The effect of flucytosine on the germination of *Candida albicans*

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**Summary**

The effect of flucytosine on the germination of 3 strains of *Candida albicans* in serum was tested. No inhibition of germination was observed and it was concluded that germination did not require the synthesis of DNA.

**Introduction**

Recent studies with *Candida albicans* have revealed that there are at least 2 fundamentally different modes of action for flucytosine (Arai *et al.*, 1977; Polak and Wain, 1977, 1978; Diasio, Myers and Bennett, 1978). The incorporation of the deaminated metabolite 5-fluorouracil (5-FU) into RNA was investigated by Giege and Weil (1970), and the implications of different malfunctionings in the pathways of uptake and incorporation have been described by Polak and Scholer (1975). Another metabolite of flucytosine, 5-fluorodeoxyuridine monophosphate has been detected by Diasio, Bennett and Myers (1978) and they also demonstrated an inhibition of thymidylate kinase. Arai *et al.* (1977) ascribed the morphological changes induced by flucytosine to an inhibitor of DNA synthesis and Polak and Wain (1977, 1978, 1979) have shown that total DNA increase, hyphal nuclear division and thymidine incorporation into DNA are all inhibited by flucytosine. Polak and Wain (1978, 1979) also showed that abnormal yeast cell shapes (Arai *et al.*, 1977; Polak and Wain, 1978) were due to abnormal, accelerated carbohydrate synthesis. These 2 modes of action have been shown to apply to both yeast and hyphal phases of *C. albicans* (Polak and Wain, 1979) and to affect budding in the yeast phase (Arai *et al.*, 1977; Polak and Wain, 1977, 1978). Polak and Wain (1977, 1978, 1979) demonstrated a clear inhibition of hyphal elongation by flucytosine and this inhibition was seen in cultures which had germinated to at least 90% before the addition of the drug. Arai *et al.* (1977) said that the germ-tube formation did not seem to be affected even in the presence of flucytosine. The control of germination by *C. albicans* on inoculation into serum or other biological fluids is still not clear and the action of flucytosine on both macromolecular synthesis and germination during the first few hours after inoculation into serum have therefore been investigated.

**Materials and methods**

Three strains of *C. albicans* sensitive to flucytosine (MIC 0.5 mg/l) and selected for their consistent ability to form hyphae in serum were maintained on Sabouraud's malt agar. Overnight cultures, termed early stationary phase cultures, were inoculated into pooled human serum at 37°C. The serum had been filtered through 0.45 μm pore size membrane filter (Millipore) and contained either flucytosine or cytosine to a concentration of 100 μg/ml. The non-budding cells were inoculated to a concentration of 1·5 × 10⁶/ml and cultured at 37°C with shaking. Samples were taken at intervals to measure germination, hyphal elongation rates and DNA, RNA, protein and carbohydrate amounts (Polak and Wain, 1977, 1979). All experiments were repeated at least twice with all 3 strains.

**Fig. 1.** The germination of *Candida albicans* in serum in the presence of flucytosine.

<table>
<thead>
<tr>
<th>Strain</th>
<th>Time (hr)</th>
<th>Cytosine</th>
<th>Flucytosine</th>
</tr>
</thead>
<tbody>
<tr>
<td>4</td>
<td>1</td>
<td>□</td>
<td>■</td>
</tr>
<tr>
<td>5</td>
<td>2</td>
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</tr>
<tr>
<td>10</td>
<td>3</td>
<td>□</td>
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Results

It was very clear from Fig. 1 that flucytosine did not affect the germination of C. albicans in serum. Moreover, the only immediate effect of flucytosine was on the increase in DNA (Fig. 2). Unlike the exponentially growing hyphae used by Polak and Wain (1977, 1978, 1979) there was not a parallel inhibition of RNA increase in the germinating cells. The effect on the RNA was seen several hours later in the growing, germinated hyphae.

Discussion

Although flucytosine is an inhibitor of both RNA and DNA synthesis in exponentially growing cells, it does not influence the germination in serum.

Investigations with synchronous cultures of both yeast and hyphal phases of C. albicans have shown that yeast budding follows the first period of DNA synthesis whereas hyphal germination precedes it (Wain et al., 1976). The slower manifestation of abnormal carbohydrate synthesis produced by flucytosine is unlikely to affect germination, which occurs within the first 2 hr of inoculation into serum. Similarly, although DNA synthesis is more rapidly affected than carbohydrate synthesis, since germination precedes DNA synthesis, any inhibition of DNA synthesis, however rapid, should not affect germination.

Germination is thus seen to be an event which does not require the synthesis of DNA and which can proceed in the presence of an RNA inhibitor. The lack of similarity between RNA and DNA in response to flucytosine during germination lends further support to the idea that the influence of flucytosine on RNA synthesis is also mediated by the action on DNA. The information for germination is immediately available from the existing DNA in response to some factor or factors which induce germination.

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**FIG. 2.** Rate of hyphal elongation and increase in DNA, RNA, protein and carbohydrate content of *Candida albicans* in the presence of flucytosine (5-FC). Control; ○○○ 5-FC added at 2 hr; ▲▲▲ 5-FC added at 0 hr.

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**References**


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