

Comparing preemptive injection of peri-articular-multimodal drug with oral celecoxib for postoperative pain management in total knee arthroplasty: A randomized clinical trial

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Background: Total knee arthroplasty (TKA) is associated with tremendous postoperative pain, and pain relief should concisely be considered. This study aims to compare the efficacy of preemptive periarticular multimodal drug injection versus placebo and oral celecoxib on postoperative pain control after TKA. **Materials and Methods:** This study is a randomized clinical trial on 146 patients candidate for TKA who were randomly allocated to three treatment groups, including (1) a cocktail consisting of bupivacaine, morphine, epinephrine, and ketorolac ($n = 48$), (2) only epinephrine (placebo group) ($n = 49$), and (3) 400 mg celecoxib orally (control group) ($n = 49$) using the Random Allocation software. The injections and oral therapy were performed within 15 min before the surgical procedure. The study's primary outcome was the Knee Society Score (KSS) calculated at baseline, within 6 weeks and 6 months postoperatively. Range of motion (ROM) and Visual Analog Scale (VAS) to assess pain intensity as the other primary outcomes were evaluated before the procedure, within 24 h, 48 h, and 6 weeks postoperatively. **Results:** The three studied groups were similar regarding demographic characteristics, including age ($P = 0.33$), gender distribution ($P = 0.65$), and involved knee side ($P = 0.94$). Baseline comparison of KSS ($P = 0.39$), VAS ($P = 0.24$), and ROM ($P = 0.37$) among the groups revealed insignificant differences. All the studied groups showed a statistically significant trend of improvement in KSS, VAS, and ROM ($P < 0.001$), while the comparison of the three groups in terms of KSS ($P = 0.001$), VAS ($P < 0.001$), and ROM ($P < 0.001$) revealed remarkable superiority of multimodal injection to the other treatments. **Conclusion:** Preemptive periarticular multimodal drug injection, including bupivacaine, morphine, epinephrine, and ketorolac, can cause considerable postoperative pain relief and better ROM achievement in comparison to placebo or oral celecoxib.

Key words: Multimodal analgesia, pain, range of motion, total knee arthroplasty

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INTRODUCTION

Total knee arthroplasty (TKA) is a painful but efficient orthopedic procedure used to treat patients suffering from degenerative disease, osteoarthritis in particular.^[1,2] TKA, which is mostly performed on the elderly, may cause considerable postoperative pain that can lead to an increased hospital stay, disability to take part in rehabilitation programs, poor outcomes, and increased

rate of referral to health-care centers as well as chronic pain in up to 35% of patients.^[3,4]

The concept of pain turned into a significant issue to the extent that it has been declared as the fifth parameter of the vital sign by the Joint Commission on Accreditation of Health-care Organizations (JCAHO).^[5] Accordingly, varieties of pain-relieving approaches have been provided to manage postoperative pain.^[6] Nevertheless, the old age of the patients who are the majority of

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patients requiring TKA can limit the alternatives for this aim.^[7]

The traditional methods include the use of analgesic agents such as nonsteroidal analgesic agents (NSAIDs), acetaminophen, gabapentin, and pregabalin, which not only has short-term effects but also may cause drug-related adverse effects and are mostly required to be administered for a long time.^[8] Nevertheless, preoperative use of NSAIDs, cyclooxygenases-2 inhibitors, in particular, has shown promising outcomes as it can reduce the amount of retrieved inflammatory factors at the site of surgery in addition to its opioid-sparing effect, which can reduce opioid-related side effects, including nausea and vomiting, which contribute to the postoperative morbidity associated with TKA.^[7]

Using opioids for postoperative pain management is a common and useful option but has complications such as nausea and vomiting, respiratory depression, drowsiness, decreased gut motility, and urinary retention.^[9] Other postoperative pain relief strategies with acceptable outcomes are continuous epidural, lumbar plexus, femoral, and sciatic nerve blocks. These approaches can effectively control postoperative pain and decrease narcotic consumption, though have several side effects such as epidural bleeding, less muscle strength, urinary retention, and neuronal damages.^[10]

The use of multimodal local analgesics at the surgical site during or by the end of the procedure is another optimal option for reducing postoperative pain with minimal side effects in TKA. Intra-articular injection of mixed analgesics following TKA surgery has been accompanied by decrease in further requirements for analgesia and led to earlier hospital discharge;^[11,12] however, studies in this regard have not achieved unified outcomes.^[13,14]

Despite various introduced strategies for postprocedural pain management among patients undergoing TKA, limited comparative evidence is available to assess the best pain controlling approach. Besides, numerous patients are suffering from chronic postoperative pain worldwide to the extent that they are dissatisfied with their procedure.^[15,16] Therefore, the current study is raised to assess the efficacy of periarticular multimodal injections versus placebo and oral analgesic therapy for post-TKA pain relief.

METHODS

Participants and study design

This is a randomized clinical trial conducted on 146 patients undergone TKA in orthopedic hospitals affiliated at Isfahan University of Medical Science (IUMS) in 2016.

The presence of osteoarthritis, making the patients candidate for primary unilateral TKA, and body mass index of 20–30 kg/m² were determined as inclusion criteria for participation in this study.

Abnormal renal function test (Cr clearance <60 ml/min), abnormal liver function test, myofascial pain syndrome or fibromyalgia, and patient's unwillingness for participation in the study were considered as unmet criteria. Patients with allergy, drug and/or alcohol abuse, inflammatory arthritis, history of previous major knee surgery, and those who were a candidate for bilateral TKA were the other conditions defined as unmet criteria, as well.^[17] American College of Rheumatology Preliminary Diagnostic Criteria were used to diagnose fibromyalgia and myofascial pain syndrome.^[18]

Failure to refer for follow-up assessments and more than 20% defects in medical records were the current study's exclusion criteria.

The study was approved in the Ethics Committee of IUMS (ethical code: IR.mui.rec. 1396.3.048), and the study protocol was registered in the ClinicalTrials.gov (NCT05324995) and also was explained entirely for the patients, they were reassured about their information confidentiality, and written consent was obtained.

This study was designed as a census study; therefore, all the patients who met the criteria for participation in this study were recruited. Participants were randomly divided into three groups using the Random Allocation software, so each patient was provided with a specific number from 1 to 146 by the software and allocated to one of the groups.

The investigator who performed the baseline and follow-up assessments and the patients were unaware of the study group; as the investigator was absent at the time of the surgery and all the patients administered a pill before the surgical procedure (one group was treated with celecoxib and the other two ones administered placebo similar in shape and color with celecoxib).

The surgical technique

All TKA surgeries were performed under spinal anesthesia using 10–15 mg Marcaine (bupivacaine). About 1 g cefazoline (EXIR®, Boroujerd, Iran) was administered for patients within an hour before the surgery.

TKA surgery was performed for all participants by a reference skilled orthopedic surgeon electively using knee prosthesis (Zimmer®, Warsaw, IN, USA).

After proper sterilization, a preemptive periarticular injection was done for participants in the first and second

groups. Then, an anterior midline skin incision was done within 8 cm above the patella to 2 cm distal to the tibial tubercle.

A pneumatic tourniquet was pumped up before incision and collapsed before the soft tissue repair at the end of surgery. A low vacuum intraarticular drain was used during soft tissue repairing for postoperative wound drainage and then removed 48 h after surgery.^[19]

For pain control following the surgery, meloxicam (15 mg daily), celecoxib (400 mg daily), acetaminophen (1000 mg every 8 h), tramadol (50 mg every 8 h), ketorolac (30 mg slow IV every 8 h, 4-dose max), and morphine (5–10 mg slow IV, PRN) were prescribed for the patients.

On the 1st day after surgery, all patients were mobilized, and daily 60 mg subcutaneous enoxaparin (Clexane®, Prefilled syringes Sanofi, Seine Martine, France) was used for deep-vein thrombosis (DVT) prophylaxis for 14 days. Prophylactic treatment of patients with a higher risk of DVT was continued with bridging therapy of warfarin, and then, warfarin was merely prescribed for a further 3 months.^[20]

After that, the patients were discharged within 3 days after the surgery if they were appropriately mobilized and had dry clean wounds. After discharge, standard rehabilitation methods were used for patients at home or in physiotherapy centers. The rehabilitation schedule consisted of strengthening lower extremity muscles by concentration on the quadriceps, and gait training with weight bearings was allowed by the reconstructed knee to perform the necessary daily life.^[21]

The interventions

The first group was treated using a preemptive periarticular injection of multimodal drugs. Therefore, a combination of drugs consisted of 50 mg bupivacaine hydrochloride 0.5% (AstraZeneca, Cenexi, France), 1 ml morphine sulfate 10 mg/ml (DarouPakhsh, Tehran, Iran), 300 mcg epinephrine (1:1000) (DarouPakhsh, Tehran, Iran), and 30 mg ketorolac (Caspian Tamin, Rasht, Iran) diluted by 0.9% sodium chloride solution to make a total 100 ml of injection drug was injected in the periarticular area. The injections were done within 15 min before the incision in seven areas of the joint as followed: 15 ml in posterolateral soft tissue and lateral femoral periosteum, 15 ml posteromedial soft tissue and medial femoral periosteum, 20 ml inferomedial capsule, 20 ml superomedial capsule, 10 ml lateral capsule, 10 ml medial subcutaneous tissues, and 10 ml lateral subcutaneous tissue.

The second group received 300 mcg epinephrine (1:1000) (DarouPakhsh, Tehran, Iran) peri-articular, similar to the first group.

The third group administered celecoxib (200 mg) orally immediately before the surgery initiation.

Outcomes

Before the surgical procedure, patients' age, gender, and the operated knee side were recorded in the study checklist.

The study's primary outcomes included measurements of Knee Society Score (KSS) and range of motion (ROM) as the determinant of knee function and Visual Analog Scale (VAS) as the means for pain assessment.

The KSS is an assessment of the knee that comprises two arms, the functional ability and clinical examination scores, each with a 100-point denominator. One-hundred points are allocated to the functional capacity, including walking distance, stair-climbing ability, and the use of walking aids. The latter 100 points are used to assess knee motion, stability, and pain. This scoring is divided into two subscales, 50 points for evaluating motion, stability, and alignment. The remained 50 points for pain intensity assessment.^[22] According to the scores 160–200, 140–159, 120–139, and <120, the KSS total score was defined as excellent, good, fair, and poor, respectively.^[19] KSS was at baseline, and then, within 6 weeks and 6 months postoperatively.

In addition, the knee ROM was evaluated using a standard goniometer.^[23]

VAS was used to evaluate the patient's postoperative pain intensity with a 0–10 rating scoring scale.^[16]

The patient's postoperative pain intensity and ROM were evaluated in the preoperative visit (baseline), within 24 h, 48 h, and 6 weeks following surgery.

Statistical analysis

The obtained data were entered into the Statistical Package for the Social Sciences (version 22, IBM Corporation, Armonk, NY, USA). The descriptive data were presented in mean, standard deviation, absolute numbers, and percentages. In order to assess the normality of data, the Kolmogorov–Smirnov test was administered. The Chi-square test was utilized to compare the categorical variables, and the quantitative variables were compared using independent *t*-test for two variables and ANOVA for more than two variables. Tukey *post hoc* test was utilized to detect the difference between two independent variables evaluated by ANOVA test. Repeated-measure ANOVA assessments were performed to compare independent quantitative variables in repeated measurements. $P < 0.05$ was considered statistically significant level.

RESULTS

In this clinical trial, 170 patients undergone unilateral TKA were assessed, among which 164 ones were eligible to participate in the study. Five patients did not meet inclusion criteria, and two ones refused to participate in this study. During the study, three patients in the second group did not receive the intervention because of unwillingness to participate in the study, and eight ones did not refer for follow-ups (2 in the first, 5 in the second, and 1 in the third group) and thus were excluded from the study. The information of one patient in the first group was excluded because of having congestive heart failure. Eventually, data about 146 patients were analyzed [Figure 1].

One-hundred and forty-six patients with a mean age of 63.38 ± 5.486 years (37–81) and female gender predominance 92.5% ($n = 135$) were assessed.

The comparison of demographic factors, including age ($P = 0.33$), gender distribution ($P = 0.33$), and side of the operated knee ($P = 0.94$), was not statistically different among the three groups. Detailed information is provided in Table 1.

Baseline KSS ($P = 0.39$), ROM ($P = 0.24$), and VAS ($P = 0.37$) assessments of the three groups revealed insignificant differences; however, further comparison of KSS measurements within 6 weeks ($P < 0.001$) and 6 months ($P < 0.001$) postoperatively represented significant differences among the groups. *Post hoc* evaluations revealed multimodal injection superiority to the other two techniques ($P < 0.001$), whereas the control and placebo groups, were similar ($P > 0.05$). The VAS and ROM assessments within 24 and 48 h after the surgical procedure, as well as within the next 6 months, were accompanied by remarkable better statuses of multimodal injection than the

Table 1: Demographic data of participants

Variable	Multimodal injection, n (%)	Placebo group, n (%)	Control group, n (%)	P
Age (mean±SD)	63.44±6.164	64.18±4.902	62.53±5.319	0.330*
Gender				
Male	5 (10.4)	3 (6.1)	3 (6.1)	0.65**
Female	43 (89.6)	46 (93.9)	46 (93.9)	
Side				
Right	26 (54.2)	28 (57.1)	28 (57.1)	0.94**
Left	22 (54.8)	21 (42.9)	21 (42.9)	

*One-way ANOVA test did not show significant differences, **Chi-square test did not show significant differences. SD=Standard deviation

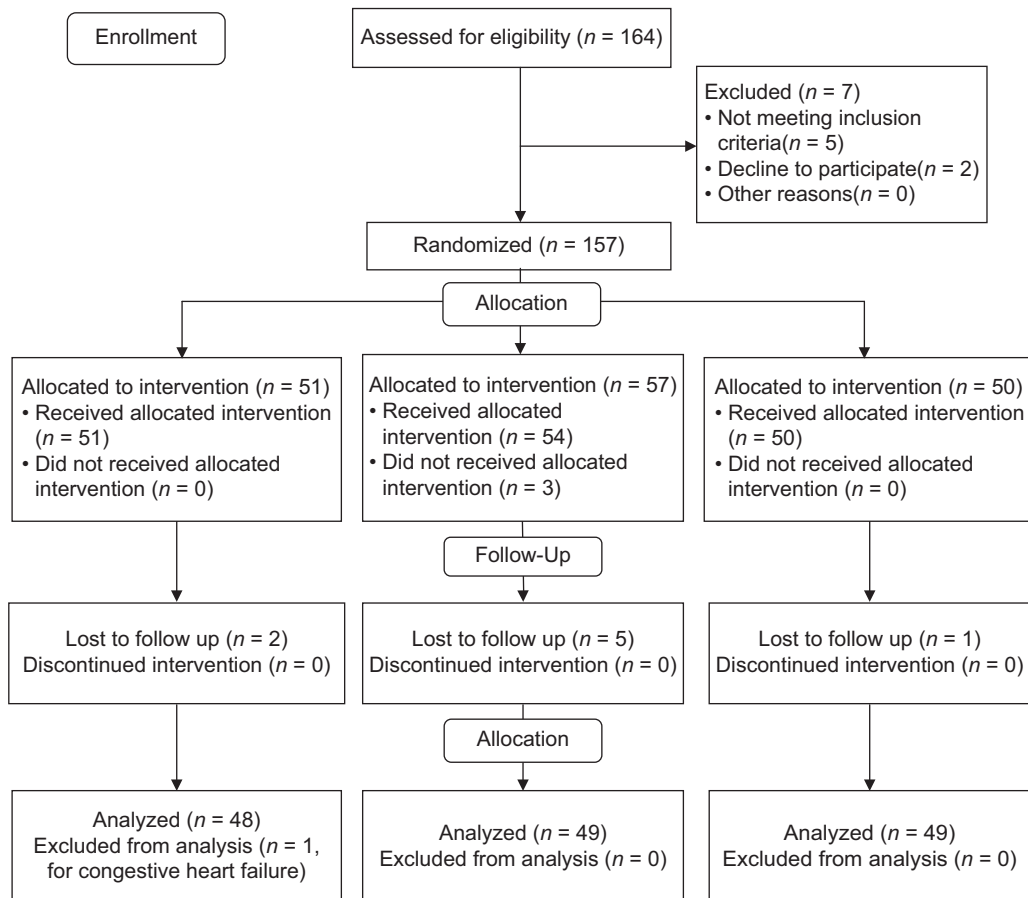


Figure 1: The flow diagram of randomization of patients

two other interventions ($P < 0.05$) in all the assessments. The *post hoc* measurements showed remarkable better outcomes of multimodal injection than the other two interventions in all assessments ($P < 0.05$), while generally, the other two groups were similar, but not in all measurements demonstrated in Table 2. The repeated measurements of KSS, VAS, and ROM revealed statistically significant improvements in all the interventions ($P < 0.001$ for all three groups represented as P_1); however, the comparison of the groups revealed remarkable superiority of multimodal injection to the other two techniques in terms of higher KSS scores ($P < 0.001$ represented as P_3), less pain intensity based on VAS ($P < 0.001$ represented as P_3), and better ROM ($P < 0.001$ represented as P_3).

DISCUSSION

This study's study population was not statistically different regarding age, gender, and side of the knee under TKA. Therefore, demographic variables could not influence the outcomes of the study. This study demonstrated that using the preemptive periarticular injection of multimodal medications, including bupivacaine hydrochloride, morphine, epinephrine, and ketorolac, can efficiently tranquilize postoperative pain. However, the two other techniques were accompanied by promising outcomes, as well. Nevertheless, the three groups' general comparison revealed the superiority of the preemptive periarticular

injection of multimodal medications to the other two techniques.

Surgery trauma during TKA decreases afferent nociceptive neuron thresholds and increases spinal neuron excitability. This is a somatic pain that may be present for months.^[24] Postoperative pain management in TKA is critical and includes systemic, local, and regional methods.^[25] Local injection of analgesic agents in the site of surgery is appropriate for controlling postoperative pain after TKA with limited side effects.^[17,24] Limited studies have demonstrated the efficacy of preemptive periarticular injection in controlling postoperative pain in TKA.

A reviewed study reported that periarticular injection of morphine in TKA surgery could significantly decrease postoperative pain compared to placebo.^[13] Another study used the preemptive periarticular injection of opioid, ketorolac, ropivacaine, and epinephrine and evaluated postoperative pain within 6 and 12 h after surgery and showed that this method could significantly decrease self-reported pain score and improve self-satisfactory presentation.^[17] Studies demonstrated that a mixture of opioids, NSAIDs, epinephrine, and long-acting local anesthetics could cause proper analgesia after TKA.^[17,26] Yue *et al.* presented that receiving a periarticular injection of multimodal drugs consisted of corticosteroids can cause earlier rehabilitation and hospital discharge, although pain

Table 2: Total contents of knee society score, visual analog scale and range of motion in three groups

Variable	Mean±SD			P	P'' (between multimodal injection and placebo)	P'' (between multimodal injection and control group)	P'' (between placebo group and control group)
	Multimodal injection	Placebo group	Control group				
KSS							
Baseline	1.49±24.894	1.94±23.942	6.84±13.801	0.39	0.918	0.224	0.26
Within 6 weeks	113.15±14.89	100.24±17.97	102.02±7.91	<0.001	<0.001	<0.001	0.53
Within 6 months	115.96±13.49	101.20±16.62	98.53±6.97	<0.001	<0.001	<0.001	0.31
P_1 (time)	<0.001	<0.001	<0.001				
P_3 (intervention)		0.001					
VAS							
Baseline	7±1.22	7.39±1.07	7.27±1.09	0.24	0.098	0.25	0.59
Within 24 h	6.39±1.45	8.80±0.88	8.10±0.65	<0.001	<0.001	<0.001	0.001
Within 48 h	5.02±1.25	6.20±1.22	6.18±0.80	<0.001	<0.001	<0.001	0.92
Within 6 weeks	3.46±1.14	4.15±1.25	3.67±0.65	0.006	0.002	0.31	0.028
P_1 (time)	<0.001	<0.001	<0.001				
P_3 (intervention)		<0.001					
ROM							
Baseline	87.89±10.41	90.10±10.02	90±3.68	0.37	0.21	0.23	0.95
Within 24 h	107.56±6.36	94.49±3.57	95±2.88	<0.001	<0.001	<0.001	0.57
Within 48 h	113.44±5.09	96.12±4.11	98.16±2.83	<0.001	<0.001	<0.001	0.015
Within 6 weeks	128.33±13.18	119.80±8.16	122.65±4.57	<0.001	<0.001	0.003	0.12
P_1 (time)	<0.001	<0.001	<0.001				
P_3 (intervention)		<0.001					

*ANOVA, ***Post hoc* Tukey test. P_1, P_2, P_3 at the 5% level error of repeated measures. KSS=Knee society score; VAS=Visual Analogue Scale; ROM=Range of motion; SD=Standard deviation

relief was not positively affected by cocktails containing corticosteroids in comparison to those without steroids.^[27] The further study presented that periarticular injection of multimodal medications had fewer side effects, caused the reduced incidence of nausea and vomiting, and also decreased the risk of wound infections.^[28]

The efficacy of multimodal medications may be attributed to its consistency. Opioid receptors are located in peripheral inflamed tissues and are expressed after surgical trauma. Mentioned receptors are responsible for sensory input to the central nervous system.^[29] NSAIDs can decrease peripheral sensitization and activation of nociceptors.^[30] Bupivacaine can block afferent peripheral nociceptive activity and also has anti-inflammatory effects.^[31] The epinephrine used in periarticular multimodal injections causes decreased local blood flow leading to local anesthesia toxicity reduction.^[32]

Several studies evaluated the effects of celecoxib on postoperative pain after TKA in comparison to placebo. However, there is no study compared this medication with the periarticular injection of multimodal medications. One study prescribed 400 mg celecoxib before surgery for patients undergoing TKA and then continued with 200 mg celecoxib for 12 days postoperatively. This method significantly improved postoperative knee ROM and decreased postoperative morphine requirement compared with the placebo.^[33] Another study showed that perioperative use of celecoxib was accompanied by less time to achieve maximal ROM and improved knee function within a year after surgical procedure.^[34]

Studies have presented the superiority of 400 mg celecoxib to the 200 mg doses.^[35] Moreover, our study results may have occurred due to ketorolac's use, as an NSAID, in our cocktail. Therefore, we want to recommend further studies prescribing preoperative 400 mg of oral celecoxib.

Limitations

Despite the appropriate population of this study, a significant limitation in the current report is to follow the patients for a short period; therefore, further studies with a larger sample population and more extended follow-up periods are strongly recommended. Another limitation was lacking hospitalization duration and postoperative opioid requirement assessment. The third limitation of this study may be attributed to the assessment of probable confounders. It is still possible that unmeasured confounders interfere with a part of the associations have been missed. Among the probable confounder, the patients' job, smoking, physical activity, and dietary status can be named that has not been assessed in the current report.

CONCLUSION

Preemptive periarticular multimodal drug injection, including bupivacaine, morphine, epinephrine, and ketorolac, can cause considerable postoperative pain relief and better ROM achievement in comparison to placebo or oral celecoxib. Thus, periarticular multimodal drug injection is recommended for patients who are candidates for TKA.

Further studies with a more extensive study population and considering more variables affecting the study outcomes are strongly recommended.

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Conflicts of interest

There are no conflicts of interest.

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