Epidural Fentanyl speeds the onset of sensory and motor blocks during epidural ropivacaine anaesthesia

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ABSTRACT

BACKGROUND

Different adjuvants are added to local anaesthetic solutions to hasten the onset of sensory and motor block in epidural anaesthesia.

OBJECTIVE

To study the effect of epidural and intravenous fentanyl on time of onset of sensory and motor blockade in epidural anaesthesia with 0.75% ropivacaine.

MATERIALS AND METHODS

After obtaining informed consent and permission from hospital scientific and ethical committee 40 patients with ASA grade I and II who are undergoing lower abdominal, anogenital, urological, gynaecological, and orthopaedic surgeries are included in this study. Patients with bleeding disorders, infections at puncture site, history of opioid dependence, pregnant ladies, age <18 years, and morbidly obese patients are excluded from this study. Patients are randomly allocated into two groups: epidural fentanyl (EF) and intravenous fentanyl (IF) with 20 patients in each group. After the test dose, Epidural Fentanyl (EF) group receive 20 mL of 0.75% ropivacaine plus 100 µg of fentanyl (2 mL) followed by IV injection of 2 mL of normal saline. The patient in the intravenous fentanyl (IF) group receive 20 mL of 0.75% ropivacaine plus 2 mL of normal saline followed by IV injection of 100 µg fentanyl (2 mL). Time of onset of sensory block, time of onset of motor block, upper level of sensory block, incidence of different complication are recorded during the study. Independent samples t - test, Kruskal - Wallis test and one way ANOVA test were used for statistical analysis.

RESULTS

The onset time for sensory block is earlier in group EF than group IF (10.90±2.81 mins <16±3.03 mins) (mean±SD). The motor blockade onset time is more rapid in group EF than group IF (14.95±5.43 mins <26.40±5.94 mins) (mean±SD). The upper level of sensory block is indifferent in both groups (1.70±0.571 = 1.75±0.716) (mean±SD). The incidence of different complications is insignificant in both groups (p<0.135 >0.05).

CONCLUSION

Epidural administration of the mixture of fentanyl and ropivacaine solution accelerated the onset of sensory and motor blocks during epidural ropivacaine anaesthesia without significant fentanyl related side effects. Fentanyl is a useful adjunct to local anaesthetic ropivacaine for epidural anaesthesia.

KEYWORDS

Ropivacaine, Fentanyl, Epidural Anaesthesia, Sensory Block, Motor Block.


INTRODUCTION

Recent trends suggest that regional anaesthesia is replacing general anaesthesia in all most all the surgeries below umbilicus mainly because its benefits such as avoidance of polypharmacy, airway manipulation, misplacement of endotracheal tube, hypo or hyperventilation, vomiting, pulmonary aspiration.¹ The delayed onset time of sensory block in epidural anaesthesia is sometimes a drawback for clinical practice. Shortening the onset time of sensory block is a practical goal to improve the quality of epidural anaesthesia. A variety of adjuvants are used for epidural infusion to enhance analgesia while minimizing side effects, like clonidine,²,³ epinephrine,⁴,⁵ ketamine,⁶ sodium bicarbonate,⁷ etc. The addition of fentanyl to a local anaesthetic solution is widely used during epidural anaesthesia,⁸ The addition of fentanyl to lidocaine,⁸ bupivacaine,⁹ and mepivacaine,¹⁰ solutions produces a rapid onset of sensory block during epidural anaesthesia. Conversely, other investigators have reported no change in the onset of analgesia with the addition of fentanyl to epidural mepivacaine.¹¹ Alkalisation of local anaesthetic solution has been used to shorten the onset time.¹² Ropivacaine, a long-acting amide type local anaesthetic is widely used in epidural anaesthesia.
AIMS AND OBJECTIVES
The aim of this prospective randomised study is to examine the effect of epidural fentanyl on the onset times of sensory and motor blocks during epidural ropivacaine anaesthesia.

MATERIALS AND METHODS
After obtaining informed consent and permission from hospital scientific and ethical committee, 40 patients with ASA grade I and II randomly allocated into two groups: Epidural Fentanyl (EF) and Intravenous Fentanyl (IF) with 20 patients in each group who are admitted into the Seven Hills Hospital, Visakhapatnam for undergoing lower extremity orthopaedic, urologic, gynaecological, and lower abdominal surgeries are included in this study.

Randomisation is done using computer-based software into two different groups. Patients with bleeding disorders, infections at puncture site, history of opioid dependence, allergic to study drugs, pregnant ladies, age < 18 years, and morbidly obese patients are excluded from this study.

Patients are monitored with electrocardiogram, non-invasive arterial blood pressure, heart rate, and pulse oximetry during surgery.

With patients in sitting position, the epidural space is identified with an 18-Gauge Tuohy’s needle by the loss of resistance technique. A test dose of 3 mL of solution containing 3 mL of 2% lignocaine with 15 μg of adrenaline in 1:2,00,000 dilutions is given for identification of intravascular or intrathecal placement of epidural needle.

After the test dose, Epidural Fentanyl (EF) group receive 20 mL of 0.75% ropivacaine plus 100 μg of fentanyl (2 mL) followed by IV injection of 2 mL of normal saline. The patient in the Intravenous Fentanyl (IF) group receive 20 mL of 0.75% ropivacaine plus 2 mL of normal saline followed by IV injection of 100 μg fentanyl (2 mL).

The sensory block is assessed by pinprick method in left anterior axillary line at 2 minutes interval for 20 mins. Onset of sensory block is defined as time from epidural injection to the occurrence of sensory block at T₁₀ dermatome. The upper level of sensory block is recorded.

The motor block is assessed at 2 minutes interval for 40 mins by modified Bromage scale, (4=no movement, 3=unable to raise extended leg or bend knee, 2=unable to raise the extended leg, but able to bend the knee, 1=able to move feet, but not able to bend the knees, 0=complete leg and foot movement). The onset of motor block is defined as the time from epidural injection to occurrence of motor block at each scale.

Hypotension (Systolic blood pressure <100 mmHg or a decrease of more than 30% from baseline) is treated with IV ephedrine as needed. Side effects such as nausea, vomiting, pruritus, respiratory depression, or shivering are recorded during surgery.

Based on a previous study, an estimated standard deviation of 5 min for the onset of sensory block during epidural ropivacaine anaesthesia is used. A decrease in the onset time of 30% is considered clinically significant. On the basis of these estimates, a sample size of 20 patients in each group is sufficient to get a two-tailed type I error of 0.05 and a power of 80%. Data is analysed using Microsoft excel - 2007 and computer-based software programme under the name of SPSS statistics version - 20 marketed by IBM corporation 2011.

RESULTS AND DISCUSSION
The two study groups are similar in demographic profile (Age, sex, height, weight) assessed with independent samples t-test and independent samples Kruskal-Wallis test as shown in table-1. The anesthetic characters are shown in table-2 and analysed with independent samples t-test. The incidence of different complication was analysed by one-way ANOVA test. Microsoft office excel - 2007 and computer-based software programme under the name of SPSS statistics version 20 marketed by IBM Corporation 2011 are used for statistical analysis.

In this randomised non-blinded study, effect of epidural fentanyl and intravenous fentanyl on onset time of sensory and motor block of epidural ropivacaine anaesthesia is analysed.

In this study, the demographic profile (Age, sex, height, and weight) between the two groups is analysed with independent sample t-test and hypothesis is tested with independent samples Kruskal-Wallis test. The p-value is > 0.05 indicating insignificant differences among the anthropometric variables among the groups. Similar results among the anthropometric variables is found by Tomar GS et al while comparing two different doses of fentanyl added to bupivacaine for intermittent epidural labour analgesia. They analysed demographic data using ANOVA test and found a p-value of > 0.05 concluding there is no significant difference in the study groups.

In this study, the mean sensory block onset time (T₁₀ dermatome) of epidural fentanyl group is found to be 10.90±2.81 (mean±SD) minutes and that of intravenous fentanyl group is 16±3.03 (mean±SD) minutes. Data is analysed using independent samples t-test with a t-value of -5.252 and p-value of 0.000 (p-value < 0.05), which is statistically significant. Similar results are obtained by Cherng CH et al while comparing the sensory block onset time (T₁₀ dermatome) with (EF group) epidural administration of 17 mL of 2% lidocaine plus 100 μg fentanyl and followed by intravenous (IV) injection of 2 mL of normal saline, (IF group) epidural administration of 17 mL of 2% lidocaine plus 2 mL of normal saline and followed by IV injection of 100 μg of fentanyl and (C group) epidural administration of 17 mL of 2% lidocaine plus 2 mL of normal saline and followed by IV injection of 2 mL of normal saline. They found the onset time of sensory block up to T₁₀ dermatome was significantly more rapid in the EF group (8.3+/−3.7 minutes) than that of the IF group (13.1+/−4.2 minutes, P<.05) or C group (14.2+/−5.4 minutes, P<.05).

In this study, the mean motor block onset time for modified Bromage scale 1 and 2 of epidural fentanyl group is found to be 14.95±4.3 (mean±SD) minutes and that of intravenous fentanyl group is found to be 26.40±5.94 (mean±SD) minutes. Analysis of data with independent samples t-test give the resultant t-value of -6.360 and p-value of 0.000 (p-value < 0.05), which is statistically significant. Chen-Hwan Cherng et al examined the onset times of sensory and motor block during epidural ropivacaine anaesthesia with and without the addition of fentanyl to the epidural solution. They found that the onset times of motor block up to Bromage scale 1 and 2 were significantly more rapid in the EF group (11.9±4.6 and 24.4±5.9 min) than in the IF group (16.7±4.7 and 30.8±5.6 min, P <0.05) or C group (18.3±4.9 and 32.7±5.7 min, P <0.05). They concluded that onset time of


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motor block to the modified Bromage scores 1 and 2 was significantly more rapid in the EF group compared with the IF and C groups.

In this study, the upper sensory level is up to T4 dermatome level in both the groups, which is analysed using ANOVA test with F value of 0.060 and p-value of 0.809 showing statistical insignificance. Similar statistical insignificance in upper sensory level is found in the study of Chen-Hwan Cherng et al[20] They analysed data using the Kruskal-Wallis test and the Dunn’s multiple comparison procedure for post hoc comparison and found a p-value >0.05.

In this study, incidence of complication like shivering, hypotension, pruritus, nausea and vomiting, respiratory depressions, urinary retention are recorded. There is no incidence of pruritus, nausea and vomiting, respiratory depression, which are more common with fentanyl. Incidence of hypotension and shivering are noted in both groups. Incidence of complications in both epidural fentanyl and intravenous fentanyl group analysed with one-way ANOVA test, which gave a p-value >0.05 (0.135) showing statistical insignificance. Similar results are found by Cherng CH et al[8] Tomar GS et al[9] Chen-Hwan Cherng et al[20] with p-value >0.05 showing statistical insignificance.

![Comparison of Sensory and Motor Block Onset Times among different groups.](image)

**Fig. 1: Comparison of Sensory and Motor Block Onset Times among different groups. EF-Epidural Fentanyl, IF-Intravenous Fentanyl, SBOT-Sensory Block Onset Time, MBOT-Motor Block Onset Time**

**CONCLUSION**

From this study, it was concluded that epidural administration of the mixture of 100 μg fentanyl and 0.75% ropivacaine solution accelerated the onset of sensory and motor blocks during epidural ropivacaine anaesthesia without significant fentanyl related side effects. Fentanyl is a useful adjunct to local anaesthetic ropivacaine for epidural anaesthesia.

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