The Effect of Fibrinogen on Blood Loss After Lumbar Surgery: A Double-Blind Randomized Clinical Trial

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Abstract

Background: Spinal surgeries often have a high risk of hemorrhage during and after surgery, thus most patients require blood transfusions and blood products. Fibrinogen is used in different forms to control hemorrhage.

Objectives: The present study aimed to evaluate the outcomes of prophylactic fibrinogen administration in reducing hemorrhage after lumbar surgery.

Methods: This was a randomized clinical trial conducted on 30 patients undergoing lumbar surgery. The levels of fibrinogen, as well as hemoglobin (HB), hematocrit (HCT), prothrombin time (PT), partial thromboplastin time (PTT), and INR, were assessed pre-operatively as the baseline values. The patients were divided into two groups: intervention (N = 15) and control (N = 15) groups. The intervention group received 1 g fibrinogen dissolved in 50 cc distilled water with surgical incision and the control group received 50 cc distilled water with the surgical incision. At the end of the operation, the volume of hemorrhage transfused blood products (fresh frozen plasma, packed cell, and platelet) was measured. In addition, at 0, 6, and 24 hours after the end of surgery and transfer to recovery, serum levels of fibrinogen, HB, HCT, INR, PT, PTT, and hemovac drain volume were measured.

Results: The hemorrhage during and after the operation in the control group was significantly higher than that of the intervention group (P < 0.05). There were no significant differences between hemoglobin and serum level of fibrinogen before and after surgery between the two groups. The postoperative hypotension showed no significant difference between the two groups.

Conclusions: The findings showed the effectiveness of fibrinogen in reducing acute hemorrhage. Considering the adverse consequences of hemorrhage and coagulopathy in patients undergoing surgery, using fibrinogen as prophylaxis is recommended in surgeries with high risks of hemorrhage.

Keywords: Fibrinogen, Hemorrhage, Lumbar Surgery

1. Background

Spinal surgery usually has a high risk of hemorrhage during and after the operation, thus most patients need blood transfusions and blood products (1-3). Hemorrhage during or after surgery is extremely important and this situation is known as a crisis in the operating room throughout the world (3, 4). Reducing hemorrhage is important in maintaining hemodynamic stability of the patient and improving the field of surgery and reducing complications such as hemolytic and non-hemolytic reactions, acute injuries, viral and bacterial infections, hypothermia, and coagulation disorders (4, 5). Previous studies have shown that fluid therapy or infusion of red blood cells is less than the amount of blood loss, leading to dilutions of the blood and a decrease in the concentration of coagulation factors such as fibrinogen (6). Different methods, including induction of controlled hypotension during surgery, injection of various medications during surgery, or change in the mechanical ventilation with a positive pressure during anesthesia have been proposed to reduce hemorrhage during and after the surgery, with positive outcomes in many cases (7-10). In addition, the results of studies on the efficacy of prophylactic fibrinogen injections have been reported to reduce the hemorrhage during surgery (11-17). Fibrinogen binds to platelet glycoproteins, enhances platelet aggregation and improves cross-linking, resulting in stabilization and clotting of the clot (18-20). Therefore, the administration of fibrinogen to restore the blood concentration level to normalize clot formation in patients who have hemorrhage is important (21-23). Allergic reactions, fever, thrombotic events e.g. cardiac infarction and
pulmonary embolism are complications of intravenous fibrinogen infusion. Fibrinogen is used in various forms such as the injection method to control the hemorrhage. In the injection method, the drug is absorbed throughout the body and may not be sufficiently damaged to the affected tissue, and a large amount of this product should be prescribed for proper hemorrhage. Considering the importance of hemorrhage during and after lumbar surgery, the role of prophylactic administration of fibrinogen in this field has not been considered.

2. Objectives

The purpose of the study was to evaluate the results of fibrinogen prophylactic infusion in reducing the hemorrhage of lumbar surgery.

3. Methods

All of the procedures of this study were approved by the local Ethics Committee of Ahvaz Jundishapur University of Medical Sciences, Ahvaz, Iran (ethics code: IR. AJUMS.REC.1396.1136). This study was designed as a double-blind, placebo-controlled, clinical trial (IRCTID: IRCT2018081104076N1). The study population consisted of 30 patients (23), aged 18 - 60 years old, with one or two levels of discopathy, candidates of lumbar disc surgery with an ASA class I-II anesthetic risk admitted in Ahvaz Golestan Hospital (September 2017 - October 2018). They were randomized to receive either fibrinogen concentrate or placebo. Following the clear explanation of the object and potential risks and benefits of the study, written informed consent was obtained from all participants. Patients in the intervention group (N = 15) received 1 g of fibrinogen dissolved in 50 cc distilled water along with surgical incision. The patients in the control group (N = 15) were injected with 50 cc distilled water as placebo along with surgical incision. The exclusion criteria were: the patients with cardiovascular problems, coagulation disorders, anticoagulant drugs, smoking, HCT, BMI > 30. The plasma levels of fibrinogen, as well as HB, HCT, INR, PT, PTT, and hemovac drain volume, were again measured at 0, 6, and 24 hours. To describe the data, the mean and standard deviation (SD) in quantitative variables and in qualitative variables frequency and percentage were used. To compare the results, t-test and chi-square test were used. All analyses were performed using the statistical package of SPSS (version 22) and the significance level for all statistical analyses was considered 0.05 (P ≤ 0.05).

4. Results

During the study period from September 2017 to October 2018, 120 patients undergoing elective lumbar disc surgery were eligible to participate in the trial (CONSORT Diagram, Figure 1). After initial screening, 60 patients agreed to participate and provided written informed consent. Among them, 30 patients did not meet the inclusion criteria. Finally, 30 patients were selected and divided into the two groups (intervention and control). There were no significant differences between them in terms of demographic characteristics (P > 0.05) (Table 1).

Although the hemoglobin level and serum fibrinogen levels were not significantly different between both groups pre- and post-operatively (P > 0.05) (Table 2), the fibrinogen level was significantly different between the intervention and control groups pre- and post-operatively (P < 0.001) (Table 3).

Intraoperative hemorrhage in the intervention group was significantly lower than the control group (225.33 ± 180.431 versus 530.00 ± 275.032, P < 0.001). In addition, postoperative bleeding in the intervention group was significantly lower than the control group (43.33 ± 41.690 versus 107.33 ± 85.228, P < 0.001) (Table 4).

The need for blood transfusion and blood products in the intervention group were significantly lower than the control group (intraoperative use unit of PC: 0.07 ± 0.258 versus 1.00 ± 1.195). However, intraoperative use unit of FFP
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Assessed for eligibility (n = 120)

Excluded (n = 60)

Denied from research (n = 60)

Analysed (n = 15)

Excluded from analysis (give reasons) (n = 0)

Lost to follow-up (give reasons) (n = 0)

Discontinued intervention (give reasons) (n = 0)

Allocated to control (n = 15)

→ Received allocated placebo (15)

Allocated to intervention (n = 15)

→ Received allocated intervention (n = 15)

Lost to follow-up (give reasons) (n = 0)

Discontinued intervention (give reasons) (n = 0)

Analysed (n = 15)

→ Excluded from analysis (give reasons) (n = 0)

Analytical Characteristics

Table 1. Demographic Characteristics of the Intervention and Control Groups

| Variable                | Intervention | Control | P Value |
|-------------------------|--------------|---------|---------|
| Age, mean ± SD          | 45.00 ± 8.418| 47.46 ± 13.289| 0.421   |
| Gender, male/female, No. (%) | 9/6 (60/40) | 8/7 (53.3/46.7) | 0.520   |
| BMI, kg/m²               | 22.57 ± 3.2  | 22.64 ± 3.29 | 0.910   |

Table 2. Comparison of Hemoglobin and Serum Fibrinogen Pre- and Post-Operatively Between the Two Groups

| Variable                | Intervention, Mean ± SD | Control, Mean ± SD | P Value |
|-------------------------|-------------------------|-------------------|---------|
| Pre-operative hemoglobin, mg/dL | 12.64 ± 1.8361 | 13.36 ± 2.1148 | 0.35    |
| Post-operative hemoglobin, mg/dL | 11.55 ± 2.0836 | 12.33 ± 1.7775 | 0.35    |
| Pre-operative serum fibrinogen, mg/dL | 293.27 ± 72.448 | 290.20 ± 86.257 | 0.383   |
| Post-operative serum fibrinogen, mg/dL | 346.67 ± 55.845 | 320.27 ± 65.952 | 0.383   |

showed no significant difference between the groups: 0.00 ± 0.000 versus 0.00 ± .000 (Table 4).

There were no significant differences between PT, PTT, and HCT levels in the two groups at preoperative, 0, 6 and 24 hours postoperatively. There was a significant difference in the rate of INR in the two groups at the preoperative
(P < 0.029), 6 (P < 0.017) and 24 (P < 0.04) hours postoperatively, but there was no significant difference at 0 time point (Table 5).

5. Discussion

This study was a double-blind, randomized controlled trial that was designed to determine the effect of the prophylactic fibrinogen injection on reducing the hemorrhage in the lumbar discopathy surgery. The results showed that the administration of prophylactic fibrinogen reduced the need for blood supply during and after surgery significantly. Most studies in the past, similar to the present study, confirmed the effect of intraoperative fibrinogen use on reducing hemorrhage during and after surgery (24). Several studies have shown that fibrinogen plasma concentration during surgery determines the volume of the hemorrhage and the need for blood transfusion (25). In the study of Ucar et al., on the patients who were candidates for coronary artery bypass graft surgery, the level of fibrinogen during surgery significantly correlated with the amount of hemorrhage 48 hours after surgery (25). Similarly, Blome et al. reported a strong direct relationship between fibrinogen concentration and 24-hour hemorrhage volume in patients with cardiac arrest (26). Our findings showed that the rate of hemorrhage in both groups was decreased after the operation. In a clinical trial conducted by Pournajafian et al., they examined the reduction in the hemorrhage during spinal surgery and found no significant difference between serum hemoglobin and fibrinogen level before and after surgery (25). In the study of Karlsson et al., the rate of hemorrhage in the fibrinogen-receiving group was 32% lower than that of the control group (13). Several studies have shown that fibrinogen plasma concentration during surgery is a strong direct relationship between fibrinogen concentration and hemorrhage 48 hours after surgery.

### Table 3. Comparison of Serum Fibrinogen Pre- and Post-Operatively in Both Groups

| Variable                        | Intervention, Mean ± SD | Control, Mean ± SD | P Value |
|---------------------------------|-------------------------|--------------------|---------|
| Pre-operative serum fibrinogen, mg/dl | 293.27 ± 72.448        | 290.20 ± 86.257    |         |
| Post operative serum fibrinogen, mg/dl | 346.67 ± 55.845        | 320.27 ± 65.952    |         |

### Table 4. Comparison of Hemorrhage and Blood Products Transfusion Intra- and Postoperatively Between the Two Groups

| Variable                        | Intervention, Mean ± SD | Control, Mean ± SD | P Value |
|---------------------------------|-------------------------|--------------------|---------|
| Intraoperative hemorrhage, cc    | 225.33 ± 10.431         | 530.00 ± 275.032   | < 0.001 |
| postoperative hemorrhage, cc     | 43.33 ± 41.690          | 107.33 ± 85.228    | < 0.001 |
| PC transfusion, unit             | 0.07 ± 0.258            | 1.00 ± 1.995       | < 0.001 |
| FFP transfusion, unit            | 0.00 ± 0.000            | 0.00 ± 0.000       | > 0.05  |

*The postoperative hypotension was not significantly different between the two groups (P = 0.083).

### Table 5. Comparison of Blood Factors Before and After the Intervention Between the Two Groups

| Variable | Intervention, Mean ± SD | Control, Mean ± SD | P Value |
|----------|-------------------------|--------------------|---------|
| PT       | Before 29.747 ± 3.2500 | 28.400 ± 3.5617   | 0.355   |
|          | 0 29.667 ± 3.310       | 29.31 ± 4.0652    | 0.562   |
|          | 6 29.867 ± 3.2566      | 28.291 ± 3.0392   | 0.348   |
|          | 24 29.931 ± 2.875      | 28.692 ± 2.8978   | 0.341   |
|          | Before 12.567 ± 0.4619 | 12.433 ± 0.759    | 0.325   |
|          | 0 12.400 ± 0.2070      | 12.692 ± 0.8191   | 0.6     |
|          | 6 12.400 ± 0.2803      | 12.385 ± 0.2096   | 0.87    |
|          | 24 12.333 ± 0.2440     | 12.423 ± 0.2774   | 0.397   |
| HCT      | Before 37.660 ± 4.300  | 39.429 ± 4.2941   | 0.348   |
|          | 0 34.953 ± 5.6140      | 33.300 ± 10.510   | 1       |
|          | 6 34.800 ± 5.3542      | 35.843 ± 3.9656   | 0.445   |
|          | 24 34.980 ± 5.3079     | 35.76 ± 2.4838    | 0.565   |
| INR      | Before 1.007 ± 0.0258  | 1.060 ± 0.0828    | 0.292   |
|          | 0 1.003 ± 0.0352       | 1.069 ± 0.1251    | 0.399   |
|          | 6 1.007 ± 0.0258       | 1.069 ± 0.1099    | 0.017   |
|          | 24 1.007 ± 0.0258      | 1.062 ± 0.0961    | 0.047   |
which was in line with the results of the present study about the requirement for PRBCs (27). Carling et al. studied a patient undergoing scoliosis in a clinical trial and found that hemorrhage in the patient had a significant correlation with fibrinogen plasma concentration but did not show significant correlation with platelet count, PT, and PTT (28). Wafaisade et al. investigated a large number of trauma patients (N = 294) and reported that the 6-hour mortality rate in the fibrinogen-receiving group was lower than the control group, and not only an increased time to death but also an increased rate of multiple organ failure was reported. A reduction of overall hospital mortality was not observed in patients receiving FC, but contrary to our findings, the need for blood and blood product administration between the intervention and control groups indicated no significant differences. This difference can be as a result of the effect of the number and variety of patients and also the effect of various factors such as the severity of lesions during trauma and the volume of hemorrhage (29). In a clinical trial performed on children with craniosynostosis and scoliosis, Haas et al. saw that receiving intracerebroventricular concentrate can reduce hemorrhage during craniosynostosis; however, does not affect scoliosis, which is consistent with the results of the present study (30). In a cohort study, Geck et al. argued that preoperative fibrinogen level had a negative logarithmic correlation with the total blood loss, and there was no significant relationship between thrombin and thromboplastin with preoperative plasma fibrinogen, which was consistent with our study (31). The limitations of the current study are the difference in the duration of surgery and the lack of follow-up of long-term complications of surgery in patients. To reduce the amount of the hemorrhage during and after surgery in spinal surgery, several measures should be taken in terms of surgical and anesthetic techniques, the use of anti-fibrinolytic drugs, such as tranexamic acid and coagulation compounds such as FFP, cryoprecipitate, and fibrinogen.

5.1. Conclusions

Our findings indicated that fibrinogen plays a key role in reducing hemorrhage and the need for blood products in patients undergoing acute spinal surgery. Therefore, this compound could be used as an important factor in preventing and stopping the hemorrhage. Considering the adverse effects of hemorrhage and coagulopathy in the patients undergoing surgery, the use of this treatment prophylactically in all surgeries with a high risk of hemorrhage is recommended. From the viewpoint of this study, it is better to consider these findings in the planning of hospital managers and policymakers to allocate funds for providing sufficient fibrinogen for the hospital operating rooms.

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Footnotes

Authors’ Contribution: Fatemeh Javaherforoosh Zadeh conceived the study, Mohsen Shafaee Tonekaboni searched the literature and collected the data, Farahzad Janatmakan and Mansoor Soltanzadeh drafted the manuscript.

Clinical Trial Registration Code: This study was designed as a double-blind, placebo-controlled, clinical trial (code: IRCTID: IRCT2018081040763N1).

Conflict of Interests: There is no conflict of interest to be declared.

Ethical Considerations: The corresponding author affirmed that this manuscript was resulted from honest, accurate, and transparent findings of the study and no important aspects of the study were omitted. In addition, any discrepancies from the study as planned (and so registered) were explained. The reporting method of this work was compliant with CONSORT guidelines. All of the procedures of this study were approved by the local Ethics Committee of Ahvaz Jundishapur University of Medical Sciences, Ahvaz, Iran (ethics code: IR.AJUMS.REC.1396.1136).

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Patient Consent: Written consent form was obtained from all participants.

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