

Original Article

Incidence and Pattern of Bacterial Growth in Propofol Vial - An In vitro Study

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Received : 01-12-2015

Review Completed : 13-05-2016

Accepted : 14-05-2016

Keywords : Propofol, Contamination, Bacterial growth

Abstract :

Introduction- Propofol vials are often used in parts or are opened and left unattended. This has lead to blood stream infections, surgical site infections and acute febrile episodes. A prospective observational study was undertaken to know the incidence and pattern of bacterial growth in samples of Propofol in tropical climate.

Materials and methods- Samples were collected from vials of propofol of different brands, both with and without edetate at different time intervals with relation to room temperature. Each sample of 1ml were inoculated in Brain Heart Infusion (BHI) and incubated for 48hours. Presence of bacterial growth and their pattern were studied.

Statistical analysis used- Paired t test for categorical variables and for non categorical variables Levine's test and Pearson correlation.

Results- Overall 42.26% of samples showed bacterial growth. The incidence was more in samples of propofol without edetate (43.75%) compared to samples with edetate (41.97 %). Most common organism was *Staphylococcus aureus*, followed by *Enterococcus*, *Acinetobacter*, *Bacillus* species, *Pseudomonas* and *Staphylococcus citreus*.

Conclusion- Propofol vial once opened favours bacterial colonisation and growth. Adding edetate to propofol has not shown much benefit in decreasing the incidence.

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Introduction

Propofol, being a lipid emulsion, is a good medium for bacterial growth. Adhering to manufacturer's recommendations of "use without delay" is often difficult. This is because of the need to use small quantity of the drug during general anaesthesia, such as to reduce the intubation response, increase the depth of anaesthesia, prevention of extubation response and laryngospasm in addition to induction of anaesthesia. This results in a delay between opening of the vial and using the last portion of the drug. Extrinsic contamination of propofol by various microorganisms has been associated with outbreaks of bloodstream infections, surgical site infections, and acute febrile episodes^{1,2,3,4}

Available literature regarding bacterial growth in propofol

as such and effect of additives such as edetate is scanty. Further, the pattern of bacteria also depends on the geographic location and temperature. Hence, a study was undertaken to find out the pattern of bacterial growth in a series of randomly selected samples of propofol.

Materials and methods

A prospective observational study was conducted in department of Anaesthesiology in association with department of Microbiology in a tertiary care hospital. The study was conducted with the aim to determine the incidence of growth of microorganisms in samples of propofol in tropical climate, the pattern of microbial growth and any difference in incidence between samples of propofol with or without edetate. The study sample was collected from different operation theatres. Vials of

propofol, both with and without edetate were used for the study. The opening time of all the vials, from which samples were obtained, was noted. Subsequently, series of samples were taken at different intervals starting from immediately after opening the vial to as long as 6hrs 30minutes. The samples were collected using 2cc syringe with aseptic precautions. Each sample of 1ml was inoculated in Brain Heart Infusion (BHI) in cork screw bottles and the theatre temperature was noted. The cork screw bottles were transported to microbiology laboratory. The incubation in the microbiology department was performed within 30 minutes of inoculation in the transport media – Brain Heart Infusion. Incubation was performed in blood agar and McConkey agar at 37 degree Celsius for 48 hours. The incubated samples were then evaluated for the presence of microorganism by standard microbiological procedures and the organism was identified. If there were no organisms grown, the samples were kept incubated and evaluated for the presence of organism after 72hours. The flora grown from routine sampling of operation theatres were Staphylococcus aureus and Bacillus species being the commonest, followed by coagulase negative Staphylococcus species and non-fermenters of gram negative bacilli respectively.

Results

Out of one hundred, 3 samples were lost while transferring from OT complex to microbiology laboratory. Final analysis included 97 samples. Statistical analysis was performed using Paired t test for categorical variables and non categorical by Levene test and Pearson correlation. No correlation was found either between openings of the propofol vial and inoculation time (Table 1 & Table 3, Graph 1) or between differences of the temperature and bacterial isolation (Table 2). Total of 41 samples among 97 studied showed bacterial growth (42.26%). In propofol vials without edetate, organisms were grown in 43.75% (7 samples in 16) whereas in propofol vials with edetate, incidence was 41.97% (34 samples in 81) (Table 4). However the difference in incidence between the samples with and without edetate was not statistically significant (p value>0.005) One sample without edetate grew two

species of bacteria, Staphylococcus and Bacillus species. Most common organism was Staphylococcus aureus (70%), followed by Enterococcus (12%), Acinetobacter (7%) and Bacillus species, Pseudomonas and Staphylococcus citreus (Table 4).

Table 1 : Correlation between opening time and inoculation time

		Opening time	Inoculation time
Opening time	Pearson correlation sig. (2-tailed)	1	0.584- 0.000
	N	100	100
Inoculation time	Pearson correlation sig. (2-tailed)	0.584-0.000	1
	N	100	100

Correlation is significant at the level 0.01 (2- tailed)

Table 2 : Correlation between Temperature difference and bacterial growth Group statistics

Temperature	Edetate	N	Mean	Standard deviation	Standard error Mean
	Present	83	23.42	3.190	0.350
	Absent	17	23.12	1.409	0.342

Independent samples test

		Levene's test of equality of variances		t-test for equality of means		
		f	Sig.	t	df	Sig.
Temperature	Equal variances assumed	4.755	0.032	0.384	98	0.702
	Equal variances not assumed			0.621	55.322	0.537

Independent samples test

		t-test for equality of means			
		Mean difference	Standard error difference	95% confidence interval of the difference	
				lower	upper
Temperature	Equal variances assumed	0.304	0.791	-1.266	1.875
	Equal variances not assumed	0.304	0.489	-0.676	1.284

Table 3 : Difference in bacterial growth depending upon inoculation time and opening time of the vial Group statistics

Difference between the inoculation time and opening time	Edetate	N	Mean	Standard deviation	Standard error Mean
	Present	83	2.6537	1.93182	0.21204
	Absent	17	2.7518	1.35054	0.32755

Independent samples test

		Levene's test of equality of variances		t-test for equality of means		
		f	Sig.	t	df	Sig. (2-tailed)
Difference between the inoculation time and opening time	Equal variances assumed	6.661	0.011	-0.199	98	0.843
	Equal variances not assumed			-0.251	31.153	0.803

Independent samples test

		t-test for equality of means			
		Mean difference	Standard error difference	95% confidence interval of the difference	
				lower	upper
Difference between the inoculation time and opening time	Equal variances assumed	-0.09803	0.49235	-1.07509	0.87903
	Equal variances Not assumed	-0.09803	0.39020	-0.89369	0.69763

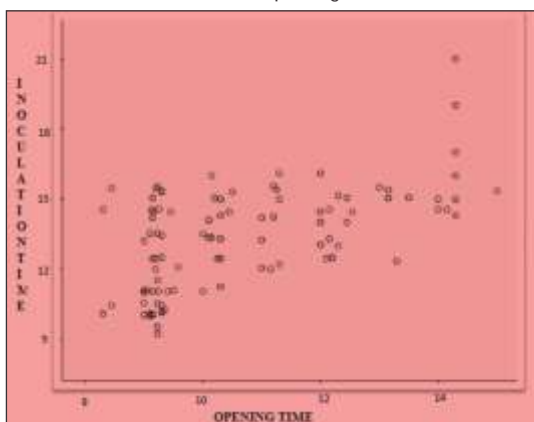
Table 4 : Difference in bacterial growth depending on whether edetate present or absent Case processing summary

Time of inoculation	Samples with edetate			Samples without edetate		
	Total	Organism present		Total	Organism present	
0 to 30min	7	Nil	-	0	nil	-
30min to 90min	25	13	11- Staph aureus 1-Enterococcus 1-Pseudomonas	4	3	3-Staph aureus 1- missing
90min to 150min	15	9	2-Enterococcus 1-Bacillus sp 1-Staph aureus 1- Staph citreus 1-Acinetobacter	5	3	1-Enterococcus 1-Staph aureus 1-Staph aureus and Bacillus species
150min to 270min	20	6	5-Staph aureus 1-Acinetobacter 1-missing	6	1	1-Staph aureus
270min to 390min	16	6	4-staph aureus 1-Acinetobacter 1-Enterococcus 1- missing	2	nil	

Samples					
valid		missing		Total	
N	Percentage	N	Percentage	N	Percentage
97	97%	3	3%	100	100%

		Organism		Total
		Present	Absent	
Edetate	Present	34	47	81
	Absent	7	9	16
Total		41	56	97

Graph 1 : Correlation between opening time and inoculation time



Discussion

The day-to-day practicalities of clinical anaesthesia dictate that some delay between the opening of propofol and its injection into the patient is unavoidable. Further, these delays are variable in both duration and causation. Our study compared the incidence of bacterial growth in propofol vials from immediately after opening of the vial up to 390 minutes after opening the vial. Bacterial contamination was observed in 42.26% of samples, significantly more compared to other studies which reported incidence rates ranging from 6.3% to 26%. One of

the reasons for lower incidence rates in the previous studies were due to immediate inoculation after opening the vial with the maximum time being up to 90 minutes⁵. However, even in our study, there was no statistically significant difference in incidence of bacterial growth with different time intervals of opening and inoculation. This finding is consistent with those of previous studies⁵.

Type of bacteria

Overall, *Staphylococcus aureus* (70%) was the most common organism, followed by *Enterococcus* (12%), *Acinetobacter* (7%) and *Bacillus* species, *Pseudomonas* and *Staphylococcus citrius* in our study. Previous studies⁵ have also shown *Staphylococcus* species was the commonest isolated organism followed by *Diphtheroids*, *Micrococcus* and *Bacillus* species. One sample had grown two species of bacteria, *Staphylococcus* and *Bacillus* species.

Staphylococcus aureus is both a commensal organism and a pathogen. The anterior nares are the main ecological niche for *S. aureus*⁶. Approximately 20% of individuals are persistently nasally colonized with *S. aureus*, and 30% are intermittently colonized. However, other sites may be colonized, including the axillae, groin, and gastrointestinal tract. Colonization provides a reservoir from which bacteria can be introduced when host defences are breached, whether by shaving, aspiration, insertion of an indwelling catheter, or surgery. Colonization clearly increases the risk for subsequent infection^{6,7,8}.

Bacillus species are common contaminating organisms of addicts' heroin and injection paraphernalia, due to their ubiquity in natural, domestic and hospital environments and their production of resistant endospores⁹. Reports of "pseudo-outbreaks" of *Bacillus* species is connected with contaminated clinical and laboratory equipment⁹.

One study¹ isolated *Moraxella osloensis*, *Enterobacter agglomerans* and *Serratia marcescens* from opened propofol vials. Studies have shown that propofol supports the growth of *Staphylococcus* species^{10,11}, *Candida albicans*^{10,11}, *Moraxella osloensis*¹⁰ and *Pseudomonas aeruginosa*¹¹. Another study¹² showed that propofol

strongly supports the growth of *E. coli* and *C. albicans* but is bacteriostatic toward *S. aureus* and weakly bactericidal toward *P. aeruginosa*.

Staphylococcus aureus, *epidermidis*, *diphtheroids* were the bacterial strains isolated from propofol by preparation technique similar to the clinical practice in a study¹³.

Enterococci are a part of the normal human faecal flora. Sites less often colonized are the oral cavity, genitourinary tract and skin especially in the perineal area. The main sites of colonization in the hospitalized patients are soft tissue wounds, ulcers and the gastrointestinal tract (GIT)¹⁴. Enterococci have emerged as an important cause of nosocomial infections. These infections are recognized by 3 ts - tough, tenacious and often troublesome^{14, 15}. The most frequent infections caused by enterococci are urinary tract infections^{14, 16}. The second most frequent enterococcal infections are intra - abdominal and intra - pelvic abscesses or post-surgery wound infections^{14, 16}. The third most frequent infection caused by these organisms is blood stream infections^{14, 17}. Other infections caused with lower frequency are central nervous system (CNS) and neonatal infections. Enterococci rarely cause respiratory tract infections, osteomyelitis, or cellulitis^{14, 18}.

The gram-negative coccobacillus, *Acinetobacter*, a pathogen once seen only in hot, humid climates, has become an increasingly common nosocomial problem even in temperate climates¹⁹. The association of *A. baumannii* with pneumonia, bacteremia, wound infections, urinary tract infections, and meningitis has been well described^{20,21}. Risk factors associated with colonization or infection (which can be difficult to distinguish) include prolonged hospitalization, intensive care unit admission, recent surgical procedures, antimicrobial agent exposure, central venous catheter use, prior hospitalization, nursing home residence, and local colonization pressure on susceptible patients^{20,21}.

The prevalence of colonisation by *P. aeruginosa* in healthy subjects is usually low, but higher colonisation rates can be encountered following hospitalisation, especially amongst

subjects treated with broad-spectrum antimicrobial agents. Colonisation is common in the respiratory tract of mechanically ventilated patients, in the gastrointestinal tract of patients receiving anticancer chemotherapy, and on the skin of burn patients²². Also, sinks, mops, disinfectant solutions, respiratory equipment, food mixers and other moist environments can act as reservoirs of *P. aeruginosa* in the hospital setting^{22,23,24}.

Staphylococcus citreus occurs as the normal flora of nose in upto 1% and normal flora of throat in 15 – 25%²⁵.

The difference in type of bacteria compared to previous studies^{1,5,13} could be explained by geographical differences and local practices between the present and previous study locations.

Effect of edetate

Our study did not show statistically significant difference in incidence (p value=1) between bacterial growth in propofol vials with and without edetate. With absence of edetate

vials organism were found in 43.75% versus 41.97%. Sticking to propofol handling guidelines²⁶ including hand wash may help in decreasing the incidence of bacterial growth in propofol vials. Edetate has proven to have inhibitory effect on bacterial growth²⁷. Our study could not establish statistically significant difference in incidence of bacterial growth at different temperatures. Further studies are required to evaluate the effect of temperature on bacterial growth.

Conclusion

Propofol being lipid emulsion facilitates bacterial growth and edetate, though has inhibitory effects, do not completely prevent bacterial growth. Hence strict aseptic precautions including hand wash and safe injection practices for administration of propofol have to be followed. The gap between what is recommended and what is actually done clinically regarding safe injection practices should be identified and rectified.

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