MEMBRANE FLUIDITY AND LIPID COMPOSITION OF FLUCONAZOLE RESISTANT AND SUSCEPTIBLE STRAINS OF CANDIDA ALBICANS ISOLATED FROM DIABETIC PATIENTS

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Submitted: May 08, 2007; Returned to authors for corrections: November 04, 2007; Approved: February 18, 2008.

ABSTRACT

Ten clinical isolates of Candida albicans, five strains belonging to each of fluconazole resistant and susceptible groups isolated from diabetic patients, were studied for the membrane fluidity and lipid composition. Compared to fluconazole susceptible strains, fluconazole resistant ones exhibited enhanced membrane fluidity as measured by fluorescence polarization technique. The increased membrane fluidity was reflected in the decreased p-values exhibited by the resistant strains. On the other hand, susceptible isolates contained higher amount of ergosterol, almost twice as compared to resistant isolates which might have contributed to their lower membrane fluidity. However, no significant alteration was observed in the phospholipid and fatty acid composition of these isolates. Labeling experiments with fluorescamine dye revealed that the percentage of the exposed aminophospholipid, phosphatidylethanolamine was highest in the resistant strains as compared to the susceptible strains, indicating a possible overexpression of CDR1 and CDR2 genes in resistant strains. The results presented here suggest that the changes in the ergosterol content and overexpression of ABC transporter genes CDR1 and CDR2 could contribute to fluconazole resistance in C. albicans isolated from diabetic patients.

Key-words: membrane fluidity, lipid composition, azole resistance, diabetic patients

INTRODUCTION

Due to the increase in the number of immunocompromised patients, fungal infections have become more common. Among various human fungal pathogens, C. albicans accounts for the majority of systemic infections (27). These pathogenic species of Candida derive their importance not only from the severity of their infection but also from their ability to develop resistance against antifungals (27,35-37). In the recent years, there has been an increase in the incidence of treatment failures in candidiasis patients receiving prolonged fluconazole therapy (3,10,19,29). Ten clinical isolates of Candida albicans, five strains belonging to each of fluconazole resistant and susceptible groups isolated from diabetic patients, were studied for the membrane fluidity and lipid composition. Compared to fluconazole susceptible strains, fluconazole resistant ones exhibited enhanced membrane fluidity as measured by fluorescence polarization technique. The increased membrane fluidity was reflected in the decreased p-values exhibited by the resistant strains. On the other hand, susceptible isolates contained higher amount of ergosterol, almost twice as compared to resistant isolates which might have contributed to their lower membrane fluidity. However, no significant alteration was observed in the phospholipid and fatty acid composition of these isolates. Labeling experiments with fluorescamine dye revealed that the percentage of the exposed aminophospholipid, phosphatidylethanolamine was highest in the resistant strains as compared to the susceptible strains, indicating a possible overexpression of CDR1 and CDR2 genes in resistant strains. The results presented here suggest that the changes in the ergosterol content and overexpression of ABC transporter genes CDR1 and CDR2 could contribute to fluconazole resistance in C. albicans isolated from diabetic patients.

Key-words: membrane fluidity, lipid composition, azole resistance, diabetic patients

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also been observed that the drug resistant ABC protein of mammalian cells (P-glycoprotein[P-gp]), MDR proteins of yeasts (Pdr5p and Yol1p in Saccharomyces cerevisiae and Cdr1p and Cdr2p in C. albicans) can translocate phospholipids between the two monolayers of the plasma membrane, while P-gp has been shown to participate in sterol homeostasis in mammalian cells (5,23,24). Additionally, these drug extrusion pumps are found to be particularly sensitive to the nature and the changes in the physical state of the surrounding lipids (9,17,34-36). The fact that the lipids could also play an important role in azole susceptibilities is becoming apparent from a host of recent studies. It has been shown that some of theazole-resistant C. albicans isolates have exhibited altered membrane sterol compositions (15, 20, 21). Such lipid changes are observed both in clinical and in vitro adapted azole-resistant isolates (13,14) of C. albicans. The Cdr1p and Pdr5p are susceptible to fluctuations in the lipid environment, where functions mediated by these drug extrusion pumps are selectively affected (17,34-37). Since the phospholipid composition of the azole resistant and susceptible strains was not altered, the observed fluctuation in ergosterol content appeared to be responsible for the observed changes in membrane order. The alteration exhibited in membrane fluidity was also associated with the changes in membrane lipid asymmetry of azole resistant and susceptible strains. Our results suggest that membrane alterations associated with fluconazole resistance should be an important consideration in delineating the complex mechanism of drug resistance in C. albicans cells.

MATERIALS AND METHODS

Ten clinical isolates of C. albicans including five azole (fluconazole) resistant, namely DR1, DR2, DR3, DR4 and DR5 with MIC (Minimum inhibitory concentration values expressed in μg/ml) values of 64, 64, 62, 64 and 62 respectively and four azole susceptible isolates, DS1, DS2, DS3, DS4 and DS5 with respective MIC values of 8, 7, 0.5, 0.2, and 2.0 μg/ml were collected from diabetic patients. MIC of the strains was determined using broth microdilution method and reading of the endpoints was taken at 48 h. The MIC test end point was defined as the lowest drug concentration that gave >80% inhibition of growth compared with drug-free controls (MIC<sub>90</sub>). For all experimental studies, the yeast cells were maintained on yeast extract-peptone-dextrose (YEPD) medium at 37ºC.

Ergosterol was extracted as described earlier by Ansari et al. (1) and quantified as described previously by Arthington-Skaggs et al. (2). The steady-state fluorescence polarization measurements on yeast cells were carried out essentially as described earlier Ansari et al. (1). Measurements were carried out on whole cells by using a fluorescent probe, 1,6-diphenyl-1, 3,5-hexatriene (DPH). Fluorescence polarization was measured at excitation and emission wavelengths of 360 and 426 nm, respectively. The measured fluorescence intensities were corrected for background fluorescence and the light scattered from the unlabeled samples.

Phospholipids were extracted separated on thin layer chromatography (TLC) plates and then estimated essentially according to the method given by Ibrahim and Ghannoum (16). Fatty acids were extracted by refluxing with ethanolic KOH as per the method described by Ibrahim and Ghannoum (16). After methanolysis the fatty acids were then estimated by the use of gas liquid chromatography (GLC) according to the procedure adopted by above authors.

Fluorescamine, a fluorescent probe which specifically labels exposed aminophospholipids (7), was used for labeling of phosphatidylethanolamine in the outer mono layer of the resistant and susceptible strains of C. albicans. Cells were harvested in mid-log phase by centrifuging at 3,000 rpm (Beckman model TJ-6) for 5 min at 4ºC and washed two to three times with buffer A (100 mM potassium phosphate-5 mM EDTA, pH 7.5). The harvested cells (0.6 g wet weight) were resuspended in 5 ml of buffer B (100 mM potassium phosphate and 600 mM KCl, pH 8.2) and kept at 4ºC with gentle swirling. Fluorescamine (15.6 mM) in dehydrated dimethyl sulfoxide was added dropwise to the cell suspension with constant gentle swirling. After 30 s the reaction was stopped by adding an equal volume of 1 M ammonia in 600 mM KCl. The cells were centrifuged and washed at 4ºC three to four times till the color of the dye disappeared from the supernatant. Resolved phospholipids and derivatized phosphatidylethanolamine were scraped off from TLC plates, and their phosphate content was estimated (16).

RESULTS AND DISCUSSION

The amount of ergosterol in fluconazole susceptible isolates was much higher, almost twice as compared to fluconazole resistant isolates (Fig.1). Azole resistance in Candida albicans is known to occur primarily through ERG11, which encodes for 14<sup>-</sup>-lanosterol demethylase, an enzyme involved in ergosterol biosynthesis. Inhibition of this step in ergosterol biosynthesis by fluconazole ultimately leads to accumulation of high levels of toxic 14<sup>-</sup>-methylated sterols, causing disruption of membrane structure (11,22). Accordingly, any alteration in Erg11p would affect the affinity of the drug for the target leading to resistance in such isolates (18,25,38,39). In view of these observations, it could be assumed that point mutation in the ERG11 gene encoding demethylase enzyme would contribute to resistance through decreased levels of ergosterol. This fact has been exploited in several reports wherein many point mutations in Erg11p has been identified in response to azoles (22).

The decreased amount of ergosterol in fluconazole resistant strains was further reflected in increased membrane fluidity (low
Fluconazole resistante and susceptible *C. albicans*

p-value) as determined by the use of the fluorescent probe DPH (Fig. 2). Contrary to this, susceptible isolates accordingly had higher p-values which indicated lower membrane fluidity, which is known to have a direct bearing on drug efflux, making itself one of the important molecular mechanisms of antifungal resistance (27,37).

In the present study also, since resistant strains had inverse relationship between ergosterol content and membrane fluidity, this further supported the contention that perhaps the resistance to fluconazole in the present clinical isolates was due to increased efflux pump activity as a result of increased membrane fluidity and decreased ergosterol. This assumption is further strengthened by recent studies where fluconazole resistant isolates have been shown to overexpress efflux pumps namely, *CDR1* and *CDR2*, encoding multidrug efflux transporter proteins belonging to ABC transporter and *CaMDR1* belonging to MFS

![Figure 1. Ergosterol content (%) in azole resistant and susceptible strains of *Candida albicans* isolated from diabetic patients. Mean ergosterol content of the cells is expressed as a percentage of the wet weight of the cells ± the standard deviation of the mean of the three sets of experiments and ergosterol quantitation was done as described in the materials and methods.](image1)

![Figure 2. Membrane fluidity of azole resistant and susceptible strains of *Candida albicans* isolated from diabetic patients as determined by fluorescence polarization and expressed as mean fluorescence polarization “p” values of the cells (p value is inversely proportional to membrane fluidity) ± the standard deviation of the mean of the three sets of experiments.](image2)
class of transporters (26, 32). Cdr1p and Cdr2p represent two major drug extrusion pumps of *C. albicans*, effluxing not only azoles and their derivatives but also a wide variety of structurally unrelated compounds (27). Cannon’s group has recently characterized allelic variations of drug extrusion proteins of *Candida*, representing an interesting possibility contributing to azole resistance in *C. albicans* (4). The phospholipids (Fig.3) and fatty acid (Fig.4) composition of the clinical isolates did not contribute to observed changes in fluidity, since they did not show significant differences between the resistant and susceptible strains (13).

Aminophospholipids are present primarily in the inner monolayer of the membrane and phospholipid translocation between the two monolayers of the lipid bilayer of the plasma membrane of *C. albicans* is maintained by an energy-dependent process mediated by the ABC transporter protein Cdr1p (8, 20, 30). Since Cdr1p functioning is also susceptible to alterations in membrane fluidity (34), we investigated the differences in the asymmetrical distribution of phospholipids by labeling with fluorescamine and quantitated aminophosphopholipids translocated to the outer monolayer in these clinical isolates (Fig.5). Since entire phosphatidylserine is localized predominantly...

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**Figure 3.** Relative percentage of phospholipids in azole resistant and susceptible strains of *Candida albicans* isolated from diabetic patients. Abbreviations used: PI, phosphatidylinositol; PS, phosphatidylserine; PA, phosphatidic acid; PC, phosphatidylcholine; PE, phosphatidylethanolamine; PG, phosphatidylglycerol. Each value in the figure is the mean of three independent experiments.

**Figure 4.** Relative percentage of fatty acids of azole resistant and susceptible strains of *Candida albicans* isolated from diabetic patients. Each value in the figure is the mean of three independent experiments.
Fluconazole resistante and susceptible *C. albicans*

in the cytoplasmic leaflet of the lipid bilayer, therefore the dye could label only externalized phosphatidylethanolamine as small percentage of this aminophospholipid is also present in outer monolayer (8). These experiments revealed that the percentage of exposed phosphatidylethanolamine was higher in the resistant strains, compared to susceptible isolates indicating a higher functional activity of Cdr1p. Higher percentage of exposed phosphatidylethanolamine has earlier been correlated with maximum expression of *CDR1*. In this study, the membrane fluidity of resistant strains was higher and it may be an important factor regulating phospholipid translocation mediated by overexpressed *CDR1*. Dogra et al. (8) have earlier demonstrated the involvement of Cdr1p in phospholipid translocation. The conclusion was based on the fact that an increased externalization of phosphatidylethanolamine in the outer leaflet of the plasma membrane of the mycelial form correlated well with *CDR1* expression which was also more pronounced in mycelia than in the bud form of *C. albicans*, further confirming causal relationship between *CDR1* expression and phospholipid translocation.

The results obtained on these isolates have demonstrated clearly that increased membrane fluidity in resistant strains is probably either due to decreased content of ergosterol or increased overexpression of *CDR1* and *CDR2* genes, and/or both, suggesting that any change in membrane lipid composition and asymmetry would have direct bearing on drug resistance in clinical isolates of *C. albicans*.

**ACKNOWLEDGEMENT**

NNM is thankful to Prof. Rajendra Prasad for providing necessary facility to carry out the entire work in Membrane Biology Lab, School of Life Sciences Jawaharlal Nehru University, New Delhi, India. Prof. Randhir Singh, CSIR Emeritus Scientist, Kurukshetra University, Kurukshetra, India is thanked for critically going through the manuscript and making valuable suggestions.

**RESUMO**

Fluidez e composição lipídica da membrana de cepas de *Candida albicans* resistentes e sensíveis ao fluconazol, isoladas de pacientes diabéticos

Dez isolados clínicos, sendo cinco resistentes e cinco sensíveis ao fluconazol, obtidos de pacientes diabéticos, foram estudados quanto à fluidez e composição química da membrana. Quando comparados aos isolados sensíveis ao fluconazol, os isolados resistentes apresentaram fluidez de membrana aumentada, conforme mensurado pela técnica de polarização fluorescente. A fluidez de membrana aumentada refletiu-se pelos valores mais baixos de p. Por outro lado, os isolados sensíveis continham quantidades mais elevadas de ergosterol, quase o dobro dos isolados resistentes, o que pode ter contribuído para a fluidez de membrana mais baixa. Entretanto, não se observou alteração significativa na composição fosfolipídica e de ácidos graxos nesses isolados. Experimentos de marcação com corante fluorescamina indicaram que a porcentagem de aminofosfolípides e fosfatidiletanolamina expostos foi mais elevada nos isolados resistentes do que nos sensíveis, indicando uma possível superexpressão dos genes *CDR1* e *CDR2* nos isolados resistentes. Os resultados aqui apresentados sugerem que alterações no teor de ergosterol e superexpressão dos genes
**ABC transportadores CDR1 e CDR2 podem contribuir na resistência ao fluconazol em isolados de *C. albicans* de pacientes diabéticos.**

**Palavras-chave:** Fluidez da membrana, composição lipídica, resistência a azois, pacientes diabéticos

**REFERENCES**

1. Ansari, S.; Gupta, P.; Mohanty, S.K.; Prasad, R. (1993). The uptake of amino acids by erg mutants of *Candida albicans*. *J. Med. Vet. Mycol.*, 31, 377-386.

2. Arthington-Skaggs, B.A.; Jradi, H.; Desai, T.; Morrison, C.J. (1999). Mechanism of resistance to azole antifungal agents in *C. albicans* strains. *Anticancer Res.*, 19, 2634-2640.

3. Boken, D.J.; Swindells, S.; Rinaldi, M.G. (1993). Fluconazole–glycoprotein mediated multidrug resistance and the lipid phase of *Candida albicans* superfamily. *J. Biol. Chem.*, 272, 1026-1031.

4. Decottignies, A.; Grant, A.M.; Nichols, J.W.; Wet, H.D.; McIntosh, C.B.; Goffeau, A. (1998). ATPase and multidrug transport activities of the overexpressed yeast ABC protein Yorlp. *J. Biol. Chem.*, 273, 12612-12622.

5. Debry, P.; Nash, E.A.; Nekalson, D.W.; Metherall, J.E. (1997). Role of translocases in the generation of fluconazole susceptibility of *Candida albicans*. *J. Clin. Microbiol.*, 35, 3332-3337.

6. Diaz, C.; Schroit, A.J. (1996). Role of translocases in the generation of phosphatidylserine asymmetry. *J. Membr. Biol.*, 151, 1-9.

7. Dogra, S.; Krishnamurthy, S.; Gupta, V.; Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

8. Dogra, S.; Krishnamurthy, S.; Gupta, V.; Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

9. Dey, P.; Nash, E.A.; Neakolson, D.W.; Metherall, J.E. (1997). Role of multidrug resistance P-glycoproteins in cholesterol esterification. *J. Biol. Chem.*, 272, 1026-1031.

10. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

11. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

12. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

13. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

14. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

15. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

16. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

17. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

18. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

19. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

20. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

21. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.

22. Dogra, S.; Krishnamurthy, S.; Gupta, V., Dixit, B.L.; Gupta, C.M.; Sanglard, D.; Prasad, R. (1999). Asymmetric distribution of phosphatidylethanolamine in *C. albicans*: possible mediation by *CDR1*, a multidrug transporter belonging to ATP binding cassette (ABC) superfamily. *Yeast*, 15, 111-121.
ergosterol interaction are important determinants of multidrug resistance in *Candida albicans*. Antimicrob. Agents Chemother, 48: 1778-1787.

36. Prasad, T.; Saini, P.; Gaur, N.A.; Vishwakarma, R.A.; Khan, L.A.; Haq, Q.M.; Prasad, R. (2005). Functional analysis of *CaIPT1*, a sphingolipid biosynthetic gene involved in multidrug resistance and morphogenesis of *Candida albicans*. Antimicrob. Agents Chemother., 49: 3442-3452.

37. Prasad, T.; Chandra, A.; Mukhopadhyay, C.K.; Prasad, R. (2006). Unexpected link between iron and drug resistance of *Candida* spp.: iron depletion enhances membrane fluidity and drug diffusion, leading to drug-susceptible cells. *Antimicrob. Agents Chemother.*, 50: 3597-3606.

38. White, T.C.; Holleman, S.; Dy, F.; Mirels, L.F.; Stevens, D.A. (2002). Resistance mechanism in clinical isolates of *Candida albicans*. Antimicrob. Agents Chemother., 46, 1704-1713.

39. Xiao, L.; Madison, V.; Chau, A.S.; Loebenberg, D.; Palermo, R.E.; McNicholas, P.M. (2004). Three-dimensional models of wild-type and mutated forms of cytochrome P450 14-sterol demethylase from *Aspergillus fumigatus* and *Candida albicans* provide insights into posaconazole binding. *Antimicrob. Agents Chemother.*, 48, 568-574.