Efficacy of Tranexamic Acid in Craniosynostosis Repair in an Infant

Goran Rakić¹  Danica Stanić¹  Anna Uram-Benka¹  Marina Pandurov¹  Jovana Simin¹  Biljana Drašković¹

¹Institute for the Healthcare of Children and Youth of Vojvodina, Novi Sad, Serbia

Address for correspondence Dr. Marina Pandurov, MD, Institute for the Healthcare of Children and Youth of Vojvodina, Hajduk Veljkova 10, Novi Sad 21000, Serbia (e-mail: pandurovmarina@gmail.com).

Introduction

Craniosynostosis (craniostenosis) is a condition where one or more cranial sutures fuse prematurely leading to focal or global growth delay of the skull.¹ Surgical treatment involves expansion and remodeling of the cranium in early infancy to prevent increased intracranial pressure, cerebral compression, and blindness.²

Because of the risk of massive bleeding, prolonged anesthesia in small children, the possibility of a difficult airway, raised intracranial pressure, venous air embolism, hypothermia, endotracheal tube displacement, positional injuries, corneal abrasions, head and neck edema, and the need of an invasive monitoring, craniosynostosis surgery represents a great challenge for the anaesthetist.²,³

Extensive blood loss is common in pediatric craniosynostosis reconstruction surgery and so are transfusion-related morbidity and mortality.²,⁴ Tranexamic acid is an antifibrinolytic agent, increasingly used in children to reduce perioperative blood loss in various settings, including during craniosynostosis surgery.

Case Report

An 8-month-old, 9-kg male infant was admitted to our hospital for craniosynostosis repair. The operation was performed under a balanced general anesthesia. Two central lines and one peripheral line were cannulated in case of a need of massive transfusion. Invasive monitoring was used, as well as prevention of hypothermia. As massive blood loss was expected, before beginning the surgery bolus of tranexamic acid as well as packed red blood cells was administered. During the operation, tranexamic acid was given continuously in an intravenous infusion. The child was hemodynamically stable throughout the operation. After the completion of surgery, which lasted for 5 hours, the patient was extubated in the operating room. Postoperatively, the patient was admitted to the intensive care unit, where he stayed for 24 hours. Hemoglobin values were stable, and there was no need for additional blood replacement.

Conclusion

Extensive blood loss is common in pediatric craniosynostosis reconstruction surgery, transfusion being unavoidable in the majority of cases. In our patient, tranexamic acid proved effective in reducing perioperative blood loss and transfusion requirement.
craniosynostosis, without intracranial hypertension or neurological deficiencies. Computed tomographic scan showed absence of large and small fontanelle, as well as a complete parietal synostosis.

The patient was premedicated with 100 µg of intramuscular atropine and 200 mg of rectal paracetamol half hour before induction. The induction was inhalational with sevoflurane and oxygen. The patient was intubated with 3.5 sized cuffed endotracheal tube. The Cormack–Lehane view was grade 1. Right internal jugular vein was cannulated with 22 gauge cannula and right femoral vein with 22 gauge cannula, guided by ultrasound. Invasive arterial blood pressure was monitored via a right dorsalis pedis artery, cannulated with 24 gauge cannula. Peripheral venous access included a 22 gauge cannula in left upper limb. The anesthesia was maintained with oxygen, air, sevoflurane, cisatracurium, and continuous infusion of remifentanyl. The patient was positioned very carefully in the prone position and all intravenous fluids were warmed, as well as the operating room and the patient himself, to prevent hypothermia. There was no possibility to use a precordial Doppler probe to notice venous air embolus, so we had to rely on capnography.

Because massive blood loss was expected, bolus of 400 mg (50 mg/kg) of tranexamic acid was administered before beginning the surgery. During the operation tranexamic acid was given continuously at a rate of 10 mg/kg/h. It was difficult to estimate the amount of bleeding; but, since the hemoglobin value was borderline (110 g/L) before the surgery, the patient received 15 mL/kg of packed red blood cells before surgical incision. The child was hemodynamically stable throughout the operation. After the completion of surgery, which lasted for 5 hours, the patient was extubated in the operating room. Postoperatively, the patient was admitted to the intensive care unit, where he stayed for 24 hours. Hemoglobin values were stable and there was no need for additional blood replacement.

**Discussion**

Craniosynostosis surgery is often complicated by massive perioperative bleeding, which can lead to severe hypotension, metabolic acidosis, coagulopathy, acute lung injury, postoperative ventilation, and even cardiac arrest and death.²,⁸ The relative amount of blood volume lost is greater in younger children. This is because the head has a proportionally greater percentage of blood volume and surface area, and therefore, it is prone to more bleeding.⁴,⁹ Complications associated with blood transfusion may occur to a greater degree and with a greater frequency in infants than in older children and adults, due to higher oxygen consumption and a higher cardiac output to blood volume ratio.¹⁰

It is important to know at which moment during the operation, one can expect an extensive blood loss to predict and prepare for hemorrhage. Our patient entered the operating room with borderline values of hemoglobin concentration. As significant hemorrhage occurs during initial scalp dissection and the raising of the periosteum,² we decided for preemptive administration of packed red blood cells (15 mL/kg) and a loading dose of tranexamic acid (50 mg/kg). Since gradual but significant blood loss can occur throughout the whole procedure and extends into the postoperative period, we decided to give tranexamic acid as an intravenous infusion of 10 mg/kg/h during surgery.

It has been proven that tranexamic acid markedly reduces perioperative blood loss and transfusion requirements in children; however, there is significant variability in dosing regimens. Optimal dosage remains uncertain, with wide variations in reported loading doses (10–100 mg/kg) and infusion rates (1–10 mg/kg/h).²,⁷

It is difficult to assess the amount of blood volume lost intraoperatively due to child’s young age and the fact that a small amount of blood is collected by the suction. Most of the blood lost is absorbed by surgical drapes and the surrounding area.⁴ Because of this, we used hemodynamic parameters, blood gas analysis as well as hemoglobin and hematocrit values as transfusion triggers, which is in accordance with literature findings.⁴ In this case, there was no need for further blood transfusion intra- or postoperatively.

Adverse events have been described in the literature, including seizures as well as thrombotic events.² No adverse events were recorded in our patient.

**Conclusion**

Extensive blood loss is common in pediatric craniosynostosis reconstruction surgery, transfusion being unavoidable in the majority of cases. In our patient, tranexamic acid proved effective in reducing perioperative blood loss and transfusion requirement.

**References**


