Intraoperative Sedation During Epidural Anesthesia: Dexmedetomidine Vs Midazolam

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Citation


Abstract

Background: This study evaluated the ability of dexmedetomidine to provide sedation during epidural anesthesia compared with midazolam, examining the cardiorespiratory variables, analgesic requirements and side effects.

Patients and Methods: Sixty patients undergoing inguinal herniorrhaphy under regional anesthesia were randomized into two groups to receive either dexmedetomidine or midazolam for intraoperative sedation. Cardiorespiratory effects, level of sedation, quality of analgesia, time to first analgesic requirement were evaluated.

Results: There were significant declines in HR and MAP values compared to baseline in both groups but the difference between groups was not significant. There were higher sedation levels in midazolam group and 16 patients receiving midazolam needed dose adjustment. The time to first analgesic requirement was significantly longer in dexmedetomidine group.

Conclusion: Supplement of intravenous dexmedetomidine in patients receiving epidural anesthesia may provide a good sedative effect and postoperative pain management without any clinically important untoward cardiorespiratory reactions.

INTRODUCTION

Central neuraxial anesthesia is a widely used method and may be associated with stress, anxiety and even embarrassment causing intraoperative discomfort. Although some patients tolerate being awake during surgical procedures without any medication, in some patients sedatives are required to limit discomfort. Administration of a sedative agent however, is associated with a risk of side effects, especially cardiorespiratory problems.

Pharmacologic agents that create an adequate level of sedation without any clinical side effects are of increasing interest to clinicians. Potential desirable effects include decreased requirements of anesthetics and analgesics, a diminished sympathetic response to stress and the potential for cardioprotective effects against myocardial ischemia with minimal effects on respiration. Although a primary indication for dexmedetomidine has been the sedation of critically ill patients, it can also be used for intraoperative sedation.

The purpose of this study was to evaluate the cardiorespiratory end-points of dexmedetomidine and midazolam in providing sedation during epidural anesthesia. Postoperative analgesia requirements and satisfactory outcomes were also investigated.

PATIENTS AND METHODS

This prospective, randomized, double-blind study was conducted with a population of patients undergoing elective inguinal herniorrhaphy. After Institutional Ethics Committee approval and written informed consent of the participants, a total of 60 adult male patients (aged 30-65 yr, American Society of Anesthesiologists-ASA-physical class I-II)
enrolled in the study. All patients have normal renal, hepatic function and no history of allergy or chronic use of medical therapy. The patients with second or third degree of heart block, current history of psychiatric disorders, history of sleep apnea, or patients with a body mass index greater than 40 kg/m² were excluded.

All patients received no premedication and monitored by non-invasive blood pressure, electrocardiogram (ECG) and pulse oxymetry on arrival to the operating room. A computer-generated randomization list was used to assign patients to one of two study groups. Before the insertion of epidural catheter, the patients in the first group (group D) received sedation with an intravenous loading dose of 1 µg.kg⁻¹ dexmedetomidine and the second group (group M) received 0.04 µg.kg⁻¹ midazolam via a syringe infusion pump over a 10-min period. After then, an epidural catheter was inserted at L₄-₅ with loss-of-resistance to saline and the patient in the lateral decubitus position. A 4 ml test dose of 2% lidocaine was given followed by 75 mg 0.5% plain bupivacaine. When the analgesia level became adequate for surgery continuous infusions of 0.5 µg.kg⁻¹ h⁻¹ dexmedetomidine and 0.04 µg.kg⁻¹ h⁻¹ midazolam were started in study groups respectively. Drug infusions were discontinued if one of the following adverse events was observed: apnea lasting longer than 20 s, hemoglobin oxygen saturation lower than 90%, decrease of heart rate (HR) below 50 beats. min⁻¹, mean arterial pressure (MAP) below 30% of the initial value. The evaluation of quality of sedation was based on a six point Ramsay Sedation Score (RSS) and according to the sedation level infusion dose was decreased to one half or increased to twice to maintain the RSS≤4. The quality of analgesia was assessed by using a 100-mm visual analog scale (VAS) in which 0 represents no pain at all and 100 represents incredible pain. If the patient reported pain exceeding 60 mm on the scale, intravenous fentanyl in doses of 0.05 mg was administered. Oxygen was delivered by a facemask 5 L.min⁻¹ to all patients throughout the procedure. Administration of any medication apart from the study protocol and occurrences of complications and side effects were recorded.

Sedation and monitoring were performed by the same anesthesiologist in all cases but assessments were performed by an individual who was blinded to the study drug. Surgeons were asked about their satisfaction with neuromuscular relaxation during the procedure. The following parameters were measured continuously: heart rate (HR), respiratory rate (RR), mean arterial pressure (MAP), hemoglobin oxygen saturation (SpO₂). The recorded data were analyzed and averaged over the following time intervals: before injection of study drug (baseline), at least 3 min later from the first injection of study drug, after epidural administration of bupivacaine and every 10 minutes from the start to the end of surgery (at which the infusions were discontinued). RSS and VAS was assessed during epidural catheter implantation, intraoperative period (VAS was assessed until RSS reached to score 4), the post-anesthesia care unit at the 30th and the 60th minutes. The patients were transferred to ward when RSS was 2 point. A 12-h follow up was made to assess the analgesic requirements of the patients and ask their willingness to undergo a repeat procedure with the same anesthetic regimen in the future if required.

Statistical analysis: Statistical analyses were performed using Statistica for Windows version 10.0 software. Results were expressed as members of occurrences, percentages and mean ± SD. With a 2-sided type I error of 5 % and study power at 80%, the number of patients required in each group to demonstrate a difference between groups was 25. Repeated measures analysis of variance was used to compare continuous variables. The difference in continuous parameters such as patient characteristics, preoperative data and amount of supplemental analgesic were analyzed using one-way analysis of variance or Kruskal Wallis test for non-parametric quantitative data. A p values less than 0.05 was considered significant.

RESULTS
In all patients the study was completed without any serious complication. The study groups were comparable regarding to ASA physical status, demographic characteristics, initial vital signs, maximum analgesia level and duration of operation (Table 1).
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Figure 1
Table 1: Patient demographics, preoperative and operative data.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Group D</th>
<th>Group M</th>
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<tbody>
<tr>
<td>ASA I (%)</td>
<td>26 (66.6%)</td>
<td>23 (76.6%)</td>
</tr>
<tr>
<td>Age (yr)</td>
<td>42.36 ± 13.14</td>
<td>46.32 ± 11.03</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>74.00 ± 11.24</td>
<td>69.16 ± 9.55</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>171.33 ± 7.83</td>
<td>170.16 ± 5.88</td>
</tr>
<tr>
<td>MAP (mmHg)</td>
<td>102.27 ± 11.74</td>
<td>98.20 ± 13.80</td>
</tr>
<tr>
<td>HR (beat min⁻¹)</td>
<td>77.36 ± 13.43</td>
<td>71.60 ± 11.07</td>
</tr>
<tr>
<td>SpO₂ (%)</td>
<td>97.27 ± 1.68</td>
<td>96.40 ± 1.82</td>
</tr>
<tr>
<td>RR (micro⁻¹)</td>
<td>16.87 ± 3.63</td>
<td>16.13 ± 3.32</td>
</tr>
<tr>
<td>Max analgesia level (T)</td>
<td>9.27 ± 1.23</td>
<td>9.15 ± 1.12</td>
</tr>
<tr>
<td>Duration of operation (min)</td>
<td>42.90 ±11.60</td>
<td>40.60 ± 6.97</td>
</tr>
</tbody>
</table>

Data were presented as n or mean ± SD.

There were significant declines in HR and MAP values compared to baseline in both groups but the difference between groups was not significant (p>0.05). Atropine requirements between the groups were not significant. MAP was significantly reduced in one patient in both groups and treated with ephedrine. The variations in SpO₂ and respiratory rate were negligible in both groups (Figure 1).

Figure 2

Figure 1: Cardiorespiratory variables during the intraoperative period. Mean values of mean arterial pressure, heart rate, SpO₂ values, respiratory rate in determined times. ABD: after bolus drug, BEB: before epidural block, DEB: during epidural block, PEB: postepidural block, ES: end of surgery, PO: postoperative period.

RSS were significantly higher in group M during the intraoperative period and dose reduction of the drug was required in 16 patients. In group D only one patient required dose adjustment. There were no differences between treatment groups at postoperative 30ᵗʰ and 60ᵗʰ min in respect to sedation scores. VAS during epidural catheter implantation and the intraoperative period was decreased significantly in both groups. The patients receiving dexmedetomidine presented lower VAS values in the postanesthesia care unit but the difference between groups throughout the postoperative period was not significant (Figure 2). No patient required supplemental fentanyl.
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**Figure 3**

Figure 2: Comparison the level of sedation according RSS and intensity of pain according to VAS.

The number of patients (n=24 in group D, n=20 in group M) and surgeon (n=22 in group D, n=13 in group M) who revealed a good satisfaction from anesthetic technique were similar. The time to first analgesic requirement was significantly longer (p<0.01) in group D (487.27 ± 201.25 min and 278.54 ± 153.48 min respectively). No other side effects or administration of medication other than those in the study protocol were recorded.

**DISCUSSION**

The present randomized, double-blinded study demonstrated that loading dose of 1 µg.kg⁻¹ dexmedetomidine followed by 0.5 µg.kg⁻¹ h⁻¹ infusion provided a good and stable sedative effect during epidural anesthesia and manifested a longer time to first analgesic requirements in patients undergoing inguinal herniography. Loading dose of 0.04 µg.kg⁻¹ midazolam followed by 0.04 µg.kg⁻¹ h⁻¹ infusion resulted in higher sedation scores in patients whose the heavy sedation was not necessary. Cardiovascular stability and respiratory function were well maintained in both study groups.

During surgical procedures, both under- and over- sedation carry inherent risk, the former increases the likelihood of recall and agitation-induced sympathetic activation, and the latter, excessive depression of vital physiologic functions (⁹). It’s important to distinguish the sedation scales used to assess the sedation during surgical procedures rather than in patients in intensive care units, because the aim of intraoperative sedation is to provide calmness more than decrease the level of consciousness (⁹). Selection of sedation agents largely depends on physician preference. A wide variety of centrally-active drugs are used to provide sedation, anxiolysis, and amnesia. There is a growing interest in the use of alpha-2 adrenoceptors agonists as sedatives. Dexmedetomidine is a currently used agent because of its short half-life, sedation, analgesic properties and favorable cardiopulmonary effects (⁹). It’s sympatholytic effect is manifested by decreases in arterial blood pressure, heart rate and norepinephrine release (w/Re11).

It has been previously reported that the use of dexmedetomidine in colonoscopy (⁰), intravenous sedation (¹), awake craniotomy (⁴,⁵), carotid endarterectomy (⁶), fiberoptic intubation (⁷) and intravenous regional anesthesia (⁸) provided satisfactory sedation, intra- and post-operative analgesia and hemodynamically stable perioperative period. However, only a limited number of reports describe the use of a-2 receptor agonists for intraoperative sedation during regional anesthesia. Systemically administration of dexmedetomidine may prolong the duration of spinal anesthesia depending on activation of a-2 adrenoceptors. Supplementation of intravenous dexmedetomidine during spinal anesthesia may be beneficial to overcome the discomfort of the patients especially in prone position (⁹). In comparison to propofol, dexmedetomidine achieved similar levels of sedation with a slower onset and offset of sedation, comparable respiratory changes and more stable hemodynamic parameters. Blood pressure and heart rate decreased in both groups of our patients but were not significant between groups. These decreases were not only depending on the drug infusions but also decreased sympathetic reflex and release from anxiety.

As previously reported, dexmedetomidine may provide better analgesia for postsurgical pain compared with widely used drugs (w/Re20). Although the patient satisfaction with their procedure is impacted by multiple variables, intra and post-operative pain control is the primary determinant. Regional anesthesia may be somewhat painful itself and require analgesic medication (⁴,⁵). Dexmedetomidine administration also reduced the discomfort during epidural catheter replacement and postoperative analgesic requirements in our patients. Therefore, it may be advantageous for the recovery and satisfaction point of view. It was considered that, dexmedetomidine, in addition to its sedative effect is a good analgesia-sparing agent.

In conclusion, during epidural anesthesia loading dose of µg.kg⁻¹ dexmedetomidine followed by continuous infusion
of 0.5 µg·kg⁻¹·h⁻¹ may be beneficial for the providing stable sedation, hemodynamics and respiration together with the good postoperative analgesia. Efficacy of intravenous dexmedetomidine during intra-operative period needs to be researched in a large number of patients.

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