A Comparative Efficacy of two Regional Techniques for Labour Analgesia: Combined Spinal Epidural Analgesia versus Epidural Analgesia

Authors

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Abstract

Background and Aim: Central Neuraxial analgesia is the most versatile method of labour analgesia and the gold standard technique for pain control in obstetrics that is currently available. The satisfaction of birth experience is greater with neuraxial techniques. Neuraxial analgesia is the mainstay analgesic, frequently administered to women in labor. We aimed at comparing efficacy of two regional techniques for labour analgesia by combined spinal epidural analgesia versus epidural analgesia evaluating by onset of analgesia, duration of analgesia, maternal satisfaction, mode of delivery, neonatal outcome, complications (maternal and fetal).

Materials and Methods: 40 parturient of ASA physical status I or II in active labor with a cervical dilatation of 3-6 cm requesting labour analgesia were motivated and enrolled were divided into two groups. Group A: Received combined spinal Epidural technique using Intrathecal dose 1 ml of solution containing 1 mg of 0.5% Hyperbaric Bupivacaine + 25 mcg Fentanyl. Followed by subsequent top ups on patient demand using 10 ml of solution containing 0.1% Bupivacaine + 2 mcg/ml of Fentanyl. Group B: Received Epidural analgesia bolus dose of 10 ml solution 0.1% Bupivacaine + 2 mcg/ml of Fentanyl. Intermittent top ups on patient demand using 10 ml solution containing 0.1% Bupivacaine + 2 mcg/ml of Fentanyl. Onset of analgesia, Mode of delivery, Neonatal outcome, Maternal satisfaction, Complications (maternal and fetal) were assessed. Pain intensity by using a numerical rating scale and motor block by using bromage scale.

Results: Eighteen parturients from CSEA had onset of analgesia in less than 5 min, twelve parturient from LEA group had onset of analgesia between 11 to 15 min and seven parturients had onset of analgesia between 16-20 min. The mean time of onset of analgesia in Group A was 5.05 ± 2.25 Min. The mean time of onset of analgesia in Group B was 15.2 ± 1.93 Min. Eighteen parturients from Group had undergone normal vaginal delivery. One parturient from Group B had undergone caesarian section for different indication. One parturient from Group B had undergone delivery in instrumental mode. All the Neonates from Group A and Group B had Apgar score of 5th min of delivery. There was no difference in neonatal outcome between two groups. Five parturients in Group A complained of Pruritis and none of the parturients in group B complained of pruritis. Two parturients from each group had Hypotension. None of the parturients in neither group had other Complications like nausea and vomiting, non progressive labor or fetal distress.

Conclusion: From the present study we conclude that both the techniques Combined spinal Epidural and Epidural for labor analgesia produce excellent analgesia. However the CSE technique had faster onset of analgesia when compared to Epidural technique. Both the technique were similar in terms of safety and efficacy.

Keywords: Labour analgesia, Combined spinal epidural analgesia, epidural analgesia.
Unrelieved stress in labour produces increased plasma cortisol and catecholamines concentrations which reduces utero-placental blood flows by 35-70% compounding the effects of hyperventilation on the oxygen supply to the foetus. Metabolic acidosis as a result of increased metabolic rates especially in the second stage of labour is transferred to the foetus. There is delayed gastric emptying and urinary emptying. Effective pain relief reduces plasma noradrenaline, prevents the rise during first and second stage of labour of 11-hydroxy corticosteroid, prevents metabolic acidosis by reducing the rate of rise of lactate and pyruvate. The pain-induced hyperventilation and hypocapnia reduces utero-placental blood flow by 25%. The respiratory alkalosis further impairs foeto-maternal gas exchange by shifting the oxyhaemoglobin dissociation curve to the left and fetal PaO2 may fall by 23%. There have been no proven scientific data analysis of the quality of pain relief offered by non pharmacological methods neither transcutaneous electric nerve stimulation nor inhalation of entonox was not adequate for pain relief in labour. Central neuraxial analgesia is the most versatile method of labour analgesia and the gold standard technique for pain control in obstetrics that is currently available. Epidural analgesia was associated with greater pain relief than non epidural methods however it is associated with longer first and second stage of labour. An increased incidence of fetal malposition and increased use of oxytocin and instrumental vaginal deliveries. Combined spinal epidural analgesia is as safe as conventional epidural technique and is associated with greater maternal satisfaction.

The satisfaction of birth experience is greater with neuraxial techniques. Neuraxial analgesia is the mainstay analgesic, frequently administered to women in labor. On this background we tried to compare both the techniques combined spinal epidural and epidural in labour analgesia in order to evaluate onset, efficacy of analgesia and safety of mother and fetus.

Aims and Objectives
The aim of the study is to compare efficacy of two regional techniques for labour analgesia using combined spinal epidural analgesia versus epidural analgesia

The parameters studied are
1. Onset of analgesia.
2. Mode of delivery.
5. Complications (maternal and fetal).

Materials and Methods
Study was conducted in the Department of Anesthesiology in association with Department of Gynecology and Obstetrics at King George Hospital, Visakhapatnam. 40 parturient of ASA physical status I or II in active labor with a cervical dilatation of 3-6cm requesting epidural analgesia were motivated and enrolled. They were divided into two groups. Group A: Received combined spinal Epidural technique using Intrathecal dose 1ml of solution containing 1mg of 0.5% Hyperbaric Bupivacaine + 25mcg Fentanyl. Followed by subsequent top ups on patient demand using 10ml of solution containing 0.1% Bupivacaine + 2ug/ml of Fentanyl.

Group B: Received Epidural analgesia bolus dose of 10ml solution 0.1% Bupivacaine+2mcg/ml of Fentanyl.Intermittent top ups on patient demand using 10ml solution containing 0.1% Bupivacaine + 2mcg/ml of Fentanyl.

Inclusion Criteria
a) Healthy Primigravida and gravida 2 patients at term.
b) ASA (American Society of Anesthesiologists) 1 and 2.
c) Maternal request for labour analgesia.
d) Age group 18-35 years.

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c) Maternal request for labour analgesia.
d) Age group 18-35 years.
e) Women in active labour with cervical dilatation in primi about 4-5 cm and gravida 2 with cervical dilatation of 3-4 cm.

Exclusion Criteria
a) Multiple pregnancies or preterm gestation pregnancies.
b) Weight >90 Kg, Age <18 yrs.
c) Allergy to any study drug.
d) Any contraindication to Epidural analgesia like patient refusal, bleeding disorders, localizes infection at the site of injection etc.

Group A:

Technique of CSEA
CSEA technique can be performed by SINGLE SPACE NEEDLE THROUGH NEEDLE TECHNIQUE.

Patient Position
• L2-L3 or L3-L4 space identified and skin was infiltrated with small amount of local anesthetic with 25G needle and skin wheal is created using standard TUHOY needle epidural space is identified. Then a fine bore long spinal needle (27G and 124mm or longer) is inserted into subarachnoid space through the epidural needle. A characteristic pop was felt when spinal needle pierces the duramater and CSF was seen dribbling out.
• A dose containing 1mg Bupivacaine + 25mcg Fentanyl making to 1ml using normal saline was injected into subdural space and needle was withdrawn. • Then the catheter was threaded into the epidural space upto 3 -4 cm length and was taped to back and patient was positioned to supine. Further top ups are given to patient demand through epidural catheter as 10ml solution containing 0.1% Bupivacaine + Fentanyl 2mcg/ml.

GROUP B:

Technique of Epidural Analgesia
• L2-L3 or L3-L4 space was identified and the skin infiltrated with small amount of local anesthesia with a 25 G needle and skin wheel is created. Using standard TUHOY needle epidural space is identified by "Hanging drop" which provides visual identification or “loss of resistance method" which gives evidence of entry into Space regardless of the technique used, it is confirmed that the needle was in Epidural Space. The epidural space is confirmed by injecting 3ml of saline or 4-5 cc of air or a test dose of local anesthetic. • Patient who did not experience symptoms from test dose received 10ml Containing 0.1% Bupivacaine +2mcg/ml of Fentanyl as 3ml increment in 5min. • Analgesia was maintained throughout labor by intermittent top ups given, on demand from the patient.

• Pain intensity by using a numeric rating scale.
  • 0=no pain
  • 10=worst pain imaginable and sensory levels to pin prick.
  • Motor block by using a Bromage scale(0-3 scale)
• side effects like Nausea, Pruritis and Respiratory depression were assessed at baseline 5min, 15 min, 30 min, 45 min, 60min, then every 2hrs until complete cervical dilatation and at delivery.
• Patients were asked to rate the pain intensity during uterine contractions.
• Patient satisfaction was assessed immediately after delivery as excellent good, fair and poor.

Analysis of Results
Statistical analysis were performed by
• Data was presented as mean SD.
• P value of less than 0.05 was considered statistically significant.
Observation and Results

Distribution of Age between CSEA and LEA

Table 1 and Figure 1 shows the age distribution in study.

<table>
<thead>
<tr>
<th>AGE YRS</th>
<th>NO. OF PARTURIENTS IN CSEA</th>
<th>NO. OF PARTURIENTS IN LEA</th>
</tr>
</thead>
<tbody>
<tr>
<td>18-24</td>
<td>16</td>
<td>17</td>
</tr>
<tr>
<td>25-30</td>
<td>4</td>
<td>3</td>
</tr>
</tbody>
</table>

Sixteen parturient in Group A and Seventeen parturient in Group B were within the age group of 19-25 yrs. Most of our parturient in the study group are in this range of age group. Four parturients in Group A and three parturients in group 3 were in the age group of 26-30yrs. It was observed that the mean age in group A was 22.65±2.45 and mean age group in Group B was 21.7±1.94

Statistical analysis of age distribution was compared between two groups using Standard error of difference between means. The P value is >0.05 so statistically no difference exists between two groups in respect to age distribution.

Table 2 and fig 2 shows height distribution in the study.

<table>
<thead>
<tr>
<th>HEIGHT IN CM</th>
<th>NO. OF PARTURIENTS IN CSEA</th>
<th>NO. OF PARTURIENTS</th>
</tr>
</thead>
<tbody>
<tr>
<td>141-150</td>
<td>2</td>
<td>3</td>
</tr>
<tr>
<td>151-160</td>
<td>12</td>
<td>12</td>
</tr>
<tr>
<td>161-170</td>
<td>6</td>
<td>5</td>
</tr>
</tbody>
</table>
It was observed that 12 parturients from each group in height range of 151-160cm. 6 parturients from group A and 5 parturients from group B were in the height range of 161-170cm and 2 parturients from group A, 3 parturients from group B were in the height range of 141-150cm. The mean height in group A was 157.25±4.8, mean height in group B was 158±5.9 with p value of >0.05 was statistically not significant.

Table 3 and figure 3 Shows the weight distribution in the study.

<table>
<thead>
<tr>
<th>WEIGHT IN KG</th>
<th>NO OF PARTURIENTS</th>
<th>NO OF PARTURIENTS IN LEA</th>
</tr>
</thead>
<tbody>
<tr>
<td>51-60</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>61-70</td>
<td>15</td>
<td>13</td>
</tr>
<tr>
<td>71-80</td>
<td>0</td>
<td>2</td>
</tr>
</tbody>
</table>

Fifteen Parturients from Group A thirteen parturients in Group B were in weight of 61-70kgs, five parturients from each group were in weight range of 51-60kgs, 2 parturients from group B were in weight range of 71-80kgs. It was observed that the mean in Group A was 62.2±2.25 and the mean in Group B was 62.15±5.53). Statistical analysis was done using Standard error of difference between means and is not significant.

Distribution between CSEA and LEA
Table no 4 and figure 4 shows the number of Nulliparous and parous parturients in each group

<table>
<thead>
<tr>
<th>PARITY</th>
<th>NO OF PARTURIENTS IN CSEA</th>
<th>NO OF PARTURIENTS IN LEA</th>
</tr>
</thead>
<tbody>
<tr>
<td>NULLIPAROUS</td>
<td>12</td>
<td>12</td>
</tr>
<tr>
<td>MULTIPAROUS</td>
<td>8</td>
<td>8</td>
</tr>
</tbody>
</table>
It was observed that twelve parturients from CSEA group and twelve parturients from LEA group were Nulliparous and eight parturients from each group were Parous.

**Onset of Analgesia between CSE and LEA**

Table no 5 and figure 5 shows onset of analgesia after intrathecal administration in Group A and, after administration of bolus Epidural dose in Group B.

<table>
<thead>
<tr>
<th>Onset of Analgesia in Min</th>
<th>No of Parturients in CSEA</th>
<th>No of Parturients in LEA</th>
</tr>
</thead>
<tbody>
<tr>
<td>0—5</td>
<td>18</td>
<td>0</td>
</tr>
<tr>
<td>6—10</td>
<td>2</td>
<td>1</td>
</tr>
<tr>
<td>11—15</td>
<td>0</td>
<td>12</td>
</tr>
<tr>
<td>16—20</td>
<td>0</td>
<td>7</td>
</tr>
</tbody>
</table>

Eighteen parturients from CSEA had onset of analgesia in less than 5 min, twelve parturient from LEA group had Onset of analgesia between 11 to 15 min and seven parturients had onset of analgesia between 16-20min.

The mean time of onset of analgesia in Group A was 5.05 ±2.25 Min. The mean time of onset of analgesia in Group B was 15.2 ± 1.93 Min. The statistical analysis was done using Standard error of difference between means and p value of <0.05 indicating significant difference in onset of analgesia between two techniques.

**Duration of Analgesia Distribution between CSEA and LEA**
### Table 4: Duration of Analgesia

<table>
<thead>
<tr>
<th>Duration in Min</th>
<th>No of Parturients in CSEA (Spinal Component)</th>
<th>No of Parturients in LEA (First Epidural Bolus)</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;30</td>
<td>3</td>
<td>0</td>
</tr>
<tr>
<td>30-60</td>
<td>10</td>
<td>11</td>
</tr>
<tr>
<td>60-90</td>
<td>7</td>
<td>8</td>
</tr>
<tr>
<td>&gt;90</td>
<td>0</td>
<td>1</td>
</tr>
</tbody>
</table>

Three parturients from group A asked for analgesia in <30min. 10 parturients from group A and 11 parturients from group B were in range of 30 to 60min. 7 parurients from group A and 8 parturients from group B and 1 parturient from group B had analgesia for >90min.

The mean duration for analgesia for spinal dose was 55 ± 17.70. Mean duration for bolus dose of group B was 65.75 ± 12.26. P value of <0.05 indicating significant difference between two techniques in duration of analgesia of first dose.

### Mode of Delivery distribution between CSEA and LEA

Table 6 and figure 6 shows mode of delivery in Group A and Group B.

<table>
<thead>
<tr>
<th>MODE OF DELIVERY</th>
<th>NO OF PARTURIENTS IN CSEA</th>
<th>NO OF PARTURIENTS IN LEA</th>
</tr>
</thead>
<tbody>
<tr>
<td>VAGINAL</td>
<td>19</td>
<td>18</td>
</tr>
<tr>
<td>INSTRUMENTAL</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>CAESEREAN</td>
<td>1</td>
<td>1</td>
</tr>
</tbody>
</table>

Nineteen parturient from Group A and Eighteen parturients from Group had undergone normal vaginal delivery. One parturient from Group B had underwent caesarain section for different indication. One parturient from Group B had undergone delivery in instrumental mode.

The statistical analysis was done using Chi-Square test, The P value is > 0.05, so statistically not significant.
Neonatal outcome distribution between CSEA and LEA

Table 7 and figure 7 shows Neonatal outcome in Group A and Group B.

**APGAR**

<table>
<thead>
<tr>
<th></th>
<th>NO OF NEONATES IN CSEA</th>
<th>NO OF NEONATES IN LEA</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;7</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>&gt;7</td>
<td>20</td>
<td>20</td>
</tr>
</tbody>
</table>

Neonatal outcome was graded according to Apgar score of Neonate at 1st and 5th min after delivery.

All the Neonates from Group A and Group B had Apgar score of at 5th min of delivery. There was no difference in neonatal outcome between two groups.

Complications distribution between CSEA and LEA

Table 8 and Figure 8 shows the complications that occurred during the period of analgesia in Group A and Group B.

<table>
<thead>
<tr>
<th>COMPLICATION</th>
<th>CSEA</th>
<th>LEA</th>
</tr>
</thead>
<tbody>
<tr>
<td>HYPERTENSION</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>PRURITIS</td>
<td>5</td>
<td>0</td>
</tr>
<tr>
<td>NON PRORESSION LABOUR</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>FOETAL DISTRESS</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>NAUSEA &amp; VOMITING</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>
Five parturients in Group A complained of Pruritis and none of the parturients in group B complained of pruritis. Two parturients from each group had Hypotension. None of the parturients in neither group had other Complications like nausea and vomiting, non progressive labor or fetal distress.

Maternal Satisfaction Distribution between CSEA and LEA
Table 9 and fig 9 shows patient satisfaction on analgesia in group A and group B

![Maternal Satisfaction Distribution between CSEA and LEA](image)

<table>
<thead>
<tr>
<th>MATERNAL SATISFACTION</th>
<th>NO OF PARTURIENTS IN CSEA</th>
<th>NO OF PARTURIENTS IN LEA</th>
</tr>
</thead>
<tbody>
<tr>
<td>EXCELLENT</td>
<td>18</td>
<td>16</td>
</tr>
<tr>
<td>GOOD</td>
<td>2</td>
<td>4</td>
</tr>
<tr>
<td>FAIR</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>POOR</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

16 parturient from group A and 18 parturients from group B quoted quality of analgesia as excellent. 4 parturients from group B quoted quality of analgesia as good.

The statistical analysis was done using chi square test ,p value is >0.05 indicating no significance in maternal satisfaction between two techniques.

Discussion

Neuraxial analgesia is now considered gold standard for providing pain relief during labour. The CSE technique has gained popularity as it provides rapidity of spinal analgesia, minimal motor blockade and flexibility of epidural catheter to extend the block for caesarean section and for providing post operative analgesia.

Despite advantages regional analgesia may have its own problems, most important of which are its effect on progress of labour and outcome. Epidural analgesia may increase, decrease or have no effect on the rate of cervical dilatation in the first stage of labour.

In the current study fourty parturient of ASA grade 1 or 2 of age group of 19-30 yrs including both primi and parous women were randomly selected and divided into groups A and B. Group A received Combined spinal epidural technique by needle through needle approach. Group B received Epidural analgesic technique. Except for spinal component both the groups were standardized in respect to drug dosages of epidural component.
Continuous hemodynamic monitoring was done throughout the process of labour analgesia. The demographic variables like age, height, weight were similar between two groups. The statistical analysis of demographic variables is done using the standard error difference between the means. The p value >0.05 is statistically not significant.

The onset of analgesia in group A who received combined spinal epidural analgesia is <5min when compared to who received epidural analgesia 15-20min.

On the contrary in the study conducted by Hepner et al to compare the CSE and low dose epidural in labour analgesia in respect to initiate and manage motor block onset of analgesia and satisfaction during labour. They observed that there was no difference in either technique motor blockade, parturient satisfaction and the onset of analgesia in epidural group was <2-5min.

Duration of analgesia after spinal dose is significantly less than duration of analgesia after onset of analgesia of bolus dose. The mean duration for analgesia for spinal dose in group A was 55+/−17.70.

The mean duration of analgesia of bolus dose in Group B was 65.75+/−12.26.

Three parturient (15%) from group A complained of pain in <30min where as none of the parturient requested for additional analgesia in Group B within 30min after the onset of analgesia.

In the study conducted by Dr. Sunanda Gupta et al. the effectiveness of CSE was compared with epidural using spinal dose of 1.25mg bupivacaine plus 25mcg of fentanyl and bolus epidural dose of 10 ml solution of bupivacaine +25 mcg of fentanyl. They observed that the interval from initial bolus dose to maternal request for additional analgesia was increased in epidural group compared to CSE group.

The mode of delivery between two Groups was similar 19 parturient from group A and the 18 parturient from Group B delivered vaginally one from each Group undergone caesarean section and one parturient from Group B needed instrumental assistance for delivery.

The Neonatal outcome, as assessed by Apgar scores was compared between two groups. Only one neonate had Apgar score of <7 at 1st min from Group A remaining all neonates from both groups had Apgar score > 7 at 1st and 5th min.

The incidence of side effects between two groups was observed two groups was observed .five parturient from Group A complaint about Pruritis and none from Group B complained of Pruritis.

Two parturient from either group had Hypotension treated Inj phenylephrine and I.V fluids. Pruritis had been mentioned as main side effect in combined spinal epidural analgesia and study supports incidence of Pruritis in Spinal Epidural group . 25% of parturient in combined spinal epidural group about pruritis and none from epidural group complained about pruritis. to assess the relative effects of CSE and epidural during labor.

The parturient response to both the technique is Excellent to Good in 100% of Parturient. The maintenance of maternal expulsive power during second stage of labor, to ability ambulate at their will shows their increased acceptance by the parturient.

**Summary**

The present study was done to compare efficacy and safety of two techniques of labour analgesia using COMBINED SPINAL EPIDURAL and EPIDURAL ANALGESIA.

The onset of analgesia, duration of analgesia of first dose (spinal or epidural bolus dose) , mode of delivery , neonatal outcome, maternal and fetal side effects and maternal satisfaction were observed , compared and analyzed statistically.

Continuous hemodynamic monitoring of the mother and fetal heart rate were monitored throughout the process of tabor. There was no statistical significance (p<0.05) regarding demographic variables, Age, Height, Weight in between two group. Likewise parity status also similar between two groups.
The onset of analgesia was faster (5min) in combined spinal epidural technique compare with Epidural technique (15-20min). This difference in onset of analgesia was statistically Significant (p value <0.05. Mode of delivery was similar and neonatal outcome was good and equal in both groups without any statistical significant difference.

The incidence of Pruritis was more in combined spinal epidural group than Epidural group. Two parturient from each group had maternal hypotension. None of the parturient from either group had other side effects like nausea, vomiting, urinary retention and respiratory depression.

Overall maternal satisfaction was also similar between two groups.

**Conclusion**

From the present study we conclude that both the techniques Combined spinal Epidural and Epidural for labor analgesia produce excellent analgesia. however the CSE technique had faster onset of analgesia when compared to Epidural technique. Both the technique were similar in terms of safety and efficacy.

The present study would have been more effective if the study had done on larger sample of parturient’s than present study and also could be done by reducing the concentration of Bupivacaine to 0.0625%

**References**