

## Comparative Study between Adductor Canal Block and Femoral Nerve Block for Postoperative Analgesia in Knee Arthroscopy

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### Abstract

**Background:** Femoral Nerve Block (FNB) is one of the easiest peripheral nerve blocks. However, prolonged motor blockade is associated with a clinically important risk of fall. With the advent of ultrasonography, the adductor canal can be easily visualized at the mid-thigh level, allowing performance of Adductor Canal Block (ACB) with a high success rate.

**Aim of the Study:** To compare the safety and efficacy of ultrasound guided adductor canal block versus ultrasound guided femoral nerve block as postoperative analgesic in patients undergoing knee arthroscopy.

**Patients and Methods:** This study was carried out in Tanta University Hospitals from September 2015 till March 2016 on 105 adult patients of both sexes with ASA physical status I/II scheduled for knee arthroscopy. Patients divided into three equal groups (Group I) received basic analgesia in the form of paracetamol and diclofenac, (Group II) received ultrasound guided FNB and (Group III) received ultrasound guided ACB.

**Results:** There were no significant differences among the three studied groups according to demographic data. Comparison of the mean value of NPS score showed no significant difference between FNB and ACB, but there were significant increase in control group in comparison to both FNB and ACB. The first time to introduce morphine and total morphine consumption showed no significant difference between FNB and ACB. There was significant decrease of BBS score in FNB till 6-8h post-operative in comparison with control group and ACB.

**Conclusion:** Ultrasound guided adductor canal block is efficient as ultrasound guided femoral nerve block in control post-operative pain in patients undergoing knee arthroscopy. Also ACB result in early mobilization with no risk of fall that renders ACB preferred.

**Key Words:** Post-operative analgesia – Adductor canal block – Femoral nerve block – Knee arthroscopy.

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### Introduction

**KNEE** arthroscopy is a very common procedure and often is performed as day-case surgery [1]. It has been reported that a significant number of patients have moderate to severe pain 24 hours after ambulatory surgery in general and knee arthroscopy in particular, and pain affects the patient's activity level and satisfaction [2]. In an effort to provide an effective, safe and long lasting post-arthroscopy analgesia, several studies using different drugs and regimes have been published during the last two decades [3,4]. Intra-articular administration of local anesthetics has been widely used but some studies have questioned their efficacy [5].

The Femoral Nerve Block (FNB) is one of the easiest peripheral nerve blocks to master because the landmarks are generally easy to identify and the nerve is usually found at a superficial depth. However, prolonged motor blockade from FNB is associated with a small but clinically important risk of fall [6,7].

With the advent of ultrasonography, the adductor canal can be easily visualized at the mid-thigh level, allowing performance of Adductor Canal Block (ACB) with a high success rate [8,9].

### Patients and Methods

After obtaining the Research Ethics Committee approval (approval code: (30342/06/15) an informed consent was taken from each patient, a prospective single blind randomized study was carried out in Tanta University Hospitals from September 2015 till March 2016 on 105 adult patients of either sex, American Society of An-

esthesiologists (ASA) class I & II scheduled for knee arthroscopy either therapeutic or diagnostic, excluding the patients who refused to share in the study, those with history of hypersensitivity to local anesthetics and those with coagulopathies.

We also planned to exclude patients who were complaining of local infection at the site of the block, preexisting neuropathy and/or femoral AV malformation on the operative limb.

The total participant number reached 105 patients, each one was randomly assigned to one of the three equal groups using opaque sealed envelope. This randomization was done by independent assistant that didn't share in the next steps of the study.

*Group I (control group): (No. 35):*

Patients received intraoperative analgesia an intravenous paracetamol 1gm which was repeated orally every 6h post-operatively and diclofnac 75mg infusion intraoperative then was repeated orally 25mg every 6h post-operatively.

*Group II (group FNB): (No. 35):*

Patients received Femoral Nerve Block (FNB) after induction of general anesthesia.

*Group III (group ACB): (No. 35):*

Patients received Adductor Canal Block (ACB) after induction of general anesthesia.

*Management:*

*In the anesthesia clinic:*

History taking include (age and sex), careful examination to exclude any neurological deficit in lower limb and laboratory investigation. Patient education about the use of 0-10 numerical pain scale (NPS: With endpoints of 0: 'no pain' and 10: 'Worst pain imaginable).

*Intraoperative:*

Basic monitoring [ECG, Non-Invasive Blood Pressure (NIBP), oxygen saturation (SPO<sub>2</sub>) using pulse oxymetry] was performed; an Intravenous (I.V) line was general anesthesia was induced using 1 µg/kg of fentanyl and 2.0mg/kg of propofol. Tracheal intubation was facilitated by 0.15mg/kg cisatracurium after mask ventilation for 5 minutes. Muscle relaxation was maintained with additional doses 0.03mg/kg. Anesthesia was maintained with isoflurane 1-2 MAC. Controlled mechanical ventilation to keep endtidal CO<sub>2</sub> between 34-37.

*Techniques of nerve block:*

*Equipment:*

- 22-gauge 100mm length, short-beveled regional block needle.
- Skin antiseptic solution (0.5% Chlorhexidine spray).
- Sterile gloves, towels and gel.
- For femoral nerve block 30mls of 0.25% Bupivacaine.
- For adductor canal block 15mls of 0.25% Bupivacaine.
- Portable ultrasound machine (Toshiba Viano U.S.A), a 6-12MHz linear type probe.

*Technique of FNB:*

- Position: Supine position.
- The skin around the femoral crease was disinfected.
- Transducer positioned to identify femoral artery and nerve.
- Once femoral nerve identified, the needle inserted in-plane in a lateral-to-medial orientation and advanced toward the femoral nerve. When the needle tip adjacent to the nerve, inject 30ml Bupivacaine 0.25%.

*Technique of ACB:*

- Position: Supine.
- Thigh abducted and externally rotated to allow access to the medial thigh.
- Skin over mid-thigh disinfected.
- The transducer placed anteromedially midway between inguinal crease and medial condyle to identify sartorius muscle.
- Probe positioned perpendicular to artery and using in-plane technique with needle directed from lateral to medial to deposit local anesthetic under sartorius and around the femoral artery. We injected 15ml Bupivacaine 0.25%.

*Post-operative:*

After recovery, all patients were transported to PACU for 2 hours then to the ward where observation was completed for 12 hours. Any patient with NPS  $\geq 4$  intravenous morphine was titrated every 5min in 3mg increments (2mg in patients weighing  $\leq 60$ kg), and pain was assessed every 5min until pain relief, defined as a NPS score less than 4.

Clinical monitoring included respiratory rate measurements, oxygen saturation measured by pulse oximetry, arterial blood pressure, and heart rate. Morphine titration was stopped if the patient had a respiratory rate lower than 12 breaths/min, had an oxygen saturation measured by pulse oximetry lower than 95%.

**Results**

Statistical presentation and analysis of the present study was conducted, using the mean, standard deviation and chi-square test by SPSS V.16.

The age in Group I range from (19-39) years with mean value  $26.9 \pm 5.65$ , in Group II range from (18-39) years with mean value  $28.5 \pm 5.6$  and Group III range from (20-40) years with mean value  $29 \pm 6.3$  with insignificant difference when compared to each other as shown in (Table 1).

According to the sex in Group I show 18 male (51.4%) and 17 female (48.6%), in Group II show 20 male (57.1%) and 15 female (42.9%) and Group III show 16 male (45.7%) and 19 female (54.3%) with insignificant difference among the three studied groups as shown in (Table 1).

ASA classes in Group I showed 23 patients ASA I (65.7%) and 12 patients ASA II (34.3%), in Group II 24 patients ASA I (68.6%) and 11 patients ASA II (31.4%) and in Group III 20 patients ASA I (57.1%) and 15 patients ASA II (42.9%) with insignificant difference among the three studied groups as shown in (Table 1).

BMI in Group I range from (18-36) min with mean value  $24.57 \pm 5.09$ , in Group II range from (17-37) min with mean value  $27.17 \pm 5.8$  and Group

III range from (16-36) min with mean value  $26.69 \pm 6.28$  which show insignificant difference when compared to each other as shown in (Table 1).

Comparison of the mean value of NPS among the three studied groups revealed that there was significant decrease in Group II & III in comparison with Group I at 30min, 1hr, 2hr, 3hr, 4hr, 5hr, 6hr, 8hr, 10hr and 12hr post-operatively as shown in (Table 2).

Comparison of the mean value of BBS (Berg Balance score) among the three studied groups revealed that there was no significant difference at the base line (preblock) but there was significant decrease in Group II in comparison with Group I & Group III at 30min after full recovery, 2hr, 4hr, 6hr and 8hr post-operatively as shown in (Table

Comparison among the three studied groups according to onset of 1st dose morphine revealed that there was significant increase in Group II & III in comparison with Group I as shown in (Table

The total dose of analgesic consumption (morphine in (mg)) showed significant increase in Group I in comparison with Group II & Group III ( $p$ -value  $< 0.001$ ) with insignificant change between Group II & Group III ( $p$ -value 0.943) as shown in (Table 5).

Duration of motor block (BBS score  $< 40$ ) in Group I & Group III was 0hr. While in Group II ranged between 2-8hr with mean value  $(4.68 \pm 1.3)$  hr. in comparison among the three groups there was significant increase in Group II in comparison with Group I & III as shown in (Table 6).

Table (1): Comparison among the three studied groups according to demographic data (age, sex, ASA and BMI).

Items	Group I (n=35)	Group II (n=35)	Group III (n=35)	Tests	
				f & $\chi^2$	p-value
<i>Sex:</i>					
Male	18 (51.4%)	20 (57.1%)	16 (45.7%)	0.915	0.633
Female	17 (48.6%)	15 (42.9%)	19 (54.3%)		
<i>Age (years):</i>					
Range	19-39	18-39	20-40	1.209	0.303
Mean $\pm$ SD	$26.91 \pm 5.65$	$28.51 \pm 5.61$	$29 \pm 6.32$		
<i>ASA:</i>					
ASA I	23 (65.7%)	24 (68.6%)	20 (57.1%)	1.072	0.585
ASA II	12 (34.3%)	11 (31.4%)	15 (42.9%)		
<i>BMI (Kg/m<sup>2</sup>):</i>					
Range	18-36	17-37	16-36	2.024	0.137
Mean $\pm$ SD	$24.57 \pm 5.09$	$27.17 \pm 5.81$	$26.69 \pm 6.28$		

Table (2): Comparison among the three studied groups according to NPS.

	NPS (post-operative)									
	30min	1hr	2hr	3hr	4hr	5hr	6hr	8hr	10hr	12hr
<i>GI:</i>										
Mean	4.31	4.23	4.26	3.89	3.77	4.06	4.51	4.34	4.0	4.46
SD	1.02	0.81	0.82	0.90	1.09	0.80	0.61	1.08	0.84	0.78
<i>GII:</i>										
Mean	1.13	1.28	1.28	1.31	1.34	1.38	1.47	1.50	1.53	1.38
SD	0.98	0.99	1.08	1.06	1.04	1.04	1.14	1.08	1.08	1.10
<i>GIII:</i>										
Mean	1.27	1.45	1.52	1.55	1.45	1.45	1.64	1.52	1.61	1.64
SD	0.91	0.90	0.91	0.87	0.94	0.90	0.90	0.83	0.97	0.93
H. test	65.950*	67.099*	66.513*	61.809*	56.820*	63.500*	68.375*	63.008*	58.485*	66.172*
<i>p</i> -value	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*
<i>p</i> <sub>1</sub>	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*
<i>p</i> <sub>2</sub>	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*
<i>p</i> <sub>3</sub>	0.492	0.409	0.241	0.224	0.757	0.804	0.463	1.000	0.571	0.273

*p*<sub>1</sub>: *p*-value for comparing between Group I & Group II.  
*p*<sub>2</sub>: *p*-value for comparing between Group I & Group III.

*p*<sub>3</sub>: *p*-value for comparing between Group II & Group III.  
 \*: Statistically significant at *p*≤0.05.

Table (3): Comparison among the three studied groups according to BBS.

	BBS						
	Pre	30m	2hr	4hr	6hr	8hr	12hr
<i>Control:</i>							
Mean	56.0	56.0	56.0	56.0	56.0	56.0	56.0
SD	0.0	0.0	0.0	0.0	0.0	0.0	0.0
<i>FNB:</i>							
Mean	56.0	25.53	34.94	43.09	51.25	55.06	56.0
SD	0.0	5.45	4.63	5.28	4.87	2.14	0.0
<i>ACB:</i>							
Mean	56.0	56.0	56.0	56.0	56.0	56.0	56.0
SD	0.0	0.0	0.0	0.0	0.0	0.0	0.0
F-test	—	1064.969*	705.235*	203.766*	32.353*	6.538*	—
<i>p</i> -value	—	<0.001*	<0.001*	<0.001*	<0.001*	0.002*	—
<i>p</i> <sub>1</sub>	—	<0.001*	<0.001*	<0.001*	<0.001*	0.002*	—
<i>p</i> <sub>2</sub>	—	1.000	1.000	1.000	1.000	1.000	—
<i>p</i> <sub>3</sub>	—	<0.001*	<0.001*	<0.001*	<0.001*	0.002*	—

*p*<sub>1</sub>: *p*-value for comparing between Group I & Group II.  
*p*<sub>2</sub>: *p*-value for comparing between Group I & Group III.  
*p*<sub>3</sub>: *p*-value for comparing between Group II & Group III.  
 \*: Statistically significant at *p*≤0.05.

Table (4): Time till need for first dose of morphine in hours.

	Group I	Group II	Group III
Min.	0.0	1.0	2.0
Max.	0.0	12.0	12.0
Mean	0.0	11.22	11.09
SD	0.0	2.28	2.57
H-test		85.455	
<i>p</i> -value		<0.001*	
<i>p</i> <sub>1</sub>		<0.001*	
<i>p</i> <sub>2</sub>		<0.001*	
<i>p</i> <sub>3</sub>		0.983	

*p*<sub>1</sub>: *p*-value for comparing between Group I & Group II.  
*p*<sub>2</sub>: *p*-value for comparing between Group I & Group III.  
*p*<sub>3</sub>: *p*-value for comparing between Group II & Group III.  
 \*: Statistically significant at *p*≤0.05.

Table (5): Comparison among the three studied groups according to total morphine consumption (in mg).

	Group I	Group II	Group III
Min.	6	0	0
Max.	25	20	20
Mean	14.11	1.37	2.29
SD	4.63	3.87	4.78
F-test		57.947	
<i>p</i>		<0.001	
<i>p</i> <sub>1</sub>		<0.001	
<i>p</i> <sub>2</sub>		<0.001	
<i>p</i> <sub>3</sub>		0.943	

*p*<sub>1</sub>: *p*-value for comparing between Group I & Group II.  
*p*<sub>2</sub>: *p*-value for comparing between Group I & Group III.  
*p*<sub>3</sub>: *p*-value for comparing between Group II & Group III.  
 \*: Statistically significant at *p*≤0.05.

Table (6): Comparison among the three studied groups according to duration of motor block (in hours).

	Group I	Group II	Group III
Min.	0.0	2.0	0.0
Max.	0.0	8.0	0.0
Mean	0.0	4.68	0.0
SD	0.0	1.3	0.0
F-test		438.56	
$p_1$		<0.001*	
$p_2$		1.00	
$p_3$		<0.001*	

$p_1$  :  $p$ -value for comparing between Group I & Group II.  
 $p_2$  :  $p$ -value for comparing between Group I & Group III.  
 $p_3$  :  $p$ -value for comparing between Group II & Group III.  
 \* : Statistically significant at  $p \leq 0.05$ .

### Discussion

Peripheral nerve blocks are associated with less pain and lower odds of unplanned hospital admission compared to systemic analgesia [10]. The decision regarding continuous versus single-injection depends on the expected surgical trauma and patient factors [11].

Benefits of ACB may include shorter hospital stays, earlier and more efficient rehabilitation, and pain control. Additionally, patients will retain the ability to report pain in neighboring distributions that can be involved when attempting to block the femoral nerve at the inguinal crease. This technique also embraces the emerging regional philosophy of selectivity or blocking only the area involved in the surgery [12].

The most important advantage of ultrasound for peripheral nerve block PNB is the ability to confirm local anesthetic spread around the target nerve. This is the difference from conventional blind techniques, which can fail because local anesthetic does not uniformly surround the target nerve [13].

In our study, there was no significant difference among the three studied groups as regard to demographic data (age, sex, ASA & BMI).

Comparison of the mean value of NPS score among the studied groups revealed that there was no significant difference between FNB and ACB ( $p$ -value 0.54), but there were significant increase in control group in comparison to both FNB and ACB ( $p$ -value <0.001).

The first time to introduce morphine and total morphine consumption showed no significant difference between FNB and ACB ( $p$ -value 0.983, 0.0943) respectively, but there were significant

increase in control group in comparison to both FNB and ACB ( $p$ -value <0.001).

While comparison of the mean value of BBS score among the studied groups revealed that there was no significant difference between control group and ACB, there was significant decrease of BBS score in FNB till 6-8h post-operatively in comparison with control group and ACB ( $p$ -value <0.001, <0.001) respectively, indicating high risk of falling.

In agreement with our results, Hanson et al., [14] stated that all patients received ACB were able to stand in the post anesthesia care unit and no study participant subjectively mentioned leg weakness or reported falls within post-operative 24 hours. However, they didn't objectively measure the quadriceps muscle strength.

Kwofie et al., [15] demonstrated that Quadriceps strength and balance scores were similar to baseline following ACB. Following FNB, there was a significant reduction in quadriceps strength.

David Kim et al., [16] estimated that the FNB would result in at least 50% decrease in motor strength in comparison with the saphenous nerve block.

Jaeger et al., [17] reported that FNB reduced 49% of quadriceps strength from baseline but ACB caused only 8% reduction in healthy young subjects.

In controversy to our results, Espelund et al., [18] concluded that there were no significant analgesic effect of the ACB after minor arthroscopic knee surgery with a basic analgesic regimen.

El-Ahl, [19] found that the VAS pain score and opioid consumption was significantly higher in patients received ACB than FNB.

### Conclusion and Recommendation:

Ultrasound guided ACB should be considered as a safe efficient alternative to ultrasound guided FNB for post-operative pain in cases of knee arthroscopy. Also ACB results in early mobilization with no risk of fall and that renders ACB preferred.

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## دراسة مقارنة بين تخدير القناة المقربة وتخدير العصب الفخذي في تسكين آلام ما بعد عمليات منظار الركبة

عادة ما تتسبب جراحات منظار الركبة في آلام شديدة فيما بعد العملية لمدة قد تصل إلى ٢٤ ساعة، لذا بعد تخدير الأعصاب الطرفية طريقة فعالة وأمنة وطويلة المدى كمسكن لتلك العمليات. وقد أدى استخدام الموجات فوق الصوتية إلى إرتفاع نجاح تخدير الأعصاب الطرفية.

وكان من شأن تخدير الأعصاب الطرفية أن توفر للمريض القدرة المبكرة على الحركة، بالإضافة إلى تسكين آلام ما بعد الجراحة.

ولما كان لتخدير العصب الفخذي تأثير فعال على قدرة المريض على الحركة فيما بعد العملية فقد إتجهت الأبحاث مؤخرا إلى استخدام تخدير القناة المقربة كتقنية بديلة لتسكين آلام ما بعد جراحات الركبة، حيث أنه يعطى نفس التأثير كمسكن مع الحفاظ على قدرة المريض على الحركة فيما بعد الجراحة.

وكان الهدف من البحث: دراسة مقارنة بين تخدير العصب الفخذي وتخدير القناة المقربة باستخدام الموجات فوق الصوتية للتحكم في آلام ما بعد عمليات منظار الركبة.

حيث تضمن البحث ١٠٥ من المرضى البالغين المصنفين من الفئة الأولى أو الثانية حسب تصنيف الجمعية الأمريكية للتخدير والذين خضعوا لجراحات منظار الركبة، وتم إستبعاد المرضى الذين يعانون من عجز عصبى فى الطرف السفلى، سيولة بالدم، إلتهابات موضع حقن التخدير الموضعى وكذلك من كان لديهم حساسية من التخدير الموضعى.

وقد أظهرت نتائجنا أن تخدير القناة المقربة آمن وفعال كتخدير العصب الفخذي فى تسكين آلام ما بعد عمليات منظار الركبة بالإضافة إلى إعطاء المريض القدرة على الحركة المبكرة فيما بعد العملية.