

Preoperative Use of 0.2% Chlorhexidine Digluconate Mouthwash does not affect the Bacteremia following Closed Dental Extraction

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Abstract: Post extraction bacteremia is transitory condition observed in healthy individuals. Aim of the study at hand is to investigate the effect of preoperative rinsing of the mouth with 0.2% chlorhexidine digluconate on post extraction bacteremia following closed tooth extraction.

Materials and Methods: This research focused on evidence of bacteremia among 58 individuals, divided equally in two study groups – without preoperative rinsing (first group) and with preoperative rinsing (second group) with Parodontax Extra (GlaxoSmithKline, Great Britain), followed by open extraction of a single tooth. Aerobic and anaerobic hemocultures (Bact/ALERT, BioMerieux, Inc., Durham, N.C.) were used to establish the bacteremia, while blood samples were obtained preoperatively, 30 seconds after and 15 minutes after the extraction was completed.

Results: Preoperative bacteremia was established in 7 (12.07%) patients in total – 4 patients from the first and 3 patients from the second group. At the 30th second after the extraction was done bacteremia was established in 6 (20.69%) patients from the first and in 8 (27.59%) patients from the second group. At the 15th minute mark bacteremia was found in 3 patients (10.34%) from each group. We failed to identify any statistically significant difference in occurrence of bacteriemia among the subjects in both groups 30 seconds (p=0.548) as well as 15 minutes after the extraction was completed (p>0.05).

Conclusions: *Rinsing the mouth with 0.2% chlorhexidine digluconate mouthwash does not appear to offer any statistically significant reduction of the occurrence of post extraction bacteremia following typical dental extraction.*

Keywords: bacteremia, dental extraction, 0.2% chlorhexidine

1. INTRODUCTION

The extraction of teeth is among the most commonly researched manipulation that can produce bacteremia. In healthy individuals the bacteremia is a transitory condition with no clinical presentation and complains. Under certain circumstances it is possible that the bacteremia produces some complications, most notably bacterial endocarditic.

It is well established, that hematogenically disseminated micro flora from the oral cavity is responsible for 10%-15% of bacterial endocarditis' occurrences. Mortality rate of

bacterial endocarditic nowadays is still high and varies between 5% and 11%. [1,2] Several antibiotic regimens are used for prevention of the complications related to post extraction bacteremia. Currently, a trend for reducing the duration of antibiotic prophylaxis can be observed - from 5 days and maximum 21 doses of antibiotic (as suggested in 1955 by the AHA) to a single dose 30-60 minutes prior to the manipulation with Amoxicillin being antibiotic of choice in various regimen. [3] Alternative methods for controlling the post extraction bacteremia are investigated in an effort to limit the use of antibiotics and therefore

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bacterial resistance. In 1977 AHA suggested disinfection of the gingival sulcus prior and in addition to antibiotic prophylaxis of bacterial endocarditic for patients at risk. [4] In 1992 The Society Antimicrobial British for Chemotherapy, BSAC, refined the type and concentration of oral disinfectant Chlorhexidine -1% gel for application on the gingival margin, or 0.2% mouthwash for rinsing the mouth for 5 minutes. [5] In 2006 BSAC recommended single rinsing with 0.2% Chlorhexidine gluconate prior to dental manipulations that can induce bacteremia in patients at risk of bacterial endocarditic. (6) In contrast, since 2007 AHA refrains from recommending ani antiseptics. [1]

Aim of the study is to investigate the effect of preoperative rinsing with 0.2% chlorhexidine digluconate on bacteremia following closed dental extraction.

2. MATERIALS AND METHODS

58 individuals were equally divided in two trial groups – first group did not rinse preoperatively, and the second group did. Inclusion criteria were: clinically healthy patient; single tooth extraction was necessary. Exclusion criteria were: lack of consent, multiple extractions, pharmaceutically controlled chronic conditions, use of antibiotics in the last 6 months, acute oral inflammation, tumors and malignancies, compromised immune system, diabetes mellitus, pregnancy, history of/upcoming radiotherapy to the head and neck region.

Immediately before surgery subjects in the second group rinsed their mouth two times with 10ml 0.2% chlorhexidine digluconate (Parodontax Extra, GlaxoSmithKline, Great Britain) for one minute each. The solution was given in two single-use chemically clean plastic cups. Patients did not rinse with water after that. Tooth extraction was conducted in the following order: 1) local anesthesia; 2) syndesmotomy; 3) luxation and/or rotation of the tooth; 4) traction; 5) revision and inspection of the wound; 6) manual compression; 7) hemostasis.

Several aerobic and anaerobic hemocultures, incubated in an automated system, were utilized for research of bacteriemia (Bact/ALERT, BioMerieux, Inc., Durham, N.C.). The site of venipuncture was disinfected with ethanol, followed by iodine solution. 5ml of venous blood for each hemoculture (aerobic and anaerobic) was collected from the cubital vein. Then another sterile needle was used to aseptically transfer the material from the syringe into the container which was timely brought to the microbiology laboratory. Three samples of paired hemocultures for aerobic and anaerobic were acquired accordingly: bacteria 1) preoperatively, prior to any manipulations in the mouth; 2) 30 seconds after the extraction was completed; 3) 15 minutes after the extraction was completed. The hemocultures were incubated in BactALERT 3D 60 (BioMerieux. Inc., Durham, N.C.) for 6 days. Positive hemocultures were transferred in solid and liquid nutrient mediums and prepared by Gram stain. Identification of the isolated strains was conducted according to the standard methodsor automatically - using Vitek 2 (BioMerieux, Inc., Durham, N.C.). Some positive hemocultures that showed no bacteria through Gram staining were automatically subcultuvated up to 6 days and were deemed false-positive if no bacterial growth was evident. Hemocultures that were not marked by the device were subjected to routine incubation and transferred to solid nutrient mediums. Evident growth marked them as falsenegative, whereas true-negative hemocultures showed no growth whatsoever.

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3. RESULTS

Out of all 58 patients, 29 were males and 29 were females. Average age in the first group was 40.76 with standard deviation of 3.63, and in the second group it was 48.17 years with standard deviation of 4.13. We failed to establish any statistically significant difference in age between both groups (p=0.183). Distribution of the extracted teeth in both groups follows in table1.

Table1: Distribution of the extracted teeth according to type

	Molars		Premolars		Canines		Incisors		All	
group	N⁰	%	N⁰	%	N⁰	%	N⁰	%	N⁰	%
1 st group (no rinsing)	24	41.38	1	1.72	2	3.45	2	3.45	29	50
2 nd group (rinsing)	16	27.59	7	12.07	1	1.72	5	8.62	29	50
All	40	68.97	8	13.79	3	5.17	7	12.07	58	100

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Preoperative bacteremia was confirmed in 7 (12.07%) patients – four from the 1st group and 3 from the 2^{nd} group. At the 30th second after completing the extraction bacteremia was evident in 6 patients (20.69%) from the first group and 8 (27.59%) from the 2^{nd} . Samples at the 15th minute after the extraction revealed 3 subjects (10.34%) with bacteremia from both study groups. We failed to recognized any statistically sound difference in occurrence of bacteremia between subjects in both groups at 30th second (p=0.548) and 15th minute (p>0.005) after the extraction. The most common finding preoperatively Coagulase was negative Staphylococcus- in 71.43% of the positive aerobic and in 28.57% of the positive anaerobic hemocultures. Its presence in both is explained with its facultative anaerobic nature, which allows it to benefit from both respiration and fermentative metabolism. At the 30th second mark after completing the extraction in the first group most common bacteria incubated from aerobic hemocultures was Streptococcus milleri (33.33%), and Streptococcus viridians (25%) from the anaerobic ones. In the second group the most common finding was Streptococcus viridians- 50% of the positive aerobic and 37.5% of the positive anaerobic hemocultures. The Coagulase negative staphylococcus was most commonly cultivated from the samples at the 15th minute mark - 33.33% of all positive hemocultures in the first group and 66.67% of the positive aerobic hemocultures and 33.33% of the positive anaerobic hemocultures in the second study group. Register of the cultivated bacteria is presented in table 2.

		First gro	up	Second group				
	n	Isolated from aerobic	Isolated from	n	Isolated from	Isolated from		
		hemoculture	anaerobic		aerobic	anaerobic		
			hemoculture		hemoculture	hemoculture		
Pre	5	Coagulase negative	Coagulase negative	3	Coagulase	Coagulase		
operat		Staphylococcus	Staphylococcus;		negative	negative		
ivelly			Streptococcus		Staphylococcus	Staphylococcus		
			mitis/oralis		Staphylococcus	Streptococcus		
					aureus	mitis/oralis		
At	6	Streptococcus viridans;	Streptococcus	8	Streptococcus	Coagulase negative		
30 th		Coagulase negative	viridans;		viridans ;	Staphylococcus;		
secon		Staphylococcus;	Streptococcus mitis;		Coagulase negative	Streptococcus		
d		Streptococcus milleri;	Streptococcus		Staphylococcus;	viridans;		
mark		Streptococcus	milleri		Streptococcus	Streptococcus		
		constelatus			milleri	milleri		
At 15 th	3	Coagulase negative	Coagulase negative	3	Coagulase	Coagulase		
minute		Staphylococcus;	Staphylococcus;		negative	negative		
mark		Streptococcus viridans	Bacillus species		Staphylococcus	Staphylococcus		

Table2: Isolated microorganisms after incubating the hemocultures

The average duration of the extraction in the first group was 16.24 min at standard deviation of 2.09, and in the second group it was 15.24 min at standard deviation of 2.27. No statistically evident difference between two groups was observed (p=0.757). We established that the post extraction bacteremia is unaffected by the duration of closed extraction at 30th second mark (p=0.289), at 15th minute mark (p=0.394), as well as the type of extracted tooth (p=0.241 at 30th second, p=0.869 at 15th second).

4. DISCUSSION

Conflicting evidence about the effect of chlorhexidine prior to dental extraction is preset in the literature. Some authors found that rinsing with 0.2% chlorhexidine prior to extraction reduces the bacteremia significantly. Barbosaet al. [7] compared the effect of preoperative

rinsing with 0.2% chlorhexidine digluconate in 50 subjects with 52 other patients who did not rinse. Blood samples were acquired 30 seconds and 15 minutes after concluding the extraction. They reported no statistical difference in the occurrence of bacteriemia between both groups at 30th second mark - 50% versus 52%, however definite difference was observed at the 15th minute mark - 4% versus 23%.

Ugwumba et al. [8] confirmed that preoperative rinsing with 0.2% chlorhexidine digluconate reduces the post extraction bacteremia after closed tooth extraction. Their study included 101 subjects, divided in two groups. Samples for hem cultures were collected at 1st and 15th minute after extraction. They found that the occurrence of bacteremia in the control group was 52.4% and in the patients who used the mouthwash it was only 27.1% (p=0.012). Most

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commonly cultivated bacteria were Staphylococcus aureus, Actinomycetes naesulendi, Prevotella, Streptococcu sspp. And Acinetobacter iwoffii. Results of Tomás et al. [9] demonstrate that preoperative application of chlorhexidine convincingly reduces occurrence of post extraction bacteremia – 96% versus 79% at 30th second and 64% versus 30% at 15th minute. Other authors confirmed our findings and did not recognize the effect of 0.2% chlorhexidine on the occurrence of bacteremia following closed dental extraction. Maharajet al. [10] reported no statistically significant difference in the occurrence of bacteremia between patients who rinsed preoperatively with 0.2% chlorhexidine digluconate and the control group who did not - 40% versus 35%. Similar announced Lockhart [11], results who randomized. double-blinded. conducted placebo-controlled study on 70 subjects. Hem cultures were positive in 94% of the control patients and in 84% of patients who rinsed with chlorhexidine, with no evidence statistically significant difference.

5. CONCLUSION

Data from the conducted study demonstrated that preoperative rinsing with 0.2% chlorhexidine digluconate has no statistically meaningful effect on reduction of bacteremia following closed extraction of a single tooth.

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