Comparative study of dexamethasone nebulisation with magnesium sulphate nebulisation in preventing post operative sore throat following endotracheal intubation

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Abstract

Introduction: Post-operative sore throat is one of the most common complications following endotracheal intubation. Though considered minor complication, it may cause significant patient dissatisfaction. Various non-pharmacological and pharmacological trials have been used with variable results.

Objectives: To compare the efficacy of nebulised dexamethasone with that of nebulised magnesium sulphate in decreasing the incidence and severity of postoperative sore throat (POST).

Materials and Methods: In this prospective double blind study 90 patients undergoing surgery under general anaesthesia with endotracheal intubation lasting <3hr were randomly assigned into two equal groups. Group D received dexamethasone 8mg (2ml) with 3ml saline nebulisation and group M received magnesium sulphate (50%W/V 2ml) with 3ml saline nebulisation 30 min before the induction of anaesthesia. Primary outcome assessed was incidence and severity of POST. Secondary outcome assessed were the incidence of post-operative hoarseness and cough.

Results: Compared to group M, significantly lesser number of patients in group D had post-operative sore throat at 0hr (p=0.0262), 4th hr (p=0.000022), 8th hr (p=0.00039) and 12hr (p=0.000657). None of the patients in group D had any hoarseness of voice at 0hr, 4th hr, 8th hr of assessment (p<0.05). Except one patient in group M, none of our patients in either of the group had cough at any point of assessment.

Conclusion: Preoperative dexamethasone nebulisation just before induction of anaesthesia is an effective method of reducing the incidence and severity of POST following endotracheal intubation. Dexamethasone nebulisation reduces the severity of sore throat more effectively than magnesium sulphate nebulisation.

Keywords: Postoperative sore throat, Nebulisation, Hoarseness, Cough, Dexamethasone, Magnesium sulphate.

Introduction

Sore throat is one of the common postoperative complaint that leads to morbidity and patient dissatisfaction. It is 5th most frequent adverse clinical anaesthesia outcome. The incidence of sore throat has been reported to be very high [21%-65%]. Irritation and inflammation of the airway, mechanical injury during intubation, damage to the mucosa due to the pressure from the endotracheal tube cuff and dehydration of the mucosa were considered the cause of postoperative sore throat (POST). Although symptoms subside without any treatment, the management for POST is still advised because it enhances patient satisfaction, acceptance of anaesthesia and improves the activities after discharge.

Prophylactic management of POST is recommended to improve the quality of anaesthesia care of duration of time a patient stays in the post-anesthesia care unit because of POST increases the cost of care. Patients with POST had a 14min longer stay in the post anaesthesia care unit and 25 min longer stay in the ambulatory care unit and were discharged 51 min later from the facility compared with those who did not complain of POST.

Different agents like ketamine gargle and nebulisation, endotracheal tube spraying with Beclomethasone, intravenous injection of dexamethasone, magnesium sulphate gargle and Lozenges, nebulized lignocaine, lignocaine jelly have been used with variable success for decreasing both the incidence and severity of POST.

Corticosteroids are used in the perioperative period to enhance the effects of analgesics and antiemics. Inhaled corticosteroids deliver the drug to the airways without systemic effects. Dexamethasone is a potent steroid that has 26.6 and 6.6 times stronger anti-inflammatory and immune-suppressant effects than cortisol and prednisone, respectively. It has been supported that it is very useful for relieving POST. Therefore, inhaling dexamethasone may be used as a method to reduce POST following general anesthesia.
to the airways with no or minimal systemic effects. Both magnesium sulphate and dexamethasone are readily available in the operation theatre. Present study was done to compare the efficacy of dexamethasone nebulisation pre-operatively with that of magnesium sulphate nebulisation. In addition we also observed for other airway related complaints like hoarseness and cough, postoperatively.

**Materials and Methods**

This was a prospective double blind randomised study conducted after obtaining approval from our hospital ethical committee (IEC NO: SIMS & RC/IECC/05/2017). Written informed consent was obtained from patients after proper explanation regarding the study. ASA1 & ASA2 physical status patients aged between 20 to 50yrs admitted for elective surgery under general anaesthesia with endotracheal intubation with mallampatti score 1-2, undergoing surgeries lasting less than 3hrs duration were selected for the study. 90 patients were chosen by simple chit picking method into two groups of 45 patients each. To show 50% reduction in the incidence of POST (which is reported to be 65%) at $\alpha=0.05$, confidence interval of 95% and a power of 90%, we required 30 patients each group. On adding 10%-15% patients loss due to multiple attempts at laryngoscopy and due to bucking during extubation, each group needed 40 to 45 patients.

Patients with history of sore throat, on long term anti-inflammatory analgesic medications, patients who have undergone insertion of devices which stimulate oral cavity, larynx, pharynx like nasogastric tube, endoscopic nasobiliary drainage tube, mallampatti grading $>2$, patients who are undergoing head and neck surgeries, patients who needed more than two attempts at endotracheal intubation and use of bougie and styllete, obese, diabetic, pregnant patients, surgeries lasting for more than 3hrs, surgeries in prone position were excluded from the study.

Group D received dexamethasone 8mg [2ml] with 3ml of normal saline for nebulisation.

Group M received MgSo4 [50% W/V 2ml] with 3ml of normal saline for nebulisation 30 min before the induction of anaesthesia in the preoperative receiving room. The patients were nebulised using mask connected to wall mounted oxygen driven source. (8lt, 50psi) A nurse blinded for the study nebulised the patient using the study drug prepared by an anaesthesiologist not involved in the study. All the intubations were done by an anaesthesiologist with >5yrs experience. All patients were kept nil oral overnight and were premedicated with table. Ranitidine150mg and table. Alprazolam 0.5mg 2hrs before the surgery.

After the nebulisation for 30 min in the receiving area, patients were shifted into the OT.

Patient premedicated with inj.glycopyrrolate 0.2mg and midazolam 1mg. Injection fentanyl 1 was given 2mcg/kg. Patient was induced with inj. propofol 2mg/kg. Intubation was done using non-depolarising muscle relaxant, inj.Vecuronium0.15mg/kg. Intubation was performed as smooth direct laryngoscopy after 4min of injecting vecuronium. Endotracheal tube size 7 was selected for females and 8 was selected for males. A single use portex high volume low pressure cuffed endotracheal tube was used for the intubation. Patient were excluded from the study if intubation failed in two attempts or O2 saturation dropped below 95% during intubation. Anaesthetic gas mixture was used to inflate the endotracheal tube cuff till no leak was heard around the tube. Cuff pressure was adjusted to 20-22CmH2O using cuff inflator/ pressure gauge PORTEX TM. Cuff pressure was monitored every half an hour to maintain the cuff pressure of 20 to 22CmH2O.

Maintenance of anaesthesia was the same in both the groups using isoflurane 1MAC and vecuronium divided doses with 50%O2+50%N2O. 20 min before the completion of the procedure inj. ondansetron 4mg and inj.pethidine0.5mg/kg mg IV was given to prevent post operative nausea and vomiting and bucking on the tube respectively. Neuromuscular blockade was reversed using neoistigmine 0.05mg/kg and glycopyrrolate 0.001mg/kg. Gentle suctioning of oropharynx was done. Extubation was performed using the same method in both the groups when the patient was fully awake. Patients who bucked or coughed during extubation were excluded from the study. In the recovery room patient received O2 by mask 5lt/min and inj. Paracetamol 1gm 8th hr.

**Assessment:** Patients were assessed in the postoperative ward by a nursing staff blinded for the study. Incidence of sore throat, discomfort, pain and other symptoms were assessed at the time of shifting to postoperative ward [0hr], 4hr, 8hr, 12hr and 24hrs according to the grading shown in Fig.1.

**SEVERITY**

| Grade | Description |
|-------|-------------|
| 0     | No sore throat at any time since the operation |
| 1     | Minimal - patient answered in the affirmative when asked about sore throat |
| 2     | Moderate - patient complained of sore throat on his/ her own |
| 3     | Severe - patient is in obvious distress |
| 0     | No cough at any time since the operation |
| 1     | Minimal |
| 2     | Moderate |
| 3     | Severe |
| 0     | No complaint of hoarseness at any time since the operation |
| 1     | Minimal - minimal change in quality of speech. Patient answers in the affirmative only when enquired about |
| 2     | Moderate - moderate change in quality of speech of which the patient complains on his/her own |
| 3     | Severe - gross change in the quality of voice perceived by the observer |

**POST - Post-operative sore throat**

Fig. 1
Total incidence of POST was calculated by dividing the number of patients who experienced postoperative throat pain at least once by the total number of patients. The patients were then examined to determine the occurrence of any other postoperative complications and the findings were recorded in detail. Patients who had POST grading more than 2 were instructed to gargle aspirin 75mg dispersible tablet.

**Statistical Analysis**

The collected data was analysed using statistical package for social sciences (SPSS, V-20) Descriptive and inferential statistical analysis has been carried out in the present study.

Results on continuous measurements are presented on Mean ± SD and results for P value were calculated using t-test. Categorical measurements are presented in number. Chi square test is used to compare the difference in each group. Significance was assessed at 5% level of significance. If P <0.005 is considered statistically significant.

**Table 1**

| Parameters          | Group M       | Group D       | P-value |
|---------------------|---------------|---------------|---------|
| Age                 | 37.65±10.060  | 36.88±9.053   | 0.228   |
| Duration of Surgery | 112.83±38.046 | 119.93±36.287 | 0.336   |
| Gender              |               |               |         |
| Male                | 25            | 26            |         |
| Female              | 15            | 14            | 0.816   |
| ASA                 |               |               |         |
| 1                   | 29            | 33            |         |
| 2                   | 11            | 7             | 0.284   |
| No of attempts      |               |               |         |
| 1                   | 34            | 33            |         |
| 2                   | 12            | 7             | 0.338   |

Incidence of POST was 27.5% in group D and 57.5% in group M and the difference was statistically significant at all times of assessment. (Table 2). Compared to group M, the number of patients in group D who had POST were significantly low at rest at 0 hr (p= 0.0262), 4th hr (p=0.00022), 8th hr (p=0.00039) and 12hr (p=0.000657). (Table 2). Also duration of pain relief was significantly longer (12hrs) in group D compared to group M (P=0.000657). Only one patient in each group had POST at 24hr which was statistically not significant. (P= 1.00) (Table 2)

Incidence of pain on swallowing was also more in patients in group M at all points of assessment compared to group D. (Table 2). Though the highest incidence of sore throat occurred at 4th hr post-extubation in both the groups, it showed declining trend. This decline was significant in group D. (Table 2)

None of the patients in group D had postoperative hoarseness which was statistically significant at 0hr, 4th hr, 8th hr of assessment. At 24hrs none of the patients in both the study groups had any hoarseness. (Table 2)

None of the patients in group D had cough at any point of assessment where as one patient in group M had cough at 4th hr and 8th hr which was clinically and statistically not significant. (P=1.00) (Table 2)
This graph shows the severity of sore throat in both the groups. From the graph it is evident that group D patients had lesser grades of POST. None of the patients in group D had more than grade 1 POST. Where as in group M 8 patients at 0hr, 15 patients at 4th hr, 14 patients at 8th hr and 2 patients at 12hr had grade 2 POST. No patients had severe POST that required further treatment.

27 patients in group M complained of vicks like taste/ smell in the throat after nebulisation. No such complaints were given by patients in group D.

Table 2: Comparison of incidence of post-operative sore throat, cough and hoarseness of voice

| Time      | Occurrence | Group M, n (%) | Group D, n (%) | P-value |
|-----------|------------|----------------|----------------|---------|
| Pain at Rest |            |                |                |         |
| O HRS     | No         | 24(60)         | 33(82)         | 0.0262* |
|           | Yes        | 16(40)         | 7(18)          |         |
| 4 HRS     | No         | 17(43)         | 33(82)         | 0.00022*|
|           | Yes        | 23(57)         | 7(18)          |         |
| 8 HRS     | No         | 19(47)         | 34(85)         | 0.00039*|
|           | Yes        | 21(53)         | 6(15)          |         |
| 12 HRS    | No         | 28(70)         | 39(97)         | 0.000657*|
|           | Yes        | 12(30)         | 1(3)           |         |
| 24 HRS    | No         | 39(97)         | 39(97)         | 1.0000  |
|           | Yes        | 1(3)           | 1(3)           |         |
| Pain on Swallowing |            |                |                |         |
| O HRS     | No         | 30(75)         | 33(82)         | 0.4122  |
|           | Yes        | 10(25)         | 7(18)          |         |
| 4 HRS     | No         | 17(43)         | 33(82)         | 0.00022*|
|           | Yes        | 23(57)         | 7(18)          |         |
| 8 HRS     | No         | 19(47)         | 34(85)         | 0.0039* |
|           | Yes        | 21(53)         | 6(15)          |         |
| 12 HRS    | No         | 29(72)         | 39(97)         | 0.000044*|
|           | Yes        | 11(28)         | 1(3)           |         |
| 24 HRS    | No         | 38(95)         | 39(97)         | 1.0000  |
|           | Yes        | 2(5)           | 1(3)           |         |
| Hoarseness |            |                |                |         |
| O HRS     | No         | 27(68)         | 40(100)        | 0.00414*|
|           | Yes        | 13(32)         | 0(0)           |         |
| 4 HRS     | No         | 24(60)         | 35(88)         | 0.00518*|
|           | Yes        | 16(40)         | 5(12)          |         |
| 8 HRS     | No         | 24(60)         | 40(100)        | 0.0041  |
|           | Yes        | 16(40)         | 0(0)           |         |
| 12 HRS    | No         | 37(92)         | 40(100)        | 0.3049  |
|           | Yes        | 3(8)           | 0(0)           |         |
Discussion

Post-operative sore throat, cough, and hoarseness of voice are common, uncomfortable, distressing sequelae after tracheal intubation. Throat irritation in the presence of a large abdominal or thoracic incision can be very annoying especially in the presence of inadequate analgesia since any attempt to cough to clear the throat causes severe pain. The contributing factors for POST include young age, females, gynaecological surgery, use of succinylcholine, larger tracheal tubes, high cuff pressure.5,24

POST can be multifactorial in origin, including mechanical injury during laryngoscopy and intubation causing aseptic inflammation, continuous pressure by the inflated tracheal tube cuff on tracheal mucosa causing damage and dehydration of the mucosa.5,13,19,20,23

Although many drugs are used through different routes to relieve the POST, these could result in unwanted side effects. Ketamine gargle might cause adverse hemodynamic effects and taste may not be acceptable. Higher doses of drugs are required when used through intravenous route causing unwanted side effects. Gargling requires patient co-operation. So we selected method of nebulisation to deliver the study drug to throat. Dose of drug used is very minimal and acts topically avoiding adverse systemic effects.

Present study shows that prophylactic dexamethasone nebulisation helps in significant reduction in incidence of sore throat at 0hr (P =0.0262), 4th hr (P =0.00022), 8th hr (P =0.00039) and 12th hr (P =0.000657) post extubation when compared to magnesium sulphate nebulisation. Severity of sore throat also was less compared to group M. (Fig. 2) There was no incidence of moderate sore throat (grade 2) in group D patients. None of the patients in group D had hoarseness of voice. Incidence of severe sore throat (grade 3) was not found among both the groups. This may be because we used well defined inclusion and exclusion criteria and intubations were done by an experienced anaesthesiologist, duration of surgery was less than 3hrs, cuff pressure was monitored through out the procedure and bucking on the endotracheal tube was avoided.

Usually steroids and nebulised adrenaline are used for the treatment of POST.18 Intravenous dexamethasone or hydrocortisone is used for this purpose. It has been shown that both, topical and intravenous dexamethasone had reduced the incidence of POST.13,14,18 But glucose intolerance, fluid retention are concerns with the use of intravenous steroids. So we used nebulised dexamethasone at a dose of 8mg. Widespread recognition of under treatment of sore throat by clinicians has led to the development of preemptive strategies for its alleviation. So we used nebulisation just before the induction of anaesthesia.

Our study showed significantly lower incidence of POST in dexamethasone group at all points of assessment. This was similar to the results of Salama et al.21

Tabari et al22 studied the effectiveness of betamethasone gel applied to the tracheal tube and intravenous Dexamethasone on POST on 225 ASAI and ASII patients undergoing elective abdominal surgery with tracheal intubation who were randomly allocated into three groups: the betamethasone gel group, the intravenous dexamethasone group and control group. They concluded that widespread application of betamethasone gel over tracheal tubes effectively reduced POST, compared to intravenous dexamethasone.

These findings are consistent with the topical application of steroid on the upper airway before endotracheal intubation.

Magnesium sulphate gargle10 and lozenges11 have been used for treating POST. We selected magnesium sulphate nebulisation to avoid large doses and for ease of administering the drug. Not much literature is available about the use of nebulized magnesium sulfate for attenuation of POST. Blitz et al.24 used nebulised magnesium sulphate for the treatment of acute asthma without any systemic and local adverse outcomes. To the best of our knowledge, there is currently no study that compares the efficacy of nebulised dexamethasone

|                | 24 HRS | 0 HRS | 4 HRS | 8 HRS | 12 HRS | 24 HRS |
|----------------|--------|-------|-------|-------|-------|--------|
|                | No     | Yes   | No    | Yes   | No    | Yes    |
|                | 40(100)| 40(100)| 39(97)| 39(97)| 40(100)| 40(100)|
|                | 40(100)| 0(0)  | 40(100)| 0(0)  | 40(100)| 0(0)  |
|                | 40(100)| 0(0)  | 40(100)| 1(3)  | 40(100)| 1(3)  |
|                | 40(100)| 0(0)  | 40(100)| 1(3)  | 40(100)| 1(3)  |
|                | 40(100)| 0(0)  | 40(100)| 0(0)  | 40(100)| 0(0)  |
|                | 40(100)| 0(0)  | 40(100)| 0(0)  | 40(100)| 0(0)  |

Intravenous Dexamethasone on 225 ASA1 and ASAII patients undergoing elective abdominal surgery with tracheal intubation who were randomly allocated into three groups: the betamethasone gel group, the intravenous dexamethasone group and control group. They concluded that widespread application of betamethasone gel over tracheal tubes effectively reduced POST, compared to intravenous dexamethasone.
with that of nebulised magnesium sulphate on the incidence and severity of POST. In Borzan et al11 study, there was a significantly decreased incidence of POST when subjects sucked on a lozenge containing magnesium preoperatively at 2nd hr and 4th hr but not immediately or 24hrs post-operatively.

Incidence of POST is more common at 4th to 6th hr is due to the gradually developing local inflammation. But in Our study, more than 50% of patients (Table 2) in magnesium sulphate group had significant POST at 4th hr and 6th hr. This shows that magnesium sulphate is not very effective in controlling POST. Even though incidence of POST was not significantly reduced with the use of nebulised magnesium sulphate, there was significantly reduced incidence of cough. In our study, group M patients who complained of hoarseness had prolonged duration of surgery (>120 min). This may be the cause, because of prolonged cuff pressure causing irritation of mucosa and edema of vocal Cords. So in our study magnesium sulphate nebulisation did not reduce the incidence of POST, but had a role in attenuating the severity of sore throat since none of the patients had grade 3 POST in group M also.

In the present study, the overall incidence of POST was 27.5% in group D and 57.5% in group M which is the same as evaluated by Macario et al.2 Even though few patients in magnesium sulphate group had relief from sore throat, it was not statistically significant.

Limitations of the Study
Firstly we did not use humidity moisture exchanger in the gas delivery circuit and dry gases are implicated in the development of postoperative sore throat 19. Secondly we don’t know how efficient is dexamethasone nebulisation in relieving sore throat in case of multiple attempts at laryngoscopy, prolonged intubation due to long procedures, use of bougies. Repetation of the nebulisation can be considered as a subjective scale and may be associated with bias.

Conclusion
Preoperative dexamethasone nebulisation just before induction of anaesthesia is an effective method of reducing the incidence and severity of POST following endotracheal intubation. Magnesium sulphate has a role in reducing the severity of sore throat. So we conclude that preoperative dexamethasone nebulisation reduces the incidence and severity of sore throat more effectively than magnesium sulphate nebulisation.

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