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Meta-Analysis on Isolation Frequencies of Selective Pathogenic Bacteria in Complete Denture Patients

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Abstract

Introduction: A number of micro-organisms have been isolated from the oral cavity from dentulous patients. The chances of microorganisms to colonize complete denture surfaces as well as edentulous oral cavity are on the rise. Bacteria namely Streptococcus spp., Staphyloccus spp., Enterobacter spp., Klebsiella spp. Pseudomonas spp. and E.coli have been implicated in various cardiovascular, respiratory and gastrointestinal infections. All these bacterial spp. have been found to inhabit complete denture surfaces. Hence their isolation frequencies from complete denture have to be studied and compared by means of metaanalysis of similar studies.

Methodology: *Pubmed, Google Scholar, Open J Gate, Web of Science data bases were searched for relevant articles.*

Result: The isolation frequencies of the selective bacteria from the highest to the lowest is as follows: Streptococcus spp.(53.3%), Staphylococcus spp.(21.3), Enterobacter spp.(16.8%), Klebisella spp.(11%), Pseudomonas spp(6.1%). and E.coli (5.5%).

Conclusion: Since the isolation frequencies of these bacteria are known, the appropriate disinfection methods of complete dentures targeting the various organisms have to be implemented to prevent systemic disease likely to be caused by their aspiration.

Keywords: pathogenic bacteria, complete dentures, isolation frequencies, meta-analysis.

Introduction

A number of micro-organisms have been isolated from the oral cavity from dentulous patients .The chances of microorganisms to colonize complete denture surfaces as well as edentulous oral cavity are on the rise. This may be due to the fact that the prostheses act as niches for the growth of the microorganisms mainly due to the rough topographic surface of the complete dentures. Several studies have been done on yeast i.e. candidal *spp*. harboring on complete denture surfaces.

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Comparatively very few studies have been done on bacteria colonizing complete dentures. Bacteria which are of considerable interest are those which are capable of being pathogenic to the systemic health of the patient. For example, Streptococcal are known causative organisms spp. of cardiovascular disease especially endocarditis¹. Other examples include Staphylococcal spp. and Klebsiella *spp*. which are potential respiratory pathogens^{1,3,4}. Pseudomonas spp. and Staphylococcus SDD. are known to cause suppurative infections. Enterobacter spp. and E. coli have a positive role in gastro-intestinal infections^{2, 5, 6}

All these bacterial spp. have been found to inhabit complete denture surfaces in both disease and healthy persons. An ecological imbalance of the micro-organisms decides whether the patient might either acquire disease or resistant to disease caused by micro-organisms ^{1,2}. These organisms could lead to endocarditis, aspiration pneumonia, pneumonitis as well as gastro-intestinal infections and hence the isolation frequencies of these pathogenic organisms have to be compared and analysed by methodological method namely metaanalysis.

Methodology

The databases searched for the present study are J Gate, Web of science, Pubmed and Google scholar. The time frame for the article search is 26 years starting January 1991 to August 2016. The key words used in searching the data base include "Denture associated bacteria", "Bacteria in complete dentures" and "Complete denture associated bacteria".

For the key word "Denture associated bacteria", the total articles found in Pubmed, J gate and web of science were 54, 36 and 40 respectively.

For the key word "Bacteria in complete dentures", search results yielded via Pubmed , J gate and web of science were 177, 31and 6 respectively.

For the keyword "Complete denture associated bacteria", search results yielded via Pubmed , J

gate and web of science were19, 8 and 6 respectively.

By Google scholar refined search for the key word "pathogenic denture plaque" 101 articles were obtained which was further refined to keywords "biofilms in edentulous oral cavity" which yielded 96 articles.

The articles were hand searched for getting the isolation frequencies of bacteria which are as follows:

- 1. Enterobacter *spp*.
- 2. Klebsiella *spp*.
- 3. Streptococcus spp.
- 4. Staphylococcus *spp*.
- 5. Pseudomonas spp.
- 6. E.coli

The analysis broadly included the isolation frequencies of bacterial species including the various bacterial genus /serotypes if so mentioned in the study.

Initially abstracts of 177 articles were read through to sort out the relevance of the articles to the topic. From the initial screening (*Level-1*) only 90 articles were related to the topic. Those articles which are based on isolation of yeast i.e. candidal *spp*. were not included in the analysis. Review articles were not included as much of data could not be collected from them. However, references from review articles were utilized to search for articles of interest.

The next level of screening (*Level-2*) included distinction of articles based on prostheses type. Only articles in which complete dentures were the sources of the samples were included in the analysis. Articles wherein samples were obtained from dentate individuals, RPDs, FPDs and Implants were not included. 55 articles were sorted out with this screening.

Level-3 screening included distinguishing articles based on whether the study was done in vitro or in vivo. This screening yielded 32 articles.

All the 32 articles were thoroughly examined for obtaining details of isolation frequencies (*Level-4*) screening. Among the 32 articles only 14 articles yielded isolation frequencies. Other articles

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mentioned colony forming units' values, probability values as well as isolation frequencies of other bacteria not relevant to the topic. The screening method is illustrated in the following Flowchart:1.

Results and Discussion

The spreadsheet showing the isolation frequencies of the selective bacteria according to various studies is tabulated in Table 1.This data was subject to Meta analysis by comprehensive metaanalysis software version 2. The analyzed data and the respective forest plot for individual bacteria are illustrated in Tables: 2, 3,4,5,6, 7 and 8.

From the Tables: 2, 3 and 4 the isolation frequencies of Enterobacter *spp*. and Klebisella *spp*. were 16.8 and 11 percent respectively. As seen from the Table:3 ,the study done by Goldberg et al^8 ,the isolation frequency of Enterobacter *spp*. was 48 percent which was

statistically insignificant (p>0.05) .But careful analysis of the journal article showed that the combined isolation frequencies of Enterobacter spp. and Klebsiella *spp*. was found to 48 percent .Hence a higher isolation frequency was found.

From Table 2 and 5 the isolation frequency of Staphylococcus *spp*. was found to be 21.3 percent which is statistically significant (p<0.001).But, the isolation frequency of Streptococcus *spp*. was found to be highest which was 53.3 percent that was statistically significant (p<0.001)(Table 2 and 6).On the contrary Jaffari et al(2013)²⁰ have indicated that the isolation frequencies of Streptococcus and Staphylococcus *spp*. to be 100 percent .

With regard Pseudomonas spp. and E.coli, the isolation frequencies were 6.1 and 5.5 percent respectively which has statistical high significance (p<0.001) (Table 2, 7 and 8).

TABLE:1 Representing The Spreadsheet Which Tabulates The Study Details, Selective Bacteria And Their Isolation Frequencies

	(N	YE	IN	SOUR	ENTEROBAC	KLEBSIELL	STAPHYLOC	STREPTOCOC	PSEUDOMON	
NAME OF STUDY	Ĵ	AR	VIVO	CE	TER <i>spp</i> .	A spp.	OCUS spp.	OUS spp.	AS spp.	E.COLI
		199	IN							
GOLDBERG ET AL	62	7	VIVO	C.D	48					
		200	IN							
DANILUK ET AL	57	6	VIVO	C.D	17.5	8.80%	21.1	93.1	8.8	3.5
		199	IN							
RITSUKO ET AL	5	9	VIVO	C.D				31		
		200	IN	<i>a</i> b	<u>_</u>					
MARCELLE ET AL	41	9	VIVO	C.D	9	1.5		4.5	7.5	
	53	201	IN	CD	17.0	10.4	22.4		<i>.</i>	1.5
DOROTA ET AL	10	4 199	VIVO IN	C.D	17.9	10.4	22.4		6	1.5
SEDGLEY ET AL	0	199 5	VIVO	C.D	16	14			7	6
SEDULET ET AL	0	201	IN	C.D	10	14			/	0
DANIELA ET AL	90	201	VIVO	C.D			34.4	53.3		
KAKLAMANOS ET	14	200	IN	C.D			54.4	55.5		
AL	8	5	VIVO	C.D	7.4		21.6			
	-	200	IN							
SUMI ET AL	50	2	VIVO	C.D	18	16	10	0	2	8
		199	IN							
HARDING ET AL	20	1	VIVO	C.D			15	31		
	10	201	IN							
LEWIS ET AL	0	5	VIVO	C.D			30			
		201	IN							
NISHI ET AL	96	2	VIVO	C.D	3.5	0.7	13.4	93.7	2.8	
OCAMPO GRACIA	16	201	IN		_					
ET AL	0	5	VIVO	C.D	2		5	5		
	<i></i>	199	IN	GD						
KANONEN ET AL	51	1	VIVO	C.D				84		

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Table: 2 Representing The Comprehensive Isoaltion Frequencies Of Selective Bacteria After Meta-Analysis

BACTERIA	ISOLATION FREQUENCIES
Enterobacter spp.	16.8 (Table: 2)
Klebsiella spp.	11 (Table: 3)
Staphylococcus spp.	21.3(Table: 4)
Streptococcus spp.	53.3 (Table: 5)
Pseudomonas spp.	6.1 (Table: 6)
E. coli	5.5 (Table: 7)

Table: 3 Representing the Forest Plot of Enterobacter *spp*.

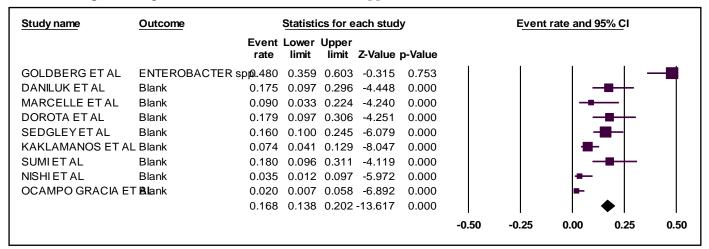


Table: 4 Representing The Forest Plot Of Klebsiella spp.

Study name	Outcome		Statisti	cs for ea	ach stud	ý	Event rate and 95% CI					
		Event rate	Lower limit		Z-Value	p-Value						
DANILUK ET AL	KLEBSIELLA	spp. 0.090	0.038	0.197	-4.999	0.000				_		
MARCELLE ET A	LBlank	0.020	0.002	0.154	-3.489	0.000				-		
DOROTA ET AL	Blank	0.100	0.043	0.214	-4.799	0.000			-			
SEDGLEY ET AL	Blank	0.140	0.085	0.223	-6.299	0.000			- I - I			
SUMI ET AL	Blank	0.160	0.082	0.289	-4.299	0.000			-			
NISHI ET AL	Blank	0.070	0.033	0.142	-6.467	0.000				-		
		0.110	0.082	0.147	-12.392	0.000						
							-0.50	-0.25	0.00	0.25	0.50	

Table : 5 Representing The Forest Plot Of Staphylococcus spp.

Study name	Outcome		Statisti	cs for ea	ach study	-	Event rate and 95% Cl					
		Event rate	Lower limit	Upper limit	Z-Value	p-Value						
DANILUK ET AL	STAPHYLOCOCUS spp.	. 0.211	0.124	0.336	-4.063	0.000			· · ·			
DOROTA ET AL		0.224	0.131	0.355	-3.771	0.000						
DANIELA ET AL	Blank	0.344	0.253	0.448	-2.909	0.004					_	
KAKLAMANOS ET AL	Blank	0.216	0.157	0.290	-6.454	0.000						
SUMI ET AL	Blank	0.100	0.042	0.219	-4.661	0.000				_		
HARDING ET AL	Blank	0.150	0.049	0.376	-2.770	0.006				-		
LEWIS ET AL	Blank	0.300	0.218	0.397	-3.883	0.000				-+	-	
NISHI ET AL	Blank	0.134	0.079	0.218	-6.228	0.000				-		
OCAMPO GRACIA ET A	AlBlank	0.050	0.025	0.097	-8.117	0.000						
		0.213	0.184	0.246	-13.738	0.000				\bullet		
							-0.50	-0.25	0.00	0.25	0.50	

Table :6 Representing The Forest Plot Of Streptococcus spp.

Study name	Outcome		Statisti	cs for ea	ach study	_	Event rate and 95% Cl				
		Event rate	Lower limit	Upper limit	Z-Value	p-Value					
DANILUK ET AL	STREPTOCOCOUS spp.	0.931	0.829	0.974	4.979	0.000					
RITSUKO ET AL	Blank	0.310	0.063	0.749	-0.827	0.408			<u> </u>	-	-
MARCELLE ET AL	Blank	0.045	0.011	0.171	-4.055	0.000					
DANIELA ET AL	Blank	0.533	0.430	0.633	0.626	0.532				-	
HARDING ET AL	Blank	0.310	0.148	0.537	-1.655	0.098					
NISHI ET AL	Blank	0.937	0.867	0.971	6.426	0.000					
OCAMPO GRACIA ET A	AlBlank	0.050	0.025	0.097	-8.117	0.000			-		
KANONEN ET AL	Blank	0.840	0.713	0.917	4.341	0.000					
		0.533	0.466	0.599	0.975	0.330				•	
							-1.00	-0.50	0.00	0.50	1.00

Table :7 Representing The Forest Plot Of Pseudomonas spp.

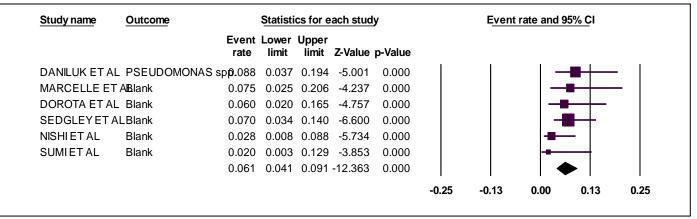
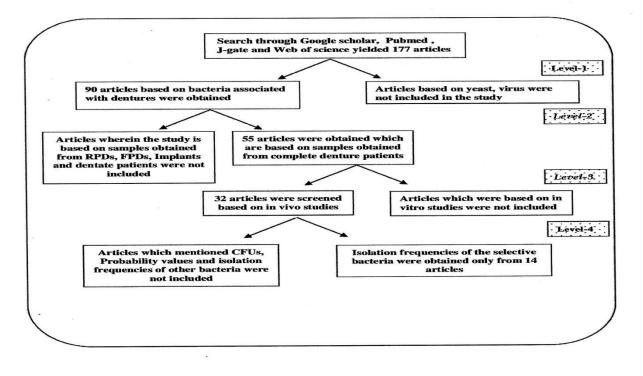


Table: 8 Representing The Forest Plot Of E.Coli

Study name	Outcome		Statisti	cs for ea	ich study	-		Event	rate and	95% CI	
		Event rate	Lower limit	Upper limit	Z-Value	p-Value					
DANILUK ET AL	E.coli	0.035	0.009	0.130	-4.602	0.000					
DOROTA ET AL	Blank	0.015	0.002	0.122	-3.703	0.000					
SEDGLEY ET AL	Blank	0.060	0.027	0.127	-6.535	0.000			-		
SUMI ET AL	Blank	0.080	0.030	0.195	-4.685	0.000			— —		-
		0.055	0.032	0.093	-9.852	0.000					
							-0.25	-0.13	0.00	0.13	0.25





Conclusion

From the meta-analysis it was found that several pathogenic bacteria were present in the complete dentures as evident from their isolation and microbiological examination. But the isolation frequencies of these selective pathogenic differed remarkably. The isolation frequencies of the selective bacteria from the highest to the lowest follows: Streptococcus are as spp., Staphylococcus spp., Enterobacter spp., Klebisella spp., Pseudomonas spp. and E.coli. Now that the isolation frequencies of these bacteria are known,the various disinfection procedures targeting these pathogenic bacteria have to be implemented to prevent possible aspiration/ engulfment that could be an etiological factor for systemic conditions such endocarditis. as respiratory infections as well as gastrointestinal infections.

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