

## A Prospective Study to evaluate the Hemodynamic Changes and Adverse reactions associated with Propofol and Etomidate during General Anesthesia

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### Abstract

**Background:** Among general anesthesia induction drugs, etomidate is the only imidazole, and it has the most favorable therapeutic index for single bolus administration. Propofol has been shown in clinical studies to be a safe, effective, hypnotic, and amnesic anesthetic agent at induction doses of 2-2.5 mg/kg and maintenance doses of approximately 9mg/kg per hour. **Aim:** To assess hemodynamic changes and complications occurring with Propofol and Etomidate during general anesthesia. **Methods:** A total of 100 subjects were enrolled in the present study and were broadly and randomly divided two study groups with 50 subjects in each group: Group A: Subjects who received 1% Propofol injection, and Group B: Subjects who received 0.3mg/kg of etomidate injection. Monitoring of the blood pressure, mean arterial pressure and heart rate was done throughout the surgery and until 10 minutes after induction. Recording of the pain during injection was done on a scale of 0 to 10 with 0 referring to no pain while 10 referring to maximum pain. **Results:** No significant difference was observed while comparing the mean arterial pressure and heart rate among subjects of both the study groups at different time intervals except for at the time of induction. Mean pain score was found to be significantly higher in group A in comparison to group B. **Conclusion:** Among patients with associated altered hemodynamic status, etomidate is an improved option. However; further studies are recommended.

**Keywords:** Etomidate, Hemodynamic, Propofol

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### Introduction

Among general anesthesia induction drugs, etomidate is the only imidazole, and it has the most favorable therapeutic index for single bolus administration. It also produces a unique toxicity among anesthetic drugs-- inhibition of adrenal steroid synthesis that far outlasts its hypnotic action and that may reduce survival of critically ill patients[1-3]. The major molecular targets mediating anesthetic effects of etomidate in the central nervous system are specific  $\gamma$ -aminobutyric acid type A receptor subtypes. Propofol has been shown in clinical studies to be a safe, effective, hypnotic, and amnesic anesthetic agent at induction doses of 2-2.5 mg/kg and maintenance doses of approximately 9mg/kg per hour[4-6]. Hence; under the light of abovementioned data, we planned the present study to assess hemodynamic changes and complications occurring with Propofol and Etomidate during general anesthesia.

### Methods

This was a prospective, cross-sectional study. The study was conducted at the Department of Anesthesia at Vardhman Institute of Medical Sciences, Pawapuri. The study was conducted over a period of 14 months from June 2020 to August 2021. The study was approved by the institutional research and ethical committee. An informed and written consent was taken from all the subjects before the commencement of the study.

The present study included assessment of hemodynamic changes and complications occurring with Propofol and Etomidate during general

anesthesia.

A total of 100 subjects were enrolled in the present study. Inclusion criteria for the present study included:

- Subjects within the age group of 20 to 60 years,
- Subjects with negative history of any other systemic illness,
- Subjects with negative history of any known drug allergy.

After meeting the inclusion criteria, all the 100 subjects were broadly and randomly divided two study groups with 50 subjects in each group:

Group A: Subjects who received 1% Propofol injection, and

Group B: Subjects who received 0.3mg/kg of etomidate injection.

Detailed demographic data of all the patients was obtained. Complete haematological and biochemical analysis of all the patients was carried out. In all the subjects, premedication was done with alprazolam tablets and ranitidine tablets. Recording of the baseline hemodynamic values was done in all the patients, after they entered operation theatre. Recording of the time of induction and patient's myoclonic activity was done. Monitoring of the blood pressure, mean arterial pressure and heart rate was done throughout the surgery and until 10 minutes after induction. Recording of the pain during injection was done on a scale of 0 to 10 with 0 referring to no pain while 10 referring to maximum pain. Recording of all the results was done in Microsoft excel sheet followed by analysis by SPSS software. Chi-square test was used for assessment of level of significance. P-value of less than 0.05 was taken as significant.

### Results

In the present study, a total of 100 subjects were enrolled and were broadly divided into two study groups with 50 patients in each group. Mean age of the patients of the group A and group B was 29.5 years and 30.1 years respectively. Mean weight of the patients of the

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group A and group B was 65.8 and 66.1 Kg respectively. There were 30 males and 20 females in the group A while there were 28 males and 22 females in the group B. [Table 1] shows the mean hemodynamic parameters at different time intervals. No significant difference was observed while comparing the mean arterial pressure

and heart rate among subjects of both the study groups at different time intervals except for at the time of induction. Mean pain score was found to be significantly higher in group A in comparison to group B.

Mean hemodynamic parameter	Group A	Group B	p- value	
Mean arterial pressure	Baseline	88.8	90.8	0.58
	Induction	79.5	87.25	0.00(Significant)
	At 10 minutes	95.8	96.4	0.44
Heart rate	Baseline	83.1	85.1	0.82
	Induction	99.1	97.5	0.46
	At 10 minutes	80.2	81.8	0.38

Parameter	Group A	Group B	p- value
Mean pain score	1.5	0.7	0.00

### Discussion

Few of the side effects that occur with Propofol are blood pressure drop, ventilation depression in a dose dependent manner and pain during injection.<sup>[4]</sup> Etomidate has a special property of its hemodynamic stability. It causes minimal respiratory depression and has cerebral protective actions[7-9]. In the present study, a total of 100 subjects were enrolled and were broadly divided into two study groups with 50 patients in each group. Mean age of the patients of the group A and group B was 29.5 years and 30.1 years respectively. Mean weight of the patients of the group A and group B was 65.8 and 66.1 Kg respectively. There were 30 males and 20 females in the group A while there were 28 males and 22 females in the group B. Mayer M et al compared the haemodynamic effects, the patients' sensations, signs of thrombophlebitis and postoperative nausea and vomiting (PONV) following injection of both drugs. Following premedication with 2 mg Lorazepam p.o. in 50 patients per group, anaesthesia was induced with either 0.51 mg etomidate in lipid emulsion or 3.04 mg propofol per kg bw. No opioid or benzodiazepine was given i.v. before induction. After injection of the tested drug, the cannula was removed. Changes in blood pressure and heart rate were recorded as well as signs of discomfort during and after injection (pain, burning, tension, cold). Venous sequelae were assessed for 5 days after injection to register signs of thrombophlebitis. Demographic data showed no difference between the two groups. After propofol more often a fall in blood pressure was seen. Pain (25 vs 1 pt), burning 19 vs 1), tension 15 vs 3), cold (35 vs 17) after injection was registered significantly more often in the propofol group, whereas myocloni predominated in the etomidate group (13 vs 6)  $P < 0.05$ , chi-squared-test). No difference was seen in PONV in either groups. Etomidate formulated in a medium chain lipid emulsion causes significant less discomfort for the patients than propofol, which is solved in a long chain formulation. Myocloni, however, occur significantly more frequently after etomidate than after propofol[10]. [Table 1] shows the mean hemodynamic parameters at different time intervals. No significant difference was observed while comparing the mean arterial pressure and heart rate among subjects of both the study groups at different time intervals except for at the time of induction. Mean pain score was found to be significantly higher in group A in comparison to group B. Aggarwal S et al compared propofol and etomidate for their effect on hemodynamics and various adverse effects on patients in general anaesthesia. Hundred ASA I and II patients of age group 18-60 years scheduled for elective surgical procedure under general anaesthesia were randomly divided into two groups of 50 each receiving propofol (2mg/kg) and etomidate (0.3mg/kg) as an induction agent. Vital parameters at induction, laryngoscopy and thereafter recorded for comparison. Adverse effect viz. pain on injection, apnea and myoclonus were carefully watched. Demographic variables were comparable in both the groups. Patients in etomidate group showed

little change in mean arterial pressure (MAP) and heart rate (HR) compared to propofol ( $p > 0.05$ ) from baseline value. Pain on injection was more in propofol group while myoclonus activity was higher in etomidate group. This study concluded that etomidate is a better agent for induction than propofol in view of hemodynamic stability and less pain on injection[11].

### Conclusion

Under the light of above obtained results, the authors conclude that among patients with associated altered hemodynamic status, etomidate is an improved option. However; further studies are recommended.

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