Efficacy of Ultrasound-Guided Quadratus Lumborum Block in Postoperative Analgesia of Total Hip Replacement

YING ZHANG*, LI ZHANG, L. P. WANG AND Z. Y. ZHANG

Department of Anesthesiology, The Second Hospital of Tangshan, Tangshan, Hebei Province 063000, China

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Moderate or even severe pain is frequently reported in patients after total hip arthroplasty, severely compromising early postoperative recovery. Previous studies have reported favorable results using anterior, posterior and lateral quadratus lumborum block for pain control during hip surgery. In this study, we evaluate the effectiveness of ultrasound-guided quadratus lumborum block in postoperative analgesia of total hip replacement. 200 patients who underwent total hip arthroplasty in our hospital between January 2020 and January 2021 were recruited and assigned via random number table method to receive either ultrasoundguided quadratus lumborum block (study group) or fascia iliaca block (observation group) for anesthesia. All patients were given self-administered intravenous analgesia postoperatively. Outcome measures included cumulative postoperative sufentanil use, resting pain, motor pain, postoperative hip mobility and incidence of nausea and vomiting. Ultrasound-guided quadratus lumborum block resulted in significantly less sufentanil use in patients at 24 h and 48 h postoperatively (52.14±5.11, 105.74±8.14) vs. fascia iliaca block (77.58±7.93, 150.18±10.58) (p<0.05). Patients in the study group showed lower visual analogue scale scores both at rest and at exercise than those in the observation group at 12, 24 and 48 h postoperatively (p < 0.05). The study group had significantly greater maximum hip flexion and abduction mobility at 24 h and 48 h postoperatively than the observation group (p<0.05). Ultrasound-guided quadratus lumborum block was associated with a lower incidence of nausea and vomiting (7.00 %) vs. fascia iliaca block (39.00 %) (p<0.05). Ultrasoundguided quadratus lumborum block reduces the use of opioids in total hip replacement patients, significantly alleviates surgical pain, promotes early functional recovery and lowers the incidence of adverse effects.

Key words: Ultrasound, quadratus lumborum block, hip joint, analgesia, fracture

Total hip arthroplasty is a common procedure for the treatment of severe hip disease and reconstruction of joint function^[1,2]. However, the severe postoperative pain seriously compromises postoperative recovery, which underlines the significance of preoperative and postoperative analgesia for hip fracture^[3]. The PROSPECT 2010 guidelines recommend various methods to minimize perioperative pain during total hip arthroplasty in elderly patients, such as intravenous analgesia, epidural analgesia, local anesthetic infiltration techniques and peripheral nerve blocks. Nevertheless, due to the complex innervation of the hip, there is a critical need for a new approach to block the surgical area of the hip while preserving muscle motility.

Existing analgesic modalities for hip fractures rely on ultrasound-guided nerve block techniques as well as intravenous and oral analgesics^[4]. Currently, peripheral nerve block is a key component of perioperative multimodal analgesia that provides site-specific and fast-acting analgesia^[5]. Quadratus Lumborum Block (OLB) is new multimodal analgesia with established postoperative analgesic effects in abdominal surgery in recent years^[6,7]. Since the muscles and skin sensation involved in total hip arthroplasty are innervated by branches of the superior knee nerve, the inferior costal nerve, the inferior ilioinguinal nerve, the ilioinguinal nerve, the femoral nerve, the foramen ovale nerve, the sciatic nerve and the lateral femoral cutaneous nerve, which accounts for the difficulty of QLB to meet the anesthetic needs in total hip arthroplasty^[8,9]. Furthermore, the effectiveness of QLB in hip arthroplasty has been marginally explored.

To this end, this study was conducted to evaluate the effectiveness of ultrasound-guided QLB in postoperative analgesia of total hip replacement and to explore a more appropriate analgesic solution for better rehabilitation surgery.

MATERIALS AND METHODS

General information:

In this retrospective study, 200 patients (127 males and 73 females, aged under 80 y) who underwent total hip arthroplasty in our hospital between January 2020 and January 2021 were recruited and assigned *via* random number table method to receive either ultrasound-guided QLB (study group) or fascia, all patients received self-administered intravenous analgesia postoperatively. Patients provided written informed consent and the study was approved by the ethics committee of our hospital.

Inclusion and exclusion criteria:

Inclusion criteria: Patients who underwent primary unilateral hip arthroplasty, regardless of gender; with American Society of Anesthesiologists (ASA) classification of II-III and, with complete clinical data and no hospital referral.

Exclusion criteria: With morbid obesity, i.e., Body Mass Index (BMI)> 35 kg/m^2 ; with hypersensitivity to study-related drugs or severe coagulation disorders; with chronic pain, long-term use of analgesics or other psychotropic drugs and with puncture site infection.

Methods:

Preoperative preparation: All patients fasted for (6-8) h before surgery and no preoperative medication was administered. After entering the operating room, peripheral venous access was established and the Electrocardiogram (ECG), invasive arterial blood pressure and pulse oximetry of the patients were monitored using an ultrasound machine (diagnostic ultrasound system, model Wisonic Navis, Shenzhen Wisonic Medical Technology Co). The depth of anesthesia was controlled with an Electroencephalography (EEG) Bispectral Index (BIS) monitor. Mask oxygenation and invasive arterial monitoring were performed under local anesthesia. All patients received intravenous inhalation compound general anesthesia.

Nerve block: In the study group, nerve blocks were performed under ultrasound guidance after hip arthroplasty. With the patients in a lateral position,

after routine disinfection, the nerve block was performed using a short 20 G beveled needle (10 cm, Braun, Germany) under the guidance of the Terason T3000 portable ultrasound device (Terason, United States of America (USA)). The ultrasound probe was first placed along the mid-axillary line between the iliac crest and the edge of the costal arch, and then slowly moved dorsally. Under the short-axis plane method, the puncture needle was inserted between the lumbar square and lumbar major muscles (at the level of lumbar 4/5), 1-2 ml of saline was injected to determine the tip position and then 30 ml of ropivacaine at a concentration of 0.375 % was injected^[10,11].

Patients in the observation group were placed in the supine position with the high-frequency linear transducer placed parallel to the inguinal region. The femoral artery was first located and then moved laterally to locate the femoral nerve, broad fascial tensor, suture muscle, iliac spine and iliac fascia. 30 ml of ropivacaine at a concentration of 0.375 % was injected into the iliac fascia gap. 13 min after the block, the effect of the block was assessed by one investigator using pinprick sensation in each cortical distribution of the closed foraminal nerve, lateral femoral cutaneous nerve and femoral nerve.

Anesthesia method: With the patient in the supine position, a subarachnoid puncture was performed at L4 and L5 after sterilization. After the presence of cerebrospinal fluid reflux, 2 ml of bupivacaine at a concentration of 5 % was injected. Midazolam and sufentanil were used intraoperative as adjunctive analgesic sedatives and 4.5 mg of tropisetron was used to prevent nausea and vomiting. Patientcontrolled analgesia was administered in all patients (150 μ g of sufentanil, 8.96 mg of tropisetron and saline supplementation to 300 ml, background dose of 4 ml/h, a self-controlled dose of 4 ml, lock time of 20 min, maximum dose of 20 ml/h).

Observation indicators:

Postoperative analgesia included oral non-steroidal anti-inflammatory drugs, patient-controlled intravenous analgesia and rescue analgesia. Self-controlled intravenous analgesia consisted of 100 μ g sufentanil and 8 mg of tropisetron, diluted to 100 ml with saline, delivered in 2 ml per dose, with a lock time of 15 min and background-free infusion.

The doses of sufentanil at 24 h and 48 h postoperatively were compared between the two

groups of patients. Pain at rest and during exercise was assessed at different time points (6 h, 12 h, 24 h and 48 h) after surgery in both groups using a Visual Analogue Scale (VAS) (total score of 10 points; the score is proportional to pain). Maximum hip flexion and abduction mobility were measured in all patients at 24 h and 48 h postoperatively. The occurrence of nausea and vomiting was recorded for all patients.

Statistical data:

GraphPad Prism 8 was used for image processing and Statistical Package for the Social Sciences (SPSS) 26.0 software was used to organize and statistically analyze the data. The measurement data were expressed by $(\bar{x}\pm s)$ and a t-test was used for analysis. Count data were expressed as rate (%) and analyzed using the Chi-square (χ^2) test. p<0.05 indicates that the difference is statistically significant.

RESULTS AND DISCUSSION

The observation group had 62 male and 38 female cases, aged 55-78 y (65.84 ± 6.37) y, with a height of 163.83 ± 7.14 cm, weight of 69.38 ± 15.98 kg, BMI of 23.21 ± 2.78 kg/m², ASA classification grade II in 44 cases and grade III in 56 cases, operative time of 80.98 ± 19.78 min, preoperative resting VAS score of 4.12 ± 0.58 and preoperative exercise VAS score of 8.02 ± 0.87 .

The study group had 65 males and 35 females, aged 58-77 y (65.53 ± 6.48) y, with a height of 163.25 ± 7.36 cm, weight of 69.11 ± 16.08 kg, BMI of 23.04 ± 2.93 kg/m², ASA classification grade II in 41 cases and grade III in 59 cases, operative time of 81.23 ± 19.65 min, preoperative resting VAS score of 4.21 ± 0.47 and preoperative exercise VAS score of 8.04 ± 0.79 . The patient characteristics between the two groups were comparable (p>0.05) as shown in Table 1.

Ultrasound-guided QLB resulted in significantly less sufertanil use in patients at 24 h and 48 h postoperatively (52.14 ± 5.11 , 105.74 ± 8.14) vs. fascia iliaca block (77.58 ± 7.93 , 150.18 ± 10.58) (p<0.05) as shown in Table 2.

The patients in the observation group had resting VAS scores of (1.51 ± 0.21) at 6 h postoperatively, (1.80 ± 0.35) at 12 h postoperatively, (2.61 ± 0.48) at 24 h postoperatively and (2.42 ± 0.45) at 48 h postoperatively. The resting VAS scores of patients in the study group were (1.53 ± 0.22) at 6 h postoperatively, (1.78 ± 0.41) at 12 h postoperatively, (2.11 ± 0.23) at 24 h postoperatively and (1.91 ± 0.33)

at 48 h postoperatively. The difference in pain at rest between the two groups at 6 h and 12 h postoperatively was not significant (p>0.05) and the VAS scores at rest in the study group were lower than those in the observation group at 24 h and 48 h postoperatively (p<0.05) as shown in fig. 1.

The motor VAS scores of patients in the observation group were (3.42 ± 1.02) at 6 h postoperatively, (8.18 ± 2.01) at 12 h postoperatively, (8.89 ± 2.23) at 24 h postoperatively and (6.87 ± 1.94) at 48 h postoperatively. Patients in the study group had (3.39 ± 1.11) at 6 h postoperatively, (4.35 ± 1.25) at 12 h postoperatively, (4.99 ± 1.37) at 24 h postoperatively and (3.62 ± 1.01) at 48 h postoperatively. The difference in pain during exercise was not significant between the two groups at 6 h postoperatively (p>0.05) and the VAS scores during exercise were lower in the study group than in the observation group at 12 h, 24 h and 48 h postoperatively (p<0.05) as shown in fig. 2.

The study group had significantly greater maximum hip flexion and abduction mobility at 24 h and 48 h postoperatively (68.88 ± 10.45 , 77.15 ± 12.95 , 25.98 ± 4.45 and 28.47 ± 5.89) than the observation group (55.41 ± 10.84 , 64.54 ± 12.45 , 19.65 ± 3.44 and 22.41 ± 4.41) (p<0.05) as shown in Table 3. Ultrasound-guided QLB was associated with a lower incidence of nausea and vomiting (7 %) vs. fascia iliaca block (39 %) (p<0.05) as shown in Table 4.

The anterior segment of the hip capsule is innervated by the femoral and foraminal nerves, and the posterior segment is innervated by the femoral square muscle nerve and occasionally by the superior gluteal nerve (poster-lateral region) and sciatic nerve (poster superior region). Intravenous sufentanil analgesia is a common clinical analgesic method for hip replacement surgery, but it is predisposed to various adverse effects such as nausea and vomiting, and respiratory depression. It was found that regional block anesthesia is effective in analgesia with fewer adverse effects^[12]. One of the regional anesthetic options for hip surgery is iliac fascia compartment block, which acts on the nerve-femoral nerve, closed foraminal nerve and lateral femoral cutaneous nerve. QLB is a posterior abdominal wall block performed entirely under ultrasound guidance^[13]. Recent findings suggest that the branches of the femoral nerve innervating the hip are located deep and that the lateral femoral cutaneous nerve has significant anatomical variability in its location under the inguinal ligament^[14]. Therefore, traditional analgesic modalities are considered insufficient to meet the analgesic needs of patients^[15,16].

Conventional analgesia is usually injected at the level of lumbar 2-4, with the blocking range covering chest 4-lumbar 1. In the present study, the injection location was shifted to the level of lumbar 4/5 (low QLB). Results of prior studies showed that the blocking range of this approach could reach the level of thorax 8-waist 2/3, theoretically fulfilling the analgesic needs of hip surgery. Moreover, QLB shows no implications for postoperative hip movement. Experimental studies have shown that ultrasoundguided QLB could significantly reduce postoperative sufentanil dosage, decrease pain scores and improve the quality of early recovery without increasing complications.

In the present study, ultrasound-guided QLB resulted in significantly less sufentanil use in patients at 24 h and 48 h postoperatively *vs.* fascia iliaca block; patients in the study group showed lower VAS scores both at rest and at exercise than those in the observation group at 12, 24 and 48 h postoperatively. QLB extends the anesthetic effects through the thoracolumbar fascia to the paravertebral space, effectively blocking partial sympathetic nerve, promoting the establishment of collateral circulation, increasing blood supply and reducing the level of inflammatory mediators, thus relieving pain^[17,18]. The results of this study showed that the study group had significantly greater maximum hip flexion and abduction mobility at 24 h and 48 h postoperatively than the observation group. Ultrasound-guided QLB was associated with a lower incidence of nausea and vomiting vs. fascia iliaca block, suggesting that patients in the study group had better pain relief, lower opioid requirements and enhanced quality of recovery. The lower drug requirement was attributable to the potentiated analgesic effect of QLB, which provides pain relief in the incisional area for patients undergoing total hip arthroplasty primarily by blocking the thoracic 10-lumbar 3 nerve region and skin tissue^[19]. The QLB provides pain relief in the incision area for patients undergoing total hip arthroplasty primarily by blocking the thoracic 10-lumbar 3 nerve region and skin tissue. In addition, all anesthetic blocks were performed before induction of anesthesia, which facilitates the correct deposition of local anesthetic and thus increases the success rate of the block. Besides, the risk of unrecognized vascular bleeding and nerve injury is reduced due to the shallow injection site of the needle^[20]. Kadane *et al.* found that QLB reduced pain scores and the need for analgesic medication at 24 h postoperatively, which is consistent with the results of the current study.

		Observation group (n=100)	Study group (n=100)	t	р
Gender	Male	62 (62.00)	65 (65.00)		
	Women	38 (38.00)	35 (35.00)		
Age (years)		55-78 (65.84±6.37)	58-77 (65.53±6.48)	0.341	0.733
Height (cm)		163.83±7.14	163.25±7.36	0.566	0.572
Body weight (kg)		69.38±15.98	69.11±16.08	0.119	0.905
BMI (kg/m²)		23.21±2.78	23.04±2.93	0.421	0.674
ASA grade II		44 (44.00)	41 (41.00)		
Grading grade III		56 (56.00)	59 (59.00)		
Surgery time (min)		80.98±19.78	81.23±19.65	0.090	0.928
Preoperative resting VAS score		4.12±0.58	4.21±0.47	1.206	0.229
Preoperative exercise VAS score		8.02±0.87	8.04±0.79	0.17	0.865

TABLE 1: PATIENT CHARACTERISTICS (x±s)

TABLE 2: SUFENTANIL DOSAGE (x±s)

Group	n	24 h (µg)	48 h (µg)
Observation	100	77.58±7.93	150.18±10.58
Study	100	52.14±5.11	105.74±8.14
t		26.967	33.291
р		<0.001	<0.001



Fig. 1: Resting VAS score Note: *p<0.05, (--): Observation group and (--): Research group



Fig. 2: Exercise VAS score Note: *p<0.05, (---): Observation group and (---): Research group

TABLE 3: POSTOPERATIVE HIP MOBILITY (x±s)

Group	n	Maximum hip flexion (°)		Outreach activity (°)	
		24 h after surgery	48 h after surgery	24 h after surgery	48 h postoperative
Observation	100	55.41±10.84	64.54±12.45	19.65±3.44	22.41±4.41
Study	100	68.88±10.45	77.15±12.95	25.98±4.45	28.47±5.89
t		8.946	7.02	11.254	8.236
р		<0.001	<0.001	<0.001	<0.001

TABLE 4: INCIDENCE OF NAUSEA AND VOMITING (%)

Group	n	Number of cases of nausea and vomiting	Incidence
Observation	100	39	39.00
Study	100	7	7.00 %
t			28.910
р			<0.001

The present study has the following limitations; due to the absence of objective indicators to quantify the effect of nerve block on muscle strength and the potential implications for motor function by severe postoperative pain, medically induced nerve injury and transient nerve palsy, subjective issues that compromise the reliability of the results of the current study exist and the postoperative decrease in motor function is not entirely ascribed to nerve block. In addition, the current study lacks parameters such as time lapses before first out-of-bed activity, length of stay and patient satisfaction, which will be further, investigated in future studies.

Ultrasound-guided QLB reduces the use of opioids in total hip replacement patients, significantly alleviates surgical pain, promotes early ultrasoundguided QLB reduces the use of opioids in total hip replacement patients, significantly alleviates surgical pain, promotes early functional recovery and lowers the incidence of adverse effects.

Conflict of interests:

The authors declared no conflict of interests.

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This article was originally published in a special issue, "Role of Biomedicine in Pharmaceutical Sciences" Indian J Pharm Sci 2023:85(2) Spl Issue "203-208"