

# Anti-*Candida albicans* activity, cytotoxicity and interaction with antifungal drugs of essential oils and extracts from aromatic and medicinal plants

## Actividad contra *Candida albicans*, citotoxicidad e interacción con antifúngicos de aceites esenciales y extractos de plantas medicinales y aromáticas

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### Abstract

**Objective:** To determine anti-*Candida albicans* activity, cytotoxicity and drug interaction of essential oils and extracts from plants collected in Colombia.

**Materials and methods:** The antifungal activity was evaluated following the AFST-EUCAST protocol. With most active samples, the inhibition of the formation of germ tubes and budding, the *in vitro* pharmacodynamics, using time-kill assays, and the interaction with itraconazole and amphotericin B following the checkerboard technique were evaluated. The cytotoxicity assay for all samples was done using MTT.

**Results:** Strong activity in 17.57% of the samples was found. The lowest MIC values were obtained with *Piper bredemeyeri* Jacq and *Lippia origanoides* Kunth (B) oils and *Morinda royoc* L extract. The three samples inhibited the formation of germ tubes and budding. *P. bredemeyeri* Jacq oil and *M. royoc* L extract samples showed fungicidal activity at 2xMIC. A synergistic effect was obtained with the combination of *P. bredemeyeri* Jacq oil and itraconazole, but not for the combination with amphotericin B. Active samples against *C. albicans* were not cytotoxic on Vero cells ATCC CCL-81, excluding *P. bredemeyeri* Jacq oil.

**Conclusions:** The results of this study suggest that Colombian medicinal and aromatic plants represent an untapped source of compounds with anti-*C. albicans* activity that could be a resource in the development of new therapeutic natural products.

**Key words:** Essential oils, extracts, *Candida albicans*, antifungal activity, cytotoxicity, synergism

### Resumen

**Objetivo.** Determinar la actividad anti-*Candida albicans*, la citotoxicidad y la interacción con antifúngicos de aceites y extractos de plantas recolectadas en Colombia.

**Materiales y métodos.** La actividad antifúngica fue evaluada siguiendo el protocolo *Antifungal Susceptibility Testing Subcommittee of the European Committee on Antimicrobial Susceptibility Testing (AFST-EUCAST)*. Con las muestras más activas se evaluó la inhibición de la formación de tubo germinal y la gemación, la farmacodinamia mediante curvas de tiempo muerte y la interacción con itraconazol y anfotericina B. Se determinó la citotoxicidad mediante la técnica MTT.

**Resultados.** Se encontró actividad en 17,57 % de las muestras. La mayor actividad se obtuvo con los aceites de *Piper bredemeyeri* Jacq y *Lippia origanoides* Kunth (B) y el extracto de *Morinda royoc* L. Las tres muestras inhibieron la formación de tubo germinal y la gemación. El aceite de *P. bredemeyeri* Jacq y el extracto de *M. royoc* L mostraron actividad fungicida con dos veces la concentración inhibitoria mínima. Se encontró un efecto sinérgico por la combinación del aceite de *P. bredemeyeri* Jacq e itraconazol, pero no con anfotericina B. Las muestras activas no fueron citotóxicas, excepto el aceite de *P. bredemeyeri* Jacq.

**Conclusión.** Los resultados de este estudio sugieren que las plantas de Colombia son una fuente no explorada de compuestos con actividad anti-*C. albicans*, útiles para el desarrollo de nuevos productos terapéuticos.

**Palabras clave:** aceites esenciales, extractos, *Candida albicans*, actividad antifúngica, citotoxicidad, sinergismo.

### Introduction

Many researchers, particularly the ones from countries with a rich biodiversity, have contributed to the detection of new antifungal com-

pounds in medicinal plants. Screening by using *in vitro* evaluation is a useful tool for the discovery of new potential antifungal agents from natural products such as essential oils and extracts derived from plants <sup>(1)</sup>. Colombia is the second ri-

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Recibido: 08/10/2010; Aceptado: 03/08/2011

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chest country in the world in *biodiversity*, and its floral diversity is estimated at 40,000 species of vascular plants <sup>(2)</sup>. Although Colombia possesses a rich tradition in the use of medicinal plants, the antifungal activity of medicinal plants' derivatives has not been deeply studied.

Candidiasis is a common infection of the skin, nails, oral cavity, esophagus, and vagina, caused by yeast of the *Candida* genus. Systemic yeast infections are a common consequence of immunosuppression, long-term indwelling catheters, and endocrinopathies. *Candida albicans* is the most common pathogen causing that fungal opportunistic infection. Additionally, it has the ability to adhere to host surfaces or to prosthesis leading to the formation of biofilms which further facilitate adhesion, infection and resistance to the antifungals <sup>(3)</sup>.

In Colombia, different studies have shown the importance of *C. albicans* as the principal agent causing bloodstream fungal infection (44.7%) in tertiary care level hospitals <sup>(4)</sup>, and as cause of invasive infection (43.6% and 54.5%) <sup>(5,6)</sup>.

The number of clinical infections worldwide by *C. albicans* has risen considerably in recent years, and the incidence of resistance to traditional antifungal therapies is also increasing <sup>(7)</sup>. In addition, drug-related toxicity, significant drug interactions and insufficient bioavailability of the conventional antifungals, have encouraged the search for new alternatives among natural products <sup>(8)</sup>.

The aim of this study was to evaluate anti-*C. albicans* activity and cytotoxicity of essential oils and plant extracts obtained from aromatic and medicinal plants of different Colombian regions. Furthermore, the combined effects of itraconazole and amphotericin B with the most active samples, and their pharmacodynamics were evaluated *in vitro* by the checkerboard method and the time-kill curves, respectively.

## Materials and methods

### Plant material

Stems and leaves of 74 plants were collected in different regions of Colombia, from 2005 to 2008, as part of a survey conducted by CENIVAM, a research center devoted to the study of aromatic plants and essential oils in Colombia. The taxonomic identification of the botanical samples was performed by José Luis Fernandez at the Herbario Nacional de Colombia, Instituto de Ciencias Naturales, Facultad de Ciencias, Universidad Nacional de Colombia at Bogotá, where voucher specimens were deposited.

The selected plants belong to the following genera: *Lippia* (Verbenaceae) (26); *Salvia* (Lamiaceae) (7); *Lepechinia* (Lamiaceae) (3); *Piper* (Piperaceae) (3); *Baccharis* (Asteraceae) (2); *Hedyosmum* (Chloranthaceae) (3); *Illicium* (Schisandraceae) (2); *Cymbopogon* (Poaceae) (3); *Minthostachys mollis* (Lamiaceae) (2); *Thymus vulgaris* (Lamiaceae) (2); and *Rosmarinus officinalis* (Lamiaceae) (2). Additionally, the following plants were also selected: *Cananga odorata* (Annonaceae), *Tagetes lucida* (Asteraceae), *Eucalyptus citriodora* (Myrtaceae), *Turnera* aff. *diffusa* Willd. ex Schult (Turneraceae), *Nectandra tomentella* (Lauraceae), *Sigesbeckia agrestis* (Asteraceae), *Lantana fucata* Lindl (Verbenaceae), *Hyptis perbullata* (Lamiaceae), *Origanum vulgare* (Lamiaceae), *Zingiber officinale* (Zingiberaceae), *Cascarilla saravena* (Euphorbiaceae), *Achyrocline alata* (Kunth) DC (Asteraceae), *Pimenta racemosa* (Myrtaceae), *Elettaria cardamomum* L (Zingiberaceae), *Aloysia triphylla* (Verbenaceae), *Curcuma monteria* (Zingiberaceae), *Chenopodium ambrosioides* L (Amaranthaceae), *Bursera graveolens* (Kunth) Triana & Planch (Burseraceae), and *Morinda royoc* L (Rubiaceae).

### Extracts and essential oils extraction

Essential oils (59) and extracts (15) were evaluated. The essential oils were extracted from dried stems and leaves (300 g) by microwave-assisted hydrodistillation as described <sup>(9)</sup>. The extracts

were obtained from 40 g of dried leaves from each plant, macerated with 200 ml ethanol and left in suspension for 7 days at room temperature (28 °C). The mixture was filtered and concentrated using a Buchi rotavapor. Stock solutions of both oils and extracts, were prepared in DMSO ( $\leq 1\%$  v/v), for subsequent bioassays.

### **Antifungal susceptibility testing**

The Minimum Inhibitory Concentrations (MIC) of all samples were determined by the Antifungal Susceptibility Testing Subcommittee of the European Committee on Antibiotic Susceptibility Testing (AFST-EUCAST) protocol <sup>(10)</sup>. *Candida albicans* ATCC 10231, *C. albicans* ATCC 90028 and a clinical blood isolate randomly selected from a group of clinical isolates of the *Candida* species collected in our laboratory were used at inoculum size of  $1-5 \times 10^5$  CFU/ml. The oils and extracts were evaluated at five concentrations of 31.25-500  $\mu\text{g/ml}$ . Oils and extracts were considered active when they exhibited MIC values equal or greater than 500  $\mu\text{g/ml}$ . Itraconazole (Sigma-Aldrich, Co., MO, USA) was used as a positive control at a range of 0.031-16  $\mu\text{g/ml}$  with the strains *C. krusei* ATCC 6258 and *C. parapsilosis* ATCC 22019. A negative control (inoculum without treatment) was also included. MIC values were expressed as a geometric mean (GM-MIC) of tests performed in duplicate in three different assays against each *Candida* strain and the clinical isolate.

### **Effect on germ tube formation and budding**

The effect on germ tube formation and budding was evaluated according to Ishida, *et al.* <sup>(11)</sup>. Different concentrations (31.25-500  $\mu\text{g/ml}$ ) of *P. bredemeyeri* Jacq, *L. origanoides* Kunth (B) oils and *M. royoc* L extract were tested with the three yeasts. A suspension of  $1-5 \times 10^5$  CFU/ml in RPMI of each yeast was prepared, and 100  $\mu\text{l}$  were added to the same volume of each sample concentration. Positive and negative controls with

nystatin at 6.25  $\mu\text{g/ml}$  (La Santé, Bogotá, Colombia) and cells without treatment, respectively, were included.

For germ tube assays, after two hours of incubation at 35 °C, the absence or presence of germ tubes was determined using a light microscopy (Nikon, Eclipse E200, Japan). Each sample was tested in duplicate using two different assays. To evaluate the effect on budding, after three hours of incubation at 35 °C, the budding cells/ml were counted using a hemacytometer.

The statistical analysis of the effect on budding, was performed by R version 2.9.1 (The R Foundation for Statistical Computing, ISBN 3-900051-07-0) using ANOVA after transforming the response variable with square root. The Tukey multiple comparison test was used to compare strains with and without treatment. A 0.05 significance level was employed.

### **Time-kill assay**

The *in vitro* pharmacodynamics of *P. bredemeyeri* Jacq oil and *M. royoc* L extract against *C. albicans* ATCC 90028 were performed by the method described by Klepser, *et al.* <sup>(12)</sup>. An initial inoculum ranging from  $1-5 \times 10^5$  CFU/ml was seeded on each sample, at concentrations of 0.25, 0.5, 1 and 2 times the MIC. The samples were incubated at 35 °C with agitation. At 0, 4, 8, 12, and 24 hours, volumes of 10  $\mu\text{l}$  were then spread onto potato dextrose agar (Oxoid, Basingstoke, Hampshire, UK) and incubated at 37 °C for 24 hours to determine the number of CFU/ml. The limit of detection was 100 CFU/ml. Time-kill curves with itraconazole and amphotericin B (Sigma-Adrich, Co, MO, USA) were used as fungistatic and fungicide controls, respectively. Experiments were carried out in duplicate in two separate experiments. Time-kill curves were constructed by plotting of mean  $\pm$  standard deviation (SD) of colony count ( $\log_{10}$  CFU/ml) as a function of time (hours) with the Prism<sup>TM</sup> (GraphPad Software, Inc., USA, 2007) sta-

tistical package. Fungicidal activity was defined as  $\geq 3 - \log_{10}$  ( $\geq 99.9\%$ ) reduction in CFU/ml from the starting inoculum ( $0.5 - 2.5 \times 10^5$  CFU/ml).

### Interaction of essential oils and drugs

Assays were performed using the checkerboard method<sup>(13)</sup>. *Candida albicans* ATCC 10231 was used at a final concentration of  $0.5-2.5 \times 10^5$  CFU/ml. The final concentrations of the itraconazole and amphotericin B ranged from  $0.004 \mu\text{g/ml}$  to  $2 \mu\text{g/ml}$  and the essential oil of *P. bredemeyeri* Jacq from  $7.813$  to  $500 \mu\text{g/ml}$ . The fractional inhibitory index (FICI) was calculated, and the interpretation was determined as follows:  $\leq 0.5$ , synergistic effect;  $>0.5$  but  $<4$  no interaction; and  $\geq 4$  antagonistic effect<sup>(13)</sup>.

### Cytotoxicity assay

*Cercopithecus aethiops* African green monkey kidney cells (Vero cell line ATCC CCL-81) were used. The cells were grown in Eagle's minimum essential medium (MEM) supplemented. The cytotoxicity of the essential oils and their components was examined *in vitro* using an MTT (dimethylthiazol-2-yl]-2,5-diphenyl tetrazolium bromide) (Sigma, New Jersey, USA) assay as described by Betancur-Galvis, *et al.*<sup>(14)</sup>. The minimal dilution of the essential oils that induced 50% growth inhibition of the cells was expressed as inhibitory concentration 50% ( $IC_{50}$ ). The  $IC_{50}$  values for each compound were obtained by linear regression analysis of the dose-response curves generated from the absorbance data with the R (Development Core Team, Vienna, Austria, 2008) statistical package.  $IC_{50}$  values were expressed as the mean  $\pm$  standard deviation ( $M \pm SD$ ) of two independent experiments done in quadruplicate.

## Results

The MICs of 13 active samples (GM-MIC range  $157.5-500 \mu\text{g/ml}$ ) tested against *C. albicans* and the respective  $IC_{50}$  values, are presented in Table 1. The most active samples were the oils from *P.*

*bredemeyeri* Jacq (GM-MIC of  $157.5$ ,  $176.8$  and  $222.7 \mu\text{g/ml}$  for *C. albicans* ATCC 10231, *C. albicans* ATCC 90028 and the clinical isolate, respectively) and *L. origanoides* Kunth (B) (GM-MIC of  $157.5$ ,  $157.5$ ,  $198.4 \mu\text{g/ml}$  for *C. albicans* ATCC 10231, *C. albicans* ATCC 90028, and the clinical isolate, respectively). Also *M. royoc* L extract was active (GM-MIC =  $250 \mu\text{g/ml}$ ) with the three evaluated yeasts (Table 1). The MIC values of the positive control itraconazole with *C. parapsilosis* ATCC 22019 and *C. krusei* ATCC 6258 ( $0.169$  and  $0.999$ , respectively), were in the acceptable range according to the standard protocol ( $0.3125-0.250$  and  $0.0625-0.250 \mu\text{g/ml}$ , respectively).

*Piper bredemeyeri* Jacq and *L. origanoides* Kunth (B) oils inhibited the formation of germ tubes in the two strains and in the clinical isolate at a  $\geq 31.25 \mu\text{g/ml}$  concentration. Also, *M. royoc* L extract inhibited the formation of germ tube in both: *C. albicans* ATCC 90028 strain and in the clinical isolate, at the same concentrations. The inhibition in *C. albicans* ATCC 10231 was at a  $\geq 125 \mu\text{g/ml}$  concentration. Germ tubes were observed in the negative control, but not in the positive control. In addition, the treatment of the strains and the clinical isolate with *P. bredemeyeri* Jacq and *L. origanoides* Kunth (B) oils, and *M. royoc* extract, inhibited the process of budding, at  $250 \mu\text{g/ml}$ . The decrease of budding in treated blastoconidia, compared with the negative control was significantly different ( $p < 0.05$ ). The positive control inhibited the budding, and in the negative control there was a normal rate of budding.

The killing activity of *P. bredemeyeri* Jacq oil and *M. royoc* L extract against *C. albicans* ATCC 90028 as well as itraconazole and amphotericin B is represented in Figure 1. At 2 times the MIC, *P. bredemeyeri* Jacq oil, after 4 hours of incubation, showed fungicidal activity similar to amphotericin B (Figure 1a and Figure 1c, respectively). It was found that at 0.5 times and 1 time the MIC concentration there was a reduction in growth until approximately 8 hours, when colony counts continued to grow near the initial starting inoculum. *M. royoc* L ex-

tract showed fungicidal activity at 2 times the MIC after 8 hours of incubation (Figure 1b). At lower concentrations (0.25, 0.5 and 1 times the MIC), we did not find significant reduction in CFU/ml ( $<3 - \log_{10}$ ) compared with the starting inoculum.

A synergistic effect was obtained for the combination of itraconazole and *P. bredemeyeri* Jacq (FICI range 0.09-0.13), but no interaction was detected for the combination of *P. bredemeyeri* Jacq and amphotericin B (FICI=1.06).

According to the American National Cancer Institute (USA) criteria samples with anti-*C. albicans* activity were not cytotoxic on Vero cells excluding *P. bredemeyeri* Jacq oil ( $IC_{50}=15.2 \pm 3$ ) (Table 1).

## Discussion

Plants, marine organisms, and microbes usually produce biologically active compounds as a defense against predators and competition with neighbors. Thus, it seems logical that most of the drugs derived from natural sources have anticancer or anti-infective properties. Important antifungal agents such as polyenes, echinocandins/pneumocandins, aureobasidins and sordarins had their origin in natural products <sup>(15)</sup>.

In this study, oils and extracts from medicinal and aromatic plants were tested against two *C. albicans* strains and a clinical isolate. Currently, there is no agreement on the acceptable level of activity for natural products when they are compared to standard drugs. Aligiannis, et al.<sup>(16)</sup>, suggested a classification of antifungal activity in plant derivatives (antifungal activity based on MIC results) as follows: strong inhibitors, MIC up to 0.5 mg/ml; moderate inhibitors, MIC between 0.6 and 1.5 mg/ml; weak inhibitors, MIC above 1.6 mg/ml. According to these criteria, we found strong anti-*C. albicans* activity in 17.57% of the evaluated samples (Table 1).

Previous studies have shown the antifungal activity in extracts and oils of plants belonging to *Piper*, *Morinda*, and *Lippia* genus <sup>(17-19)</sup>. López, et al.<sup>(19)</sup>, determined a MIC value of 100.000 µg/ml against *C. albicans* of the methanolic extract from the plant *Piper lanceafolium*, using an agar diffusion method. In contrast, in this study, the microdilution standard method AFST-EUCAST was used, and lower MICs with *P. bredemeyeri* Jacq were found (GM-MIC of 157.5, 176.8, and 222.7 µg/ml, with the three evaluated yeasts) (Table 1).

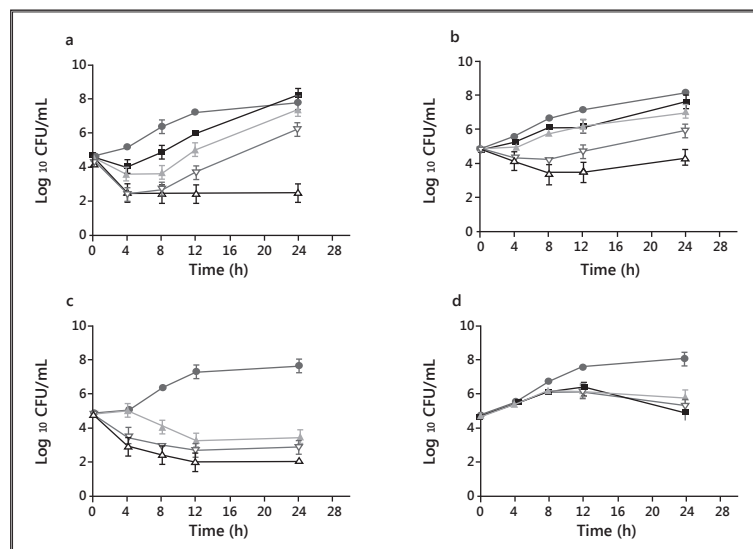


Figure 1. Time-kill plots against *C. albicans* ATCC 90028 of a) *P. bredemeyeri* Jacq oil, b) *M. royoc* L y *P. bredemeyeri* Jacq extract, c) amphotericine B and d) itraconazole. ● Growth control; ■ 0.25x MIC; ▲ 0.5x MIC; ▼ 1x MIC, Δ 2X MIC; □ 4x MIC



Anti-*Candida albicans* effect, cytotoxicity and interaction with antifungal drugs of essential oils and extracts from aromatic and medicinal plants

**Table 1.** Geometric means of minimal inhibitory concentration (GM – MIC, µg/ml) and inhibitory concentration 50% (IC<sub>50</sub>) of essential oils and extracts actives against *Candida albicans*

Plant	Voucher specimen	Sample	GM - MIC (µg/ml)			IC50 (M ± SD)	R2
			<i>C. albicans</i> ATCC 10231	<i>C. albicans</i> ATCC 90028	<i>C. albicans</i> (Clinical isolate)		
<i>Lippia alba</i> 007	484650	Oil	*	353.6	420.4	32.8 ± 3.6	0.89
<i>Lippia origanoides</i>	512271	Oil	396.9	500	500	52.3 ± 11.5	0.8
<i>Lippia origanoides</i>	512075	Oil	250	250	315	74.6 ± 16.9	0.75
<i>Lippia origanoides</i>	512075	Oil	500	500	500	60.4 ± 11.2	0.83
<i>Piper bredemeyeri</i> Jacq.	516939	Oil	157.5	176.8	222.7	15.2 ± 3	0.81
<i>Turnera aff. diffusa</i> Willd. ex Schult.	516293	Oil	*	353.6	500	52.2 ± 5.2	0.93
<i>Lippia origanoides</i> Kunth (A)	517741	Oil	500	500	500	31.4 ± 5.6	0.75
<i>Lippia origanoides</i> Kunth (B)	517741	Oil	157.5	157.5	198.4	31.4 ± 5.6	0.75
<i>Lippia origanoides</i> Kunth (C)	517741	Oil	500	396.9	396.9	34.4 ± 5.9	0.76
<i>Morinda royoc</i> L.	512222	Extract	250	250	250	≥200	NA
<i>Piper hispidum</i> Sw.	519969	Oil	250	280.6	250	51.7 ± 9.3	0.8
<i>Lippia origanoides</i>	512087	Oil	500	500	500	104.4 ± 5.9	0.98
<i>Cymbopogon citratus</i>	531013	Oil	*	500	500	≥200	NA

\* MIC > 500 µg/ml; R<sup>2</sup>: linear regression coefficient; NA: not applicable.

This is one of the few studies focused on evaluating the anti-*C. albicans* activity of *L. origanoides* oils using a standard method for determination of MIC by broth dilution of fermentative yeast. Oliveira, *et al.*<sup>(20)</sup>, also found important activity against *C. albicans* in an oil of *L. origanoides* using an agar diffusion method; however, we cannot compare our results with theirs, because they used a technique that determined inhibition zone (mm) but not MIC (µg/ml).

*Candida albicans* is a dimorphic fungus, able to grow as yeast or filamentous form. The yeast to hyphal transition begins with the formation of a germ tube and it is the initial stage of hyphal formation<sup>(21)</sup>. The oils from *P. bredemeyeri* Jacq and *L. origanoides* Kunth (B) and the *M. royoc* L extract inhibited the germ tube formation. It is possible that those oils and extracts are acting on an important process in the morpho-transformation of *C. albicans*, as wall or membrane integrity or reorganization of the cytoskeleton<sup>(22, 23)</sup>. This characteristic could make a new antifungal more selective towards the infecting germ than towards the host cells.

The concentration at which *P. bredemeyeri* Jacq and *L. origanoides* Kunth (B) oils were able to inhibit the germ tube formation of *C. albicans* (≥31.25 µg/ml or ≥125 µg/ml) are within the concentration ranges published for this activity with *Melaleuca alternifolia* oil (tea tree oil) (0.25 and 12.5% (v/v, approximately 2.25 and 112.5 µg/ml, respectively)<sup>(24)</sup>. Those results make our findings promising if we consider that tea tree oil is used as commercial product for treatments of fungal infections as oropharyngeal and vaginal infections by *C. albicans*<sup>(24-26)</sup>.

*Candida albicans* has isotropic growth by budding and several antifungal drugs and plant extracts can inhibit budding of yeast cells<sup>(21, 27)</sup>. In this study the treatment of the strains and the clinical isolate with *P. bredemeyeri* Jacq and *L. origanoides* Kunth (B) oils, and *M. royoc* extract, decreased blastoconidia budding at 250 µg/ml compared with the negative control (p < 0.05). Previous studies have suggested that the deleterious effect of the essential oil on the cell wall of the fungus could be the main reason for the decrease in the rate of yeast budding, because the cell wall is necessary for cell division<sup>(27)</sup>.

The time-kill studies are useful for the evaluation of the pharmacodynamics' characteristics of new antimicrobial agents<sup>(28)</sup>. The oil of *P. bredemeyeri* Jacq and the extract of *M. royoc* L showed a fungicidal activity on *C. albicans* at 2 x MIC after 4 and 8 hours, respectively. It is clinically more important to find fungicidal compounds than fungistatic, particularly in HIV patients, because the prophylactic use of fungistatic drugs has been associated with an increased frequency of innate or acquired drug resistance in clinical isolates<sup>(29)</sup>.

Chemical analyses of most active essential oils against *C. albicans* have been carried out in our laboratory. The main compounds of *P. bredemeyeri* Jacq oil were terpenes *alpha-pinene* (20.3%) and *beta-pinene* (32.3%) (unpublished data). Other studies have demonstrated that those terpenes act on cellular integrity, inhibition of the respiration and ion transport processes, and increase membrane permeability in *C. albicans*<sup>(30, 31)</sup>. Hence, it is possible that the antifungal activity of *P. bredemeyeri* Jacq oil could be explained by a higher concentration of those *monoterpene hydrocarbons*. On the other hand, chemical analyses of oil from *L. organoides* Kunth (B) have also been carried out in our laboratory. The main components were the oxygenated monoterpenes *thymol* (43.8%) and *carvacrol* (17.3%). The antimicrobial activities of those terpenes have also been demonstrated, and their mechanism of action has been associated with membrane permeability<sup>(20)</sup>.

To the best of our knowledge, this is the first time that antifungal activity of *M. royoc* L extract is described. *Morinda* genus contains substantial amounts of anthraquinones, especially in the roots. The antifungal activity of those quinones has been demonstrated<sup>(32)</sup>, so it is possible that anti-*C. albicans* activity of *M. royoc* L extract may be produced by the presence of such molecules.

In our laboratory, the antifungal activity of *L. organoides* Kunth (B) and *M. royoc* L extract against *Aspergillus fumigatus*, *A. flavus*, *C. krusei*

and *C. parapsilosis* has been demonstrated. Unlike *P. bredemeyeri* Jacq oil only showed activity against *C. krusei* (unpublished data).

Potential synergy of essential oils with antibiotics has been previously considered with the aims of increasing the rate of fungal killing, shortening the duration of therapy, avoiding the emergence of drug resistance, expanding the spectrum of activity, and decreasing drug-related toxicity by allowing lower doses of antifungal agents to be administered<sup>(33)</sup>. In this study, a synergistic effect was obtained when itraconazole and the oil from *P. bredemeyeri* Jacq were combined (FICI range, 0.09-0.13). No interaction was detected with the combination of the oil of *P. bredemeyeri* Jacq and amphotericin B (FICI=1.06).

The criteria of cytotoxic activity for the crude extracts as established by the American National Cancer Institute (USA), is an  $IC_{50}$  of less than 30  $\mu\text{g/ml}$ <sup>(34)</sup>. According to these criteria, we consider that oils with anti-*C. albicans* activity and *M. royoc* L extract were not cytotoxic on Vero cells ( $IC_{50} \geq 200 \mu\text{g/ml}$ ) excluding the oil of *P. bredemeyeri* Jacq ( $IC_{50} = 15.2 \pm 3 \mu\text{g/ml}$ ) (Table 1). González, et al.<sup>(35)</sup>, orally administered an extract of *M. royoc* in rats and they did not find toxic effects. Both our results of *in vitro* cytotoxicity, as well as the toxicity *in vivo* demonstrated by González, et al.<sup>(35)</sup>, suggest that *M. royoc* L extract may be a candidate for developing an antifungal of natural origin against *C. albicans*.

The discovery of a novel natural product for therapeutic use is a slow process. For example, Taxol, an anticancer agent developed from the plant *Taxus brevifolia*, was discovered after a random screening of 35,000 plant samples that took more than 25 years<sup>(36, 37)</sup>.

In conclusion, the results of this study suggest that Colombian medicinal and aromatic plants represent an untapped source of compounds with anti-*C. albicans* activity that could be a resource in the development of therapeutically natural products.

## Acknowledgements

Grant RC-245-2011 (Fondo Nacional de Financiamiento para la Ciencia, la Tecnología y la Innovación, Francisco José de Caldas).

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