

# Journal of Global Pharma Technology

Available Online at: www.jgpt.co.in

**RESEARCH ARTICLE** 

# Distribution of Both Fungi and Bacteria Detected from the Nasal and Oral Cavity (Statistical Survey in Kirkuk)

# Sanaa H. Mohammad<sup>1\*</sup>, Muhsin H. Edham<sup>1</sup>, Hawazin A. Abid<sup>2</sup>

<sup>1.</sup> College of Sciences, Department of Biology University of Kirkuk, Iraq.

<sup>2.</sup> College of Science, Department of Biology, Tikrit University, Iraq.

\*Corresponding Author: Email: Sanaabio411@gmail.com

#### Abstract

The study was carried out at the period from 1<sup>st</sup> march 2018 to 31<sup>st</sup> june 2018 in order to determine the relationship of the allergic fungi with the worker in services and restaurant also university student and study the occurrence of fungi and bacteria detected from nasal and oral cavity among University Students , Restaurant workers and services workers with mean age  $(47.40\pm22.14)$  subjects from different areas of Kirkuk City(107) isolates of bacteria and fungi isolated from the nasal cavity in the same time(45) isolates of bacteria and fungi isolated from the oral cavity the isolates represented by : *A. niger, A. flavus, A. fumigatus, A. nidulans, A. terreus, A. japonicas, A. japonicas, Alternaria spp, C. albicans, C. parapsilosis, Penicillium* spp, *Rhizopus* spp, *S.pyogen, S.aureus* .The statistical analysis, existence of significant differences at the level of probability (P< 0.05) between the type of isolates the high frequency isolation represented by, *A.niger, and S.aureus* in the oral and nasal cavity which appeared with high occurrence . The subjects in this study were divided into five age groups. On the other hand the statistical analysis of the existence significant differences (P< 0.05) between the fungal and bacterial isolation infection and age for all age groups, it had been found that the age group 26-35 years was the most influential of these infections, compared with the other age groups.

Keywords: Fungi, Bacteria, Nasal and oral cavity.

#### Introduction

Normal nasal and oral flora organisms which includes bacteria, and fungi, the most important organism belong to the genus *Aspergillus* and Candida [1]. Human infections caused by *Candida albicans* and *Aspergillus fumigatus* and other related species range from the more common nasal and oral thrush to fatal, systemic super infections in patients who are afflicted with other diseases [2].

Fungal infection by species may be consider from up to one-third of the nasal and oral cavity of normal individuals and are considered inhabitants of the normal flora of oral and gastrointestinal tract [3]. C. *albicans* and A. *fumigatus* the main fungi associated with human nasal and oral mycoses and is the most fungi have virulence factors during infections especially in the students in the school and university [4]. The abilities of *A. fumigatus* and *C. albicans* to transform from blastopores to the hyphal phase specially in the restaurant workers and building workers found in most of them form germ tubes regarding *C. albicans*, while *A.fumigatus* mark the onset of hyphal growth of *Aspergillus* [5].

The most main factors in the fungal infections in the crowded university students and workers advent of the human immunodeficiency the immunosuppressive therapy and increasing incidence of diabetes, may be consider the global scenarios that have resulted in the fungal infections in the workers and students[6].

This spectrum of fungal infections has paved way for the increased incidence of nasal infections, and oral candidiasis (OC) [7]. The present study aimed to isolation and identification of fungi from nasal and oral cavity in University Students and Restaurant workers and services workers.

### Methods

Samples collection: Swap samples were collected from 290 patients ages ranged between (<10 >56 years old) in order to detect infection with asthma and aspergillosis, during the period of  $1^{st}$  march 2018 to  $31^{st}$  june 2018 from nasal and oral cavity in University Students and Restaurant workers and Services workers and divided it according to smoker and nonsmoker.

The samples were examined directly under the microscope using 10% KOH solution and culturing on the SDA agar and examined for after 7 days [8, 9, 10, 11]. The isolates were identified according to [3, 4, 12]. Results subjected were to statistical analysis. The significant differences are determined in rate of probability 5% as the statistical analysis includes one wav variance (ANOVA). Also analysis of significant differences are examined between means using test of less significant difference LSD [13].

### **Results and Discussion**

The current study investigates the both fungi and bacteria detected from nasal and oral cavity among University Students . Restaurant workers and services workers with mean age  $(47.40\pm22.14)$  subjects from different areas of kirkuk City Table (1) have shown Mean difference of Age and Distribution of both fungi and bacteria and oral cavity.. the detected from nasal of statistical analysis results and ล significant difference at the level of probability (P < 0.005).

#### Statistical Analysis

Type of	Total	%	% Total % Total Positive						t-test
isolate	NO.		+		Nasal	%	oral	%	
University Students	70	24.1	38	25	25	23.3	13	28.8	2.89
Restaurant workers	100	34.4	50	32.8	33	30.8	17	37.7	22.14
services workers	120	41.3	64	42.1	49	45.7	15	33.3	11.47
total	290	100	152	100	107	100	45	100	20.13

Table 1: Distribution of both fungi and bacteria detected from nasal and oral cavity

P.value significant < 0.005

Distribution of both fungi and bacteria detected from the nasal and oral cavity according to the type of isolates there's different isolates of bacteria and fungi isolated during the study 107 isolates of bacteria and fungi isolated from the nasal cavity in the same time 45 isolates of bacteria and fungi isolated from the oral cavity the isolates represented by : A. niger, A. flavus, A. fumigatus, A. nidulans, A.terreus, A. japonicas, A. japonicas, Alternaria spp, C. albicans, C. parapsilosis, Penicillium spp, Rhizopus spp, S. pyogen, S.aureus.as show in table 2 and Figure 1.

Table 2: Distribution of	both fungi and	l bacteria	detected	from the	e nasal	and ora	l cavity	according to	the type of
<u>isolates</u>									

Organism isolate	nasal	%	mouth	%	Total	%
A. niger	17	15.8	8	17.7	25	16.4
A. flavus	12	11.2	5	11.1	17	11.1
A. fumigatus	7	6.5	4	8.8	11	7.2
A. nidulans	6	5.6	4	8.8	10	6.57
A.terreus	1	0.9	0	0	1	0.65
A. japonicas	4	3.7	0	0	4	2.6
Alternaria spp	5	4.6	4	8.8	9	5.9
C. albicans	14	13.0	8	17.7	22	14.4
C.parapsilosis	7	6.5	6	13.3	13	8.5
Penicillium spp.	2	1.8	0	0	2	1.3
Rhizopus spp	7	6.5	1	2.2	8	5.2
S.pyogen	0	0	5	11.1	5	3.2
S.aureus	25	23.3	0	0	25	16.4
Total	107	100	45	100	152	100

P. value significant <0.05>

The statistical analysis, existence of significant differences at the level of probability (p < 0.05) between the type of

isolates and this percentage agreed with results of other researchers.

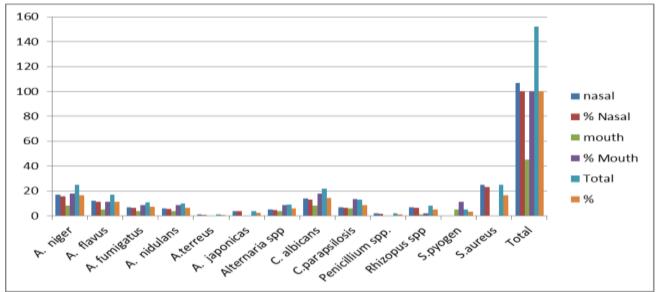


Figure 1: Distribution of both fungi and bacteria detected from the nasal and oral cavity according to the type of isolates

When divided the isolates distribution according to the site of isolation there's only fungi isolated from University Students as summarized in Table 3, The statistical analysis, existence of significant differences at the level of probability (p < 0.05) between

the type of isolates which represented by A. flavus, A. niger, A. nidulans, A. terreus, A. japonicus, Alternaria spp., Candida albicans and S.aureus.

Table 3: Distribution of both fungi and bacteria detected from nasal and oral cavity in University Students

Type of isolate		Total	Positive	
	Nasal	%	mouth	%
A. flavus	5	20	2	15.3
A.niger	4	16	1	7.6
A. nidulans	6	24	4	30.7
A.terreus	1	4	0	0
A.japonicus	1	4	0	0
Alternaria spp.	1	4	2	15.3
Candida albicans	2	8	4	30.7
S.aureus	5	20	0	0
Total	25	100	13	100

p. value significant <0.05

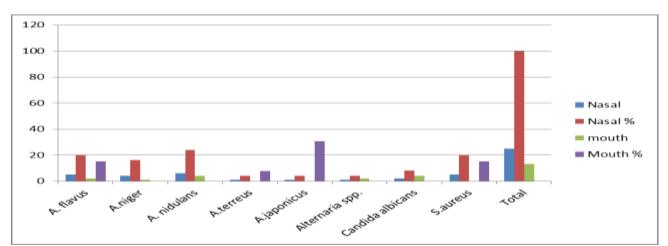


Figure 2: Distribution of both fungi and bacteria detected from nasal and oral cavity in University Students

#### Distribution of Both Fungi and Bacteria Detected from Nasal and Oral Cavity in Restaurant Workers

When divided the isolates distribution according to the site of isolation there's different type of fungi and bacteria isolated from Restaurant workers as summarized in table 4, and figure 3 The statistical analysis, existence of significant differences at the level of probability (p < 0.05) between the type of isolates which represented by A. flavus, , A. fumigatus A. niger A. Japonicas Candida. albicans C. parapsilosis Rhizopus spp. Penicillium spp. Alternaria spp. S. pyogen and S. aureus.

Table 4: Distribution of both fungi and bacteria detected from nasal and oral cavity in Restaurant workers

Type of isolate	Total Positive						
Γ	Nasal	%	mouth	%			
A. Flavus	2	6.06		5.8			
			1				
A.fumigatus	1	3.03		23.5			
			4				
A.niger	5	15.1	3	17.6			
A. japonicas	3	9.0	0	0			
Candida.albicans	4	12.1	0	0			
C.parapsilosis	7	21.2	6	35.2			
Rhizopus spp.	2	6.06	0	0			
Penicillium spp.	2	6.06	0	0			
Alternaria spp.	2	6.06	0	0			
S.pyogen	0	0	3	17.6			
S.aureus	5	15.1	0	0			
Total	33	100	17	100			

P. value significant <0.05

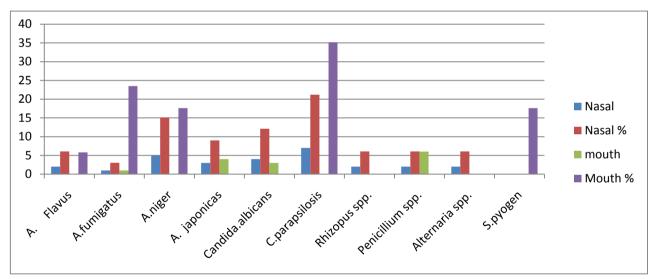


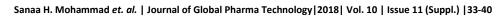
Figure 3: Distribution of both fungi and bacteria detected from nasal and oral cavity in Restaurant workers

The fungi and bacteria isolated from nasal cavity its more than oral cavity in the services worker as summarized in table 5, and Figure 4 The statistical analysis, existence of significant differences at the level of probability (p < 0.05) between the type of isolates the high frequency isolation represented by, *A. niger, and S. aureus* in the oral and nasal cavity which appeared with high occurrence.

Table 5: Distribution of both fungi and bacteria detected from nasal and oral cavity in services workers

Type of isolate	Total Positive							
	Nasal	%	mouth	%				
A. fumigatus	6	12.2	0	0				
A. niger	8	16.3	4	26.6				
A. flavus	5	10.2	2	13.3				

Alternaria spp	2	4.0	2	13.3
Rhizopus sp	5	10.2	1	6.6
Candida. albicans	8	16.3	4	26.6
S.pyogen	0	0	2	13.3
S.aureus	15	30.6	0	0
Total	49	100	15	100



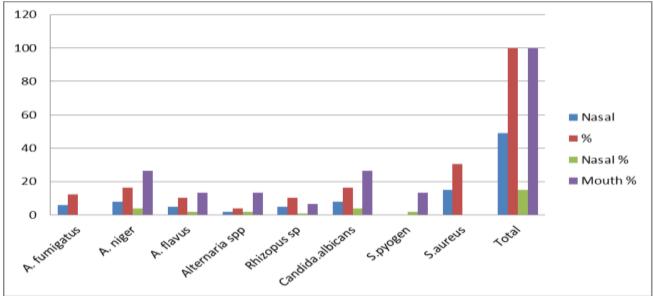


Figure 4: Distribution of both fungi and bacteria detected from nasal and oral cavity in services workers

The subjects in this study were divided into five age groups. On the other hand the statistical analysis of the existence significant differences (p< 0.05) between the fungal and bacterial isolation infection and age for all age groups, it had been found that the age group 26-35 years was the most influential of these infections, compared with the other age groups. Table (6).

	Positive Cases						
%	М	%	N	(years)			
11.1	5	9.3	10	<10			
20	9	18.69	20	25-16			
33.3	15	56.0	60	35-26			
24.4	11	9.3	10	45-36			
11.1	5	6.5	7	56-45			
100	45	100	107	Total			

Table 6: Distribution of both fungi and bacteria detected from nasal and oral cavity according to age

 $\rm X^{2}{=}$  10.182, indexed = 7.815 and a significant difference between age group (p< 0.05)

In total 290 individual included in the study university student, restaurant worker , and services worker , Table 7 and figure 5 show 62% from the total number smoker and 37.9 % were nonsmoker , The statistical analysis,

existence of significant differences at the level of probability (p < 0.05) between the Smoker and nonsmoker individual included in the current study.

 Table 7:
 Distribution of both fungi and bacteria detected from nasal and oral cavity according to smokers or nonsmokers

	Total NO	%	Positive NO	%	Negative NO.	%
smoker		62.0		78.9	60	43.4
	180		120			
Un smoker		37.9				56.5
	110		32	21.0	78	
Total	290	100		100	138	100
			152			

P. value significant <0.05

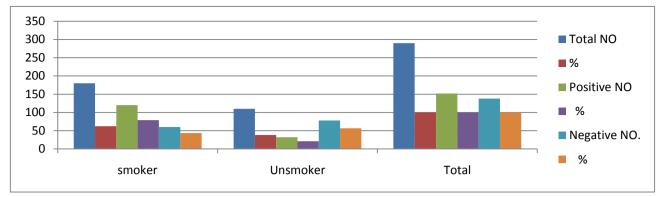


Figure 5: Distribution of both fungi and bacteria detected from nasal and oral cavity according to smokers or nonsmokers

Few reports have been published on the Distribution of both fungi and bacteria detected from the nasal and oral cavity. We assessed the the correlation between fungal and bacterial infection and work or living specially the correlation between fungal and bacterial infection and work or living especially in the restaurant and building worker [16, 17]. We analyzed the prevalence of the given fungal and bacterial species (the percentage) and the proportions in which the were colonised palate and by each microorganism. Aspergillus and S.aureus were the most frequently detected in subjects with university student [18].

significantly higher frequency of А methicillin-sensitive Staphylococcus aureus (S. aureus MSSA) was observed in the university student was predominant on the palate. The development of the fungal infection subjects was characterized by a significant increase in the prevalence of pathogenic bacteria [19]. This paper is dedicated to study the relationship between fungal and bacterial infection and the mode of work the oral cavity, which remains sterile throughout prenatal development, becomes a diverse ecosystem colonized by numerous microorganisms during the work in restaurant or building [20].

The resident microbiota in the period work depends mainly on external factors, around university student and worker in the restaurant. These conditions are favorable for the development of a diverse ecosystem based on the interactions between bacteria, fungi and the host environment [21]. The early oral detection occurring within several hours following working in crowded place or smoking is composed of mixed infection bacteria and fungi which are commensals permanently colonising the oral and nasal cavity [22]. Along with other bacteria, they participate in the formation of a "colonization cascade" that determines future indigenous micro biota [23]. Neonates with complete cleft lip and palate (CLP) are characterized by the existence of communication between oral and nasal cavities extending from the upper lip and nasal vestibule to the end of the soft palate. This condition adversely affects natural sucking or even impairs the ability to restaurant food [24].

Moreover oral infection with or facial cleft require specialized care to maintain proper hygiene of the incisive bone, nasal passages, and the oral cavity with special attention paid to preparation for future surgical procedures [13]. Distribution of Bacteria and fungi according to their age The distribution of fungi showed higher infection of A.niger in age group ranged >50 vears while the lower infections were at ages lower than 5 years old. In general, the higher infection was starting from 25 years old of age and above (Table 6). Infection in worker with bacteria and fungi was found to be more prevalent in >30 years old [25]. These results are in concomitant with the present study.

The prevalence of the infection was proportional with certain cases i.e. impaired associated with immune state the malignancy, management of organ transplantation, autoimmune and inflammatory conditions; critically in the school and university student [26].

Microscopic and culturing examination show that theirs higher bacterial and fungal contamination in outdoor 87 (63.5%); 18(2.6%); ,as well as recorded the highest percentage in *A.niger28* (23.6%)followed by *Penicillium spp* 19(6.5%); *A. flavus*14 (3.8%); *Alternaria spp* 5(10.1%);*Microsporium spp*  and Fusarium spp10(5.1%): A. fumigatus 4(2%) respectively as show the figure (3) shows the image which the fungi isolates during this study isolated from nasal and mouth cavity during the present study .several study according to fungal and bacterial infections showed the frame material in building and in the crowded sit like university campus had more importance on fungal concentrations than moisture damage, the contaminations in the present study, detected the contamination the study with bacteria and fungi may be related to different cause highly moisture, overcrowded and in a poor condition, which all encourage fungi and to grow quickly.

In addition, existence of some nearby plants i.e. Kirkuk Cement Factory, North Gas Company and North Oil Company have added bonus factors to over contaminate the surrounding air.

Consequently, the inhaled air would ease infection of the respiratory system and may

#### References

- 1. Aira MJ, Rojas TI, Jato V(2002) Fungi associated with three houses in Havana (Cuba).Grana ,41:114-118.
- 2. Al- Ameed, AIM (2008) Isolation of Aspergillusfumigatus from Human Being and the study of immune response on rabbits immunized by their antigens .MS. Thesis. College of Veterinary Medicine-Baghdad University, Iraq.
- 3. AL-Malikey HS (2009) Effect of alcoholic extract of callistemon viminalis (Sol. ex Gaertn) G.D on fungi cause aspergillosis in albino mice. College of science - Baghdad University, Iraq.
- Anderson MJ, Gull K, Denning DW (1996) Molecular typing by random amplification of polymorphic DNA and M13 southern hybridization of related paired isolates of *Aspergillus fumigatus*. J. Clin. Microbiol., 34 (1): 87-93.
- 5. Arruda LK, Mann BJ, Chapman MD (1992a) Selective expression of amajor allergen and cytotoxin, AspfI, in *Aspergillus fumigatus*. Implications for the immunopathogenesis of *Aspergillus*related diseases. J. Immunol., 149: 3354-3359.
- Awosika SA, Olajubu FA, Amusa NA (2012) Microbiological assessment of indoor air of a teaching hospital in

affect the body immunity [3, 4]. In terms of the biosafety levels suggested by [6], *A.flavus* and *A.fumigatus* would be considered the species of most harmful to human health; however, *A.niger*, *A.terreus*, *A.tamarii and C.cladosporioides* have also been implicated in a range of pathologies [13] Appropriate measures should therefore be implemented to reduce fungal density in the environments tested with a view to improving air quality and avoiding potential adverse effects [27].

## Conclusion

In the present study clear that bacteria and fungi contaminate the nasal and oral cavity especially in the worker indifferent age and the university student.

Nigeria. Asian Pacific Journal of Tropical Biomedicine, 465-468.

- Borrego S, Guianmet P, Gomez de, Saravia S, Batistini P, (2010) The quality of air at archives and the biodeterioration of photographs. International Biodeterioration and Biodegradation, 64:139-145.
- 8. Cho SJ, Ramachandran G, Banerjee S, Ryan AD, Adgate JL (2008) Seasonal variability of culture able fungal genera in the house dust of inner-city residences, J. Occupy Environ. Hyg., 5: 780-789.
- 9. Deeb OB (2013)Study on the pathogenicity, cytotoxicity and virulence factors of some clinical of Aspergillus fumigatus isolates and the use of RAPD markers to distinguish them. Ph. D. Thesis College of Science. Tikrit. University, Iraq.
- Eichner RD, Al-Salami M, Wood PR, Mullbacher A (1986) The effect of gliotoxin upon macrophage function. Int. J. Immunopharmacol, 8:789-797.
- 11. Ellis M (1999) Therapy of Aspergillus fumigatus-related diseases. Contrib. Microbiol., 2:105-129.

- Forbes B, Sahm D, Weissfeld A (1998) Bailey and Scott's diagnostic microbiology, 10<sup>th</sup> Ed., Mosby, Inc. St., Louis.
- Horner WE, Helbling A, Salvaggio JE, Lehrer SB (1995) Fungal allergens. Clin. Microbiol. Rev., 8: 161-79.
- 14. Hsu NY, Lee CC, Wang JY, Li YC, Chang HW, Chen CY, Bornehag CG, Wu PC, Sundell J, Su HJ (2012) Predicted risk of childhood allergy ,asthma, and reported symptoms using measured phthalate exposure in dust and urine .Indoor Air,22:186-199.
- Jawetz E, Melnick J, Adelberg EA (1991) Medical Microbiology. 20<sup>th</sup> Ed., Middle East edition, 532-546.
- 16. Madbouly AK, IMI Mohamed, FS Ahmed, AA Mosaad (2012) CO- Occurence of mycoflora ,aflatoxins and fumonisins in maize and rice seed from markets of different districts in Cairo. Egypt, Food Additives and Contaminants, B 5 (2): 112-120.
- Suleiman RK, Rosentrater C Bern (2013) Effects of Deterioration Parameters on Storage of Maize. A Review. Journal of Natural Sciences Research. 3(9): 147-165.
- 18. Kamei K, Watanabe A (2005) Aspergillus mycotoxins and their effect the host. Med .Mycol. 43 (1): S95-99.
- 19. Kirk PM, Cannon PF, David JC, Stalpers JA (2008) Ainsworth an Biby's dictionary of the fungi. 9th edition (CABI).Bio. Science., Oxon, UK, 452.
- 20. Komase R, Nakamura Y, Watari T (2007) Phylogenic analysis of 8 lower respiratory tract infection species using chitin

synthase 1 gene sequences. Mycoses, 40:411-414

- 21. Kowalski W, Bahnfleth WP (1998) Airborne respiratory diseases and technologies for control of microbes. Heating Piping Air Conditioning,70 (6): 34-48
- Larone DH (1993) Medically important fungi - a guide to identification. 2<sup>nd</sup> edition, American Society for Microbiology, Washington DC, USA.
- 23. Larone DH (1995) Medically important fungi: Aguide Fitzpatrick's Dermatology in General Medicine. McGraw-Hill, New York., 133-144.
- 24. Latgé JP (2001) The pathobiology of Aspergillus fumigatus. Trends Microbiol., 9: 382-389.
- 25. Latgé JP (1999) Aspergillus fumigates and aspergillosis .Clin .Microbiol. Rev., 12:310-350.
- 26. Latgé JP (2001) The pathobiology of Aspergillus fumigatus. Trends. Microbiol., 9:382-389.
- 27. Magnuson J, Lasure L (2004) Organic acid production by filamentous fungi. In: Tkacz J, Lange L (eds). Advances in Fungal Biotechnology for Industry, Agriculture, and Medicine. New York: Kluwer Academic and Plenum Publishers, 307-340.
- 28. Maiz L, Cuevas M, Lamas A, Sousa A, Quirce S, Suarez L (2008) Aspergillus fumigatus and Candida albicans in cystic fibrosis :Clinical significance and specific immune response involving serum immunoglobulins G,A, and M. Arch. Bronconeumol., 44(3):146-51.