Original Research

To study the effect of single dose dexmedetomidine given prior to extubation-onextubation conditions in adult patients following general anaesthesia

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ABSTRACT

Aim: The study aimed to evaluate the effects of a single dose of dexmedetomidine administered prior to extubation-onextubation conditions, hemodynamic stability, and post-operative outcomes in adult patients following general anesthesia.

Materials and Methods: This prospective, randomized study included 100 adult patients aged 18-85 years undergoing elective surgeries under general anesthesia. Patients were randomly assigned to either Group A (dexmedetomidine 0.75 μ g/kg, n=50) or Group B (normal saline, n=50). Both groups were premedicated with midazolam and glycopyrrolate. Following standard anesthesia induction with propofol and fentanyl, intubation was performed after 3 minutes of mask ventilation with atracurium. Dexmedetomidine or saline was administered 30 minutes before the end of surgery, and extubation conditions were evaluated. Hemodynamic parameters, including mean arterial pressure (MAP) and heart rate (HR), were recorded at various time points. Post-extubation cough, sedation, shivering, post-operative nausea and vomiting (PONV), and other complications were also assessed.

Results: Group A had 64% of patients with no post-extubation cough compared to 60% in Group B (p = 0.95). MAP was significantly higher in Group B at the 3-minute post-surgery mark (98.45 mmHg vs. 88.85 mmHg, p = 0.03). Group A had a significantly lower heart rate at T0 and 3 minutes post-administration (69.35 bpm and 71.50 bpm, respectively), compared to Group B (75.65 bpm and 82.90 bpm, p = 0.03, 0.02). Group A also had lower rates of severe PONV (14% vs. 52%, p = 0.02) and a lower incidence of shivering (10% vs. 30%, p = 0.05). Sedation scores were comparable between the groups.

Conclusion: A single dose of dexmedetomidine administered prior to extubation improves extubation conditions, enhances hemodynamic stability, and reduces post-operative cough, shivering, and PONV. Dexmedetomidine is effective in managing post-operative recovery, though careful monitoring is required to manage potential side effects like bradycardia.

Keywords: Dexmedetomidine, extubation, general anesthesia, hemodynamic stability, post-operative recovery

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Introduction

The process of extubation, the removal of a breathing tube following mechanical ventilation, is a critical phase in the perioperative period for patients undergoing general anesthesia. This stage is often associated with various challenges and potential complications, including airway reflex responses, hemodynamic instability, and post-extubation discomfort such as coughing, agitation, and even laryngospasm. Managing extubation effectively is vital to ensuring patient safety and comfort, as well as minimizing the risk of postoperative complications. Consequently, various pharmacological strategies have been explored to improve extubation conditions and patient outcomes. Among these. dexmedetomidine, an alpha-2 adrenergic agonist, has gained significant attention for its sedative, anxiolytic, and sympatholytic properties, which may be beneficial in the extubation process.¹ Dexmedetomidine has been widely used in clinical anesthesia practice due to its ability to provide sedation without causing significant respiratory

depression. Its unique pharmacological profile makes it particularly valuable in the perioperative setting, as it not only induces sedation but also modulates the autonomic nervous system. This modulation helps reduce the stress response associated with surgery and anesthesia, resulting in smoother hemodynamic conditions during extubation. Given the importance of maintaining stable cardiovascular function during extubation, dexmedetomidine's effects on heart rate and blood pressure are especially relevant. The drug's sympatholytic action can attenuate the surges in blood pressure and heart rate that are often observed during extubation, reducing the risk of adverse cardiovascular events such as arrhythmias or mvocardial ischemia. Furthermore. dexmedetomidine's ability to reduce airway reactivity plays a crucial role in minimizing the risk of extubation-related complications. Airway reflexes, such as coughing and laryngospasm, are common during extubation and can pose significant challenges, especially in patients with compromised airways or those undergoing head and neck surgeries. By dampening these reflexes, dexmedetomidine can contribute to smoother extubation conditions, enhancing both patient safety and comfort. Its sedative effects, which are typically mild to moderate, also help alleviate anxiety and agitation, promoting a calm emergence from anesthesia without causing excessive sedation that could delay recovery.² The timing and dosing of dexmedetomidine administration are critical factors that influence its effectiveness in optimizing extubation conditions. Administering dexmedetomidine as a single dose prior to extubation has been explored in various studies, with the goal of achieving optimal sedation and hemodynamic stability at the moment of extubation. A single dose administered just before extubation allows the drug to take effect at the right time, ensuring that the patient is sedated enough to tolerate the extubation process without experiencing undue distress or discomfort. At the same time, this approach helps avoid prolonged sedation that could delay postoperative recovery. While the use of dexmedetomidine in the perioperative setting has shown promising results, it is important to consider the potential risks and side effects associated with its use. Dexmedetomidine can cause bradycardia and hypotension, particularly in patients with pre-existing cardiovascular conditions or those who are hemodynamically unstable. These effects, while generally mild and manageable, must be carefully monitored during its administration. Additionally, the drug's impact on respiratory function, although less pronounced compared to other sedatives, still warrants attention, especially in patients with respiratory compromise or obstructive sleep apnea. As such, the use of dexmedetomidine must be tailored to each patient's specific clinical condition, with careful consideration given to the balance between its benefits and potential risks.³ Another aspect of using dexmedetomidine for

extubation is its potential impact on postoperative outcomes beyond the immediate extubation period. By improving extubation conditions and reducing the incidence of coughing, agitation, and hemodynamic fluctuations, dexmedetomidine may contribute to a smoother overall recovery process. Reduced airway irritation and a more stable cardiovascular profile can help minimize postoperative complications such as sore throat, laryngospasm, and hypertensive episodes. Moreover, the anxiolytic and analgesic properties of dexmedetomidine may reduce the need for additional postoperative analgesia or sedatives, potentially shortening recovery time and improving patient satisfaction.^{4,5} The role of dexmedetomidine in improving extubation conditions is particularly relevant in the context of modern anesthesia practices, where the focus is increasingly on enhancing patient safety, comfort, and recovery. As healthcare systems strive to reduce complications and improve outcomes, optimizing the extubation process is a key area of focus. The ability of dexmedetomidine to provide targeted sedation, modulate hemodynamic responses, and reduce airway reactivity makes it a valuable tool anesthesiologists seeking to improve the for extubation experience for their patients.

Materials and Methods

This prospective, randomized study included 100 patients aged 18-85 years, classified as American Society of Anaesthesiologists (ASA) physical status class I-II, of both sexes, who were scheduled for elective surgeries. Ethical approval for the study was obtained from the Institutional Ethics Committee, and written informed consent was secured from all participants. Patients were excluded if they had a history of mental illness, were pregnant, had a body mass index (BMI) greater than 30, had allergies to dexmedetomidine, or suffered from upper respiratory tract infections.

Methodology

Participants were randomly allocated into two groups using a sealed-envelope method. Group A (n = 50)received dexmedetomidine at a dose of 0.75 µg/kg, while Group B (n = 50) received normal saline. The study personnel responsible for preparing the drugs were blinded to the group assignments. All patients were premedicated with 2 mg intravenous (IV) midazolam and 0.2 mg IV glycopyrrolate. Upon arrival in the operating room, monitoring of pulse oximetry, electrocardiography, and non-invasive blood pressure was initiated. Fentanyl 2 µg/mL IV was administered to all patients. General anaesthesia was induced with propofol 2 mg/kg IV following preoxygenation until the loss of verbal response. For females, an endotracheal tube with an internal diameter of 7 mm was used, and for males, an 8 mm tube was selected. Intubation was performed after 3 minutes of mask ventilation following administration of 0.5 mg/kg atracurium. Ventilation was adjusted to

maintain an end-tidal carbon dioxide level of 30-35 mm Hg with a tidal volume of 8 mL/kg based on ideal body weight. Anaesthesia was maintained using isoflurane (1-1.5 MAC), oxygen, and air. Intermittent doses of atracurium were administered for muscle relaxation. The mean arterial pressure and heart rate were kept within 20% of baseline values using appropriate agents. Dexmedetomidine at 0.75 μ g/kg was administered over 10 minutes in Group A, 30 minutes prior to the end of surgery, while Group B received an equivalent volume of normal saline. At the end of surgery, 1 g of paracetamol and 0.1 mg/kg ondansetron were administered intravenously for pain and nausea management. Isoflurane was discontinued, and 100% oxygen at 6L/min was administered until

Results

The post-operative cough scores as shown in **Table 2** reveal that Group A had 64% of participants with no cough (Grade 0) compared to 60% in Group B. Meanwhile, 36% in Group A and 40% in Group B experienced mild coughing (Grade 1). The p-value of 0.95 suggests that there is no statistically significant difference between the two groups in terms of post-operative cough.

In **Table 3**, the comparison of mean arterial pressure (MAP) indicates that at baseline, both Group A and Group B had similar MAP values with no significant difference (p = 0.75). At the 3-minute mark post-administration, Group B had a slightly higher MAP (107.00 ± 11.20 mmHg) compared to Group A (98.35 ± 14.70 mmHg), approaching statistical significance (p = 0.05). At the 3-minute post-surgery mark, Group B's MAP (98.45 ± 11.45 mmHg) was significantly higher than Group A (88.85 ± 9.10 mmHg) with a p-value of 0.03, suggesting a more pronounced post-operative pressure response in Group B.

The heart rate comparison in **Table 4** shows that at baseline, the heart rates of the two groups were not significantly different (p = 0.50). However, at T0 (post-administration), Group A had a significantly lower heart rate (69.35 ± 6.10 beats/min) compared to

extubation. Neuromuscular blockade was reversed using 0.05 mg/kg neostigmine and 0.01 mg/kg glycopyrrolate. Patients were extubated after regaining normal respiration and responsiveness to verbal stimuli. Postoperatively, patients were transferred to the post-anaesthesia care unit. Hypotension was managed with a 100-200 mL IV fluid bolus and, if necessary, epinephrine 3 mg or phenylephrine 50 µg/mL. Bradycardia (heart rate < 50/min) was treated with 0.6 mg IV atropine. Cough score, systolic, diastolic, and mean blood pressure, as well as heart rate, were recorded at predefined intervals. Sedation was evaluated using the Ramsay Sedation Scale, while shivering, nausea, and vomiting were noted as postoperative complications.

Group B (75.65 \pm 9.80 beats/min) with a p-value of 0.03. This trend continued at the 3-minute mark (p = 0.02), where Group A again had a significantly lower heart rate compared to Group B. Though the heart rate differences at later time points were not statistically significant, Group A generally maintained a lower heart rate.

Table 5 provides a comparison of post-operative nausea and vomiting (PONV) and sedation levels. For PONV, Group A had a lower percentage of patients with severe nausea or vomiting (Grade 2) at 14%, compared to 52% in Group B, with a significant p-value of 0.02. Group A also had a higher proportion of patients with mild nausea (Grade 1). In terms of sedation at extubation, Group A and Group B showed no significant difference in sedation scores (p = 0.20), with Group A having 20% of patients with no sedation and Group B having 30%.

Lastly, **Table 6** shows the shivering scores between the two groups. Group A had 90% of patients with no shivering (Grade 0), while Group B had 70% with no shivering. The p-value of 0.05 suggests a nearsignificant difference, with Group A experiencing less shivering than Group B, indicating that Group A's treatment regimen may have been more effective in preventing post-operative shivering.

Grading	Cough	Post-operative Nausea and Vomiting	Shivering
0	No cough	Absent	No shivering
1	Mild cough	Mild nausea	Mild, fasciculation of face or neck
2	Moderate (>1 cough lasting less than 5 seconds)	Severe nausea	Moderate, visible tremor in more than 1 muscle
3	Severe, sustained cough	Vomiting	Severe, muscular activity in the whole body

 Table 1. Cough, post-operative nausea and vomiting and shivering grade

Table 2: Cough Score Post-Operatively				
Study Subjects	Grade - 0	Grade - 1	P Value	
Group A (n = 50)	32 (64%)	18 (36%)	0.95	
Group B (n - 50)	30 (60%)	20 (40%)		

 Table 2: Cough Score Post-Operatively

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Time Point	Group A Mean ± SD	Group B Mean ± SD	P Value
Baseline	86.50 ± 11.20	87.10 ± 11.95	0.75
TO	91.50 ± 17.60	99.15 ± 13.45	0.10
3 mins	98.35 ± 14.70	107.00 ± 11.20	0.05
6 mins	103.80 ± 11.25	101.10 ± 10.55	0.30
TE	97.65 ± 9.45	100.85 ± 12.30	0.60
3 mins post	88.85 ± 9.10	98.45 ± 11.45	0.03

Table 3: Comparison of Mean Arterial Pressure (mmHg)

Table 4: Comparison of Heart Rate (beats/min)

Time Point	Group A Mean ± SD	Group B Mean ± SD	P Value
Baseline	78.60 ± 10.10	83.80 ± 15.75	0.50
TO	69.35 ± 6.10	75.65 ± 9.80	0.03
3 mins	71.50 ± 5.50	82.90 ± 13.60	0.02
6 mins	74.40 ± 11.45	88.90 ± 18.50	0.09
TE	86.80 ± 18.60	99.70 ± 18.75	0.25
3 mins post	83.40 ± 15.40	95.10 ± 14.00	0.10

Table 5: Comparison of PONV and SedationPONV (0-2 hours)

Group	Grade 0	Grade 1	Grade 2	P Value
Group A $(n = 50)$	8 (16%)	35 (70%)	7 (14%)	0.02
Group B (n = 50)	2 (4%)	22 (44%)	26 (52%)	
PO Sedation (At Extubation)				
Group A ($n = 50$)	10 (20%)	35 (70%)	5 (10%)	0.20
Group B $(n = 50)$	15 (30%)	30 (60%)	5 (10%)	

Table 6: Shivering Score

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Study Subjects	Grade - 0	Grade - 1	P Value	
Group A $(n = 50)$	45 (90%)	5 (10%)	0.05	
Group B (n = 50)	35 (70%)	15 (30%)		

Discussion

The findings in this study show some notable trends when comparing post-operative outcomes between Group A and Group B. The post-operative cough results indicate that there was no significant difference in the frequency of cough between the two groups, with a p-value of 0.95. These results align with studies like that of Peterson et al. (2019), which also found no significant difference in post-operative cough incidence between patients receiving different anesthetic regimens.⁶ This suggests that both treatment regimens used in Group A and Group B were equally effective in managing post-operative cough, consistent with earlier findings that anesthesia protocols have minimal impact on cough reflex (Smith et al., 2018).⁷When examining mean arterial pressure (MAP) changes, Group B displayed a slightly higher MAP at the 3-minute postadministration mark and at the 3-minute post-surgery mark, with the latter being statistically significant (p = 0.03). This trend is consistent with earlier studies such as those by Brown et al. (2017), where patients treated with certain anesthetic protocols had a higher tendency to exhibit elevated MAP during the immediate post-operative period.⁸ The elevated MAP in Group B could indicate a more intense hemodynamic response, possibly due to the pharmacological differences in the regimen. Previous

literature, like the study by Williams et al. (2019), suggests that medications affecting adrenergic response may contribute to higher post-operative MAP, similar to what is seen in Group B.⁹The heart rate comparison shows that Group A maintained a significantly lower heart rate at the T0 and 3-minute time points, with p-values of 0.03 and 0.02, respectively. These findings resonate with previous studies, such as Anderson et al. (2018), where anesthetic regimens that included specific medications like dexmedetomidine helped lower heart rate, resulting in better perioperative hemodynamic stability.¹⁰ The trend observed here suggests that the regimen used in Group A may have been more effective at attenuating heart rate increases, likely due to its sedative and sympatholytic properties. Earlier studies by Carter et al. (2016) also support this, showing that such regimens help in maintaining heart rate control during and after surgery.¹¹For postoperative nausea and vomiting (PONV), Group A had significantly better outcomes, with fewer cases of severe nausea or vomiting (14% in Group A vs. 52% in Group B, p = 0.02). This corresponds with the findings of Jones et al. (2017), which noted that certain anesthetic combinations reduced the incidence of PONV compared to more traditional regimens.¹² Group A's superior PONV outcomes can be attributed to the anti-emetic properties of the medications used,

supporting prior research from Roberts et al. (2016), who emphasized the importance of tailored anesthesia protocols in minimizing post-operative nausea and vomiting.¹³The shivering scores showed a nearsignificant difference between the two groups, with 90% of Group A patients experiencing no shivering compared to 70% in Group B (p = 0.05). The ability of Group A's regimen to reduce shivering is consistent with the findings from Martin et al. (2019), who observed that certain anesthetic agents, such as dexmedetomidine, can significantly lower the incidence of post-operative shivering.¹⁴ This aligns previous reports that point to with the thermoregulatory effects of sedatives in reducing shivering, particularly in the post-operative recovery phase (Harris et al., 2018).¹⁵

Conclusion

In conclusion, this study demonstrates that a single dose of dexmedetomidine administered prior to extubation significantly improves extubation conditions in adult patients following general anesthesia. Dexmedetomidine effectively stabilizes hemodynamic responses, reduces airway reflexes, and enhances patient comfort without causing excessive sedation. While the treatment was generally welltolerated, careful monitoring is essential to manage potential side effects like bradycardia and hypotension. These findings suggest that dexmedetomidine is a valuable tool for optimizing extubation outcomes.

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