

COMPARISON OF HAEMODYNAMIC PROFILE AFTER SPINAL ANAESTHESIA IN PATIENTS ON REGULAR TREATMENT WITH CALCIUM CHANNEL BLOCKERS AND β BLOCKERS

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ABSTRACT

BACKGROUND

Hypertensive patients develop wide swings in blood pressure intraoperatively, especially after spinal anaesthesia. Long term antihypertensive agents can modify this effect by controlling blood pressure. This study was undertaken to evaluate the haemodynamic effect in hypertensive patients on regular treatment with calcium channel blockers and beta-blockers who are undergoing elective surgery under spinal anaesthesia and compared with normotensives.

AIMS

To study the intraoperative changes of blood pressure and heart rate in patients on calcium channel blockers and beta-blockers undergoing surgery under spinal anaesthesia and to compare the data with normotensive patients.

MATERIALS AND METHODS

90 patients were included in the study; 30 patients were normotensives (Group N) and 30 patients were hypertensives who were regularly on calcium channel blockers (Group C) and 30 patients on beta-blockers (Group B). Both the groups continued the drug on the day of surgery. The baseline blood pressure and heart rate were recorded. After spinal anaesthesia, the blood pressure and heart rate were noted at 2 min, 4 min, 6 min, 8 min, 10 min, 15 min and thereafter with 5 minutes interval till the end of the procedure.

STATISTICAL ANALYSIS

Baseline data was analysed using descriptive statistics. Intragroup variation in parameters analysed by one way ANOVA and within the different time intervals by post-hoc test. Intergroup comparison by unpaired "T" test. P value <0.05 considered statistically significant.

RESULTS

The hypotension was evident in Group C compared to other groups, but bradycardia (heart rate <60) more seen in Group B and were statistically significant (P<0.001). The usage of rescue medication was more in study group and was statistically significant (P<0.002).

CONCLUSION

Incidence of hypotension was more with patients on calcium channel blockers and required vasopressors and fluids, but incidence of bradycardia was seen more in patients on beta blockers who needed atropine. Anaesthesiologists should anticipate and be adequately prepared for any untoward consequences.

KEYWORDS

Hypertension, Spinal Anaesthesia, Calcium Channel Blockers, Hypotension, Beta Blockers, Bradycardia.

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INTRODUCTION

Spinal anaesthesia is the most commonly used procedure for lower abdominal and limb surgeries and common side effects being hypotension and bradycardia due to sympathetic denervation after spinal anaesthesia. Reason being maximal venodilatation causing peripheral pooling and decreased preload to the heart, however, smooth muscles in arteries maintain autonomic tone after spinal also.

Hypertensive patients can develop wide swings in blood pressure intraoperatively, which increases the risk of post-operative cardiac and renal complications such as myocardial ischemia, cerebrovascular accidents and acute renal failure, independent risk factors.^{1,2} were identified which are necessary for the prevention of perioperative cardiac events causing both morbidity and mortality. Structural changes in arteriolar walls play a primary role in haemodynamic response to anaesthesia and explain greater changes in systemic vascular resistance and arterial pressure in hypertensive patients than normotensive patients with similar degree of sympathetic blockade.³

Studies have been done to determine whether it is beneficial to continue Calcium channel blockers and Beta blockers.⁴ in patients undergoing spinal anaesthesia. Effect of continuation of angiotensinogen converting enzyme inhibitors (ACE I).⁵ in patients undergoing spinal anaesthesia and found

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that continuation of ACE I did not affect early hypotension severity compared to normal individuals. There is inadequate evidence on the effects of Calcium channel blockers and Beta blockers on Blood pressure in hypertensive patients undergoing Spinal anaesthesia. Because of end organ changes in hypertensive patients, post spinal hypotension could have detrimental effect.

So this study was an effort to compare the incidence of hypotension and bradycardia in hypertensive patients on regular medication with standardized doses of calcium channel blockers and beta-blockers with that in normotensive patients after spinal anaesthesia.

MATERIALS AND METHODS

The study was conducted on 90 patients. Group N (Normotensive group), Group C (Calcium channel blockers group), Group B (Beta blocker group) undergoing elective surgeries under spinal anaesthesia at Chigateri Government Hospital and Bapuji Hospital attached to JJM Medical College, Davangere, from 2013 to 2014.

Inclusion Criteria

Patients undergoing elective surgeries under spinal anaesthesia who are between 30 and 60 years old (ASA 1 and ASA 2) and diagnosed cases of essential hypertension who are on anti-hypertensive medications at least 1 month prior to surgery.

Exclusion Criteria

1. Patients having contraindications for spinal anaesthesia.
2. Patients with other co-existing diseases like diabetes, coronary artery disease or other cardiac diseases, severe hypovolemia, sepsis, pregnant patients.

The Institutional Ethical clearance was obtained before beginning the study. Informed written consent about procedure was obtained prior. During preanaesthetic evaluation, patients were assessed and explained about the anaesthetic procedure and perioperative monitoring. History regarding anti-hypertensive medications, duration of treatment, other co-existing diseases and any other medications were taken. In operating room, after inserting the 18G IV cannula patients were coloaded with 15 mL/kg of Ringer Lactate (RL) solution, maintenance fluid continued throughout procedure. Total volume infused were recorded. Standard monitoring included continuous ECG, pulse rate, oxygen saturation with a pulseoximeter and automated Non-Invasive Blood Pressure (NIBP) Systolic (SBP) and Diastolic (DBP) and Mean Blood Pressures.

Baseline values were taken Spinal anaesthesia was given in the left lateral position at L3-L4 interspace with all the aseptic precautions using a 25-gauge Quincke-Babcock spinal needle. After noticing a free flow of the cerebrospinal fluid, 3 mL of 0.5% hyperbaric bupivacaine was administered over a 10 second period. The patient was put in a supine position. The level of sensory blockade was determined by a pin-prick at 5 min and 15 min after the drug administration. NIBP and pulse rate were recorded every 2 min for the first 10 min and every 5 min till the end of the surgical procedure.

Hypotension was defined as a decrease of mean arterial pressure of more than 20% from baseline and was intervened with fluids or vasopressors within 20 min. Mephentermine 6

mg/dose and IV fluids 5 mL/kg were used for Hypotension until both Systolic Arterial Pressure (SAP) and Mean Arterial Pressure (MAP) increased above the threshold level. The time of rescue dose and the total dose for hypotension was recorded.

Bradycardia was defined as heart rate <60 beats/min and was treated with Inj. atropine 0.6 mg if not responding to intravenous fluids.

STATISTICAL ANALYSIS

Baseline data was analysed using descriptive statistics. The changes of systolic blood pressure, diastolic blood pressure, mean arterial pressure and heart rate were analysed as follows:

1. The changes of the parameter within the group was analysed using one way ANOVA and the variation within the different time intervals was analysed with the post-hoc tests.
2. Intergroup comparison at the 2 min, 4 min, 6 min, 8 min, 10 min and 15 minutes from the data of controls and cases respectively was done by unpaired 't' test.

'P' value <0.05 was considered significant and <0.01 highly significant.

RESULTS

Demographic Data

The demographic characteristics of both the groups were comparable as shown in Table 1.

Average Sensory Block

The average height of the block after spinal anaesthesia was at the 7th to 8th thoracic level Table 2.

Baseline Haemodynamics (Table 3)

The average baseline systolic blood pressure was 134.53±5.42 mmHg in group C and 135.93±4.91 mmHg in group B, which was on an average 6.5 mmHg and 8 mmHg higher in the hypertensive group C and group B respectively than the normotensive group. The baseline diastolic blood pressure was 82.93±6.54 mmHg in group C and 80.73±9.23 in group B, which was on an average 6.86 mmHg and 4.66 mmHg higher in the hypertensive group C and group B respectively than the normotensive group. The baseline mean blood pressure was 100.18±5.95 mmHg in group C and 99.27±7.41 in group B which was on an average 6.8 mmHg and 5.87 mmHg higher in the hypertensive group C and group B respectively than the normotensive group. The baseline heart rate was almost the same in normotensive group N and the hypertensive group C (70 bpm) and it was on lower side in the hypertensive group B (67 bpm).

Systolic blood pressure after spinal Anaesthesia (Table 4, Table 5, Fig. 1): Hypotension was observed in all groups, statistically significant fall in systolic blood pressure occurred from 4 minutes after spinal up to 15 minutes. Fall from baseline SBP (Table 4) in control group at 4, 6, 8, 10, 15 minutes were 16±8.2 (mmHg), 20.6±10.3 (mmHg), 20±11 (mmHg), 21±10 (mmHg), 17.8±10.4 (mmHg). In group C at 4, 6, 8, 10, 15 minutes were 27±15 mmHg, 30.27±17.28 mmHg, 28±18 mmHg, 31±18 mmHg, 27±16.95 mmHg (P<0.001) statistically significant fall in systolic blood pressure occurred compared to control group and group B. Fall in SBP was

observed in group B at 4, 6, 8, 10, 15 minutes, but not significant from control group (Table 5).

Diastolic blood pressure and mean blood pressure (Table 6, Table 7, Fig. 2): Statistically significant fall in Diastolic Blood Pressure in group C at 4, 6, 8, 10, 15 minutes compared to other two groups were 21.2±4.5, 23.3±4.1, 22.6±4.18, 23.4±6.7, 22.7±6.7 mmHg from the baseline (P<0.001) (Table 6 and 7). Changes in mean arterial pressures showing significant fall at 4, 6, 8, 10,15 minutes observed more in group C (Table 8 and 9) Fig. 3, 14/30 patients in group C required Mephentermine for hypotension (Table 11).

Heart rate variation: Fall in heart rate observed significantly in group B compared to other groups at 2, 4, 6, 8, 10, 15 minutes were 5.2±2.4, 6.5±1.8, 7.7±1.7, 10.2±0.58, 10.4±0.81, 10.5±0.8 rate/minute from the baseline (Table 10), 16/30 of patients in group B required atropine for bradycardia (Table 11).

Variables		Group N N=30	Group C N=30	Group B N=30
Age (Mean & SD)		47.9±10.27	49.76±8.93	55.1±7.87
Age Categories	< 50 yrs.	10	16	9
	51 - 60	8	11	15
	61 & Above	12	3	6
Sex	Male	21	21	19
	Female	9	9	11
Group N= Normotensive group, C= Calcium channel blocker group, B= Beta blocker group Data expressed by Mean Standard deviation and Number				

Table 1: Demographic Data

Variables		Group N N=30	Group C N=30	Group B N=30
Block height	T ₆	10	8	4
	T ₈	14	12	4
	T ₁₀	6	10	22
Average Level - T7 to T8				

Table 2: Sensory Levels in Different Groups

Variables	Group N N=30	Group C N=30	Group B N=30
Baseline SAP (mmHg)	128.06±5.41	134.53±5.42	135.93±4.91
Baseline DAP (mmHg)	76.07±7.03	82.93±6.54	80.73±9.23
Baseline MAP (mmHg)	93.38±4.83	100.18±5.95	99.27±7.41
Baseline HR (beats/min)	75.0±5.47	72.46±5.81	67.53±8.87
Data expressed as Mean±SD			

Table 3: Baseline Haemodynamics

	Group N		Group C		Group B		Statistical Analysis	
	Mean Difference	Std. Deviation (SD)	Mean Difference	Std. Deviation	Mean Difference	Std. Deviation	One Way ANOVA	Significance P
Base & 0	4.66	5.4	2.6	5.63	4.5	5.7	1.24	Not Sig
Base & 2 min	10.3	9.12	13.1	10.7	8.87	5.78	1.77	Not Sig
Base & 4 min	16	8.2	27	15	15	16	7.04	P<0.001
Base & 6 min	20.6	10.3	30.27	17.58	20.4	9.5	5.66	P<0.005
Base & 8 min	20	11	29	18	20	11	3.77	P<0.02
Base & 10 min	21	10	31	18	21	9.9	5.60	P<0.005
Base & 15 min	17.8	10.4	27	16.95	23.3	6.8	2.23	Not Sig
Decrease in SBP after SA in each group is expressed as Mean fall ±SD and 'P' value								

Table 4: Systolic Blood Pressure (SBP) Change After Spinal Anaesthesia (SA)

Bonferroni Multiple Comparisons			
	Normotensive Vs Calcium Channel	Normotensive Vs Beta Blocker	Calcium Channel Vs Beta Blocker
Base & 0	0.60, NS	1.0, NS	0.46, NS
Base & 2 min	0.23, NS	1.0, NS	0.66, NS
Base & 4 min	P<0.004	1.0, NS	P<0.006
Base & 6 min	P<0.01	1.0, NS	P<0.04
Base & 8 min	0.07, NS	1.0, NS	P<0.04
Base & 10 min	P, 0.01	1.0, NS	P<0.01
Base & 15 min	1, NS	0.23, NS	0.19, NS
'P' value significant between Group N and Group C & Group B and Group C			

Table 5: Intergroup Comparison of 'P' Values in SBP

	Group N		Group C		Group B		Statistical Analysis	
	Mean Difference	Std Deviation	Mean Difference	Std Deviation	Mean Difference	Std Deviation	One Way ANOVA	Significance
Base & 0	1.5	0.4	3	2	2.1	1.3	.58	Not Sig
Base & 2 min	4.6	0.48	11.27	0.4	8.33	0.48	7.04	P<0.001
Base & 4 min	8.93	1.1	21.2	4.5	13.6	2.3	10.31	P<0.000
Base & 6 min	14.5	4.9	23.3	4.1	14.7	2.8	4.94	P<0.009
Base & 8 min	14.5	4.9	22.6	4.18	16.4	1.79	4.13	P<0.01
Base & 10 min	13.47	2.88	23.4	6.37	17.13	0.21	7.39	P<0.001
Base & 15 min	13.42	2.78	22.7	6.17	17.03	0.2	7.22	P<0.01

Drop in DBP is expressed as Mean±SD and statistically significant throughout

Table 6: Diastolic Blood Pressure (DBP) Change After Spinal Anaesthesia (SA)

Bonferroni Multiple Comparisons			
	Normotensive Vs Calcium Channel	Normotensive Vs Beta Blocker	Calcium Channel Vs Beta Blocker
Base & 0	NS	NS	NS
Base & 2 min	P<0.02	NS	P<0.001
Base & 4 min	P<0.02	NS	P<0.000
Base & 6 min	P<0.02	NS	P<0.02
Base & 8 min	P<0.02	NS	P<0.02
Base & 10 min	P<0.05	NS	P<0.001
Base & 15 min	P<0.05	NS	P<0.001

P value is insignificant between Group N and Group B and Group C and Group B

Table 7: Intergroup Comparison of 'P' Values in DBP

	Group N		Group C		Group B		Statistical Analysis	
	Mean Difference	Std. Deviation	Mean Difference	Std. Deviation	Mean Difference	Std. Deviation	One Way ANOVA	Significance
Base & 0	3.7	0.9	2.8	1	3.1	1.1	.36	Not Sig
Base & 2 min	7.82	0.63	11.94	1.06	8.6	0.75	3.54	P<0.03
Base & 4 min	7.83	0.8	23.11	6.3	14.4	3.18	10.11	P<0.000
Base & 6 min	11.9	0.87	25.73	6.36	16.74	2.35	5.31	P<0.007
Base & 8 min	18.32	5.38	25	6	18	3	2.98	P<0.05
Base & 10 min	18	6	26	8	18	1	6.97	P<0.002
Base & 15 min	17.9	6	25.83	6	17.8	1	6.96	P<0.002

Drop in MAP is expressed as mean±SD and p value is significant throughout

Table 8: Mean Blood Pressure (MAP) Change After Spinal Anaesthesia

Bonferroni Multiple Comparisons			
	Normotensive Vs Calcium Channel	Normotensive Vs Beta Blocker	Calcium Channel Vs Beta Blocker
Base & 0	Ns	NS	Ns
Base & 2 min		NS	P<0.02
Base & 4 min	P<0.004	P<0.02	P<0.000
Base & 6 min	P<0.009	NS	P<0.04
Base & 8 min	P<0.009	NS	NS
Base & 10 min	P<0.01	NS	P<0.02
Base & 15 min	P<0.01	NS	P<0.02

Table 9: Intergroup Comparison of 'P' Values for MAP

	Group N		Group C		Group B		Statistical Analysis	
	Mean Difference	Std. Deviation	Mean Difference	Std. Deviation	Mean Difference	Std. Deviation	One Way ANOVA	Significance
Base & 0	0.4	1.4	0.2	0.1	3.53	2.28	7.53	P<0.001
Base & 2 min	1.93	1.9	0.73	0.1	5.2	2.4	6.15	P<0.003
Base & 4 min	2.66	1.28	1	0.14	6.5	1.8	10.98	P<0.000
Base & 6 min	3.2	2	2.3	0.82	7.7	1.7	9.20	P<0.000
Base & 8 min	2.4	2	3.2	0.9	10.2	0.58	14.16	P<0.000
Base & 10 min	2.5	2.3	2.67	0.52	10.4	0.81	17.05	P<0.000
Base & 15 min	2.4	2.2	2.7	0.54	10.5	0.8	16.84	P<0.000

Table 10: Pulse Rate (PR) Changes After Spinal Anaesthesia

Bonferroni Multiple Comparisons			
	Normotensive Vs Calcium Channel	Normotensive Vs Beta Blocker	Calcium Channel Vs Beta Blocker
Base & 0	P<0.005	P<0.003	P<0.005
Base & 2 min	NS	P<0.04	P<0.003
Base & 4 min	NS	P<0.006	P<0.000
Base & 6 min	NS	P<0.004	P<0.000
Base & 8 min	NS	P<0.000	P<0.000
Base & 10 min	NS	P<0.000	P<0.000
Base & 15 min	NS	P<0.000	P<0.000
'P' value is significant between Group N & Group B and also between Group C and Group B			
Table 11: Intergroup Comparison of P Values for PR			

Variables		Group N N=30	Group C N=30	Group B N=30
Hypotension	Present	12	18 [†]	10
	Absent	18	12	20
Mephentermine	Required	4	14 [†]	4
	Not required	26	16	26
Bradycardia	Present	0	0	16 [†]
	Absent	30	30	14
Atropine	Required	0	0	16 [†]
Data represented as absolute numbers, important one is indicated by †				
Table 12: Intergroup Incidence of Side Effects (Hypotension, Bradycardia) and Need of Treatment				

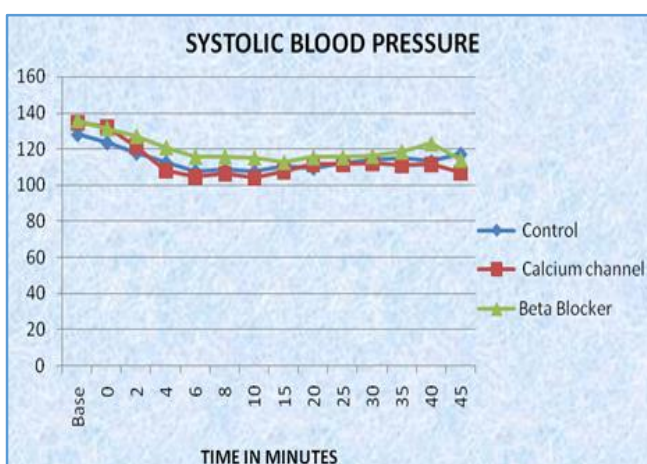


Fig. 1: Systolic Blood Pressure Changes Overtime in Different Study Groups

X-axis – duration, Y axis – systolic blood pressure in mmHg. The maximum fall of SBP is seen at 8 min in both the groups.

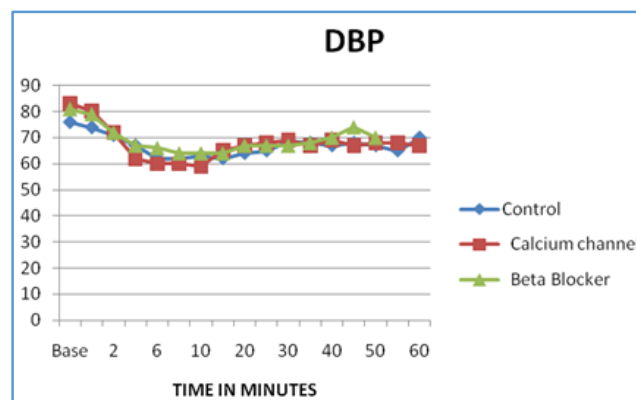


Fig. 2: Diastolic Blood Pressure Changes

X-axis – Time, Y axis – DBP in mmHg. The maximum fall of DBP is seen at 8-10 min in both the groups.

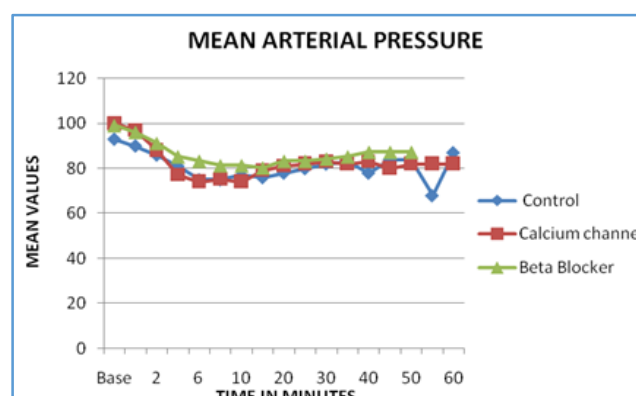


Fig. 3: Mean Arterial Pressure Changes

X-axis –time, Y axis –mean arterial pressure in mmHg. The maximum fall of MAP is seen at 8 min in both the groups.

DISCUSSION

Hypotension is the most frequent side effect associated with spinal anaesthesia, predisposing the individual to myocardial and brain ischaemia. Identification of risk factors for prediction of early hypotension were age, female sex, body mass index >30 kg/m², history of hypertension, diabetes mellitus, anaemia, high baseline heart rate, high systolic and diastolic blood pressure, pulse pressure, sensory level of blockade higher than or equal to T6. Association of hypotension was more in patients with h/o hypertension (OR -1.739) and incidence reduced in patients on anti-hypertensives (1.012).¹

Risk factors must be easily obtainable and known before the blockade to meet their preventive aim.² Alek Rook et al³ found out that an exaggerated decrease in blood pressure occurs in elderly patients with cardiac disease and hypotension was mainly a result of decrease in systemic vascular resistance. Hence, the knowledge of risk factors for exaggerated hypotension after spinal anaesthesia may help the anaesthesiologist in adopting preventive measures to minimize the catastrophe. We chose and compared the haemodynamic changes in hypertensive patients without any other cardiac morbidity on antihypertensive drugs for >1-month duration with that of normotensives.

Our study compared standardized doses of calcium channel blocker (Amlodipine 10 mg) (Group C), B blocker (Atenolol 50 mg) (Group B) on Haemodynamics after spinal anaesthesia with normotensive patients since the effect on cardiac and vasculature is dose dependent and previous study⁴ had shown effect of different doses of antihypertensives within the group on haemodynamics. But our study observations also correlated well with this study, which showed that hypotension is more common in patients who are on CCBs (Group C) (18/30) and 80% of them required fluids and vasopressors more than those on B blocker or normal patients, effect could be due to peripheral vasodilatation already existing in pts on CCBs and could have exaggerated by decreased vascular resistance after spinal anaesthesia.

Studies have been done on effect of B blocker and CCB during induction of general anaesthesia.⁶, effect of the same on laryngoscopy and intubation.⁵ hypotension during induction is primarily due to decrease in SVR, which is also one of the primary mechanism with sympathetic blockade after spinal and effect of ACE I on post spinal blood pressure.⁷ and found hypotension >25% can be detrimental to vital organ circulation.

Mojica JL et al⁸ have studied regarding the timing of IV fluids during spinal anaesthesia and found coloadng (IV fluids during spinal anaesthesia) is more advantageous than preloading in prevention of severe hypotension or Arndt JO et al⁹ compared effectiveness of fluids and prophylactic vasopressors in prevention of severe hypotension. We used 15 mL/kg of Ringer's lactate to reason out hypotension was mainly the effect of vasodilatation after spinal and precipitated effect of antihypertensives and not due to absence of coloadng.

Parida S.¹⁰ described in case series regarding the severity of Hypotension following spinal anaesthesia in patients on amlodipine 10 mg even after withholding morning dose and preloading before spinal, our study correlated with the findings of the series but degree of hypotension was less probably because study group involved young patients also where the vasomotor tone is preserved compared to older individuals.¹¹

Structural changes (Medial hyperplasia and hypertrophy) in arteriolar walls play a primary role in haemodynamic response to anaesthesia and explain greater changes in systemic vascular resistance and arterial pressure in hypertensive patients than normotensive patients with similar degree of sympathetic blockade.¹² In Singla et al¹ study there were patients not taking any treatment for hypertension, they all fell in hypotensive group. Regular antihypertensive treatment was associated with reduced risk of hypotension. Prolonged antihypertensive treatment may induce regression of the structural changes in arterioles and consequently a more normal functional response to vasodilatation or constriction.

Vukalic J.¹³ et al studied elderly hypertensives on B blocker medication and found continuation of B blockers in the pre-operative period was associated with hypotension, bradycardia requiring atropine and nausea after spinal anaesthesia compared to other group which discontinued the drug, so they concluded discontinuation of B blockers in pre-operative period is prudent, but other studies.¹⁴ however found that cessation of beta blocker therapy leads to rebound

hypertension, arrhythmias and this followed by various cardiovascular consequences, it is beneficial to continue the B blockers pre-operatively. Our study correlated with the above study except lesser degree of hypotension, which could be due to inclusion of young individuals in the study even after continuation of B blockers, but bradycardia was significant (16/30) requiring atropine in all patients.

CONCLUSION

Our study showed exaggerated hypotension in patients on calcium channel blockers, which could be partly explained by the vasodilator property of the drug which could have additive effect after spinal anaesthesia inducing exaggerated hypotension. This exaggerated fall in blood pressure after spinal anaesthesia warrants the discontinuation of calcium channel blockers before surgery. Bradycardia following spinal anaesthesia was entirely limited to the beta blocker group in our study. The knowledge of these risk factors could be useful in increasing vigilance in those patients most at risk for hypotension, in allowing timely therapeutic intervention and taking preventive measure to minimize the spinal hypotension.

REFERENCES

1. Singla D, Kathuria S, Singh A, et al. Risk factors for development of early hypotension during spinal anaesthesia. *J Anaesthesiol Clin Pharmacol* 2006;22(4):387-93.
2. Hartmann B, Junger A, Klasen J, et al. The incidence and risk factors for hypotension after spinal anaesthesia induction: an analysis with automated data collection. *Anaesth Analg* 2002;94(6):1521-9.
3. Rooke GA, Freund PR, Jacobson AF. Haemodynamic response and change in organ blood volume during spinal anaesthesia in elderly men with cardiac disease. *Anaesth Analg* 1997;85(1):99-105.
4. Kaimar P, Sanji N, Upadya M, et al. A comparison of hypotension and bradycardia following spinal anaesthesia in patients on calcium channel blockers and beta-blockers. *Indian journal of pharmacology* 2012;44(2):193-6.
5. Swear JW, Jewkes C, Tellez JC, et al. Does the choice of antihypertensive therapy influence haemodynamic responses to induction, laryngoscopy and intubation. *British Journal of Anaesthesia* 1994;73(3):303-8.
6. Samad K, Khan F, Azam I. Haemodynamic effects of anaesthetic induction in patients treated with beta and calcium channel blockers. *Middle East J Anaesthesiol* 2008;19(5):1111-28.
7. Hohne C, Meier L, Boemke W, et al. Angiotensin convertase inhibitor inhibitors do not exaggerate the blood pressure decrease in the early phase of spinal anaesthesia. *Acta Anaesthesiologica Scandinavica* 2003;47(7):891-6.
8. Mojica JL, Meléndez HJ, Bautista LE. The timing of intravenous crystalloid administration and incidence of cardiovascular side effects during spinal anaesthesia: the results from a randomized controlled trial. *Anaesth Analg* 2002;94(2):432-7.
9. Arndt JO, Bömer W, Krauth J, et al. Incidence and time course of cardiovascular side effects during spinal anaesthesia after prophylactic administration of

- intravenous fluids or vasoconstrictors. *Anaesth Analg* 1998;87(2):347-54.
10. Parida S, Nawaz M, Kundra P. Severe hypotension following spinal anaesthesia in patients on amlodipine. *J Anaesthesiol Clin Pharmacol* 2012;28(3):408-9.
 11. Alecu C, Cuignet-Royer E, Mertes PM, et al. Pre-existing arterial stiffness can predict hypotension during induction of anaesthesia in the elderly. *Br J Anaesth* 2010;105(5):583-8.
 12. Folkow B. Cardiovascular structural adaptation : its role in the initiation and maintenance of primary hypertension. *Clin Sci* 1978;55:3s-22s.
 13. Vukalic J, Rakaric MP. Adverse effects of beta adrenergic blockers during spinal anaesthesia with 0.5% bupivacaine. *Crit Care* 2010;14(Suppl 1):P480.
 14. Ponten J, Biber B, Bjuro T, et al. Beta receptor blockade and spinal anaesthesia. Withdrawal versus continuation of long term therapy. *Acta Anaesthesiol Scand Suppl* 1982;76:62-9.