



COMPARING TWO DOSES OF DEXMEDETOMIDINE TO ATTENUATE EXTUBATION RESPONSE IN ELECTIVE LAPAROSCOPIC ABDOMINAL SURGERY: A PROSPECTIVE RANDOMIZED DOUBLE-BLIND CONTROLLED TRIAL

Dr. Komal Kishanrao Mutkhede^{1*}, Dr. Megha Tajne², Dr Tanisha Uikey³, Dr Avishkar Gaikwad⁴, Dr. Pratiksha Gosavi⁵, Dr. Saikat Dhar⁶

^{1*}Junior Resident, Department of Anesthesiology, Government Medical College, Nagpur, Maharashtra, India.

²Professor & HOD, Department of Anesthesiology, Government Medical College Nagpur, Maharashtra, India.

³Undergraduate Student, Datta Meghe Institute of Higher Education and Research, Nagpur, Maharashtra, India.

⁴Undergraduate Student, Datta Meghe Institute of Higher Education and Research Nagpur, Maharashtra, India.

⁵Junior Resident, Department of Anesthesiology, Government Medical College Nagpur, Maharashtra, India.

⁶Junior Resident, Department of Anaesthesiology, Government Medical College Nagpur, Maharashtra, India.

***Corresponding Author:** Dr. Komal Kishanrao Mutkhede

Junior Resident, Department of Anesthesiology, Government Medical College, Nagpur, Maharashtra, India.

ABSTRACT

Background

Extubation during general anesthesia is associated with significant sympathetic stimulation, causing transient but potentially harmful hemodynamic and airway reflex responses. Dexmedetomidine, an α_2 -adrenoceptor agonist, is increasingly used to attenuate these stress responses, but the optimal dose for elective laparoscopic abdominal surgery remains under-investigated.

Objective

To compare the efficacy and safety of two intravenous dexmedetomidine doses 0.75 $\mu\text{g/kg}$ and 0.5 $\mu\text{g/kg}$ —in suppressing extubation-related hemodynamic and airway responses and sedation score in patients undergoing elective laparoscopic abdominal surgery.

Methods: In a prospective, double-blinded, randomized controlled trial, 74 ASA I-II adult patients (18–60 years) were assigned to receive either 0.75 $\mu\text{g/kg}$ (Group A) or 0.5 $\mu\text{g/kg}$ (Group B) of dexmedetomidine intravenous infusion over 10 minutes, 15 minutes prior to extubation. Hemodynamic parameters (heart rate, systolic/diastolic blood pressure, mean arterial pressure), oxygen saturation, extubation quality, and sedation scores were recorded pre-extubation and up to two hours post-extubation.

Results: The groups were comparable demographically. Group A demonstrated significantly better attenuation of heart rate and blood pressure responses post-extubation ($p < 0.001$).

Smooth extubation without coughing (Extubation Quality Scale score 1) occurred in 59.5% of Group A versus 8.1% in Group B ($p<0.01$). Group A patients showed higher Ramsay Sedation Scores immediately post-extubation, indicating deeper but clinically safe sedation. Oxygen saturation remained above 98% with no respiratory compromise in either group. No episodes of bradycardia or hypotension requiring intervention occurred.

Conclusion: Dexmedetomidine at 0.75 $\mu\text{g/kg}$ provides superior hemodynamic stability and smoother extubation compared to 0.5 $\mu\text{g/kg}$ dose in elective laparoscopic abdominal surgery. This dose optimally balances efficacy and safety, making it preferable for clinical use.

Keywords: Dexmedetomidine, extubation response, laparoscopic surgery, hemodynamic stability

INTRODUCTION

Laparoscopic abdominal surgeries have revolutionized surgical care by minimizing tissue trauma, reducing postoperative pain, and shortening hospital stays¹. Despite the benefits, these procedures involve pneumoperitoneum with CO₂ insufflation, which elevates intra-abdominal pressure and provokes significant sympathetic activation². This neurohumoral response, combined with airway manipulation during extubation, often results in marked cardiovascular responses including tachycardia and hypertension³.

Tracheal extubation itself is a critical and potentially hazardous phase in anesthesia, especially in patients with cardiovascular comorbidities⁴. The laryngeal and pharyngeal stimulation during extubation leads to tachycardia, hypertension, coughing, bucking, and at times laryngospasm or bronchospasm⁵. Such hemodynamic perturbations elevate myocardial oxygen demand and can precipitate ischemia or arrhythmias^{6,7}.

Numerous agents have been explored to mitigate these responses: opioids, beta blockers, calcium channel blockers, local anesthetics, and alpha-2 agonists^{1,8}. Dexmedetomidine, a highly selective α_2 -adrenoceptor agonist, acts centrally to decrease norepinephrine release and sympathetic outflow, providing sedation, analgesia, sympatholysis, and anxiolysis without significant respiratory depression⁹⁻¹¹. Its efficacy in attenuating extubation stress responses has been demonstrated across a range of doses (0.25 to 1.0 $\mu\text{g/kg}$), but an optimal dose balancing hemodynamic stability and sedation remains uncertain, especially in laparoscopic abdominal surgery¹²⁻¹⁴.

This study aims to directly compare two commonly used dexmedetomidine doses—0.75 $\mu\text{g/kg}$ and 0.5 $\mu\text{g/kg}$ —administered as infusions before extubation, to evaluate their relative effectiveness in attenuating extubation-related responses in patients undergoing elective laparoscopic abdominal surgery.

MATERIALS AND METHODS

This prospective, randomized, double-blind controlled trial was conducted at a tertiary care teaching hospital between January 2023 and July 2025. Institutional Ethics Committee approval and written informed consent were obtained.

Inclusion criteria

Age 18–60 years, ASA physical status I or II, Scheduled for elective laparoscopic abdominal surgery under general anesthesia, Provided informed consent.

Exclusion criteria

Refusal to participate, Reactive airway disease, Significant cardiovascular or respiratory comorbidities, Pregnant or lactating women.

Study procedure

Seventy-four eligible patients were allocated equally into two groups by computer-generated block randomization with allocation concealment. Both patients and investigators involved in data collection were blinded to group assignment, with dexmedetomidine doses prepared by an

independent anesthesiologist.

Group A Dexmedetomidine 0.75 µg/kg diluted to 10 ml, infused over 10 minutes starting 15 minutes before planned extubation.

Group B Dexmedetomidine 0.5 µg/kg, similarly diluted and infused.

Standard anesthesia induction and maintenance included Premedication with glycopyrrolate 0.2 mg, ondansetron 4 mg, pantoprazole 40 mg iv. Induction with fentanyl (2 µg/kg), propofol (2–2.5 mg/kg), suxamethonium (2 mg/kg) to facilitate intubation. Maintenance with isoflurane (0.5–1.5%) in 60% N₂O and 40% O₂, supplemented with intermittent vecuronium. End-tidal CO₂ maintained between 30–35 mmHg. Analgesia with intravenous diclofenac 75 mg and paracetamol 1 g infusion intraoperatively. Dexmedetomidine infusion started 20 minutes prior to extubation as per group. Neuromuscular blockade reversed once TOF ratio >0.9 using neostigmine and glycopyrrolate. Extubation performed within 10 minutes after dexmedetomidine infusion.

Parameters were recorded at baseline and multiple time points including immediately pre-extubation and post-extubation at 1, 2, 3, 4, 5, 15, 30, 45, 60, 75, 90, 105, and 120 minutes: Heart rate (HR), Systolic blood pressure (SBP), Diastolic blood pressure (DBP), Mean arterial pressure (MAP), Oxygen saturation (SpO₂), Extubation quality scored by a 5-point scale (1 = no coughing, 5 = poor/extreme responses), Sedation assessed using Ramsay Sedation Score (RSS).

Statistical Analysis

Data were analyzed with STATA 10.1. Continuous variables were tested for normality; expressed as mean ± SD. Independent t-test compared means between groups, and chi-square or Fisher's exact tests were used for categorical data. P < 0.05 was considered statistically significant.

RESULTS

Patient Demographics

Demographic characteristics were comparable with no statistically significant differences:

Parameter	Group A (0.75 µg/kg)	Group B (0.5 µg/kg)	P-value
Age (years)	43.74 ± 10.39	40.73 ± 9.12	0.19
Weight (kg)	53.78 ± 5.86	54.16 ± 4.73	0.76
Gender (M:F)	21:16	19:18	0.81
ASA Grade I/II	26/11	28/9	0.79

Table 1: Demographic Characteristics of Study Groups

Extubation Quality Assessment

Quality Score	Description	Group A (%)	Group B (%)	P-value
1	No coughing	59.5	8.1	<0.01
2	Minimal coughing (1-2 times)	40.5	73.0	<0.01
3	Moderate coughing (3-4 times)	0	18.9	<0.01
4	Severe coughing (5-10 times)	0	0	-
5	Poor extubation (>10 times)	0	0	-

Table 2: Extubation Quality Distribution Between Groups

Group A showed significantly smoother extubation with fewer coughing episodes.

Hemodynamic Parameters Analysis

Parameter	Time Point	Group A	Group B	P-value
-----------	------------	---------	---------	---------

HR (bpm)	Pre-extubation	78.2 ± 8.4	82.1 ± 9.2	0.06
	1 min post	82.5 ± 8.9	95.3 ± 11.2	<0.001
	5 min post	80.1 ± 7.6	91.7 ± 10.8	<0.001
	15 min post	78.9 ± 8.1	87.2 ± 9.4	<0.001
SBP (mmHg)	Pre-extubation	118.3 ± 12.1	120.7 ± 11.8	0.38
	1 min post	125.2 ± 13.4	138.9 ± 15.6	<0.001
	5 min post	122.8 ± 12.9	135.1 ± 14.2	<0.001
	15 min post	120.5 ± 11.7	131.8 ± 13.5	<0.001
DBP (mmHg)	Pre-extubation	72.1 ± 8.9	74.3 ± 9.2	0.28
	1 min post	76.8 ± 9.4	87.2 ± 11.3	<0.001
	5 min post	75.2 ± 8.7	84.6 ± 10.8	<0.001
	15 min post	73.9 ± 8.3	82.1 ± 9.9	<0.001
MAP (mmHg)	Pre-extubation	87.5 ± 9.8	89.8 ± 10.2	0.31
	1 min post	92.9 ± 10.5	104.4 ± 12.8	<0.001
	5 min post	91.1 ± 9.9	101.4 ± 11.9	<0.001
	15 min post	89.4 ± 9.2	98.7 ± 11.1	<0.001

Table 3: Hemodynamic Response Comparison at Key Time Points

Heart Rate: Group A showed significantly lower HR at all intervals up to 15 minutes post-extubation ($p < 0.001$). Post 30 minutes, HR values approximated each other, indicating stabilization.

Systolic Blood Pressure: Lower in Group A during the initial 45 minutes post-extubation, with significant differences at all early time points ($p < 0.001$).

Diastolic Blood Pressure: Significantly attenuated in Group A compared to Group B up to 1 hour post-extubation ($p < 0.001$).

Mean Arterial Pressure: Consistently lower in Group A post-extubation ($p < 0.001$), with differences diminishing after 90 minutes.

Oxygen Saturation: Both groups maintained $SpO_2 > 98\%$ throughout, with no significant intergroup differences.

Sedation

Ramsay Sedation Scores indicated deeper sedation in Group A immediately post-extubation:

Score	Description	Group A (%)	Group B (%)	P-value
1	Anxious, agitated	5.4	64.9	<0.01
2	Cooperative, oriented	56.8	35.1	NS
3	Responds only to commands	37.8	0	<0.01

Table 4: Ramsay Sedation Score Distribution Post-Extubation

No excessive sedation or respiratory depression observed.

No episodes of clinically significant bradycardia, hypotension, respiratory depression, laryngospasm, or other adverse events requiring intervention were reported in either group.

DISCUSSION

This study confirms that dexmedetomidine at a 0.75 µg/kg dose provides superior attenuation of the extubation-associated sympathetic surge compared to the 0.5 µg/kg dose in patients undergoing elective laparoscopic abdominal surgery. Patients receiving the higher dose had significantly better hemodynamic stability, reduced cough and airway reflexes, and smoother extubation quality,

without prolongation of recovery or oxygen desaturation.

The findings are consistent with previous literature that highlights the dose-dependent sympatholytic, sedative, and analgesic effects of dexmedetomidine¹⁵⁻¹⁷. The drug's action on central α_2 -adrenoceptors reduces norepinephrine release, blunting tachycardia and hypertension during stress^{18,19}. Its sedative profile, resembling natural sleep, allows cooperative sedation without compromising respiratory drive^{20,21}.

Our results align with multiple studies investigating optimal dexmedetomidine dosing for extubation. Jamal et al. (2018) demonstrated that 0.75 $\mu\text{g/kg}$ offered optimal stability with minimal side effects compared to 0.5 $\mu\text{g/kg}$ and 1.0 $\mu\text{g/kg}$ doses⁴. Similarly, Mohamed Ali et al. (2021) found 0.75 $\mu\text{g/kg}$ significantly better at stabilizing all hemodynamic parameters with comparable sedation levels²².

Bhardwaj et al. (2021) studied controlled hypertensive patients and concluded that 0.75 $\mu\text{g/kg}$ was optimal for extubation, facilitating smooth extubation while maintaining hemodynamic stability without excessive sedation²³. Manickam et al. (2021) revealed that 0.75 $\mu\text{g/kg}$ yielded the best combination of cough suppression and hemodynamic stability in cholecystectomy patients²⁴. Recent studies by Jain et al. (2022) and Ayyanagouda et al. (2023) further support our findings, demonstrating superior hemodynamic control with 0.75 $\mu\text{g/kg}$ compared to lower doses^{25,26}. The consistency across diverse patient populations and surgical procedures strengthens the evidence for 0.75 $\mu\text{g/kg}$ as the optimal dose.

Mechanism and Clinical Implications

The superior efficacy of 0.75 $\mu\text{g/kg}$ can be attributed to more complete α_2 -adrenoceptor occupancy, leading to enhanced sympatholysis^{27,28}. This dose achieves better balance between hemodynamic control and sedation without triggering adverse effects often observed at higher doses (≥ 1 $\mu\text{g/kg}$), such as profound bradycardia or delayed emergence^{2,29}.

The higher number of patients with no coughing in Group A emphasizes enhanced airway reflex suppression at this dose, which is particularly beneficial in laparoscopic surgery where pneumoperitoneum already stresses the cardiovascular system^{30,31}.

LIMITATION

Limitations of the study include restriction to ASA I-II adults and elective laparoscopic procedures, which may limit generalizability to other populations or emergency surgeries. Further investigations in elderly or high-risk cardiac patients could expand clinical understanding.

CONCLUSION

For elective laparoscopic abdominal surgery, dexmedetomidine administered at 0.75 $\mu\text{g/kg}$ intravenously over 10 minutes, approximately 15 minutes before extubation, effectively attenuates the hemodynamic and airway reflex responses compared to 0.5 $\mu\text{g/kg}$. It provides smoother extubation, better cardiovascular stability, adequate sedation, and maintains respiratory function without significant adverse events. This dose is recommended as optimal for anesthetic management in similar patient populations and surgical settings.

REFERENCES

1. Rao S, Somasekharam P, Dinesh K, Ravi M. Effect of bolus dose of dexmedetomidine on hemodynamic responses and airway reflexes during tracheal extubation. *World J Pharm Pharmacol Sci*. 2015;04:731-40.
2. Liu Z, et al. Optimal dexmedetomidine dose to suppress airway and cardiovascular responses to extubation. *J Perioper Sci*. 2023.
3. Turan G, Ozgultekin A, Turan C, Dincer E, Yuksel G. Advantageous effects of dexmedetomidine on haemodynamic and recovery responses during extubation for intracranial surgery. *Eur J Anaesthesiol*. 2008;25:816-820.

4. Jamal MK, Ahmad S, Ahmad F. A comparative study of three dexmedetomidine doses for extubation response. *J Med Sci Clin Res.* 2018;6(3):1521.
5. Bindu B, Pasupuleti S, Gowd UP, Gorre V, Murthy RR, Laxmi MB. To study the effect of dexmedetomidine on hemodynamic and recovery responses during tracheal extubation. *J Anaesthesiol Clin Pharmacol.* 2013;29:162-67.
6. Agarwal M, Malliwal A. Effect of dexmedetomidine on hemodynamic responses during tracheal extubation. *Ind J Clin Anaesth.* 2019;06:23-29.
7. Kotak N, Mamde R, Desai PM. Prospective randomised comparative trial of dexmedetomidine versus esmolol for attenuation of extubation response. *Med J DY Patil Vidyapeeth.* 2019;12:131-35.
8. Sharma A, Gupta M, Kapoor BB. Comparative evaluation of dexmedetomidine, fentanyl, magnesium sulphate and control group to attenuate pressor responses and airway reflexes to intubation during general anaesthesia. *J K Sci.* 2018;20:120-27.
9. Standing S, ed. *Gray's Anatomy: The Anatomical Basis of Clinical Practice.* 41st ed. London: Elsevier; 2016. p. 503-521, 688-698.
10. Miller RD, Eriksson LI, Fleisher LA, Wiener-Kronish JP, Cohen NH, Young WL. *Miller's Anesthesia.* 9th ed. Philadelphia: Elsevier; 2020. Chapter 56: Extubation and Emergence, p. 1495-1512.
11. Precedex (Dexmedetomidine) Package Insert. Abbott Park, IL: Abbott Laboratories; 2004.
12. Drummond JC, Dao AV, Roth DM, Cheng C, Atwater IB, Minokadeh A, et al. Effect of dexmedetomidine on cerebral blood flow velocity, cerebral metabolic rate and carbon dioxide response in normal humans. *Anesthesiology.* 2008;87:684-90.
13. Panzer O, Moitra V, Sladen RN. Pharmacology of sedative-analgesic agents: dexmedetomidine, remifentanyl, ketamine, volatile anesthetics, and the role of peripheral mu antagonists. *Critical care clinics.* 2009;25(3):451-69.
14. Dogru K, Arik T, Yildiz K, Bicer C, Madenoglu H, Boyaci A. The effectiveness of intramuscular dexmedetomidine on hemodynamic responses during tracheal intubation and anesthesia induction of hypertensive patients: a randomized, double- blind, placebo-controlled study. *Current therapeutic research.* 2007;68(5):292-302.
15. Fragen RJ, Fitzgerald PC. Effect of dexmedetomidine on the minimum alveolar concentration of sevoflurane in adults aged 55-70 years. *Journal of Clinical Anesthesia.* 1999;11:466-70.
16. Kanazi GE, Aouad MT, Jabbour-Khoury SI, Al-Jazzar MD, Alameddine MM, Al- Yaman R, et al. Effect of small dose dexmedetomidine or clonidine on the characteristics of bupivacaine spinal block. *Acta Anaesthesiol Scand.* 2005;50:222-7.
17. Khan ZP, Munday IT, Jones RM, Thornton C, Mant TG, Amin D. Effects of dexmedetomidine on isoflurane requirement in healthy volunteers: 1 Pharmacodynamics and pharmacokinetic interactions. *Br J Anaesth.* 1999;83:372-80.
18. Talke P, Tayefeh F, Sessler DI, Noursalehi M, Richardson C. Dexmedetomidine does not alter the sweating threshold but comparably and linearly decreases the vasoconstriction and shivering thresholds. *Anesthesiology.* 1997;87:835-41.
19. Shehabi Y, Ruettimann U, Adamson H, Innes R, Ickeringill M. Dexmedetomidine infusion for more than 24 hours in critically ill patients: sedative and cardiovascular effects. *Intensive Care Med.* 2004;30:2188-96.
20. Aho M, Erkola O, Kallio A, Scheinin H, Korttila K. Comparison of dexmedetomidine and midazolam sedation and antagonism of dexmedetomidine with atipamezole. *Journal of Clinical Anesthesia.* 1993;5:194-203.
21. Ebert TJ, Hall JE, Barney JA, Uhrich TD, Colino MD. The effects of increasing plasma concentrations of dexmedetomidine in humans. *Anesthesiology.* 2000;93:382- 94.
22. Mohamed Ali M, Sharma L, Patel D. Comparative study of two dexmedetomidine doses on extubation response. *Saudi J Anaesth.* 2021;15:89-95.
23. Bhardwaj V, Singha D, Pathania A, Chaudhary U, Chaudhary S. Comparing different doses of

- dexmedetomidine in attenuating extubation response in hypertensive patients undergoing laparoscopic cholecystectomy. *Bali J Anaesthesiol.* 2021;5:72-7.
24. Manickam V, Kumar P, Singh S, et al. Dose-dependent dexmedetomidine effects on extubation in cholecystectomy. *Int J Surg Anesth.* 2021;52:75-80.
 25. Jain D, et al. Hemodynamic and coughing response to extubation: Dose-response evaluation of dexmedetomidine. *Indian J Clin Anaesth.* 2022;9(1):10-6.
 26. Ayyanagouda B, Dhulkhed VK, Mudigere SS. Comparison of 0.5 µg/kg and 0.75 µg/kg dexmedetomidine in attenuating hemodynamic response during extubation. *Indian J Anaesth.* 2023;67:43-316.
 27. Luthra A, Prabhakar H, Rath GP. Alleviating stress response to tracheal extubation in neurosurgical patients: A comparative study of two infusion doses of dexmedetomidine. *J Neurosci Rural Pract.* 2017;8:S49-56.
 28. Singh R, et al. Comparison of dexmedetomidine 0.5 vs 1 mcg/kg for smooth extubation in neurosurgical cases. *Asian J Neurosurg.* 2020;15(2):332-8.
 29. Ranjan A, Kumar M. Comparison of two different doses of dexmedetomidine in attenuation of haemodynamic response during endotracheal extubation. *Int J Pharm Clin Res.* 2024;16(6):2487-90.
 30. Suresh R, Prakash S, Rao M, et al. Efficacy of dexmedetomidine 0.75 µg/kg for extubation hemodynamics. *Ann Anesth Res.* 2020;41(2):12-18.
 31. Kotak S, Phalgune DS. Comparison of two dexmedetomidine doses in elective surgery. *Indian J Anaesth Crit Care.* 2022;8(3):145-150.