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Comparison of Ultrasound-Guided Femoral and Infrapatellar Nerve Block Effects after Anterior Cruciate Ligament Repair Surgery.

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Abstract

Despite the minimally-invasive nature of knee arthroscopic surgery, compared to traditional surgical methods, postoperative pain in these patients may be significantly high. These patients sometimes require high doses of opioids. Furthermore, after the utilization of such medications, there will be a possibility of some complications, such as respiratory depression, nausea, and vomiting. Peripheral nerve blocks have shown high efficiency and can decrease the requirement for opioids administration. It is worth mentioning that adequate postoperative pain relief is a significant factor in the early ambulation and rehabilitation of patients after knee surgery.

Keywords: ultrasound; femoral and infrapatellar nerve block; anterior cruciate; ligament repair; surgery

1. Background

Despite the minimally-invasive nature of knee arthroscopic surgery, compared to traditional surgical methods, postoperative pain in these patients may be significantly high. These patients sometimes require high doses of opioids (1, 2). Furthermore, after the utilization of such medications, there will be a possibility of some complications, such as respiratory depression, nausea, and vomiting. Peripheral nerve blocks have shown high efficiency and can decrease the requirement for opioids administration (3-6). It is worth mentioning that adequate postoperative pain relief is a significant factor in the early ambulation and rehabilitation of patients after knee surgery (7). Lower extremity innervation has arisen from the lumbar and lumbosacral plexus. The lumbar plexus is normally formed by L1-L4 roots and occasionally by branches of T12 or L5 (8,9). The femoral nerve is composed of L2, L3, and L4 branches, which are the largest and most significant branches of the lumbar plexus (9). The femoral nerve block (FNB) is an easy technique with low side effects. This method has been followed for anterior leg surgeries and pain management after pelvic and knee surgeries (10). The anterior cruciate ligament (ACL) reconst- ruction surgery can cause the weakness of the quadriceps muscle mainly due to reflex inhibition which is secondary to

joint effusion, or the pain that prevents the easy movement of the knee joint (11,12). The preservation of the strength of quadriceps muscles is the aim of post-operative rehabilitation and a significant factor that decides about the return of the affected person to daily activities and professional performance after ACL reconstruction surgery (12-14). Despite the high efficiency of FNB in decreasing the pain intensity after the knee surgery, it has led to some complications, such as quadriceps weakness (15-23). Multiple methods have been surveyed for pain control after ACL reconstruction, such as multimodal analgesia, intra-articular injection of morphine, and sciatic nerve block (24-29). However, FNB is still the method that is mostly used due to its simplicity and effectiveness.

2. Objectives

This study aimed to compare the femoral and infrapatellar nerve block effects in terms of pain score, quadriceps force, and satisfaction after the performance of these two techniques. It should be mentioned that the infrapatellar nerve is a sensory nerve and one of the terminal branching of the femoral nerve.

3. Methods

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This single-blind clinical trial was performed on patients who were candidates for elective arthroscopy ACL surgery and referred to Rasoul Akram Hospital, Tehran, Iran, during 2019. The study protocol was approved by the Ethical Committee of Iran University, Tehran, Iran (IR.IUMS.FMD.REC. 1398.377) and registered in the Iranian Registry of Clinical Trials (IRCT20120814010599N22). The inclusion criteria were: 1) age range between 20 and 45 years, 2) physical status of I-II according to American Society of Anesthesiologists classification, and 3) patients' willingness. The primary outcome variable in this study is the pain score (classified from 0-10), and more than a 2- point decrease in this score is considered clinically significant. Moreover, the confidence and power levels in this study were 95% and 80%, respectively. Considering the sample attrition, the minimum number of patients in each group was estimated at 46 cases. On the other hand, the patients with coagulation disorders, infection at the injection site, neurological disorders, neuropathies at the site of the operation, sensitivity to local anesthetics, drug addiction, liver or kidney failure, and body mass index (BMI) ≥35 were excluded from the study. Pulse oximetry, electrocardiography process, and non-invasive blood pressure measurement of the patients were controlled upon entering the operating room, and they were then hydrated with Ringer's lactate solution. Subsequently, the spinal anesthesia at the L4-L5 level was conducted in the lateral position using a 25-gauge needle containing 3 cc bupivacaine 0.5%. Following the surgery with regression of at least three dermatomes from the spinal block, the patients were assigned to ultrasound-guided FNB or infrapatellar nerve block (IPNB) groups using a computer-generated random sequence. Afterward, the area was prepared for the FNB, and the linear ultrasound probe was located over the inguinal ligament in a shortaxis view. In the next stage, the femoral nerve was placed lateral to the femoral artery below the iliac fascia. A 5-7 cm acoustic needle was inserted in-plane from the outside of the artery and directed toward the femoral nerve, followed by the perineural injection using 15 cc of ropivacaine 0.2%. The patients' thighs in the IPNB group were placed at an external rotation. The linear ultrasound probe was positioned in a short-axis view in mid-thigh exactly medial to the sartorius muscle. In this position, the saphenous nerve is more superficial than the femoral artery. After the saphenous nerve was observed, it was moved more distally to determine the infrapatellar nerve being isolated (1 to 2 cm lower the saphenous nerve). Furthermore, an injection was made using a 5-7 cm sonovisible needle containing 15 cc of ropivacaine 0.2%. Patientcontrolled intravenous analgesic pump containing 10 cc of fentanyl in 100 cc normal saline was connected to continuous flow amount of 2 cc/h with a lockout interval of 15 min in both groups. The first drug request and the overall drug dose was calculated in 24 h. Patients were evaluated in recovery, as well as at 1, 3, 6, 12, and 24 h after surgery in terms of pain intensity at rest and knee flexion according to a visual analog scale (VAS), satisfaction level according to a 5-point Likert scale (strongly satisfied=5, strongly dissatisfied=1), and muscle force grading (no contraction=grade 0, normal=grade 5). The evaluation was conducted by one of the authors who was blind to the patient grouping.

3.1 Statistical Analysis

The data were analyzed using SPSS software (version 22) through the Kolmogorov-Smirnov test to determine the normality of quantitative variables. Furthermore, the quantitative and qualitative variables were assessed using the t-independent or Mann-Whitney-U test and chi-square or Fisher's exact test, respectively. A p-value less than 0.05 was considered statistically significant.

4. Results

This study included 92 patients who were referred to Rasoul Akram Hospital, Tehran, Iran, for elective ACL surgery. The patients were assigned to the IPNB (n=46) and FNB groups (n=46). Figure 1 illustrates the consort flow chart.

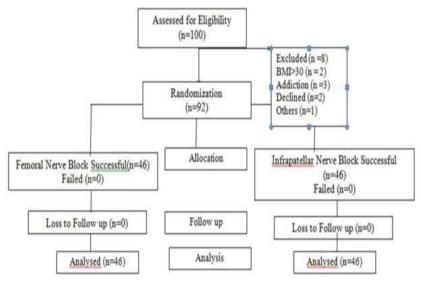


Figure 1. CONSORT flowchart.

According to the results, there was no significant difference between the two groups regarding age, gender, BMI, and duration of surgery (P>0.05) (Table 1). The mean VAS scores at rest, 1 and, 3 h was significantly lower in the FNB group, compared to those in the IPNB group (P<0.05) (Table 2).

Moreover, the evaluation of pain score at rest using repeated ANOVA showed no significant difference between groups regarding the next 24 h

after the NB (P=0.082). No significant difference was also observed between the two groups in terms of the mean VAS scores in flexion during recovery times, as well as 3, 6, 12, and 24 h after NB (P>0.05). Flexion pain scores 1 h after NB in the FNB group was significantly lower, compared to those in the IPNB group (P<0.05) (Table 3). Additionally, the evaluation of pain score trends during flexion using repeated ANOVA revealed a significant difference between the groups in terms of the next 24 h after NB (P=0.004) The mean satisfaction score one h after the block

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in the FNB group was significantly higher than that in the IPNB group (P<0.05). At other times, there was a significant difference between the two groups in this regard (P>0.05) (Table 4). Similarly, the results of repeated ANOVA regarding the evaluation of satisfaction score trends indicated a significant difference between the two groups in terms of the next 24 h after the NB (P=0.03). In addition, the mean muscle force of patients during the first day after the surgery in the FNB group was

significantly lower than that in the IPNB group (P<0.05) (Table 5). Moreover, the FNB group obtained a higher mean time required for the first dose of analgesic request, compared to the IPNB group (P<0.05). The mean dose of used opioids during the first and second 12 h in the FNB group was significantly lower than that in the IPNB group (P<0.05) (Table 6). In the second 12 h, the used opioid was significantly higher in both groups (P<0.05).

Table 1. Demographic characteristics of the two groups			
Variables	Groups		P-value
	Infrapatellar	Femoral nerve	1-value
Age(year) Mean±SD	34.15±6.42 ⁺	33.11±6.18	0.429
Gender Male Female	6 (13%) [#] 40 (87%)	10 (21.7%) 36 (78.6%)	0.271
Body mass index (kg/m²) Mean±SD	26.16±1.94	26.1±2.92	0.752
Duration of Surgery (hours) Mean±SD	123.26±14.46	126.84±14.99	0.246

+: Mean±SD, #: Frequency (%)

Table 2. Comparison of pain scores at rest bweteen the two groups			
Time Measurement	Groups		P-value
Time Weasurement	Infrapatellar	Femoral nerve	- 1-value
Recovery	3.47±0.58+	3.67±0.66	0.139
1 h after surgery	2.84± 0.41	2.41±0.58	0.001
3 h after surgery	2.5±0.5	2.17±0.38	0.001
6 h after surgery	2.45±0.5	2.47±0.54	0.843
12 h after surgery	2.89±0.64	2.95±0.51	0.592
24 h after surgery	3.11±0.67	2.95±0.55	0.241

+: Mean±SD

Table 3. Comparison of pain scores at flexion between the two groups			
Time Measurement	Groups		P-value
	Infrapatellar	Femoral nerve	1 -value
Recovery	5.74±0.77 ⁺	5.56±0.8	0.294
1 h after surgery	4.58± 0.65	4.17±0.64	0.003
3 h after surgery	4.08±0.55	3.97±0.49	0.322
6 h after surgery	4.28±0.58	4.33±0.66	0.74
12 h after surgery	4.84±0.94	4.56±0.65	0.098
24 h after surgery	4.78±1.03	4.37±0.85	0.069

+: Mean±SD

Table 4. Comparison of satisfaction scores between the two groups			
Time Measurement	Groups		P-value
	Infrapatellar	Femoral nerve	1-value
Recovery	2.28±0.54 ⁺	2.11±0.7	0.189
1 h after surgery	2.93 ± 0.71	3.54±0.72	0.001
3 h after surgery	3.78±0.47	3.89±0.31	0.194
6 h after surgery	3.91±0.28	3.95±0.2	0.404
12 h after surgery	3.63±0.53	3.71±0.45	0.402
24 h after surgery	3.21±0.51	3.32±0.52	0.315

Table 5. Comparison of muscle force score between the two groups		
	Group	

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Time Measurement	Infrapatellar	Femoral nerve	P-value
1 h after surgery	1.21±0.41+	1±0	0.001
3 h after surgery	2.56± 0.5	1±0	0.001
6 h after surgery	3.28±0.45	1.11±0.31	0.001
12 h after surgery	4.45± 0.54	2.34±0.52	0.001
24 h after surgery	5±0	2.91±0.28	0.001

Table 6. Comparison of the first analgesic request time and the prescribed dose of opioids during the first and second 12 h between the two groups.

Variables	Group		P-value
v arrables	Infrapatellar	Femoral nerve	1 value
Time required for the first analgesic request (hour)	3.39±0.49 ⁺	3.76±0.52	0.001
Used opioids during the first 12 h (μg)	97.22±9.97	61.52±10.48	0.429
Used opioids during the second 12 h (µg)	154.44±8.42	100.32±10.02	0.429

5. Discussion

Arthroscopic ACL reconstruction is one of the most common orthopedic procedures. Multiple protocols have been utilized for the pain relief following this surgery while trying to keep quadriceps muscle force. In this regard, the present study was conducted to compare the effects of FNB and IPNB in terms of reduction in pain severity, patient satisfaction, and maintenance of muscle force. It was indicated that FNB had superiority to IPNB in terms of reduction in pain score at rest, knee flexion (during early hours after the procedure), and patient satisfaction score. The femoral group required needed less analgesia after surgery; however, the use of opioids was higher in the second 12 h after the surgery, which may be due to the termination of block duration time. The IPNB had superiority to FNB in terms of the maintenance of the quadriceps force, which is in line with the findings of other studies. As previously noted, the utilization of the FNB leads to some complications, such as the delayed return of quadriceps muscle function (15-21). This complication is sometimes observed up to six months following the FNB (22, 23). The delayed return of muscle force in the long term can lead to patient dissatisfaction. Therefore, a balance should be maintained between the good analgesia outcome of the FNB and its complications after ACL surgery. According to a study conducted by Marit Lundblad et al., patients experienced less pain and improvements in sleep after ACL arthroscopy surgery with IPNB. It should be mentioned that this study did not compare the groups regarding muscle force (30). In the current study, the reduction of pain in the FNB group was significantly more noticeable during the early hours after the surgery; however, the IPNB group showed improvements in muscle force. In a study performed by Rahimzadeh et al., the analgesic impact of the FNB and the adductor canal block was compared after arthroscopic knee surgery. The results revealed a decrease in the mean VAS scores from 5.6 to 4 immediately after both blocks. The pain scores three hours after the block were reduced to 2 and 3.4 in the FNB and adductor block groups, respectively. This study indicated that patients in the FNB group required analgesics less than the other group; in addition, they showed a higher satisfaction rate. It is worth mentioning that this study mentioned no muscle force (31). In the same vein, Ahl concluded that the patients with adductor canal block had higher VAS scores and morphine consumption, compared to the FNB group. Nonetheless, the adductor group obtained less muscle weakness, which was in line with the results of the current study (32).

Magnussen et al. compared the impacts of FNB and placebo block; moreover, they revealed that the patients who underwent FNB obtained a

mean quadriceps femoris-limb symmetry index that was 13.4% lower than that in the control group. In addition, they had a poorer knee injury and osteoarthritis outcome scores six weeks after the surgery, compared to controls. This decline in quadriceps force was quite similar to that in the present study following FNB (33). Guirro et al. compared the patients who underwent spinal anesthesia and FNB with a group that only had spinal anesthesia. According to the results, pain control after surgery was more efficient in the first group, compared to the other group. However, there was no difference between the two groups in terms of tramadol requests for pain relief (34). Based on the results of the present study, it seems that FNB is associated with a greater pain relief and satisfaction score in patients undergoing arthroscopic ACL repair, compared to the IPNB technique. However, a further delay in the recovery of quadriceps muscle force is evident in the FNB technique which did not lead to dissatisfaction in our patients. Regarding the limitations of this study, one can name the limited follow-up time and the number of patients. Therefore, it is recommended that multi- centric studies be conducted with a larger sample size for a longer time.

6. Conclusion

The FNB is associated with greater pain relief and satisfaction in patients undergoing arthroscopic ACL reconstruction, compared to the IPNB technique; however, a further delay was observed in the return of quadriceps force.

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Footnotes

Authors' Contribution: Study concept and design: SHRF and PR; analysis and interpretation of data: PR and SSK; drafting of the manuscript SHRF, FI, and PR; critical revision of the manuscript for important intellectual content: SHRF, PR, and SSK; statistical analysis: SHRF, PR, KH, and AS

Conflict of Interests: The authors declare that they have no competing interests.

Ethical Approval: The study protocol was approved by the Ethics Committee of Iran University of Medical Sciences, Tehran, Iran (IR.IUMS.FMD.REC.1398.377) and registered in the Iranian Registry of Clinical Trials (IRCT20120814010599N22).

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Informed consent: Informed consent was obtained from the patients.

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