

Oral Carriage Rate of *Candida* Species in Diabetic Patients

*Faris Abdul Kareem Khazal Mb Chb Dm Cabm ** Adnan Mahran PhD
*** Huda Hadi Al-Hasnawi MSC

ABSTRACT

Background: It is well known that oral carriage of *Candida* species increase in many situations, like obesity, debility, leukemia, viral infection, use of certain drugs in addition to diabetes mellitus.

Objective: find the relation between diabetes and its control on oral carriage of *Candida*.

Methods: Thirty four hundred oral swabs from diabetic patients 67% are females and 33% are males, 41.7% are type 1 diabetes and 58.3% are type 2. different culture media are used.

Results: we found that 37.9% of diabetics had oral carriage, older age group had more but the

difference is not significant statistically $P > 0.05$, in addition females carry more *Candida* than males $P < 0.05$, while type of diabetes had no effect on oral carriage we found that control of diabetes had significant effect $P < 0.01$.

Conclusion: There is high carriage rate of *Candida* species in oral cavity of diabetic patients 37.9%, the rate of carriage increase in females and those with poor control. *Candida albicans* is the most common type with all the associated factors.

Key Words: Diabetes, oral carriage, *Candida*

Al-Kindy Col Med J 2006; Vol.3 (1): P9-12

Introduction

Candidiasis is an acute or chronic, superficial or disseminated mycosis caused by *Candida* species⁽¹⁾.

They are opportunistic fungi which fail to induce disease in most normal hosts but may do so in those with impaired defense system⁽²⁾.

The most common predisposing factors are, under-nutrition, debility, poor sanitation, iron deficiency anemia, obesity, age, hematological malignancies, viral infections HIV, HSV, CMV, neutropnia, pregnancy, drugs e.g. antibiotics, steroid, oral contraceptive drugs, cytotoxics, chronic infections, tropical areas, trauma of the skin, and diabetes mellitus⁽³⁾.

The history of Candidiasis dates back to fourth century BC when Hippocrates describe oral thrush, the initial discovery of the organism was made in 1839 by Langenback⁽⁴⁾.

The incidence of Candidiasis was increased since 1940 with the advent of antibiotics and other drugs and later on AIDS infection.

In diabetes increase carriage rate is recognized and was thought due to increase glucose availability⁽⁵⁾, glycation induced alteration in T cell function⁽⁶⁾, affected intracellular killing of granulocyte⁽⁷⁾, and impair opsonisation⁽⁸⁾.

This can be aggravated by wearing of artificial denture (mechanical damage)⁽⁹⁾.

There is many stereotypes of *Candida*: *albicans*, *tropicalis*, *krusei*, *glabrata*, *kefyr*, *parapsilosis*, *clustitaniae*, *norvegensis*, and *viswanathii*⁽¹⁰⁾.

Candida proliferates in superficial layer of squamous mucosa; extra-cellular aspartic proteinase may play a role in virulence. Toxin

production, tissue invasion and hypersensitivity state are assumed disease processes⁽¹¹⁾.

C. albicans is a normal commensal of the gastrointestinal and genitourinary tracts of the human being.

The frequency of carriage rate of *Candida* was highest in the mouth followed by vulvovaginal and then anorectal regions. *C. albicans* is the most common isolates followed by *tropicalis* and *glabrata*⁽¹⁰⁾.

Methods

Specimens of 340 mouth swabs were taken from diabetic patients, 228(67%) are females and 112 (33%) are males with an age between 10 and 60 years, attending diabetic clinic in the teaching hospital in Najaf from Jan. 2002 till Sept. 2002.

The type of diabetes was stated according to WHO criteria of diagnosis of diabetes 1998.

41.7% of those are type 1 diabetes and 58.3% are type 2.

Swab collection done by using disposable cotton swab rubbed over the tongue, palate, and buccal mucosa, then dipped in sterile brain heart infusion broth transported to the lab.

State of control of diabetes calculated as follow; Good when fasting plasma glucose less than 115 mg/dl. Accepted when less than 150mg/dl and poor when above 150mg/dl.

Culture media:

The following types are used:

1. Brain heart infusion broth.
2. Sabourauds-dextrose broth with chloramphenicol.
3. Eosin methylene blue agar.
4. Urea agar base.

5. Corn meal agar.

6. Sugar fermentation basal medium.

Swab incubated at 37°C for 24 hours, next day incubated on both Sabouraud's dextrose agar and eosin methylene blue agar for 1-3 days for visible growth of *Candida* colonies, if no growth more incubation for 5-10 days, if no growth discarded as negative result.

Identification and confirmation of type of *Candida* species done according to conventional culture and biochemical tests

Statistical analysis:

Performed using chi square (χ^2) and standard normal distribution (z) tests

Results

Positive culture results found in 129 patients (37.9%) and the types of isolates found are *albicans*, *tropicalis*, *krusei* and *glabrata*.

In **Table-1** prevalence rate of oral carriage of *Candida* species in different age groups was 37.9%, highest in age group 41-60 (41%) and least in age group less than 20 years (24%), statistical analysis show that difference is not significant statistically ($P > 0.05$).

In **Table-2** influence of sex on oral carriage rate of *Candida* species show that (46%) of females had positive culture result in comparison to (21%) in males and this result is significant statistically ($P < 0.01$).

Table-3 show that in both males and females *Candida albicans* is the most common isolate (13% and 32% respectively) and *Candida tropicalis* came second (7.1% and 5.2%) and *Candida glabrata* (0% and 3.9%), and the difference between carrier rate of *albicans* and other types in females is significant ($P < 0.05$).

Table-5 shows no difference of carriage rate between type 1 and type 2 diabetes ($P > 0.05$).

In **Table-5** a comparison between different species of *Candida* in both type 1 and type 2 diabetes show that *Candida albicans* is still the most common in both (24% and 27%) than the *tropicalis*, *krusei* and *glabrata* with no significant difference in both types of diabetes ($P > 0.05$).

that *Candida albicans* is still the most common in both (24% and 27%) than the **Table-6** show relation of glycemic control to oral carriage rate

of *Candida* and its different comparison to 11% and this difference is significant ($P < 0.01$).

Species in different age groups types. Those patients with poor control had much more possibility to carry *Candida* orally than those with good control (45.8% in *ropicalis*, *krusei* and *glabrata* with no significant difference in both types of diabetes ($P > 0.05$)).

Discussion

Oral carriage rate of *Candida* species in normal individuals is about 25-30%⁽¹⁰⁾ while in Iraq lower results was noticed by alhussaini 14%⁽¹²⁾ and higher results by amin 36%⁽¹³⁾. Several previous studies had shown that the prevalence of *Candida* species infection to be greater among diabetics than normal persons⁽¹⁴⁾, as reported in **Table-2**, prevalence rate of oral carriage is 37.9% with no effect of age on carriage rate ($P > 0.05$), in spite that higher rate of carriage in age group 41-60, this is also found by tapper-jones *et al* 1981⁽⁹⁾, while smits B.J. *et al*

1996⁽¹⁵⁾ claim increase prevalence of *Candida albicans* with advancing age.

Influence of sex on oral carriage rate of *Candida* species show that females are more susceptible to oral carriage than males in our study ($P < 0.01$) This is also shown by Barlow⁽¹⁴⁾ and Anendorf⁽¹⁶⁾ and the possible cause is hormonal difference.

Prevalence rate of different species of *Candida* in both sexes show that *Candida albicans* is the most common of them in both females and males, then *tropicalis* and after that *krusei* and *glabrata*. and the difference of carriage rate of *Candida albicans* to other species in females is significant ($P < 0.05$) while it is not significant in males and we could find no studies to compare.

Both type 1 and 2 diabetes had the same carriage rate of *Candida* species ($P > 0.05$). *Candida albicans* still is more common than other types of *Candida* in both types of diabetes while *Candida tropicalis* is the second and then *krusei* and *glabrata* which are less common, so the type of diabetes had no effect on oral carriage rate of *Candida* ($P > 0.05$), a significant difference was reported between *Candida albicans* and other species of *Candida* ($P < 0.05$)

(Table 1) The Prevalence Rate of Oral Carriage of Candida

Age group	no. of patients	no. of isolates	%	P value
<=18	29	7	24	P>0.05
21-40	69	23	33	
41-60	203	85	41	
>60	40	14	35	
Total	340	129	37.5	

(Table 2) The

the Oral Carriage Rate of Candida Species

Influence of Sex on

Sex	no.	Oral Carriage	%	P value
Female	228	105	46.0	P<0.01
Male	112	24	21.4	
Total	340	129	37.9	

(Table 3) The Number and Percentage of Different Species of Candida Isolates in Males and Female

Candida Species	Female		Males		P value
	No.=22	%	No.=11	%	
	8		2		
C.albicans	75	32.9	15	13.4	P<0.05
C.tropicali	12	5.3	8	7.1	
C.krusei	9	3.9	1	0.9	
C.glabrata	9	3.9	0	0	

(Table4) Influence of Type of DM on Oral Carriage Rate of Candida Species

Type of DM	No.	Oral carriage	%	P value
I	142	52	36.9	P>0.05
II	198	77	38.9	

(Table 5) A Comparison between Type1, Type2 Diabetes in the Percentage of Carriage of Different Species of Candida Isolates

Candida Species	Type1 no.=142	%	Type1 no.=198	%	P value
C.albicans	35	24.7	55	27.8	p>0.05
C.tropicali	11	7.8	9	4.6	
C.krusei	1	0.7	9	4.6	
C.glabrata	5	3.6	4	2.0	

(Table6) The Relationship between the Degrees of Glycemic Control and the Oral Carriage Rate of Candida Species

Glycemic Control	no.of patients	no. of isolates	%	P value
Good FPG<=115mg\dl	42	5	11.9	P<0.01
Acceptable FPG>=110mg\dl & <=150mg\dl	130	47	36.2	
Poor FPG>=150mg\dl	168	77	45.8	

in both types of diabetes, other studies do not cover this subject. Glycemic control affects the oral carriage rate of *Candida* where better control leads to less carriage rate ($P < 0.01$), similar results reported by Odd F C et al 1978 where 60 % of poorly controlled diabetics carried *Candida* orally⁽¹⁷⁾, but Fisher BM et al 1987⁽¹⁸⁾ report no significant relation between glycemic control and oral carriage rate of *Candida*, we think that high oral glucose level in poorly controlled diabetics predispose to this infection.

*From the Department of Medicine, Alkindy college of Medicine, University of Baghdad

** From the Department of Microbiology

Alkufa College of Medicine, University of Kufa

***** From the Department of Microbiology

Alkufa College of Medicine, University of Kufa

References

1. Emmonds, C W: Binford, C H, and UTZ, J.P. (1974), medical mycology .Lea and Feiberger, Philadelphia.
2. Bodey GP: , candidiasis in cancer patients,(1984) *American J Med.*,77:13-19, cited by Chung, K.J.K. and *Bennett J E*(1992)
3. Bougnoux ME:resolutive candida utilis fungemia in a non neutropenic patient. *J of clinical microbiology.*(1993) 31:1644-1645.
4. Bennett JH:(1844) on the parasitic vegetable structure found growing in living animals. In *Trans R Soc Edin* . 15:277-294. Cited by Chung,K.J.K. and *Bennett* (1992).
5. Chandler, PT: Chandler, S.D.Pathogenic carrier rate in diabetes mellitus. *The American J of medical Science* (1977) 273:3.
6. Shilton, BH: Walton DJ : Site of glycation of human and horse liver alcohol dehydrogenase in vitro.(1991) *J Bio Chim* 266.
7. Cech P Papathanassion A: Hereditary myeloperoxidase ficincy.(1979),*Blood*,53(3):403-411.cited by the midline.
8. Elliot, JR: Mark J.A.,: Infection and diabetes: the case for glucose control.(1982) *American J of Medicine* .72.
9. Tapper-Jones, L.M.:*Candida* infection and population of *candida albicans* in mouths of diabetics.(1981)*Jof clinical pathology*.,34:706-11
10. Chung,K .J .K and Bennett,J.E. (1992) : *Candidiasis.*, *Med Mycology*.
11. Ruchel, R. : *Candida* acid proteinases.(1991) *J Med Vet Myc...*,29..
12. Al-Hussaini A.M.Microbiological study of dental caries,gingivitis and periodontitis.M Sc. thesis,(2002)Kufa University ,College of Science.
13. Amin K.Microbiological study of chronic paronychia in house wives; isolation, identification and treatment.M Sc. Thesis (2002),Kufa University, College of Medicine.
14. Barlow A.J.E. and Chattaway F.W.: Observations on the carriage of *C. albicans* in man.(1969) *Br J of Dermatology*, 81:103-106.
15. Smits BJ: Prior AP:and Arblaster P.G., Incidence of *Candida* in hospital inpatients and the effect of antibiotic therapy.(1966) *BMJ* 1:208-210.
16. Arenfold TM:and Walker DM : The prevalence and intraoral distribution of *C. albicans* in man.(1980) *Arch oral Biol*. 15:1-10.
17. Odds FC Evans E.G.V. Taylor M.A.R. and Wales J.K.: Prevalence of pathogenic yeast and humoral antibodies to *candida* in diabetic patients.(1978) *J Clin Path*31: 840.
18. Fisher B.M. : Carriage of *Candida* species in the oral cavity of diabetic patients ,relationship to lucose control.(1987) *J Oral Path*. 16:282-284.

Al-Kindy Col Med J 2006; Vol.3 (1): P 8-11