Sparing Effects of Sufentanil on Epidural Ropivacaine in Elderly Patients Undergoing Transurethral Resection of Prostate Surgery

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Purpose: This study examined the sparing effect of sufentanil on the median effective concentration (EC50) of epidural ropivacaine in elderly patients undergoing elective transurethral resection of prostate (TURP).

Materials and Methods: This was a prospective randomized double-blind dose-response study. Fifty eight elderly patients with American Society of Anesthesiologists physical status I–II who were scheduled for TURP surgery under epidural anesthesia were randomly allocated to a group receiving 15 mL of ropivacaine (group R) or a group receiving ropivacaine plus 5 μg of sufentanil (group RS). The concentration of ropivacaine was determined by a Dixon’s up-and-down sequential allocation. The first participant received 0.3% of ropivacaine in both groups and subsequent concentrations were determined by the response of the previous patient in the same group. The EC50 of epidural ropivacaine was analyzed using the Dixon and Massey method.

Results: The EC50 of epidural ropivacaine during TURP surgery was 0.186% (95% confidence interval, 0.173–0.200%) in group R, and 0.136% (95% confidence interval, 0.127–0.144%) in group RS (p<0.01). No differences in Bromage scale of motor block examination and the onset time of sensory block were observed.

Conclusion: Administration of 5 μg of epidural sufentanil caused a 37% reduction in the EC50 of epidural ropivacaine in elderly patients who underwent TURP surgery. Thus, sufentanil addition during surgery of TURP can decrease the dose of ropivacaine anesthesia required.

Key Words: Ropivacaine, sufentanil, elderly patients

INTRODUCTION

Lumbar epidural anesthesia is the most widely used anesthetic regimen in elderly patients during transurethral urological surgery. However, the use of epidural nerve blockade could result in shorter onset and longer duration of anesthesia as well as more intense motor blockade in elder patients.1,2 Ropivacaine is a long-acting amide local anesthetic agent that exerts effects via reversible inhibition of sodium ion influx in nerve fibres with reduced lipophilicity, which is also associated with decreased potential for central nervous system toxicity and cardiotoxicity.3 Furthermore, recent clinical data have shown that ropivacaine is a proper alternate local...
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anesthetic for spinal anesthesia in elderly patients with co-existing systemic disease for transurethral resection of prostate (TURP) operations. However, our previous study demonstrated that the \( EC_{50} \) of ropivacaine required to establish motor block decreases significantly with advancing age. Also, ropivacaine can still cause obvious cardiotoxicity when used in high concentrations. Therefore, co-administration of other pain relievers with local anesthetics has been tested in the clinic to reduce the concentration of ropivacaine used.

Previous studies have demonstrated that the addition of an opioid allows a lower concentration of local anesthetics to be used, which is beneficial during labour analgesia. Intrathecal addition of a lipophilic opioid to local anesthetics markedly improves the quality of anesthesia, prolongs the duration of analgesia, reduces the dose requirement of local anesthetics, and shortens the onset time of block. Sufentanil is a more lipophilic opioid with higher affinity for opioid receptors, has lower cephalad spread, and exhibits much higher analgesic potency compared to fentanyl or morphine. Several previous studies have shown that addition of sufentanil to ropivacaine, bupivacaine, or levobupivacaine for labor analgesia or analgesia reduced minimum local analgesic concentrations (MLAC) of epidural local anesthetics and minimized motor block. However, no study has been carried out to determine the sparing effect of sufentanil on the \( EC_{50} \) of epidural ropivacaine for elderly patients.

This study investigated the local-anesthetic-sparing efficacy of sufentanil on the epidural anesthesia of ropivacaine for elderly patients during TURP.

MATERIALS AND METHODS

Subject selection
Sixty-six American Society of Anesthesiologists physical status I–II patients scheduled for transurethral prostate and bladder procedures were enrolled in this study from May 2010 to Jun 2012. All patients were between 65 and 85 years of age with body mass index less than 25. The protocol (reference number 201006) was approved by the Ethics Committee of the First Affiliated Hospital, College of Medicine, Zhejiang University (Hangzhou, China). Signed informed consent was acquired from each participant. The exclusion criteria were: 1) patient is hypersensitive to amide local anesthetics or sufentanil, 2) patient exhibits contraindication to epidural anesthesia, and 3) patient received opioids or sedatives during the previous 12 hours.

Sample size determination
Sample size was estimated based on an averaged SD of 0.025%. Power was given at 0.80 to detect a 10% difference of 1 SD (0.025%) at \( p<0.05 \). Therefore, a minimum of 12 subjects would be needed for each of the two groups. To estimate the \( EC_{50} \), the sample size was doubled and 33 subjects were recruited for each group.

Study design
This was a prospective, randomized, double-blinded study.

Anesthesia procedures
Participants were randomly allocated to one of two groups by a random allocation table to receive 15 mL of ropivacaine (group R) (Naropin®; AstraZeneca, Sodertalje, Sweden) alone or ropivacaine with 5 \( \mu \)g of sufentanil (group RS) before insertion of the epidural catheter. The anesthesiologist performing the procedure and subsequent assessment was blinded to the concentration of anesthetics. All patients were not premedicated before anesthesia. After IV prehydration with 500 mL of lactated Ringer’s solution, patients were placed in the flexed lateral position. The epidural space at the L2‒3 or L3‒4 level was identified with the loss of resistance to air injection. Every puncture was performed with a 16-gauge Tuohy epidural needle. A motorized epidural catheter was advanced 3 cm into the epidural space in the cephalad direction. The test dose was omitted due to the study design. After negative aspiration for cerebrospinal fluid or blood, 15 mL of ropivacaine (without epinephrine) was injected within 2 min through the catheter. The patient was then placed in the horizontal supine position.

The first patient in each group received 0.3% weight/volume (w/v) of ropivacaine based on our previous study. Thereafter, the concentration of ropivacaine for each subsequent patient was determined by the response of the previous patient in the same group using an up-down sequential allocation technique. The testing increment or decrement of ropivacaine was set to 0.025% (w/v) in both groups. The study solution was freshly prepared using 0.9% saline. In the group RS, 1 mL of sufentanil (5 \( \mu \)g/mL) was used instead of equivalent saline volume to achieve the desired ropivacaine concentration in the same final 15 mL volume. Effective or ineffective responses were determined for each
Sensory changes were recorded bilaterally along the anterior axillary line by pinprick using a short beveled 25-gauge needle. Sensory level was assessed by the Hollmen scale: 0=ability to appreciate a pin pricking as sharp; 1=ability to appreciate a pin pricking as less sharp; 2=ability to appreciate a pin pricking as sharp (analgesia); 3= inability to appreciate a pin touching (anesthesia). Onset time was defined as the period from full injection of the dose to the patient showing a Hollmen scale grade 2 at T10, bilaterally. The level of sensory anesthesia was assessed by an independent observer, who was blinded to the allocations.

Motor block in the lower limbs was assessed bilaterally by the same anesthetist 30 minutes after epidural injection with a Bromage scale 27 (4=no motor block: complete flexion of knee and foot; 3=partial motor block: reduced flexion of knee, complete flexion of foot; 2=almost complete motor block: no flexion of knee, flexion of foot; and 1=complete motor block: no flexion of knee, no flexion of foot).

**Outcomes**

Effective: there was no pain and discomfort of the penis when pulling 30 minutes after epidural injection without the need for an epidural rescue bolus. This result elicited a decrement of 0.025% (w/v) ropivacaine for the next patient in the same group.

Ineffective: there was pain or discomfort of the penis when pulling 30 minutes after epidural injection, but the patient responded to a rescue bolus of 10 mL of 2% lidocaine. An increment of 0.025% (w/v) ropivacaine was used for the next patient in the same group.

Rejected: there was pain or discomfort of the penis when pulling 30 minutes after epidural injection, and the patient failed to respond to a rescue bolus. This outcome caused this patient to be excluded from subsequent analysis, and the same concentration of anesthetics was used for the next patient in the same group.

**Cardiovascular measurements**

The parameters of pulse-oximetry, electrocardiogram, and non-invasive blood pressure measurements were recorded before epidural injection and at 5-min intervals during the study, and then every 5 min throughout surgery. Hypotension was defined as a decrease in systolic blood pressure by more than 30% of the pre-anesthetic value or a systolic blood pressure <90 mm Hg. Hypotension was treated by intravenous injection of 6 mg of ephedrine and crystalloid fluid. Bradycardia (<55 bpm) was treated by intravenous injection of 0.5 mg of atropine.

**Statistical analysis**

Statistical analysis was performed using SPSS 11.5 software (SPSS Inc., Chicago, IL, USA). Student’s t-test was used for the normally distributed data, and the Mann-Whitney U test was used for non-normally distributed data. A $p<0.05$ (two sided) was considered statistically significant. The EC$_{50}$ and its 95% confidence interval of the up-down sequences were analyzed using the formula of Dixon and Massey. 28-30

**RESULTS**

A total of 66 subjects were recruited and 8 patients (4 in each group) were excluded from this study because of vascular puncture during block performance, unilateral block, and no anesthetic effect. Fifty-eight patients were included in the data analysis. No significant differences in pre-test variables such as demographic and surgical data were observed between the two groups (Table 1). No incidence of pruritus, nausea or vomiting was observed in the two groups. No significant differences in sensory block, the onset time to T10, and time to highest level of block were observed between the two groups.

**Table 1. Demographic Data of Patients and Characteristics of Epidural Anesthesia Administration**

|                      | Group R (n=29) | Group RS (n=29) |
|----------------------|---------------|-----------------|
| **Age (yrs)**        | 73 (71–77)    | 71 (69–75)      |
| **Weight (kg)**      | 66 (60–70)    | 68 (60–70)      |
| **Height (cm)**      | 169 (167–170) | 168 (164–170)   |
| **Operating time (min)** | 55 (37–74) | 63 (35–75) |
| **Sensory block to pinprick** |             |                 |
| Onset time to T10 (min) | 8 (7–9)     | 8 (7–9)         |
| Highest level of block | T6 (T6–T10) | T6 (T6–T10) |
| Time to highest level of block (min) | 11 (7–15) | 15 (8–20) |

Data were presented as median (25th–75th percentile).
between two groups.

Figs. 1 and 2 show the sequences of effective and ineffective anesthesia for patients in group R and RS. The EC$_{50}$ of ropivacaine without sufentanil for anesthesia during TURP surgery in elderly patients was 0.186% (95% confidence interval, 0.173–0.200%), while the EC$_{50}$ of ropivacaine with sufentanil was 0.136% (95% confidence interval, 0.127–0.144%). A significant difference in EC$_{50}$ was observed between the two groups ($p < 0.01$).

Hemodynamic data are shown in Fig. 3. No significant differences in hypotension, bradycardia, and respiratory depression were observed between the two groups. The motor block was determined by the modified Bromage score, and no patient in the two groups developed motor block.

**DISCUSSION**

Previous studies have demonstrated that addition of intrathecal sufentanil to ropivacaine reduced local ropivacaine concentrations or ED$_{50}$ during labor or caesarean delivery.$^{21-23}$ However, there has been no study on the effect of sufentanil on the EC$_{50}$ of epidural ropivacaine in elderly patients undergoing TURP. This study demonstrated that the EC$_{50}$ of epidural ropivacaine coadministered with 5 μg of sufentanil was 0.136% in elderly patients during TURP surgery, which was significantly lower than the EC$_{50}$ of ropivacaine without sufentanil (0.186%). To our best knowledge, this is the first study to provide evidence that 5 μg of sufentanil can significantly reduce the amount of epidural ropivacaine required for elderly patients undergoing TURP.

TURP is most often performed on elderly patients who have hypertension, other cardiovascular problems, problems in breathing, and problems in kidney functions. The mortality rate was 0.2% in the patients who underwent TURP, and increased morbidity was found in elderly patients.$^{28}$ Local anesthesia is highly preferable in elderly patients undergoing TURP due to the observation that general anesthesia causes more hemodynamic differences.$^{29,30}$ However, epidural local anesthetics can still produce high sensory and sympathetic block in elderly patients.$^{1,2}$ Although reducing the dose of local anesthetic can decrease the hemodynamic side effects during cesarean delivery,$^{14}$ a stable anesthesia regimen would be more beneficial for these patients. The addition of opioids to local anesthetics has been demonstrated to be an alternative method to establish sufficient sensory and motor block, while reducing hemodynamic side effects in elderly patients.
patients and patients with cardiovascular disease. In this study, the addition of sufentanil to ropivacaine provided sufficient anesthesia, but decreased the EC$_{50}$ of epidural ropivacaine in elderly patients during TURP surgery.

The addition of opioids to local anesthetics has become a well-accepted practice in the management of spinal anesthesia for caesarean delivery. Sufentanil is a lipophilic opioid. Epidural sufentanil injected can penetrate the lipid filled subarachnoid and epidural space and enhances the local anesthetic effect by blocking the pain impulses. Co-administration of sufentanil with epidural ropivacaine exhibits rapid onset, long duration of analgesia, and motor block in a short time. Previous studies have shown that the local anesthetic sparing ability of sufentanil can be established using epidural infusions of local anesthetic with varying doses of sufentanil ranging from 2.5 to 20 μg. However, the effect of sufentanil on the EC$_{50}$ of epidural ropivacaine in elderly patients undergoing TURP has not been reported. In this study, 5 μg of sufentanil was selected based on previous studies in caesarean delivery. However, a significant limitation of the current study is that only one dose of sufentanil was investigated, and this may not be the most effective dose. Therefore, studies on the effect of different doses of epidural sufentanil on EC$_{50}$ of epidural ropivacaine for the transurethral procedure are needed to provide more accurate choices for clinicians.

In this study, the EC$_{50}$ for ropivacaine is significantly lower than that of another study center. It is always difficult to compare EC$_{50}$ values from different centers because the patient characteristics, anesthetic practice, and surgical practice may vary. Also, differences in MLAC methodology and clinical endpoints may contribute to the observed variations. Our previous study revealed that the EC$_{50}$ of ropivacaine for motor blockade in elderly patients who underwent transurethral procedure was 0.383%. The EC$_{50}$ was higher in our previous study than in this study (0.186%). This difference may be due to the fact that the protocol in the present study focused on exploring the EC$_{50}$ of anesthesia, which might have less anesthetic requirements.

In this study, the dose of ropivacaine was adjusted according to up-down sequential allocation. The main advantage of this study design is the accuracy in determining the EC$_{50}$. However, the starting ropivacaine concentration was 0.3% based on our previous study. This ropivacaine concentration is significantly different from the actual observed value of EC$_{50}$. It would be ideal to start at a concentration close to the actual value of EC$_{50}$, such as 0.2% to improve the validity of the results. This is because ropivacaine begins to be effective at 0.125% in patients who were administered ropivacaine and sufentanil and 0.175% in patients who were administered ropivacaine only.

In conclusion, the addition of 5 μg of sufentanil significantly decreased the EC$_{50}$ of epidural ropivacaine for transurethral surgery without motor block on the lower limbs. More studies are needed to further investigate the ideal dose of sufentanil that should be coadministered epidurally with ropivacaine in elderly patients.

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