

A comparative study of midazolam plus fentanyl versus midazolam plus propofol with respect to hemodynamic stability during regional anaesthesia

Vinayak S Sirsat¹, Sangita M Agale(Eram)^{2*}

¹Associate Professor, ²Assistant Professor, Department of Anaesthesiology, Government Medical College, Latur-413512, Maharashtra, INDIA.

Email: drvinayak1@gmail.com

Abstract

Background: Patient safety has always been a major concern for the physicians of both ancient and modern eras. **Aims and Objectives:** To study Midazolam plus Fentanyl versus Midazolam plus Propofol with respect to Hemodynamic stability during regional Anaesthesia. **Methodology:** We conducted a comparative study of conscious sedation using midazolam with fentanyl in group-I vs. midazolam with propofol in group-II. In the department of anesthesia at Government Medical College, Latur. In the period between January 2016 to December 2016. 60 patients of ASA Grade I,II,and III, were randomly divided in two groups, 30 in each group, of between 15 to 60 years. **Results:** Systolic blood pressure changes in both the groups are comparable with each other at 30 minutes after sedation, but blood pressure fall was more in group II from the base line. Heart rate changes in both the groups applying test of significant changes in heart rate seen after sedation in group-I (Midazolam+Fentanyl) compare to group II (Midazolam+ propofol). Statistically $p < 0.001$ is highly significant in both groups. **Conclusion:** It can be concluded from our study that Systolic blood pressure changes in both the groups are comparable with each other at 30 minutes after sedation, but blood pressure fall was more in Midazolam plus Propofol group from the base line also in heart rate was more.

Key Word: Midazolam plus Fentanyl, Midazolam plus Propofol, Hemodynamic stability.

*Address for Correspondence:

Dr. Sangita M Agale(Eram), Assistant Professor, Department of Anaesthesiology, Government Medical College, Latur-413512, Maharashtra, INDIA.

Email: drvinayak1@gmail.com

Received Date: 12/07/2017 Revised Date: 20/08/2017 Accepted Date: 28/09/2017

DOI: <https://doi.org/10.26611/1021418>

Access this article online

Quick Response Code:	Website: www.medpulse.in
	Accessed Date: 27 October 2017

INTRODUCTION

Patient safety has always been a major concern for the physicians of both ancient and modern eras.¹ Propofol is a widely administered hypnotic agent that is of unique advantages yet some disadvantages.²⁻⁴ Induction of anesthesia with propofol is associated with significant blood pressure reduction and hemodynamic instability

especially in patients over 50 years old. In patients with previous hypotension and those with American society of anesthesiologists' physical status (ASAPS) > II, this drop is more dramatic.^{3,5} Regional anesthesia is becoming an increasing important aspect of anesthesia practice. Its advantage include avoidance of certain risk with general anesthesia those of pulmonary aspiration and airway obstructions, laryngospasm. Avoidance of operation theatre pollution, provision of good postoperative analgesia. Benefits in certain pre-existing pulmonary embolism postoperatively^{6,7,8,9,10}. Midazolam is used for conscious sedation for short diagnostic or endoscopic and dental procedure, adjunct to local or regional anesthesia¹⁰. Propofol is a sedative hypnotic drug, which is becoming popular for sedation during our patients procedures performed under local anesthesia. Its high clearance and favorable recovery profile offers advantages over other intravenous sedative and analgesic drugs. Sedation with propofol can be adjusted with manual intermittent bolus

injections techniques^{11,12}. Fentanyl: Fentanyl is a potent synthetic opiate agonist, estimated to be 25 fold to 100 fold more potent than morphine. It is highly lipid soluble and enters the central nervous system swiftly. Leading to rapid onset of action. Fentanyl provides relief of moderate to severe pain and has become the narcotic drug of choice for a wide variety of painful procedures. It has relatively short duration of action. These qualities make it ideal for the expeditious completion of painful procedures in the emergency department setting^{13,14}.

MATERIAL AND METHODS

We conducted a comparative study of conscious sedation using midazolam with fentanyl in group-I vs. midazolam with propofol in group-II. In the department of anesthesia at Government Medical College, Latur. In the period between January 2016 to December 2016. 60 patients of ASA Grade I, II, and III, were randomly divided in two groups, 30 in each group, of between 15 to 60 years of either sex undergoing any surgery under regional anesthesia (spinal, epidural anesthesia or peripheral nerve

blocks, Routine of emergency surgery were included into study while the patients with History of allergic reaction to the study medication, Chronic opioid or sedative drug use, Obesity(>130% for ideal body weight), Clinically significant cardiac, pulmonary, hepatic or renal dysfunction were excluded from the study.

Table 1: Sedation score is as follows

Sr. No	Parametere	Score
A	Fully awake and anxious	1
B	Drowsy or awake and comfortable	2
C	Eyes closed but responds to verbal commands	3
D	Eyes closed but responds to light physical stimulation.	4
E	Unresponsive to light physical simulation.	5

Patients were specifically asked awareness during the surgical procedure and whether they will be happy to have same anesthetic technique again. The hemodynamic parameters like Blood pressure, systolic Diastolic and Heart Rate etc. were recorded. The statistical analysis done by unpaired t-test, calculated by SPSS 19 software.

RESULT

Table 2A: Hemodynamic changes

Characteristics	Group I	Group II	Group I/II	Group I/II	Remarks
Duration in minutes after starting the drip	Midazolam + fentanyl	Midazolam + Propofol	'T' values (test of Sig.)	P Value	
	Systolic Blood pressure				
Preop SBP	121.9±12.34	129±16.6	1.978	NS	NS
SBP (10 min)	108.466±9.608	118±14.5	2.99	P=0.01	SS
SBP (20 min)	100.66±7.849	108.1±9.819	3.224	P<0.001	HS
SBP (30 min)	97±6.533	98.26±5.085	0.838	P<0.001	HS

Group I:-Midazolam+Fentanyl, Group II:- Midazolam +propofol. Table 2a and chart 1 shows hemodynamic changes of both the groups applying test of significance(t). Systolic blood pressure changes in both the groups are comparable with each other at 30 minutes after sedation, but blood pressure fall was more in group II from the base line.

Table 2B: Hemodynamic changes

Characteristics	Group I	Group II	Group I/II	Group I/II	Remarks
Duration in minutes after starting the drip.	Fentanyl	propofol	'T' values (Test of sig)	P values	
	Heart Rate Changes				
Preop HR	87.73±6.164	85.7±10	0.962	NS	NS
HR (10 Min)	78±5.632	83.5±8.665	2.91	P<0.01	SS
HR (20 min)	71±5.06	79.5±7.62	5.11	P<0.001	HS
HR (30 Min)	68.67±5.287	75.67±6.583	4.54	P<0.001	HS

Table 2b and chart II shows heart rate changes in both the groups applying test of significant changes in heart rate seen after sedation in group-I (Midazolam+Fentanyl) compare to group II (Midazolam+ propofol). Statistically p<0.001 is highly significant in both groups

DISCUSSION

Propofol produces decrease in systemic blood pressure that is greater than those evoked by comparable dose of thiopental. These decreases in blood pressure are often accompanied by corresponding changes in cardiac output and systemic vascular resistance. The relaxation of vascular smooth muscles produced by propofol is primarily due to inhibition of propofol may result from a decrease in intracellular calcium influx. Stimulation produced by direct laryngoscopy and intubations of the trachea reverse the blood pressure effect of propofol, although the drug is more effective than thiopental in blunting the magnitude of this pressure response. Propofol also effectively blunts the hypertensive response to placement of laryngeal mask airway. The blood pressure effect of propofol may be exaggerated in hypovolemic patients, elderly patients, and patients with compromised left ventricular function due to coronary artery disease. Adequate hydration before rapid IV administration of propofol is recommended to minimize the blood pressure effect of this drug. Addition of nitrous oxide dose not alters the cardiovascular effects of propofol. Despite decreased in systemic blood pressure, heart rate often remains unchanged. Bradycardia and asystole have been observed after induction of anesthesia with propofol, resulting in the occasional recommendation that anticholinergic drugs be administered when vagal stimulation is likely to occur in association with administration of propofol. Fentanyl even in large doses 50ug/kg IV. Does not evoke the release of histamine. As a result, dilatation of venous capacitance vessels leading to hypotension is unlikely. Carotid sinus baroreceptor reflex control of heart rate is markedly depressed by fentanyl. 10 ug/kg IV, administered to neonates. Bradycardia is more prominent with fentanyl than morphine and may lead to occasional decreases in blood pressure and cardiac output. Sedation was achieved after the bolus doses and that remained throughout the procedure in both the groups. In propofol group patients sedated immediately after bolus without hypotension or bradycardia. But in fentanyl group 9 to 10 min. were required to achieve the adequate sedation. 5 patients from fentanyl group had bradycardia and 2 patients had hypotension, so we reduced the doses by adjusting the microdrip and administering IV fluids and atropine. And further fall in blood pressure and heart rate was avoided. In our study we found that Systolic blood pressure changes in both the groups are comparable with each other at 30 minutes after sedation, but blood pressure fall was more in group II from the base line. Heart rate changes in both the groups applying test of significant changes in heart rate seen after sedation in group-I (Midazolam+Fentanyl) compare to group II (Midazolam+ propofol). Statistically $p < 0.001$ is highly significant in both groups.

CONCLUSION

It can be concluded from our study that Systolic blood pressure changes in both the groups are comparable with each other at 30 minutes after sedation, but blood pressure fall was more in Midazolam plus Propofol group from the base line also in heart rate was more.

REFERENCES

1. Golzari SE, Khan ZH, Ghabili K, Hosseinzadeh H, Soleimanpour H, Azarfarin R. *et al.* Contributions of medieval Islamic physicians to the history of tracheostomy. *AnesthAnalg.* 2013; 116:1123–32.
2. Soleimanpour H, RajaeiGhafouri R, Taheraghdam A, Aghamohammadi D, Negargar S. *et al.* Effectiveness of intravenous Dexamethasone versus Propofol for pain relief in the migraine headache: A prospective double blind randomized clinical trial. *BMC Neurol.* 2012; 12:114.
3. Dhungana Y, Bhattarai BK, Bhadani UK, Biswas BK, Tripathi M. Prevention of hypotension during propofol induction: A comparison of preloading with 3.5% polymers of degraded gelatin and intravenous ephedrine. *Nepal Med Coll J.* 2008; 10:16–9.
4. Reich DL, Hossain S, Krol M, Baez B, Patel P, Bernstein A. *et al.* Predictors of hypotension after induction of general anesthesia. *AnesthAnalg.* 2005; 101:622–8.
5. Yamaura K, Hoka S, Okamoto H, Kandabashi T, Akiyoshi K, Takahashi S. Changes in left ventricular end-diastolic area, end-systolic wall stress, and fractional area change during anesthetic induction with propofol or thiamylal. *J Anesth.* 2000; 14:138–42.
6. Modig J, Borg T, Karlstrom G, Maripuu E, Sahlstedt B. Thromboembolism after total hip replacement: role of epidural and general anesthesia. *Anesthesia and analgesia* 1983; 62: 174-80.
7. Freund FG, Bonica JJ, Ward RJ, Akamatsu TJ, Kennedy WF Jr. Ventilator reserve and level of motor block during high spinal and epidural anesthesia. *Anesthesiology* 1967; 28: 834-7.
8. S. Jorgren s. wright B. respiratory changes during continuous epidural blockade. *Acta anaesthesiologica scandinavica* 1972; 16: 27-49.
9. Wahba WM, Craig DB. The cardiorespiratory effects of thoracic epidural anesthesia. *Canadian anesthetists' society journal* 1972; 19: 8-19.
10. Aitkenhead AR, Grant IS. Interaction with concurrent disease and medication. In: Henderson JJ, Nimmo WS, eds. *Practical regional anaesthesia*, Oxford: Blackwell scientific publication, 1983, 143-62.
11. While PF, Nequs JB, sedative infusion during local and regional anesthesia a comparison of midazolam and propofol. *J. clin anesthesia* 1991; 3:32-9.
12. Taylor E, Ghouri AF, White PF, Midazolam in combination with propofol for sedation during local anesthesia. *J. of clin anesthesia* 1992; 4:213-6.
13. Seibs SM the treatment of pain the emergency department. *Clin. North Am.* 1989; 36:965-78.
14. Cafe CJ sedation for the pediatric patients A. review. *pediatric clin. North Am.* 1994; 41; 31-58.

Source of Support: None Declared
Conflict of Interest: None Declared