CHEMICAL COMPOSITION AND ANTIMICROBIAL ACTIVITY OF Piper auritum KUNTH AND Piper holtonii C. DC. AGAINST PHYTOPATHOGENIC FUNGI

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The growth of fungi during pre- and postharvest of fruits may cause spoilage and result in a reduction in quality and quantity. The aim of the present work was to investigate the efficacy of four extracts (n-hexane, dichloromethane, ethyl acetate, and methanol) and the essential oil of Piper auritum Kunth and P. holtonii C. DC. on the growth inhibition of three important postharvest pathogens of fruits (Colletotrichum acutatum, C. gloeosporioides, and Botryodiplodia theobromae). The in vitro antifungal activity was assayed following the poisoned food technique. The results showed that both n-hexane extract (HE) as essential oil (EO) displayed a relative good control against the three pathogens, being the originating materials of P. holtonii the most actives. Mycelial growth of C. acutatum, C. gloeosporioides, and B. theobromae was significantly inhibited at 400 µg mL⁻¹. Additionally, the chemical composition of the bioactive materials was analyzed by gas chromatography-mass spectrometry (GC-MS). Safrole (64.54/56.88%) and apiol (64.24/57.20%) were the major constituents of the EO/HE from P. auritum and P. holtonii respectively. Structural identification was also confirmed by nuclear magnetic resonance. Both compounds exhibited significant antifungal properties. It can be concluded that EOs/HEs from P. auritum and P. holtonii, and their major constituents, have interesting applications to control plant pathogenic fungi.

Key words: Piperaceae, essential oil, apiol, Colletotrichum, Botryodiplodia.

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tem-end rot (causal agent Botryodiplodia theobromae) and anthracnose (causal agent Colletotrichum acutatum and C. gloeosporioides) are two serious diseases that contribute significantly to pre- and postharvest loss of economically important fruits around the world (Verma and Sharma, 1999). These three pathogens have been reported to attack a wide range of hosts including tamarillo (Solania betaceum Cav.), avocado (Persea americana Mill.), mango (Mangifera indica L.), papaya (Carica papaya L.), citrus, apple (Malus domestica Borkh.), and peach (Prunus persica L.), among others. The primary method of control of both diseases involves the use of synthetic fungicides (Eckert and Ogawa, 1985). The pre- and postharvest treatments with Cu, dithiocarbamates, benzimidazole, and triazole compounds has been highly effective for controlling of these postharvest diseases in various fruits. However, effectiveness of chemical treatment has been decreasing in recent years due to the development of tolerance by C. acutatum, C. gloeosporioides, and B. theobromae (Gutiérrez et al., 2003). As a result, it has been necessary to increase the quantities of these compounds that have to be used in the agriculture, with the consequent additional problems of residue persistence, and serious economic damage arising from the decreased quality of fruits (Tripathi and Dubey, 2004). Therefore, public opinion and fruit industry urgently demand new alternatives to combat major postharvest decays that are free of synthetic fungicides (Sharma and Pongener, 2010). In such way, the use of essential oils, plant extracts, and some of their constituents can be a very attractive method for pre- and postharvest disease control of fruits, due of their relatively safe status and wide acceptance by consumers (Ormancey et al., 2001).

Thus, the aim of the current work was to screen the antifungal activity of extracts and essential oils from Piper auritum and P. holtonii, and their major constituents, against the fungi C. acutatum, C. gloeosporioides, and B. theobromae. In addition, for bioactive materials, chemical composition using gas chromatography-mass spectrometry (GC-MS), along with the toxicity on fruit peel, were evaluated.

MATERIALS AND METHODS

Microorganisms and plant material

Colletotrichum acutatum, C. gloeosporioides, and B. theobromae were respectively isolated from infected tamarillo, papaya, and avocado fruits. Fungi were characterized and kindly provided by the Laboratory of

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Phytopathology (Universidad Nacional de Colombia, Medellín). To evaluate the antifungal activity, previously sterile Petri dishes measuring 9 cm in diameter were inoculated with 1 mL of a spore suspension of the fungus and incubated at 25 °C for 48 h. Mycelial disc of 7 mm of diameter was used for antifungal test.

The plant material of *P. auritum* and *P. holtonii* were collected at the Campus of Universidad Nacional de Colombia, Medellín (6°15′ N, 75°34′ W), in October 2010, and identified by Dr. R. Callejas, Universidad de Antioquia, Colombia. Voucher specimens (*P. holtonii*, catalogue number 407594, number of collection 06355; *P. auritum*, catalogue number 494711, number of collection 569) were deposited at the Herbarium “Gabriel Gutiérrez V”-Medelín of the Universidad Nacional de Colombia (Medellín).

**Isolation of essential oil sand preparation of organic extracts**

The fresh aerial part (900 g) of *P. auritum* and *P. holtonii* were submitted to hydrodistillation for 3 h. The distilled water was extracted using ethyl acetate (EtOAc), dried over anhydrous sodium sulfate, and the solvent evaporated. The essential oils (EOs) were preserved in sealed vials at 4 °C until further analysis.

Fresh plant aerial material (900 g) of *P. auritum* and *P. holtonii* were extracted with different solvents in increasing order of polarity, by percolation at room temperature during 1 wk, to obtain hexanic (HE), dichlorometane (DE), ethyl acetate (AE), and methanol (ME) extracts. Then, extracts were dried, the solvents evaporated, and stored in sealed vials at 4 °C until further analysis.

**Determination of antifungal activity of essential oil and organic extracts**

The poisoned food technique described by Grover and Moore (1962), with slight modifications, was used. Essential oils and extracts were mixed with ethanol (0.2% EtOH), and tested at 100, 200, and 400 µg mL⁻¹. Samples were incorporated into the autoclaved and cooled (50 °C) potato dextrose agar (PDA) medium and then poured into sterilized Petri dishes. Subsequently, a mycelial disk of *C. acutatum*, *C. gloesporioides*, and *B. theobromae*, and measuring 7 mm in diameter was placed in the center of the agar plate. Control treatments consisted of Petri plates with the mycelial disc but treated with sterile distilled water (control) and EtOH (solvent control). Thymol (2-isopropyl-5-methylphenol) and carbendazim (methyl benzimidazol-2-ylcarbamate) at 100 µg mL⁻¹ were used as positive controls. Petri dishes were incubated at room temperature and the diameter of the mycelial growth was measured every 24 h for *Colletotrichum* species, and every 12 h for *B. theobromae*. The incubation was stopped when the mycelial mass of control treatment reached the edges of the plates (ca. 11 d for *C. acutatum*, and *C. gloesporioides*, and 6 d for *B. theobromae*). The plates were used in triplicates for each treatment. The relative growth inhibition of the treatment compared to the control was calculated as percentage, using the following formula:

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\text{Inhibition} (\%) = \frac{(C - T)/C \times 100}{},
\]

where *C* and *T* are the radial growth (mm) of fungus in control and treated plates, respectively.

In addition, for bioactive materials and their major constituents, the concentration that inhibits 50% fungal mycelial growth (IC₅₀) was preliminary determined for *C. acutatum* and *B. theobromae* (Rivillas-Acevedo and Soriano-García, 2007).

**Toxicity of bioactive materials on fruit peel**

Extracts and/or essential oils that show a promissory antifungal activity were tested for their phytotoxicity on the avocado (var. Lorena) and tamarillo (var. Naranja) peel. Mature fruits were washed with tap water, air-dried at room temperature (23-25 °C) and its surface permeated with the treatment solution, using a fine brush. Phytotoxicity has been defined as the extent of the apparent burning damages (discoloration) induced at 1% (Khwakhom and Shangchote, 2009). Therefore, bioactive materials were dissolved in ethanol to reach that concentration. Control treatments consisted of fruits treated with sterile distilled water (control) and EtOH (solvent control). Further, fruits were stored in plastic baskets at room temperature for 8 d, and every 24 h a new application was made. Phytotoxicity was evaluated visually using a scale indicating the extent of surface on which color changes (discoloration, necrosis) are observed (Amiri *et al*., 2008). Assays were performed in triplicate for each treatment.

**In vivo assay on natural infected papaya fruit**

The *in vivo* assay was made as described previously by García *et al.* (2008) with some modifications. Briefly, papaya fruits (cv. Hawaiian) free from defects were washed, disinfected with sodium hypochlorite solution (0.05%), rinsed with distilled water, and air-dried. Then, fruits were wounded with a sterile nail in the equatorial zone and inoculated with a micelal mass of *C. gloesporioides*. After 12 h, fruits were sprayed with 2% of extract, EO, or compound that showed a promissory antifungal activity. Fruits were stored in plastic baskets at room temperature (24 ± 2 °C), and the incidence of infected wounds evaluated after 7 d. All experiments were performed with five replicates per treatment and untreated fruits were used as controls.

**Gas chromatography-mass spectrometry (GC-MS) analysis**

The GC-MS analysis of the bioactive materials was performed using a Hewlett-Packard 6890 (Agilent Technologies, Palo Alto, California, USA) equipped with a HP 5973 MSD (Mass selective detector-Quadrupole
type) detector and ZB-35 column (Phenomenex, length: 30 m, film thickness: 0.25 μm, column ID: 0.25 mm). Chromatographic conditions were as follows: the oven temperature was programmed from 50 to 300 °C at 5 °C min⁻¹, and then held isothermally for 10 min; injector temperature, 175 °C; detector temperature, 250 °C; carrier gas was helium (constant flow at 1.0 mL min⁻¹) and the average gas velocity was 36 cm s⁻¹. Samples of 3 µL were injected in the splitless mode. Electron impact mass spectra were obtained at 70 eV of electron energy, and monitored from mass to charge ratio (m/z) 30 to 400. The ion source and quadrupole analyzer temperatures were fixed at 230 and 150 °C, respectively. Relative composition of the individual constituents was determined from the peaks area relative to the total peak area.

Isolation and identification of the major compounds
Bioactive materials were subjected to column chromatography with silica gel 60 (0.063-0.200 mm, Merck) and eluted with n-hexane:EtOAc mixtures of increasing polarity to yield six fractions. Fractions 2 and 3 (n-hexane:EtOAc; 9:1 and 8:2) were purified by preparative thin-layer chromatography (TLC, Si 60 F254, 0.25 mm, Merck) (n-hexane: EtOAc; 9:1) to yield two major compounds, safrole (from *P. auritum*) and apiol (from *P. holtonii*). The molecular structures of these compounds were identified by MS, ¹H and ¹³C Nuclear magnetic resonance (NMR) analysis, and comparison with the NIST 2002 computerized mass spectral library. NMR spectra were recorded in deuterated chloroform (CDCl₃) on an AMX 300 NMR spectrometer (Bruker, Germany, ¹H NMR, 300.12 MHz; ¹³C NMR, 75.42 MHz). Chemical shifts, δ, were expressed in ppm units downfield from tetramethylsilane and coupling constant J in hertz (Hz). 2D experiments (JMOD, ¹H-¹H COSY, HMQC and HMBC) were carried out for structure determination.

RESULTS AND DISCUSSION

Antifungal activity of essential oils and organic extracts
Essential oils and extracts yields of fresh aerial parts of *P. auritum* and *P. holtonii* are shown in Table 1. In general, yields of EO and extracts of *P. holtonii* were higher than those of *P. auritum*. Furthermore, the DE of both Piper species showed the highest yield (0.45 and 0.53% for *P. auritum* and *P. holtonii*, respectively). However, the percentage of EO from *P. auritum* was lower than those reported for leaves (2.3%) and inflorescences (2.4%) using microwave-assisted hydrodistillation and a Clevenger apparatus.

The inhibitory effects of EO and organic extracts of *P. auritum* and *P. holtonii* on the growth of *C. acutatum*, *C. gloeosporioides*, and *B. theobromae* are presented in Figures 1, 2 and 3. In general, the growth of fungal mycelium depends on the EO and extract in the culture medium, and the tested Piper species. The EO and HE of both Piper species exhibited moderate to high antifungal activity against all the tested fungi. At 400 µg mL⁻¹, EO and HE of *P. auritum* inhibited the radial growth of *C. acutatum* completely during the first 48 h. Then, growth inhibition descended to reach 40% after 264 h. Even at 100 µg mL⁻¹, inhibition percentages of *C. acutatum* ranged from 87 (between 24 and 48 h) to 20% (from 192 to 264 h) for both treatments (data not shown). For *P. holtonii*, EO and HE exhibited the strongest antifungal activity. In fact, fungal colony development initiated only after 96 and 144 h when *C. acutatum* was treated respectively with HE and EO at 400 µg mL⁻¹. Indeed, growth inhibitions were

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Table 1. Yields of essential oils and extracts for the aerial parts of *Piper* species.

| Extract/Essential oil | *Piper auritum* | *Piper holtonii* |
|-----------------------|-----------------|-----------------|
|                       | Crude products a | Crude products a |
| n-Hexane              | 2.18            | 2.91            |
| Dichloromethane       | 4.08            | 4.80            |
| Ethyl acetate         | 2.09            | 2.09            |
| Methanol              | 2.22            | 2.45            |
| Essential oil         | 1.08            | 1.17            |

aCrude weight results from 900 g of fresh aerial parts.
bYield (%) = (crude weight/900 g)×100.
even higher than those reached by *P. auritum*, ranging from 100 to 90% (after 264 h) for EO, and 100 to 80% (after 264 h) for HE, at 400 µg mL$^{-1}$. A similar behavior of extracts and EOs against *C. gloeosporioides* was observed. Essential oil exhibited the highest antifungal activity against this fungus followed by HE. Additionally, *P. holtonii* displayed a mycelial growth inhibition higher than *P. auritum*. In the case of *B. theobromae*, EO of both *Piper* species also exhibited the highest mycelial growth inhibitions, followed by HE. During the first 72 h, the fungus was completely inhibited by EO from *P. auritum*. Overall, EO of *P. holtonii* appeared to possess the strongest antifungal activity; it inhibited 100% the fungal growth of *B. theobromae* during 96 h. On the other hand, HE from *P. holtonii* showed a moderated effect against *B. theobromae*, causing growth inhibitions between 80 and 60% after 96 h. As can be seen, ME, AE, and DE showed a weak antifungal activity against all the tested fungi. Thymol and carbendazim presented a significantly higher antifungal effect than all evaluated extracts. Carbendazim is commonly used for control anthracnose disease both in the field and postharvest. However, some plant pathogens, such as *C. acutatum* and *C. gloeosporioides* possess resistant strains to carbendazim (Zhang and Huang, 2007). Meanwhile, thymol has been recognized for its potent antifungal activity (Numpaque et al., 2011).

**Chemical composition of bioactive materials**

The chemical composition of EOs and HEs from fresh aerial parts of *P. auritum* and *P. holtonii* was analyzed by means of GC-MS. The detected constituents (relative amounts higher than 0.1%) are shown in Table 2. Analysis by GC-MS led to the identification and quantification of about 50 components, among them 26 in *P. auritum* and 38 in *P. holtonii*, comprising between 82-100% of the total composition of the EOs/HEs. Phenylpropanes were the main components detected in EOs and HEs for both species, being safrole (69.54 and 56.88% in EO and HE) and apiole (64.24 and 57.20% in EO and HE) the major constituents on *P. auritum* and *P. holtonii*, respectively. The phenylpropanes eugenol, myristicin, isosafrole, elemicin, and asarone were also found in low proportion. The presence of safrole in EO of *P. auritum* as major compound is consistent with some previous reports (Gupta et al., 1985). Nevertheless, there are differences regarding the amount of safrole found in *P. auritum*
Table 2. Chemical composition of the essential oils (EOs) and hexanic extracts (HEs) from *Piper auritum* and *P. holtonii*.

| Nº | Compound              | Rt' | GC Area (%) | Mass spectrometry†† | Main fragments ions (m/z,% abundance) |
|----|-----------------------|-----|-------------|----------------------|--------------------------------------|
|    |                       | EO  | HE          | EO                   | HE                                  |
| 1  | p-Cymene              | 8.55| 3.52        | 1.58                 |                                      |
| 2  | α-Terpinene           | 9.10| 0.97        | 5.11                 |                                      |
| 3  | Citral                | 9.38|             | 0.83                 |                                      |
| 4  | Terpinoleno           | 9.96| 1.42        | 1.46                 |                                      |
| 5  | Myrcene               | 10.11|            | 1.46                 |                                      |
| 6  | Linalool              | 10.53|            | 0.98                 |                                      |
| 7  | α-Terpinene           | 11.18|            | 2.61                 |                                      |
| 8  | 5,6-Dihydro-6-pentyl-2H-pyran-2-one | 11.49|         |                      |                                      |
| 9  | Camphor               | 12.95|            |                      |                                      |
| 10 | Eugenol               | 13.53|            | 0.73                 |                                      |
| 11 | 2,4-Decadienoic acid methyl ester | 14.28|     | 0.82                 | 1.31                                 |
| 12 | p-Cymen-8-ol          | 14.82|            |                      |                                      |
| 13 | β-Isoborneol          | 15.16|            | 1.68                 |                                      |
| 14 | α-Amorphene           | 15.75|            | 0.34                 |                                      |
| 15 | Cubenol               | 16.77|            |                      |                                      |
| 16 | α-Copaene             | 17.55|            |                      |                                      |
| 17 | α-Terpinene           | 18.12|            |                      |                                      |
| 18 | Myrcene               | 18.77|            |                      |                                      |
| 19 | Caryophyllene oxide   | 19.08|            |                      |                                      |
| 20 | Safrole               | 19.72|            |                      |                                      |
| 21 | Isoafrole             | 20.11|            |                      |                                      |
| 22 | Spatululenol          | 20.80|            |                      |                                      |
| 23 | α-Caryophyllene       | 21.52|            | 1.04                 | 0.79                                 |
| 24 | 3-Hydroxy-β-damascone | 21.95|           | 0.82                 | 1.31                                 |
| 25 | α-Humulene            | 22.17|            | 1.03                 | 0.39                                 |
| 26 | α-Ionone              | 22.37|            | 0.17                 |                                      |
| 27 | 3-Oxo-α-ionol         | 22.69|            |                      |                                      |
| 28 | α-Cadinene            | 22.91|            |                      |                                      |
| 29 | α-Cubebene            | 23.42|            |                      |                                      |
| 30 | B-Cubebene            | 23.95|            |                      |                                      |
| 31 | 2-Tridecanone         | 24.37|            |                      |                                      |
| 32 | α-Muurolene           | 24.54|            |                      |                                      |
| 33 | Apion                 | 25.41|            |                      |                                      |
| 34 | t-Cadinene            | 25.71|            |                      |                                      |
| 35 | 3-Oxo-β-ionol         | 26.22|            |                      |                                      |
| 36 | β-Cadinene            | 26.63|            |                      |                                      |
| 37 | Calamenene            | 27.14|            |                      |                                      |
| 38 | Palmiticacidmylester  | 27.65|            |                      |                                      |
| 39 | t-Nerolidol           | 28.15|            |                      |                                      |
| 40 | Unknown               | 28.45|            |                      |                                      |
| 41 | Unknown               | 28.75|            |                      |                                      |
| 42 | Palmitic acid         | 29.24|            |                      |                                      |
| 43 | Elemicin              | 29.54|            |                      |                                      |
| 44 | Asarone               | 29.85|            |                      |                                      |
| 45 | α-Cadinol             | 30.15|            |                      |                                      |
| 46 | Phenylpropenes        | 30.45|            |                      |                                      |
| 47 | Monoterpenes          | 30.75|            |                      |                                      |
| 48 | Cyclic                | 31.05|            |                      |                                      |
| 49 | Sesquiterpenes        | 31.35|            |                      |                                      |
| 50 | Monocyclic            | 31.65|            |                      |                                      |
| 51 | Bicyclic              | 31.95|            |                      |                                      |
| 52 | Tricyclic             | 32.25|            |                      |                                      |
| 53 | Others                | 32.55|            |                      |                                      |

- No detected. †Retention times (Rt) in minutes. ††Molecular and fragment ions (m/z) were determined by mass spectrometry, electronic ionization at 70 eV, total ion monitoring was done in a scan mass range from 30-400 m/z, and match comparison with the NIST-MS library. †††Optical rotations were measured in CHCl₃ solution at 20 °C with a JASCO P-2000 digital polarimeter.
According to Castañeda et al. (2007), the EO obtained by hydrodistillation from aerial parts of *P. auritum* presented among 64.0 and 91.4% of safrole. It is well known, however, that the variability on proportion of each component and composition of EO depends on several parameters including geographical origin, vegetative stage of plant, storage condition and extraction method, among others (Chatzopoulou and Katsiotis, 1995).

Minor components such as monoterprenes and sesquiterpenes were also detected. In the EO and HE of *P. auritum* the proportion of monoterprenes was respectively 8.39 and 15.69%, mainly related to cyclic monoterprenes (*p*-cymene, γ-terpinene, terpinolene, α-terpinene, and camphor). Some of these components possess antifungal and synergistic properties (Rattanachaikunsopon and Phumnakhachorn, 2010); furthermore, several studies have shown that monoterprenes cause damage to the microbial cell membrane (Cristani et al., 2007). For *P. holtonii*, however, monoterprenes were not detected. Instead, several sesquiterpenes were found in oil (12.47%) and extract (10.79%). *Piper holtonii* showed the presence of acyclic (*t*-nerolidol), monocyclic (humulene), bicyclic (β-caryophyllene and their oxide), and tricyclic (α-copaene, β-bourbonene, among others) sesquiterpenes. Several fatty acids and their esters, rose ketones (*α*-ionone, 3-oxo-*α*-ionol, 3-hydroxi-β-damascone), and other minor components were classified as “others” comprising between 0.3% and 10-23% of the total composition of EOs and HEs respectively. To the best of our knowledge, this is the first report of the EO/HE composition and antifungal activity of *P. holtonii*. However, some earlier papers have reported cytotoxic, leishmanicidal, and antimalarial activity of extracts and fractions of *P. holtonii* and the isolation of 5-(3',4'-methyleneoxyphenyl)-2E,4E-pentadienoic acid isobutyl amide (Garavito et al., 2006; Correa et al., 2010; Sánchez-Suárez et al., 2010). Overall, HEs from *P. auritum* and *P. holtonii* were more effective against plant pathogenic fungi than DEs, AEs, and MEs, indicating that the potential fungitoxic compounds were present in the low-polarity extracts. These fractions (HEs) possess as major components the phenylpropenes, being the most abundant safrole and apiol. Similarly, EOs have a high proportion of phenylpropanes and show a relatively good antifungal activity against the assayed microorganisms. For these EOs the observed IC₅₀ values were minor in all the assayed microorganisms (Table 3). These results indicate that the antifungal activity of HEs/EOs could be associated to the presence and amount of the phenylpropenes. Safrole has been found in other plants and is the major component of the EO of *Sassafras albidum* (Lauraceae), which has shown antifungal activity.

Table 3. Mean inhibitory concentrations (IC₅₀) of hexanic extracts (HEs), essential oils (EOs), safrole, and apiol against *Botryodiplodia theobromae* and *Colletotrichum acutatum*.

| Fungus        | *Piper auritum* | *Piper holtonii* |
|---------------|----------------|-----------------|
|               | HE | EO | Safrole | HE | EO | Apiol |
| *B. theobromae* | 540.85 | 254.40 | 891.20 | 126.49 | 50-100 | <50 |
| *C. acutatum*  | 766.50 | 364.70 | 349.50 | 139.40 | 70.38 | 36.16 |

Figure 4. In vivo antifungal activity of essential oil (EO), n-hexane extract (HE), and apiol from *Piper holtonii* at 2% in papaya cv. Hawaiian, 7 d post-inoculation with *Colletotrichum gloeosporioides*. (a) and (f) are the untreated control; (b) and (g) are controls treated with 2.5% lecithin; (c) and (h) are fruits treated with EO; (d) and (i) are fruits treated with HE; and (e) and (j) are fruits treated with apiol.
are believed to be associated with the presence of safrole. Against properties of HE/EO from lower than those found for HE or EO. Thus, the antifungal activity/potent antifungal activity against B. theobromae with IC50 μg mL⁻¹ was lower than that from P. auritum. This activity can be associated with the kind of phenylpropene presents in the EO/HE. Thus, some structural requirement could be involved in the antifungal activity.

In view of the results presented by EOs and HEs from P. auritum and P. holtonii, these were selected for a phytochemical study. Bioactive materials (EOs and HEs) were subjected to column chromatography on silica gel leading to isolation of safrole and apiol. Their structures were confirmed by NMR data, including bidimensional analysis and comparison with spectroscopy data reported in the literature (Benevides et al., 1999; Chu et al., 2011). The interpretation of bidimensional spectra indicated that compound obtained from P. holtonii corresponds to isomer 2,3-dimethoxy-4,5-methylenedioxy-1-allylbenzene.

According to the results given in Table 3, both compounds displayed significant antifungal activity. However, apiol was found to be more effective against both fungi as compared to safrole. Apiol exhibited a potent antifungal activity against C. acutatum and B. theobromae with IC50 values < 50 μg mL⁻¹, being 36.2 μg mL⁻¹ for the first. These results are in agreement with previous investigations in which it has been demonstrated that apiol possesses antifungal activity against some agriculturally important fungi (Meepagala et al., 2005). Furthermore, IC50 values for apiol are lower than those of their corresponding HE and EO. These results demonstrate that is possible to think the antifungal effect of HE/EO of P. holtonii is mainly due to the presence of apiol. On the other hand, although safrole exhibited a moderated activity against C. acutatum, the IC50 value (349.5 μg mL⁻¹) was lower than those found for HE or EO. Thus, the antifungal properties of HE/EO from P. auritum against C. acutatum are believed to be associated with the presence of safrole. Remarkably, however, the IC50 value for safrole (891.2 μg mL⁻¹) against B. theobromae was higher than those of their corresponding HE/EO (749.0 and 475.7 μg mL⁻¹). It suggests that the effectiveness of HE/EO from P. auritum against this pathogen could be due to some other minor constituents or synergistic interactions between the constituents of each bioactive material.

It becomes evident that a relationship exists between the structural characteristics of the phenylpropenes and the antifungal activity. The presence of two methoxyl groups in positions 2, 3 of the benzene ring of apiol led to a notorious increase of the antifungal activity when is compared to safrole. These results indicate that lipophilicity and/or electronic and steric factors on phenylpropenes might be important for the antifungal activity. Although not much is known about the mechanism of antifungal action of phenylpropenes, it has been reported that their lipophilic character enables them to alter the fluidity and permeability of cell membranes (Devi et al., 2010). In addition, these compounds inhibit specific cellular processes or enzymes (Wendakoon and Sakaguchi, 1993). Thus, the two methoxyl groups (a bulky electron-donating group) on apiol may lead to a correct balance of hydrophilicity-lipophilicity, which is a determinant factor for the antifungal effect in this compound. On the other hand, evidence of electronic and steric factors in the fungitoxic mechanism of organic compounds has been previously reported (Gershon et al., 2004). According to the results, EO/HE from P. holtonii and apiol were selected as the most promising materials. Their toxicity on fruit peel and antifungal activity in whole fruits were evaluated.

In vivo assay on natural infected papaya fruit
In order to study the in vivo effectiveness of materials as fungistatic agents, papaya fruits inoculated on a wound with C. gloeosporioides were treated with the more promising materials according to in vitro assay: HE, EO from P. holtonii and apiol, at 2%. There was 11, 47, and 54% reduction of the lesion diameter on treatments with HE, EO, and apiol, respectively, when compared to controls (untreated fruits or fruits treated only with lecithin). Compact white mycelial mass surrounded by small water-soaked spots were present over the lesions on the surface of fruits (Figure 4). Internal tissue in the infected area showed a grayish white discoloration which was deeper in both controls. The progress of the lesion into the parenchyma of the fruit was lower in the treatment with apiol followed by EO and HE.

Despite relatively good antifungal activity of the EO/HE and apiol, and low phytotoxicity on the fruit peel, it is necessary further studies on the persistence and toxic effects in order to develop new antifungal agents for the control of these fungi.

CONCLUSIONS
The antifungal activity of EOs and hexane, dichloromethane, ethyl acetate, and methanol extracts obtained from Piper auritum and P. holtonii revealed that only the hexanic extracts and EOs showed a relatively good antifungal activity against C. acutatum, C. gloeosporioides, and B. theobromae. In addition, oil and extract from P. holtonii were more active than those from P. auritum. Furthermore, the antifungal activity observed for the hexanic extract and EO suggests that the active metabolites of both Piper species are mostly lipophilic compounds. Active oils/extracts were found to be rich

(Simic et al., 2004). Meanwhile, apiol is present at high concentration (40 mg g⁻¹ dry seed) in seeds of Ligusticum hultenii. It has shown to possess antifungal properties (Meepagala et al., 2005). On the other hand, it is also possible that some minor components might be involved in some synergistic and/or antagonistic interactions with other active compounds (Gould, 1996). It is noteworthy that although the proportion of safrole and apiol in all materials was close, HE and EO from P. holtonii were the most active against C. acutatum, C. gloeosporioides, and B. theobromae than that from P. auritum. This activity can be associated with the kind of phenylpropene presents in the EO/HE. Thus, some structural requirement could be involved in the antifungal activity.

According to the results given in Table 3, both compounds displayed significant antifungal activity. However, apiol was found to be more effective against both fungi as compared to safrole. Apiol exhibited a potent antifungal activity against C. acutatum and B. theobromae with IC50 values < 50 μg mL⁻¹, being 36.2 μg mL⁻¹ for the first. These results are in agreement with previous investigations in which it has been demonstrated that apiol possesses antifungal activity against some agriculturally important fungi (Meepagala et al., 2005). Furthermore, IC50 values for apiol are lower than those of their corresponding HE and EO. These results demonstrate that is possible to think the antifungal effect of HE/EO of P. holtonii is mainly due to the presence of apiol. On the other hand, although safrole exhibited a moderated activity against C. acutatum, the IC50 value (349.5 μg mL⁻¹) was lower than those found for HE or EO. Thus, the antifungal properties of HE/EO from P. auritum against C. acutatum are believed to be associated with the presence of safrole. Remarkably, however, the IC50 value for safrole (891.2 μg mL⁻¹) against B. theobromae was higher than those of their corresponding HE/EO (749.0 and 475.7 μg mL⁻¹). It suggests that the effectiveness of HE/EO from P. auritum against this pathogen could be due to some other minor constituents or synergistic interactions between the constituents of each bioactive material.

It becomes evident that a relationship exists between the structural characteristics of the phenylpropenes and the antifungal activity. The presence of two methoxyl groups in positions 2, 3 of the benzene ring of apiol led to a notorious increase of the antifungal activity when is compared to safrole. These results indicate that lipophilicity and/or electronic and steric factors on phenylpropenes might be important for the antifungal activity. Although not much is known about the mechanism of antifungal action of phenylpropenes, it has been reported that their lipophilic character enables them to alter the fluidity and permeability of cell membranes (Devi et al., 2010). In addition, these compounds inhibit specific cellular processes or enzymes (Wendakoon and Sakaguchi, 1993). Thus, the two methoxyl groups (a bulky electron-donating group) on apiol may lead to a correct balance of hydrophilicity-lipophilicity, which is a determinant factor for the antifungal effect in this compound. On the other hand, evidence of electronic and steric factors in the fungitoxic mechanism of organic compounds has been previously reported (Gershon et al., 2004). According to the results, EO/HE from P. holtonii and apiol were selected as the most promising materials. Their toxicity on fruit peel and antifungal activity in whole fruits were evaluated.

In vivo assay on natural infected papaya fruit
In order to study the in vivo effectiveness of materials as fungistatic agents, papaya fruits inoculated on a wound with C. gloeosporioides were treated with the more promising materials according to in vitro assay: HE, EO from P. holtonii and apiol, at 2%. There was 11, 47, and 54% reduction of the lesion diameter on treatments with HE, EO, and apiol, respectively, when compared to controls (untreated fruits or fruits treated only with lecithin). Compact white mycelial mass surrounded by small water-soaked spots were present over the lesions on the surface of fruits (Figure 4). Internal tissue in the infected area showed a grayish white discoloration which was deeper in both controls. The progress of the lesion into the parenchyma of the fruit was lower in the treatment with apiol followed by EO and HE.

Despite relatively good antifungal activity of the EO/HE and apiol, and low phytotoxicity on the fruit peel, it is necessary further studies on the persistence and toxic effects in order to develop new antifungal agents for the control of these fungi.

CONCLUSIONS
The antifungal activity of EOs and hexane, dichloromethane, ethyl acetate, and methanol extracts obtained from Piper auritum and P. holtonii revealed that only the hexanic extracts and EOs showed a relatively good antifungal activity against C. acutatum, C. gloeosporioides, and B. theobromae. In addition, oil and extract from P. holtonii were more active than those from P. auritum. Furthermore, the antifungal activity observed for the hexanic extract and EO suggests that the active metabolites of both Piper species are mostly lipophilic compounds. Active oils/extracts were found to be rich
in phenylpropenes being safrole and apiol the major compounds from *P. auritum* and *P. holtonii*, respectively. Apiole can effectively inhibit the mycelial growth of the evaluated pathogenic fungi. In addition, the application of active oils/extracts and their major constituents on the peel of avocado and tamarillo fruits show no apparent damages, or odor or taste changes. So, EOs and hexanic extracts or their major compounds could become a natural and effective alternative to synthetic fungicides for use in the treatment of plant pathogens and develop of novel types of selective and natural fungicides.

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**Composición química y actividad antifúngica de *Piper auritum* Kunth y *Piper holtonii* C. DC. contra hongos fitopatogénicos.** El crecimiento de hongos durante pre- y poscosecha de frutas puede causar el deterioro y resultar en una reducción en la calidad y cantidad. El objetivo del presente trabajo fue investigar la eficiencia de cuatro extractos (n-hexano, diclorometano, acetato de etilo, y metanol) y el aceite esencial de *Piper auritum* y *P. holtonii* C. DC. en la inhibición del crecimiento de tres importantes patógenos de pre- y poscosecha de frutas (*Colletotrichum acutatum*, *C. gloeosporioides* y *Botryodiplodia theobromae*). La actividad antifúngica in vitro se evaluó siguiendo la técnica del alimento envenenado. Los resultados mostraron que tanto el extracto en n-hexano (HE) como el aceite esencial (EO) exhibieron un relativo buen control con ambos patógenos, siendo los materiales provenientes de *P. holtonii* los más activos. El crecimiento micelial de *C. acutatum*, *C. gloeosporioides* y *B. theobromae* se inhibió significativamente a 400 μg mL⁻¹. Adicionalmente, se analizó la composición química de los materiales bioactivos mediante cromatografía de gases-espectrometría de masas (CG-EM). Safrole (64.54/56.88%) y apiole (64.24/57.20%) fueron los componentes mayoritarios del EO/HE para *P. auritum* y *P. holtonii*, respectivamente. La identificación estructural fue adicionalmente confirmada por resonancia magnética nuclear de 1H. Ambos compuestos exhibieron propiedades antifúngicas significativas. Se puede concluir que EOs/HEs de *P. auritum* y *P. holtonii*, y sus componentes mayoritarios tienen interesantes aplicaciones para el control de hongos patogénicos de plantas.

**Palabras clave:** *Piperacea*, aceite esencial, apiole, *Colletotrichum*, *Botryodiplodia*.

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