

Ultrasound-guided fascia iliaca compartment block in orthopedic fractures: Bupivacaine 0.2% or 0.3%?

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Abstract

Background: Postoperative pain is one of the major problems in patients with femoral or hip fracture. Current study investigates the effect of 0.2% and 0.3% bupivacaine in ultrasound-guided fascia iliaca compartment block (FICB) on postoperative pain.

Methods: This randomized clinical trial study was conducted in Rasoul Akram Hospital of Iran University of Medical Sciences. Forty-eight patients with femoral or hip fractures who were candidates for surgery underwent FICB, divided into two groups receiving bupivacaine in 0.2% and 0.3% concentrations. Pain was evaluated with Visual Analog Scale (VAS) at times 2, 6, 12, 24, and 48 hours after surgery. Need for opioids, nausea and vomiting after surgery, patients' satisfaction with pain control and motor block were also recorded. T-test or Mann-Whitney U test and Repeated measure ANOVA was used for analysis.

Results: Pain score after surgery was significantly lower in groups receiving 0.3% than the group receiving 0.2% concentration. Patients' satisfaction with pain control in 0.2% group was significantly higher ($p=0.04$). Time to analgesic onset in both groups had no significant differences ($p=0.5$). The incidence of nausea and vomiting ($p=0.5$) and opioid consumption ($p=0.1$) between the two groups showed no significant difference.

Conclusion: In femoral or hip fracture, bupivacaine with 0.3% concentration in fascia iliaca compartment block can cause lower pain score compared to bupivacaine with 0.2% concentration, but patients' satisfaction with pain control and severity of motor block is higher in bupivacaine 0.2%.

Keywords: Postoperative pain, Bupivacaine, Fascia iliac block.

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Introduction

In fractures of the femur and pelvic bones, reduced postoperative pain and patient comfort are concerned (1,2). The consequences of post-operative pain include physiological stress which may lead to an increase in the hemostatic system, deep vein thrombosis resulting from venous stasis and impaired immune system which result in increased infections, fatigue and delay in return of muscle function (3).

Various postoperative pain control methods include intravenous and oral anti-inflammatory drugs, opioids, epidural and peripheral nerve block. Each of mentioned

methods has some advantages and disadvantages. The use of opioids is accompanied by side effects such as nausea, vomiting, constipation, itching, dizziness and respiratory depression. Epidural anesthesia is along with the risks such as epidural hematoma (4).

Attempts for creating effective and secure anesthesia have led to design of various methods for local anesthesia (5). Peripheral nerve block is highly considered due to its effective role in reducing postoperative pain, faster release, and movement of patients' movement (6- 9).

The previous limitation in the application

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of blocks was their limited efficacy which has been eliminated today with the presence of catheters and pumps, though it brings about such risks as catheter infection and toxicity of anesthetic drugs and prolonged block (10-12).

One of the peripheral nerve block methods, which is currently considered in orthopedic surgery is Fascia Iliaca Compartment Block (FICB). Some studies have investigated its efficacy and security on pain reduction in patients (13-17).

Meanwhile, bupivacaine is an amino acid anesthetic local drug which its efficacy has been emphasized in various studies considering its low price and availability (18). Since the volume and concentration of the local anesthetic drug, factors affecting the absorption and tissue distribution of the drug and injection site influence its efficacy (19), efficacy of bupivacaine in different concentrations has been studied as an adjuvant in multimodal analgesia approach in controlling postoperative pain (3,20).

Some studies had shown that the use of 0.5% bupivacaine or 0.25% for peripheral nerve blocks provides longer duration of postoperative analgesia for lower extremity surgery than the other long-acting local anesthetics for example ropivacaine or mepivacaine. Also, bupivacaine may exhibit a longer delay of onset of surgical anesthesia. Despite this, there is concern about the use of higher concentrations of bupivacaine because of the potential for systemic toxicity with larger doses, and there is the potential for prolonged nerve block (20,21).

According to our knowledge, no randomized double-blind trial study has been investigated the effect of bupivacaine in different concentrations in FICB method. Thus, we investigated the effect of bupivacaine in FICB in patients with femoral and hip fractures. Two difference concentrations of bupivacaine were used for achieving maximum pain control so that optimal level of drug is determined which is free from consequences and prolonged motor block in FICB.

Methods

This study was a Double-Blind Randomized Clinical Trial. Research population included 18–70 years old people with a femoral or hip fracture which underwent elective surgery. For calculation of sample size, according to the results of the pilot study (10 cases), difference and SD of pain with Visual Analog Scale (VAS) in the prescription of bupivacaine in doses 0.2% and 0.3% was 1.5. Thus, considering alpha error as 0.05 and power as 90% for this study, and given 10 percent probability of drop out, the sample size was calculated as 24 for each group.

Samples were selected using convenient sampling among patients referred to Rasoul Akram Hospital in Tehran. Informed written consent was taken from the patients. The study was done from March 2014 to April 2015.

Inclusion criteria were: 1) Patients with femoral or hip fracture at age range 18 to 70 years old 2) Patients with American Society of Anesthesiologist (ASA) 1 and 2 3) Patients with verbal and written communication abilities who could consciously provide the consent for participation in the study

Exclusion criteria were: 1) Multiple fractures in different parts simultaneously (multiple trauma) 2) Sensitivity to amid anesthetic drugs 3) Addiction to alcohol, drugs or cigarettes 4) Body Mass Index (BMI) < 18 and morbid obesity (BMI> 40) 5) Pregnancy 6) Peripheral neuropathy and mental disorders that are potentially effective on pain perception

Patients were randomly assigned to two groups (n=24) based on computer randomized list. One group received 0.2% bupivacaine (Milan, Italy, Markaeen) and the other group received 0.3% bupivacaine. For making study as double-blind, the patient and anesthesiologist who injected drugs were not aware of the prescribed medication.

Patients underwent standard monitoring including ECG, pulse oximetry and noninvasive blood pressure in operation room.

All patients experienced general anesthesia in a similar way. It included reception of midazolam 0.02mg/kg and fentanyl 2mcg/kg as premedication. For induction of anesthesia, thiopental 5mg/kg and atracurium 0.5mg/kg were used. For maintenance of anesthesia, propofol 100mcg/kg/min was used. After the end of the operation and transfer of the patient to the recovery room and necessary monitoring for patients (EKG- NIBP- POM), ultrasound-guided fascia iliaca compartment block was applied on patients using a linear probe. For performing the block, the patient was placed in Supine position. Ultrasound probe was placed in the inguinal region, and following observation of femoral arteries, the probe was moved a little toward the lateral so that iliopsoas muscle is specified as a hypo echo part in lateral to the artery and femoral nerve. After dermal anesthesia with 2 cc lidocaine 1% at the place of needle entry using spinal needle No. 23 as in line, 30 cc of local anesthetic was injected between Fascia Iliaca layer and Iliopsoas muscle after passing Fascia Iliaca layer. Distribution of local anesthetic as inward and outward of injection location and beneath Fascia location is proper. In recovery, pain severity was evaluated using VAS scale every 5 minutes (22) and pain severity below 3 was considered as the initiation of onset of analgesia. In addition, pain severity was measured at rest state at hours 2, 6, 12, 24, and 48 hours after operation based on VAS tool. In the case of non-pain control, pethidine 20 mg was injected intravenously in conditions of pain scores above 4. Also, sedation score was recorded using Ramsay Sedation Scale during 48 h follow-up (23). The frequency of nausea and vomiting was evaluated and recorded. In the case of nausea and vomiting in recovery, metoclopramide 10 mg was used. Satisfaction with the pain control and motor block severity (knee flexion and ability to lift leg), after 48 hours was recorded. Patient satisfaction level from pain control after 48 hours were measured as excellent, good, moderate and poor.

Data collection was done using a questionnaire. Information was collected through an author-made questionnaire and was carried out by one of the main authors.

Collected data were entered into SPSS software. Frequency distribution was used for qualitative variables and mean and Standard Deviation (SD) was used for quantitative variables. Analysis of qualitative data was done using chi-square test. Quantitative data analysis was done using t-test or Mann-Whitney U test. Repeated measure ANOVA was used for comparing pain severity at recorded times in two groups. In statistical analysis, P-value less than 0.05 was considered as significant. In this study, ethical principles of the Helsinki were pursued. The study was approved by Ethical Committee of Iran University of Medical Sciences, and it is registered at IRCT Center coded as IRCT5N2014081014199.

Results

Fifty-six patients were investigated; one of whom was excluded due to an addiction history. Two patients were excluded due to BMI above 40 and 3 patients were excluded due to age above 70 years, and 2 patients did not consent to participate in the study. Finally, information for 48 patients in two groups (n=24) was analyzed (Fig. 1).

18 patients (37.5%) were female and 30 patients (62.5%) were male. Mean±SD age of the patients was 43.4±17.4 years. Demographic information of patients, blood pressure and heart rate before the blocks and type of fractures are given in Table 1 in terms of two groups. Statistically, significant difference was not observed between the two groups and it may suggest the right patient randomization in two groups (p<0.05).

Overall pain score after the operation was lower in the group receiving 0.3 doses, and this difference was statistically significant (p<0.001) (Fig. 2).

Pain score of patients after 2, 6, 12, 24, and 48 hours of the operation are given in Table 2 for two groups. After 24 hours, no

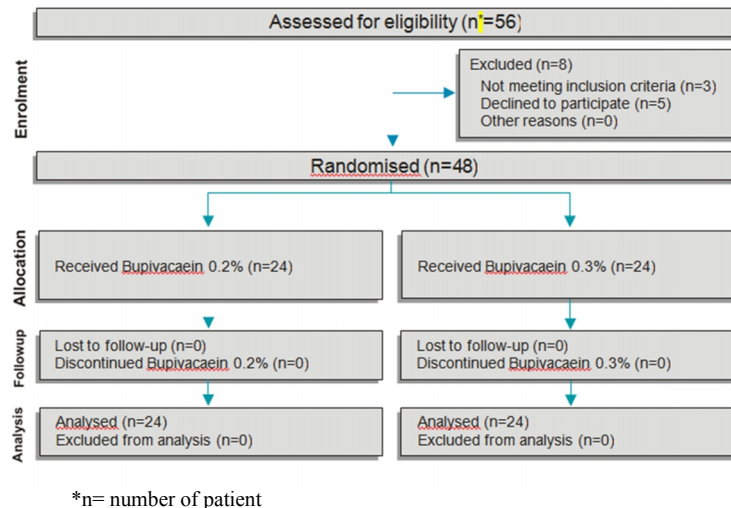


Fig. 1. Flow diagram of patients in the trial

Table 1. Demographic data type of fracture in two groups of patients

Variable	Bupivacaine 0.2% (n = 24)	Bupivacaine 0.3% (n = 24)	p
Age (year)	40.7±17.1*	45.1±15.7	0.4
Gender (female/male)	8.16	10.14	0.5
Body Mass Index (Kg/m2)	23.5±2.3	23.5±2.8	0.99
Type of fracture (Femur / Hip)	22.2	22.2	0.99

*Data presented as mean ± SD

significant different pain score was observed in two groups.

As observed in Table 3, time to analgesic onset was not significantly different between the two groups ($p=0.5$). The occurrence of nausea and vomiting was higher in the group receiving 0.3 doses, but there was no statistically significant difference ($p=0.5$). The number of patients requiring opioid consumption (analgesic) ($p=0.1$) and opioid dose ($p=0.6$) was lower in the group receiving 0.3 doses, but with

no statistically significant difference. The number of patients with high satisfaction (excellent score) of postoperative pain control and motor block severity (knee flexion and ability to lift leg) was significantly higher in the group receiving 0.2 doses ($p=0.04$) (Table 3).

Discussion

Local anesthesia techniques are preferred for pain control in orthopedic operations (4). FICB is mainly used for analgesia in hip and femoral surgery (13). Bupivacaine is a long-acting local anesthetic which is used as an adjuvant in local anesthesia techniques (24). Peripheral nerve block techniques can reduce postoperative pain and drug use, though they may cause long-term numbness and weakness and thus delays in the ability to movement in the affected patient and the patient's dissatisfaction (4). Hence, it is necessary that mentioned anesthesia techniques are investigated in terms of advantages and disadvantages by performing more accurate studies. Several studies have demonstrated the effectiveness of FICB method in pain

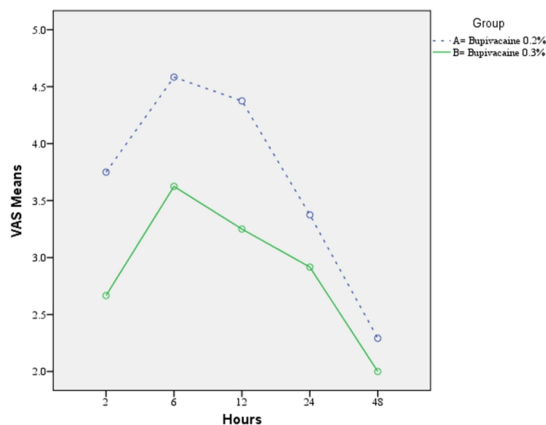


Fig. 2. Comparison of pain score changes in different times in two groups (VAS = Visual Analog Scale (0-10))

Table 2. Pain score in term of times 2, 6, 12, 24 and 48 hours after operation

Time	Mean VAS (0-10)		p
	Bupivacaine 0.2% (n = 24)	Bupivacaine 0.3% (n = 24)	
2	3.7±0.9*	2.7±0.9	< 0.001
6	4.6±1.4	3.6±1.1	0.01
12	4.4±1.3	1.3±0.9	0.002
24	3.4±2.9	1.5±0.8	0.02
48	2.3±1	2±0.4	0.04

*Data presented as mean ± SD

Table 3. Incidence of nausea-vomiting, opioid consumption, patient's satisfaction and time to analgesic onset

Variable	Bupivacaine 0.2% (n=24)	Bupivacaine 0.3% (n=24)	p
Number of patient with nausea-vomiting	8 (31.3%)*	10 (41.7%)*	0.5
Number of patients requiring opioid consumption	19 (79.2%)*	14 (58.3%)*	0.1
Consumption opioid dose (mg)	24.3±11.6**	22.1±6.3**	0.6
Number of patients with high satisfaction	15 (62.5%)*	8 (33.3%)*	0.04*
Time to analgesic onset (minute)	20.2±6.3**	18.9±5.4**	0.5

*N(%), ** Mean±SD

control in patients with orthopedic operations (13–16). However, the efficacy of bupivacaine with different concentrations, as an adjunct in local anesthesia techniques has been studied in other peripheral nerve blocks.

In the current study, effect of 0.2% and 0.3% bupivacaine in FICB on postoperative pain in femoral or hip fractures was studied. Current study indicates though pain score of patients with 0.3% bupivacaine was lower than 0.2% ($p < 0.001$), satisfaction patients with pain control and motor block severity was significantly lower in the group receiving 0.3% bupivacaine ($p = 0.04$). Although block anesthesia time was not directly measured in the current work, it seems lower satisfaction of patients was due to lengthened motor block in higher doses of bupivacaine and occurrence of numbness, weakness and delay in the walking (4).

Although some previous studies have compared bupivacaine with the other long-acting local anesthetics, few studies compared two difference concentrations of bupivacaine as an adjuvant in multimodal analgesia approach in controlling postoperative pain (20).

Mulroy studied the effect of femoral nerve block along with 0.25 and 0.5% bupivacaine on the duration of anesthesia time in patients undergoing an arthroscopic operation to repair the anterior cruciate liga-

ment of the knee. Findings showed bupivacaine in 0.25% and 0.5% concentrations is accompanied by acceptable anesthesia, and duration of anesthesia showed no difference with two different concentrations of bupivacaine (20).

Xie investigated the effect of peripheral nerve blocks (femoral, obturator and lateral femoral transcutaneous) with two different concentrations of bupivacaine in pain after total knee arthroplasty. Research findings suggested that patients receiving 0.25% bupivacaine had lower pain and higher satisfaction compared to the group receiving 0.5% bupivacaine (4).

In the study by Krych, use of multi-modal analgesia with fascia iliaca block fascia in acute pain control in patients undergoing hip arthroscopy was investigated. This study indicated that multi-modal analgesia with FICB in patients undergoing hip arthroscopy is secure and effective. The quality of initial analgesia after the operation is great by FICB leading to the lower use of opioid, high pain control quality and high patient satisfaction (14).

In another by Fujihara, FICB effect on pain control in patients with femoral fracture was compared with NSAID prescription after the operation. Finally, the study concluded that FICB is secure and effecting in developing analgesia and it is suggested for old patients with femoral fracture (15).

Pain score in the current study showed a

reduction in both groups after 24 hours, but there was no significant difference between two groups. In a study by Xie, femoral nerve block with bupivacaine showed a significant effect on early pain reduction after the operation, but it showed lower effect on pain after one day (4). In the study by de Lima E Souza, femoral nerve block using 0.25% ropivacaine or 0.5% bupivacaine was effective for postoperative analgesia after total knee replacement or anterior cruciate ligament reconstruction, particularly for the first 10 hours after spinal anesthesia (25).

Findings in the current study are similar to findings by Xie (4) indicating no significant difference in opioid consumption in both high and low dose of bupivacaine.

There was no significant difference in terms of occurrence of nausea and vomiting in both groups in this work which is consistent with findings by Xie (4).

Limitations of this study include lack of investigation of pain in patients in different motor states, lower numbers of patients with hip fracture and lack of measurement of analgesia duration of the block.

It is recommended that effect of peripheral nerve block procedures such as FICB with auxiliary local anesthetic drugs on the duration of anesthesia and pain in various motor states and other long-term consequences of the femoral and hip operations are studied in the future works.

Conclusion

In patients with femoral or hip fracture, bupivacaine with 0.3% concentration in fascia iliaca block can cause lower pain score than bupivacaine with 0.2% concentration after the operation, but the satisfaction of patients with pain control and motor block severity is higher in 0.2% concentration.

Thus, use of 0.2% concentration in fascia iliaca block is recommended for postoperative pain control in patients with femoral or hip fracture.

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Conflict of interest

The authors have no conflicts of interest to declare.

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