

# Effects of chemically synthesized Azole compounds on Clinical isolates of Vaginal Candidiasis, in comparison with commercially available anti-fungal drugs

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## Abstract

Candida species is observed as the most common cause of 'Opportunistic Mycoses' worldwide. Candida species is a member of normal flora of skin, mouth, vagina, and stool. Infections caused by Candida species are in general referred to as Candidiasis. Vaginal Candidiasis is a fungal or yeast infection of the vulva and/or vagina, which causes a smelly, thick, white-yellow discharge, which might be accompanied by itching, burning and swelling. It can also can make walking, urinating or sex very painful.

The Genus Candida includes around 154 species. While Candida albicans is the most abundant and significant species; importantly, there has been a recent increase in infections due to non-albicans Candida species along with increasing drug resistance.

The present study includes isolation of Candida species from the patients suffering from vaginal candidiasis followed by identification and differentiation between different Candida species by their morphological studies, microbiological analysis and biochemical activities, which are: C. albicans, C. tropicalis, C. glabrata & C. krusei.

The present study also includes a comparative study of anti-microbial activities of commercially available anti-fungal drugs as well as 22 novel chemically synthesized 1% DMSO soluble compounds (belonging to Azoles) by "Disc Diffusion Method" according to NCCLS Guidelines. Out of which 5 compounds were found to be remarkably effective, especially AntiC-15 was found to be the most potent and excellent among all; which exhibits its inhibitory effects against some drug resistant isolates also. It is sent for further study in order to use it at commercial level.

## INTRODUCTION

Candida species is yeast and the most common cause of opportunistic mycoses worldwide. It is also a frequent colonizer of human skin and mucous membranes. Candida species is a member of normal flora of skin, mouth, vagina, and stool. Infections caused by Candida species are in general referred to as Candidiasis. The clinical spectrum of Candidiasis is extremely diverse. Almost any organ or system in the body can be affected. Candidiasis may be superficial and local or deep-seated and disseminated. Candidiasis is mostly an endogenous infection, arising from overgrowth of the fungus inhabiting in the normal flora. However, it may occasionally be acquired from exogenous sources such as catheters or prosthetic devices or by person-

to-person transmission such as Oral Candidiasis in neonates of mothers with Vaginal Candidiasis or Endophthalmitis following corneal transplantation from an infected donor.

Vaginal Candidiasis is a fungal or yeast infection of the vulva and/or vagina. It causes a smelly, thick, white-yellow discharge that might be accompanied by itching, burning and swelling. It can also can make walking, urinating or sex very painful.

The Genus Candida includes around 154 species. Among these, very few are most frequently isolated in human infections. While Candida albicans is the most pathogenic and most commonly encountered species among all due to its ability to adhere to host tissues, produce aspartyl

proteases & phospholipase enzymes and transformation from yeast to hyphal phase, which are the major determinants of its pathogenicity. *Candida tropicalis*, *Candida glabrata*, *Candida parapsilosis*, *Candida krusei* and *Candida lusitanae* are also isolated as causative agents of Candidiasis.

Importantly, there has been a recent increase in infections due to non-albicans *Candida* species along with increasing drug resistance. For the same, the present study also includes a comparative study of anti-microbial activities of commercially available anti-fungal drugs as well as 22 newer chemically synthesized 1% DMSO soluble compounds (belonging to Azoles).

## **MATERIAL & METHODS**

### **STUDY GROUP**

The study group includes different patients suffering from Vaginal Candidiasis with varying degree of severity.

### **SPECIMEN COLLECTION:**

- Vaginal Swab
- Vaginal discharge

### **DIRECT EXAMINATION:**

- Gram staining
- Lacto phenol cotton blue stain

### **ISOLATION, IDENTIFICATION AND DIFFERENTIATION BETWEEN SPECIES:**

#### **ISOLATION FROM OTHER YEASTS:**

- Morphology on Corn-meal Tween 80 agar, Capsule production, Urease activity, Growth pattern in Sabouraud broth and Fermentation or Assimilation profiles of different simple carbohydrates help in identification & differentiation of *Candida* species from other yeasts.

### **CULTURAL TECHNIQUES:**

- Use of different media like; Saboraud's Dextrose Chloramphenicol Agar, Blood agar, Corn-meal Tween 80 agar, Hi-Chrom *Candida* agar
- Urease test

### **GERM – TUBE TEST:**

- (REYNOLDS-BRUDE PHENOMENON)

### **CARBOHYDRATE UTILIZATION PROFILES:**

- Fermentation of simple carbohydrates
- Assimilation of simple carbohydrates

### **ANTI-MICROBIAL ACTIVITY**

In present study, we have analyzed 22 heterocyclic chemical compounds (1% DMSO soluble); synthesized at Chemistry Department, Saurashtra University at Rajkot by preparing discs of these compounds with different concentrations [5µg/disc, 10µg/disc, 15µg/disc] using sterile discs from Hi-Media. It is followed by analyzing their anti-microbial activities on clinical isolates by “Disc Diffusion Method” according to NCCLS guidelines in comparison with commercially available anti-fungal drugs [Anti-fungal antibiotic discs were used]. Prepared discs of Azoles: Fluconazole, Itraconazole, Ketoconazole; Polyenes: Amphotericin B, Nystatin were used.

### **NOVEL HETEROCYCLIC COMPOUNDS**

**Effects of chemically synthesized Azole compounds on Clinical isolates of Vaginal Candidiasis, in comparison with commercially available anti-fungal drugs**

**Figure 2**

Table: 1 indicates identification of clinical isolates of different species, isolated from patients suffering from Vaginal Candidiasis.

No.	Isolate Name	Biochemical characteristics														Candida species
		Morphology			Fermentation				Assimilation							
		PH	GT	Chs.	G	S	L	M	Gal	G	S	L	T	R	C	
1	ClnY-1	+	-	-	+	✓	-	+	+	+	-	+	-	+	C tropicalis	
2	ClnY-2	+	-	-	+	✓	-	+	+	+	-	+	-	+	C tropicalis	
3	ClnY-3	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
4	ClnY-4	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
5	ClnY-5	+	-	-	+	✓	-	+	+	+	-	+	-	+	C tropicalis	
6	ClnY-6	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
7	ClnY-7	-	-	-	+	-	-	-	-	+	-	-	+	-	C glabrata	
8	ClnY-8	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
9	ClnY-9	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
10	ClnY-10	+	-	-	+	✓	-	+	+	+	-	+	-	+	C tropicalis	
11	ClnY-11	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
12	ClnY-12	+	-	-	+	✓	-	+	+	+	-	+	-	+	C tropicalis	
13	ClnY-13	+	-	-	+	✓	-	+	+	+	-	+	-	+	C tropicalis	
14	ClnY-14	+	-	-	+	✓	-	+	+	+	-	+	-	+	C tropicalis	
15	ClnY-15	-	-	-	+	-	-	-	-	+	-	-	+	-	C glabrata	
16	ClnY-16	-	-	-	+	-	-	-	-	+	-	-	+	-	C glabrata	
17	ClnY-17	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
18	ClnY-18	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
19	ClnY-19	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
20	ClnY-20	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
21	ClnY-21	+	-	-	+	✓	-	+	+	+	-	+	-	+	C tropicalis	
22	ClnY-22	-	-	-	+	-	-	-	-	+	-	-	+	-	C glabrata	
23	ClnY-23	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
24	ClnY-24	+	+	+	+	-	-	+	✓	+	+	-	+	-	Calbicans	
25	ClnY-25	+	-	-	+	✓	-	+	+	+	-	+	-	+	C tropicalis	
26	ClnY-26	+	-	-	+	-	-	-	-	+	-	-	-	-	C krusei	
27	ClnY-27	-	-	-	+	-	-	-	-	+	-	-	+	-	C glabrata	

Note: The basic structure remains the same in all 22 compounds; while 'R' is variable.

During the experiment, One disc having 1% DMSO/disc was kept in each plate as a standard to check any inhibitory effect of the solvent as well as one plate kept as a control inoculated with the respective isolate only, without any disc.

**OBSERVATION & RESULTS**

The aim of the present study was to investigate pathogenic yeast organisms, mainly Candida albicans and other non-albicans group from patients suffering Vaginal Candidiasis.

Total 27 isolates of Candida spp. were analyzed for isolation, identification & differentiation using different media as well as the biochemical activities (like; Germ-tube formation, Urease test, Sugar fermentation tests, Sugar assimilation profile), along with the sensitivity profile of each isolate against commercial drugs as well as synthetic chemical compounds. [All the sets were performed in duplicates for each isolate].

Out of 22 heterocyclic compounds [belonging to Azoles], 5 compounds were found to be remarkably efficient; especially, "AntiC-15" was found to be the most potent and excellent among all, giving anti-microbial activity even at 10µg/disc.

Many isolates were found to be relatively more sensitive to commercial drugs, were also found to be fairly good susceptible to the heterocyclic compounds. Certain isolates have been found to be relatively more resistant to commercially used drugs but still found to be susceptible at least to AntiC-15.

Table: 1 includes identification of clinical isolates of different Candida species, isolated from patients suffering from Vaginal Candidiasis.

Graph: 1 indicates a comparative profile of different Candida species isolated from patient suffering from vaginal candidiasis.

Table: 2.1 & 2.2 includes a comparative analysis of anti-microbial effects of commercially available drugs as well as 1% DMSO soluble synthetic heterocyclic chemical compounds (belonging to Azoles) respectively against clinical isolates of different Candida species from vaginal candidiasis.

{image:2}

Abbreviations used in the table:

- PH : Pseudohyphae
- GT : Germ-tube formation
- CHS : Chlamyospore
- G : Glucose
- S : Sucrose
- L : Lactose
- M : Maltose
- Gal : Galactose
- T : Trehlose
- R : Reffinose

C : Cellobiose

+ : Positive, - : Negative, : Variable

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Abbreviation used:

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Abbreviations used:

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Photographs

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## DISCUSSION & CONCLUSION

Out of 27 isolates; *C. albicans* constitutes 44.44%, *C. tropicalis* constitutes 33.33%, *C. glabrata* constitutes 18.51%, *C. krusei* constitutes 3.70%; with slightly increasing non-*albicans* species [normally, *C. albicans* constitutes 60-70% and *C. krusei* constitutes only >1%].

Along with increasing number of non-*albicans* species; interestingly, the drug resistance to commercially available drugs also increases as well as each drug is having certain mild to severe side effects; so, now there is extreme need to evaluate and to use newer and more potent drugs.

Among all isolates, the overall general sensitivity gradation is as follows:

*C. albicans* > *C. tropicalis* > *C. glabrata* > *C. krusei*

However, among the anti-fungal drugs, the overall sensitivity gradation is as follows:

Amphotericin-B > Itraconazole > Fluconazole > Nystatin > Ketoconazole

Out of 22 chemical compounds, 5 compounds were found to be remarkably efficient; especially, AntiC-15 was found to be the most potential and excellent among all.

CanV-16 (*C. glabrata*), CanV-24 (*C. albicans*) and CanV-27 (*C. glabrata*) were found to be not sensitive to all the anti-fungal drugs; still found to be sensitive to the chemical compounds, at least to AntiC-15. However, CanV-26 (*C. krusei*) was found to be resistant to all; i. e., commercially

available drugs as well as heterocyclic compounds; still, AntiC-15 exhibited slight inhibition even at 10µg/disc (4 mm) and 15µg/disc (9 mm).

AntiC-15 is sent for further analysis in order to use it at commercial level against candidiasis.

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