In vitro Adhesion of Oral *Candida dubliniensis* Isolates to Acrylic Denture Surfaces following Brief Exposure to Sub-Cidal Concentrations of Polyenes, Azoles and Chlorhexidine

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**Key Words**

*Candida dubliniensis* · Nystatin · Amphotericin B · Ketoconazole · Fluconazole · Chlorhexidine · Acrylic dentures

**Abstract**

**Objectives:** We aimed to investigate the effect of brief exposure to sub-cidal concentrations of nystatin, amphotericin B, ketoconazole, fluconazole and chlorhexidine gluconate on the adhesion of oral *Candida dubliniensis* isolates to the surface of acrylic dentures. **Methods:** After determining the minimum inhibitory concentration of each drug, 20 oral isolates of *C. dubliniensis* were exposed to sub-cidal concentrations of the drugs for 1 h. The drugs were then removed by dilution, and the adhesion of the isolates to denture acrylic strips was assessed by an in vitro adhesion assay. **Results:** Compared to the controls, exposure to nystatin, amphotericin B, ketoconazole, fluconazole and chlorhexidine gluconate suppressed the ability of *C. dubliniensis* isolates to adhere to acrylic denture surfaces with a reduction of 74.68, 74.27, 57.31, 44.57 and 56.53% (*p* < 0.001 for all drugs), respectively. **Conclusions:** Brief exposure to sub-cidal concentrations of anti-mycotics suppressed the adhesion of *C. dubliniensis* oral isolates to acrylic denture surfaces.

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**Introduction**

Oral candidosis is considered to be the most common human fungal infection, which manifests in a variety of clinical guises. *Candida*-induced denture stomatitis (CDS) is one form of oral candidosis. Acrylic dentures are an important predisposing factor for CDS as they are usually ill-fitting and, with sub-optimal hygiene, can act as reservoirs of infection [1]. CDS is seen as an inflammatory candidal lesion of the oral mucosa, occurring particularly under complete or partial removable dentures and it affects up to 65% of denture wearers [2]. High salivary counts of *Candida* are also much more common in full-denture wearers than in dentate individuals [3]. It is believed that
the ability of Candida sp. to adhere to acrylic surfaces may be important in the pathogenesis of CDS, as adhesion is apparently the initial step in the colonization and subsequent infection of host surfaces by Candida.

Candida dubliniensis is now well-recognized as an opportunistic fungi attendant with recurrent oral candidosis in AIDS patients [4]. It has also been isolated from the oral cavity of diabetic patients and the sputum of cystic fibrosis patients [5]. The fact that C. dubliniensis has been isolated from the upper respiratory tract specimens, from the blood and from a case of endocarditis involving a prosthetic aortic valve suggests that it can disseminate to other sites as well [4, 6–8]. Interestingly, when Candida sp. colonization and CDS were assessed in complete denture wearers in Iran, C. dubliniensis was recovered from 10.9% of the samples analyzed [9]. C. dubliniensis isolates were also detected in the oral environment of denture wearers in association with C. albicans isolates in 10% of CDS cases in Brazil [10]. Moreover, when Candida sp. isolated from 50 individuals with CDS were analyzed, C. dubliniensis was recovered from 4 of these individuals [2]. It has thus been suggested that the association between C. dubliniensis and CDS may play an important role in the establishment and persistence of C. dubliniensis-induced CDS [10].

Many therapeutic agents with anti-fungal properties are obtainable for the treatment of CDS [11, 12]. These agents include polyene anti-fungals such as nystatin and amphotericin B, azoles such as ketoconazole (an imidazole) and fluconazole (a triazole) as well as chlorhexidine gluconate mouthwash [11, 12]. Despite the availability of these agents for the treatment of CDS, failure of therapy is not uncommon [11, 12]. For instance, the diluent effect of saliva and the cleansing effect of the oral musculature in the niches of the oral milieu tend to reduce the availability of anti-fungal agents below that of effective therapeutic concentrations, thereby lowering their efficacy. Hence, intra-orally, the pathogenic Candida may undergo a brief exposure to anti-fungal drugs, but after that the drug concentration is likely to become sub-therapeutic [11, 12]. The ability of C. dubliniensis isolates to adhere to denture acrylic after such brief exposure to anti-fungal drugs has not been studied so far. Based on the above information and the results of a recent study conducted in Kuwait where oral C. dubliniensis isolates had the highest prevalence among non-C. albicans oral Candida sp. [13], our objective was to determine the adherence of oral C. dubliniensis isolates to denture acrylic following brief exposure to sub-cidual concentrations of nystatin, amphotericin B, ketoconazole, fluconazole and chlorhexidine gluconate.

### Materials and Methods

#### Organisms

We made use of 20 oral isolates of C. dubliniensis recovered from the oral-rinse samples of patients attending the Kuwait University Dental Clinic in a previous study [13]. Initially, all the yeast isolates were tested for germ tube (GT) formation. The colony characteristics were then observed using CHROMagar Candida medium (Becton Dickinson & Co., Sparks, Md., USA), and carbohydrate assimilation profiles were obtained using the VITEK 2 yeast identification system (BioMérieux, Marcy-l’Etoile, France). The identity of C. dubliniensis was confirmed by the production of rough colonies with hyphal fringes and chlamydospores on simplified sunflower seed agar and by using semi-nested PCR amplification of the internally transcribed spacer 2 region of rDNA followed by direct DNA sequencing of the internally transcribed spacer region of rDNA [13].

#### Anti-Fungal Agents and Media

Anti-fungal agents were prepared as in previous similar studies [14, 15]. Briefly, nystatin, amphotericin B, ketoconazole and fluconazole (Sigma, St. Louis, Mo., USA) were dissolved in dimethyl sulphoxide. Initially, 10,000 μg/ml of each anti-mycotic agent was prepared and stored at −20°C as previously described [14, 15]. For 1 h of exposure to the yeasts, each agent was suspended/diluted in RPMI 1640 medium buffered with 0.165 M MOPS [3-(N-morpholino)propanesulphonic acid] containing L-glutamine and lacking sodium bicarbonate (Sigma), which was dissolved in 1 litre of sterile distilled water and adjusted to a pH of 7.2 and then filter sterilized. Chlorhexidine gluconate (0.2%; GlaxoSmithKline, Brentford, UK) was dissolved in sterile phosphate-buffered saline (PBS) at a pH of 7.2 and diluted to obtain a concentration of 0.005% prior to each experiment as previously described [16, 17].

#### Determination of Minimum Inhibitory Concentration

Anti-fungal susceptibility values of C. dubliniensis isolates were determined (as in a previous study [13]) against amphotericin B, ketoconazole and fluconazole by Etest and performed according to the manufacturer’s recommendations (AB BIODISK, Solna, Sweden). Briefly, each test isolate was freshly sub-cultured. Five isolated colonies were uniformly suspended in sterile saline, and turbidity was adjusted to a 0.5 McFarland standard, which is approximately 1–5 x10⁶ cells/ml according to the 2004 Clinical and Laboratory Standards Institute (CLSI) document M44-A. This inoculum was swabbed onto the agar plates (150 mm in diameter) and allowed to dry for 10–15 min before the Etest strips were applied. RPMI 1640 agar supplemented with 2% glucose and buffered with MOPS (0.165 μM; pH 7.0) was used for susceptibility testing according to the method recommended by the CLSI document M27-A2. The plates were incubated at 35°C, and the minimum inhibitory concentration (MIC) values were recorded after 24–48 h of incubation. The point where inhibition ellipses intercepted the scale on the anti-fungal strip was taken as the MIC for each test isolate. There was complete inhibition (100%) of growth for amphotericin B, and a marked decrease in growth intensity (80%) for fluconazole and ketoconazole. C. albicans ATCC 90028 and C. parapsilosis ATCC 22019 were the reference strains used for quality control of the susceptibility testing. The interpretive susceptibility break points for fluconazole were those recommended by the CLSI document M27-A2. Due to the lack of defined susceptibility breakpoints...

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break points for amphotericin B and ketoconazole, an isolate was considered susceptible when the MIC break point was ≤1.0 μg/ml for amphotericin B and 0.125 μg/ml for ketoconazole [13].

The MIC values of nystatin and chlorhexidine gluconate were determined by the broth dilution technique as done previously [14, 18] by performing 2-fold serial dilutions of the drug in microtitre plates using an inoculum of 1–5 × 10^5 CFU/ml. The MIC was determined visually after 24 h of incubation at 37°C [18–20]. It was defined as the lowest concentration of the drug that inhibited the growth of Candida cells, as indicated by the absence of turbidity (i.e. optically clear). It was read independently by 2 laboratory personnel, with C. albicans ATCC 90028 being used as a reference strain.

Preparation of Denture Acrylic Strips
The acrylic strips for the adhesion assay were prepared as described previously [1, 19]. In brief, transparent self-polymerizing acrylic powder (1.5 g polymethyl methacrylate powder) was spread on an aluminum foil-covered glass slide (2.5 × 7.5 cm). Monomer liquid (1 ml; DENTSPLY Ltd., Weybridge, UK) was poured on to the surface of the slide, a second slide similar to the first was immediately placed on top of the polymerizing mixture and the slides were then firmly secured at both ends with two binder clips. After bench-curing for 30 min, the glass slides were separated. The resultant acrylic strips were cut into 5 × 5 mm squares, immersed in distilled water for 1 week to leach any excess monomer and then washed in running water for 3 h. The strips were disinfected by dipping in 70% alcohol and then washed with sterile distilled water. They were then ultrasonicated for 20 min to remove any contaminants and artefacts from the surfaces, washed again in sterile distilled water, dried and then used for the adhesion assay.

Preparation of the Cell Suspension for the Adhesion Assay
A previously described method was used [1, 14–19]. Briefly, yeast cells that had been maintained on Sabouraud dextrose agar were inoculated onto fresh plates and incubated overnight at 37°C for 24 h prior to use. The organisms were harvested, and a cell suspension was prepared in sterile PBS at 520 nm to an optical density of 1.5. From this cell suspension, 0.5 ml was added to tubes containing 2 ml of RPMI (control) and 2 ml of RPMI/drug (test), in which the sub-cidal drug concentrations were 2 times the MIC values as in previous studies [1, 14–19]. This gave a cell suspension of 10^6 cells/ml in each assay tube. The tubes were then incubated at 37°C for a period of 1 h. Following this limited exposure, the cells were washed twice in PBS and centrifuged for 10 min at 3,000 g. The supernatant was decanted, and the pellets were resuspended in 6 ml of sterile PBS. This procedure has been used previously for drug removal and has been shown to reduce the concentration of the drug as much as 10,000 times, thereby minimizing any carry-over effects of the drug following its removal [1, 14–19]. Viable counts of the control and the test were performed after removal of the drugs. As the procedure of drug removal effectively eliminated any carry-over effect, there was virtually no difference on the viable counts of the control and the tests following exposure to the already-diluted sub-therapeutic drug concentrations as observed in previous studies [1, 14–19].

Adhesion Assay
A previously applied method was used [1, 19]. With the help of aseptic techniques, acrylic strips were placed vertically in the wells of a sterile serological plate. Thereafter, 400 μl of Candida cell suspension was added to each well, until it completely covered the acrylic strips. The whole assembly was then placed in an incubator for 1 h at 37°C with gentle agitation at 120 rev/min. The strips were then recovered aseptically from the wells and washed 3 times, by dipping gently into sterile PBS which helped to dislodge the loosely attached Candida cells. They were then dried and stained using a modified Gram stain without the counter-stain as done previously. After air-drying at room temperature, they were mounted on glass slides with glycerol, and the adherent Candida was quantified as previously described [20]. Adherent Candida cells in 20 fields of view for each strip (0.25 mm²/field) were determined using a light microscope at ×400 magnification, and the results were expressed as Candida cells/mm². The majority of the attached Candida cells were in the blastospore stage (cells with a rounded format), some with daughter cells and only very few with hyphae or pseudohyphae. The following previously used parameters [20] were used to standardize the counts: a budding yeast cell was considered as a unit cell if the daughter cell was smaller than the mother cell, and a hyphum was counted as a single cell.

All experiments were repeated on three separate occasions with duplicate determinations on each occasion.

Statistical Analysis
As for previous studies [1, 19], the data obtained from the adhesion assay were analyzed using the ANOVA Dunnet t test, which treats one group as a control (unexposed to drugs) and compares all other groups (exposed to drugs) against this. Variations due to the different drugs on the suppression of adhesion to the denture acrylic were analyzed using the Tukey-Kramer multiple-comparisons tests. A p value of <0.05 was considered statistically significant.

Results
The MIC of the 20 isolates of C. dubliniensis to amphotericin B ranged from 0.002 to 0.125 μg/ml. For ketoconazole, the range was 0.002–0.012 μg/ml. For fluconazole, it was 0.016–0.38 μg/ml. The MIC of the 20 isolates to nystatin ranged from 0.09 to 0.78 μg/ml. The MIC of chlorhexidine gluconate for all C. dubliniensis isolates tested was at 0.00125% dilution.

The candidal adhesion of the unexposed C. dubliniensis isolates to the acrylic denture surfaces was 45.05 cells/mm². The adhesion to the denture acrylic following brief exposure to nystatin, amphotericin B, ketoconazole, fluconazole and chlorhexidine gluconate was 11.34, 11.50, 19.14, 24.85 and 19.51 cells/mm², respectively (table 1). Hence, compared to the controls, brief exposure to nystatin, amphotericin B, ketoconazole, fluconazole and chlorhexidine gluconate inhibited the ability of the isolates to adhere to the denture acrylic with a reduction of 74.68% (range 64.75–82.09%), 74.27% (range 63.21–83.10%), 57.31% (range 47.58–67.40%), 44.57% (range 34.19–54.19%) and 28.14% (range 18.23–38.05%), respectively.
30.19–57.05%) and 56.53% (range 45.23–66.60%), respectively (table 2). These values were all highly significant (p < 0.0001).

Analysis of the variations between the 5 drugs on the suppression of candidal adhesion to acrylic denture surfaces using the Tukey-Kramer multiple-comparisons tests is shown in table 3. It was revealed that the suppressive effect elicited by nystatin and amphotericin B was significantly higher than and statistically different from that of ketoconazole, fluconazole and chlorhexidine gluconate (p < 0.001). However, the suppressive effects elicited by the 2 polyenes did not differ statistically. The sup-

<table>
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<th>Isolate</th>
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<th>NYS</th>
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<th>KETO</th>
<th>FLU</th>
<th>CHX</th>
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<td>26.1±2.41</td>
<td>20.6±1.33</td>
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Mean 45.05 11.34 11.50 19.14 24.85 19.51
SEM 0.68 0.43 0.46 0.55 0.66 0.61
SD 3.04 1.93 2.04 2.48 2.93 2.74
p value <0.001 <0.001 <0.001 <0.001 <0.001

All values indicate the average of experiments repeated on 3 separate occasions with duplicate determinations on each occasion. AMP-B = Amphotericin B; CHX = chlorhexidine; FLU = fluconazole; KETO = ketoconazole; NYS = nystatin.

Table 2. The reduction of adhesion of C. dubliniensis isolates (cells/mm²) to acrylic denture surfaces following 1 h exposure to and subsequent removal of therapeutic agents with anti-fungal properties compared to the unexposed controls

<table>
<thead>
<tr>
<th></th>
<th>NYS</th>
<th>AMP-B</th>
<th>KETO</th>
<th>FLU</th>
<th>CHX</th>
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<tr>
<td>Reduction, %</td>
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<td>74.27</td>
<td>57.31</td>
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<tr>
<td>SEM</td>
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<td>1.21</td>
<td>1.41</td>
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<tr>
<td>SD</td>
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<td>5.40</td>
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<td>p value</td>
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AMP-B = Amphotericin B; CHX = chlorhexidine; FLU = fluconazole; KETO = ketoconazole; NYS = nystatin.
pressive effect elicited by ketoconazole (imidazole) was significantly different from that of the 2 polyenes as well as from fluconazole (p < 0.001) but not from chlorhexidine gluconate. The suppressive effect elicited by fluconazole (triazole) was significantly different from that of the 2 polyenes, ketoconazole and chlorhexidine gluconate (p < 0.001).

**Discussion**

In this study, the MIC values of the polyene and azole anti-fungal agents for *C. dubliniensis* isolates were within the range of previous studies with the identical *C. dubliniensis* isolates [14, 15]. Based on CLSI standard values, all the isolates were susceptible to nystatin, amphotericin B, ketoconazole and fluconazole. An MIC at a dilution of 0.00125% for chlorhexidine gluconate was observed for *C. dubliniensis* isolates, which is similar to the MIC value for *C. albicans* isolates in a previous study for this anti-septic [18]. Studies have also demonstrated dynamic changes in the ultra-structural features of the cell wall during morphogenic transformation to GTs, and have shown that the cell wall of GTs possesses stratification comparable to that of the blastospore wall [25]. Therefore, it is reasonable to speculate that polyene-induced changes in the structure of the cell wall of *Candida* would affect active budding and multiplication, thus suppressing not only its adhesion to denture acrylic as seen in our study, but also its adhesion to BECs, GT formation and CSH following brief exposure to nystatin and amphotericin B.

Limited exposure to the 2 azole drugs, ketoconazole and fluconazole, also significantly reduced *C. dubliniensis* adhesion to the denture acrylic. Similar suppressive effects on other *C. dubliniensis* adhesion attributes such as adhesion to BECs, GT formation and CSH following brief exposure to ketoconazole and fluconazole have been reported before [15, 21]. Again, these adhesion-related suppressive effects may be related to the mode of action of the azoles. These drugs alter the fungal cell membranes by blocking the 14-α-demethylation step in the biosynthesis of ergosterol. The consequent depletion of ergosterol and accumulation of 14-α-methyl sterols leads to alterations in a number of membrane-associated functions [11, 12]. Such alterations could be the reason for the reduced ability to adhere to acrylic denture surfaces that we observed, as well as for the suppressive ef-
fects on *C. dubliniensis* isolates and other candidal adhesion traits seen in previous studies [15, 21] after limited exposure to azoles.

Chlorhexidine gluconate is used as an adjunct in the treatment of oral candidosis. For instance, 0.2% chlorhexidine gluconate has been used successfully as an oral rinse in the treatment of CDS, while the 2% suspension is used as an overnight denture disinfectant [11, 12, 26]. Chlorhexidine gluconate has a dual action on *Candida* sp.: it is fungicidal at therapeutic concentrations and it significantly suppresses yeast adhesion to both inorganic and organic substrates [11, 12, 26]. Our results also indicated that limited exposure of oral *C. dubliniensis* isolates to 0.005% chlorhexidine gluconate could suppress their ability to adhere to denture acrylic. Similar suppressive effects on other *C. dubliniensis* adhesion attributes such as GT formation and CSH following brief exposure to sub-therapeutic concentrations of chlorhexidine gluconate have been reported previously [16, 27]; these effects may be related to the pharmacodynamics of the antiseptic on the cell wall of *Candida*. Scanning and transmission electron micrographic studies have demonstrated that the anti-fungal effect of this antiseptic is most likely due to a loss of cytoplasmic components, coagulation of nucleoproteins and the associated morphological changes in the structure of the cell wall [28]. Therefore, it is reasonable to speculate that chlorhexidine gluconate, by affecting the cell wall structure, yields an anti-fungal effect and suppresses the ability of *C. dubliniensis* isolates to adhere to the denture acrylic.

In this study, brief exposure to the polyene anti-fungal agents resulted in a highly significant suppression (approx. 75%) on adhesion to denture acrylic, compared to that of ketoconazole, fluconazole and chlorhexidine gluconate (approx. 58, 45 and 57%, respectively). Such differential activity between the polyenes and the azoles may be due to the fundamental dissimilarity in their modes of action. Polyenes are fungicidal and azoles are fungistatic, which may be the reason for the difference in sensitivity of *Candida* cell surface structures to polyenes. In addition, there was a statistically significant difference between the 2 azoles in terms of their suppressive effect on adhesion to denture acrylic; a low susceptibility was elicited by fluconazole (45%) compared to ketoconazole (58%). Although fluconazole is markedly more effective than ketoconazole in the management of candidal infection, its growth inhibitory activity against *Candida* isolates in vitro is less than that of ketoconazole [11, 12]. Furthermore, there could be subtle differences between the 2 azoles, as ketoconazole is an imidazole and fluconazole is a triazole. It can thus be speculated that these differences may be the reason for the reduced ability of fluconazole to suppress adhesion to denture acrylic as we observed. Notwithstanding this significant in vitro observation, fluconazole is the drug of choice in the management of chronic *Candida* infections such as those seen in HIV-infected individuals and debilitated patients [11, 12]. The anti-candidal pharmacodynamics of chlorhexidine gluconate have not yet been fully elucidated, but our results provide a glimpse of its potency, which is still to be fully evaluated.

This study reports the suppression of adhesion of oral *C. dubliniensis* isolates to denture acrylic by a number of therapeutic agents that possess a wide spectrum of anti-candidal pharmacological properties. We included polyenes, imidazole, triazole and chlorhexidine gluconate, which covered a large number of oral *C. dubliniensis* isolates. Our findings lend further credibility to the use of these anti-fungal agents in the management of CDS. Further investigations, with a larger number of sessile *Candida* isolates encased in a biofilm as opposed to the planktonic isolates used in this study, are warranted to further augment these findings.

**Conclusion**

The findings of this study showed that a brief exposure to sub-therapeutic concentrations of nystatin, amphotericin B, ketoconazole, fluconazole and chlorhexidine gluconate suppressed the adhesion ability of *C. dubliniensis* isolates to denture acrylic in vitro. When used in vivo, there is the possibility that such exposure could reduce the pathogenicity of *Candida*.

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