

Effect of magnesium sulphate on quality of subarachnoid block in terms of onset and duration of motor and sensory block, APGAR score of the neonate and haemodynamic status of the patient

*Shahadat Hossain¹, Montosh Kumar Mondal², Beauty Rani Roy³, Jesmin Akter⁴, AKM Akhtaruzzaman⁵, Wahiuddin Mahmood⁶

¹Dental Hospital, Dhaka, ^{2,5}Department of Anaesthesia, Analgesia and Intensive Care Medicine, Bangabandhu Sheikh Mujib Medical University, Dhaka, ³ Department of Obstetric and Gynaecology, OGSB Hospital and Reproductive Centre, Mirpur, Dhaka, ⁴ OSD DG Health, Dhaka, ⁶ Square Hospital, Dhaka.

Corresponding author: E mail – montoshmondal@yahoo.com

Abstract

Background In obstetrics, pregnancy induced hypertension is still a burning question and complicates a large number of pregnancies in developing countries. Chance of hypotension is more in patients getting magnesium sulfate with subarachnoid block but it may be managed with adequate preloading and by pressor agent ephedrine.

Objectives This study was designed to observe the effect of magnesium sulphate on quality of subarachnoid block in terms of onset and duration of motor and sensory block, APGAR score of the neonates and haemodynamic status of the patients.

Methods Sixty parturients undergoing caesarian sections under subarachnoid block were enrolled for the study. They were divided into two groups. Group-A include normal parturient undergoing caesarian section and group-B include pre-celamptic parturient treated with magnesium sulphate within 1 to 2 hours before block. After recording of base line haemodynamic status (BP, HR, SPO₂) all patients received subarachnoid block with 2 ml (10 mg) hyperbaric bupivacaine at L₃₋₄ level. Onset of sensory block was assessed by using pinprick, onset of motor block was assessed by onset time of weakness of lower limb and onset time of complete paralysis of lower limb after SAB. Duration of motor block was assessed by modified bromage scale. Height of the block was assessed by using pin prick at the intercostals space in the mid axillary line after 5 minute of SAB. Neonatal assessment was done by using apgar score in 1 and 5 minutes after delivery of baby. Blood pressure was recorded normally at 2 min interval until 15 minutes then every 5 minutes interval till the surgical procedure is completed.

Results Duration of motor block in group B is significantly higher 276 ± 44.92 min compared with group A which was 197.96 ± 24.25 min ($P = 0.000$). Duration of sensory block in group B also significantly higher with 308.76 ± 61.43 min compared with group A which was 264 ± 30.57 min, and ($P = 0.001$). Changes in systolic blood pressure in group B patient is more and highly significant ($P < .05$), for upto 60 min. But changes in diastolic blood pressure in-group B was only highly significant with group A for upto 9 minutes. APGAR score was significantly low both in 1 minute and 5 minutes, in group B patients which was $5.80 \pm .61$ at 1 minute and $7.73 \pm .827$ at 5 minutes and in group A which was $6.60 \pm .85$ at 1 minute and $8.30 \pm .595$ (mean \pm SD) at 5 minutes. Onset of sensory block and onset of motor block revealed on significant difference between groups.

Conclusions Chance of hypotension is more in patients getting magnesium sulfate but it may be managed with adequate preloading and by pressor agent ephedrine.. APGAR score of baby of magnesium sulfate getting patient is low but it is acceptable.

Keywords Subarachnoid block, caesarian section, magnesium sulfate and PIH.

Introduction

Magnesium is one of the important and second most common intracellular cation after potassium. Magnesium plays an important role in nearly every physiological system by calcium antagonism. Magnesium is involved in several processes including hormone receptor binding, gating of calcium channels, transmembrane ion flux & regulating adenylate cyclase-muscle contraction, neuronal activity, control of vasomotor tone, cardiac excitability and neurotransmitter release.

Magnesium is used as an anticonvulsant by its depressant effect at synapses¹. It may decrease catecholamine release¹⁹ and antagonize bronchospasm. In cardiology magnesium decreases frequency of both atrial & ventricular arrhythmia and causes vasodilatation⁶.

In obstetrics eclampsia & pre eclampsia is still now a burning question and complicates a large number of pregnancies in developing countries. This two remains a major cause of maternal & foetal mortality, primarily by causing cerebral haemorrhage & heart failure. Mortality directly correlates with the severity of hypertension. So anesthetic management should be directed towards avoidance of exacerbation of maternal hypertension. In this group of patient airway management and intubation may be difficult due to distortion of upper airway anatomy by oedema⁷ and chance of life threatening gastric acid aspiration and exaggerated intubation reflex is more¹⁸. Moreover general anaesthesia may decrease placental blood flow²¹ and may increase the risk of maternal stroke and heart failure in severely eclamptic & pre eclamptic patient.

Spinal anaesthesia causes blockade of motor, sensory and sympathetic nervous system by blocking sodium channels in peripheral nerves. At the motor end plate magnesium inhibits neurotransmitter release in peripheral nerves by competitive calcium antagonist for membrane channels on the pre-synaptic terminal²⁰. Direct neuromuscular block has also been suggested as a mechanism of action of magnesium in pre-eclampsia and eclampsia. So motor block due to spinal anaesthesia may be potentiated by magnesium sulphate. Marked Haemodynamic changes (i.e. $\bar{B.P}$) occur after spinal anaesthesia. On the other hand, magnesium sulphate acts by

calcium antagonism via calcium channels¹⁶ and it decreases systemic vascular resistance⁹ and reverses increase calcium ion mediated vasospasm¹⁷. Magnesium inhibits catecholamine release¹⁹ and basal myogenic and hormone induced smooth muscle contraction and also has direct vasodilator effect. So chance of hypotension is more in magnesium sulphate treated patient getting spinal anaesthesia. Spinal anaesthesia is often discouraged in patients with preeclampsia and eclampsia because of the risk of severe hypotension¹⁰, leading to maternal, foetal and neonatal morbidity. The first study to call this into question was by Wallace et al². In a prospective randomized trial of anaesthesia in parturients with severe preeclampsia, they compared general anaesthesia with epidural and CSE anaesthesia for caesarean section. The need for ephedrine due to hypotension was similar between the spinal and epidural groups. A retrospective study by Hood et al³ and a prospective study by Sharwood smith et al⁴ agreed with this findings.

Kerinen J. et al have studied the neonates born from pre-eclamptic patient under spinal anaesthesia. They did not find any major effect on clinical condition of the neonates assessed by apgar score and umbilical artery pH values¹².

Comparing general and regional anaesthesia, general anaesthesia is neither contraindicated nor regional anaesthesia indicated exclusively in women with severe pre-eclampsia. Some investigator has also concluded that the use of spinal anaesthesia in cases of severe pre-eclampsia should be reconsidered⁵. When caesarean section is indicated in preeclamptic and eclamptic parturient, a large number of patients remain under magnesium sulphate therapy because it is superior to diazepam or phenytoin in controlling seizure¹. But there is lack of studies on effect of magnesium sulphate therapy on quality of spinal anaesthesia. This study reveals outcome of spinal anaesthesia in patient getting magnesium sulphate therapy by assessing the effect of magnesium sulphate therapy on quality of spinal anaesthesia.

Methods

This randomized prospective study was carried out in the department of Anaesthesiology, Dhaka medical college hospital. With approval from the hospital ethical committee and written informed

consent, a total of 60 parturients undergoing caesarean section with sub arachnoid block were included in the study. Patients aged between 20-30 years, body weight 50-70 kg, ASA class I and II scheduled for caesarean section. Any one who had relative contraindication for regional anaesthesia were dropped from the study. Patients were divided into two groups: Group A, normotensive parturient undergoing C/S and Group B, parturient getting magnesium sulfate undergoing C/S. Volume preloading done with Hartman's solution 15 ml/kg over 20 to 30 minutes in all patients before giving SAB. Normal fluid balance was however maintained at a rate 4-5ml/kg-hr.

Spinal injections were made with a 25 G Quincke Babcock spinal needle with the patient in lateral position through L 2-3 or L3-4 inter space. In both group of patients 2 ml of 0.5% hyperbaric bupivacaine (10 mg) was injected. Following the end of injection skin patch was applied quickly and patients were immediately placed to supine position.

Preoperative and intraoperative pulse rate was monitored by pulse oxymetry. Blood pressure was recorded normally at 2 min interval over the right arm until 25 minutes than every 5 minutes interval till the surgical procedure was completed. Attempt was made to maintain systolic arterial pressure(SAP) > 90 mm Hg. For this purpose if hypotension occurs (SAP<90 mm Hg) intravenous infusion of crystalloid as necessary and injection ephedrine was given 5 mg intravenously(repeated as necessary). Onset of sensory block was assessed by using pinprick and asking question about tingling, onset of motor block was assessed by onset time of weakness of lower limb and onset time of complete paralysis of lower limb after SAB. Duration of motor block was assessed by modified bromage scale. Height of the block was assessed by using pin prick at the intercostals space in the mid axillary line after 5 minute of SAB. Neonatal assessment will be done by using apgar score in 1 and 5 minutes after delivery.

Severe hypo tension following SAB was defined as a fall of Systemic Arterial Pressure to or below 80 mm Hg and bradycardia as heart rate below 60/minute.If either severe hypotension (even with usual crystalloid infusion, with or without ephedrine) or bradycardia or both ensued, rescue treatment was to be initiated. An additional intravenous channel(18 G canula) was also to be opened for rapid infusion of colloid(500 ml dextran

40 over 15-20 minutes). Bradycardia was treated with atropine(0.3 mg I/V).

Statistical analysis

The results were compiled and analysed using unpaired t-test, Chi-square ??or ANOVA as appropriate. Results are considered statistically significant if $p < 0.05$.

Results

Demographic data was statistically matched (Table I) between the groups. There was significant difference of pulse rate (Fig1) between two groups from 3 min to 23 min. ($P < 0.05$). Systolic blood pressure (Fig2) significantly varied ($P < 0.05$) during whole period and diastolic blood pressure(Fig3) ($P < 0.05$) during initial period. Onset of sensory and motor block was (Table2 and Table 3) statistically significant ($P=0.009$). Duration of sensory(Table4) and motor block (Table5) was also significant APGAR score of the neonates in one and five minutes was highly significant ($p=0.004$).

Table-I: Demographic data:

Group/Variable	Group-A	Group-B
n	30	30
Age in years	24.53±3.38	24.43±4.31
Height in cm	156.21±3.12	155.35±3.26
Weight in kg	58.30±7.27	56.66±7.82

Values are expressed as mean± SD. Values are considered statistically significant if $p < 0.05$.

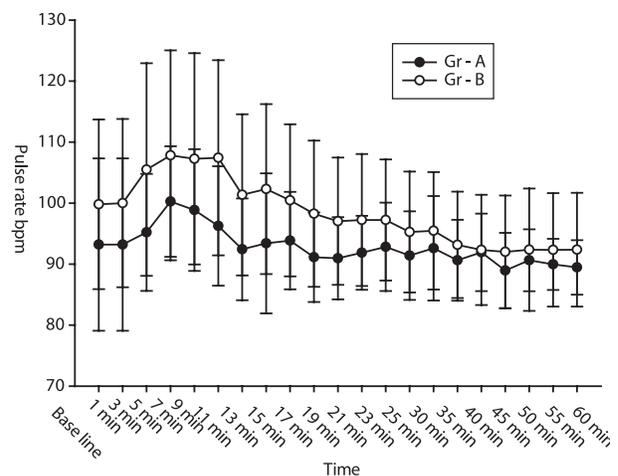


Fig 1 Changes in pulse rate in different time of two studied groups

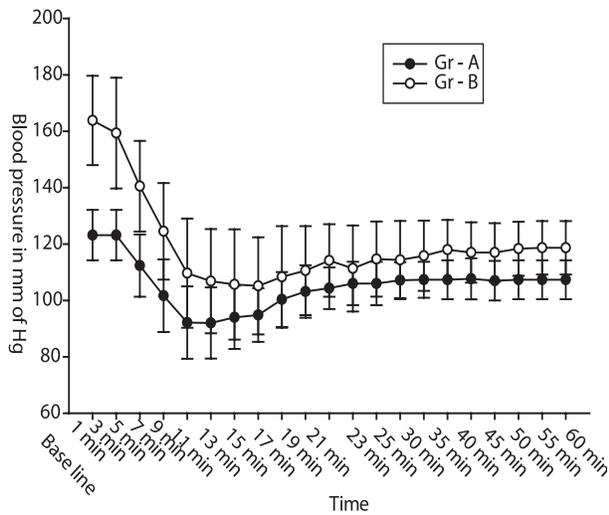


Fig.-2: Changes of systolic blood pressure in mm of Hg of two studied group

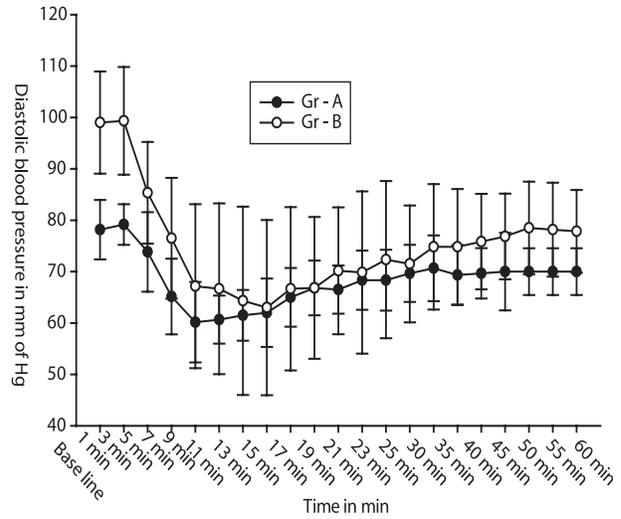


Fig.-3: Changes of diastolic blood pressure in mm of Hg of two studied group.

Table II : Onset of sensory block as indicated by tingling in the leg

Group / Variable	Group-A	Group-B	p value	Significant level
n=	30	30		
Tingling sensation in the leg in min	1.066±0.25	1.333±0.479	0.009	S

Values are expressed as mean± SD. Between groups analyses were done by student t test (unpaired). Values are expressed as significant if P<0.05 (CI-95%). S- Significant.

Table III : The onset of motor block as assessed by weakness of lower limb

Group/Variable	Group-A	Group-B	t	P value	Significant difference
Number of Patient	30	30			
Duration of Motor Block	197.96±24.25	276.26±44.92	-8.40	0.000	S

Values are expressed as mean± SD. Between groups analyses were done by student t test (unpaired). Values are expressed as significant if P<0.05 (CI-95%). S- Significant.

Table IV : Duration of sensory block

Group/Variable	Group-A	Group-B	t	P value	Significant Difference
Number of Patient	30	30			
Duration of Sensory Block	264.16±30.57	308.76±61.43	-3.560	0.001	S

Values are expressed as mean± SD. Between groups analyses were done by student t test (unpaired). Values are expressed as significant if P<0.05 (CI-95%). S- Significant.

Table V : Duration of motor block .

Group/Variable	Group-A	Group-B	t	P value	Significant difference
Number Of Patient	30	30			
Duration of Motor Block	197.96±24.25	276.26±44.92	-8.40	0.000	S

Values are expressed as mean± SD. Between groups analyses were done by student t test (unpaired). Values are expressed as significant if P<0.05 (CI-95%). S- Significant.

]

Table VI APGAR score in one and five minutes

Group/Variable	Group-A	Group-B	t	P value	Significant difference
Number Of Patient	30	30			
Weakness of Lower Limb	2.033±0.319	2.400±0.770	-2.408	0.019	S

Values are expressed as mean± SD. Between groups analyses were done by student t test (unpaired). Values are expressed as significant if P<0.05 (CI-95%).

Discussion

Pre eclampsia & eclampsia are the most common direct causes of pregnancy related death mainly due to stroke & heart failure. Mortality directly correlated with severity of hypertension. The mortality rate varies 2 to 5 %⁸. They are often treated with magnesium sulphate and antihypertensives. Because sound clinical research has shown beyond doubt that magnesium is superior to either diazepam or phenytoin for the prevention of recurrent convulsions¹. magnesium sulphate also decreases systemic vascular resistance⁹ and is beneficial for controlling acute hypertension.

Anaesthetic management is very critical in pre-eclamptic & eclamptic patients. Because general anaesthesia has several potential adverse effect like difficulties in airway management & intubation due to distortion of upper airway anatomy by edema, increase chance of aspiration, maternal stroke & heart failure & decreases placental blood flow. On the other hand, spinal anaesthesia is discouraged in pre eclamptic & eclamptic patients because of the risk of severe hypotension¹⁰.

Comparing general & regional anaesthesia, general anaesthesia is neither contra indicated nor regional anaesthesia indicated exclusively in women with severe pre eclampsia. Some investigator has also concluded that the use of spinal anaesthesia in cases of severe pre eclampsia should be reconsidered.⁵ The first study to call this in to question was by Wallace et al² in a prospective randomized trial of anaesthesia in parturients with severe pre eclampsia. They compared general anaesthesia with epidural & CSE anaesthesia for caesarean section. They found that the need for ephedrine was similar between the spinal & epidural groups and there was no significant difference in maternal or neonatal morbidity among the three groups. A retrospective study by Hood et al³ and a prospective study by sharwood-smith et al⁴ agreed with this findings.

In this study it had been shown that changes in both systolic and diastolic blood pressure were significantly higher (P<0.05) in group B patient than group A. So, pressor agent ephedrine was more used in group B patients. The result was similar to the retrospective study be Hood and Boese¹¹. Although in majority of the patients in group B systolic blood pressure fall below 100 mmHg

,however in no case this was below 80 mmHg and accordingly,none required rescue treatment. As such the present protocol did not exclude any patient from the study.

Adequate pre loading decreases the chance of hypotension. In this study preloading with crystalloid between group A & group B was not significant ($P= .421$) but in group B maternal systolic arterial pressure decreased significantly. This finding is consistent with karinen J et al study¹².

Pressor agent epinephrine should not be used in pre eclamptic patient because pre eclamptic has a markedly increased sensitivity to vasopressors^{10,13} and in advertent intravascular injection of epinephrine would further exacerbate maternal hypertension and further decreased placental blood flow¹⁴. So, the most commonly used vasopressor in obstetric anaesthesia are the predominantly b agonist drugs. Ephedrine is widely used because of its predominantly b and mild a sympathomimetic action. Ephedrine increases cardiac output & therefore flow to the maximally dilated utero placental vessels¹⁵. Incremental I/V bolus dose of ephedrine counteract hypotension. In this study it had been shown that frequency of systolic hypotension were more in-group B patients which was managed well by I/V ephedrine.

Karinen J. et. al have studied the neonates born from pre eclamptic patient under spinal anaesthesia. They did not find any major effect on clinical condition of the neonates assessed by Apgar score & umbilical artery p^H values¹². This study did not match with Karinen J. et. al study. In this study APGAR score is significantly low in group B patient both in 1st min and 5th min than group A patient. In both 1st and 5th min $P<0.05$. This study matched with the research work of Monika Sharma who found that APGAR was more than 7 in 68.75% of babies and NICU admission and foetal loss were significantly less in study group. She concluded that magnesium sulfate is safe for mother and has no adverse effect on babies.

In this study duration of motor and sensory block in Group B was increased and was statistically highly significant ($P<0.05$). Onset time of sensory block – Tingling in the leg was more in-group –B patient and was statistically significant ($P<0.05$) but in case of pinprick it was not significant ($P=0.084$).

Onset time of motor block – weakness of lower limb was more in-group –B patients and was statistically significant ($P=0.019$). On the other hand onset of motor block complete paralysis of lower limb in-group –B patients and was not significant ($P<0.06$).

So study concluded that spinal anaesthesia is safe for pre eclamptic parturient, treated with magnesium sulphate. Although chance of hypotension is more in patients getting magnesium sulfate but it may be managed with adequate preloading and by pressor agent ephedrine.. APGAR score of baby of magnesium sulfate getting patient is low but it is acceptable.

References

1. The Eclampsia Trial Collaborative Group. Which anticonvulsant for women with preeclampsia? Evidence from the collaborative Eclampsia Trial. *Lancet* 1995; 345: 1455-63
2. Wallace DH, Leveno KJ, Cunningham FG, Giesecke AH, Shearer VE, Sidawi JE. Randomized comparison of general and regional anaesthesia for Cesarean delivery in pregnancies complicated by severe preeclampsia. *Obstet Gynecol* 1995; 86: 193-9
3. Hood DD, Curry R. Spinal versus epidural anesthesia for cesarean section in severely preeclamptic patients. *Anesthesiology* 1999; 90: 1276-82
4. Sharewood-Smith G, Clark V, Watson E. Regional anesthesia for caesarean section in severe preeclampsia: spinal anesthesia is the preferred choice. *Int J Obster Anes* 1999; 8: 85-9
5. Writer D. Hypertensive disorders is chestnut D'ed, obstetric, anaesthesia principles & practice' st' Louis Mosby year book Inc, 1994: 871-2.
6. Less MM, Scot DB, et al: Haemodynamic changes associated with labour. *Journal of Obstetrics and Gynaecology of the British common wealth* 1970; 77: 29-36.
7. Brock-Utne JG, Downing JW, Seedat F. laryngeal oedema associated with pre eclamptic toxemia. *Anaesthesia* 1977; 32: 556-8.

8. Douglas KA, Redman CWG. Eclampsia in the United Kingdom. *BMJ* 1994; 309: 1395-4000.
9. Scardo JA, Hogg BB, Newman RB. Favorable hemodynamic effects of magnesium sulfate in preeclampsia. *Am J Obstet Gynecol* 1995; 173: 1249-53
10. Gutsche BB. Anesthetic considerations for preeclampsia Eclampsia. In : Shnider SM, Levinson G, eds. *Anesthesia for obstetrics*. Baltimore : Williams & Wilkins, 1979; 224-34
11. HOOD DD, Boesc PA Epidural & Spinal anaesthesia for caesarean section in severely preclamptic patients (abstract) *Region of Anaesthesia* 1992;17;35
12. Karinen J. Maternal and uterine haemodynamic state in pre-eclamptic patients during spinal anaesthesia for caesarean section. *British Journal of Anaesthesia* 1996; 76: 616-620
13. Marx GF, Hodgkinson R. Special considerations in complications of pregnancy. In : Marx GF, Bassel GM, eds. *Obstetric analgesia and anesthesia*. New York : Elsevier-North- Holland Biomedical Press, 1980; 297-334
14. Crawford JS. Epidural analgesia in pregnancy hypertension. *Clin Obstet Gynaecol* 1977; 4: 735-44
15. Ralston DH, Shnider SM, deLorimier AA : Effects of equipotent ephedrine, metaraminol, mephenteramine and methoxamine on uterine blood flow in pregnant ewe. *Anesthesiology* 40: 354, 1974
16. Volpe P, Vezu L. Intracellular magnesium and inositol 1,4,5-triphosphosphate receptor: molecular mechanisms of interaction, physiology and pharmacology. *Magnes Res* 1993; 6: 267-74
17. Altura BM, Altura BT. Magnesium ions and contraction of vascular smooth muscles; relationship to some vascular diseases. *Fed Proc* 1981; 40: 2672-9
18. Connell H, Dalgeish JG, Downing JW. General Anaesthesia in mothers with severe pre-eclampsia/ eclampsia. *Br J Anaesth* 1987; 59:1375-80
19. Douglas WW, Rubin RP. The mechanism of catecholamine release from the adrenal medulla and the role calcium in stimulus-secretion coupling. *J Physiol (Lond)* 1963; 167:288-310
20. Jenkinson DH. The nature of the antagonism between calcium and magnesium ions at the neuromuscular junction. *J Physiol (Lond)* 1957; 138: 434-44
21. Jouppila P, Kuikka J, Jouppila R, Hollman A , Effect of induction of general anaesthesia for caesarean section on intervillous blood flow. *Acta Obstetrica et Gynecologica Scandinavica* 1979; 58: 249-253