Myrciaria Glazioviana Leaves Anti-Inflammatory Antinociceptive And Hepatoprotection Potential and Chemical Profile by Mass Spectrometry

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ABSTRACT

Plants represent an ancient source of medicines for humanity and an important source of active substances. Myrciaria glazioviana belongs to Myrtaceae family with antibacterial and antifungal activities described. The objective of this study is to evaluate the anti-inflammatory and antinociceptive activity of the dichloromethane extract of the leaves of M. glazioviana through in vivo models (Carrageenan-induced paw edema model and abdominal constriction test induced by acetic acid, respectively) and to identify compounds present in the crude extract by mass spectrometry. Histopathological analyzes of the animal samples used in the studies were also performed. It was possible to observe the presence of antinociceptive, anti-inflammatory and hepatoprotective activities in M. glazioviana extract, as well as the presence of phenols and flavonoids in its composition and Gallic acid and Quinic acid were identified.

Abbreviations: MGD: Dichloromethane Extract of the Leaves of Myrciaria Glazioviana, GAE: Gallic Acid Equivalent, NAL-UFF: Laboratory Animal Nucleus of the UFF, HE: Hematoxylin Eosin

Introduction

Plants are used for many years in treatment various pathologies representing one of the most effective sources of active substances. Products of natural origin have already been described as having anti-inflammatory, analgesic and hepatoprotective properties [1,2]. Myrciaria glazioviana is a Brazilian species belonging to the Myrtaceae family, popularly known as yellow jabuticaba and hairy fruit. It has botanical synonyms such as Eugenia cabeluda and Plinia glomerata [3,4]. This species is popularly used as anti-inflammatory and analgesic [5]. Previously published articles confirm the antimicrobial activities [3], analgesic [5] and antioxidant [4] of the species. Currently, most anti-inflammatory drugs are non-steroidal drugs, which have many adverse effects, especially when used continuously by patients with chronic diseases such as rheumatoid arthritis, Crohn’s disease and some cancers [6-9]. The side effects of commonly observed nonsteroidal anti-inflammatory drugs are gastrointestinal injury, peptic ulcer and renal dysfunction [8,9]. In the search for new bioactive substances that can be anti-inflammatory drugs with fewer side effects base of existing classes, the goal of this study was to evaluate the anti-nociceptive and anti-inflammatory potential of the dichloromethane extract of the leaves of M. glazioviana and analyze its content of flavonoids and phenols. Also, was evaluated through histological analysis, changes of any type, including toxicity or protective of tissue of the extract in the induced inflammatory process.

Material and Methods

Plant Material and Extract Preparation

The leaves of M. glazioviana were collected in their natural habitat, in the city of Nova Friburgo (22° 17’02.6”S 42° 31’29.9”W) identified by the taxonomist Dr. Gustavo Hiroaki Shimizu of the Institute of Biology of State University of Campinas - São Paulo. The leaves of the plant were stabilized and dried, passed through a pulverization process and maceration with dichloromethane. After removal of all solvent, the dichloromethane extract of M. glazioviana (MGD) was obtained.
Phenolics and Flavonoids Quantification of M. glazioviana extract

For analysis of the content of soluble phenolic compounds of the extract, the Folin-Ciocalteu method was used. In this method, gallic acid was used as the standard sample. An analytical curve was made at different concentrations. Then, the absorbances of the samples and standard sample were measured in a spectrophotometer at 730 nm. The result is given as μg of gallic acid equivalent (GAE) per milligram of extract (μg of GAE/mg). For determination of total flavonoids was used for spectrophotometric assay based on formation of a complex between the ion Al+3 and the carbonyl group present in the flavonoid molecule. The absorbance of the sample and standard sample (quercetin) were accomplished in a spectrophotometer at 415 nm. The results were expressed as mg of quercetin equivalent per milligram of extract (μg quercetin/E/mg).

Both experiments were performed in triplicate.

Analysis of The Chemical Profile of The Extracts By ESI-MS/MS

Preliminary analysis of the extracts of the plant species were performed by direct infusion in a TQD Acquity mass spectrometer (Micromass-Waters, Manchester, England) using an ESI source in the negative ionization mode. Samples (1mg/ml) were dissolved in a solution containing 50% (v/v) chromatographic grade methanol and 50% of a water-alcohol solution 0.5% ammonium hydroxide (Merck, Darmstadt, Germany). The instrument parameters were: source temperature 100ºC, capillary 3.0 kV and 30 V cone voltage, collision energy 30 V. The constituents were identified by comparison of their ESI-MS/MS fragmentation spectra with data from the database (www.massbank.eu).

Animals

Adult female Swiss mice, 25-30g, from the Laboratory Animal Nucleus of the UFF (NAL-UFF) were used and kept in the Multisuser Laboratory of Biomedical Research of Health Institute of Nova Friburgo. For the beginning of the experiments, the study had already been submitted and approved to the Committee of Ethics in Animal Experimentation of UFF (nº 614).

Analysis of Antinociceptive Activity In Vivo

Analysis of the antinociceptive activity was performed by the abdominal constriction test induced by acetic acid. Initially, the animals were divided into 4 groups (n = 6) and received by gavage treatments: MGD at doses of 125mg/kg and 250mg/kg, Indomethacin 10mg/kg and saline 0.9% for positive and negative controls, respectively. After 30 minutes of treatment, the edema is induced through injection of carrageenan (30µg/paw) in the subplantar tissue of the right hind paw. The paw thickness is measured using a digital Vernier caliper before and 1, 2, 3 and 4 hours after carrageenan injection. The anti-inflammatory activity is analyzed by the percentage increase of the paws in different groups.

Histopathological Analysis of Liver and Kidney Samples

At the end of the previous experiment, the animals were euthanized by cervical dislocation and samples of liver and kidney of each animal were collected for histological analysis. The samples were subjected to routine processing, included in Paraplast®, sectioned and colored with hematoxylin eosin (HE). The histological blades were observed and photographed under a Diagtech® microscope linked to a video camera DCM510 using 100x magnification for samples of liver and 40x magnification for samples of kidney. For quantification of altered cells, was used the software STEPanizer® stereology tool, Version 1.0 and for the statistical analysis was used the software GraphPad Prism® Version 6.

Results and Discussion

Through the methods for quantification of total phenols and flavonoids of the extract of M. glazioviana was obtained a concentration of total soluble phenols of 13.97μg GAE/mg and a concentration of total flavonoid and 12.35μg quercetin E/mg. This result agrees with published studies that relate the presence of these compounds with species of the same family and genus [10-12]. The extract of M. glazioviana was analyzed by ESI-MS/MS in negative ion mode. Thus, by comparing its MS/MS fragments with data from the literature and database, it was possible to identify Quinic acid and Gallic acid. Their m/z and fragmentation are described in Table 1. Through the model of abdominal constriction induced by acetic acid, it was possible to determine the number of abdominal contractions of different groups per 30 minutes (Figure 1). The result of the analysis of variance (ANOVA) followed by Tukey test (p <0.05) showed that the positive control, the drug Indomethacin (10 mg/kg) and MGD extract at both doses showed statistically significant differences when compared to the negative control (Figure 1).

Table 1: Identified substances in the ethanolic extract of S. microglossa flowers by ESI-MS / MS and collision induced dissociation at 30 eV and MS/MS fragments.

<table>
<thead>
<tr>
<th>Precursor Ion [M-H]⁻ m/z</th>
<th>MS/MS Fragments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Substances</td>
<td></td>
</tr>
<tr>
<td>Quinic acid</td>
<td>191</td>
</tr>
<tr>
<td>Gallic acid</td>
<td>169</td>
</tr>
</tbody>
</table>

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The Indomethacin was able to decrease 68.18% of abdominal constriction, MGD 250 mg/kg decreased 60.61% and MGD 125 mg/kg inhibited 46.10% of abdominal constriction. Therefore, it was possible to determine the presence of anti-nociceptive activity of PGD extract, this result agrees with studies that relate this species with analgesic properties [5]. Furthermore, there is evidence that the presence of flavonoids and phenolic compounds is often related to this biological activity [5,13,14]. In the paw edema model induced by carrageenan, the edema produced is a sign of inflammatory response and is the result of increased vascular permeability. This edema was progressive in all groups during the 4 hours experimental observation, varying the increase from 33.05 to 71.91% in the negative control group and 26.48 to 34.76% in the positive control group. In the experimental groups, MGD 125 mg/kg and 250 mg/kg, the increase in paw thickness varied between 24.06% to 71.21% and 21.77% and 48.03%, respectively (Figure 2). The groups, MGD 250 mg/kg, at time 1 and 4, and Indomethacin 10 mg/kg, at time 3 and 4, showed statistical difference when compared to the negative control. For data analysis, we used the analysis of variance (ANOVA) followed by Tukey test, considering $p < 0.05$. From this result, it was possible to observe the anti-inflammatory activity of MGD 250mg/kg.

**Figure 1:** Number of abdominal contusions of different groups (Indomethacin 10mg/kg, MGD 125mg / kg and MGD 250mg / kg) during 30 minutes, where * represents the groups that had a statistical difference when compared to the negative control. It was considered $p < 0.05$ and 95% confidence (ANOVA followed by Tukey test).

**Figure 2:** M. glazioviana activity in carrageenan-induced paw edema. The effect of extract of *M. glazioviana* 250mg/kg and 125mg/kg on carrageenan-induced paw edema is expressed as a percentage of increase in paw thickness (%) per hour after induction of edema. The animals were treated with saline 0.9% (negative control), Indomethacin 10 mg/kg and *M. glazioviana* 250 mg/kg and 125mg/kg. The edema volume was evaluated using a digital caliper 1, 2, 3 and 4 hours after induction of inflammation, where

A) Indomethacin 10mg/kg, and

B) *M. glazioviana* 250mg/kg differed statistically from the saline group. It was considered $p < 0.05$ and 95% confidence (ANOVA).
The model used is capable of evaluating the effect of non-steroidal anti-inflammatory drugs and the participation of inflammatory mediators such as histamine, serotonin and bradykinin. During the first 4 hours of the experiment, there is release of the mediators previously mentioned, as well as phospholipase-A2, kinins and metabolites of arachidonic acid [15,9]. Thus, it is suggested that the extract of M. glazioviana was able to inhibit the action of these mediators. However, this model is considered an initial screening, requiring the induction with other phlogistic agents to accurately determine the performance of the extract in the inflammatory process. Through histological analysis of samples of liver and kidney collected, it was possible to observe the presence of hyperemia above in all animals in both samples, once the vessels and arteries showed dilated and an increased amount of erythrocytes in inside. The animals were subject to an acute inflammation model, and consequently there is increase of vascular permeability and vasodilatation that gives rise to edema and hyperemia, respectively [16,17]. Another observation made during histological analysis, was the presence of hydropic degeneration of hepatocytes since these had greater volume and large vacuoles in their cytoplasm. Further along the experiment was observed during handling, the animals showed a hyperthermia, probably induced using phlogistic agent. It is known that hyperthermia is a cause potential of the hydropic degeneration, since there is an increase in consumption of ATP by electrolytic pumps, leading to isosmotic expansion of the cell and consequently the balloon aspect present in this type of degeneration [16,18]. It can affirm then, that such a change is not related to extract toxicity, as all groups are involved.

Through the quantification of cells with hydropic degeneration in experimental groups was noted that MGD group 250 mg/kg had fewer altered hepatocytes when compared to the other groups. The result of the analysis of variance (ANOVA) followed by Bonferroni test (p < 0.05) showed that the negative control group showed 15.38% of hepatocytes with hydropic degeneration, and the positive control showed 13.33% of hepatocytes changed. In the groups MGD 125mg/kg and 250mg/kg, this change was 11.12% and 4.67%, respectively. Therefore, when comparing the MGD group 250mg/kg with the other groups, this showed a statistical difference (Figure 3). This result suggests a potential hepatoprotective of the MGD extract. The hepatoprotective activity is, in many studies, directly associated to the presence of flavonoids [19-21] and antioxidant activity [22,23], which has already been described in M. glazioviana [4]. Liver diseases represent a global health problem given the importance of liver in the homeostasis. Many synthetic drugs currently used in the treatment of these diseases have undesirable adverse effects, so this result obtained in this study is extremely important because it opens a new possibility for study with M. glazioviana [24-26].

**Conclusion**

Therefore, it is concluded that M. glazioviana extract showed antinociceptive and anti-inflammatory activity in vivo models and hepatoprotective activity in cases of hydropic degeneration as well as presented between their compounds and flavonoid phenols. These results show the phytotherapeutic potential of M. glazioviana, which may influence the increase of the studies involving this species.

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**References**


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