

# Typing and Antifungal Susceptibility of *Candida* Spp. Isolated from Clinical Samples

Senay Ozturk Durmaz\*

Department of Infectious Diseases and Clinical Microbiology, Turkey

\*Corresponding author: Senay Ozturk Durmaz, Antalya Kepez State Hospital, Department of Infectious Diseases and Clinical Microbiology, Turkey



## ARTICLE INFO

Received: 📅 January 30, 2019

Published: 📅 February 13, 2019

**Citation:** Senay Ozturk Durmaz. Typing and Antifungal Susceptibility of *Candida* Spp. Isolated from Clinical Samples. Biomed J Sci & Tech Res 14(4)-2019. BJSTR. MS.ID.002570.

## ABSTRACT

### Summary

In our study, we aimed to identify 100 yeast species fungus isolated from clinical samples sent to microbiology laboratory of our hospital and to determine in vitro activity of various antifungals to these species. 55 of the *Candida*s were isolated from urine, 18 from respiratory tract, 12 from body fluids, 11 from blood, and 4 from other (3 wound 1 vaginal) samples. 46 of them were *C. albicans*, 19 were *C. glabrata*, 14 were *C. parapsilosis*, 11 were *C. kefyr*, 9 were *C. tropicalis*, and 1 was *C. krusei*. *C. albicans* is still the most commonly isolated type of *Candida*. Amphotericin B remains one of the most effective antifungal agents. There is increased resistance to fluconazole. Increased resistance in all *Candida* species is an important treatment problem, especially in people with low immune resistance. For this reason, identification of species, antifungal testing and antifungal agent should be treated carefully.

**Keywords:** *Candida*; Antifungal Susceptibility; Fluconazole Resistance

## Introduction

*Candida* species are the most common fungal pathogen affecting humans. These organisms have a wide spectrum of disease ranging from noninvasive superficial infections to deep-tissue infections. The frequency of both out-of-hospital and nosocomial *Candida* infections increases, especially due to broad-spectrum antibiotic use, intravascular devices and the population of immunosuppressive patients [1,2]. In *Candida* species, blastoconidia, pseudohyphae, chlamydospore and tube germ formation, ascospore formation are important in the definition of species. Colonies of *Candida* species form colonies smelling yeast, with a smooth or wrinkled edge, moist, creamy appearance. They reproduce by means of sexual and asexual spores and are classified based on reproductive patterns [3-5]. In the ranking of nosocomial fungal infections, urinary tract infections are the first and fungemia is the second. It is known that urinary catheter application increases the fungus [6]. Diabetes mellitus, elevated glucose level increases fungal growth without tissue invasion. *Candida* colonization in severe burn may develop into tissue invasion. Increased estrogen and vaginal glycogen levels

during pregnancy cause vaginal colonization. Hyperalimentation fluids facilitate the intravenous hyperglycemic environment and facilitate *Candida* infections. Intravascular catheters, pressure monitoring devices, prosthetic heart valves, and pacemaker placement may lead to disseminated candidiasis. The suppression of flora bacteria with broad spectrum antibiotics can lead to *Candida* infections [7-9].

## Materials and Methods

One hundred *Candida* strains isolated from different clinical samples from SSK Izmir Training Hospital Microbiology Laboratory were included in the study. *C. albicans* ATCC 90028, *C. krusei* ATCC 6258, *C. parapsilosis* ATCC 90018 were used as standard strain. Resistance states of the identified *Candida* species to amphotericin B, fluconazole, voriconazole, itraconazole RPMI 1640 (1.5% agar) medium (ANGUS) and E test strips 7.0 with buffered pH of MOPS (3-N-morpholinopropanesulfonic acid) and E test strips (AB Biodisk, Solna), Sweden). Patients with yeast were also evaluated for risk factors. Specimens were planted in Sabouraud's Dextrose

Agar (SDA) for primary isolation. From the cultures that were understood to be pure, passages were made to the slanted SDA tubes for use in the advanced stage of the identification. Yeasts were added in 0.5 ml human serum to a small portion of the colony to be tested and mixed. It was incubated at 37 °C for 2.5-3 hours. The specimens were examined by microscope at +40 magnification and the filament shaped structures were evaluated as germ tube. The germ-producing yeast strains were defined as *C. albicans* [10-12].

## Results

Of the 100 species of yeasts identified from the samples, 55 were isolated from urine, 12 from various body fluids, 18 from respiratory tract, 11 from blood and 4 from other specimens (3 from wound, 1 from vagina). 46 of the yeast species were named *C. albicans*, 19 were *C. glabrata*, 14 were *C. parapsilosis*, 11 were *C. keyfr*, 9 were *C. tropicalis* and 1 were *C. krusei* (Table 1). Germ tube test

was found to be positive (100%) in all yeasts that were typed as *C. albicans*. In 90 of 100 licensed yeasts, sensitivity tests were tested by E test method. The distribution of abstracted fungi according to predisposing factors; 4% had diabetes + catheters, 6% had fluconazole use, 17% had antibiotic use + had an underlying disease, urinary catheter or any catheter and received immunosuppressive treatment. In urine samples, fluconazole 41%, itraconazole 47%, and voriconazole resistance 75% were found in *C. albicans*. 34% fluconazole, 47% itraconazole and 6% voriconazole resistance were found in the Candidias detected in urine except *C. albicans*. Fluconazole was isolated in 52% of the *C. albicans* isolated from the urine samples and itraconazole resistance was found in 30% and voriconazole resistance was 73%. In the non-*C. albicans* yeasts isolated from the urine samples, 22% fluconazole and 33% itraconazole and 5% voriconazole resistance were found. Amphotericin B resistance was not detected in any of the cases (Table 2).

**Table 1:** Distribution of candidias according to their location.

Distribution of Candidias According to their Location					
Species	Urine (n: 55)	Respiratory Tracts (n: 18)	Body Fluids (n: 12)	Blood (n: 11)	Other (n: 4)
<i>C.albicans</i>	21	11	5	6	3
<i>C. glabrata</i>	15	2	2	0	0
<i>C.tropicalis</i>	6	0	0	3	0
<i>C.parapsilosis</i>	5	3	3	2	1
<i>C.keyfr</i>	8	1	2	0	0
<i>C. krusei</i>	0	1	0	0	0
<i>Toplam</i>	55	18	12	11	4

**Table 2:** Candida species and resistance distributions according to samples.

Candida Species and Resistance Distributions According to Samples			
Albicans-Non Albicans	Fluconazole	Itraconazole	Voriconazole
<i>C.albicans</i> (n: 40)			
Urine: 17	7(%41)	8(%47)	12(%75)
Other than urine (n: 23)	12(%52)	7(%30)	17(%73)
Other than <i>C. albicans</i> (n: 50)			
Urine: 32	11(%34)	15(%47)	2(%6)
Other than urine (n: 18)	4(%22)	6(%33)	1(%5)

## Discussion

Nosocomial candida infections are an important cause of morbidity and mortality in immunocompromised patients. Candida species are isolated from 15% of nosocomial infections, 86% of all nosocomial fungal infections and 8-10% of nosocomial blood circulatory infections. In a study of candida cases in urine; 60.8% *C. albicans*, 12.5% *C. glabrata*, 11.2% *C.tropicalis*, 6% *C. parapsilosis* isolated (6). In our study, 38% of the 55 candida detected in urine were *C.albicans*, 27% *C.glabrata*, 15% *C.keyfr*, 10% *C. topicalis* and 10% *C. albicans* and *C. glabrata*. It was reported that *C. glabrata* and *C. parapsilosis* fungemias increased 4-6 times in Taiwan in 1981-1993 related to candidiasis. According to the results of one

year study of 8 European countries; *C. albicans* were shown to be isolated in 55%, *C. glabrata* in 20%, *C. krusei* in 0-20% (10,11). In our study, 54.5% *C. albicans*, 27% *C. topicalis* and 18% *C. glabrata* were found in candidemia and the results were similar. In 46 of 40 *C.albicans* cases, susceptibility to E test was evaluated and resistance to voriconazole was found in 16 (40%), itraconazole in 12 (30%) and voriconazole in 29 (70%).

Of the total of 90 strains, 28 (31%) were resistant to fluconazole and 22 (24%) to itraconazole and 32 (35%) were resistant to voriconazole, while no resistance to amphotericin B was detected. Yilmaz et al. [13] described the candida isolated from various clinical specimens using API CAUX system and

conventional methods and studied the antifungal sensitivity E test method; They found 44% itraconazole, 42% fluconazole and 2% amphotericin B resistance. Degerli et al. [14] have performed the antifungal sensitivity of 60 strains, including 38 *C. albicans* and 22 nonalbicans, by E test method. In this study, *C. albicans* was reported to be resistant to 34% fluconazole, 32% itraconazole, 3% amphotericin. Zer et al. [15] described 115 (56%) of 205 candida strains as *C. albicans* with API and found 23% of *C. albicans* with E test and 27% of all Candida isolates in the study found resistant to fluconazole.

## Conclusion

In our study, the most common type of candida was *C. albicans*. In addition, five different species were identified except *C. albicans*. This is a sign of infections that may occur with other candida species other than *C. albicans*, especially *C. glabrata* in the future. The determined fluconazole resistance demonstrates the need for more detailed studies to be conducted in clinical-laboratory cooperation in order to be more careful in the use of this agent and to determine the in vivo response.

## References

- Warren NG, Hazen KC (1999) Candida, Cryptococcus and Other Yeast of Medical importance. In: Murray PR, Baron EJ, Pfaller MA, Tenover FC, Tenover FC (Eds.), Manual of clinical Microbiology. American Society for Microbiology, Washington DC, USA, pp. 1184-1199.
- Maenza JR, Merz WG (1998) Candida albicans and related species. In: Gorbach SL, Bartlett JG, Blacklow NR (eds), Infectious Diseases WB, Saunders Company, Philadelphia, USA, pp. 2313-2322.
- Koneman EW, Allen SD, Janda WM (1977) Mycology Color Atlas and Textbook of Diagnostic Microbiology (5<sup>th</sup> edn.), Lippincott Co, Philadelphia, USA, pp. 983-1069.
- Warren NG, Shadomy HJ (1995) Candida, cryptococcus and other yeast of medical importance. In: Murray PR, Baron EJ, Phaller MA, Tenover FC, Tenover FC (Eds.), Manual of Clinical Microbiology (6<sup>th</sup> edn.), ASM press, Washington DC, USA, pp. 723-737.
- Dixon DM, Rhodes JC, Framtling RA (1999) Taxonomy, Classification or Morphology of the fungi. In: Murray PR, Baron EJ, Phaller MA, Tenover FC, Tenover FC (Eds.), Manual of Clinical Microbiology. American Society for Microbiology, Washington DC, USA, pp. 1161-1166.
- Kiraz N, Akgün Y (2000) The distribution of Candida spp. Causing urinary tract infections, Mycosis pp. 43-237.
- John E, Edwards JR (1995) Candida species. In: Mandell GL, Bennett JE, Dolin R (Eds.), (4<sup>th</sup> edn.), Principles and practise of infectious disease. Churchill Livingstone, New York, USA, pp. 2289-2306.
- Erbakan N (1989) Fungal diseases of the skin. Publisher Clinics Turkey, Ankara, Turkey, p. 1-90.
- Kaufman R (1988) Establishing a correct diagnosis of vulvovaginal infection. Am J Obstet Gynecol 158(4): 986-988.
- Grillot R (1992) Epidemiological survey on candidemia in Europe, mycology newsletter 2: 11-13.
- Yhu chening H, Tzou Yien L, Rey-in I, Yi Hong C (1999) Candidemia in special care nurseries: Comparison of albicans and parapsilosis infection trends in invasive fungal infections 5. Abstract book 153: 14-16.
- Helvacı S, Gedikoğlu S, Mıstık R (1992) Germ tube test for diagnosis of candida albicans. Infec Derg 6: 141-143.
- B Yılmaz, A Botrel, S Gençer, S Özer Çeşitli klinik örneklerden (2002) Isolation of isolated candida strains and detection of amphotericin B, fluconazole and itraconazole sensitivities by E test. XXX. Türk Mikrobiyoloji kongresi, 30 Eylül-05 Ekim 2002 Antalya Başak matbaacılık pp. 313.
- S Sürücüoğlu, K Değerli, S Kuzutepe, Ö Tünger, E Aktaş, B Özbakkaloğlu (2002) Examination of antifungal resistance in candida isolates which are the causative agents of invasive infection XXX. Türk Microbiology Congress 30 September-05 October 2002 Antalya Başak, pp. 314.
- Yasemin Zer (2002) The identification and antifungal susceptibility of the candida strains isolated from patients in the Iclal intensive care unit Turkish Journal of Microbiology 32: 230-234.

ISSN: 2574-1241

DOI: 10.26717.BJSTR.2019.14.002570

Senay Ozturk Durmaz. Biomed J Sci & Tech Res



This work is licensed under Creative Commons Attribution 4.0 License

Submission Link: <https://biomedres.us/submit-manuscript.php>



### Assets of Publishing with us

- Global archiving of articles
- Immediate, unrestricted online access
- Rigorous Peer Review Process
- Authors Retain Copyrights
- Unique DOI for all articles

<https://biomedres.us/>