Original Research Article

Comparison of hemodynamic effects of etomidate and propofol as an induction agent in patients with LVEF $\leq 40\%$ undergoing elective coronary artery bypass grafting surgery – A prospective and randomized study

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ABSTRACT

Background: This study aims to compare the hemodynamic response of two commonly used induction agents Propofol and Etomidate in patients with left ventricular ejection fraction $\leq 40\%$ undergoing Coronary Artery Bypass Grafting surgery.

Aim: To compare the effects of Propofol and Etomidate on hemodynamics in terms of heart rate, rhythm, blood pressure, and central venous pressure.

Setting: Narayana Multispecialty Hospital, Jaipur.

Design: Prospective, double-blinded, randomized, hospital based study.

Materials and Methods: 100 patients with LVEF $\leq 40\%$ scheduled for elective CABG, were randomly assigned to one of the two groups receiving either of the inducing agents, group A (PROPOFOL 2mg/kg) and group B (ETOMIDATE 0.2mg/kg).

Statistical Analysis: Unpaired t-test and Chi square test/Fisher exact test, $p < 0.05$ was taken as significant.

Results: HR, SBP, DBP and MAP decreased from post induction 1 min. to post induction 3 min. (fall greater in propofol, $p > 0.005$) then increased from post intubation 1 min. to post intubation 5 min. in both the groups (rise greater in etomidate group, $p < 0.005$). Need of drugs to control hypotension (62% and 26%, $p = 0.001$) and to control pressor response (10% and 38%, $p = 0.002$) was observed in both the groups.

Conclusion: Etomidate is superior to propofol in providing hemodynamic stability before and after laryngoscopy and intubation, but less effective in controlling the pressor response to intubation. Therefore, Etomidate can be used as an induction agent with suitable adjuvants to control pressor response to tracheal intubation in patients undergoing CABG with low LVEF.

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1. Introduction

Incidence of coronary artery disease is steadily on rise and coronary artery bypass grafting (CABG) surgery is the commonest performed cardiac surgery. Patients with low left ventricle ejection fraction (LVEF) undergoing CABG constitute a high risk group.¹⁻³

Anaesthetic induction agents produces variable degree of hypotension while laryngoscopy and endotracheal intubation produces hypertension and tachycardia. These changes in hemodynamics may alter the balance between

myocardial oxygen supply and demand which can be detrimental in this high risk group of patients undergoing CABG.⁴⁻⁶

Various anaesthetic agents like Thiopentone, propofol, midazolam and Etomidate are in current use as an induction agents but no single anaesthetic agent is suitable for all patients as all of these agents have their advantages and disadvantages.

Propofol,⁷⁻⁹ an alkylphenol derivative, provides rapid onset and short duration of action. It causes considerable reduction in systemic vascular resistance and arterial pressure 15% to 40% after iv induction with 2mg/kg.

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Its effect on HR is variable. It causes direct myocardial depression at doses above 0.75mg/kg.

Etomidate^{3,7,10} is a carboxylated imidazole derivative, has a rapid onset (10-60 sec), a brief duration of action (3-5 min), and hydrolyses primarily in liver. It provides hemodynamic stability in both noncardiac and cardiac disease patients after dosage of 0.15 to 0.30 mg/kg. It directly inhibits 11-beta hydroxylation, which results in temporary reduction in biosynthesis of cortisol and aldosterone.^{11–14}

Considering the paucity of information in patients with left ventricular dysfunction undergoing CABG, this study aims to compare the hemodynamic response of two commonly used induction agents Propofol and Etomidate for anesthetic induction in patients with LVEF $\leq 40\%$ who are undergoing CABG.

2. Materials and Methods

The study was conducted after prior approval of Institutional Ethical Committee (IEC).

2.1. Study design

Hospital based, Prospective, Randomized, Double blind and comparative study.

2.2. Inclusion criteria

1. Age 35 to 65 years of either sex.
2. Patient with LVEF $\leq 40\%$ scheduled for elective CABG.

2.3. Exclusion criteria

Patients with-

1. Malampatti grade 3 or 4
2. LVEF $< 30\%$
3. Allergy to these drugs
4. Coexisting valvular heart disease
5. Preoperative persistent arrhythmia and congestive cardiac failure.
6. Preoperative requirement of inotropes, intra aortic balloon pump and mechanical ventilation.
7. history of adrenal insufficiency, chronic steroid use and severe systemic non cardiac disease other than hypertension and diabetes.

2.4. Intervention

Randomization - randomization was done using computer generated random numbers and blinding was done by sealed envelope method. This study was conducted on 100 patients which were randomly allocated into two groups, 50 in each group

Group A received Propofol 2mg/kg (10mg/ml)

Group B received Etomidate 0.2mg/kg (1mg/ml)

2.5. Data collection

After approval from IEC and written informed consent, patient was taken in operation theatre. Initial monitoring inside the theatre included five lead ECG, non-invasive blood pressure, and pulse oximetry were applied. Under local anaesthesia 16G arterial line and 7Fr central venous line were placed in the right femoral artery and the right internal jugular vein, respectively. Patients were preloaded with ringer lactate solution 5ml/kg and thereafter intravenous fentanyl 2 mcg/Kg and midazolam 0.04mg/kg were administered over a period of one minute to all patients. After a period of five minutes, the baseline data in the form of heart rate, systolic, diastolic, mean systemic arterial pressures and CVP were recorded during the study period in all the patients. Subsequently, general anaesthesia was induced with one of the agents, depending on the group. The induction agent was administered slowly over a period of 60 seconds. Thereafter, an additional dose of Fentanyl 2mcg/kg and Vecuronium bromide 0.1 mg/Kg were administered to facilitate tracheal intubation, which was done three minutes after the end of induction.

The patients were ventilated by mask and ventilator with tidal volume 8ml/kg till intubation as manual ventilation might lead to development of PEEP which might cause the changes in hemodynamics. Hemodynamic data were recorded at baseline and post induction 1 min.(T1), 3 min.(T2) and post intubation 1min.(T3), 3min.(T4), 5min.(T5) and 10 min.(T6) Throughout that period the lungs were mechanically ventilated with 100% oxygen, to maintain an end-tidal carbon dioxide between 30 and 35 mmHg.

Hypotension (MAP ≤ 60 mm Hg) was treated with incremental doses of Phenylephrine upto three times and thereafter infusion of Noradrenaline (0.05 mcg/kg/min) was started.

Hypertension (MAP ≥ 100 mm Hg) was treated with fentanyl 1 μ g/kg up to three times and then with incremental dose of Nitroglycerine infusion (5 mcg/min at 3min interval). Bradycardia (HR ≤ 50 min) was treated with atropine 0.6 mg iv. Tachycardia (HR ≥ 100 min) was treated with repeated doses of fentanyl 1 μ g/kg upto three times and thereafter with institutional protocol.

2.6. Outcome measures

Primary objective was to evaluate and compare the effects of Propofol and Etomidate on hemodynamics in terms of heart rate, rhythm, mean blood pressure, and central venous pressure.

Secondary objectives to compare the effect of Propofol and Etomidate on hemodynamics in terms of-

Diagram 1: Consort 2010 flow diagram

1. Need of inotrope or vasopressor to control hypotension.
2. Need of drugs to control pressor response to laryngoscopy and intubation.
3. Any side effects.

2.7. Data management and statistical analysis

Continuous variables were summarized as mean and standard deviation, whereas nominal/Categorical variables as proportions(%). Unpaired t-test was used for analysis of continuous variables. While Nominal/Categorical variables were analysed by using Chi square test/Fisher exact test. P value <0.05 was taken as significant. MedCalc.16.4 version software was used for all statistical calculation.

3. Results

3.1. Patient characteristics across the groups (Table 1 and Table 2)

There was no significant difference in patients of two groups with respect to age, weight, BMI, sex, comorbid conditions (hypertension and diabetes mellitus), left ventricle ejection fraction, disease status (left main disease) and baseline hemodynamic parameters.

3.2. Hemodynamic parameters (Table 3)

HR slightly decreased post induction in both the groups and this fall is higher in propofol group but comparison is statistically insignificant. HR at T₃ increased in both the groups but slightly higher in Group B and comparison of these data were statistically insignificant. (p = 0.163)

HR at T₄ increased in both the groups but higher in Group B and comparison was statistically significant. ($p = 0.045$) Thereafter HR decreased slowly and returned around baseline at T₆ in both the groups.

MBP decreased from T₁ to T₂ in both the groups but fall in SBP was slightly higher in Group A and comparison of these data were statistically insignificant. MBP increased from T₃ to T₄ in both the groups but rise being higher in Group B and comparison was statistically significant. Thereafter MBP decreased slowly and returned around baseline at T₆ in both the groups. (Figure 1)

In Group A, 62% patients had hypotensive episode and required drug to control hypotension according to our study protocol while in Group B, 26% patients had hypotension and required vasopressor. Comparison of these data were statistically significant. ($p = 0.001$) (Figure 2)

In Group A, 10% patients had post intubation pressor response which required adjuvants to control pressor response according to our study protocol while in Group B, 38% patients had pressor response and required drug. Comparison of these data were statistically significant. ($p = 0.002$)

Fig. 1: Comparison of Mean Blood Pressure (MBP) in Group A and Group B

4. Discussion

Propofol and Etomidate are two commonly used intravenous induction agents. Hypotension is common with propofol induction due to vasodilatation caused by reduction in sympathetic activity, direct effect on intracellular calcium mobilization and inhibition of prostaglandin synthesis in endothelial cells.¹⁵ Etomidate does not cause this hypotension. Hemodynamic stability seen with etomidate may be partially due to lack of effect on the sympathetic nervous system and on baroreceptor function.

Various authors have used different dosages of propofol and etomidate for induction in patients undergoing cardiac

Fig. 2: Comparison of secondary outcome in Group A and Group B

surgery with the range of 1.5 -2 mg/kg for propofol and 0.2-0.3 mg/kg for etomidate. For ease of blinding, we selected an induction dose of 2mg/kg for propofol and 0.2 mg/kg of etomidate for our study. (Pandey et al.¹² and Kaushal et al.¹¹)

4.1. Hemodynamic parameters

HR slightly decreased post induction in both the groups and this fall was greater in propofol group but comparison is statistically insignificant. Kamath MN et al.¹⁶ also concluded that fall in HR was more in propofol group. The observation was similar to our study. However, this observation is consistent with most of the studies comparing etomidate and propofol in cardiac as well as non cardiac surgery settings irrespective of cardiac function.

HR increased post intubation in both the groups but the rise was greater in etomidate group and comparison was statistically significant at 2 min post intubation. ($p = 0.045$) Thereafter HR returned slowly around baseline at post intubation 15min in both the groups. This observation was consistent with Singh et al.¹⁴ which shows etomidate group was the least effective of all the groups in minimizing stress response, with statistically significant increase from baseline in both heart rate ($P = 0.001$) and mean arterial pressure ($P = 0.001$) at 1 minute after intubation. However, this observation was contrary to Young et al.¹⁷ which shows no group differences in SBP, DBP, and HR following intravenous anesthetic drug injection and endotracheal intubation. Probably the reason behind this is the dose of etomidate (0.3 mg/kg) used in the study which is relatively higher than dose (0.2 mg/kg) used in our study.

SBP, DBP and MAP decreased from post induction 1 min. to post induction 3 min. in both the groups but fall in BP was slightly higher in propofol and comparison of these data were statistically insignificant. This fall in hemodynamics is attributable to study drugs and other

Table 1: Demographic parameters, comorbid conditions and disease status

S. No.	Parameters	Group A N= 50	Group B N=50	P value
1	Mean Age (Yr)	61.54 ± 7.55	61.82 ± 8.96	0.866
2	Mean Weight (Kg)	68.3 ± 6.9	67.16 ± 6.12	0.384
3	Mean Height (cm)	162.52 ± 4.41	163.00 ± 5.05	0.614
4	BMI	25.85 ± 2.22	25.30 ± 2.28	0.228
5	Sex Male	45(90%)	44(88%)	0.760
	Female	5(10%)	6(12%)	
6	Hypertension	30(60%)	31(62%)	1.000
7	Diabetes Mellitus	22(44%)	25(50%)	0.689
8	Left Main Disease	16(32%)	15(30%)	1.000
9	Left Ventricular ejection fraction 30-33	8(16%)	6(12%)	0.782
	34-36	18(36%)	17(34%)	
	37-40	24(48%)	27(54%)	

Table 2: Comparison of baseline parameters

S.No	Vitals	Group A (N= 50) Mean ± SD	Group B (N=50) Mean ± SD	P value
1	Heart Rate (per minute)	77.44 ± 8.87	76.10 ± 8.63	0.446
2	Systolic Blood pressure (mm Hg)	125.32 ± 12.20	124.15 ± 10.70	0.621
3	Diastolic Blood pressure (mmHg)	77.12 ± 7.72	76.50 ± 6.80	0.671
4	Mean Arterial Pressure (mmHg)	93.22 ± 8.86	92.38 ± 7.58	0.612
5	Central Venous Pressure (mmHg)	9.7 ± 1.93	9.68 ± 1.77	0.957

Table 3: Comparison of hemodynamic parameters

Parameters	T1			T2			T3		
	Group A mean±sd	Group B mean±sd	P value	Group A mean±sd	Group B mean±sd	P value	Group A mean±sd	Group B mean±sd	P value
HR (/ min.)	76.86±7.70	77.06±8.62	0.903	75.04±7.85	76.52±7.56	0.339	79.08±11.38	82.12±10.23	0.163
MBP (mmHg)	87.96±7.49	88.26±6.33	0.829	82.70±10.83	86.02±10.67	0.126	83.56±14.13	91.96±17.52	0.010
CVP (mmHg)	9.62±1.98	9.60±1.82	0.958	9.36±2.02	9.26±2.02	0.805	9.24±2.03	9.28±1.76	0.916
Parameters	T4			T5			T6		
	Group A mean±sd	Group B mean±sd	P value	Group A mean±sd	Group B mean±sd	P value	Group A mean±sd	Group B mean±sd	P value
HR (/min.)	79.06±8.80	82.16±12.54	0.156	78.82±8.30	81.10±8.92	0.189	78.42±7.84	80.34±8.41	0.240
MBP (mmHg)	86.74±10.60	92.60±12.05	0.011	87.70±10.94	91.38±8.88	0.068	89.70±8.93	91.14±7.35	0.381
CVP (mmHg)	8.90±1.88	8.84±1.86	0.873	8.78±1.75	8.86±1.92	0.828	8.74±1.56	8.60±1.86	0.685

drugs used in anaesthesia technique as well as positive pressure ventilation during induction of general anaesthesia. SBP, DBP and MAP increased from post intubation 1 min. to post intubation 5 min. in both the groups but rise being higher in etomidate and statistically significant as compared to propofol, this could be due to stress response to laryngoscopy and endotracheal intubation. Thereafter SBP, DBP and MAP returned slowly around baseline at post intubation 15 min. in both the groups. These observations are consistent with Kaushal RP et al.,¹¹ Kamath MN et al.,¹⁶ Sivanna S et al.,¹⁸ and Shukla N et al.¹⁹ The observations of Singh R et al.¹⁴ are dissimilar to the present study as

they concluded that both anesthetic agents were acceptable for induction in patients with coronary artery disease and left ventricular dysfunction despite a 27-32% decrease in the mean arterial pressure and the outcome of anesthetic induction may depend on factors such as the speed of injection, route, dose and experience of the clinician, other than the property of the agent itself.

CVP was comparable between two groups at all time intervals. This observation is similar to most of the studies done in cardiac settings.

4.2. Secondary outcome variable

In our study, we observed that hypotensive episodes occurred more in propofol group and required drugs to control these episodes, and this comparison was statistically significant ($P=0.001$). This observation was similar to Shah et al.²⁰ study. Hemodynamic trend in the Afshin G B et al.²¹ study is similar to our study but the ephedrine prescription rate due to hypotension is 5% (2 patients) in etomidate. This rate is lower than rate observed in our study. Probably this may be due to more contribution to hypotension by higher dose of fentanyl in our study.

We observed that post intubation pressor response occurred more in etomidate group and required drug to control this response, and this comparison was statistically significant ($p=0.002$). This observation is consistent with Singh R et al.¹⁴ and Shivanna S et al.¹⁸ study.

Etomidate causes adrenocortical suppression. Kaushal RP et al.¹¹ and Pandey AK et al.¹² found that etomidate can be safely used in these patients with better hemodynamic stability and without cortisol suppression lasting more than 24 hours. Whereas Cuthbertson BH et al.²² did a study to check steroid suppression in critically ill patient. They found that use of bolus dose etomidate is associated with an increased incidence of inadequate response to corticotropin, but is also likely to be associated with an increase in mortality.

5. Conclusion

Etomidate is superior in providing hemodynamic stability prior to and after laryngoscopy and tracheal intubation as compared to propofol, but is lesser effective in controlling the pressor response to tracheal intubation. Therefore, Etomidate can be used as an induction agent with suitable adjuvants to control pressor response to tracheal intubation in patients undergoing CABG with low LVEF.

6. Limitations

1. We did not compare adrenocortical suppression between two drugs.
2. In our study, we did not compare advanced hemodynamic parameters.
3. We maintained the hemodynamics within a range with the use of rescue drugs.

7. Source of Funding

None.

8. Conflict of Interest

The authors declare that there is no conflict of interest.

References

1. Moffitt EA, Sethna DH. The coronary circulation and myocardial oxygenation in coronary artery disease: effects of anesthesia. *Anesth Analg*. 1986;65:395–410.
2. Merin RG. Is anaesthesia beneficial for the ischemic heart? *Anesthesiology*. 1980;53:439–40.
3. Bovill JG. Intravenous Anesthesia for the Patient with Left Ventricular Dysfunction. *Semin Cardiothorac Vasc Anesth*. 2006;10:43–8. doi:10.1177/108925320601000108.
4. Mogo EB, Goren S, Korfali G, Turker G, Kaya FN. Induction of anesthesia in coronary artery bypass graft surgery: the hemodynamic and analgesic effects of ketamine. *Clinics*. 2010;65(2):133–8. doi:10.1590/s1807-59322010000200003.
5. Habibi MR, Baradari AG, Soleimani A. Hemodynamic responses to etomidate versus ketamine-thiopental sodium combination for anesthetic induction in coronary artery bypass graft surgery patients with low ejection fraction: a double-blind, randomized, clinical trial. *J Clin Diagn Res*. 2014;8:1–5.
6. Kanchi M, Banakal S, Murugesan C, Nair HC, Murthy K. Haemodynamic response to endotracheal intubation in coronary artery disease: Direct versus video laryngoscopy. *Indian J Anaesth*. 2011;55(3):260–5. doi:10.4103/0019-5049.82673.
7. Lester L, Mitter N, Daniel, Dan EB. Pharmacology of anesthetic drugs. In: Kaplan JA, David LR, John GT, Gerard R, Maus T, editors. *Kaplan's Cardiac Anesthesia*. Philadelphia: Saunders Elsevier; 2017. p. 269–70.
8. Reves JG, Glass P, Lubarsky DA, Mcevoy MD, Martinez-Ruiz R. Anesthesia. In: Miller RD, editor. *Intravenous anesthesia*. New York: Churchill Livingstone; 2010. p. 719–58.
9. Stephan H, Sonntag H, Schenk HD, Kettler D, Khambatta HJ. Effects of propofol on cardiovascular dynamics, myocardial blood flow and myocardial metabolism in patients with coronary artery disease. *Br J Anaesth*. 1986;58(9):969–75. doi:10.1093/bja/58.9.969.
10. Massaut J, Hollander A, Barvais L, Dubois-Primo J. Cardiovascular and pulmonary responses following etomidate induction of anesthesia in patients with demonstrated cardiac disease. *Acta Anaesthesiol Scand*. 1983;27:299–302.
11. Kaushal RP, Vatal A, Pathak R. Effect of etomidate and propofol induction on hemodynamic and endocrine response in patients undergoing coronary artery bypass grafting/mitral valve and aortic valve replacement surgery on cardiopulmonary bypass. *Ann Card Anaesth*. 2015;18(2):172–8. doi:10.4103/0971-9784.154470.
12. Pandey AK, Makhija N, Chauhan S, Das S, Kiran U, Bisoi AK, et al. The Effects of Etomidate and Propofol Induction on Hemodynamic and Endocrine Response in Patients Undergoing Coronary Artery Bypass Graft Surgery on Cardiopulmonary Bypass. *World J Cardiovasc Surg*. 2012;02(03):48–52. doi:10.4236/wjcs.2012.23011.
13. Basciani RM, Rindlisbacher A, Begert E, Brander L, Jakob SM, Etter R, et al. Anaesthetic induction with etomidate in cardiac surgery. *Eur J Anaesthesiol*. 2016;33(6):417–24. doi:10.1097/eja.0000000000000434.
14. Singh R, Choudhury M, Kapoor PM, Kiran U. A randomized trial of anesthetic induction agents in patients with coronary artery disease and left ventricular dysfunction. *Ann Card Anaesth*. 2010;13(3):217–23. doi:10.4103/0971-9784.69057.
15. Miller RD, Miller RD. *Anesthesia*. 2005.
16. Kamath MR, Kamath S, Patla KP. Propofol or etomidate: Does it genuinely matter for induction in cardiac surgical procedures? *Indian J Clin Anaesth*. 2016;3(4):551–5.
17. Ko YK, Kim YH, Park S, Chung WS, Noh C, Lee JU. Comparison of etomidate and propofol on intubating conditions and the onset time associated with cisatracurium administration. *Korean J Anesthesiol*. 2015;68(2):136–40. doi:10.4097/kjae.2015.68.2.136.
18. Shivanna S, Priye S, Jagannath S, Kadli C, Mayuri, Subhash V, et al. A Comparative Study of Haemodynamic Effects of Propofol and Etomidate as an Induction Agent in Coronary Artery Surgery". *J Evol Med Dent Sci*. 2015;4:598–607.
19. Shukla N, Kushwaha BB, Kausal D, Singh V, Singh SK, Devenraj V, et al. Comparative Study of Etomidate and Midazolam with Fentanyl as Inducing Agents in Patients Undergoing Off-pump Coronary

- Artery Bypass Graft Surgery. *J Clin Diagn Res*. 2018;12(5):7–10. doi:10.7860/jcdr/2018/29105.11499.
20. Shah SB, Chowdhury I, Bhargava AK, Sabbharwal B. Comparison of hemodynamic effects of intravenous etomidate versus propofol during induction and intubation using entropy guided hypnosis levels. *J Anaesthesiol Clin Pharmacol*. 2015;31(2):180–5. doi:10.4103/0970-9185.155145.
21. Baradari AG, Alipour A, Habibi MR, Rashidaei S, Zeydi AE. A randomized clinical trial comparing hemodynamic responses to ketamine-propofol combination (ketofol) versus etomidate during anesthesia induction in patients with left ventricular dysfunction undergoing coronary artery bypass graft surgery. *Arch Med Sci*. 2017;5:1102–10. doi:10.5114/aoms.2016.63193.
22. Cuthbertson BH, Sprung CL, Annane D, Chevret S, Garfield M, Goodman S, et al. The effects of etomidate on adrenal responsiveness and mortality in patients with septic shock. *Intensive Care Med*. 2009;35:1868–76. doi:10.1007/s00134-009-1603-4.

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