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Efficacy of Femoral Nerve Block Compared to Adductor Canal Block for Post Operative Pain Management After Total Knee Arthroplasty

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KEYWORDS

Adductor canal block Femoral nerve block Numeric rating scale Postoperative pain Straight leg raise Total knee arthroplasty

DECLARATIONS

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ABSTRACT

Background: After total knee arthroplasty, regional anaesthetic methods such as femoral nerve and adductor canal blocks are frequently used to ease the pain. Objective: To assess how femoral nerve and adductor canal blocks affect post-operative mobility, healing time, and pain management following total knee arthroplasty. Methodology: From October 8, 2023, to September 25, 2024, a tertiary care hospital hosted this cross-sectional study. We randomly assigned 160 patients receiving unilateral primary total knee replacement to either the adductor canal or femoral nerve block (80 in each group). Before surgery, each group had their nerve block. Postoperative pain was measured at 6, 12, 18, and 24 hours using the numeric rating scale. Mobility on postoperative days 1–7 and the time until straight leg raise were recovery markers. **Results:** Age (femoral nerve block: 65.4±8.3 years; ACB: 64.8±7.9 years), gender distribution, body mass index, and ASA physical state did not significantly differ across groups. At every time point, the femoral nerve block group's pain scores were significantly lower: at 6 hours, this score was 3.8±1.2, compared to the adductor canal block's 4.2±1.3 (p=0.04). These tendencies continued at later evaluations (p<0.03). About 35% of the femoral nerve block group and 67.5% of the ACB group, respectively, attained straight leg raise greater than 30° on Day 1 (p<0.001). By Day 7, 93.75% of the femoral nerve block group and 100% of the ACB group had pain-free straight leg raise (p=0.05). According to multiple regression analysis, femoral nerve block was substantially linked to quicker recovery and lower pain scores (B=-0.5, p=0.001). **Conclusion**: After total knee arthroplasty, femoral nerve block provides better postoperative pain management and a faster recovery than adductor canal block, although adductor canal block allows for earlier leg mobilization and improves straight leg raise recovery.

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INTRODUCTION

Total knee arthroplasty (TKA) is a common and successful surgical procedure for treating osteoarthritis in the knee. TKA surgeries have become the most popular surgical surgery in industrialized nations, exceeding all other procedures, in the previous ten years. The healing process may be hampered by the moderate to severe postoperative discomfort that TKA is known to cause. People who experience discomfort after TKA are more likely to experience a number of postoperative complications, including infections. ioint loosening, reflex sympathetic dystrophy, and immobility-related problems such as deep vein thrombosis (DVT).^{2,3}

Because of its high effectiveness in lowering pain and its ability to spare opioids, femoral nerve block (FNB) is the gold standard percutaneous neuromodulation (PNM) treatment for patients undergoing TKA.4,5 Since performing TKA as an outpatient procedure has reduced the length of stay (LOS) in the hospital, it is becoming more widely acknowledged that strong analgesia that maintains motor strength during rehabilitation is a crucial component of the current perioperative protocol after procedure. An adductor canal block (ACB), which produces a sensory block with minimal motor involvement, has been shown to be a useful part of a multimodal approach for post-TKA pain management.6

Peripheral nerve blocks (PNBs) are a major method of pain relief after TKA. In addition to reducing pain, nerve blocks also shorten hospital stays and lower the risk of readmission.⁷⁻⁹ A frequent nerve block operation that has been shown to reduce opioid usage and decrease hospital stays following TKA is an FNB. However, if FNB causes a reduction in quadriceps muscle strength, which makes postoperative ambulation difficult, patients may be at increased risk of falls after surgery. 10-12 ACB is another nerve block technique that is currently attracting the attention of scientists since it may be superior to FNB. Several studies have demonstrated that is better than FNB in maintaining postoperative quadriceps muscle strength, facilitating postoperative ambulation. and functional encouraging recovery sacrificing pain control. 13-15 However, in terms of quadriceps strength, analgesic effect, and

postoperative functional recovery, two recent trials did not find a statistically significant difference between ACB and FNB. Thus, the purpose of this study is to evaluate the effectiveness of FNB and ACB following TKA in terms of pain management, recovery time, and post-procedure mobility.

METHODOLOGY

This cross-sectional study examined the efficacy of ACB and FNB in reducing postoperative pain following TKA. The study was cross-sectional and conducted at a tertiary care facility from October 8, 2023, to September 25, 2024. All participating patients provided written informed permission, and the study was approved by the institutional review board. After enrollment, 160 patients were split into two groups at random: FNB and ACB (80 in each group). Participants were to be between 45 and 80 years old and free of any illnesses that would preclude them from undergoing regional anaesthesia for a unilateral primary total knee arthroplasty. Participants were excluded if they had a history of opioid use disorders, peripheral vascular disease, local anaesthetic allergies, neurological or conditions.

Every patient received a comprehensive which included preoperative evaluation, gathering demographic information. categorizing their physical condition accordance with the American Society of Anaesthesiologists (ASA), and searching for any co-occurring conditions such as osteoarthritis. diabetes, or hypertension. We first used the Numeric Rating Scale (NRS) to quantify the patient's pain at baseline in order to develop a thorough anesthesia plan. General anaesthesia was administered to both groups prior to surgery. Using ultrasonography to guide the injection, 20 mL of 0.25% bupivacaine was injected into the femoral nerve at the inguinal crease in the FNB group to perform a femoral nerve block. The ACB group received an adductor canal block, with the needle aimed at the saphenous nerve inside the adductor canal, using the same quantity and dosage of local anesthetic and a similar technique. The block was administered before to the incision to ensure adequate analgesia before the procedure started. Using the NRS to measure

Table 1. Demographic characteristics

| | Variables | FNB Group (n=80) | ACB Group (n=80) | p-value |
|--------------------------|----------------------|---------------------|---------------------|---------|
| Age (years) | | 65.4±8.3 | 64.8±7.9 | 0.64 |
| BMI (kg/m ²) | | 30.2±5.4 | 29.8±5.1 | 0.57 |
| Gender | Male | 40 (50.0%) | 38 (47.5%) | 0.76 |
| | Female | 40 (50.0%) | 42 (52.5%) | |
| Residence | Urban | 45 (56.3%) | 43 (53.8%) | 0.75 |
| | Rural | 35 (43.7%) | 37 (46.2%) | |
| ASA Physical Status | ASA I | 12 (15.0%) | 10 (12.5%) | 0.68 |
| | ASA II | 58 (72.5%) | 60 (75.0%) | |
| | ASA III | 10 (12.5%) | 10 (12.5%) | |
| Comorbidities | Hypertension | 40 (50.0%) | 42 (52.5%) | 0.78 |
| | Diabetes Mellitus | 18 (22.5%) | 20 (25.0%) | 0.72 |
| | Osteoarthritis | 70 (87.5%) | 68 (85.0%) | 0.67 |
| Operative Side | Left | 38 (47.5%) | 40 (50.0%) | 0.74 |
| | Right | 42 (52.5%) | 40 (50.0%) | |

pain at 6, 12, 18, and 24 hours after surgery was one the numerous methods used to assess postoperative pain management. Additionally, measures were made to assess mobility and measure the time required to perform a straight leg raise (SLR) on days 1 through 7 following the procedure. Mobility was defined as the ability to sit up, transfer from a bed to a chair, and walk with assistance. Mobility, SLR times, pain scores, and group allocations were recorded and assessed by unaffiliated observers. Secondary outcomes including opioid intake (morphine equivalents), postoperative complications (e.g., nausea, vomiting, hemorrhage, infection), and the need for further analgesia were associated with these primary outcomes.

R Studio and SPSS version 26.0 were used for statistical analysis, descriptive statistics were calculated for variables both before and after surgery. The independent t-test was employed for between-group comparisons with continuous data. Categorical data were compared using the chi-square or Fisher's exact test. In statistical

terms, a p-value of less than 0.05 was considered significant. To find out if nerve block type had an independent effect on pain scores and recovery outcomes, the researchers of the study employed multiple regression analysis after adjusting for factors like age, body mass index (BMI), and baseline pain levels.

RESULTS

There were no notable differences between the ACB and FNB groups in terms of clinical characteristics or demographics. Both groups were identical in terms of body mass index $(30.2\pm5.4 \text{ kg/m}^2 \text{ for FNB and } 29.8\pm5.1 \text{ kg/m}^2)$ for ACB, p=0.57), average age (65.4±8.3 years for FNB and 64.8 ± 7.9 years for ACB, p=0.64), and gender distribution (50 percent male in Regarding residence, ASA both groups). and comorbidity profiles physical state, mellitus. (osteoarthritis, diabetes hypertension; p-values ranged from 0.67 to

Table 2: Pain assessment at different intervals (NRS)

| Time (Hours) | FNB Group (n=80) | ACB Group (n=80) | p-value | |
|-----------------|------------------------|---------------------|---------|--|
| 6 Hours | 3.8±1.2 | 4.2±1.3 | 0.04 | |
| 12 Hours | 4.1±1.3 | 4.6±1.4 | 0.03 | |
| 18 Hours | 4.3±1.4 | 4.8±1.5 | 0.02 | |
| 24 Hours | 4.5±1.5 | 5.1±1.6 | 0.01 | |

0.78), no significant differences were seen between the groups. There was no appreciable variation in the percentage of left- and rightsided procedures between the two groups, which were nearly equal (p=0.74). Patients in the FNB group experienced significantly less discomfort at 6, 12, 18, and 24 hours following surgery than those in the ACB group. At six hours, the ACB group recorded a slightly higher level of pain $(4.2\pm1.3; p=0.04)$ than the FNB group, which had an average of 3.8±1.2. The difference continued after 12 hours (FNB: 4.1±1.3 vs ACB: 4.6±1.4, p=0.03), 18 hours (FNB: 4.3±1.4 vs ACB: 4.8±1.5, p=0.02), and 24 hours (FNB: 4.5±1.5 vs ACB: 5.1±1.6, p=0.01). These findings suggest that the FNB group experienced significantly less pain during the trial than the ACB group. The table compares the leg mobilization after surgery on different days for the ACB and FNB groups. The proportion of patients in the FNB group who were able to move their legs was significantly lower (37.5%) than in the ACB group, which had 70% mobility on the first day (p<0.001). Over time, both groups improved, but the disparity persisted. Only 60% of FNB patients were able to mobilize by the second day, compared to 85% of patients in the ACB group (p=0.01). By day seven (p=0.04), leg mobility had increased to 90% in the FNB group and 98.75% in the ACB group. By the end of the week, 93.75% of the FNB group and 100% of ACB group had achieved full leg the mobilization (p=0.05). Although both groups shown significant improvement in leg mobility over time, the ACB group's percentage of patients who were able to mobilize their legs at each interval was consistently higher.

Table 3 compares the SLR test outcomes between the FNB and ACB groups at various intervals after surgery. The FNB group had a much lower rate of 35% (p<0.001) than the ACB group, which saw 67.5% of patients achieve an SLR greater than 30° on the first day. With a p-value of 0.002, by day two, 85 percent of the ACB group and 56.25 percent of the FNB group had SLRs greater than 45%. By day three, 92.5% of the ACB group had an SLR higher than 60%, compared to 75% of the FNB group (p=0.01). On day 4, 97.5% of the subjects in the ACB group and 87.5% of the subjects in the FNB group obtained a complete SLR (p=0.04). By the conclusion of the week, 92.5% of the FNB group reported no pain during SLR, while 100% of the ACB group did the same (p=0.05). The ACB group demonstrated more consistent and superior results, particularly in reaching higher levels of SLR earlier than the FNB group, despite the fact that both groups made progress over time.

FNB and ACB were compared in this study, and the results of the regression analysis show a relationship between a number of variables and the study's conclusions. With a p-value of 0.001 and a coefficient of -0.5 (95% CI: [-0.8, -0.2]), the group variable (FNB vs. ACB) indicates a statistically significant negative correlation between the two variables. The outcome marginally increases with each year, according to the minor positive coefficient for age (0.01, 95% CI: [0.00, 0.02]), which is statistically significant (0.046). Additionally, there is a significant gender component; males perform better (0.3, p=0.003). There was no significant effect of BMI on the outcome (p=0.183). Lastly, the substantial positive correlation between the two variables (coefficient: 0.6, p=0.001) indicates that a higher result is associated with a higher baseline pain level.

The correlation matrix presented above shows how many variables connected to post-operative recovery are related to one another. The positive connection between the two scores (0.72) indicates that higher pain ratings at 6 hours are associated with higher pain ratings at 24 hours. There were significant correlations between the number of days needed to achieve an SLR and pain scores at 6 and 24 hours (0.45 and 0.6, respectively), suggesting that longer recovery durations are associated with greater pain scores at these intervals. Mobility and the number of

Table 3: Leg mobilisation following surgery on different days

| Days | FNB Group (n=80) | ACB Group (n=80) | p- value |
|-----------------|---------------------|---------------------|-------------|
| 1st Day | 30 (37.5%) | 56 (70%) | <0.001 |
| 2nd Day | 48 (60%) | 68 (85%) | 0.01 |
| 3rd Day | 58 (72.5%) | 72 (90%) | 0.02 |
| 4th Day | 64 (80%) | 75 (93.75%) | 0.03 |
| 5th Day | 68 (85%) | 78 (97.5%) | 0.01 |
| 6th Day | 70 (87.5%) | 78 (97.5%) | 0.02 |
| 7th Day | 72 (90%) | 79 (98.75%) | 0.04 |
| After 1 Week | 75 (93.75%) | 80 (100%) | 0.05 |

days to SLR have a substantial negative correlation on days 3 and 7 (-0.65 and -0.7, respectively), indicating a relationship between delayed SLR recovery and decreased mobility. Mobility on day 3 and 7 have a strong positive correlation (0.88), suggesting that patients who exhibit greater mobility on day 3 also typically exhibit greater mobility on day 7.

DISCUSSION

Total knee arthroplasty is a surgical surgery with a very high success rate and a very good postoperative survival rate. Since the main objective of TKA is to decrease pain and improvefunctional mobility, there must be any leeway in handling this aspect of the patient's complaint. In addition to causing physical suffering, severe, untreated postoperative pain prolongs hospital stays, hinders recovery, and raises the possibility of long-term pain issues. Prior research indicated that a greater number of patients experienced severe pain following TKA, and that the procedure was poorly Approximately managed. 16,17 44-57% individuals have pain within the first three days after TKA. About 19% of TKA patients report feeling unhappy as a result of the vicious cycle that occurs when sleep deprivation reduces the pain threshold. Therefore, it appears that postoperative discomfort persistent and disturbed sleep are significant predictors of chronic functional deficits one and three months after total knee arthroplasty. 18

Table 4: Straight leg raise (SLR) test at different time points following surgery

| Days | FNB Group (n=80) | ACB Group (n=80) | p- value |
|--------------------------------------|---------------------|---------------------|-------------|
| 1st Day SLR >30° | 28 (35%) | 54 (67.5%) | <0.001 |
| 2nd Day SLR >45° | 45 (56.25%) | 68 (85%) | 0.002 |
| 3rd Day SLR >60° | 60 (75%) | 74 (92.5%) | 0.01 |
| 4th Day Full SLR | 70 (87.5%) | 78 (97.5%) | 0.04 |
| After 1 Week Pain-free SLR | 74 (92.5%) | 80 (100%) | 0.05 |

Our results demonstrate that, in terms of pain management and opioid use during TKA, neither anesthesia method is better than the other, which is in line with earlier studies. A recent meta-analysis found that the effects of ACB and FNB on patients undergoing TKA were comparable. 4 Both groups reported similar levels of discomfort and morphine intake during perioperative period at 1, 24, and 48 hours following surgery. 19,20 The ACB and FNB groups were also shown to be similar in terms of opioid use and pain scores by Jaeger et al. and Kim et al. In terms of both muscle strength and mobility, ACB performs better than FNB, which is in line with previous study findings. By obstructing the motor fibers of the femoral nerve distally, where they split off, ACB is thought to spare the quadriceps. Studies by Jaeger et al. and Kwofie et al. showed that ACB maintained quadriceps strength differently than FNB. Patients receiving continuous ACB had quadriceps strength that was 52% of the baseline value, according to Jaeger et al.'s study, but patients receiving continuous FNB alone had strength of 18%.^{21,22}

The meta-analysis model showed no statistically significant difference in fall risk between the two regimens. In any case, after administering FNB, Kwofie et al. found a higher risk of quadriceps muscle weakness and falls using the Berg Balance Scale. After 48 hours, Elkassabany et al. discovered that the FNB group experienced a higher rate of falls using the Tinetti Scale for gait and balance. Furthermore, it was demonstrated

Heatmap of Variable Correlations

1.0

0.8

Pain Scores at 24h - 0.72

1.00

0.85

0.60

1.00

0.85

0.70

1.00

0.88

1.00

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Figure 1: Heatmap, highlighting the relationships between variables

by Thacher et al. that FNB weakens the quadriceps; during physiotherapy, 13% of FNB patients and 2% of ACB patients suffered nearfall occurrences (knee-buckling), a statistically significant difference.^{23–25} The two peripheral nerve blocking techniques being examined in this study have a lot of potential advantages, but they might also have significant disadvantages.

Peripheral nerve blockades carry several dangers, such as the possibility of nerve damage and systemic and local toxicities because of the large amount of local anesthetic used. Although ACB helps maintain muscular strength following surgery, there are some hazards associated with it, including the possibility of infection, neuropathy, and myositis due to the local anesthetic that is injected into the adductor canal close to the surgical site. Additionally, constriction tourniquet mav result ischemia.²⁶⁻²⁸ The operation will require a highly experienced physician if the nerves are deep or small, or if the patient has subcutaneous emphysema, edematous tissues, or a high body mass index, all of which are known to diminish the ultrasound's visibility and make nerve blockade difficult. This is yet another significant barrier to the use of peripheral nerve blocking procedures guided by ultrasonography.^{29,30}

The single-center approach and homogeneous sample population may limit the generalizability

of the findings. Rapid surgical analgesia is the main focus, with long-term consequences like functional recovery or chronic pain management usually ignored. Bias may result from subjective pain assessments and inconsistent nerve block administration by different professionals.

CONCLUSION

In conclusion, in terms of pain management, FNB was found to be more effective than ACB for postoperative pain management after total knee arthroplasty, as evidenced by lower pain scores at multiple time intervals. The adductor canal block was found more efficacious for faster restoration of leg mobilization and SLR in the ACB group. Additionally, FNB facilitated reduced post-operative discomfort, contributing to a more efficient and pain-free postoperative period. Multiple regression analysis indicated that FNB significantly improved outcomes with baseline pain scores. Age and gender also play important roles. These findings suggest that FNB is a preferred technique for postoperative pain total management in knee arthroplasty. enhancing both pain control and reduced postoperative restlessness.

DECLARATIONS

Consent to participate: Written consent had been taken from patients. All methods were

Table 5: Regression analysis

| Variables | Coefficient (B) | 95% CI | Std. Error | t-value | p-value |
|----------------------------|--------------------|---------------|------------|---------|---------|
| Groups | | | | | |
| (FNB = 1, ACB = 0) | -0.5 | [-0.8, -0.2] | 0.15 | -3.33 | 0.001 |
| Age | | | | | |
| (years) | 0.01 | [0.00, 0.02] | 0.01 | 2 | 0.046 |
| Gender | | | | | |
| (Male = 1, Female = 0) | 0.3 | [0.1, 0.5] | 0.1 | 3 | 0.003 |
| BMI | | | | | |
| (kg/m^2) | 0.02 | [-0.01, 0.05] | 0.015 | 1.33 | 0.183 |
| | | | | | |
| Baseline Pain Score | 0.6 | [0.4, 0.8] | 0.1 | 6 | 0.001 |

performed following the relevant guidelines and regulations.

Availability of data and materials: Data will be available on request. The corresponding author will submit all dataset files.

Competing interests: None

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