Comparison of intrathecal bupivacaine (17.5mg) with preservative free Midazolam (1mg) versus intrathecal Bupivacaine (17.5mg) for postoperative pain relief in lower limb surgeries

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Abstract

Background: Spinal anaesthesia is a common anaesthetic technique for lower limb surgery. Many adjuncts have been tried to prolong the duration of analgesia provided by local anaesthetics when administered intrathecally. Midazolam has been shown to prolong the duration of analgesia when used as an adjuvant, providing the added advantages of mild sedation and amnesia, while being devoid of neurotoxicity, and the adverse effects of opioids. This study was designed to evaluate the effect of 1 mg preservative-free intrathecal midazolam added to spinal bupivacaine during postoperative analgesia, and the incidence of adverse effects, if any, in patients undergoing knee arthroscopies.

Method: 60 consenting American Society of Anesthesiologists (ASA) physical status I or II patients of either gender (men = 48, women = 12), aged between 20-60 years, were randomly allocated to two groups (30 each). Group B received 0.5% hyperbaric bupivacaine with saline intrathecally, and Group BM received 0.5% hyperbaric bupivacaine with preservative-free midazolam 1 mg intrathecally. Peak sensory level, total duration of analgesia, duration of motor blockade, pain score using the Visual Analogue Scale, and sedation score using the Observer Assessment Score of Sedation were assessed, along with vital parameters, namely heart rate and systolic, diastolic and mean blood pressure.

Results: The total duration of analgesia observed was significantly higher in Group BM (336.2 ± 37.50 minutes) vs. Group B (151.17 ± 15.41 minutes), and the pain score was lower in Group BM (33.6 ± 4.68 mm) vs. Group B (56.6 ± 8.64 mm).

Conclusion: The addition of preservative-free midazolam 1 mg to intrathecal 0.5% hyperbaric bupivacaine prolongs the duration of analgesia without any observed adverse effects in patients undergoing lower limb surgeries.

Keywords: Midazolam, Spinal Anaesthesia, post operative analgesia

1. Introduction

One of the primary aims of anaesthesia is to alleviate the patient’s pain and agony, there by permitting the performance of surgical procedures without any discomfort. Relief of postoperative pain has gained real importance in recent years considering the central, peripheral and immunological stress response to tissue injury. Any expertise acquired in this field should be extended into the postoperative period, which is the period of severe, intolerable pain requiring attention. So there is need of extended analgesia without any side effects to achieve this goal.

Intrathecal 0.5 % bupivacaine is routinely used for neuraxial blockade. Many authors have suggested the addition
of Midazolam to bupivacaine to extend the period of analgesia. Many of these studies were based on the animal models. So this study was carried out to study the effect of addition of 1mg intrathecal Midazolam to bupivacaine in humans with the objectives to compare the mean period of analgesia for intrathecal Midazolam 1mg plus bupivacaine and bupivacaine alone and to compare the side effects of intrathecal Midazolam (1mg) plus bupivacaine and bupivacaine alone.

2. Material and Methods

Patients belonging to American Society of Anesthesiologists (ASA) physical status I or II, aged between 20-60 years of either gender, and undergoing elective lower limb surgeries, were included in the study. Patients with contraindications to central neuraxial blockade, for example gross spinal deformity, spinal tenderness, local pathological conditions in the spinal area, known sensitivity to the drugs used in the study, or the presence of peripheral neuropathy, were excluded from the study.

The study was carried out in Dept. Of Anaesthesiology in J. N. Medical college, Sawangi (Meghe), Wardha from 01/05/2011 to 30/09/2012. The study was a prospective, randomised, double-blind placebo-controlled study. A prior power analysis revealed that 30 patients were needed in each group to provide a power of 80% with a significance level (p-value) taken as 0.05. A total of 60 patients were enrolled in the study.

All the patients enrolled in the study were randomly assigned to either of the two groups by drawing lots. The two groups were termed Group B and Group BM.

Group B - 3.5ml of 0.5% Bupivacaine (heavy) (17.5mg)+ 0.2 ml normal saline intrathecally.

Group BM - 3.5ml of 0.5% Bupivacaine (heavy) (17.5mg) + 0.2ml of preservative free Midazolam (1mg) intrathecally.

On the day of surgery, patients were transferred to the operating room. Standard monitors, which included an electrocardiograph (monitoring lead II and V5), a noninvasive blood pressure monitor and a pulse oximeter, were connected, and baseline vitals were recorded before the induction of spinal anaesthesia. All patients received 15 ml/kg intravenous Ringer’s lactate before induction of the spinal anaesthesia.

The spinal anaesthesia drug solutions were prepared by Observer 1, a consultant anaesthesiologist, or anaesthesiology resident, with more than one year’s experience in the field of anaesthesiology, and who did not partake in subsequent study observations. Spinal blockade was performed in all patients at the L3-4 interspace by Observer 2 (a primary investigator), adhering to the standard technique in the right lateral position, utilising a 25G Quincke spinal needle.

After confirmation of the free flow of clear cerebrospinal fluid (CSF), each patient received either a combination of 3.5 ml of 0.5% hyperbaric bupivacaine with preservativefree midazolam 1 mg (0.2 ml) (Group BM), or 3.5 ml 0.5% hyperbaric bupivacaine with 0.2 ml 0.9% saline (Group B). The time of intrathecal injection was noted, and all the patients were turned supine immediately. Supplemental oxygen (2 l/minute by nasal prongs) was administered to all the patients. The level of spinal anaesthesia was confirmed by loss of sensation of cold. Vital parameters including heart rate, blood pressure, respiratory rate and oxygen saturation were recorded every two minutes for the first 20 minutes, and subsequently, every 10 minutes until the end of the procedure. The time taken for sensory spinal level regression by the two segments was noted. Pain scores using the VAS were recorded in the postoperative period by Observer 2 every 30 minutes after the end of the procedure for the first six hours, and subsequently, every two hours for the next six hours, or until the patient received rescue analgesia, thereby ending the study. Rescue analgesia was provided with the injection of tramadol 1 mg/kg with an antiemetic (ondansetron 4 mg) on patient request, or when the VAS score was 40 mm or more, and the time was noted. The duration of analgesia (painfree period) was recorded as the time from completion of the spinal blockade to the time when rescue analgesia was administered in the postoperative period. Sedation was assessed in the perioperative period using the Observer Assessment Score of Sedation (OASS):
0 - Awake.
1 - Sleeping comfortably, but easily arousable.
2 - Deep sleep, but arousable.
3 - Deep sleep, and not arousable.

Motor blockade was assessed by the Bromage Scale, as shown below:
1 - Able to lift and extend leg.
2 - Decreased knee flexion, but full extension of feet and ankles.
3 - Flexion of ankle and feet only.
4 - Cannot move knee, leg, ankle or toes.

After administration of the spinal anaesthesia, hypotension (defined as a decrease in systolic blood pressure more than 20% below baseline) was treated by an increase in the rate of intravenous fluid administration and/or the intravenous injection of mephenetermine 3 mg boluses. Bradycardia (defined as a heart rate of less than 60 beats per minute) was treated with an intravenous injection of atropine sulphate 0.6 mg. The incidence of nausea, vomiting, shivering, urinary incontinence, respiratory depression, and any other neurological sequelae during the observation period, was recorded.

3. Results

The data were analysed using the SPSS version 10.0 for Windows®. No patients experienced complications during the study or observation period, and no patients were excluded from the study.

Demographic data were compared using the chi-square test for gender and age, and Student’s t-test for weight and height. Haemodynamic data were compared using Student’s t-test. Patients with a 20% or more baseline change in haemodynamic parameters, sedation score and peak level of sensory blockade, were compared using the chi-square test. Sedation score ≥ 1, two-segment regression time, total duration of analgesia, total duration of motor blockade, and rate of complications between the two study groups, were compared using Student’s t-test. P-values < 0.05 were considered as significant, < 0.01 as highly significant, and < 0.001 as very highly significant.

There was no statistical difference in the patient characteristics relating to age, weight, height and gender distribution between the two groups (see Table I).

<table>
<thead>
<tr>
<th>Variable</th>
<th>Group B Mean ± S.D</th>
<th>Group BM Mean ± S.D</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>34.13 ± 12.60</td>
<td>36.13 ± 13.3</td>
<td>0.364</td>
</tr>
<tr>
<td>Weight</td>
<td>55.97 ± 5.20</td>
<td>56.10 ± 6.23</td>
<td>0.269</td>
</tr>
<tr>
<td>Height</td>
<td>164.34 ± 7.15</td>
<td>169.32 ± 7.97</td>
<td>0.240</td>
</tr>
<tr>
<td>Male/Female</td>
<td>23/7</td>
<td>25/5</td>
<td></td>
</tr>
</tbody>
</table>

The addition of 1 mg midazolam to intrathecal bupivacaine prolongs the duration of postoperative analgesia (p-value 0.001) (see Table II).

<table>
<thead>
<tr>
<th></th>
<th>Group B Mean ± S.D.</th>
<th>Group BM Mean ± S.D.</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total duration of analgesia (min)</td>
<td>151.17 ± 15.41</td>
<td>336.2 ± 37.50</td>
<td>&lt; 0.001</td>
</tr>
</tbody>
</table>

The pain score using the VAS between two groups (see Table III) was clinically significant, and statistically very highly significant (p-value < 0.001).
Table III: Pain Visual Analogue Scale scores (mm)

<table>
<thead>
<tr>
<th></th>
<th>Group B Mean + S.D.</th>
<th>Group BM Mean + S.D.</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>VAS score</td>
<td>58.8 ± 8.64</td>
<td>34.7 ± 4.68</td>
<td>&lt; 0.001</td>
</tr>
</tbody>
</table>

The two-segment regression time comparison (see Table IV) indicated a significantly prolonged regression in Group BM. The difference between the groups was statistically very highly significant (p-value < 0.001).

Table IV: Two-segment regression time (minutes)

<table>
<thead>
<tr>
<th></th>
<th>Group B Mean + S.D.</th>
<th>Group BM Mean + S.D.</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Two segment regression</td>
<td>70.20 ± 5.21</td>
<td>110.12 ± 8.26</td>
<td>&lt; 0.001</td>
</tr>
</tbody>
</table>

The addition of 1 mg midazolam to intrathecal bupivacaine did not have any effect on the peak level of sensory blockade achieved.

The peak sensory levels achieved in both the groups were comparable without any statistical difference (see Table V).

Table V: Peak level of sensory blockade

<table>
<thead>
<tr>
<th>Peak level of sensory blockade</th>
<th>Group B Mean + S.D.</th>
<th>Group BM Mean + S.D.</th>
<th>Total number of patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>T4</td>
<td>4</td>
<td>6</td>
<td>10</td>
</tr>
<tr>
<td>T6</td>
<td>10</td>
<td>9</td>
<td>19</td>
</tr>
<tr>
<td>T8</td>
<td>8</td>
<td>6</td>
<td>14</td>
</tr>
<tr>
<td>T10</td>
<td>5</td>
<td>6</td>
<td>11</td>
</tr>
<tr>
<td>T12</td>
<td>3</td>
<td>3</td>
<td>8</td>
</tr>
<tr>
<td>Total</td>
<td>30</td>
<td>30</td>
<td>60</td>
</tr>
</tbody>
</table>

P-value = 0.354 for systolic blood pressure; p-value = 1 for diastolic blood pressure, mean blood pressure and heart rate. Therefore, there was no difference between the haemodynamic changes (≥ 20% change) when comparing the two groups (see Table VI).

Student’s t-test: p-value = 1, so no statistical significant difference is present between the two groups regarding adverse effects (see Table VII).

Table VI: Twenty per cent change of haemodynamic parameters from the baseline

<table>
<thead>
<tr>
<th></th>
<th>Group B Mean + S.D.</th>
<th>Group BM Mean + S.D.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean blood pressure (yes/no)</td>
<td>23/7</td>
<td>23/7</td>
</tr>
<tr>
<td>Heart rate (yes/no)</td>
<td>23/7</td>
<td>22/8</td>
</tr>
<tr>
<td>Systolic blood pressure (yes/no)</td>
<td>20/10</td>
<td>21/9</td>
</tr>
<tr>
<td>Diastolic blood pressure (yes/no)</td>
<td>18/12</td>
<td>19/11</td>
</tr>
</tbody>
</table>

Table VII: Adverse effects

<table>
<thead>
<tr>
<th></th>
<th>Group B Mean + S.D.</th>
<th>Group BM Mean + S.D.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Shivering</td>
<td>3</td>
<td>2</td>
</tr>
<tr>
<td>Nausea and vomiting</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Desaturation</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Urine retention and neurological sequel</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>
4. Discussion

In this study, it was found that the analgesic effect of intrathecal heavy bupivacaine 0.5% was augmented by the addition of intrathecal preservative-free midazolam. The addition of 1 mg midazolam prolonged the anaesthetic duration for lower limb surgeries by approximately 3-3.5 hours, without an increase in the sensory level of blockade compared to the control group. In addition, the midazolam treated group did not exhibit an increased incidence of haemodynamic instability, or observable adverse effects.

Spinal anaesthesia is the most commonly used regional anaesthetic technique. Local anaesthetic agents used for this purpose provide good intraoperative analgesia. However, they provide a very limited postoperative duration of action. In order to overcome this problem and to maximise the duration of analgesia, many adjuvants, for example opioids, neostigmine, ketamine and clonidine, have been tried increasingly in the last two decades to relieve postoperative pain. However, side-effects in the postoperative period such as nausea, vomiting, pruritus, urinary retention and respiratory depression, render most adjuvants as less than ideal. The rationale for the use of intrathecal midazolam focuses on the awareness that it is an agonist at the benzodiazepine binding site, a subunit of the pentameric gammaaminobutyric acid (GABA)A receptor. Agonist occupancy of the benzodiazepine binding site enhances the activity of GABA at the GABAAR receptor. This receptor is a chloride ionophore that, when activated, typically stabilises the transmembrane potential at, or near, the resting potential. In neurons, this typically serves to decrease excitability.

Intrathecal benzodiazepine-induced analgesia is spinally mediated. Binding sites are GABA receptors, abundantly present in the dorsal root nerve cells, with the maximum concentration found within lamina II of the dorsal nerve cells, a region that plays a prominent role in processing nociceptive and thermoceptive stimulation. The present cumulative experience with intrathecal midazolam across species broadly confirms the safety thereof, the analgesic activity of the molecule and its benzodiazepine pharmacology, and the lack of irreversible effects.

The reduction in the pain scores observed in this study is comparable to the results of a previous study by Bhattachary et al. In a study by Kim et al, VAS scores were lower in the midazolam group, but this was not found to be statistically significant.

We observed the sedation scores between the two groups at various time intervals to be comparable. The sedative effect peaked at 30 minutes, and none of the patients had a sedation score above 2. No clinically significant sedation was recorded postoperatively in either of the groups. In a study by Adam et al, the addition of midazolam to intrathecal bupivacaine was associated with sedation and tranquility. The paucity of studies on intrathecal midazolam warrants caution in elderly patients, the obese, and those who are already on other sedative medication. When intrathecal midazolam is used, all patients should be closely monitored intra- and postoperatively. The findings of this study are consistent with those of earlier reported studies by Bharti et al and Batra et al, who found that the duration of sensory blockade was significantly prolonged after the addition of midazolam to intrathecal bupivacaine. The addition of intrathecal midazolam did not significantly alter the peak level of sensory blockade that was achieved after the onset of spinal anaesthesia. This is considered a significant finding regarding the safety aspects of the combination of these two drugs, and is consistent with other studies done earlier in this regard. The duration of motor blockade was comparable in both groups in this study. There is a paucity of and variation in the literature reports with respect to the duration of the motor blockade. Many earlier studies did not find an increased duration of motor blockade after the addition of midazolam to intrathecal bupivacaine. Further studies on larger samples are needed to evaluate and confirm this finding. The present study is consistent with other studies conducted by Goodchild et al and Batra et al, who found no added haemodynamic changes, such as hypotension and bradycardia, or shivering, respiratory depression, desaturation, pruritus, urinary retention, nausea or vomiting from the baseline when intrathecal midazolam was added to intrathecal bupivacaine. The addition of 1 mg midazolam to intrathecal bupivacaine causes no significant haemodynamic disturbances, and is relatively free from common side-effects. Erdine et al conducted neurotoxicologic studies in animals by studying histologic and vascular lesions in animal spinal cord samples, indicating the neurotoxic effects of intrathecal midazolam. Therefore, they advised against the use of intrathecal midazolam in humans. Subsequent studies in humans, by Tucker et al, Valentine et al and Anjana et al, found no adverse neurological symptoms in those who had received intrathecal midazolam. In agreement with these studies, the present study observed no significant adverse neurological effects in any patient during the study period.
5. Conclusion

So this study has shown that addition of 0.2 ml midazolam (1 mg, preservative free) to bupivacaine significantly increases the period of analgesia without increase in the side effects.

References


