Introduction

Spinal anesthesia is the most commonly used technique for cesarean section as it is very economical and easy to administer. It reduces mortality rate associated with cesarean section by sixteen times when compared with general anesthesia. Spinal anesthesia avoids the risks of general anesthesia such as aspiration of gastric contents, difficulty with airway management, infant respiratory distress, and mothers’ awareness during operation [1].

Increasing the duration of local anesthetic drugs often is a matter of concern for anesthesiologists. Epinephrine, phenylephrine, clonidine, opioids, etc. are used to increase the duration of spinal anesthesia [2, 3]. Adding epinephrine to the local anesthetics may lead to tachycardia, pallor, and hypertension [1]. Opioids control postoperative pain well; however, they are followed by nausea, vomiting, drowsiness and pruritus. At present, there is no drug able to control pain specifically without having side effects [2, 3]. The use of corticosteroid compounds increases duration of anesthesia and analgesia in peripheral nerve blocks [4, 5]. In addition, intravenous (IV) and oral dexamethasone considerably alleviate postoperative pain [6-8]. Epidural and intrathecal steroids are used to reduce chronic pain [9-11]. In some studies, intrathecal dexamethasone increased duration of sensory block and postoperative analgesia [12-14]. Although intrathecal dexamethasone is used to control chronic pain; few studies have been conducted on the effects of sensory block and postoperative pain in patients undergoing surgery. The purpose of this investigation was to evaluate the effect of conjugation of dexamethasone or epinephrine with lidocaine on the duration and onset time of spinal anesthesia.

Materials and Methods

In this double-blind clinical trial research, 90 ASA class I female patients aged 18-35 years who were candidates for cesarean section were randomly divided into three groups of control group, epinephrine group, and dexamethasone group. To do so, a table was created by a computer using random numbers. Patients with history of long-term steroid therapy, allergy to the drugs, uncontrolled hypertension, neurological or psychological disorders, spinal column surgery, low back pain, opium addict or using any drug that modifies pain perception were excluded from the study.

After ethic committee approval, written informed consent was obtained from each patient preoperatively. In order to compare the sensory duration of dexamethasone or epinephrine added to lidocaine, the sample size was determined to be 30 persons in each group. We used α=5% and β=80% [12, 13]. After intravenous line preparation, a 5 cc/kg lactated ringers solution was infused to all patients. Patients received no premedication, and upon arrival of patients into the operating room, ECG, peripheral oxygen...
saturation (SPO 2), and noninvasive arterial blood pressure (NIBP) were monitored and recorded at 5-minute intervals until the end of surgery and vital signs were recorded every 15 minutes in the Post Anesthesia Care Unit (PACU).

Spinal anesthesia was performed in a sitting position using 25G Quincke needle in L3-L4 or L4-L5 intervertebral disk space. In the control group, 75 mg lidocaine and 1ml saline was injected. In the epinephrine group, 75 mg lidocaine (made in Finland) and 100 μg epinephrine (made by Iranian Darou Pakhsh Holding Company) was injected. In the dexamethasone group, 75 mg lidocaine and 4 mg dexamethasone (2.5 ml) (made by Iran Hormone Company) was injected into intrathecal space. Immediately after injection of the drugs, the patients were laid in the supine position and a pillow was put under the right buttock of each patient. Five to six liters per minute of oxygen was given by mask. The sensory block level was assessed by a pin prick test by a short bevel needle along the mid-axillary line bilaterally. The sensory block level was controlled every 30 seconds for 20 minutes; then it was evaluated every 5 minutes until a 4 sensory level regression from highest level or to the end of the surgery. Onset time was defined from the time of injection of drugs into the intrathecal space to the peak of sensory block (highest dermatome level) and the duration of sensory block was defined from peak of sensory block up to 4 sensory level regressions or when the patients feel pain in the field of surgery.

After the end of sensory block, pain level was measured with respect to Visual Analogue Scale (VAS) (zero=no pain, 10=the most severe pain the individual has suffered) per hour. If the postoperative VAS was higher than 4, it was treated by morphine 2 mg IV. Analgesia duration was measured by one-way analysis of variance (ANOVA) and Tukey post hoc test. Sensory block duration and postoperative analgesia were measured by χ² test.

Results

One patient in dexamethasone group was excluded from the study, as she was put under general anesthesia due to uterine atony and needed hysterectomy. The information obtained from 89 patients was statistically analyzed. There were no differences between the means of age, weight and height in the three groups (Table 1). Onset of sensory block in lidocaine group was considerably shorter than dexamethasone and epinephrine groups. Sensory block duration in dexamethasone and epinephrine groups was longer than that in lidocaine group (Table 2). Postoperative analgesia in epinephrine group was longer than lidocaine and dexamethasone groups. In lidocaine group, duration of postoperative analgesia was shorter than that in epinephrine and dexamethasone groups (Table 2). Twenty-five patients (86%) in dexamethasone group, 15 patients (50%) in lidocaine group, and 14 patients (46.7%) in epinephrine group had considerable hypotension (p=0.003). Figure 1 shows changes of blood pressure in the three groups. Need for ephedrine in the control group, epinephrine group and dexamethasone group were 7.67±6.89, 7.5±8.67, 9.48±8.59 mg, respectively (p=0.58). Three patients in dexamethasone group and one patient in the control group suffered from bradycardia and were received atropine injection. Incidence of nausea and vomiting in dexamethasone group was significantly lower than the other two groups (p<0.05) and no neurological and infectious complication in patients were reported (Table 3).

<table>
<thead>
<tr>
<th>Table 1. Comparison of characteristics of patients in the three groups</th>
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<td>Age (yr)</td>
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<td>Weight (kg)</td>
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<td>Height (cm)</td>
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<th>Table 2. Comparison onset of sensory block, duration of sensory block and analgesia in the three groups</th>
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<td></td>
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<tr>
<td>Onset of sensory block</td>
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<tr>
<td>Sensory block duration (min)</td>
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<td>Analgesia duration (min)</td>
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Table 3. Distribution of nausea, vomiting, in the three study groups

<table>
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<tr>
<th></th>
<th>Lidocaine N(%)</th>
<th>Lidocaine+dexamethasone N(%)</th>
<th>Lidocaine+epinephrine N(%)</th>
<th>p-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nausea</td>
<td>16(53.3)</td>
<td>5(17.2)</td>
<td>10(33.3)</td>
<td>0.014</td>
</tr>
<tr>
<td>Vomiting</td>
<td>12(40)</td>
<td>3(10.3)</td>
<td>5(16.7)</td>
<td>0.016</td>
</tr>
</tbody>
</table>

* p<0.001one-way analysis of variance (ANOVA)
** p<0.001 Compared with lidocaine (Tukey post hoc test)
*** p<0.05 Compared with lidocaine (Tukey post hoc test)
Figure 1. Comparing changes of blood pressure in the three groups

Discussion

In this study, adding 100 µg epinephrine or 4 mg dexamethasone to 75 mg intrathecal lidocaine increased sensory block duration; however, it did not delay the onset of sensory block.

Many studies have confirmed analgesic effects of steroids in peripheral and neuroaxial blocks [10-14]. Movafegh et al. reported that the addition of dexamethasone (8 mg) to lidocaine for spinal anesthesia provided significant prolongation of sensory and motor block in comparison with plain lidocaine and there is no difference between dexamethasone-lidocaine 5% and epinephrine (0.2 mg) - lidocaine 5% in sensory and motor block duration. Indeed, lidocaine-dexamethasone is not followed by any kind of hemodynamic, neurological complications or delay in the onset of sensory and motor block [13]. In another study, adding 8 mg dexamethasone to plain bupivacaine increased the duration of sensory block and postoperative analgesia and had no effect on the onset of sensory block [12]. However, in a study carried out by Abdel-Aleem et al., adding 8 mg dexamethasone to intrathecal bupivacaine had no effect on VAS and postoperative consumption of narcotics in patients candidate for cesarean section [14]. Mirzaei et al. showed that injecting epidural corticosteroid and bupivacaine considerably reduces back pain after laminectomy surgery [9].

The study conducted by Kotani et al. revealed that intrathecal methylprednisolone and bupivacaine create excellent analgesia after restoring burns and injuries to organs [15]. Moreover, 5 mg intrathecal dexamethasone reduces postoperative anesthesia and morphine consumption after laparoscopy surgery [16]. Dexamethasone and bupivacaine in supraclavicular and axillary block significantly increase duration of analgesia [17, 18]. Analgesic mechanism of corticosteroids is not known well. It seems that controlling inflammatory or immune system has a role in increasing the duration of analgesia, as intrathecal steroids reduce production of prostaglandins by controlling phospholipase A2 and cyclooxygenase [19]. In a study conducted by Drager et al., it was hypothesized that local injection of dexamethasone could control the inflammatory response and reduce the release of inflammatory mediators (prostaglandin E2, thromboxane A2 and other prostanoids) in the affected area, possibly increasing analgesia duration of local anesthetics in this way [20]. Some researchers believe that the analgesics effects of corticosteroids are caused by their systematic effects [4]. Prolongation of the block may be due to their direct effect on nerve fiber.

In this study, intrathecal dexamethasone reduced nausea and vomiting considerably. In a research conducted in 2012, injection of 8 mg intrathecal dexamethasone reduced nausea and vomiting in the women undergoing cesarean section [14]. In the study conducted by Movafegh, there was no difference between the incidence of nausea and vomiting in lidocaine-dexamethasone group, and plain lidocaine and lidocaine-epinephrine group. The mechanism of dexamethasone effect in preventing nausea and vomiting is not clear. Receptors of glucocorticosteroids are in postrema area, which regulates nausea and vomiting. Dexamethasone might prevent nausea and vomiting by regulating neurotransmitters or density of receptor in this center [21].

No neurological complications following injection of intrathecal steroids have been reported by previous studies [22]. A study reported histopathologic changes after repeated injections of intrathecal betamethasone [23]. Kotani et al. did not report any complication following injection of intrathecal methylprednisolone [15]. Sugita did not report any complication after injection of 8 mg intrathecal dexamethasone [24]. This study shows that, similar to epinephrine, intrathecal dexamethasone increases sensory block duration of lidocaine; however, it has no effect on postoperative analgesia.

The present study shows that adding dexamethasone to intrathecal lidocaine increases sensory block duration. This can be useful for the patients with no contraindications to epinephrine.

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Authors’ Contributions

All authors had equal role in design, work, statistical analysis and manuscript writing.

Conflict of Interest

The authors declare no conflict of interest.

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References


