

Comparison Between the Efficacy of Saline, Alkalinised Lignocaine 2% and Alkalinised Lignocaine 4% in the Cuff of Endotracheal Tube to Reduce the Incidence of Cough and Sorethroat During Extubation

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To compare the efficacy of saline, lignocaine 2% and Lignocaine 4% in the endotracheal tube cuff to decrease the incidence of post operative cough and sore throat. A unique, efficient and easy method designed to decrease cough and sore throat during emergence. Hence there was a need for the study. Methods: After approval from institutional Ethical Committee, Kasturba Medical college, Mangaluru and written informed consent from 120 patients of A.S.A class I & II, aged between 18-60 years posted for various surgeries were included. Randomization was done into three groups of 40 patients each. Group 1 – Normal saline, group 2 - 2% Lignocaine and group 3 - 4% Lignocaine. Postoperatively patients were assessed for incidence of cough and sore throat pain based on visual analogue scale (VAS) Results: 65% of patients in group 2 and 72% of patients in group 3 were extubated smoothly, whereas only 20% of patients in group 1 had smooth extubation, $p < 0.001$. There was significant difference in sore throat pain and cough in group 2 and 3 when compared to group 1. Conclusion: The incidence of sore throat and cough was significantly reduced with Lignocaine in the endotracheal tube cuff when compared to saline. Lignocaine 4% was found to be far more better than 2% in reducing sore throat.

Keywords: Alkalinised Lignocaine 2%, 4%; Cough; Emergence; Saline; Sore Throat.

Airway management with cuffed endotracheal tube in General Anaesthesia is an important part of an anaesthesiologist's responsibilities towards patient. Common complications like cough and sore throat are noted at emergence from general anaesthesia due to inflation of the endotracheal tube cuff. Coughing during emergence results in tachycardia, increased blood pressure, increased intracranial and intraocular pressure, surgical blood loss. This can be of significance in vascular, ophthalmic and

neurosurgical procedures. Postoperative sore throat is seen in upto 39-86% of patients¹. It is due to various factors like inflammation, haemorrhage, ciliary loss and vocal cord oedema. Topical anaesthetics like lignocaine administered in the endotracheal tube cuff represents an innovative technique to decrease the post-operative cough and sore throat². Hence there was a need for this study. The airway mucosa coming in direct contact with the ET tube cuff can be anesthetised locally with higher doses of lignocaine (200-500 mg) without compromising supra glottic airway reflex^{3,4}.

MATERIALS AND METHODS

A prospective, randomised control study was done after obtaining Institutional Scientific and Ethics committee approval. 120 ASA class I or II patients, posted for surgical procedure (minimum 90 minutes of surgical duration or anaesthesia time of > 120 minutes), aged between 18- 60 years were approved for the study. Patients not willing to participate in the study, asthma, severe pulmonary disease like COPD or emphysema, anticipated difficult intubation, subjects on cough suppressants medication, allergic to lignocaine, pregnancy and H/O previous airway surgeries were eliminated from the study. All patients were visited day before surgery for pre anaesthetic evaluation (PAC) and written consent was obtained for the study. On the day of surgery before shifting to Operation Theatre (OT), pre-anaesthetic (PAC) orders were reconfirmed. In the OT monitors like electrocardiography (ECG), non-invasive blood pressure (NIBP), pulse oxymetry (SpO₂), were connected and Baseline parameters like heart rate-HR, blood pressure (SBP, DBP, MAP) were documented and intravenous access secured. Patients were pre oxygenated with 100% oxygen for 3 minutes before inducing the patient. Premedication was done using glycopyrrolate 0.2mg/kg, midazolam 0.03 mg/kg along with fentanyl 2 µg/kg. All the Patients were induced with propofol, and muscle relaxant atracurium 0.5 mg/kg. Endotracheal intubation was done with proper sized cuffed oral endotracheal tube and the cuff was filled with saline or lignocaine according to respective groups. The Quantity of substance filled in the high volume low pressure cuff were adequate not to cause any leak when ventilated.

120 patients were randomly divided into three groups. Endotracheal tube cuff was filled with either saline or lignocaine or lignocaine with sodium bicarbonate. Group 1: 6.5 ml normal saline, Group 2: 6 ml 2% lignocaine Hydrochloride + 0.5 ml 7.5% sodium bicarbonate, and Group 3: 6 ml 4% lignocaine Hydrochloride + 0.5 ml 7.5% sodium bicarbonate. Anaesthesia was managed with nitrous oxide oxygen mixture with isoflurane. After the surgery, glycopyrrolate 0.02mg/kg and neostigmine 0.05 mg/kg was used to reverse residual blockade. Suctioning was done, cuff deflated and patients were extubated once criterias

for extubation is met. The cardiovascular responses at extubation were recorded. Just before extubation, patients were assessed for the incidence of cough. Post-operative sore throat pain were assessed based on visual analogue scale (VAS score) during immediate post op, 2nd hour, 4th hour and at 24 hour following extubation.

Statistical Analysis

With 95% confidence interval and 80% power with respect to p1 = 3.3 and p2 = 23.3, the sample size was n=120.

$$n = \frac{(Z\alpha\sqrt{2PQ} + Z\beta\sqrt{p1q1 + p2q2})^2}{(p1 - p2)^2}$$

Zα = 1.96 at 95% Confidence level, Zβ = 0.84 at 80% power, p1 = 3.3; p2 = 23.3, P = p1 + p2/2 = 13.3, q = 100 - p

SPSS version 17 was used for analysis. Using ANOVA, Bonferroni's post hoc test, Chi-square test and Fisher's exact test, Values were expressed as mean (SD) or median (range). 'p' value of < 0.05 was considered as statistically significant.

RESULTS

Factors like age, sex and also the surgery duration were indistinguishable between the groups. 20% of patients in group 1, 65% of patients in Group 2 and 72% of patients in Group 3 were extubated smoothly, p < 0.0001. We experienced laryngospasm in 1 patient in group 1, which was managed appropriately.

Data suggests a significant difference in sore throat pain between group 3, group 2 when compared with group 1. Cough was significantly higher in group 1, when compared to group 2 and 3. Post-operative sore throat pain was found to be greater in the group 1 (saline) compared to group 2 (lignocaine 2%) and group 3 (lignocaine 4%) during the post-operative 4th hour and increased to reach maximum percentage during the 6th hour and gradually decreased to 5% at the end of 24 hour. There was no much variation between group 2 (lignocaine 2%) and group 3 (lignocaine 4%) (Graph 1). Visual Analogue Scale (VAS) score was maximum in Group 1 (saline) when compared to Group 2 and 3 in the immediate, 2nd, 4th and 24 hour

postoperative period. Kruskal- Wallis test is a non-parametric test, performed to see the differences in the sore throat pain scores in the three groups which showed a significant differences with p value less than 0.001 (Table 1).

Pairwise comparison by Mann-Whitney test shows a highly significant difference between

the group 1 versus group 2 and group 3 during the immediate post-operative period, post op 2nd hour, 4th and 24 hour. (Table 1). Heart rate recorded at the time of extubation, 5 min, 10 min and at the end of 30 min shows a significant difference among the 3 groups. There is considerable decrease in the heart rate in the group 2 (lignocaine 2%) and group 3

Bar diagram showing the difference in the post-operative sore throat pain between the three groups.

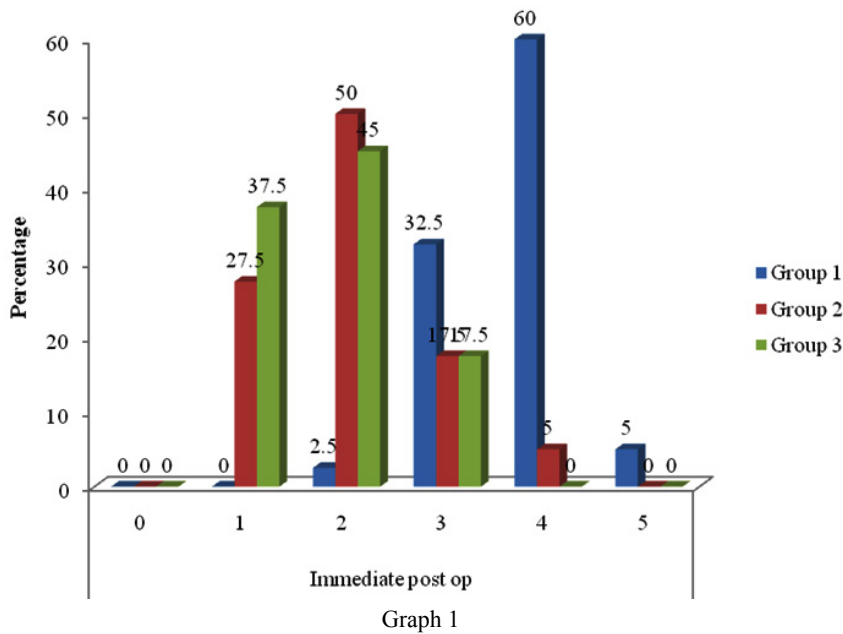


Table 1. Kruskal-Wallis Test and Pairwise comparison by Mann Whitney test comparing post-operative sore throat

Time Point	Group	Kruskal-Wallis Test p value	Significance	Pairwise comparison by Mann Whitney test p value	
				Comparison	p value
Immediate post op	Group 1	0	HS	Group 1 VS Group 2	.000
	Group 2			Group 1 VS Group 3	.000
	Group 3			Group 2 VS Group 3	.307
Post op 2 hour	Group 1	0	HS	Group 1 VS Group 2	.000
	Group 2			Group 1 VS Group 3	.000
	Group 3			Group 2 VS Group 3	.110
Post op 4 hour	Group 1	0	HS	Group 1 VS Group 2	.000
	Group 2			Group 1 VS Group 3	.000
	Group 3			Group 2 VS Group 3	.080
Post op 24 hr	Group 1	.000	HS	Group 1 VS Group 2	.000
	Group 2			Group 1 VS Group 3	.000
	Group 3			Group 2 VS Group 3	.003

VS = Very significant HS= Highly significant

(lignocaine 4%) compared to the heart rate in group 1 (saline). P value in this ANOVA test is less than 0.001% (Table 2). Results showed that there was a significant decrease in cough in group 2 (lignocaine 2%) and group 3 (lignocaine 4%) compared to group 1 (saline). Number(%) of patients having cough in group 1 (saline), 2 (lignocaine 2%) and 3 (lignocaine 4%) was 57.5%, 7.5% and 0% respectively (Table 3).

DISCUSSION

The main aim of this study was to compare the best inflating agent between saline, lignocaine 2%, lignocaine 4% that were used to inflate the cuff during anaesthesia. Administration of lignocaine into the endotracheal tube cuff resulted in the

diffusion of lignocaine out of the cuff on the tracheal mucosa. During general anaesthesia with nitrous oxide, the pressure inside the cuff increases as the nitrous oxide easily diffuses into the cuff made of polyvinyl chloride than diffusing out of the cuff because of the difference in the pressure across the cuff membrane⁵. When the endotracheal tube cuff pressure exceeds the capillary perfusion pressure (22mmHg), there is decreased blood supply to the underlying mucosa of the tracheal rings and posterior wall of the trachea resulting in ulceration and postoperative sore throat. By administering liquid medium (saline/lignocaine), consequences related to hyperinflation can be avoided^{5,6}. Rapidly acting stretch receptors are present throughout the mucosa of tracheal wall. Mechanical stimuli of these receptors are the presumed mechanism for the

Table 2. One way ANOVA Test, a parametric test was done to compare the differences between the heart rate in the three groups at extubation, 5 minute, 10 minute and 30 minute post extubation

		N	Mean	Std. Deviation	95% Confidence Interval (Mean)		ANOVA test p value	
					Lower Bound	Upper Bound		
Extubation	Group 1	40	100.20	5.774	98.35	102.05	.000	HS
	Group 2	40	94.75	5.513	92.99	96.51		
	Group 3	40	95.10	4.378	93.70	96.50		
5 min	Group 1	40	103.80	6.236	101.81	105.79	.000	HS
	Group 2	40	98.05	5.203	96.39	99.71		
	Group 3	40	101.95	4.579	100.49	103.41		
10 min	Group 1	40	105.13	6.794	102.95	107.30	.000	HS
	Group 2	40	96.25	5.471	94.50	98.00		
	Group 3	40	101.53	4.200	100.18	102.87		
30 min	Group 1	40	101.43	7.171	99.13	103.72	.000	HS
	Group 2	40	90.63	5.182	88.97	92.28		
	Group 3	40	91.38	5.143	89.73	93.02		

HS= Highly significant

Table 3. Comparison of group A (saline), group B (lignocaine 2%) and group C (lignocaine 4%) based on the presence of cough in the post operative period

		Group						Total	
		Group 1		Group 2		Group 3		Count	%
		Count	%	Count	%	Count	%	Count	%
cough	Absent	17	42.5%	37	92.5%	40	100.0%	94	78.3%
	Present	23	57.5%	3	7.5%	0	.0%	26	21.7%
Total		40	100.0%	40	100.0%	40	100.0%	120	100.0%

p=0.001, HS = Highly significant

production of cough. Endotracheal tube intubation, inflation of the cuff and the resulting hyperinflation stimulates the stretch receptors and produces cough during extubation in normal patients. Lignocaine acts on these stretch receptors and blocks the cough reflex. Comparing the present study to an earlier conducted study where intravenous and topical lignocaine were used to decrease the cough during extubation^{2,4}. Lignocaine is a CNS depressant agent and it acts on the cough centre in the medulla and depresses the cough reflex. When given intravenously plasma concentration of 0.3µg/ml is sufficient to suppress cough under general anesthesia and it lasts for a duration of 15-20 minutes. Intravenous lignocaine also results in delayed recovery from anaesthesia⁵.

In another study saline, alkalised lignocaine and air were compared in decreasing postoperative cough and sore throat where alkalised lignocaine showed better results which was comparable with our study^{5,7,8}. Alkalised lignocaine also known to decrease analgesic and sedative requirements in mechanically ventilated patients which was proved in another study⁹. One other study concluded that there was no lignocaine overdose, systemic toxicity and endotracheal tube cuff rupture when intracuff lignocaine was used¹⁰. Some other studies showed lignocaine when used with dexamethasone produced better results when compared to lignocaine used alone^{11,12}. In the present study we found that there was a statistically significant decrease in the post-operative sore throat and sore throat pain in patients in immediate postoperative period, 2nd hour, 4th hour and 24 hour belonging to group 2 (6ml 2% lignocaine Hydrochloride + 0.5 ml 7.5% sodium bicarbonate) and group 3 (6ml 4% lignocaine Hydrochloride + 0.5ml 7.5% sodium bicarbonate) compared to group 1 (normal saline). The results obtained were similar to the earlier conducted study. The incidence of cough is significantly reduced in the group 2 (6 ml 2% lignocaine Hydrochloride + 0.5 ml 7.5% sodium bicarbonate) and group 3 (6ml 4% lignocaine Hydrochloride + 0.5ml 7.5% sodium bicarbonate) compared to group 1 (saline). Studies previously have compared intravenous, intracuff lignocaine and saline, but alkalised 2% and 4% lignocaine had never been studied. Hence there was scope for this study. Filling buffered lignocaine into the endotracheal tube resulted in

diffusion of uncharged lignocaine across the cuff^{1,13,14}. Lignocaine being a weak basic, lipophilic drug binds ardently to the respiratory mucosa to produce the effect.

CONCLUSION

The incidence of post-operative sorethroat is significantly reduced when lignocaine is administered into the endotracheal tube cuff compared to the saline. On comparing lignocaine 2% and lignocaine 4%, lignocaine 4% was found to be better in reducing the postoperative sorethroat. Administration of lignocaine into the cuff also resulted in significant reduction of cough.

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