The Influence of Usage of Tourniquet in Total Knee Arthroplasty on Serum Ischemia Modified Albumin Levels

Total Diz Arthroplastisinde Turnike Kullanımının Serum İskemi Modifiye Albümin Seviyeleri Üzerine Etkisi

ABSTRACT

Objective: Ischaemia-modified albumin (IMA) has frequently been studied in recent years in ischaemia-based studies. In this study, IMA changes and postoperative pain were investigated in patients who underwent tourniquet and non-tourniquet knee arthroplasty, and the effect of the tourniquet on pain was investigated quantitatively.

Methods: After obtaining institutional review board approval, 64 patients who underwent total knee arthroplasty were divided into two groups: tourniquet and non-tourniquet using the block randomisation technique. Blood was collected at the 15th minute and 24th hour postoperatively. A visual analogue scale (VAS) was used by patients to evaluate their pain.

Results: According to the results, a significant correlation was found between VAS scores and IMA levels in the blood taken at the postoperative 15th minute (p<0.001). In addition, IMA levels were significantly higher in the tourniquet group (p<0.001).

Conclusion: Less frequent loss of quadriceps muscle strength, less postoperative pain and avoidance of ischaemia-related complications are the main advantages of non-tourniquet knee arthroplasty. IMA is effective in demonstrating these advantages and can be used for problems such as compartment syndrome, which may be accompanied by muscle damage in the extremity.

Keywords: Ischemia modified albumin, knee arthroplasty, tourniquet

ÖZ

Amaç: İskemi modifiye albümin (IMA) son yıllarda iskemi temelli çalışmalarında sıkça incelenmiştir. Bu çalışmada turnikeli ve turnikesiz diz artroplastisi uygulanan hastalarda IMA değişimleri ve postoperatif ağrı incelenmiş olup turnikenin diz arthroplastisinde ağrı üzerine etkisi kantitatif olarak araştırılmıştır.

Yöntemler: Kurumsal Etik kurul onayının alınmasının ardından diz artroplastisi uygulanan 64 hasta blok randomizasyon tekniği ile turnikeli ve turnikesiz olarak iki gruba ayrıldı. Preoperatif, postoperatif 15. dakika ve 24. saatlerde kan alınarak VAS skorlaması kullanılırak ağrı düzeyleri değerlendirildi.

Bulgular: Elde edilen verilere göre postoperatif 15. dakika alınan kanda ve IMA düzeyi ve VAS puanı arasında anlamlı korelasyon tespit edilmiştir (p<0.001). Ayrıca IMA değerleri turnikeli grupta anlamlı olarak yüksek bulunmuştur (p<0.001).

Sonuç: Quadiceps kas gücünün daha az olması, postoperatif ağrıının daha az olması, iskemiye bağlı kompleksiyonlardan kaçınılmış olması turnikeli diz artroplastisi uygulamasının temel avantajlarından biridir. IMA bu avantajların gösterilmesinde etkin olduğuna, ekstremitede kas hasarının eşlik edebildiği kompartman sendromu gibi problemlerde de kullanılabilirliğini göstermektedir.

Anahtar Sözcükler: İskemi modifiye albümin, diz artroplastisi, turniket

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**Introduction**

Knee arthroplasty is a surgical procedure with successful results in over 90% of cases when pain and functional recovery are considered based on cases of gonarthrosis during advanced age (1). The use of a tourniquet in knee arthroplasty has been examined regarding pain, function, muscle strength and prosthesis survival. It is unquestionable that a tourniquet facilitates surgical exposure; however, studies indicate that there is no benefit of tourniquet use other than showing a positive effect in cement adhesion (2-4). It is also thought that tourniquets increase postoperative pain due to ischaemia and nerve compression. However, studies on this subject have not shown a quantitative value that would be a significant ischaemia criterion (2,5,6).

The use of a tourniquet in knee arthroplasty has been associated with increased risk of deep venous thrombosis (7), wound healing impairment (8) and postoperative pain (8,9). A limited number of markers are significant in the literature to investigate the presence and possible consequences of ischaemia. Ischaemia-modified albumin (IMA) is a derivative of albumin. It is known that a reduction occurs in the metal binding capacity of albumin by the modification of the N-terminal end of the albumin caused by hydroxyl ions formed as a result of oxidative stress (10,11). IMA, which is known to be elevated in the event of ischaemia, has been accepted as a quantitative marker for ischaemia and investigated in many pathologies, including cardiac ischaemia, abdominal compartment syndrome and asthma attacks (12).

The aim of this study is to investigate the effect of ischaemia and oxidative stress on pain occurring after knee arthroplasty.

**Method**

Approval was obtained for this prospective, randomised controlled trial from the local Ethics Committee with decision numbers 2017/161. Sixty-four patients who underwent unicompartmental knee arthroplasty were divided into two groups, tourniquet and non-tourniquet, using the block randomisation technique; homogeneity was provided between the groups.

Patients with a history of symptomatic diabetic microangiopathy, cardiovascular disease, peripheral arterial disease and previous extremity surgery within the last three months were excluded from the study.

In all patients, 2 g cefazolin was administered perioperatively, and 4*1 g perioperatively over 24 hours as perioperative antibiotic prophylaxis. In all patients, standard posterior cruciate ligaments cutting posterior-stabilised designs were preferred. All procedures were performed by two different surgeons with at least five years’ experience in knee arthroplasty. Hemovac drains were placed as a standard procedure to prevent hemarthrosis. Five hundred milligrammes of tranexamic acid was injected into the joint in all patients during the postoperative period to reduce blood loss. Blood was collected from the cubital fossa veins before tourniquet application at the 15th minute (min) and 24th hour postoperatively by the same person.

A visual analogue scale (VAS) was used by patients to evaluate their pain (Figure 1).

The blood taken from patients for IMA measurement was placed into serum-separating tubes and centrifuged at 3000 rpm for 10 min using a Sigma centrifuge. Serum samples were placed in Eppendorf tubes for 1 hour and kept at -80 °C until the time of the study. From the dissolved patient serum, 200 gL was placed into the sample tube and 200 gL into the blank tube. Fifty microliters of 1 g/L CoCl₂ (Cobalt-2-chloride)(Merck, K102539.0100, Darmstadt, Germany) was added to both the sample tube and the blank tube. The prepared mixture was mixed with a vortex and allowed to incubate for 10 minutes at room temperature. After 10 min, 50 gL 1.5 g/L DTT (dithiothreitol) (Merck, K111474.0005, Darmstadt, Germany) was added to the sample tube. This was kept at room temperature for two min to observe the colour change. After two min, 1 mL of normal saline (0.9% NaCl) was added to both the sample and blank tubes. Absorbances were measured at 470 nm in a Perkin Elmer Lambda 25 Spectrophotometer. The sample and blank tube results were recorded separately. The results were recorded as Absorbance Units (ABSU). In addition, IMA values of each patient were calculated according to their operation group and albumin. Albumin determination was performed on a Beckman Coulter Au 5800 device using the bromoresol green method. The results are presented as mg/dL.

Corrected IMA (ABSU): IMA x patient’s albumin/Albumin average of the group.

**Statistical Analysis**

The IBM SPSS Statistics Version 21 programme was used for statistical analyses. The values to be examined were subjected to normality analysis using the Shapiro-Wilk test. Student’s t-test (independent sample t-test) was used to compare values between the tourniquet and non-tourniquet groups that were evaluated as parametric (sig. >0.05) in the normality test. The Mann-Whitney U test was used for the groups that were evaluated as nonparametric (sig. <0.05). In addition, for correlations between numeric values, Pearson’s test was used for the groups that were evaluated as parametric in the normality test. Spearman’s test was used for the groups that were evaluated as nonparametric. For the results, assessments of p<0.05 were considered statistically significant. In the correlation evaluations, the power of significance was determined by the value of r.

**Results**

A total of 64 patients, 58 females and six males were included in the study. The mean surgical duration was determined as
Regarding surgical duration, the mean time of the tourniquet group was 80.4±8.7 min, and the non-tourniquet group was 66.8±9.6 min, and this difference was statistically significant (p<0.001) (Table 1).

Patients had significantly higher VAS scores in the tourniquet group than the non-tourniquet group at the 12th postoperative hour (p<0.001) (Table 1). A significant correlation was found between the postoperative 15th min IMA and corrected postoperative 15th min IMA values and 12th hour VAS scores (p<0.001) (Table 3). A significant positive correlation was found between VAS scores and surgical duration (p=0.002) (Table 3).

When IMA levels were measured preoperatively and at the 15th min and 24th hour postoperatively, it was determined that the postoperative 15th min IMA and corrected IMA (13) values were significantly higher in the tourniquet group (Graph 1). However, no significant relationship was found between the two groups in the preoperative and postoperative 24th hour IMA values. There were no significant correlations observed between patient-linked factors, such as body mass index (BMI) and age, between both groups (Table 1).

When the correlation between IMA values and surgical duration was examined, IMA values at the postoperative 15th min and corrected postoperative 15th min were found to be highly correlated with surgical duration (Table 2).

**Discussion**

Regarding the difference in surgical duration between the tourniquet and non-tourniquet groups, this may have been caused by the wrapping of the tourniquet to prevent infection in patients, draining the limb blood with a non-sterile esmarch bandage, the time from starting the tourniquet to inflation, and the time lost during the case preparation. No studies could be found in the literature investigating the differences in surgical duration in patients undergoing knee arthroplasty with and without tourniquets. However, Willis et al. (14) reported that long surgery time and long tourniquet time were associated with an increased infection rate. No infections were detected in our patients, and we believe that these data can be supported with more extensive case studies.

The use of a tourniquet in knee arthroplasty is an application that depends on the surgeon’s preference and is reliable. Refaai et al. found that in 23 knee arthroscopy cases in which a tourniquet was used, IMA was significantly higher compared with preoperative levels in correlation with myoglobin in the material taken after the tourniquet was lowered (15). In our study, IMA and corrected IMA levels in the blood taken at the postoperative 15th min was found to be significantly higher in the tourniquet group. In a prospective study, Wakanhar et al. (16) reported that only postoperative early flexion movements were better in knee arthroplasties performed without tourniquet application. However, there was no long-term difference, and there was no significant difference in blood volume, analgesic requirement, and postoperative pain (16). In a meta-analysis by Zhang et al. (8), non-tourniquet knee arthroplasty was found to be superior regarding thromboembolic events and associated complications. It has been reported that tourniquet use may prevent postoperative exercise, but no significant difference was found in blood loss [8]. Also, the use of tourniquets was reported to improve surgical conditions, surgical vision and reduce blood loss in different studies (5,6,17-19). Although the

**Table 2. Statistical examination of correlation between surgical duration and postoperative IMA levels**

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<tbody>
<tr>
<td>Surgical duration-postoperative 15th min IMA</td>
<td>0.435</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Surgical duration corrected postoperative 15th min IMA</td>
<td>0.298</td>
<td>0.017</td>
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IMA: Ischemia modified albumin

**Table 3. Examination of the correlation between postoperative 15th min IMA, VAS and surgical duration values**

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<tr>
<td>Postoperative 15th min IMA-VAS</td>
<td>0.790</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Corrected postoperative 15th min IMA-VAS</td>
<td>0.782</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>VAS-Surgical duration</td>
<td>0.380</td>
<td>&lt;0.002</td>
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IMA: Ischemia modified albumin, VAS: Visual analog scale

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| Surgical duration (minutes) | 80.4±8.7  
Preoperative IMA | 1.06±0.03  
Postoperative 15th min IMA | 1.25±0.21  
Postoperative 24th hour IMA | 1.08±0.28  
VAS 12th hour | 8.85±0.65  
IMA First change | -0.19±0.20  
Corrected IMA First change | 0.20±0.27  
BMI | 32.7±6.9  
Age | 64.7±7.8  

BMI: Body mass index
use of a tourniquet provides better adhesion of cement, studies also reported that there is no difference (2-4). Estebe et al. (5) stated that the most important complication of a tourniquet is pain caused by mechanical compression or ischaemia-reperfusion injury. However, Spruce reported that tourniquets could be used safely when in compliance with the appropriate conditions of use. In addition, complications would be minimised to ensure the best possible visualisation of the surgical environment, reduction of blood loss, and optimisation of surgical conditions (6). Kumar et al. (19) examined pain in patients undergoing total knee arthroplasty with and without a tourniquet, and the group without a tourniquet reported less early postoperative pain and early functional return. In our study, the early VAS score and IMA values were significantly higher in the tourniquet group, which implies less postoperative early pain in the non-tourniquet group.

In the present study, a significant positive correlation was found between the postoperative 15th min IMA and the corrected postoperative 15th min IMA values and surgical duration. It is reported in the literature that surgical duration is important, especially for pain. The duration of ischaemia and IMA correlations have been examined in abdominal compartment syndrome in the literature, and IMA levels were found to be elevated with increased ischaemic time (12). However, there are no data in the literature on limb ischaemia time and IMA levels.

IMA was found to be higher in obese cases linked to the disruption of adipose tissue oxygenation (20). In our cases, IMA levels taken 24 hours after the end of ischaemia were not significantly different between the two groups. Since there were no significant differences between the two groups regarding BMI and age values, patient characteristics, such as BMI and age, did not have a significant negative effect.

**Study Limitations**

The limitations of our study include the functional results, the short postoperative follow-up period regarding the evaluation of the relationship between stability and IMA, the patients' BMIs being above normal values, and working with two different surgeons. We believe that more extensive case studies can be conducted by increasing the number of patients, and perhaps more accurate results can be obtained. We believe that tourniquet application is an appropriate, accessible, and effective method for the future to examine the early effects of compartment syndrome. The less frequent loss of quadriceps muscle strength, less postoperative pain and avoidance of ischaemia-related complications are the main advantages of non-tourniquet knee arthroplasty. IMA is effective in demonstrating these advantages and can be used for problems such as compartment syndrome, which may be accompanied by muscle damage in the extremity.

**Conclusion**

This prospective, randomised, controlled study evaluated the effects of the use of a tourniquet in knee arthroplasty concerning IMA, which is a quantitative value. This study is unique because it is a controlled study of IMA in orthopaedic surgery. When we evaluate our findings, it is important to remember that the use of a tourniquet in knee arthroplasty procedures is a safe method if the rules are followed. However, postoperative pain on the first day is more common in patients with tourniquets compared with those without tourniquets. In addition, we think that the tourniquet forms a controlled compartment syndrome model. Therefore, IMA is an easy marker to reach and examine during the early stages in patients with suspected compartment syndrome. However, new studies are needed to increase the value of IMA in the diagnosis of compartment syndrome and to increase its use in orthopaedic surgery.

**Ethics**

**Ethics Committee Approval:** Approval was obtained for this prospective, randomised controlled trial from the local Ethics Committee with decision numbers 2017/161.

**Peer-review:** Internally and externally peer reviewed.

**Authorship Contributions**


**Conflict of Interest:** No conflict of interest was declared by the authors.

**Financial Disclosure:** The authors declared that this study received no financial support.

**References**


