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EFFICACY OF TRANEXAMIC ACID IN REDUCING THE BLOOD LOSS DURING CESAREAN SECTION IN PRIMIGRAVIDA PATIENTS WITH BREECH PRESENTATION

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ABSTRACT:

OBJECTIVES:

The objective of the study was to evaluate the efficacy of tranexamic acid in decreasing blood loss at the caesarian section (CS).

METHODOLOGY:

A descriptive case series study was conducted in the Department of Obstetrics and Gynecology, Lady Reading Hospital, Peshawar, Pakistan. Data were collected from September 24, 2018, to March 24, 2019. A total of 114 term women (18-35 years age) with singleton term breech cases were analyzed for a period of six months

RESULTS:

In this study the mean age was 30 years with SD±2.341 and the mean period of gestation was 38 weeks with SD±4.76. Moreover, tranexamic acid was effective in 91% of patients and was not effective in 9% of patients.

CONCLUSION:

Our study concludes that the efficacy of tranexamic acid was 91% in reducing the blood loss during CS in primigravida patients with breech presentation.

KEYWORDS: Tranexamic Acid, Cesarean Section, Primigravida, Breech Presentation

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INTRODUCTION:

Planned CS was found to be safer for singleton term breech babies than planned vaginal birth, managed according to a clinical protocol, but was related to more complications for mothers¹. Breech presentation is delineated as a fetus in a longitudinal line with the buttocks or feet closest to

the cervix. Breech deliveries have been reported in 3-4% of all types of deliveries. Its frequency decreases with advancing gestational age from 22-25% of births prior to 28 weeks gestation to 7-15% of births at 32 weeks gestation to 3-4% of births at term². Cesarean section (CS) mentions the delivery of a fetus, placenta, and membranes via an abdominal and uterine incision³. CS can give rise to more complications than normal vaginal delivery⁴. Postpartum hemorrhage (PPH) is one of the most common complications of CS, which is accountable for 20% of maternal deaths worldwide⁵. In the developed world rates of CS have increased by 23.8% to 50%, whereas in developing countries, this rate is falling to >10%, and in Pakistan the rate of CS is 25%⁶. According to the World Health Organization PPH is described as "blood loss from the birth canal in excess of 500

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mL during the first 24 hour after delivery". Clinically, it is related to sweating, tachycardia, and weakness, with hemodynamic collapse, which takes place at losses of between 35 and 45% of blood volume. The use of pro-hemostatic drugs such as Tranexamic acid may go a long way in the prevention of PPH in SC. Tranexamic acid (TA) is mainly an antifibrinolytic agent that has been indicated to decrease blood loss and transfusion requirements in different elective surgeries⁷. It is a synthetic derivative of the amino acid lysine that puts in its anti-fibrinolytic effect via the reversible barricade of the lysine binding sites on plasminogen molecules⁸. TA also prevents the transformation of plasminogen to plasmin by plasminogen activator⁸. It has been found to be very useful for lowering blood loss and necessary for blood transfusion⁹. In Pakistan, limited literature is available on the effects of TA in primigravida patients with breech presentation. The objective of this study is to assess the efficacy of prophylactic injection of tranexamic acid in the reduction of blood loss during CS in primigravida patients with breech presentation.

METHODOLOGY:

A descriptive case series study was carried out among 114 patients (18-35 years of age) with singleton pregnancy from September 2018 to March 2019 at the Department of Obstetrics and Gynecology, Lady Reading Hospital, Peshawar, Pakistan. The protocol was approved by the ethical committee of the Department of Obstetrics and Gynecology, Lady Reading Hospital, Peshawar. Informed consent was taken before data collection. Sample Size was calculated by using the following formula:

$$\frac{\mathbf{z}^2 \mathbf{pq}}{\mathbf{d}^2}$$

Efficacy=92%

q=1-p and d=5% with 95% confidence interval level. Inclusion criteria were women aged 18-35 years age, singleton pregnancy and primigravida, gestational age >36 weeks on last menstrual period (LMP), breech presentation as per operational definition. Patients who had allergy to tranexamic acid, thrombolic disorders, abnormal placenta (placenta previa, placental abruption), diabetes, hypertension, were excluded from the study. In the present study 114 patients from the indoor Department of Obstetrics and Gynaecology of Lady Reading Hospital, Peshawar have included the study. After informed consent, basic

demographics were noted and all women were given TA 10 minutes before giving incision injection. TA 1gm were diluted with 20ml 5% dextrose and it infused IV slowly over 5 minutes. After delivery of baby oxytocin, 20 units IV Drip were given. Efficacy was recorded as per operational definition i-e estimated blood loss <500 ml following placental delivery till the end of the surgery. Blood was measured by using the soaked gauzes, pads (measured by subtracting preuse weight from post-use), and blood clots, which were weighed by standardizing one milliliter of blood to one gram and noted on specially designed proforma. Data were analyzed with IBM SPSS (version. 22). Frequency and percentage were computed for qualitative variables like age groups and efficacy. Mean±SD were presented for quantitative variables like age, gestational age, duration of the procedure, weight, height, and BMI. Effect modifiers like age, gestational age, duration of the procedure, and BMI were controlled by stratification. Post-stratification Chisquare test was applied and p ≤0.05 was considered statistically significant.

RESULTS:

Table 1: Demographic Features of Patients

Characteristics	N (%)	Mean±SD	
Age (114)			
18-25 Years	65 (57%)		
26-35 Years	49 (43%)	30±10.341	
BMI			
<25 Kg/m2	52 (46%)	215.11	
>25 Kg/m2	62 (54%)	2±5.11	
Gestational Period	82		
37-39 Weeks	(72%)	29 4.76	
40-41 Weeks	32 (28%)		
Duration			
<45 Minutes	71 (62%)	45.7.12	
>45 Minutes	43 (38%)	45±7.12	

Table 2: Efficacy of TA among Patients

Efficacy	N (%)
Effective	104 (91)
Not Effective	10 (9)
Total	114 (100)

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Table 3: Stratification of Efficacy w.r.t Age, Gestational Age, BMI and Duration of Procedure

Stratification of Efficacy	Efficacy		P- Value
w.r.t Age Distribution	Effective	Not Effective	
18-25 Years	59 (52%)	45 (39%)	0.8419
26-35 Years	6 (5%)	4 (4%)	
Stratification	Efficacy		
of Efficacy w.r.t Gestational Age	Effective	Not Effective	0.8869
37-39 Weeks	75 (66%)	29 (25%)	
40-41 Weeks	7 (6%)	3 (3%)	
Stratification	Efficacy		
of Efficacy w.r.t BMI	Effective	Not Effective	
<25 Kg/m2	48 (42%)	56 (49%)	0.7090
>25 Kg/m2	4 (4%)	6 (6%)	
Stratification	Efficacy		
of Efficacy w.r.t Duration of Procedure	Effective	Not Effective	0.5979
<30 Minutes	64 (56%)	40 (35%)	
>30 Minutes	7 (6%)	3 (3%)	

DISCUSSION:

TA is a competitive inhibitor of plasminogen activation and can decrease blood loss by preventing the breakdown of fibringen and fibrin clots¹⁰. TA has been extensively used to cure heavy menstrual bleeding and to decrease blood loss in elective surgery where it decreases blood transfusion by about one-third 11,12. For many years it has been used for the cure of several types of bleeding e.g. menorrhagia, postoperatively, or intra-operatively 12,13. WHO recommended TA to decrease death due to bleeding in women with PPH regardless of cause, and with no adverse maternal effects¹⁴. In the present study, 114 patients were investigated. In this study the mean age of patients was 30 years with SD±2.341 and the mean POG was 38 weeks with SD±4.76. Moreover, the outcome clearly depicted that the blood loss was decreased by the usage of TA before CS. Similar results have been reported in a study that was carried out on 100 women undergoing lower segment cesarean section (LSCS). TA significantly reduced the quantity of blood loss from the end of LSCS to 2 hours postpartum which was 86.5 ml in the study group versus 142.70 ml in the control group $(<0.001)^{15}$. Similarly, in the Indian population, Bhatia et al. (2015) and Singh et al. (2014) also investigated and revealed a reduction in blood loss by use of TA 16,17, 30% reduction in blood loss reported by Ahmed et al. in Ismailia Egypt¹⁸, and 34% reported by Maged in Cairo Egypt¹⁹. Similarly, in other similar studies carried out in India, Turkey, and

Iran it is reported reduction was between 39.1 to 43.7%²⁰⁻²². In another study conducted by Shahid et al. (2013) TA significantly reduced the amount of blood loss during the lower segment cesarean section (LSCS), but it did not decrease the blood loss significantly after the CS²³. These differences in findings of blood loss might be due to the different time intervals employed in evaluating the blood loss and the different approaches used in the valuation of blood loss in different studies.

CONCLUSION:

This study demonstrated that pre-operative intravenous tranexamic acid significantly reduced blood loss (91%) during CS in primigravida patients with breech presentation, but it did not significantly reduce the blood loss after the cesarean section.

LIMITATIONS:

Beside the strength of this study there are also some limitations of the study. First sample size was smaller. Data should be collected from a larger number in order to make results more authentic which will have a far reaching effect in future perspective.

CONFLICT OF INTEREST: None

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