Antinociceptive Action and Redox Properties of Citronellal, an Essential Oil Present in Lemongrass


1Department of Physiology, Federal University of Sergipe, Aracaju, Sergipe, Brazil.
2Center for Studies in Oxidative Stress, Federal University of Rio Grande do Sul, Porto Alegre, Rio Grande do Sul, Brazil.

ABSTRACT Citronellal (CT) is a monoterpenoid and the major constituent of the mixture of terpenoids that give the citronella oil its lemon scent. Citronella oil is widely used around the world for various purposes and is mainly obtained from plants of the Cymbopogon genus, which are known as “lemongrass.” Considering these plants have been used worldwide for various medicinal purposes, the interest of researchers to understand the biological activities of monoterpenoids related to the Cymbopogon genus has been increasing. In the present work, we investigated the antinociceptive action and the redox properties of CT. Our results indicate that intraperitoneal injection of CT was effective in reducing nociceptive face-rubbing behavior in both phases of the formalin test, which was also naloxone-sensitive. CT also evoked antinociceptive response in the capsaicin and glutamate tests. The total radical-trapping antioxidant parameter and total antioxidant reactivity assays indicate that CT at doses of 0.1 and 1 mg/mL exerts a significant antioxidant activity, which is probably related to its ability to scavenge superoxide and nitric oxide, but not H2O2 or hydroxyl radicals, as evaluated separately by specific in vitro tests. These results show for the first time the antinociceptive potential of CT and indicate that the antioxidant properties of this compound may rely on its mechanism of biological actions because CT-containing natural products are used to treat various diseases related to oxidative stress and reactive species.

KEY WORDS: • antinociception • citronellal • free radicals • oxidative stress

INTRODUCTION

Epidemiological studies have been suggested an inverse correlation between the consumption of certain natural products and the risk of degenerative diseases, cancer, cardiovascular diseases, and chronic pain.1–4 The presence of secondary metabolites with antioxidant activity or other redox-related properties is generally associated with such properties, as many of the conditions in which these natural products have been observed to exert a preventive/therapeutic action are strongly related to oxidative stress and increased reactive species production.5,6 The increase in reactive species production may result from multiple endogenous (i.e., aging, mitochondrial metabolism) and exogenous (smoke, ultraviolet radiation, pollution) factors and induce oxidative damage to biomolecules such as DNA, proteins, and lipids.

The essential oils are natural products that exhibit various biological properties, such as analgesic,7 anticonvulsant,8 and anxiolytic9 effects. Those effects are attributed to the monoterpenes, which are the major chemical components of these essential oils. Citronellal (CT) is a monoterpene product from plant secondary metabolism. It is typically isolated as a non-racemic mixture of its R and S enantiomers by steam distillation or solvent extraction from the oils of Corymbia citriodora Hill and Johnson (former Eucalyptus citriodora Hook), Cymbopogon nardus, Cymbopogon citratus, and Cymbopogon winterianus (“Java citronella”).5,10 It is also found in more than 50 other essential oils. Plants from the Cymbopogon genus are known under various names, the most popular being “lemongrass” and “citronella.”11 Cymbopogon leaves are used as infusion preparations for use as spices, medicinal and recreational teas, soups, and curries.11 Other CT-containing plants, such as members of the genera Eucalyptus, Melissa, Mentha, Allium, and Cinnamomum, among others, also present notable nutritional value and are important food/spice sources in different parts of the world.12 In Cymbopogon (lemongrass) and Melissa officinalis (balm), CT is considered a major

Manuscript received 11 May 2010. Revision accepted 3 August 2010.
Address correspondence to: Daniel Pens Gelain, Ph.D., Departamento de Bioquímica, Universidade Federal do Rio Grande do Sul, Rua Ramiro Barcelos, 2600, anexo, CEP 90035-003, Porto Alegre, RS, Brazil, E-mail: dgelain@yahoo.com.br
constituent of the leaves, contributing to a great extent of the aromatic scent of these plants. The use of lemongrass in curries in Asia and in spices in Africa is also related to prevention of food spoilage, and the preservative properties of these plants are probably related to their high content of essential oils.\textsuperscript{13–16} Nonetheless, isolated properties of the different essential oil components of these medicinal plants, such as CT, have been poorly studied.

CT is produced by many plants traditionally used to treat different conditions by widespread populations around the world, such as South America and Asia.\textsuperscript{17–19} Although CT has been identified, purified, and chemically well defined, its biological and antioxidant properties have been very poorly explored. For instance, plants of the \textit{Cymbopogon} genus, which are rich in mono-, di-, and triterpenes, are widely used to treat different forms of pain.\textsuperscript{20–24} Holanda Pinto \textit{et al.}\textsuperscript{25} demonstrated the analgesic effect of the triterpenoid \textit{x,\beta}-amyrin, a compound closely related to CT. Also, CT and related monoterpenes were reported to inhibit cytochrome P-450 CYP2B1 and other dehydrogenases,\textsuperscript{26} indicating that such compounds may exert a diverse array of biological actions.

Pain is a complex, multidimensional experience especially relevant to the orofacial region because the face and mouth have a particular biological and psychological meaning for each individual. Many of the most common types of pain occur in the orofacial region, and most difficulties in the management of conditions related to acute and chronic orofacial pain arise from a lack of understanding of its mechanisms.\textsuperscript{27} Free radicals, especially nitric oxide (NO), are also involved in the control of processes related to pain because such species evoke cell signaling cascades related to modulation of neuropathic and inflammatory pain.\textsuperscript{28} In the present work, we investigated the antinociceptive effect of CT and its redox properties.

\textbf{MATERIALS AND METHODS}

\textbf{Animals}

Male Swiss mice (weighing 30–36 g) and male Wistar rats (weighing 230–260 g), 2–3 months of age, were used throughout this study. The animals were randomly housed in appropriate cages at 22 ± 2 °C on a 12-hour light/dark cycle (lights on 6:00 a.m. to 6:00 p.m.) with free access to food and water. All experiments were carried out between 9:00 a.m. and 2:00 p.m. in a quiet room. Experimental protocols were approved by the Animal Care and Use Committee (CEPA/UFS number 12/08) at the Federal University of Sergipe, Aracaju, Brazil.

\textbf{Chemicals}

2,2′-Azobis(2-methylpropionamidine) dihydrochloride (AAPH), luminol (5-amino-2,3-dihydro-1,4-phthalazinedione), 2-deoxyribose, glycine (aminoacetic acid), Griess reagent, sodium nitroprusside (SNP), ascorbic acid, 2-thiobarbituric acid (TBA) (4,6-dihydroxypyrimidine-2-thiol), hydrogen peroxide, adrenaline, catalase, and superoxide dismutase (SOD) were purchased from Sigma Chemical Co. (St. Louis, MO, USA). Trolox (6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid) was purchased from Aldrich Chemical Co. (Milwaukee, WI, USA). Water was purified using a Milli-Q\textsuperscript{®} system from Millipore (Milford, MA, USA). All other reagents used in this study were of highest analytical or high-performance liquid chromatography grade. For all \textit{in vivo} experiments the following agents were used: CT [(R5)-(±)-CT, 98% purity, Dierberger, Barra Bonito, Brazil], morphine hydrochloride (União Química, São Paulo, Brazil), naloxone hydrochloride (Neouquímica, Anapolis, Brazil), 37% formaldehyde (Vetec, Rio de Janeiro, Brazil), Tween 80 (polyoxyethylene-sorbitan monolate), and capsaicin (Sigma). Vehicle for animals was 1 drop of Tween 80 (0.2%) dissolved in 0.9% NaCl solution. In those protocols the agents were injected intraperitoneally at a dose volume of 1 mL/10 g. The physiologic solution used for \textit{in vivo} tests was composed of 150 mM NaCl, 4 mM KCl, 2 mM CaCl\textsubscript{2}, 1 mM MgCl\textsubscript{2}, 10 mM glucose, and 10 mM HEPES, adjusted to pH 7.4 with NaOH.

\textit{Formalin test}

Orofacial nociception was induced in mice by subcutaneous injection of 20 \textmu L of 2% formalin into the right upper lip (perinasal area), using a 27-gauge needle.\textsuperscript{27,29} This volume and percentage concentration of formalin were selected from our pilot studies that showed a pain-related biphasic behavioral response (face-rubbing) of great intensity at periods of 0–5 minutes (first phase) and 15–40 minutes (second phase). Pain was quantified at those periods by measuring the time (in seconds) that the animals spent face-rubbing in the injected area with their fore- and hindpaws.\textsuperscript{27} To assess the effects of test drugs, groups of rats (\(n = 8\) per group) were pretreated systemically with vehicle (1 drop of Tween 80 [0.2%] in distilled water, the solvent for CT) or CT (50, 100, and 200 mg/kg, i.p.), 0.5 hours before the local injection of formalin. Morphine (3 mg/kg, i.p.), administered 30 minutes before the algogen, was included as the positive control. In separate experiments, the possible involvement of opioid mechanism was assessed in the antinociception produced by CT or morphine with naloxone (1.5 mg/kg, i.p.), an opioid antagonist, injected simultaneously.

\textit{Capsaicin test}

The orofacial pain was induced by capsaicin administration. Mice (\(n = 8\)) were injected with capsaicin (20 \textmu L, 2.5 \textmu g) subcutaneously into the right upper lip (perinasal area), using a 27-gauge needle. Capsaicin was dissolved in ethanol, dimethyl sulfoxide, and distilled water (1:1:8 by volume). In pilot studies, rodents manifested pain-related face-rubbing behavior following the injection of capsaicin with a high intensity for a 10–20-minute period. Therefore, pain quantification was performed at this period measuring the time (in seconds) that the animals spent face-rubbing the injected area with their fore- and hindpaws. CT (50, 100, and
described by Beirith et al. 

Glutamate-induced nociception

In an attempt to provide more direct evidence concerning the interaction of the CT with the glutamatergic system, we separately investigated whether or not the CT was able to antagonize glutamate-induced orofacial nociception in mice. The procedure used was similar to that previously described by Beirith et al.,30 with some alterations. A volume of 20 μL of glutamate (25 μM per paw prepared in phosphate-buffered saline) was injected in the right upper lip (perinasal area), using a 27-gauge needle. Animals were observed individually for 15 minutes following glutamate injection. Pain quantification was performed at this period measuring the time (in seconds) that the animals spent face-rubbing the injected area with their fore- or hindpaws. Animals (n = 8 per group) were treated with CT (50, 100, and 200 mg/kg, i.p.) or vehicle (1 drop of Tween 80 [0.2%] in distilled water, the solvent for CT) 30 minutes before glutamate injection.

Leukocyte migration to the peritoneal cavity

The leukocyte migration was induced by injection of carrageenan (500 μg per cavity, i.p., 500 μL) into the peritoneal cavity of rats 1 hour after administration of CT (50, 100, and 200 mg/kg, i.p.), vehicle (1 drop of Tween 80 [0.2%] in distilled water, the solvent for CT), or dexamethasone (2 mg/kg, s.c.) by modification of the technique previously described by Bastos et al.31 Mice (n = 6 per group) were euthanized by cervical dislocation 4 hours after carrageenan injection. Shortly after, phosphate-buffered saline containing EDTA (1 mM, i.p., 10 mL) was injected. Immediately a brief massage was done for further fluid collection, which was centrifuged (750 g for 5 minutes) at room temperature. The supernatant was disposed of, and 1 mL of phosphate-buffered saline was added to the precipitate. An aliquot of 10 μL from this suspension was dissolved in 200 μL of Turk’s solution, and the total cells were counted in a Neubauer chamber, under optic microscopy. The results were expressed as the number of neutrophils/mL. The percentage of the leukocyte inhibition was calculated as (1 – T/C)×100, where T represents the treated group’s leukocyte counts and C represents the control group’s leukocyte counts.

Motor activity evaluation

To investigate if the treatments could influence the motor activity of the animals and consequently impair the assessment of the nociceptive behavior in experimental models, the motor activity was evaluated in a Rota-rod apparatus.32,33 Initially, the mice able to remain on the Rota-rod apparatus (AVS², São Paulo) longer than 120 seconds (9 rpm) were selected 24 hours before the test.34 Then the selected animals were divided into five groups (n = 8 per group) and treated intraperitoneally with vehicle (1 drop of Tween 80 [0.2%] in distilled water, the solvent for CT), CT (25, 50 and 100 mL/kg, i.p.), or diazepam (3 mg/kg). Thirty minutes later, each animal was tested on the Rota-rod apparatus, and the time (in seconds) it remained on the bar for up to 120 seconds was recorded after 30, 60, and 90 minutes.

Total radical-trapping antioxidant parameter

An adapted method of total radical-trapping antioxidant parameter (TRAP) assay was used to determine the capacity of CT to trap a flow of water-soluble peroxyl radical produced at constant rate, through thermal decomposition of AAPH.35 In brief, the reaction mixture containing 4 mL of the free radical source (10 mM AAPH) in 0.1 M glycine buffer (pH 8.6), 10 mL of the test samples at different concentrations, and 10 mL of luminol (4 mM) as the external probe to monitor radical production was incubated at 25°C. Trolox (a water-soluble vitamin E analog) was used as reference peroxyl radical scavenger molecule. The chemiluminescence produced was directly proportional to the radical generation and measured in out-of-coincidence mode (Wallac 1409 DSA liquid scintillation counter, Wallac Oy, Turku, Finland) as counts per minute. The TRAP of CT was evaluated for instantaneously inhibition of chemiluminescence as the area under curve (AUC). The total antioxidant reactivity (TAR) was calculated as the ratio of light intensity in absence of samples (I/) light intensity right after CT addition (I) and expressed as a percentage of inhibition. CT vehicle was 1% dimethyl sulfoxide. AUC and radical basal production were acquired by software (GraphPad Software Inc., San Diego, CA, USA) analysis of data.

TBA-reactive substances assay

TBA-reactive species (TBARS) assay was used for quantify lipid peroxidation,36 and an adapted TBARS method was used to measure the antioxidant capacity of CT using egg yolk homogenate as the lipid-rich substrate.35 In brief, egg yolk was homogenized (1% wt/vol) in 20 mM phosphate buffer (pH 7.4), and 1 mL of homogenate was sonicated and then homogenized with 0.1 mL of CT at different concentrations or controls in different concentrations prepared immediately before use. Lipid peroxidation was induced by addition of 0.1 mL of AAPH solution (0.12 M). Trolox was used as the reference antioxidant molecule (positive control); the negative control was only CT vehicle (1% dimethyl sulfoxide). Reactions were carried out for 30 minutes at 37°C. After cooling, samples (0.5 mL) were centrifuged with 0.5 mL of trichloroacetic acid (15%) at 1,200 g for 10 minutes. A 0.5-mL aliquot from supernatant was mixed with 0.5 mL of TBA (0.67%) and heated at 95°C for 30 minutes. After cooling, samples absorbance was measure using a spectrophotometer at 532 nm. The results were expressed as percentage of TBARS formed by AAPH alone (induced control).
Scavenging activity of NO

NO was generated from spontaneous decomposition of SNP in 20 mM phosphate buffer (pH 7.4). Once generated, NO interacts with oxygen to produce nitrite ions, which were measured by the Griess reaction. The reaction mixtures (1 mL each) containing 10 mM SNP in phosphate buffer and vehicle (1% dimethyl sulfoxide) or CT at different concentrations were incubated at 37°C for 1 hour. A 0.5-mL aliquot was taken and homogenized with 0.5 mL of Griess reagent. The absorbance of chromophore was measured at 540 nm. Percentage inhibition of NO generated was measured by comparing the absorbance values of negative controls (only 10 mM SNP and vehicle) and assay preparations. Results were expressed as percentage of nitrite formed by SNP alone.

Hydroxyl scavenging activity

The formation of -OH (hydroxyl radical) from the Fenton reaction was quantified using 2-deoxyribose oxidative degradation. The principle of the assay is the quantification of the 2-deoxyribose degradation product, malondialdehyde, by its condensation with TBA. In brief, typical reactions were started by the addition of Fe²⁺ (6 mM final concentration) to solutions containing 5 mM 2-deoxyribose, 100 mM H₂O₂, and 20 mM phosphate buffer (pH 7.2). To measure CT antioxidant activity against hydroxyl radical, different concentrations of CT and vehicle (1% dimethyl sulfoxide) were added to the system before Fe²⁺ addition. Reactions were carried out for 15 minutes at room temperature and were stopped by addition of 4% (vol/vol) phosphoric acid followed by 1% (wt/vol) TBA (in 50 mM NaOH). Solutions were boiled for 15 minutes at 95°C and then cooled at room temperature. The absorbance was measured at 532 nm, and results were expressed as malondialdehyde equivalents formed by Fe²⁺ and H₂O₂.

Catalase-like activity

The ability of CT to scavenge H₂O₂ (“catalase-like activity”) was measured as described previously. In brief, H₂O₂ diluted in 0.02 M phosphate buffer (pH 7.0) to obtain a 5 mM final concentration was added to microplate wells in which different concentrations was placed. The microplate was immediately placed to monitor the rate of H₂O₂ decomposition in the microplate reader set at 240 nm. Dimethyl sulfoxide at 1% was used as the vehicle.

SOD-like activity

The ability of CT to scavenge superoxide anion (“SOD-like activity”) was measured as previously described. CT was mixed with native purified catalase (100 U/mL stock solution) in glycine buffer (50 mM, pH 10.2). CT vehicle was 1% dimethyl sulfoxide. Superoxide generation was initiated by addition of 2 mM adrenaline, and adrenochrome formation was monitored at 480 nm for 5 minutes at 32°C. Superoxide production was determined by monitoring the reaction curves of samples and measured as a percentage of the rate of adrenaline autoxidation into adrenochrome.

Statistical analysis

Data obtained were evaluated by one-way analysis of variance followed by Dunnett’s test. Differences were considered to be statistically significant when P < .05. Data were analyzed with GraphPad Prism version 5.01 software (1992–2007, GraphPad Software Inc.).

RESULTS

The TRAP and TAR assays indicate that CT presents a significant antioxidant activity at the highest doses tested. At 100 µg/mL and 1 mg/mL, CT inhibited the pro-oxidant pulse generated by the free radical-producing system at the TRAP assay, and TAR calculation indicates a significant antioxidant potential at 1 mg/mL (Fig. 1A and B). On the other hand, all doses of CT were not able to prevent lipo-peroxidation induced by AAPH in vitro in a lipid-enriched system (Fig. 1C).

To establish the profile of CT antioxidant activity, we tested the scavenger activity of CT against different reactive species. CT was able to significantly inhibit superoxide radical production at the doses of 100 µg/mL and 1 mg/mL, as assessed by the SOD-like activity assay (Fig. 2A). However, at these same doses, CT inhibited spontaneous degradation of H₂O₂, as determined by the catalase-like activity assay (Fig. 2B). CT also demonstrated inhibition of SNP-induced nitrite production, indicating a scavenging activity of CT toward NO (Fig. 2C). On the other hand, no activity against hydroxyl radicals was observed in the 2-deoxyribose degradation assay (Fig. 2D). Altogether, these results suggest that the mechanism of CT antioxidant effects observed in TRAP and TAR assays relies on its ability to scavenge NO, superoxide, and H₂O₂ molecules, but not hydroxyl radicals.

Intraperitoneal administration of CT to mice produced a dose-dependent reduction in formalin-induced face-rubbing behavior (Fig. 3A). Antinociception in both the first and second phases was significantly enhanced. We tested the effect of morphine, which reduced the nociceptive behavior in both phases as well, to compare with the effect of CT and evaluate the involvement of the opioid system in the effect of CT. The effects of both CT and morphine were inhibited by naloxone, a specific antagonist of morphinomimetic receptors.

CT significantly (P < .001) reduced, in a dose-dependent manner, the face-rubbing behavior induced by administration of capsaicin to mice (Fig. 3B). All doses produced an effect similar to that of morphine (3 mg/kg). Also, at doses of 100 and 200 mg/kg, i.p., CT significantly decreased the face-rubbing behavior induced by glutamate (Fig. 4A). In the Rota-rod test, CT-treated mice did not show any significant motor performance alterations, indicating the results shown here are not due to inhibition of motor activity (Fig. 4B).

Carrageenan (500 µg per cavity) induced leukocyte migration to the peritoneal cavity 4 hours after stimulus. Figure 5...
shows the inhibitory effect of CT on carrageenan-induced responses in higher doses (45.9% and 65.7% at doses of 100 and 200 mg/kg, respectively). The results obtained with the control group support the effect of CT because the vehicle presented no activity, and the positive control drug, dexamethasone (2 mg/kg, s.c.), inhibited (92.2%, $P < .001$) the carrageenan-induced leukocyte migration to the peritoneal cavity.

**DISCUSSION**

CT is a monoterpenoid and the major constituent of the mixture of terpenoids that give the citronella oil its lemon scent. Citronella oil is widely used around the world for various purposes and is mainly obtained from plants of the genus *Cymbopogon*, which are collectively known as “lemongrass.” Such plants are used in teas, soups, and curries in Asia and are also suitable for poultry, fish, and seafood. Lemongrass is a tall perennial grass, and the lower part of the stem is used as a component of different curry dishes in Asia. Ground pastes of lemongrass are also a common component of curry powders. In Thailand, lemongrass is also used to make a refreshing cold drink called “Nam Takrai,” while in Brazil infusions containing lemongrass are commonly used in folk medicine as a regulator of the functions of digestive and circulatory systems. CT and related compounds are also major constituents in *M. officinalis* (“balm,” “lemon-balm,” “Cidreira”), which is extensively used as tea for both medicinal and recreational purposes and as a flavoring agent in Brazil, England, and Southern Europe. Citronella oil is used in food, in soaps, as an antiseptic, and also as an insect repellent. Recently, a species of lemongrass (*C. citratus*) was observed to selectively induce apoptosis in malignant cancer cells, and this property was attributed to citral. This has led to an increase in the interest of researchers to explore the biological activities of monoterpenoids related to the *Cymbopogon* genus, considering these plants have been used for various medicinal and other purposes. Nonetheless, there are very few works concerning the effects of CT for internal use.

**FIG. 1.** Antioxidant activity and prevention of lipoperoxidation in vitro by citronellal. (A) Total radical-trapping antioxidant parameter (TRAP) of citronellal at different concentrations. Bars represent the area under the curve (arbitrary units) generated by 2,2’-azobis(2-methylpropionamidine) dihydrochloride (AAPH), a free radical source, in the presence of different concentrations of citronellal, as described in Materials and Methods. The system is the area generated during reading of luminescence by AAPH alone. *$P < .05$, **$P < .001$ by one-way analysis of variance with Duncan’s post hoc analysis. (B) Total antioxidant reactivity (TAR) was calculated as the ratio of light intensity in the absence of samples and the light intensity right after citronellal addition and expressed as a percentage of inhibition ($I_0/I$) using the same raw data. *$P < .05$ by one-way analysis of variance with Duncan’s post hoc analysis. (C) Thiobarbituric acid-reactive substances (TBARS) in vitro. Lipid extracted from egg yolk was subjected to oxidative damage by incubation with AAPH, and the ability of different concentrations of citronellal to prevent TBARS formation was analyzed. Control means basal lipid peroxidation; the AAPH-alone group is considered 100% of oxidative damage. *$P < .05$ relative to the control group by one-way analysis of variance with Duncan’s post hoc analysis. Data in all experiments represent mean ± SEM values ($n = 5$) from experiments performed in triplicate.
It was observed that isolated CT or CT-enriched essential oils/plant extracts inhibited P-glycoprotein-mediated transport in breast cancer cells, exhibiting anticonvulsant activity, and inhibited acetyl- and butyrylcholinesterases. However, there are no studies on the redox properties of CT. Considering the involvement of free radicals and related species in the molecular mechanisms underlying the pathogenesis and progression of different forms of diseases, such as cancer and neurodegenerative conditions, the study of the redox-related properties of CT may help to understand the mechanisms of its action and its efficacy to treat such conditions. Considering also the wide use of CT-containing plants and extracts in popular medicine to treat diseases related to chronic and acute forms of pain, we found interesting to evaluate its antinociceptive effect in an animal model.

Acute administration of CT on formalin-induced orofacial pain caused pronounced antinociception, suggesting a central analgesic effect. The antagonistic effect of naloxone suggests participation of the opioid system in the modulation of pain by CT. Also, the inhibitory effect observed with CT on capsaicin- and in the second phase of formalin-induced face-rubbing behavior may be a result of substance P release inhibition or a direct blocking action on its receptor, neurokinin-1 (NK-1), by CT. In this regard, a previous study provided evidence for tonic...

**FIG. 2.** In vitro characterization of citronellal antioxidant activity. (A) Superoxide dismutase-like activity. Control means spontaneous superoxide degradation. (B) Catalase-like activity. Control means spontaneous H$_2$O$_2$ degradation. (C) Nitric oxide scavenging activity. SNP means nitrite formation induced by sodium nitroprusside alone, considered as 100% of nitric oxide formation. (D) Hydroxyl radical formation. Control is 2-deoxyribose degradation to malondialdehyde (MDA) induced by hydroxyl radicals generated by H$_2$O$_2$ and iron, considered 100% of hydroxyl radical formation in this system. Data are mean ± SEM values (n = 5) from experiments performed in triplicate. *P < 0.05 in relation to control, **P < 0.05 in relation to *groups by one-way analysis of variance with Duncan’s post hoc analysis.
activation of NK-1 receptors, through administration of the NK-1 receptor antagonist SR14033, which blocked the second phase of the orofacial formalin test response in rats. Additionally, our results also show that CT inhibited the nociceptive response caused by right vibrissa pad injection of glutamate into mice. This nociceptive response caused by glutamate seems to involve peripheral, spinal, and supraspinal sites, and its action is mediated by \(N\)-methyl-\(d\)-aspartate and non-\(N\)-methyl-\(d\)-aspartate receptors. Several studies have demonstrated that those excitatory amino acid receptors are involved in nociceptive primary afferent transmission, in both the development and maintenance of painful response. Thus, the suppression of glutamate-induced nociception by CT treatment may be associated with its interaction with the glutamatergic system.

The results presented here indicate that CT exhibits a significant antioxidant activity, as seen in TRAP/TAR assays, which are widely accepted methods to reliably establish the ability of isolated molecules to act as general antioxidants \textit{in vitro}. Also, we demonstrated that CT is able to scavenge NO and superoxide molecules. These are relevant species to the development of oxidative and nitrosative stress, which in turn trigger diverse deleterious processes in cells. Antioxidant activities by natural products are generally thought to be responsible for the health-promoting effects observed in diets and supplemenations using vegetables and other plant-derived products. Prevention of cardiac diseases, cancer, and neurodegenerative conditions observed in epidemiologic studies is ascribed to the presence of secondary metabolites in such diets and/or supplemenations. Our results also have shown that CT had no
preventive action on free radical-induced lipid oxidation when incubated in a lipid-rich system, as seen in the TBARS assay (Fig. 1C). This result suggests that CT is not able to act directly as an antioxidant agent by directly interacting with lipids, which is the case of other lipophilic antioxidants such as α-tocopherol. It is possible that lipid peroxidation to cell membranes in vivo may be prevented in the presence of CT, especially in stress conditions related to increased peroxyl and NO radicals. CT antioxidant action against these species may lower cellular oxidative stress levels and consequently diminish the attack of free radicals on lipid membranes, protecting these lipids against oxidation by an indirect mechanism.

We also found that CT presented a pronounced antinociceptive effect in all tests, and the results from the Rotarod test demonstrated that CT did not influence motor activity, which could impair the assessment of the nociceptive behavior. It is possible that such an antinociceptive effect of CT is at least in part mediated by its antioxidant actions because reactive species may play a relevant role in pro-inflammatory processes. During acute and chronic inflammations, superoxide is produced at rates that overwhelm the capacity of the endogenous SOD enzymes to remove it, resulting in superoxide-mediated injury. Important pro-inflammatory roles for superoxide include endothelial cell damage and increased microvascular permeability, up-regulation of adhesion molecules such as intercellular adhesion molecule-1 and P-selectin that recruit neutrophils to sites of inflammation, lipid peroxidation, and oxidation, and DNA damage. Superoxide also activates redox-sensitive transcription factors including nuclear factor-κB and activator protein-1, which in turn regulate genes encoding various pro-inflammatory and pro-nociceptive cytokines.

NO is a key anti-inflammatory and cytoprotective agent when acting as a signaling molecule. We observed that CT is able to inhibit NO production by SNP in vitro, which could indicate that NO quenching by CT would enhance inflammation and nociception. However, it is important to point out that the interaction of NO with other reactive species results in a general loss of its “beneficial” effects; for instance, NO-mediated activation of cyclooxygenase and subsequent release of beneficial and anti-inflammatory prostaglandins is lost when superoxide production is also enhanced because interaction between superoxide and NO leads to in situ formation of peroxynitrite, a potent cytotoxic and pro-inflammatory reactive species. Thus, blockade of peroxynitrite formation by preventing NO or superoxide excessive formation and/or removal of peroxynitrite also results in cytoprotective and anti-inflammatory effects.

Cell recruitment during inflammation depends on the orchestrated release of local mediators that are responsible for local vascular and tissue changes as well as for the recruitment of host defense cells. The inflammation induced by carrageenan involves cell migration, plasma exudation, and production of mediators, such as nitric oxide, prostaglandin E₂, interleukin-1β, interleukin-6, and tumor necrosis factor-α. Those mediators are able to recruit leukocytes, such as neutrophils, in several experimental models. A putative mechanism associated with this activity may be inhibition of the synthesis of many inflammatory mediators whose involvement in the cell migration is well established. However, the exposure to inflammatory mediators such as carrageenan, cytokines, and endotoxin leads to the induction of both cyclooxygenase and NO synthase (inducible NO synthase or type 2 NO synthase). This observation is also supported by several studies showing that the anti-inflammatory activities of natural products, as monoterpenes (linalool and geranial), were a result of inhibiting NO production in carrageenan-induced inflammation on rodents. The experiments with carrageenan clearly showed that CT (100 and 200 mg/kg, i.p.) inhibits leukocyte mobilization to the peritoneal cavity.

In conclusion, we showed here evidence for the first time that CT has antioxidant properties and modulates neuropathic and inflammatory pain in the test models of orofacial pain induced by formalin (through a naloxone-sensitive mechanism), capsaicin, and glutamate. The antioxidant properties of CT may be involved in the mechanism of its antinociceptive action and also possibly in other biological actions of CT. Nonetheless, future studies should address redox properties and their involvement in the mechanism of CT action in biological systems.

ACKNOWLEDGMENTS

This work was funded by CNPq, IBN-Net number 01.06.0842-00, CAPES, FAPERGS-RS, and FAPITEC-SE.
AUTHOR DISCLOSURE STATEMENT

The authors declare they have no competing financial interests.

REFERENCES


32. Caz RL, Filho VC, Yunes RA, Calixto JB: Antinociceptive action of 2-(4-bromobenzoyl)-3-methyl-4,6-dimethoxy benzofuran, a novel xanthoxyl derivative on chemical and thermal models.
53. Beckman JS, Beckman TW, Chen J, Marshall PA, Freeman BA: Apparent hydroxyl radical production by peroxynitrite: implications for endothelial injury from nitric oxide and superoxide. Proc Natl Acad Sci U S A 1990;87:1620–1624.