Is there a Role of Routine Use of Topical Tranexamic Acid in Hemiarthroplasty?

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ABSTRACT

Background: Blood loss is one of the major perioperative problems in elderly patients undergoing hemiarthroplasty. Such patients, if they require blood transfusion, may be exposed to the complications associated with the same. Intravenous tranexamic acid (TXA) has been used with success in preventing blood loss in total knee arthroplasty and total hip arthroplasty. Concern about thromboembolic complications has led to the topical use of TXA in both TKA and THA. We prospectively evaluated the efficacy of topical TXA for reduction of blood loss in patients undergoing hemiarthroplasty and the related need for blood transfusion.

Materials and methods: Fifty-nine patients aged more than 60 years who presented with a fresh femoral neck fracture and were planned for hemiarthroplasty were randomly divided into two groups. Twenty-nine patients (study group) had local use of TXA during surgery, and 30 patients (control) had only placebo. Postoperative blood loss estimation was compared, which was estimated by hemoglobin (Hb) balance formula based on the postoperative fall of hemoglobin and hematocrit when compared to preoperative values. Complications such as postoperative bleeding, rehabilitation delays, and infection rates were documented.

Results: The Hb decline on the fifth postoperative day in the study group was 1.15 ± 0.40 g/dL in contrast to 1.97 ± 0.60 g/dL in the control group (p value 0.001). The blood loss in the study group was 694.14 mL when compared to 1000.2 mL in the control group patients (p value 0.001).

Conclusion: Despite inherent limitations, the study demonstrated effectiveness of topical use of TXA for decreasing blood loss and reducing need of blood transfusion in hemiarthroplasty patients.

Keywords: Blood loss, Fracture neck of femur, Hemiarthroplasty, Hip fracture, Topical, Tranexamic acid.

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INTRODUCTION

Hip fractures are a serious threat to the quality of life and survival. These fractures with or without treatment are associated with high morbidity and mortality within the first year after injury. The incidence of hip fractures in the age-group more than 35 years is estimated to increase from 1.66 million in 1990 to 6.26 million by 2050.1 According to a 2017 report from the National hip fracture database, the incidence of hip fractures in the United Kingdom is 77,000 per year, and hemiarthroplasty is the commonest surgical procedure for treatment of fracture neck femur in elderly patients.2 Surgery in these frail patients is limited by the possibility of blood loss during surgery and the overall poor physical condition of the patient.

Inspired by the success of intravenous tranexamic acid (TXA) in preventing blood loss, various surgeons have resorted its topical use in total knee arthroplasty (TKA). It has been topically used in a dosage of 1–4 g without any complications.3–5 In TKA, tourniquet is also being used, and there is increase in fibrinolysis on release of the tourniquet, thus the requirement for antifibrinolytics.6 In hip surgery, a tourniquet cannot be used, and the utility of antifibrinolytics may be different. Many surgeons have studied the role of intravenous TXA in total hip arthroplasty (THA) and found it useful. With the aim of minimizing any possible complications such as thromboembolic phenomena, there is an increasing interest in topical use. Topical use has been found effective in THA, but it remains to be seen whether it can be recommended for routine use in hemiarthroplasty (HA). The present study was designed to study the role of topical TXA in patients undergoing HA.

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MATERIALS AND METHODS

The study was a placebo-controlled prospective study on patients undergoing cemented HA of hip joint at a tertiarycare institution after obtaining approval from the institutional ethics committee. Patients with preexisting coagulopathies or those who presented with concomitant inflammatory, pyogenic, or tubercular arthritis of the hip joint and patients with any disease condition that precluded the use of TXA were excluded from the study. Out of a total of 69 patients, 10 were excluded because of the same.

The patients were divided in two groups. The study group (29 patients) were given a total of 3 g of topical TXA diluted with normal saline in two divided doses, 1.5 g made to a total volume of 50 mL poured at the site after incision of deep fascia but just before

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dividing the muscles and was left for 5 minutes, and 1.5 g made similarly poured into the surgical site after implant placement and wound wash and kept for 5 minutes after each pour and before proceeding further. The control group (30 patients) received an equal amount of saline used in the same way. The operating surgeon was unaware whether placebo or TXA had been used, the solution having been prepared under the supervision of the anaesthesiologist in a random manner.

The blood volume was taken to be as 7% of the body weight as reported by Guyton and Hall. The assessment of blood loss was based purely on the fall of hemoglobin (Hb) and hematocrit levels on the first and fifth postoperative days when compared to their levels immediately before surgery.

The blood volume on the fifth postoperative day is expected to have recovered to the preoperative levels, so the difference in the preoperative and the fifth postoperative Hb values was used to calculate Hb loss using the Hb balance formula of Gross. A unit of banked blood was considered to contain 52 g Hb. All patients were mobilized with aid on first postoperative day after obtaining satisfactory X-rays. Appropriate chemoprophylaxis for venous thromboembolism was given in all cases.

Comparison of quantitative variables was done using Student t test and paired t test. For comparing categorical data, Chi-square ($\chi^2$) test was performed.

Results

Out of the 59 patients analyzed, 36 were females and 23 were males. In our study, there was a slight preponderance of female (61%) patients when compared to male patients (39%), with 29 left and 30 right hips.

Most (60.8%) of the fractures occurred due to fall at home. Age range of cases was 60- to 96-year-olds, and 17.4% of the patients were more than 80 years old.

The mean preoperative estimated blood volume of patients in the study group was 4,622.41 mL, while in the control group it was 4,769.33 mL.

The Hb level fell to 9.90 (±1.48) g/dL on the first postoperative day in the study group, while it fell to 9.80 (±1.26) g/dL in the control group when compared to the preoperative Hb levels of 11.07 (±1.69) g/dL and 11.49 (±1.59) g/dL in study and control group, respectively. On the fifth postoperative day, Hb levels were 9.92 (±1.25) g/dL in the study group, while in the control group they were 9.51 (±1.09) g/dL. The decline in the Hb levels in the control group on fifth postoperative day was 1.59 ± 0.40 g/dL, whereas the decline in the Hb levels of the study group was 1.15 ± 0.40 g/dL. The patients who received TXA had a significantly less decline ($p$ value 0.001) in the Hb levels when compared to those who received placebo.

The total Hb Loss on fifth postoperative day was 77.75 g in the study group which was less than that in the control group where it was 117.88 g.

In the study group, the hematocrit fell to 29.90 (±4.07)% on the first postoperative day and 30.01 (±3.66)% on the fifth postoperative day from 32.76 (±7.39)% preoperatively. In the control group, it fell to 30.05 (±4.05)% on the first postoperative day and 30.01 (±3.66)% on the fifth postoperative day from 34.75 (±4.75)% preoperatively. Thus, the fall in the Hct labels in the study group were significantly less than that in the control group.

We observed in the study group that the mean blood loss to 694.14 mL which was less than the mean blood loss of 1,000.27 mL in the control group. The association of decreased blood loss in the study group (topical tranexamic acid use) when compared to the blood loss in the control group was statistically significant ($p$ value = 0.001).

Blood transfusion was given to 17 patients, of which 9 were in the study group and 8 in control group. However, there were some factors that confounded these data (explained below).

Complications observed were a superficial seroma of the wound in four patients (two in each) and low-grade fever in three patients (2 in study and 1 in control group). One patient (control group) had prolonged fever, superficial wound infection and soaking, and had to be taken for wound lavage after 1 month, and the patient recovered subsequently. There was no difference in the complication rate in both the groups, and there was no incidence of any thromboembolic phenomenon or any other systemic side effects related to TXA in either group.

Discussion

Pharmacological agents to reduce the operative blood loss have been a matter of intense study. Out of these, TXA has proved its worth over a period of many years. Initially, it had been used only intravenously in a large number of surgeries such as, cardiac, gynecological, dental, and urogenital procedures.

The use of TXA in orthopedic surgeries has gained prominence over the last few years, and it has now become an essential tool for reducing the blood loss and also the morbidity and mortality associated with blood transfusions. Concerns about systemic side effects such as venous thromboembolism have led to an interest in topical use and determining the optimum dose. A lot of work has been done on the use of TXA in TKA and THA, and indeed surgeons have started using intravenous TXA routinely for these surgeries. However, there is no current recommendation for its routine use in HA patients considering that they are expected to have less blood loss than THA. Patients undergoing HA are elderly and frail and are already at risk of thromboembolism and complications associated with blood transfusions. So, we evaluated topical use of TXA instead of I/V in these patients to address both these issues.

Various authors have compared the preoperative Hb levels with postoperative Hb levels in their studies on different postoperative days to calculate blood loss (as shown in Table 1). In our study, we have used fifth postoperative day Hb levels for comparison with the preoperative ones, considering the Hb level on postoperative day 5 to be a better indicator of actual Hb values; hemodilution due to intravenous fluids is expected to normalize by then, and blood volume is restored back to normal values.

The mean pre-op Hb levels in the study group (11.07 ± 1.69 g/dL) and in the control group (11.49 ± 1.59 g/dL) was different, and this was not significant. Even the fifth postoperative day Hb levels in the study group were 9.92 ± 1.25 g/dL and in the control group were 9.51 ± 1.09 g/dL and were similar statistically. However, the decline in Hb levels in the control group was 1.97 ± 0.60 g/dL, whereas the decline in the Hb levels of study group was 1.15 ± 0.40 g/dL which was significant ($p$ value 0.001).

We found a lower mean blood loss of 694.14 mL ± 270.54 mL in the patients in who TXA was used (study group) when compared to 1,000.27 ± 409.17 mL of blood loss in the patients, where TXA was not used (control group). The difference was statistically significant ($p$ value 0.001). This result has been compared to other studies in Table 2. It seems prudent to conclude that topical use of TXA is efficacious for decreasing blood loss in patients undergoing HA.
There is no consensus among studies in transfusion rates. From application via gauze,\textsuperscript{13} injection of TXA solution in the joint\textsuperscript{14} to application in femoral canal and acetabulum,\textsuperscript{20} the methods of application are varied. Yet, all seem to be effective in reducing blood loss. We used a simple method of soft tissue application and obtained similar results as others in their studies. From application via gauze,\textsuperscript{13} injection of TXA solution in the joint\textsuperscript{14} to application in femoral canal and acetabulum,\textsuperscript{20} the methods of application are varied. Yet, all seem to be effective in reducing blood loss. We used a simple method of soft tissue application and obtained similar results as others in their studies. Some of the authors have used Hb values in g/L rather than g/dL. To enable proper comparison the Hb values have been changed to g/dL wherever necessary in this table.

### Table 1: Comparison of Hb levels in patients who received TXA with controls

| Authors          | Operations | No. of cases study/control group | TXA regime                   | Mean blood loss in mL |
|------------------|------------|----------------------------------|------------------------------|-----------------------|
| Yue et al.\textsuperscript{13} | THA        | n = 52/n = 49                    | 3 g in 150 mL NS             | 945.5 ± 331.7         |
| Gilbody et al.\textsuperscript{18} | THA        | n = 86/n = 88                    | 3 g in 100 mL NS             | 1312 ± 367            |
| Xu et al.\textsuperscript{19}    | THA        | n = 113/n = 111                  | 3 g in 40 mL NS              | 730 ± 296             |
| Emara et al.\textsuperscript{15} | HA         | n = 20/n = 20                    | 1.5 g in 100 mL NS           | 625 ± 35              |
| Kang et al.\textsuperscript{16}  | HA         | n = 40/n = 40                    | 3 g in 100 mL NS             | 600 ± 53              |
| Our study        | HA         | n = 29/n = 30                    | 3 g in 100 mL NS             | 694.14 ± 270.54       |

The regime of topical TXA was different in different previous studies. From application via gauze,\textsuperscript{13} injection of TXA solution in the joint\textsuperscript{14} to application in femoral canal and acetabulum,\textsuperscript{20} the methods of application are varied. Yet, all seem to be effective in reducing blood loss. We used a simple method of soft tissue application and obtained similar results as others in their studies. Some of the authors have used Hb values in g/L rather than g/dL. To enable proper comparison the Hb values have been changed to g/dL wherever necessary in this table.

### Table 2: Comparison of mean blood in patients who received TXA with controls & the TXA regimens

| Authors          | Operations | No. of cases study/control group | TXA regime                   | Mean blood loss in mL |
|------------------|------------|----------------------------------|------------------------------|-----------------------|
| Yue et al.\textsuperscript{13} | THA        | n = 52/n = 49                    | 3 g in 150 mL NS             | 945.5 ± 331.7         |
| Gilbody et al.\textsuperscript{18} | THA        | n = 86/n = 88                    | 3 g in 100 mL NS             | 1312 ± 367            |
| Xu et al.\textsuperscript{19}    | THA        | n = 113/n = 111                  | 3 g in 40 mL NS              | 730 ± 296             |
| Emara et al.\textsuperscript{15} | HA         | n = 20/n = 20                    | 1.5 g in 100 mL NS           | 625 ± 35              |
| Kang et al.\textsuperscript{16}  | HA         | n = 40/n = 40                    | 3 g in 100 mL NS             | 600 ± 53              |
| Our study        | HA         | n = 29/n = 30                    | 3 g in 100 mL NS             | 694.14 ± 270.54       |

The complications observed as discussed earlier were similar in the two groups. Yue et al. in their study found postoperative superficial infections in one case and wound secretions in five patients where topical TXA was used and in seven patients where TXA was not used.\textsuperscript{13} Wei et al. could not find any significant difference in wound complications with topical TXA as previously feared that the increase in surgical time would lead to more complications.\textsuperscript{20}

In our study, we did not find any clinical evidence of thromboembolism, and hence tests like d-dimer and Doppler sonography for DVT did not become necessary. Workers like Emara et al.,\textsuperscript{15} Liu et al.,\textsuperscript{17} Kang et al.,\textsuperscript{16} and Konig et al.\textsuperscript{22} also found no thromboembolic complications in their study. However, Xu et al.\textsuperscript{18} have reported cases of DVT in their study (one each in the study and control group). Alshryda et al.\textsuperscript{13} have reported two cases in each
group in their study of 161 patients. Similarly, Gilbody et al. 18 have reported one case of pulmonary embolism in the TXA group and one case of DVT in the patients who did not receive TXA. Thus, we conclude that topically applied TXA seems safe and at the same time retains the efficacy in lowering blood loss.

**CONCLUSION**

The use of topical TXA in HA of hip joint is both safe and efficacious in reducing perioperative blood loss and need of blood transfusion. We recommend that surgeons routinely use topical TXA for the same regardless of the regimen and method of application.

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