) and other Brazilian plants used as antioxidants.²⁵ The hydroxyl groups from phenolic compounds as flavonoids, phenolic acids, tannins, and other compounds can be responsible for these antioxidant activities.³¹⁻³³

LPS-induced NO production inhibition

According to traditional medicine, the species *Endopleura uchi* is mainly used for the treatment of menstrual disorders and uterine inflammation, therefore its effect on the aqueous extracts, chloroform and ethyl acetate fractions inflammatory response through production of inflammatory mediators by LPS-induced in murine macrophage cells (J774), has been ascertained. It measured the ability to inhibit inflammatory mediators NO production.²³ Macrophage cells were exposed to LPS for NO-stimulation and treated with extracts and fractions in the concentration of 25 µg mL⁻¹. The tested samples showed cytotoxicity at concentrations above 100 µg mL⁻¹, so 25 µg mL⁻¹ was considered safe to test the samples. The negative control (cells + LPS) produced 8.5 µM of NO and the positive control (PC) (cells) produced 0.1 µM of the inflammatory mediator. The results showed cells treated with the extracts and fractions of *Endopleura uchi* to produce a significant inhibition (Figure 2). The best results were observed in leaves chloroform fractions (FEMC), twigs ethyl acetate (GEMAc) and chloroform (GEMC) fractions and bark chloroform fraction (CEMC) resulting in 0.1, 0.5, 0.6 and 0.6 µM of NO respectively in LPS-induced NO production. The other extracts and fractions obtained NO production between 1.4-3.5 µM, below the negative control (8.5 µM of NO). When assessing the inhibitory capacity one finds the chloroform fractions of the leaves (FEMC) and bark (CEMC) de *Endopleura uchi* to be the most active in producing IC₅₀ = 3.2 and 4.8 µg mL⁻¹ respectively. The remaining extracts and fractions showed values between IC₅₀ = 7-12 µg mL⁻¹ (Figure 3).

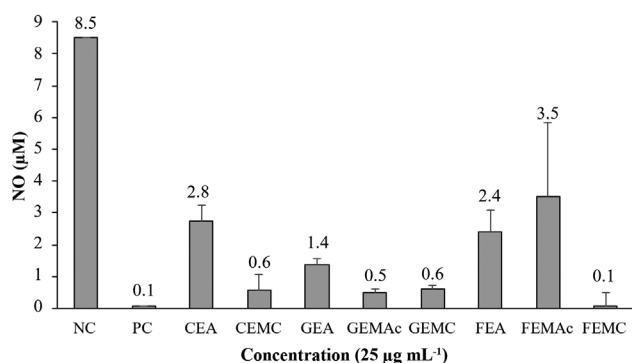


Figure 2. Effect of concentration 25 µg mL⁻¹ of the extracts and fractions of *Endopleura uchi* in the production of NO in culture of macrophages stimulated by LPS. Values are expressed as mean ± standard deviation obtained from analyzes in triplicate (n=3). NC = negative control; PC = positive control; CEA = aqueous barks extract; CEMC = barks chloroform fraction; GEA = aqueous twigs extract; GEMAc = twigs ethyl acetate fraction; GEMC = twigs chloroform fraction; FEA = aqueous leaves extract; FEMAc = leaves ethyl acetate fraction; FEMC = leaves chloroform fraction

Studies show evidence that the NO molecule, when produced in excess, can contribute to certain pathological conditions such as rheumatoid arthritis,³⁴ Alzheimer's and Parkinson's³⁵ diseases among others, therefore there is growing demand for substances capable of attenuating the NO production, which may have estimable therapeutic value in the treatment of excess nitric oxide-induced pathophysiological conditions.³⁶ The results indicate the extracts and fractions of *Endopleura uchi* to show NO inhibitory effect, therefore, it can be regarded as sources of active substances in the treatment of

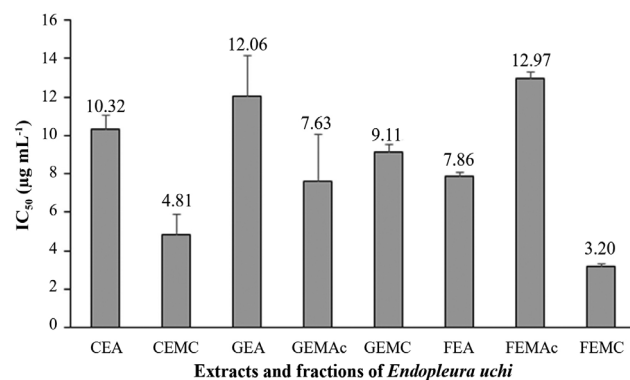


Figure 3. Inhibitory capacity (IC₅₀) of NO induced by LPS in J774 murine macrophage extracts and fractions. The data are represented as mean ± standard deviation. CEA = aqueous barks extract; CEMC = barks chloroform fraction; GEA = aqueous twigs extract; GEMAc = twigs ethyl acetate fraction; GEMC = twigs chloroform fraction; FEA = aqueous leaves extract; FEMAc = leaves ethyl acetate fraction; FEMC = leaves chloroform fraction

inflammatory processes. These findings should be taken into account for the isolation and identification of these active substances.

CONCLUSIONS

Since the findings showed to be positive for antioxidant activity by iron reduction, DPPH radical scavenging capacity methods, as well as an anti-inflammatory activity through NO suppression, they suggest that species *Endopleura uchi* can be considered an important plant material bearing therapeutic potential. Quantification showed a significant amount of bergerin in its leaves and twigs, indicating that the substance bearing different biological activities may be attained from other plant parts apart from the barks. These findings can serve as the basis for preparing antioxidant and anti-inflammatory compounds that may contribute to the safer use of this plant species along with the development of possible herbal products.

SUPPLEMENTARY MATERIAL

Figures related to the results of some experiments used in this article are in the supplementary material that is available in <http://quimicanova.sbq.org.br> in the form of an open PDF file.

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ERRATUM

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Where it reads:

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