Does prophylactic use of tranexamic acid reduce PPH from cesarean delivery when coupled with uterotonics?

Yes, there is a statistical effect, but the clinical implications are limited. In a randomized controlled trial involving 4,551 patients undergoing cesarean delivery (CD), women who received tranexamic acid in combination with prophylactic uterotonics had lower rates of postpartum hemorrhage (PPH) than women who received uterotonics and placebo. Clinically relevant hemorrhage-related outcomes, such as blood transfusions and need for additional medications or procedures to control bleeding, did not differ between groups.


EXPERT COMMENTARY
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Postpartum hemorrhage is the leading cause of maternal mortality worldwide.1 Many preventive strategies, including tranexamic acid administration, have been studied in an attempt to reduce the risk of PPH. Tranexamic acid prevents the conversion of plasminogen to plasmin, preventing the breakdown of fibrin, and ultimately stabilizing the fibrin matrix of clot.2 It has been shown to be an effective approach to treating hemorrhage in patients after trauma as well as cardiac surgery.3,4 The use of tranexamic acid in obstetric hemorrhage has reduced mortality in previous trials,5 but its prophylactic use has had mixed results in preventing obstetric hemorrhage.6-8

Recently, Sentilhes and colleagues published the largest prospective study to date addressing the efficacy of tranexamic acid for the primary prevention of PPH.

Details of the study
Multiple hospitals throughout France participated in the investigators’ double-blind randomized, placebo-controlled trial. Women undergoing CD at 34 or more weeks’ gestation (N = 4,551) were randomly assigned to receive 1 g of intravenous (IV) tranexamic acid or placebo after cord clamping. Both groups received IV prophylactic uterotonics.

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The primary outcome was PPH, defined by estimated blood loss (EBL) greater than 1 L or receipt of red blood cell transfusion within the first 2 days after surgery.

Results. The rate of PPH was significantly lower in women who received tranexamic acid compared with those who received placebo. Yet, the mean EBL between the 2 groups differed by only 100 mL. The rates of blood transfusions, additional uterotonic administration, arterial embolization, and hysterectomy did not differ between groups.

The clinicians responsible for the care of these patients did not observe a difference in the rate of “clinically significant” PPH between those who received tranexamic acid and those who received placebo. Women who received tranexamic acid were more likely to experience nausea and vomiting, but they did not have any increased risk of venous thromboembolic disease.

Study strengths and limitations
Sentilhes and colleagues’ study findings contradict those of an earlier meta-analysis on the topic. This may be due to the effect of publication bias on meta-analyses, which makes them prone to supporting the findings of published positive trials while missing data from negative trials that did not reach
publication. The gold standard for addressing a research question such as this is a randomized controlled trial (RCT). The study reviewed here is an excellent example of a well-designed and executed RCT.

There may be a benefit to prophylactic tranexamic acid in certain populations not well captured among these study participants. The inclusion criteria were broad, including both prelabor and intrapartum CDs, making the results generalizable. However, the population studied, with a mean body mass index of 26 kg/m² and age of 33, may not resemble some readers’ patient population. Prespecified subgroup analyses did not find a benefit to tranexamic acid in patients considered at high risk for PPH or in those undergoing intrapartum CD.

References