ORIGINAL ARTICLE

Fluid Preloading Versus Ephedrine in the Management of Spinal Anesthesia-Induced Hypotension in patients undergoing Cesarean Delivery

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ABSTRACT

Objective: To compare the efficacy of prophylactic ephedrine versus fluid preload (Hartmann's solution) in preventing hypotension in women undergoing cesarean section under spinal anesthesia at a teaching hospital

Methodology: Study was conducted in the Department of Anesthesiology at Liaquat University Hospital, Hyderabad from January 2023 to September 2023. Patients were randomly divided into two groups (Group A and Group B) using a draw method, where they selected sealed envelopes marked A or B. Group A received prophylactic ephedrine (0.25 mg/kg intravenous infusion immediately after bupivacaine injection), while Group B received prophylactic fluid preload (20 mL/kg Hartmann's solution over 10–15 minutes before spinal anesthesia).

Results: The study enrolled a total of 62 patients, equally divided into Group A and Group B, with 31 patients in each group. The two groups were comparable in terms of age (32.32 ± 5.62 years in Group A vs. 30.35 ± 3.31 years in Group B, p = 0.197) and BMI (26.96 ± 2.88 kg/m² in Group A vs. 27.03 ± 3.31 kg/m² in Group B, p = 0.964). In terms of efficacy, Group A demonstrated a significantly higher success rate compared to Group B (67.7% vs. 38.7%, p = 0.022).

Conclusion: The frequency of hypotension is lower in women receiving prophylactic ephedrine during spinal anesthesia for cesarean section compared to those given a Hartmann's solution preload.

Keywords: Spinal anesthesia, Fluid preload, Hypotension, Ephedrine, Cesarean section.

INTRODUCTION

Spinal anesthesia is widely used for cesarean deliveries due to its rapid onset and effective pain control¹. However, a significant complication is spinal anesthesia-induced hypotension (SAIH), which occurs in 50-80%² of parturients without prophylactic intervention. This abrupt drop in blood pressure results from sympathetic blockade, leading to reduced systemic vascular resistance and venous return³. Hypotension can compromise uteroplacental perfusion, increasing the risk of fetal acidosis and maternal nausea or vomiting. Given its high incidence and potential adverse effects, effective prevention and management strategies are essential in obstetric anesthesia⁴.

Two common approaches to mitigate SAIH are fluid preloading and vasopressor administration, such as ephedrine⁵. Fluid preloading, typically with crystalloids or colloids, aims to expand intravascular volume and counteract vasodilation⁶. However, its efficacy remains debated, as rapid fluid shifts and limited duration of action may reduce its effectiveness. Conversely, ephedrine, a mixed alpha- and beta-adrenergic agonist, increases blood pressure by enhancing cardiac output and peripheral resistance⁷. While effective, ephedrine may cross the placenta and cause fetal tachycardia or acidosis, raising concerns about its safety^{8,9}.

The optimal strategy for preventing SAIH in cesarean deliveries remains controversial. Some studies advocate for combined fluid and vasopressor therapy, while others suggest that vasopressors alone may suffice¹⁰. Fluid preloading is non-pharmacological and avoids drug-related side effects, but its benefits may be limited in parturients with rapid hemodynamic changes. On the other hand, ephedrine provides a reliable pressor response but requires careful dosing to minimize fetal effects. Understanding the comparative efficacy and safety of these interventions is important for improving maternal and neonatal outcomes¹¹.

The findings of this study will contribute to evidence-based guidelines for managing SAIH in obstetric anesthesia. If fluid

Received on 10-10-2023 Accepted on 26-12-2023 preloading proves non-inferior to ephedrine, it could reduce reliance on vasopressors and minimize fetal drug exposure. Conversely, if ephedrine demonstrates superior hemodynamic control, its judicious use may be prioritized. Ultimately, this research will help optimize anesthesia protocols to enhance outcomes for both mothers and newborns undergoing cesarean delivery.

METHODOLOGY

Study was randomized controlled trial in design and conducted in the Department of Anesthesiology at Liaquat University Hospital, Hyderabad from January 2023 to September 2023. Study was started after obtaining approval from the Ethics Committee. Sample size of 62 patients was calculated using 95 CI, margin of error 0.5% and efficacy of ephedrine 74%² and fluid preload 50%². Participants were women aged 20 to 40 years, both nulliparous and multiparous, with a gestational age greater than 37 weeks, undergoing cesarean section under subarachnoid block. Exclusion criteria comprised patients with local anesthetic allergies, hypertension, shock, bleeding disorders, spinal injection site infections, spinal deformities, or known valvular heart disease.

Before the procedure, all patients were thoroughly briefed about the study protocol, potential risks, and benefits, and written informed consent was obtained from each participant. Baseline demographic and clinical data, including weight, height, body mass index (BMI), age, type of cesarean section (elective or emergency), and baseline blood pressure measurements, were meticulously recorded. Using a randomized draw method, patients were allocated into two groups Group A and Group B by selecting sealed, opaque envelopes labeled either "A" or "B." Group A received prophylactic intravenous ephedrine at a dose of 0.25 mg/kg, administered immediately after the intrathecal injection of bupivacaine, while Group B received a prophylactic fluid preload consisting of 20 mL/kg of Hartmann's solution, infused over 10–15 minutes prior to the administration of spinal anesthesia.

Spinal anesthesia was performed by a senior consultant anesthetist using a standardized technique with a 27-gauge pencil-point spinal needle at the L3-L4 or L4-L5 interspace, with correct positioning confirmed by free cerebrospinal fluid (CSF) flow.

Following the induction of spinal anesthesia, blood pressure was continuously monitored at regular intervals for the first 15 minutes to promptly detect any episodes of hypotension, which was defined as a reduction of more than 20% in systolic and/or diastolic blood pressure compared to the patient's baseline values. This rigorous monitoring protocol ensured timely intervention if hypotension occurred, thereby maintaining maternal and fetal hemodynamic stability throughout the procedure.

The researcher recorded all relevant data, which were analyzed using SPSS-27. Numerical variables are presented as mean \pm SD, to compare the two means (Student's t-test) was applied . Categorical variable are presented as N (%) , to check the association (chi-square test) was applied. P value less than or equal to 0.05 was considered as significant.

RESULTS

The study enrolled a total of 62 patients, equally divided into Group A and Group B, with 31 patients in each group. The two groups were comparable in terms of age (32.32 \pm 5.62 years in Group A vs. 30.35 \pm 3.31 years in Group B, p = 0.197) and BMI (26.96 \pm 2.88 kg/m² in Group A vs. 27.03 \pm 3.31 kg/m² in Group B, p = 0.964). However, Group A had a significantly higher proportion of urban residents (64.5% vs. 38.7%, p = 0.042), while Group B had more rural participants (61.3% vs. 35.5%, p = 0.042). No significant difference was observed in procedure type (emergency/elective) between the groups (p = 0.442). Table-1

In terms of efficacy, Group A demonstrated a significantly higher success rate compared to Group B (67.7% vs. 38.7%, p = 0.022). Table-2

Table 1: Demographics and baseline profile

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Variable	Group A 31 (50.0%)	Group B 31 (50.0%)	Test of sig.	
Age (years)	32.32±5.62	30.35±3.31	t=1.31, d.f=60, p=0.197	
BMI (kg/m²)	26.96±2.88	27.03±3.31	t=-0.045, d.f=60, p=0.964	
Area of residence				
Urban	20 (64.5)	12 (38.7)	χ ² =4.13, d.f=1, p=0.042	
Rural	11 (35.5)	19 (61.3)		
Procedure type				
Emergency	12 (38.7)	15 (48.4)	χ ² =0.59, d.f=1, p=0.442	
Elective	19 (61.3)	16 (51.6)		
N (%), chi-square test was applied.				

Table 2: Comparison of efficacy among the study groups

Efficacy	Group A 31 (50.0%)	Group B 31 (50.0%)	Test of sig.	
Yes	21 (67.7)	12 (38.7)	2 5 05 4 5 4 = 0.000	
No	10 (32.3)	19 (61.3)	χ ² =5.25d.f=1, p=0.022	
N (%), chi-square test was applied.				

DISCUSSION

The demographic similarity between the two groups in this study aligns with previous research, where age and BMI were well-matched in trials comparing vasopressors and fluid preload for hypotension management. A study by Prakash et al¹² also reported no significant differences in age or BMI between groups receiving ephedrine or crystalloid preload (p>0.05), supporting the validity of our findings.

Our findings on ephedrine's effectiveness are consistent with earlier studies demonstrating its reliability in preventing spinal anesthesia-induced hypotension. Ngan Kee et al¹³ found that prophylactic ephedrine significantly reduced hypotension incidence compared to placebo (p<0.001). However, some studies suggest that phenylephrine may be superior due to fewer fetal acid-base disturbances, indicating a need for further comparative studies¹⁴.

The fluid preload group's results are comparable to those reported in previous studies, where crystalloid preloading showed moderate efficacy but was often inferior to vasopressors in sustained blood pressure control. A meta-analysis by Banerjee et al¹⁵ concluded that while fluid preload reduces hypotension, it is less effective than vasopressors in preventing severe drops in

blood pressure. This aligns with our observations, suggesting that fluid preload alone may be insufficient in high-risk patients.

Desalu et al¹⁶ compared fluid preload using one liter of normal saline alone versus one liter of normal saline mixed with 30 mg of ephedrine and found that the systolic blood pressure was higher in the group that received normal saline with ephedrine. After the subarachnoid block, the preload group most frequently experienced a drop in blood pressure at 5 minutes, whereas in the ephedrine group, this occurred at 10 minutes.

The study by Kamat et al¹⁷ demonstrated that a prophylactic strategy involving an initial 6 mg ephedrine bolus administered at the onset of subarachnoid block, followed by a continuous 24 mg ephedrine infusion, was significantly more effective in preventing spinal-induced hypotension during cesarean delivery compared to crystalloid preloading alone. These findings align with the results reported by Chan et al¹⁸ and Lee et al¹⁹, who similarly observed a higher incidence of hypotension in patients who received crystalloid preloading as opposed to those managed with prophylactic ephedrine administration. The consistency across these studies suggests that vasopressor-based interventions, such as ephedrine bolus and infusion, provide superior hemodynamic stability in parturients undergoing spinal anesthesia for cesarean delivery when compared to volume expansion with crystalloids alone. This may be attributed to ephedrine's direct alpha- and beta-adrenergic agonist effects, which counteract the sympathetic blockade-induced vasodilation, whereas crystalloid preloading alone may be insufficient to compensate for the rapid and profound decreases in systemic vascular resistance following spinal anesthesia.

A survey conducted by Burns et al²⁰ in the United Kingdom among obstetric anesthetists revealed that over 95% still prefer using ephedrine as the primary vasopressor during cesarean deliveries. Despite the growing popularity of alternative vasopressors, concerns persist regarding their safety and potential adverse effects on uterine artery and blood flow. Both alpha-1 agonists and ephedrine effectively maintain normotension in pregnant women undergoing cesarean sections without significantly compromising fetal well-being.

CONCLUSION

The frequency of hypotension is lower in women receiving prophylactic ephedrine during spinal anesthesia for cesarean section compared to those given a Hartmann's solution preload; therefore, we recommend the routine use of prophylactic ephedrine during spinal anesthesia induction in general practice to prevent spinal-induced hypotension.

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This article may be cited as: Khan A, Siddiqui MA, Asad H, Hussain S, Taqi M, Kumar K: Fluid Preloading Versus Ephedrine in the Management of Spinal Anesthesia-Induced Hypotension in patients undergoing Cesarean Delivery. Pak J Med Health Sci, 2024; 18(1): 302-304.