

# **Outcome in primary cemented total knee arthroplasty** with or without drain

A prospective comparative study

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# ABSTRACT

**Background:** Suction drain insertion is a common practice in orthopedic surgery, especially after joint arthroplasty to prevent the formation of a hematoma. Theoretically the use of a drain should diminish the volume of hematoma; however the literature has conflicting data. Some authors state that drainage evacuates fluid from a limited area only and can be a cause of infection due to retrograde migration of bacteria. It can also impair the early postoperative rehabilitation. The aim of this study was to evaluate the clinical outcome (especially postoperative pain) and intake of analgesics in patients who had undergone primary cemented total knee arthroplasty (TKA) with or without a postoperative drain.

**Materials and Methods:** A prospective comparative study of 108 consecutive patients (121 knees) was conducted. They were divided into two groups: A study group, with no drainage and a control group with drain inserted at the end of surgery. A total of 121 patients were recruited into two groups. A study group consisted of 59 knees, in which we did not use drainage after TKA and a control group with 62 knees, in which drain was inserted post surgery. Both groups were comparable in terms of preoperative characteristics. The indication for TKA was osteoarthritis (n = 105) and rheumatoid arthritis (n = 16).

**Results:** In patients without drainage we observed lower need for opioids, higher blood loss on the 1<sup>st</sup> postoperative day and a lower need for change of dressings. There were no statistically significant differences in terms of total blood loss, hidden blood loss, transfusion rate, range of motion, length of hospital stay or incidence of complications between the two groups. In 1 year observation there were no differences in clinical outcome between the two groups.

**Conclusions:** The present study conclude that there is no rationale for the use of drain after primary TKA. There are benefits in terms of lower opioid intake, lower blood loss on the first postoperative day and lower need for dressing reinforcement during hospitalization.

Key words: Blood loss, drain, dressing reinforcement, opioids, total knee arthroplasty MeSH terms: Arthroplasty, replacement, knee, blood loss, surgical drainage suction

# INTRODUCTION

The use of drain in joint arthroplasty is a matter of controversy. Suction drains had been traditionally used to prevent the formation of a hematoma

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which, in turn, may decrease joint mobility, reduce local tissue perfusion and increase the likelihood of an infection. However, there is a paucity of available studies supporting these traditional beliefs. Some authors state that drainage evacuates fluid only from a limited area and does not prevent infection if retrograde migration of bacteria occurs.<sup>1,2</sup> It can also impair the early postoperative rehabilitation. In a recent metaanalysis, there is no clear advantage of the use of suction drains, apart from reduced need for change of dressing after total knee arthroplasty (TKA).<sup>3,4</sup> Moreover, in patients who had undergone TKA with drainage, the risk of transfusion was higher.<sup>3</sup>

Majority of authors focus on blood loss and complication rate following TKA with or without drain insertion and there is a paucity of studies evaluating its impact on postoperative pain and intake of analgesics. Thus, we undertook a prospective, comparative study to assess the clinical outcome after primary cemented TKA in patients with or without postoperative drainage.

# MATERIALS AND METHODS

124 elective primary TKA were performed in consecutive 111 patients (of them 13 subjects underwent non simultaneous bilateral surgery). The inclusion criteria were knee arthritis impeding daily activities. Exclusion criteria included significant bone loss that required augmentation, previous thromboembolism and intake of opioids preoperatively. We also excluded three patients for other reasons (a woman suffering from Alzheimer's disease, one man speaking a foreign language and one man who refused to participate in the study). The followup rates at 6 and 12 months were 100% and 96%, respectively.

A total of 121 patients were recruited into two groups. A study group consisted of 59 knees, in which we did not use drainage after TKA and a control group with 62 knees, in which drain was inserted post surgery. Both groups were comparable in terms of preoperative characteristics [Table 1]. The indication for TKA was osteoarthritis in 105 and rheumatoid arthritis in 16 subjects.

The regional ethical committee approved the study protocol and all patients provided informed consent.

# **Operative procedure**

All operations were performed by the senior surgeon (DW). We used two prosthesis systems: Genesis II (Smith and Nephew, Memphis, TN, USA) and Search Evolution (Aesculap, Tuttlingen, Germany). We implanted 100 posterior stabilized prostheses and 21 cruciate retaining (CR) prostheses, all stabilized with cement. Similar numbers of posterior stabilized and CR prostheses were implanted in

Table 1	I: Clinical	details	of the	patients
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Baseline characteristics	Study	Control
	group	group
N (% subjects)	59 (15)	62 (13)
Age, mean (SD)	66.7 (9.8)	69.9 (8.7)
BMI (kg/m <sup>2</sup> ), mean (SD)	29.1 (4.4)	30.6 (4.5)
Duration of symptoms (years), mean (SD)	9.8 (6.3)	9.9 (4.8)
Etiology		
Osteoarthritis/rheumatoid arthritis	52/7	53/9
Type of prosthesis (% PS)		
Genesis II	31 (71)	21 (57)
Search evolution	28 (89)	41 (100)
Preoperative laboratory tests, mean (SD)		
RBC (×10 <sup>6</sup> /µL)	4.5 (0.4)	4.4 (0.4)
Hb (g/dL)	13.2 (1.1)	12.9 (1.2)
HCT (%)	39.5 (2.8)	38.7 (3.6)
Preoperative clinical examination (°), mean (SD)		
Knee flexion	104 (14)	104 (14)
Knee extension deficit	-6 (7)	-6(7)
EBV (L), mean (SD)	4.4 (0.6)	4.5 (0.6)
SD-Standard doviation BMI-Rody mass index BS-Restorior	stabilized BBC	-Dod blood

SD=Standard deviation, BMI=Body mass index, PS=Posterior stabilized, RBC=Red blood cell, Hb=Hemoglobin, HCT=Hematocrit, EBV=Estimated blood volume

both groups. A pneumatic tourniquet was applied to each patient. It was deflated after applying compression dressings at the end of the surgery.

Knee arthrotomy was performed through a midline skin incision and a medial parapatellar capsular incision. Patella was everted. In the control group, drainage was placed intraarticularly. It was removed within first 24 h postoperatively.

Apart from spinal anesthesia (119 patients) and general anesthesia (2 patients), in all patients we injected 0.25% bupivacaine solution with epinephrine intraoperatively in the joint capsule, rectus femoris tendon and the infrapatellar fat pad. Skin was closed with intracutaneous continuous sutures. After surgery all patients were monitored in the recovery room for 24 h, where the pain was controlled with intravenous morphine pump infusion. Analgesics were administered on patient's demand under the supervision of a doctor, starting with nonsteroidal anti inflammatory medication. When it was necessary, opioids were given to provide pain relief.

All patients received low molecular weight heparin, starting 12 h before surgery. They also wore compression stockings from the  $2^{nd}$  postoperative day. Antibiotics (cefuroxime 1.5 g and amikacin 0.5 g) were administered intravenously 30 min before surgery. If the operation had extended, antibiotics were continued up to 3 days. Patients were transfused, if their hemoglobin (Hb) level decreased to less than 7.5 g/dL and symptoms of anemia developed.

In each case proper knee alignment was restored. Rehabilitation protocol was the same in both groups. On the  $1^{st}$  postoperative day patients stood up with a walker, performed active flexion of the operated knee up to 90°. Exercises with continuous passive motion were commenced. From the  $2^{nd}$  postoperative day, patients were allowed to walk on crutches with full weight bearing as tolerable.

#### Assessment

#### **Primary outcome factors**

The primary outcome factors in our study were pain intensity and analgesic intake. Pain intensity was measured with the help of a visual analog scale (VAS). Intake of analgesics per patient were determined. The administered drugs were divided according to their membership in the World Health Organization analgesic ladder.

# **Blood loss and transfusions**

In early postoperative period, we recorded Hb and the hematocrit (HCT) levels (preoperatively, then 8 h, 1 day and 2 days after surgery). We assessed the calculated blood loss (CBL), hidden blood loss (HBL), total measured blood

loss (TMBL), transfusion rates. CBL was calculated using the Gross formula,<sup>5</sup> which uses the maximum reduction in Hb and estimated blood volume, depending on patient's height, weight and gender.<sup>6</sup> TMBL was the sum of the intraoperative blood loss noted by the anesthetists and drainage blood loss. HBL was determined by subtracting the TMBL from the CBL.

# **Dressing reinforcement**

We evaluated requirement for dressing reinforcement and the length of hospital stay. During hospitalization all patients had dressings covering the site of drain exit in the control group and similar expected site on skin in the study group.

# Radiographic examination

All patients were clinically and radiologically evaluated preoperatively, during hospitalization, then at followup, approximately 6 and 12 months after surgery. Radiographs in the anteroposterior and the lateral view were performed using the Knee Society Roentgenographic Evaluation System, any radiological findings suggesting prosthesis loosening at postoperative visits were determined.<sup>7</sup>

Range of movement of operated knee, wound problems and complications were assessed postoperatively and on control visits. A doctor performing followup examination was blinded to information about drain usage in a particular patient.

#### Questionnaires

In addition, patients were assessed with questionnaires, such as Knee Injury and Osteoarthritis Outcome Survey (KOOS) and SF-36 Health Survey version 2 (SF-36 v2) (preoperatively and at followup examinations).

# **Statistics**

Statistical analysis was performed using the Student's *t*-test, Chi-squared test and Mann-Whitney test, depending on the nature of the variables. Results are expressed as mean and standard deviation (SD) or as mean with range. P < 0.05 was considered to be significant.

# RESULTS

# **Primary outcome factors**

On the day of surgery the intake of analgesics, including i.v. morphine and any supplemental opioids was comparable between both the groups. From the 1<sup>st</sup> postoperative day up to discharge, we noted lower demand for opioids in the study group compared with the control group [Figure 1]. Patients in the study group required approximately three times less opioids than patients in the control group (mean analgesia request amounted 0.15 and 0.44/patient, respectively) (P < 0.01).

We observed differences in the mean VAS values between both groups, the biggest was seen on the 1<sup>st</sup> postoperative day, but these results were not significant. VAS values presented a decreasing trend up to discharge [Figure 2].

# **Blood loss and transfusions**

The mean of blood collection in the postoperative drain in the control group was 229 mL (SD 200 mL). In both groups, the values of Hb and HCT decreased during the 1<sup>st</sup> 2 postoperative days (P < 0.01). In addition, on the 1<sup>st</sup> postoperative day, we noted statistically significant average reduction in Hb and HCT levels in the control group compared with the study group (P < 0.05). There were no significant differences between both groups in CBL, HBL and transfusion rates. Evaluation of blood loss is shown in Table 2. CBL in patients who needed a blood transfusion compared with those, who did not need it, was 1542 mL (SD 557 mL) and 1438 mL (SD 384 mL) in the study group, whereas in the control group it was 1542 mL (SD 538 mL) and 1625 mL (SD 420 mL), respectively.

# **Dressing reinforcement**

We recorded a significant difference in dressing reinforcement (P < 0.01). In the study group every patient needed the dressing to be changed at an average 4.5 ± 1.1 times during hospitalization compared with 5.0 ± 1.2 times in the control group. Each patient had at least three dressing reinforcements. We also noted that 7 patients from the control group required regular dressing changes due to prolonged oozing from the wound after drain removal.

# **Range of motion**

There were no significant differences between the groups in the range of motion or length of hospital stay. All

Table	2:	Blood	loss and	l transfusion	requirements	

Characteristic	group	group
Hb decrease (g/dL), mean (SD)		
8 h after surgery	2.6 (0.9)	2.4 (0.8)
Day 1	0.7 (0.6)*	1.1 (0.8)*
Day 2	1.2 (0.9)	1.5 (1.7)
HCT decrease (%), mean (SD)		
8 h after surgery	7.6 (2.1)	7.1 (2.5)
Day 1	2.2 (2.1)*	3.2 (2.4)*
Day 2	4.0 (2.6)	4.4 (5.0)
CBL (mL), mean (SD)	1479 (459)	1580 (484)
HBL (mL), mean (SD)	1444 (476)	1320 (485)
TMBL (mL), mean with range	24 (0-1300)†	259 (0-1560) <sup>†</sup>
Blood unit transfused (autotransfusions)	32 (9)	55 (10)
No. of patients transfused (% of patients transfused)	23 (39)	33 (53)
Total no of transfused blood units per patient, mean with range	1.4 (0-4)	1.7 (0-5)

SD=Standard deviation, Hb=Hemoglobin, HCT=Hematocrit, CBL=Calculated blood loss, HBL=Hidden blood loss, TMBL=Total measured blood loss. \*P<0.05, \*P<0.001

patients achieved full extension of the operated knee at discharge. The knee flexion was comparable between the two groups [Table 3]. Patients from both groups were discharged after 10 days.

#### **Complications**

Complications were observed in six patients from the study group namely; four wound related: prolonged wound healing (n = 3), prolonged healing of injured scar (n = 1) and two general: gastrointestinal hemorrhage and respiratory tract infection. There were 11 complications in the control group. We observed four wound-related problems: Superficial wound infection (1), prolonged wound healing demanding secondary suture (1), persistent leg edema (2) and seven general: Cerebrovascular accident (1), myocardial infarction (1), erysipelas (1), respiratory tract infection (1), and urinary tract infection (3). No knee required any aspiration. There were no other complications 6 and 12 months after surgery. In three patients from the control group, the drain was accidentally removed shortly after its insertion.

#### Questionnaires

The two groups were comparable in terms of preoperative KOOS and SF-36 outcomes, apart from KOOS activity of daily living subscale (ADL) and SF-36 bodily pain score (BP), which were significantly higher in the study group (P < 0.05). Average functional outcome in both groups improved during followup. The results obtained did not differ significantly between the two groups. Changes in questionnaires scores are displayed in Figures 3 and 4.

#### DISCUSSION

Waugh and Stinchfield are the first authors who advocated the use of drains in modern orthopaedics.<sup>8</sup> Most of the recent prospective randomized studies have shown no benefits with its use.<sup>9-14</sup> Nevertheless, according to Chandratreya, 94% of British Orthopedic Association members use drains after TKA in their practice.<sup>15</sup> Much of available reports concentrate on blood loss in the presence of a drain,<sup>10-12,14,16</sup> while very few assess its influence on pain and analgesics requirement.<sup>1,17,18</sup> In this study, we observed significantly higher need for opioids in patients with drain, which is distinct from prior reports. Slightly higher mean pain scores were seen in the control group, especially on

Table 3: Range of motion in operated knee					
Knee flexion (°), mean (SD)	Study group	Control group			
At discharge	86 (6)	84 (10)			
After 6 months	106 (9)	108 (10)			
After 12 months	110 (10)	112 (11)			
SD=Standard deviation					

the 1<sup>st</sup> postoperative day, when the drain was present but the difference was not significant.



Figure 1: Bar diagram showing intake of opioids per patient during hospitalization



Figure 2: Graph showing postoperative pain intensity







**Figure 4:** A line graph showing change over time in SF-36 scores. PF = physical functioning, RP = Role physical, BP = Bodily pain, GH = General health, VT = Vitality, SF = Social functioning, RE = Role emotional, MH = Mental health

To the best of our knowledge, this is the first study to show that patients after TKA who had a drain, need higher doses of opioids. A study done by Holt *et al.* observed no statistically significant difference in postoperative opioid consumption after TKA, regardless of the presence of a drain.<sup>17</sup> Confalonieri *et al.* evaluated patients after unicompartmental knee arthroplasty and noted lower analgesic requirements on the 1<sup>st</sup> postoperative day in patients without drain.<sup>19</sup> However, the authors did not specify what kind of analgesics was applied.

Disruption of continuity of skin and deeper tissues along with the application of the drain causes peripheral sensitization, resulting in a decrease of nociceptors threshold. <sup>20</sup> In addition concentration of local inflammatory mediators increases and secondarily induces central sensitization. This two level action causes pain hypersensivity and persistent decrease in the pain threshold at the site of injured as well as surrounding uninjured tissues. Moreover, opioid receptors are present in inflamed tissues.<sup>21</sup> It is possible that weaker analgesics are insufficient to control this pain. In contrast, opioid analgesics may sometimes cause adverse effects.<sup>22</sup>

Yiannakopoulos and Kanellopoulos emphasized the often neglected fact, that drain tube removal causes pain and discomfort.<sup>23</sup> It is especially important, because proper pain management is considered to be fundamental for minimizing postoperative stress and improving the quality of life and comfort after surgery.

Although some articles suggest that a suction drain does not increase blood loss after TKA,<sup>10,16,18</sup> other authors demonstrated higher blood loss when it was used.<sup>1,3,14</sup> We confirm these results since we noted significant average reduction of Hb and HCT in drained patients on the first postoperative day. The possible explanation of this is that the presence of a drain reduces the tamponade effect.

The decrease of Hb concentration can lead to a higher probability of blood transfusion.<sup>12,14,16,18</sup> Although we observed no significant differences in the overall transfusion rates in both groups, paradoxically in the drained group we have seen lower CBL in transfused patients compared to those who did not necessitate it. This may be associated with the fact that decision regarding transfusion was made not only by the orthopedic surgeon, but also by anesthetist. It seems that the presence of the drain was automatically treated by them as a risk factor for greater blood loss and regardless of indications, resulted in blood transfusion. It was not only our remark. Audet et al. established that blood transfusion after elective orthopedic surgery was reasonable in 68% cases on the day of surgery and in the next days it decreased to 35%.24 The drained blood volume recorded in our analysis was smaller compared with data from other authors.<sup>11,12</sup> It may be a result of techniques to reduce blood loss which we used. These consist of tourniquet use, cementing of the prosthesis, local epinephrine infusion, minimally invasive manipulations on soft tissues.

In our study, we observed that significantly more changes of dressing were made in drained patients. Dora *et al.* had similar findings after total hip arthroplasty.<sup>13</sup> Until date, most authors assessing this issue reported that in the absence of drain the need for dressing reinforcement was higher or at least did not differ significantly.<sup>12,17</sup> Minnema *et al.* revealed that drainage is an independent risk factor for infection after TKA.<sup>2</sup> On the other hand, Ovadia *et al.* evaluated 58 patients following TKA and found significantly higher serous wound discharge when the drain was not used.<sup>16</sup> Nevertheless, it did not affect the complication rate. Our analysis of range of motion, length of hospital stay and other outcomes as assessed with KOOS and SF-36 questionnaires showed no differences between both the groups.

Our study has some limitations. The number of subjects were small but comparable to other studies.<sup>1,17</sup> The difference in opinions on transfusion criteria between orthopedic surgeons and anesthetists might have affected the transfusion rate.

We conclude that there is no rationale for the use of drain after primary TKA. There are benefits in terms of lower opioid intake, lower blood loss on the first postoperative day and lower need for dressing reinforcement during hospitalization.

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