We report anesthetic management of a 7-day old, 3-kg male infant admitted to our hospital with complaints of poor feeding, lethargy (Glasgow coma scale [GCS] E1V1M5), and seizure. The baby was born via normal vaginal delivery and was apparently healthy at birth. There was no significant antenatal history or history of any drug intake during pregnancy. Noncontrast computed tomography (NCCT) of the head at admission revealed a large fronto-temporo-parietal subdural hematoma (SDH) in the left side, warranting urgent surgical intervention (►Fig. 1). Preoperative international normalized ratio (INR) was 1.03 and the platelet count was 1,06,000/mm³. In operating room, the patient was received with tracheal tube of 3.5-mm ID in situ and a respiratory rate of 30 breaths/min, heart rate (HR) of 150 beats/min, and blood pressure (BP) of 80/40 mm Hg. Anesthetic induction was done with fentanyl, oxygen, sevoflurane, and rocuronium. Right internal jugular vein and right femoral artery were cannulated for intravenous access and invasive BP monitoring, respectively. Intraoperative monitoring included 5-lead electrocardiogram (ECG), invasive BP, central venous pressure, pulse oximetry, temperature, urine output, and arterial blood gas (ABG) analysis. Fentanyl 1.5 µg/h and cisatracurium 0.3 mg/h with sevoflurane were used for maintenance of anesthesia. Normocarbia and normothermia were targeted. Intraoperative ABG analysis showed normal gas exchange with hemoglobin (Hb) of 5 g/dL and blood glucose of 158 mg/dL. There were two episodes of intraoperative hypotension, which were managed with blood or fluid bolus and mephentermine 0.3 mg. A total of 180 mL of crystalloid was given intraoperatively. Blood loss was estimated to be 200 mL and was replaced with equal amount packed red blood cells (RBCs), platelets, and fresh frozen plasma (FFP) (15 mL/kg). Toward the end of surgery (decompressive craniotomy), BP dropped again (40/28 mm Hg) and an infusion of noradrenaline 1 µg/min was started. The patient was shifted to the intensive care unit (ICU) for elective ventilation with HR 160 beats/min, BP 80/46 mm Hg, and temperature 35.2°C on noradrenaline infusion. Removal of SDH alone could not ensure adequate recovery due to complex multisystem effect of neurological injury. The patient was kept sedated and mechanically ventilated in postoperative period. Monitoring in ICU included 5-lead ECG, invasive BP, central venous pressure, pulse oximetry, temperature, urine output, and ABG analysis, along with monitoring of neurological status and postoperative hematological and biochemical investigations. The patient’s BP dropped further, and dopamine infusion was started as well. Investigations revealed coagulopathy with deranged prothrombin time, INR of 3, and platelet count of 62,000/mm³. Both platelets and FFP were transfused. Postoperative CT revealed removal of hematoma with evolving infarct. Despite all the steps taken to maintain hemodynamics and correct coagulopathy, the patient could not make it and succumbed to his injuries on first postoperative day.

Incidence of SDH in newborns is reported to be as high as 48%; however, SDH requiring surgical decompression is
Letter to the Editor

Meticulous attention to fluid management, blood loss, temperature control, and analgesia is essential during neonatal neurosurgery. Surgery involving massive blood loss such as in our case results in replacement of entire blood volume and leads to hypothermia and coagulopathy, which may adversely affect the outcome apart from the injury itself. Appropriate anesthesia and analgesics are necessary even in neonates. We used volatile agent along with opioid to maintain anesthesia. Total intravenous anesthesia with propofol is better in tight brain, but there is a little evidence for its use in neonates. The use of propofol in neonates is off-label. The Food and Drug Administration (FDA) has approved it for maintenance of anesthesia only in children ≥ 2 years of age. Strict cutoff values regarding physiological parameters in neonates are limited. Hypotension and cerebral perfusion pressure limits in neonates are not very well defined. Though the new guidelines for management of severe pediatric traumatic brain injury (TBI) suggests a minimum cerebral perfusion pressure of 40 mm Hg, there may be age-specific thresholds with infants at the lower end and adolescents at or above the upper end of this range. Similarly, there are limited recommendations regarding choice of hyperosmolar therapy in neonates. Hypertonic saline is recommended for treatment of raised intracranial pressure in children with traumatic brain injury. Although mannitol is a widely used agent for this purpose, no studies meeting inclusion criteria were identified for use as evident from recent guidelines for pediatric TBI guidelines. Massive blood transfusion, tissue hypoxia due to hypotension, severe brain injury, and coagulopathy resulted in fatal outcome, but since only few neonates present for such emergencies, regular auditing and reporting are required to strengthen the evidence for management of such cases.

Massive SDH in a neonate for emergency neurosurgery is challenging, and it becomes more difficult when the cause is not certainly known. Strict vigilance and anticipation of complications, especially coagulation abnormalities in large SDH without any apparent history, should be kept in mind. Ruling out all probable causes of SDH, proper screening for neonatal coagulopathies, and prompt management are essential for favorable outcome.

Conflict of Interest
None declared.

References
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