



A Comparative Study of the Effect of Fentanyl 25 MCG with Bupivacaine 0.5% Verses Buprenorphine 60 MCG with Bupivacaine 0.5% in Spinal Anaesthesia for Elective Caesarean Section

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Abstract

Introduction: *Pregnancy is accompanied by physiologic changes in multiple organ systems that may influence maternal responses to anaesthesia and the choice of anaesthetic techniques. The rate of caesarean sections is on the rise. Caesarean section in a conscious patient is a challenging test of regional anaesthesia. Spinal anaesthesia is perhaps the most efficient approach to this challenge. In modern scientific era, there has been a curiosity in using opioid analgesic adjuvants to subarachnoid local anaesthetics. Arrival of a new synthetic lipophilic opioid, fentanyl, has revolutionized its use in the past three decades. Fentanyl has a shorter duration of action and fast onset as compared to pethidine and morphine. Buprenorphine is a long acting, highly lipophilic opioid. It has proved to be a promising analgesic by intra thecal route^{1,2}. Buprenorphine is twenty five times more potent than morphine. In this study, an effort is made to compare the perioperative and postoperative analgesic efficiency of these two lipophilic opioid drugs along with bupivacaine in caesarean section.*

Aim: *To evaluate the efficacy of the combination of intrathecal fentanyl 25 mcg and 7.5 mg of 0.5% hyperbaric bupivacaine in comparison with buprenorphine 60 mcg and 7.5 mg of 0.5% hyperbaric bupivacaine used for lower segment caesarean section*

Materials and Methods: *A total of 50 patients who underwent elective caesarean section were taken up for the study. Patients were randomised into two groups each. In group A, patients received 1.5 ml of 0.5% hyperbaric bupivacaine (7.5mg) with 0.5ml of fentanyl (25 mcg) and in group B, patients received 1.5 ml of 0.5% hyperbaric bupivacaine (7.5mg) with 0.2 ml of buprenorphine (60 mcg) and 0.3 ml of normal saline. The final volume of the injected solution is 2 ml on both groups.*

Results: *In the current study, onset of analgesia was significantly earlier due to the addition of buprenorphine. This may be attributed to high lipid solubility and highest affinity for opiate receptors of buprenorphine. Both the groups had the same mean time to achieve motor blockade. Both groups maintained hemodynamic stability which was statistically insignificant. The mean duration of effective analgesia was 200.32 minutes (3.33 hours) in Group A and 491.28 minutes (8.1 hours) in group B which was highly significant statistically(p<0.01).*

Conclusion: *We observed that anaesthesia was superior when buprenorphine is mixed with bupivacaine (0.5%) as compared to bupivacaine with fentanyl. Addition of buprenorphine to bupivacaine 0.5% augments the sensory blockade of local anaesthetics without affecting the sympathetic activity. Thus it is concluded that intrathecal buprenorphine is suitable drug for post operative analgesia for caesarean section .*

Keywords: *Caesarean section, Spinal anaesthesia, bupivacaine, fentanyl, buprenorphine, post operative analgesia.*

Introduction

Pregnancy is accompanied by physiologic changes in multiple organ systems that may influence maternal responses to anaesthesia and the choice of anaesthetic techniques. According to a recent analysis of national health data, the rates of average caesarean sections done in our country has gone up from five percent to 18 percent over the last few decades.

Caesarean section in a conscious patient is a challenging analysis of regional anaesthesia. The most efficient approach to this challenge is spinal anaesthesia. The advantages are

1. Small needle
2. Profound anaesthesia,
3. Minimal amount of drug and
4. Excellent operating conditions, that can be readily provided for the major surgeries like LSCS.

Hypotension following spinal anaesthesia is the most clinically significant aspect, that can occur rapidly and may have a significant aspect on the neonatal outcome. In the last few decades, there has been a curiosity in using opioid analgesic adjuvants to subarachnoid local anaesthetics to decrease the local anaesthetic dose, hence reducing the occurrence and degree of hypotension, at the same time without compromising intra operative analgesia and to enable faster recovery also providing efficient post operative analgesia.

The discovery of opioid receptors in the spinal cord and intrathecal opioid administration has opened a new horizon in pain management during perioperative period, and it has gained significance in the past three decades.

Arrival of a new synthetic lipophilic opioid, fentanyl, has revolutionized its use in the past three decades. Fentanyl has a shorter duration of action and fast onset as compared to pethidine and morphine. It acts as an agonist to mu receptors. Fentanyl is more specific, shorter acting and about hundred times more potent than morphine. It is less hydrophilic and has little rostral spread which

causes lesser respiratory depression when compared with morphine.

Buprenorphine is an opioid with high lipophilic property and longer duration of action. It has proved to be a favourable analgesic by intrathecal route^{1,2}. Buprenorphine is twenty five times more potent than morphine. Buprenorphine is an agonist – antagonist with lipid solubility about five times greater than that of morphine and has a low level of physical dependence³. Buprenorphine is associated with lower incidence of respiratory depression because there is no rostral spread. But it has been associated with urinary retention⁴.

In the study, an effort is made to compare the perioperative and postoperative analgesic efficiency of these two lipophilic opioid drugs along with bupivacaine in caesarean section.

Aim

In this context, the present study was undertaken to evaluate the efficacy of the combination of intrathecal fentanyl 25 mcg and 7.5 mg of 0.5% hyperbaric bupivacaine in comparison with buprenorphine 60 mcg and 7.5 mg of 0.5% hyperbaric bupivacaine used for lower segment caesarean section with respect to

1. Time of onset of analgesia and motor blockade
2. Duration of sensory and motor block
3. Quality of intra operative anaesthesia
4. Incidence of hypotension
5. Ephedrine requirement to combat hypotension
6. Foetal outcome
7. Duration of post operative analgesia

Materials and Methods

After getting proper concurrence from ethics committee, The present study was conducted in Rajah Muthiah medical college and hospital, Annamalai university. A total of 50 patients who underwent elective caesarean section were selected for the study. The age of the patients ranged from 20-37 years weighing 40-65 kg and height ranging from 140-167 cms. All patients were thoroughly examined pre-operatively. Patients belonging to ASA grade I and grade II

were alone taken up for the study. Initially, patients were reassured and counselled to gain confidence. The procedure was explained and an Informed consent was obtained.

Inj. Ranitidine 50mg was given intravenously as premedication 45 minutes before surgery and patients were randomised into two groups each.

GROUP A: Patients received 1.5 ml of 0.5% hyperbaric bupivacaine (7.5mg) with 0.5ml of fentanyl (25 mcg)

GROUP B: Patients received 1.5 ml of 0.5% hyperbaric bupivacaine (7.5mg) with 0.2 ml of buprenorphine (60 mcg) and 0.3 ml of normal saline.

The final volume of the injected solution is 2 ml on both groups.

In preoperative assessment clinic, baseline investigations like haemoglobin, urine analysis for albumin and sugar, blood sugar, urea, creatinine and ECG were checked. Vital parameters like pulse rate, blood pressure respiratory rate were recorded. Thorough examinations of all the systems and airway assessment was done. The patients were educated about Visual analogue scale (VAS) and its interpretations.

In the operating room, appropriate equipment for airway management and emergency drugs were kept ready. Patients were shifted to the operating room. The horizontal position of the operating table was checked and the patient was placed on it. The noninvasive blood pressure monitor, pulse oximeter and electro cardiogram leads were connected to the patient. In the anaesthesia chart, proper recording of preoperative baseline systolic and diastolic blood pressure, pulse rate, respiratory rate and oxygen saturation was documented. 18G intravenous cannula was secured to the patients and preloading was done with 1000ml of Ringers lactate. The patient was placed in left lateral position. The skin over the back were cleaned with swabs and antiseptic solution and draped with sterile towel. The L3 – L4 interspaces was identified and 25G Quincke Babcock spinal needle was introduced in this

space through midline approach. After confirming free flow of CSF, the prepared solution was injected. The patients were made to lie supine immediately after injection and left lateral tilt was provided by wedge under right buttock.

The time of onset and duration of sensory block, motor block were noted. The grading of motor block was done according to modified Bromage scale.

Modified Bromage scale

0 – No block. Able to raise extended leg against gravity

1 – Unable to raise extended leg, just able to flex knees

2 – Unable to flex knees, but able to flex ankle

3 – Total block. Inability to flex ankle / move leg.

The systolic and diastolic blood pressure, pulse rate, respiratory rate and oxygen saturation were noted every minute for the first 10 minutes and thereafter every 5 minutes until the immediate post operative period.

Side effects such as nausea, vomiting, hypotension, respiratory depression, pruritus and allergic reaction were looked and were reported, if any.

Time interval between subarachnoid block and the time to reach $VAS \geq 4$ is defines as the “ duration of effective analgesia”.

In the post anaesthesia care unit, pain assessment using VAS was done every 15 minutes till VAS score ≥ 4 was reached. The APGAR scores of the newborn were recorded at 1 minute and 5 minute intervals after delivery of the baby.

Observations and Results

A total of 50 patients participated in the study and the statistical data were analysed. The average age in both groups were similar. The mean weight and height of the patients were comparable in both groups and they were statistically insignificant.

Sensory block

The mean time of onset of sensory block at T_{10} was 138.6 ± 22.61 seconds in Group A with a range of 105-180 seconds and 112 ± 23.4 seconds

in Group B with a range of 80-154 seconds. This was statistically significant which was confirmed by unpaired Student's test ($p < 0.05$) as shown in the bar diagram.

Maximum level of sensory block

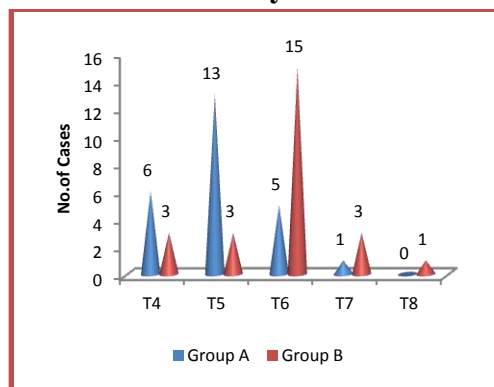
The range of maximal level of sensory block was T₄-T₇ in Group A and in Group B, it was between T₄ - T₈.

Level	Number of patients	
	Group A	Group B
T ₄	6	3
T ₅	13	3
T ₆	5	15
T ₇	1	3
T ₈	0	1

Time to regression to L₁

The mean time to regression of sensory blockade to L₁ was 200.32 ± 9.1 minutes, with a range of 173 - 234 minutes in Group A. In Group B, it was 491.28 ± 153.97 minutes, with a range of 420 - 540 minutes. This was statistically significant ($p < 0.01$).

Maximum level of sensory block



Motor block

Onset of Grade I motor block

The time taken to achieve Grade I motor block was 159 ± 20.31 seconds in Group A with a range of 120 - 195 seconds. In Group B, it was 160.8 ± 22.3 seconds with a range of 135 - 210 seconds. This was found to be statistically insignificant ($p > 0.1$).

Maximum degree of motor block

The maximum degree of motor block ranged between grade 3 and grade 2 in both the groups.

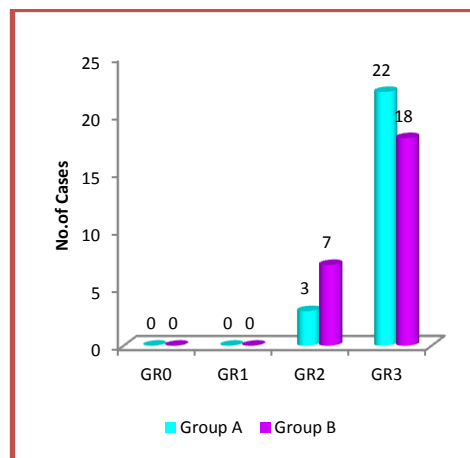
The distribution of patients in each grade is shown in the table.

Degree of Motor Block	Number of Patients	
	Group A	Group B
GR0	0	0
GR1	0	0
GR2	3	7
GR3	22	18

Duration of motor block

The mean duration of motor block was 70.8 ± 11.06 minutes in Group A with range of 60 - 90 minutes. In Group B, it was 69.54 ± 20.23 minutes with a range of 60 - 90 minutes. This was found to be statistically insignificant ($p > 0.1$).

Degree of motor block

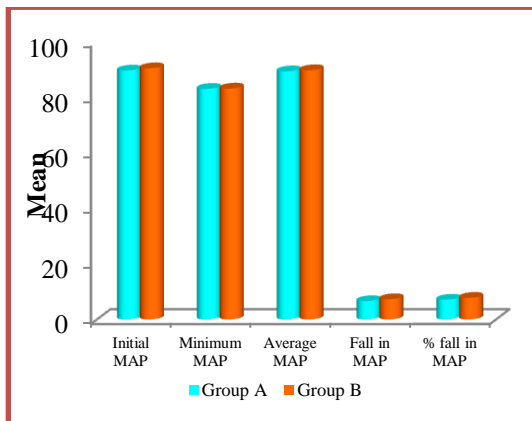


Mean Arterial Pressure variation in study groups

MAP	Group A		Group B	
	Mean	SD	Mean	SD
Initial MAP	89.8	6.6	90.5	7.5
Minimum MAP	83.2	7.5	83.2	6.7
Average MAP	89.5	5.5	89.8	4.9
Fall in MAP	6.5	8.1	7.3	66.4
% fall in MAP	7.2	7.8	7.8	6.8

'p' value = 0.152, which is insignificant.

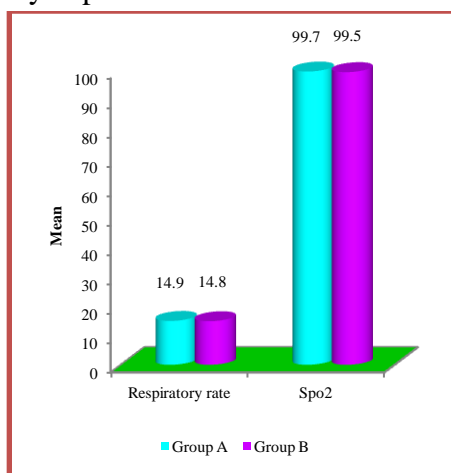
The fall in Mean Arterial Pressure was insignificant in all 2 groups. Thus, addition of low dose fentanyl and buprenorphine gives an added advantage of hemodynamic stability when compared to plain bupivacaine alone.



Respiratory rate and SpO2 among the study groups

	Respiratory rate		Spo ₂	
	Group A	Group B	Group A	Group B
Mean	14.8	14.9	99.8	99.6
SD	1.1	1.2	0.54	0.69
'p' for 2 groups	0.5526, not significant		0.7826, not significant	

There was no significant difference in the two groups, with respect to Respiratory rate and SpO₂. Almost both the groups are stable in this aspect. There are no reports of desaturation and respiratory depression.



Duration of effective analgesia

The mean duration of effective analgesia was 200.32 ± 9.1 minutes (3.33 hours) with a range of 173 – 234 minutes in Group A, but in Group B, it was 491.28 ± 153.97 minutes (8.1 hours) with a range of 420 – 540 minutes. This was statistically highly significant. (p< 0.01)

Quality of surgical anaesthesia

In this study, the quality of surgical anaesthesia was graded as 'excellent' in all, but 3 patients in Group A complained of discomfort intraoperatively and required 10µgm fentanyl intravenously.

Incidence of hypotension

In Group A, 12 patients had hypotension whereas in Group B, only 7 patients had hypotension. The incidence is 48% with Group A against 28% in Group B. This was tested to be statistically significant. (p<0.001)

Mean ephedrine requirement

The mean ephedrine required to counter hypotension was 12 mg in Group A, whereas it was 3.12 mg in Group B.

Duration of post operative analgesia (in minutes)

Post operative analgesia (mins)	Group A	Group B
Mean (mins)	200.32	491.28
SD	9.1	153.97
'p' value for 2 groups	0.0001, significant	

The post operative period is, till the patient demands systemic analgesic (i.e., VAS score ≥4) from the initiation of subarachnoid blockade. The mean duration of effective analgesia was 200.32 ± 9.1 minutes (3.33 hours) with a range of 173 – 234 minutes in Group A, but in Group B, it was 491.28 ± 153.97 minutes (8.1 hours) with a range of 420 – 540 minutes. This was statistically highly significant. (p< 0.01)

Side effects

Side Effects	Number of patients	
	Group A	Group B
Pruritus	4	0
Nausea	2	6
sedation	7	17

The incidence of pruritus was 16% in Group A (4 patients) whereas it was nil with Group A. Pruritus in Group A was mild and settled with reassurance.

6 patients complained of nausea in Group B and 2 in Group A. This was attributed to the addition of

opioids to bupivacaine. Bradycardia and respiratory depression did not occur in any of the patients involved in the study.

In our study, most of the patients were catheterized before shifting to operation theatre, hence urinary retention could not be assessed and compared.

Sedation of grade I was seen with 17 patients in Group B.

Assessment of the fetus

All the babies showed 1 minute APGAR of 8 and above and 5 minute APGAR of 9 and above in both the groups. The difference was statistically insignificant.

Discussion

“Pain is a more terrible lord of mankind than death itself”. Pain is a complicated subjective experience, which is challenging to measure in a reproducible way⁵. Operative pain is more extreme after surgery which thereafter gradually tapers over the next 24 hours⁶. Existence of pain has been a led to the discovery of both newer drugs and procedures of pain relief.

The principal aim of this study was to analyze the efficacy of the combination of fentanyl with bupivacaine verses buprenorphine with bupivacaine in spinal anesthesia for elective caesarean section regarding incidence of hypotension and mean ephedrine requirements apart from other usual parameters and to compare the duration of postoperative analgesia.

Intrathecal opioids were first clinically used by Wang et al⁷. Postural hypotension and exaggerated sympathetic blockade is absent with use of opioids, hence parturient are allowed to ambulate early and mother can breastfeed child effectively, thereby improving interaction between mother and child⁸. Risk of thromboembolic disease is increased during pregnancy. Excellent pain relief is provided postoperatively by intrathecal buprenorphine, hence it enhances early ambulation, thereby decreasing chances of thromboembolic phenomenon.

Without affecting motor block, buprenorphine increases duration of sensory block and gives preferable hemodynamic stability⁹. In the present study, the buprenorphine group had significant early onset of analgesia. This is by virtue of high lipid solubility and highest affinity for opiate receptors of buprenorphine^{10,11}.

The findings in our study also correlates with the study done by Singh H, Yang J, Thornton K and Giesecke¹² AH who concluded that addition of intrathecal fentanyl 25µg to hyperbaric bupivacaine did not hasten the onset of sensory block.

The duration of motor block was not prolonged much by the addition of fentanyl/buprenorphine to 0.5% hyperbaric bupivacaine intrathecally. With increase in dose of bupivacaine, the duration of motor block was increased. Hence, by adding opioid adjuvants, we reduce the dose of bupivacaine thereby reducing the undesirable long duration of motor blockade which causes delay in ambulation in post operative period.

The mean duration of effective analgesia was 200.32 minutes (3.33 hours) in Group A and 491.28 minutes (8.1 hours) in group B which was highly significant statistically. In our study the duration of analgesia was increased due to addition of buprenorphine, the same theory was analysed by capogna etal¹³, who concluded that the duration of analgesia is dose dependent. Longer duration of action and analgesic efficacy of intrathecal buprenorphine can be explained by its high affinity for spinal receptors. Smaller doses of buprenorphine produce a high concentration of the drug at spinal receptors. Higher lipid solubility of buprenorphine favours its diffusion to spinal cord. The diffusion from the spinal cord in to the bloodstream is slow and does not approach the bulbar centres with bulk of CSF. Hence high lipid solubility, strong opiate receptor binding and intense and prolonged activity was responsible for its longer duration of action¹⁴.

In our study, 3 patients in group A complained of discomfort intraoperatively and required supplementation with intravenous fentanyl 10µg.

The quality of surgical anaesthesia was excellent in all other patients.

In our study, the incidence of hypotension was 48% in group A and 28% in group B. This was statistically significant. Our study confirmed the fact that the decrease in sympathetic efferent activity after spinal anaesthesia with bupivacaine was dose related and that intrathecal opioids caused neither by itself nor in combination with bupivacaine, any further depression of efferent sympathetic activity. This correlated with the study of Ben David et al (1984)¹⁵.

Incidence of nausea, vomiting, pruritus were present in both the groups which may be contributed due to addition of opioids, yet, all the side effects were mild and mostly settled with reassurance. None of the patients developed respiratory depression or bradycardia.

With use of opioids, the incidence of Postural hypotension and exaggerated sympathetic blockade is absent. It allows parturient to ambulate early and mother can breastfeed child effectively thereby improving interaction between mother and child¹⁶. A good pain relief is provided in postoperative period by intrathecal buprenorphine, it improves mobility thereby reducing chances of thromboembolic phenomenon which is one of the risks associated with pregnancy and in postpartum period.

Intrathecal fentanyl and buprenorphine did not adversely affect the neonatal outcome.

Conclusion

In this comparative study, an effort was made to study the analgesic efficacy of fentanyl and buprenorphine with 0.5% bupivacaine intrathecally for elective caesarean section.

- 1) There was no significant hemodynamic changes in either of the groups.
- 2) Anaesthesia was superior when buprenorphine is mixed with bupivacaine (0.5%) as compared to bupivacaine with fentanyl.
- 3) Addition of buprenorphine to bupivacaine 0.5% enhances the sensory blockade of local

anaesthetics without affecting the sympathetic activity.

Thus it is concluded that intrathecal buprenorphine is suitable drug for post operative analgesia for caesarean section.

References

1. Celleno D, Capogna G. Spinal buprenorphine for postoperative analgesia after caesarean section. *Acta anaesthesiol Scand* 1989;33:236-8
2. Miwa Y, Yonemura E, Fukushima K, Epidurally administered buprenorphine in the preoperative period. *Can J Anaesth* 1996;43:907-13
3. Lalla RK. Low dose intrathecal buprenorphine for postoperative analgesia. *Indian J Anaesth* 1997;41:38-9
4. Bromage PR. The price of intraspinal narcotic analgesia: Basic constraints (editorial) *Anaesth Analg*. 1981 July; 60(7): 461-463
5. R.G. Wheatly, S.A.Schug, D.Watson. Safety and efficacy of postoperative analgesia. *Br. J Anaesth* 2001 Feb; 87(5): 47-61.
6. Parkhouse, J.Lambrechts, R.w. Simpson. Incidence of postoperative pain. *Br J Anaesth*. 1961 Apr; 33(4): 576-581.
7. Wang JK, Nauss LA, Thomas JK. Pain relief by intrathecally applied morphine in man. *Anaesthesiology* 1979; 50: 149-51.
8. Jeff Gasden, Stuart Hart and Alan C. Sandos. Post – cesarean delivery analgesia. *Anaesth Analg* 2005; 101: 62-69.
9. Atweth SF, Kuhar MJ. Autoradiographic localization of receptors in rat brain, spinal cord and lower medulla. *Brain Research* 1977; 124: 53-67.
10. Dikenson AH. Spinal cord pharmacology of pain. *British Journal of Anaesthesia* 1995; 75: 193.
11. Chang HM, Berde CB, Holz GG et al. sufentanil, morphine metenkephalin and K agonist (U-50, 488H) inhibit substance P

release from primary sensory neurons: A model for presynaptic spinal opioid actions. *Anaesthesiology* 1989; 70:672

12. Singh H, Yang J, Thornton K, Giesecke AH. Intrathecal fentanyl prolongs sensory bupivacaine spinal block. *Canadian Journal of Anesthesia* 1995; 42(11):987-91.
13. Singh H, Yang J, Thornton K, Giesecke AH. Intrathecal fentanyl prolongs sensory bupivacaine spinal block. *Canadian Journal of Anesthesia* 1995; 42(11):987-91.
14. Cahill J, Murphy D, O'Brain D, Mulhall J. Epidural buprenorphine for pain relief after major abdominal surgery. A controlled comparison with epidural morphine. *Anaesthesia*. 1983 Aug; 38(8): 760-764.
15. Ben-David B, Frankel R, Arzumonov T, Marchevsky Y, Volpin G. Minidose bupivacaine- fentanyl spinal for surgical repair of hip fracture in the aged. *Anaesthesiology* 2000; 92: 6-10
16. Wang JK, Nauss LA, Thomas JE. Pain relief by intrathecally applied morphine in man *Anaesthesiology* 1979; 50; 149-5.