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The efficacy of local infiltrative analgesia in total knee arthroplasty regarding pain and movement in early postoperative period

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Abstract

Background: Total knee arthroplasty is a successful procedure for relieving pain and restoring function in cases with severe rheumatoid arthritis and osteoarthritis. Local infiltration analgesia is becoming more commonly used owing to the excellent pain relief, the low frequency of complications, and the anti-inflammatory effect.

Objective: To evaluate the efficacy of local infiltrative analgesia when it's uses in total knee arthroplasty patients for decrease the pain score and increase range of motion in early post-operative rehabilitation. **Methods:** a comparative prospective study carried in the period from October 2016 to September 2018, was applied for 36 patients in Medical City, Hospital of Specialized Surgeries and Private Nursing Home Hospital.

Results: In group A, means of ROM score one day and two days postoperatively were decreased than that before operation (105.12 versus 89.37, P=0.001 and 105.12 versus 87.37, P=0.001 respectively). Ingroup B, means score one day and two days postoperatively were decreased than that before operation (105.75 versus 88.50, P=0.001 and 105.75 versus 82.75, P=0.001 respectively). During rest, means of VAS score four hours, 12 hours, one day, and two days after operation were significantly decreased in (group A) compared to group B. During movement, insignificant differences in means of VAS score between study groups four hours, 12 hours, and two days after operation (P=0.258, 0.057, and 0.284 respectively).

Conclusion: Local infiltration analgesia effectively reduce pain scores during rest and movement also reduces hospital stay in patients undergoing total knee arthroplasty, and better physiotherapy outcome.

Keywords: Total knee arthroplasty, ROM score, VAS score, local infiltration analgesia

Introduction

Anatomy of knee joint

The knee joint is the largest and one of the most complex joints in the human body. A unique interaction of bones, muscles, menisci, and ligaments results in a compromise between stability and mobility ^[1]. Basically, it consists of two condylar joints between the medial and lateral condyles of the femur and the corresponding condyles of the tibia, and a gliding joint, between the patella and the patellar surface of the femur. Above are the rounded condyles of the femur; below are the condyles of the tibia and their cartilaginous menisci; in front is the articulation between the lower end of the femur and the patella. The articular surfaces of the femur, tibia, and patella are covered with hyaline cartilage ^[2].

Innervation of the knee joint

The neural innervation of the knee joint is complex which innervated by the articular branches of the muscles which move the joint. The articular nerves are derived from the femoral, obturator, tibial, common peroneal, and recurrent peroneal nerves. The femoral nerve, through its saphenous branch and also via its branches to the vast us medial is, intermedium, and lateralis muscles, supplies the suprapatellar recess, the patellar periosteum, and the anteromedial and anterolateral portions of the joint capsule. The medial, lateral, and posterior aspects of the joint capsule, the infrapatellar fat pad, tibial periosteum, and the superior tibio-

fibular joint are supplied by the tibial nerve. The common peroneal nerve supplies the anterolateral portion of the capsule, the infrapatellar fat pad, and the tibial periosteum. The obturator nerve supplies the superior part of the posteromedial capsule and the anteromedial aspect of the capsule. Cutaneous innervation of the anterior aspect of the knee is supplied by the femoral nerve ^[3].

Total Knee Arthroplasty (TKA)^[4].

Total knee arthroplasty (TKA) is the surgical treatment of choice for advanced osteoarthritis after failure of other modes of conservative management in properly selected patients. There are two designs of implants used for TKA:

•Unconstrained: which include cruciate retaining, Cruciate substituting and mobile bearing knees.

•Constrained: which include hinged and non-hinged. However unconstrained type is the most common type, used for uncomplicated knee problems, artificial components inserted into the knee are not linked to each other, have no stability built into the system, and relies on the patient's own ligaments and muscles.

Goal of TKA

- 1. Pain relief.
- 2. Restoration of normal limb alignment.
- 3. Restoration of a functional range of motion.

Technical Goals of TKA

- 1. The restoration of mechanical alignment.
- 2. Preservation (or restoration) of the joint line.
- 3. Balanced ligaments.
- 4. Maintaining or restoring a normal Q angle.

Absolute contraindications for TKA

- 1. Recent or current knee sepsis.
- 2. Remote source of ongoing infection.
- 3. Extensor mechanism discontinuity or sever dysfunction.
- 4. Recurvatum deformity secondary to muscular weakness.
- 5. Presence of painless well functioned knee arthrodesis

Post-operative analgesia

Postoperative Total knee arthroplasty (TKA) analgesia remains a challenging issue. It is reported that more than half of the patients undergoing knee replacement would experience severe pain in the early postoperative period Considerable postoperative pain interferes with patients' participation in physiotherapy, prolongs inpatient stay, lowers patient satisfaction, and leads to chronic pain and dysfunction. Thus, effective pain control in the immediate postoperative period is crucial for patients' convalescence after TKA ^[5].

Local infiltration analgesia (LIA) is becoming more commonly used owing to the excellent pain relief, the low frequency of complications, and the anti-inflammatory effect ^[6]. Options for postoperative pain control include patient administered narcotics, epidural anesthetics, and spinal anesthetics with adjuncts such as long-acting morphine and peripheral nerve blocks (with and without catheters). These concepts are widely used, but there are reports of multiple side effects secondary to parenteral opioids and problems associated with motor blockade after nerve blocks, which can lead to delays in rehabilitation ^[7].

Local infiltration analgesia is simple, practical, safe, and effective for pain management after knee surgery. We describe a multimodal technique for control of acute postoperative pain following lower limb joint replacement surgery. The technique is based on systematic infiltration of a mixture of ropivacaine, ketorolac, and adrenaline (RKA) around all structures subject to surgical trauma. The technique, known as local infiltration analgesia (LIA), was developed specifically to avoid sedation and facilitate rapid physiological recovery after lower limb arthroplasty in order to enable early mobilization and discharge ^[8].

Postoperative pain is usually severe after knee arthroplasty, In the last few years, several studies supporting the benefits of the LIA technique or the modified LIA technique in knee and hip arthroplasties have been published ^[9].

Visual Analog Scale (VAS)

The pain VAS is a uni-dimensional measure of pain intensity, which has been widely used in diverse adult populations, including those with musckelo-skeletal diseases ^[10].

The scale consists of one horizontal or vertical line, usually 10 centimeters in length, that is anchored with verbal descriptors of "no pain "and "pain as bad as it could be". Equipment needed. Pencil and paper. The VAS is also available as a plastic slide ruler and on colored cards that can be given to the respondent ^[11].

The pain VAS is self-completed by the respondent. The respondent is asked to place a line perpendicular to the VAS line at the point that represents their pain intensity $^{[2, 8, 9]}$.

A higher score indicates greater pain intensity, based on the distribution of pain VAS scores in postsurgical patients, who described their postoperative pain intensity as none, mild, moderate, or severe, the following cut points on the pain VAS have been recommended; no pain (0–4 mm), mild pain (5–44mm), moderate pain (45–74 mm), and severe pain (75–100 mm), The VAS takes one minute to complete ^[10].

Aim of study: To describe the efficacy of local infiltrative analgesia in TKA patients for decrease the pain score and increase range of motion in early post-operative rehabilitation.

Patients and Method

Study design and setting

A comparative prospective study carried in the period from October 2016 to September 2018 in Medical City/ Hospital of Specialized Surgeries and Private Nursing Home Hospital. (36) Patients enrolled in the current study. Total knee arthroplasty done by the same surgical team and then followed-up to November 2018, (11) of them were males and (25) were females (mean age, 64, 13 years), The objective is to evaluate the local infiltrative analgesia efficacy.

Inclusion criteria: All patients those indicate for primary TKA.

Exclusion criteria

- 1. Patients with a history of deep venous thrombosis or pulmonary embolism or on long-term warfarin therapy.
- 2. Patients undergoing revision procedure.
- 3. History of allergy or intolerance to 1 of the study drugs.
- 4. Patient with rheumatoid arthritis.
- 5. Abnormal liver enzymes.

Operative technique

Surgery done under general anesthesia or spinal anesthesia according to the senior anesthetist opinion and patient condition and comorbidities. Intra-operative mid-thigh pneumatic tourniquet used with pressure setting (300-350 mm Hg) and assistive leg holder had been used for all the patients.

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Intravenous antibiotics infused half an hour before induction of anesthesia and continued for further two days. Then 1gm of tranxamic acid given i.v before starting the procedure.

Anti-Thrombotic given 12 hours after surgery and continued for further 21 days in the form of subcutaneous low molecular weight heparin 4000 International units. The same surgical team performed the whole surgeries that eliminate technical differences and selection bias. In this study same implant design for all patient which was posterior-cruciate substitution. In this study the approach was for all patients, a Midline skin incision in slight knee flexion extended proximally about (3 cm) above the upper pole of patella and down to tibial tuberosity and then the arthrotomy done through medial Para patellar approach. We used the femur first technique so we start femur first by setting the IM (intramedullary) angle according to the x-ray measurement and cutting 9 mm from the more distal condyle then start sizing and chamfering the distal femur.

At this stage, not cut the notch and then shift to finish the tibia bone cut. We used intramedullary guide for tibia cut. The entry point near the base of the anterior tibial spine (anterior 1/3 of the tibial articular surface), reaming the entry hole, before resection confirm the amount of resection by using of stylus (2 mm) from the affected side and insure proper rotational alignment of the component After cutting the tibia 7-degree posterior slope and perpendicular to the mechanical axis, we used tibial tuberosity as marker to obtain proper rotational alignment of tibial component (medial 1/3).After finishing the extension gap, we start to cut the flexion gap, Used posterior referencing in sizing of femur implant. To obtain rotational alignment of the femoral component we chose 30 external rotations. Then start anterior-posterior cut and then chamfering the distal femur. Then started soft tissue balancing, before started to any soft tissue release, removed the osteophyte then carefully examine the flexion gap and extension gap by using spacer block and laminar spreaders, if obtain target (rectangular flexion and extension gap), so continued the procedure. In the LIA group a mixture of 150 mg (5 mg/ml) of Bupivacaine and 30 mg (1 ml) of Ketorolac and 0.5 mg epinephrine (0.5 ml), these were mixed with sterile normal saline solution to make up a combined volume of 150 mL in the operating room.

Fifty ml of this solution where injected in the subcutaneous tissue in the line of skin incision prior to start the surgery together with infiltration of arthrotomy site before doing the arthrotomy, another fifty ml of the solution was injected in flexed position into the posterior part of the capsule, the intercondylar area, and around the collateral in ligaments just before cementing the implants by using 50 ml 23-G needle.

Special care was taken to avoid infiltration of the common peroneal nerve and popliteal fossa to avoid injury to vessels

and sciatic nerve. The remainder was injected throughout the various soft tissue layers prior to wound closure, in this way, all tissues that were traumatized received the analgesic solution. Then intra articular (deep) drain was placed in the lateral gutter of the knee joint, another (superficial) drain was placed in the subcutaneous tissue. The wound was then closed in layers and adhesive dressings applied, Compression bandages were then applied finally. Postoperatively, regular 1 gm of paracetamol were given three times daily to both groups together with celecoxib 200 mg orally was given twice daily unless contraindicated. First assessment of pain score was four hours after surgery, then 12 hours after surgery, then once daily until the discharge. The assessment of pain according the VAS score was done during the rest and movement. The range of motion measured once daily until discharge and the last measure done six weeks after the surgery by advise the patient to sit on the edge of the bed, the straight leg raising test done at time of discharge, the straight leg raising test was successful if the participant could hold the lower limb 10 cm from the bed with fully extended knee for 10 seconds. The time to fulfillment of discharge criteria was recorded by a physician who was unaware of the group randomization. The discharge criteria were: mild pain (VAS < 3 at rest) sufficiently controlled by oral analgesics, ability to walk with elbow crutches, ability to eat and drink, and no evidence of any surgical complications. Both groups of patients are involved in same rehabilitation protocol; involving range of motion exercise by using continuous passive motion machine in day one postoperative, then the patient can sit in the bed, get out of bed, and start to walk for many steps by using standard walker, then continue the rehabilitation program.

Statistical analysis

The data analyzed using Statistical Package for Social Sciences (SPSS) version 25.Categorical data presented by frequencies and percentages. Independent t-test (two tailed) was used to compare the continuous variables among study groups accordingly. Paired t-test was used to compare means of ROM scores pre and postoperatively. Pearson's Chi–square test was used to assess statistical association between study group and straight leg raise test. A level of P –value less than 0.05 was considered significant.

Results

As shown in table (1), statistically insignificant differences were seen in means of ROM score between study groups preoperatively, one day, two days, and six weeks after operation (P= 0.141, 0.329, 0.80 respectively). While statistically significant difference was seen in means of this score on discharge from hospital (P=0.001).

 Table 1: Comparison in means of ROM score preoperatively, one day, two days, after discharge, and six weeks postoperatively between study groups

Variable	ROM	P - Value		
variable	Group A Mean ± SD	Group B Mean ± SD	P - value	
Preoperatively	95.75 ± 27.57	105.75 ± 10.03	0.141	
One Day Postoperatively	89.37 ± 11.89	88.50 ± 7.27	0.329	
Two Days Postoperatively	87.18 ± 6.04	82.75 ± 8.18	0.080	
On Discharge	92.50 ± 6.04	78.75 ± 6.25	0.001	
Six Weeks Postoperatively	107.81 ± 7.29	105 ± 5.61	0.215	

Association between LIA and straight leg raise after operation

Table (2) shows the association between LIA injection and ability to performing straight leg raises after knee

arthroplasty. It was clear that there was no statistically significant association between administration of LIA and ability for raising leg straightly after operation (P > 0.069).

Table 2: Association between LIA and straight leg raises following knee arthroplasty

Variable	Study Groups Total (%) n=36		Total	D volue			
variable	Group A (%) n= 16	Group B (%) n= 20	Total	P- value			
Straight leg raise							
Yes	12 (57.9)	9 (42.1)	21 (58.3)	0.069			
No	4 (26.7)	11 (73.3)	15 (41.7)	0.009			

Comparison in means of VAS score between study groups four hours, 12 hours, one day, and two days, following total knee arthroplasty

During rest, we found that means of this score four hours, 12 hours, one day, and two days after operation were significantly decreased in patients managed with LIA (group A) compared to means of patients in group B. (2.70 versus 1.87, P= 0.016; 3.45 versus 2.25, P=0.003; 3.90 versus 3.12,

P=0.026; and 4.50 versus 3.50, P=0.002 respectively). During movement, statistically insignificant differences in means of VAS score between study groups four hours, 12 hours, and two days after operation (P= 0.258, 0.057, and 0.284 respectively). While mean of this score one day postoperatively was decreased in patients of group A compared to that in patients of group B and this difference in means was statistically significant (P=0.016).

Table 3: Comparison in means of VAS score between study groups four hours, 12 hours, one day, and two days, after operation

Variable	VAS Score		
variable	Group A Mean ± SD	Group B Mean ± SD	value
During Rest 4 Hours Postoperatively	1.87 ± 0.95	2.70 ± 0.97	0.016
12 Hours Postoperatively	2.25 ± 1.23	3.45 ± 1.05	0.003
One Day Postoperatively	3.12 ± 1.2	3.90 ± 0.78	0.026
Two Days Postoperatively	3.5 ± 1.09	4.50 ± 0.68	0.002
During Movement 4 Hours Postoperatively	2.75 ± 1.48	3.20 ± 0.83	0.258
12 Hours Postoperatively	3.31 ± 1.30	4.05 ± 0.94	0.057
One Day Postoperatively	3.81 ± 1.16	4.65 ± 0.81	0.016
Two Days Postoperatively	4.65 ± 1.20	4.95 ± 0.75	0.284

Discussion

In group A and group B in the current study, at one and two days postoperatively, means of this score were significantly decreased than pre-operatively (P<0.05). Significant difference was seen in means on discharge from hospital (P=0.001). Also, no significant association between LIA and ability for raising leg straightly after operation observed (P> 0.069). In Moghtadaei *et al.* study in 2014, mean of ROM score at the time of discharge was improved in comparison to three months after surgery (69.5° to 114.4°), in which no significant association found (P>0.05) ^[11].

Also in Carli *et al.* study in 2010, maximal knee extension improved in group of patients managed with LIA (Group I) and other one managed with nerve block (Group F) at six weeks with no difference between them. Maximal knee flexion decreased significantly in Group I (-7.8%) but not significantly in Group F (-6.0%), with no differences between groups regarding absolute values ^[12].

In Chaumeron *et al.* study in 2013 on patients managed with ILA and other with nerve block, the pattern of postoperative knee-assisted flexion between groups significantly differed over days one through five (p = 0.020). Also there was a significant greater knee assisted flexion on the day of surgery in the LIA group (p = 0.001) but none at other follow-ups (*P*>0.05). The pattern of free knee flexion between groups differed significantly over days one through five (p=0.001). Furthermore, the LAI group had increased free knee flexion on the day of surgery (p = 0.042) but none at other follow-ups (*P*>0.05)^[13].

Mullaji *et al.* in 2010, conducted a study of bilateral TKAs in which one knee received LIA and the other knee did not. Patients had greater active flexion up to four weeks and superior quadriceps recovery up to two weeks in the LIA in knee ^[14].

Relief of acute pain after TKA represents a major therapeutic challenge as post-operative pain hinders early mobilization and rehabilitation with subsequent consequences on mobility, duration of hospitalization and overall recovery. It offers the benefits of blocking pain influx at its origin and maximizing muscle control ^[22]. The new rehabilitation protocols emphasizing ROM restoration in out clinic form (outpatients and patients outside the clinic) which bases the discharge criteria on the physical ability not pain control. Furthermore, good perioperative analgesia facilitates rehabilitation, improves patient satisfaction ^[15].

Visual analogue scale (VAS) score results among study groups shows that the means of this score during rest were significantly decreased in group A than in group B in all periods. There is significantly decreased in one day in-group A compared to B (P=0.016).

Andersen *et al.* study in 2010, found that VAS at rest and during mobilization were statistically significantly lower in patients received LIA than those not received it during the whole study period, with the exception of pain scores during mobilization at 24–48 h after surgery (p = 0.05)^[16].

Busch and his colleagues in a study at 2006, found that patients who had received the LIA used significantly less patient-controlled analgesia at six, at twelve, and over the 1sttwenty-four hours after the surgery. In addition, they had higher VAS scale for patient satisfaction and lower VAS scale for pain during activity in the post-anesthetic-care unit and four hours after the operation, concluded that intraoperative LIA with multimodal drugs can significantly reduce the requirements for analgesia and improve patient satisfaction, with no apparent risks, following TKA ^[17].

Furthermore, Vendittoli *et al.* study in 2006 stated that multimodal perioperative analgesia protocol that included LIA, offered improved pain control and minimal side effects to patients undergoing TKA. It also confirmed the safety of the protocol ^[18].

In another study, Toftdahl *et al.* study in 2007, found that peri and intra-articular application of analgesics by infiltration and bolus injections can improve early analgesia and mobilization for patients undergoing TKA, when noticed that patients with LIA could significantly walk < 3 meters on the 1stpostoperative day (p< 0.001). Same patients had significantly lower pain scores during activity and lower consumption of opioids on the 1stpostoperative day ^[19].

In Albrecht *et al.* study in 2016, a reduction in analgesia consumption at 12 hours postoperatively observed and the difference between groups reached statistical significance (P<0.0001). Also they found no significant differences in pain scores at rest (P=0.80) or pain scores on movement (P=0.64) on postoperative day one. Similarly, length of stay was statistically different but without direct clinical relevance. They concluded that LIA provides similar postoperative analgesic efficacy to nerve block after TKA, but requires a higher dose of local anaesthetic and reported incidence of complications does not differ between groups ^[20].

In comparison to other studies, Andersen and colleagues found in their study in 2010, that mean of age of the patients was 67 years (in range of 63–72), a male predominance observed (57%) with Male: Female ratio was 1.3:1 and the mean of BMI level was 28 Kg/m2(in range of 25–32 Kg/m2). The median length of hospital stay was 4 days (Between3–5) days in patients received LIA; however, discharge criteria were significantly earlier in same group 3 days (between 3–3.5) days) than those not received LIA 4 days (between 3–5 days) (p < 0.004)^[15].

Moghtadaei *et al.* study in 2014, observed the mean and SD of the patients age was 64 ± 6.9 years, and the mean and SD of the BMI was 27 ± 2.5 Kg/m2, with male represented the highest proportion, when constituted 72% of them, with male: female ratio was 2.6:1. Concerning number of hospitalization days in those managed with LIA were 5.8 days and those without LIA, they were 5.3 days on average with no significant differences between both groups ^[11]. Finally, Yoon *et al* study in 2010, found that mean of the participant's age was 70 years (ranged from 34-83 years) and a female represented the vast majority of their study, when constituted 94.1% of them, with female to male ratio was 16:1, also mean of BMI was 26.4 Kg/m2(ranged from 19.1-34.2 Kg/m2) ^[21].

Conclusion

LIA is a simple effective method to reduce pain and improve physiotherapy outcome also reduces hospital stay in patients undergoing TKA.

Recommendations

- 1. A larger sample size and longer period of study to decrease the chance of error and bias.
- 2. Long period of follow-up to assess the effectiveness of LIA on long-term functional outcome.
- 3. Prepare and mix the content of LIA by one of surgical team in the field of surgical sets to ensure complete aseptic technique.
- 4. Availability of drugs and anesthetist are mandatory for continuous using this method.

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