Alterations in growth and branching of Neurospora crassa caused by sub-inhibitory concentrations of antifungal agents

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ABSTRACT

Six antifungal agents at subinhibitory concentrations were used for investigating their ability to affect the growth and branching in Neurospora crassa. Among the antifungals herein used, the azole agent ketoconazole at 0.5 µg/ml inhibited radial growth more than fluconazole at 5.0 µg/ml while amphotericin B at 0.05 µg/ml was more effective than nystatin at 0.05 µg/ml. Morphological alterations in hyphae were observed in the presence of griseofulvin, ketoconazole and terbinafine at the established concentrations. The antifungal agents were more effective on vegetative growth than on conidial germination. Terbinafine markedly reduced growth unit length (GU) by 54.89%, and caused mycelia to become hyperbranched. In all cases, there was a high correlation between hyphal length and number of tips (r > 0.9). All our results showed highly significant differences by ANOVA (p < 0.001, α = 0.05). Considering that the hyphal tip is the main interface between the fungus and its environment, through which enzymes and toxins are secreted and nutrients absorbed, it would not be desirable to obtain a hyperbranched mycelia with inefficient doses of antifungal drugs.

Key words: antifungal drugs, apical growth, Neurospora crassa

RESUMEN

Alteraciones de crecimiento y ramificación en Neurospora crassa provocadas por concentraciones subinhibitorias de agentes antimicóticos. Se investigó el efecto de seis agentes antimicóticos en concentraciones subinhibitorias sobre el crecimiento y la ramificación en Neurospora crassa. El agente azólico ketoconazol a la concentración de 0,5 µg/ml inhibió el crecimiento radial más que el fluconazol a 5,0 µg/ml, y la anfotericina B a 0,05 µg/ml fue más eficiente que 0,05 µg/ml de nistatina, entre los agentes poliénicos usados. En presencia de griseofulvina, ketoconazol y terbinafina a las concentraciones establecidas se observaron alteraciones morfológicas en las hifas. Los agentes antimicóticos fueron más eficientes sobre el crecimiento vegetativo que sobre la germinación conidial. La terbinafina redujo marcadamente (54.89%) la longitud de la unidad de crecimiento y provocó la hiperramificación del micelio. En todos los casos, existió gran correlación entre la longitud y el número de ápices de las hifas (r > 0.9). Todos los resultados mostraron diferencias altamente significativas de acuerdo con ANOVA (p < 0.001, α = 0.05). Considerando que el ápice de la hifa es la principal interfase entre el hongo y su ambiente, a través de la cual las enzimas y las toxinas son secretadas y los nutrientes son absorbidos, un micelio hiperramificado resultante de dosis ineficientes de agentes antimicóticos sería perjudicial.

Palabras clave: drogas antimicóticas, crecimiento apical, Neurospora crassa

INTRODUCTION

Due to several correlated factors like the emergence of AIDS and other highly infectious diseases, organ transplant and cancer chemotherapy, the nosocomial immunocomprised population has noticeably increased over the last decades, bringing along the spectre of opportunistic fungal infections (5, 21, 25). It is important to call attention to the limited number of effective antifungal compounds available and the need to develop new clinical options (3, 9). Fungi are eukaryotic organisms that share a large set of potential drug targets with their hosts, hindering the development of new agents (21). In spite of this, new azoles (voriconazole and posaconazole), a class of echinocandins, especially caspofungin and anidulafungin, have recently been examined (8, 15, 24).

The aim of the present work was to study the effects of subinhibitory concentrations of six antifungal drugs on different phases of the development of N. crassa, used as fungal model, by analyzing changes in conidial germination, vegetative growth and morphology of hyphal branching.

MATERIALS AND METHODS

Microorganism

Neurospora crassa wild type 74OR 8-1a (ATCC) kept on slants of Vogel’s Minimum Medium (VMM) supplemented with 2% (w/v) glucose and stored at 4 °C, was used throughout this work.
For the experiments, fungus was grown in this medium at 30 ± 1°C for 3 days, and at room temperature for 4 days.

Antifungal drugs

Sub-lethal concentrations of six antifungal agents from four classes of antibiotics having different mechanisms of action were tested. Griseofulvin, a carbamate, exerts its fungistatic action by inhibiting cell division, leading to the disruption of the mitotic spindle structure (22). Fluconazole and ketoconazole are azole agents that decrease ergosterol synthesis by inhibiting cytochrome P450 enzymes (16). Terbinafine, an agent from the allylamines class, acts by inhibiting the enzymes responsible for squalene conversion (17). Polyenic macrolides (nystatin and amphotericin B) have in common a mechanism of action by means of which they selectively bind to ergosterol (7). In earlier experiments, different concentrations of antifungals have been tested in order to establish those that would allow a certain fungus growth degree. Lethal concentrations of antifungals would not fulfill the aims of this study. It is worth pointing out that the excipient used, usually lactose and/or magnesium stearate contained in the tablets or capsules wherefrom the employed antifungals were extracted, are innocuous for man. Tablets or capsules were pulverized and dissolved in dimethylsulphoxide [DMSO 99.5% (v/v)] to yield drug solutions at the established concentrations, after adequate dilution. Samples were aseptically added to the VMM at 45 °C. The final medium DMSO concentration was adjusted to a maximum of 0.5% (v/v). Controls contained the same final DMSO concentration, but no antifungal drug was added.

Evaluation of antifungals on fungal growth and morphology

To evaluate the effect of antifungal drugs on N. crassa growth, 7-day old conidia were inoculated on VMM containing the antifungals, in Petri dishes and incubated at 30 ± 1 °C for 24 h. Growth was followed by measuring colony Radii after 7; 15; 20 and 24 h in three replicates for each concentration. Fungus growth rate was determined using 50.00 cm race tubes containing VMM supplemented or not with each antifungal. Media at the end of the race resulted in a 2-fold increase. Among the polyene antifungals, in Petri dishes and incubated for an additional 24 hours.

Statistical Analysis

ANOVA and Student’s t tests for significance were performed at the level of $\alpha = 0.05$. Correlation between hyphal length and number of tips (GU) was determined by regression analysis.

RESULTS

Radial growth

In previous experiments, several concentrations of each antifungal had been tested to determine the sub-lethal concentration. We observed a significant colony radial growth inhibition compared to controls, but less than 50% (data not shown). A concentration was established for each drug according to its effects (5.00 µg/ml griseofulvin; 5.00 µg/ml fluconazole; 0.50 µg/ml ketoconazole; 0.25 µg/ml terbinafine; 0.05 µg/ml nystatin; 0.05 µg/ml amphotericin B). At equal concentration, fluconazole caused an inhibition approximately 4-fold higher than that produced by griseofulvin (Table 1). At one-half the concentration of ketoconazole, the inhibition by terbinafine resulted in a 2-fold increase. Among the polyene antifun-

Table 1. Standard concentration of antifungals and inhibitory effect on colony radius.

<table>
<thead>
<tr>
<th>Drug</th>
<th>Concentration (µg/ml)</th>
<th>Inhibition (%)</th>
<th>ANOVA</th>
</tr>
</thead>
<tbody>
<tr>
<td>Griseofulvin</td>
<td>5.00</td>
<td>10.57</td>
<td>-</td>
</tr>
<tr>
<td>Fluconazole</td>
<td>5.00</td>
<td>43.99</td>
<td>++</td>
</tr>
<tr>
<td>Ketoconazole</td>
<td>0.50</td>
<td>15.99</td>
<td>+</td>
</tr>
<tr>
<td>Terbinafine</td>
<td>0.25</td>
<td>31.84</td>
<td>++</td>
</tr>
<tr>
<td>Nystatin</td>
<td>0.05</td>
<td>0</td>
<td>-</td>
</tr>
<tr>
<td>Amphotericin B</td>
<td>0.05</td>
<td>33.54</td>
<td>++</td>
</tr>
</tbody>
</table>

(-) non significant ($p > 0.05$); (+) significant ($p < 0.05$); (++) extremely significant ($p < 0.001$). Antifungals were added to VMM, and conidia were inoculated over dialysis membranes. Plates were then incubated at 30 °C for 24 h and colony radius determined. Values are the average of at least 3 replicates and were tested for significance (ANOVA, $p < 0.001$, $\alpha = 0.05$)

Evaluation on conidial germination and vegetative growth

Approximately 100 conidia were inoculated on dialysis membranes, overlaying VMM with antifungals, and incubated for 5 h at 30 °C. The total numbers of conidia, as well as the amounts germinated were evaluated under a Zeiss Axioskop microscope. To assess the effect of drugs on vegetative hyphae, conidia were allowed to grow for 24 h on VMM. Mature hyphae were then sectioned about 1.0 cm below the apex, allowing to rest on the agar for 1 h to achieve osmotic stabilization. Next, the pieces of dialysis membranes covered with hyphae, were transplanted onto VMM supplemented with different antifungal drug concentrations and incubated for an additional 24 hours.
The effect of antifungals on *N. crassa* growth

- **Antifungal Effect on Growth**
  - Amphotericin B led to a 33% growth inhibition, whereas, nystatin at the same concentration, did not result in inhibition. Neither DMSO used to prepare the test solutions nor the pharmaceutical excipients presented significantly inhibited growth at the doses of antifungals employed (Student's *t*, *p* < 0.001, α = 0.05).

- **Alterations in Hyphal Morphology**
  - It is known that in its normal growth, *Neurospora crassa* displays highly polarized and regularly spaced branches emerging from a main hypha (Figure 1D). In the presence of griseofulvin (Figure 1A), the branches became curled denoting a loss of the polarized growth. Supplementation of VMM with ketoconazole resulted in shorter and bud-like branches (Figure 1B); terbinafine induced unusual multipolar germination of conidia leading to short hyphae (Figure 1C). No alterations on hyphal morphology were detected in nystatin, amphotericin B and fluconazole presence.

- **Percentages of Conidial Germination and Inhibition of Vegetative Growth**
  - All drugs caused a reduction of approximately 30 – 40% of conidial germination after 5 h incubation, with the exception of terbinafine, in whose presence less than 20% of the total conidia inoculum, germinated (Figure 2). Terbinafine was more effective on conidial germination than on inhibiting vegetative growth (80% against 50%, respectively). The inhibitor effect of griseofulvin, fluconazole and ketoconazole was stronger on vegetative growth than on conidial germination. Polynye drugs did not significantly inhibit vegetative growth.

- **Effect on Growth Rate**
  - Terbinafine caused a strong reduction of *N. crassa* growth rate (Table 2), whereas none of the polynye anti-
fungals showed this effect to a significant extent (ANOVA, \( \alpha = 0.05 \)). The other agents tested only exhibited moderate inhibition effects. Under the conditions employed, an initial 24-hour adaptation period post inoculation was shown, followed by a constant growth rate. Adaptation to terbinafine required approximately 48 h.

**Table 3. Effect of antifungals on the GU of *N. crassa.***

<table>
<thead>
<tr>
<th>Condition</th>
<th>n</th>
<th>GU (µm)</th>
<th>SEM</th>
<th>Variation (%)</th>
<th>( r )</th>
<th>Dunnett’s test</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>16</td>
<td>243.76</td>
<td>12.2</td>
<td>- 0.45</td>
<td>0.9008</td>
<td>-</td>
</tr>
<tr>
<td>Griseofulvin</td>
<td>10</td>
<td>242.67</td>
<td>10.1</td>
<td>+ 5.65</td>
<td>0.9324</td>
<td>-</td>
</tr>
<tr>
<td>Fluconazole</td>
<td>14</td>
<td>258.27</td>
<td>9.7</td>
<td>- 11.43</td>
<td>0.9184</td>
<td>-</td>
</tr>
<tr>
<td>Ketoconazole</td>
<td>8</td>
<td>215.89</td>
<td>17.8</td>
<td>- 54.89</td>
<td>0.9443</td>
<td>+</td>
</tr>
<tr>
<td>Terbinafine</td>
<td>13</td>
<td>109.96</td>
<td>5.8</td>
<td>+ 45.10</td>
<td>0.9448</td>
<td>+</td>
</tr>
<tr>
<td>Nystatin</td>
<td>15</td>
<td>353.75</td>
<td>18.0</td>
<td>+ 54.25</td>
<td>0.9585</td>
<td>+</td>
</tr>
<tr>
<td>Amphotericin B</td>
<td>8</td>
<td>376.02</td>
<td>23.8</td>
<td>- 54.89</td>
<td>0.9448</td>
<td>+</td>
</tr>
</tbody>
</table>

\( n \), number of replicates; SEM, standard error of mean; \( r \), correlation coefficient; -, non significant; +, significant (\( p < 0.01 \)). Values obtained were submitted to Dunnett’s multiple comparison tests of significance.

**DISCUSSION**

It has been reported that DMSO concentrations greater than 2.0% may inhibit growth and disrupt *Neurospora crassa*’s apical actin cap (4, 11). The concentration used in the present work (0.5% v/v) caused none of these effects when radial growth was evaluated. A direct relationship between the antymycotic concentration and radial growth inhibition was observed for all drugs studied. However, agents acting on similar metabolic fungal pathways affected them to a different extent. For instance, the same concentration of amphotericin B and nystatin (0.05 µg/ml) caused a 33.54% inhibition and no inhibition, respectively. Drugs acting on the same molecular targets, like fluconazole and ketoconazole, respectively inhibited fungal growth in different ways. Such differences may be explained by the drug’s structural variety leading to different binding affinities to affected enzymes. Terbinafine’s inhibition of squalene epoxidase has been proven to be more effective than azole’s action on 14\( \alpha \)-demethylase (25). It is also known that fungi present distinct physiological features according to their various developmental stages (18). Therefore, in this study, the antymycotic effect was evaluated during both conidial germination and vegetative growth phases and as a result, a variety of effects were observed. Thus, although unable to cause a significant inhibitory effect during vegetative growth, macrolides reduced conidia germination by approximately 30%. Although azole drugs are known to act on identical molecular targets, they affected fungal growth differently according to their phase. Fluconazole reduced vegetative growth by 52.26%, but ketoconazole caused only a 28.66% inhibition. In turn, ketoconazole reduced conidial germination by 42.20% while fluconazole reduced this parameter by 32.74%. Structure-activity relationships could account for such differences (20). Irrespective of the growth phase, terbinafine proved to be the most ef-

![Figure 3. Effects of antifungals on Growth Unit (GU) of *N. crassa.* Parameters were obtained from the ratios between total length of a given hypha and its respective number of tips, shown on digitally enhanced micrographs of at least 8 determinations. Controls (ctrl) contained no antifungal; standard errors remained below 10%. Griseofulvin (grs), fluconazole (flz), ketoconazole (ktz), terbinafine (terb), nystatin (nys), and amphotericin B (AmB).](image-url)
fective agent of all those studied, as it decreased vegetative growth by 53.5% and conidial germination by 79.58%. With the exception of nystatin and amphotherin B, antifungals added to VMM, on average, significantly reduced the growth rate between 8.58% for fluconazole and 65.45% for terbinafine. It has been previously established for a variety of fungal species, that different growth conditions or the presence of inhibitors can lead to alterations in hyphae morphology, growth and branching patterns (23). Determining the hyphal growth unit could bring about new data concerning these alterations. The present results show that griseofulvin, fluconazole and ketoconazole, kept the growth unit and hyphal density unaltered relative to controls. On the other hand, terbinafine significantly reduced the growth unit length by over 50%. This could be explained as a result of the higher number of branches per length of hypha (i.e. the number of branches in the terbinafine-supplemented medium could practically be doubled with respect to the controls).

Hyphae morphology on non-supplemented VMM showed polarized growth and regularly spaced branches, in agreement with previous reports (1). Under the effect of griseofulvin, hyphae became rugged and curled. This phenotype is similar to that of the N. crassa dynein-deficient roppy mutant (19). Fluconazole caused slight increase in mycelium density and branches emerged in irregular fashion from the main hypha. Also, at closer examination, apex frequently showed a dichotomic pattern very similar to that observed in hyphae submitted to cold shock (12) or in the presence of the calcium channel blocker, verapamil (6). The incorporation of ergosterol precursors into the plasma membrane may also lead to a barrier of altered fluidity, reflecting effect on the function of calcium-dependent ATPases and other integrating proteins (13). The hyperbranched growth pattern observed under the effect of terbinafine closely resembled that of spray and frost N. crassa mutants (6). It is generally accepted that for filamentous fungi, life is at the tip (10), since the main biological processes such as growth, reproduction, morphogenesis, substrate absorption, environment recognition, and protein secretion occur in the apical region. It is to be noted that previous reports had shown that protein secretion in filamentous fungi is restricted to the growing tips and that conditions that increase growing surface lead to an increase in the amount of protein secretion (14).

In this work, we demonstrated that at least two of the antymycotics tested, induced morphological alterations. Griseofulvin disorganized polarized growth, while terbinafine led to hyperbranching. Considering that increased branching could result in increased active growth area, it may be speculated that for pathogenic fungi this could be translated into increased pathogenicity. Misuse of medicines, especially of antibiotic agents, is a well known clinical occurrence and the main cause of bacterial (and, more recently demonstrated) fungal resistance. Early interruption of treatment, often times prior clinical cure, can result in the selection of resistant strains. Eventually, plasma levels of these drugs could decrease to levels similar to those shown in our work to have caused hyperbranching.

In conclusion, a potentially more resistant strain may be induced to increasing branching that in turn, may lead to a higher level of protein and toxin secretion.

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