Antifungal and post-antifungal effects of chlorhexidine, fluconazole, chitosan and its combinations on *Candida albicans*

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Abstract

Objective: The aim of this work was to assess the antifungal and post-antifungal effects of chlorhexidine, fluconazole, chitosan and its combinations on virulence factors of *Candida albicans*.

Study Design: Ten isolated strains of *Candida albicans* obtained from 10 patients with oral candidiasis and a collection strain of *C. albicans* were treated with antifungal agents in different concentrations or combinations of them. Virulence factors analyzed were the cell surface hydrophobicity, the germinative tube development, the phospholipase activity and the post-antifungal effect of that exposure.

Results: Virulence factors of the isolated strains obtained from patients together with the collection strain showed significant decreases with the different antifungal treatments, except for hydrophobicity and phospholipase activity. The development of germinative tube was the most sensitive factor to all the antifungal agents used. Untreated strains as well as the ones treated with antifungal agents showed a positive correlation among the virulence factors analyzed. No synergic effects arose from the combinations of the used drugs.

Conclusions: *C. albicans* isolated strains from patients showed high phospholipase activity and germinative tube production, which corroborates their capacity to infect the oral mucosa and the high prevalence of species. As a whole, our results imply that short exposures to sub-inhibitory concentrations of the antifungal agents under analysis, isolated or combined, can modulate the way virulence factors get manifested, thus decreasing their pathogenicity.

**Key words:** Candida albicans, chitosan, chlorhexidine, fluconazole, antifungal effect.
Introduction
Candida genus is composed of more than 150 different species, but *Candida albicans* is the main responsible species for oral candidiasis. An increase in the percentage of colonized people has been observed according to their age and the presence of predisposing local or general factors (1,2). The first stage of the infection includes adhesion to cellular surfaces. Adhesion of yeast has been studied using buccal epithelial cells, different cell lines, and plastic materials (3,4), with the cell surface hydrophobicity of fungus (CSH) as the main force implied in this process (5). Besides, *C. albicans* is capable of secreting exoenzymes, such as phospholipase (PL), and of developing hyphae, which allows penetration in oral tissues and, in turn, the infection process (6,7). Another form of measuring the yeast virulence is through the post antifungal effect (PAFE), which measures the growth recovery capacity after a limited exposure to antifungal agents (8).

In recent years, an increase of resistance to some antifungal agents has been observed and this has caused concern due to the eukaryotic character of Candida as well as of host cells, and due to a smaller number of available antifungal drugs compared to antibiotics (9). Some clinical research has used some chlorhexidine-based (CLX) oral rinses as an alternative to antifungal agents used regularly, obtaining good results (10); while other studies have shown an increase of pathogenicity in strains treated with fluconazole (FLZ) in a model of systemic infection in rats (11). Other authors have shown an important FLZ effect on the capacity of adhesion and production of exoenzymes (12). One of the new biomaterials being used in Dentistry is chitosan (HMWC), a cationic polymer obtained from chitin (13). The effect on inhibition of *C. albicans* fungal adhesion to epithelial cells and to hydroxyapatite pearls has been demonstrated in vitro, as well as its use as a vehicle or adjuvant in antifungal action (3, 14-16). Seyfarth et al. showed an existing positive correlation between chitosan molecular weight and its effectiveness as an antifungal agent in an in vitro study (17). In this sense, the combination of treatments could reduce the employed dose, thus preventing the development of antifungal resistance in the therapeutic treatment of this pathology.

In this work, antifungal and post-antifungal effects of CLX, FLZ, HMWC and its combinations were assessed on some virulence factors of *C. albicans* after a short exposure to these drugs in sub-inhibitory concentrations. Besides, the existing correlation between the mentioned factors was analyzed in the presence and in the absence of the antifungal agents under study.

Materials and Methods
Isolation and identification of 10 isolated wild *C. albicans* strains
The strains were isolated from the saliva of 10 patients with oral candidiasis, having complete or partial dentures, who were assisted at Cathedral of Prosthodontics III, Faculty of Dentistry, National University of Córdoba, Argentina; a collection strain was isolated as well (*C.albicans* serotype A NCPF 3153 - CA 3153 - kindly provided by Dr. Jose Ponton, University of the Basque Country, Spain). The patients agreed to sign a consent form prior to their participation in the study. Patients who were under treatment with antifungal drugs, xerostomic drugs, or under a prolonged treatment with anti-inflammatory drugs 10 days prior to exam were excluded. The protocol was approved by the Ethics Committee, Polo Hospitalario, Córdoba, Argentina. Samples were harvested in Sabouraud glucose agar (SGA) with 1% chloramphenicol (Britania, Argentina); identification of *C. albicans* species was carried out through germinative tube development test, sugar assimilation and fermentation (Candidfast, International Microbe, France), and growth within a chromogenic medium (CHROMagar Candida, France) (18). The isolated yeasts were kept in sterile water at 4 °C.

**Antifungal agents**
Chlorhexidine solutions were used, 12.5-25-50 µg/ml chlorhexidine (CLX, Microsules Bernabó, Argentina), as well as 50-100-150 µg/ml fluconazole (FLZ, Pfizer, Argentina), 0.25 g% high-molecular-weight chitosan (HMWC 0,25% in acetic acid buffer pH 4, 90% deacetylated, 280 cps, PM 300 kDa, Unifarma Argentina) and their combinations (QAPM 0.25g% / CLX 50 µg/ml and QAPM 0.25 g% / FLZ 150 µg/ml) in phosphate saline buffer (PBS) pH 7.4. These concentrations, which are sub-inhibitory for yeasts, were previously determined in the isolated strains in our laboratory, and proved to coincide with the ones employed in our previous work and in other authors’ work (16,19,20). Minimum inhibitory concentration (MIC) was determined using the agar dilution method, which defines MIC as the lowest antifungal drug concentration where a considerable decrease in the colony size is observed compared to control growth. This method has shown a good correlation with the reference method (21).

**Yeast suspension preparation for the trials**
Before the trials, isolated yeast strains were inoculated on SGA plates and incubated during 24 h at 37°C. Subsequently, suspensions were prepared in sterile PBS (absorbance: 1.500 at 520 nm). Then, 1 mL of suspension was added to 4 mL of PBS (control) or to 4 mL of PBS-antifungal agents in the aforementioned concentrations, resulting in a 1x10⁶-1x10⁷ cells/mL concentration. Tubes were incubated during 60 min at 37°C with gentle agitation. Cells were washed twice with sterile
PBS (10 min at 3,000 rpm); later, they were resuspended in sterile PBS and adjusted to Abs520 nm = 0.800. After this, the viability was verified on SGA plates. Besides, viability of all strains was verified after treatment with acetic acid buffer pH 4; inhibition of growth was not observed.

**Determination of cell surface hydrophobicity (CSH)**

Adhesion to hydrocarbons method was used to determine CSH. Yeast suspensions were mixed with chloroform (volume relationship 5:1); the mixture was agitated for 1 minute and put to rest during 10 min at 37°C. CSH was defined as the difference between absorbances at 520 nm of the aqueous phase before and after the confronting C. albicans suspension and the organic phase, ΔAbs (5).

**Germinative tube development (GT)**

In order to induce GT development, 250 µL yeast suspensions treated with antifungal drugs were mixed with 1 mL of human serum, and incubated at 37°C during 2.5 hours. Once that time elapsed, 300 cells in continuous fields (X 400) in Neubauer chamber were counted and GT cells were expressed as a percentage (6).

**Determination of phospholipase activity (PL)**

PL activity of C. albicans strains was determined by a modified version of Vidotto’s technique (6). A 0.075 g% soy lecithin suspension (Herbaccion Isa, Argentina) with 0.056 g% CaCl2 and 5.84 g% NaCl was used in each plate of SGA. 10 µL of suspensions were inoculated in SGA supplemented with lecithin and incubated for 72-96 hours at 37°C. Each strain was tested in triplicate. The value of PL activity was determined as the quotient between the colony diameter plus halo and the colony diameter, Pz = (d col + d halo) / dcol (7).

**Determination of post antifungal effect (PAFE)**

In order to study the inhibitory effect on the recovery of fungal growth of the analyzed compounds, namely PAFE, 800 µL of C. albicans suspensions were incubated with 1.2 mL of RPMI 1640 (American Biorganics, USA) during 24 h at 37°C in a thermostatic bath with gentle agitation. Changes of absorbance at 520 nm were observed from the beginning of trial and after 24 h (ΔAbs520nm). Thus, a greater value indicates a better recovery of the fungal growth and, therefore, a smaller PAFE (8). A period of 24 hours was selected as final time for absorbance record, after an experiment where the absorbance at 520 nm was recorded each hour, from time 0 to 32 h. All experiments were carried out in triplicate and data were statistically processed by ANOVA, Spearman’s correlation coefficient and Student’s t-test for independent samples (p<0.05).

**Results and Discussion**

**Germinative tube (GT) development**

In isolated C. albicans strains obtained from patients and treated with CLX, FLZ and HMWC, alone or combined with the aforementioned compounds, a significant decrease of GT was observed (Fig. 1-a). The collection strain, CA 3153, showed significant decreases in GT after treatment with antifungal drugs and its combinations, except for FLZ. The most significant decreases of GT were observed in the presence of CLX, which proved to be the most efficient antifungal agent regarding proportional reductions of GT with concentrations (data not shown), in agreement with results observed by Ellepola et al. (22).

**Cell surface hydrophobicity (CSH)**

A significant CSH increase in isolated strains from patients was observed (Fig. 1-b) and also in CA 3153 after treatment with HMWC, HMWC/CLX and HMWC/FLZ. No significant differences were observed after treating strains only with CLX or FLZ. These results do not agree with those observed by Anil et al. (20) and ours on previous works (3), although employing the same concentrations (50 µg/ml). This could be explained by the differences in the experimental conditions, since in the present study yeast suspensions are within a solution in contact with antifungics in short times, which implies contact on cell surface and would represent a similar situation as the one occurring during the use of oral rinses. Goldberg et al. (23) found results similar to ours in the presence of HMWC, assigning the CSH increase to the polycationic character of chitosan, and this would cause a decrease of superficial negative charge of yeasts and the subsequent increase of cellular hydrophobicity (17,23). Lack of effect on CSH when strains were treated with antifungics in the absence of HMWC indicates that the polycationic polymer would be responsible for the observed effect, without synergism when combined with CLX and FLZ.

**Determination of post antifungal effect (PAFE)**

Isolated strains from patients and CA 3153 showed a significant decrease in the growth recovery when they were treated with CLX (Figure 1-c). FLZ produced a significant decrease of CA 3153 growth recovery. In the presence of HMWC, alone or combined with FLZ, an increasing tendency in the growth recovery was observed. These results show that the significant decrease observed in the presence of CLX or FLZ is counteracted in the presence of HMWC, the effect of which prevails when antifungal drugs are combined with it. Previous results have shown the antifungal effect of dissolved chitosan within the solid medium of yeast growth (3); nevertheless, the experimental conditions of this work have been different, as it has been discussed in a previous section; this is likely to be this way due to the short period contact, similarly to what happens in the case of an oral rinse.
Determination of phospholipase activity (PL)

All isolated *C. albicans* strains obtained from patients and CA 3153 exhibited PL activity. When they were treated with the analyzed antifungal agents, CA 3153 showed a significant decrease of PL activity in the presence of CLX, which proved directly proportional to concentrations, in accordance with Kadir et al.’s findings (7). In isolated strains from patients no differences were observed in PL activity with the different treatments (Fig. 1-d). This difference in the behavior between isolated strains from patients and CA 3153 may arise from the fact that these, due to their wild condition, could be adapted more easily through other mechanisms to overcome environmental difficulties or host immunity mechanisms, resulting in more resistant strains when treated with antifungal drugs.

Correlation among the analyzed virulence factors

A strong positive correlation among the analyzed virulence factors was observed in isolated untreated strains ($r^2 > 0.73$ and $p < 0.03$). These results corroborate the cooperative effect of these factors, which is a breeding ground for the development of yeast virulence and, in turn, the illness, in accordance with Vidotto et al. (6). Other authors (24) have shown the coordinated action of virulence factors, such as the contribution to fungal nutrition and the micro-environmental modification produced by fatty acid liberation from fungal lipase activity. Such liberation benefits exoenzymes proteinases activity, producing a pH decrease to values between 5.5 and 2.8. Additionally, fatty acid liberation improves *C. albicans* adhesion to host surfaces. Besides, Masuoka et al. correlated the composition of *C. albicans* cell wall (long of chain of the acid-labile portion) with change in the fungal hydrophobicity (25).

When treating yeasts with the analyzed antifungics, CLX produced positive correlations among all the factors ($p < 0.002$); and FLZ produced the same effect on GT, PAFE and PL activity ($p < 0.02$). HMWC produced strong positive correlations ($p < 0.02$), except between PAFE and PL activity. When combining HMWC with CLX, a strong positive correlation was observed between the factors ($r^2 > 0.7 - p <0.002$), except for GT and PAFE, while HMWC combined with FLZ did not produce significant associations. The combined use of antifungal drugs has been evaluated by many authors. Senel et al. (in vitro studies) (26) and Giunchedi et al. (in vivo studies) (27) obtained good results, showing that chitosan produced an adjuvant effect on liberation and effectiveness of drugs to which the former was associated in those studies. In accordance with our results, Ellepola et al. showed a strong correlation between virulence factors after treating *C. albicans* strains with CLX (22). Nevertheless, CLX is not habitually used by clinicians in our country for oral candidiasis treatment, considering its effects on the sense of taste and the staining of oral tissues and acrylic dentures (28). In this work, after combining HMWC with FLZ, a drastic effect on the capacity for GT development was observed. Considering the fact that GT development implies not only morphological changes but also a greater capacity for infection and penetration of oral soft tissues, this work contributes evidences to the results obtained by other authors in this area.

The use of virulence factors as gravity predictors of fungal infections or as in vitro models for studying *C. albicans* behavior should be carefully analyzed. Though our results show that GT development is very sensitive to the antifungal treatment, it should not be considered as an only parameter or predictor of fungal virulence. The strong correlation observed in the fungus virulence factors involved in infections shows that an individual evaluation of them would be a simplification of the analysis, considering that not all the factors are expressed in infections or necessarily in a particular stage of infection (29).

Ellepola et al. have shown that a combined treatment with antifungics would act as adjuvants in denture-related oral candidiasis and would allow diminishing...
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On the other hand, this study confirms that antifungal candidiasis or the status of patients as healthy carriers. Fungi are known to be important etiologic agents of oral candidiasis. Our results indicate that short exposures to sub-inhibitory concentrations of antifungals would be able to modulate the expression of the virulence factors studied and therefore, to facilitate the elimination of oral cavity microorganisms, and to decrease Candida’s pathogenicity, in spite of the brief exposure period in the oral environment.

Conclusions

The present work was based on the effect that short exposures to sub-inhibitory concentrations of antifungals have on virulence factors of Candida albicans, the most important etiologic agent of oral candidiasis. Our results suggest that adhesion, represented by the evaluation of CSH, exoenzymes production, dimorphism and growth recovery after treatment with antifungals in short periods, can be factors that add to the predisposing conditions of host and, in turn, determine the development of candidiasis or the status of patients as healthy carriers. On the other hand, this study confirms that antifungals used would be able to modulate the development of candidiasis.

References

24. Hube B, Stehr F, Bossenz M, Mazur A, Kretschmar M, Schäfer U. Expression of the virulence factors studied and therefore, diminish the infection capacity of C. albicans, represented by virulence factors, in accordance with other authors (7). In clinical terms, our results would indicate that short exposures to sub-inhibitory concentrations with the analyzed drugs would be able to modulate the expression of the virulence factors studied and therefore, to facilitate the elimination of oral cavity microorganisms, and to decrease Candida’s pathogenicity, in spite of the brief exposure period in the oral environment.

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