Anesthetic management of spontaneous cervical epidural hematoma during pregnancy: a case report

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Abstract

Background: Spontaneous spinal epidural hematoma during pregnancy is a quite rare event requiring emergent decompressive surgery in the majority of cases to prevent permanent neurological damage. Therefore, there is little data in the literature regarding anesthetic management of cervical localization during pregnancy. The potential for difficult airway management with the patient under general anesthesia is one of the major concerns that needs to be addressed to prevent further cord compression. Anesthetic management should also include measures to maintain the mean arterial pressure to improve spinal cord perfusion. Furthermore, spine surgery in pregnant patients needs special consideration in terms of positioning and in the postoperative period.

Case presentation: We present a case of a 35-year-old white woman at 21 weeks of gestation with a spontaneous cervical epidural hematoma. Fiberoptic bronchoscope-guided nasal intubation was a safe option to ensure a higher rate of successful endotracheal intubation while minimizing the risk of aggravating the injury. Her care posed other multiples challenges that required a multidisciplinary team approach.

Conclusions: The case of our patient serves as a reminder of this rare condition and its implications regarding anesthesia.

Keywords: Spontaneous cervical epidural hematoma, Pregnancy, Anesthesia, Fiberoptic bronchoscope intubation

Background

Spontaneous cervical epidural hematoma (SCEH) during pregnancy is a rare condition; only ten cases have been reported in the literature to date. The aim of this case report was to present an anesthetic approach in surgery performed in a pregnant woman at 21 weeks of gestation with an SCEH.

Case presentation

A 35-year-old white woman (weight 75 kg, height 160 cm), gravida 3 para 2, presented to our hospital at 21 weeks of gestation with paraplegia. She had no significant medical or anesthetic history; in particular, she had no history of trauma, and she had not been taking aspirin or anticoagulants. She had been receiving regular care from an obstetrician during pregnancy, which had progressed normally. Twenty-four hours before her admission, the patient had developed a sudden onset of severe interscapular pain radiating to her neck with sudden weakness and paresthesia in both the upper and lower limbs. The patient was confined to bed, and she did not experience any improvement, so she decided to consult in our hospital.

Her physical examination revealed flaccid paralysis in both legs. Her lower limb power bilaterally was grade 0/5. In her right and left higher limbs, power was grades 2/5 and 3/5, respectively. Her upper level with sensitivity was C7–T1. All her deep tendon reflexes were absent. Her plantar response was bilaterally mute. She also had urinary sphincter disturbances. The rest of her physical examination was unremarkable.

The results of hematological investigations were within normal ranges: serum potassium 4 mEq/L, hemoglobin 14 g/dl, platelets 180,000/mm³, international normalized ratio 1.2, and prothrombin time 11 seconds. Obstetrical
ultrasound revealed a normal pregnancy at 21 weeks of
gestation and a fetal heart rate (HR) of 140 beats per mi-
minute. Magnetic resonance imaging of the cervical and
dorsal spine revealed a cervical spine epidural hematoma
extending from levels C3 to C6 with severe spinal cord
compression (Fig. 1).

An anesthesiologist, a neurosurgeon, and an obstetri-
cian urgently reviewed the case. After discussion with
the patient, it was decided to perform spinal cord de-
compression with the patient under general anesthesia.
After explanation of the anesthesia plan, the patient ver-
balized understanding and consented to undergo awake
fiberoptic intubation. Her American Society of Anesthesi-
ologists Physical Status classification was I, her Mallampati
classification was I, and she had a good mouth opening.
On arrival in our operating room (3 hours from admis-
sion), noninvasive blood pressure (BP) monitoring with
three-lead electrocardiography and pulse oximetry was
established. The patient’s basal values were BP 120/60
mmHg, HR 75 beats per minute and peripheral capillary
oxygen saturation of 100% while breathing room air. Intra-
venous access was secured with two 18-gauge intravenous
cannulae, and the patient was given 10 ml/kg of isotonic
saline solution. Cefazolin 2 g was also administered
intravenously. After 5 minutes of preoxygenation with
100% oxygen and nebulization of lidocaine 2%, we car-
rried out a superior laryngeal block, which was followed
by nasal fiberoptic intubation. Tracheal intubation was
easily performed with a 7.0-mm armored tube. Once
the patient’s end-tidal carbon dioxide level confirmed
placement, general anesthesia was induced with propofol
2.5 mg/kg, fentanyl 4 μg/kg, and rocuronium 0.6 mg/kg.
Immediately after intubation, the patient’s BP was 130/70
mmHg, and her HR was 85 beat per minute. She was
ventilated with a target tidal volume of 450 ml and a
respiratory rate of 13 breaths per minute. Sevoflurane
(1–1.5%) in a mixture of oxygen and air (1:2) was used
for maintenance of anesthesia. A radial arterial line was
placed for continuous monitoring of arterial BP and
collection of arterial blood for blood gas analysis during
surgery. We also inserted a urinary catheter.

The patient was turned to the prone position to allow
a posterior cervical approach to the hematoma with
appropriate padding and careful checking for pressure
points. The endotracheal tube was rechecked for correct
position by chest auscultation. The patient’s airway pres-
sure was 22 cmH₂O. Intraoperative fetal monitoring was
not possible with the patient in this position. The
patient’s mean BP was maintained between 85 and 95
mmHg. Perioperatively, blood gas analysis showed pH
7.4, partial pressure of carbon dioxide 36 mmHg, partial
pressure of oxygen 135 mmHg, and bicarbonate 22.3
mmol/L. A laminectomy with evacuation of the epidural
hematoma and decompression of the thecal sac was
performed, followed by surgical hemostasis (Fig. 2).
The operative findings did not reveal any arteriovenous

![Fig. 1](https://example.com/fig1.png)

**Fig. 1** T2-weighted sagittal magnetic resonance image of the cervical spine showing an epidural hematoma located dorsal to the cord extending from C3 to C6 levels, causing obliteration of the spinal subarachnoid space and pushing the cord anteriorly against the vertebral bodies

![Fig. 2](https://example.com/fig2.png)

**Fig. 2** Operative finding: a hematoma on the dorsal aspect of the cervical spinal cord
malformation. The patient’s hemodynamic parameters remained stable during the procedure, and no vasopressors were required. Estimated blood loss was 350 ml, and the patient received a total of 1000 ml of isotonic crystalloid with adequate urine output.

At the end of the surgery, anesthetic gases were discontinued, and the patient was awakened. Immediately following surgery, a neurological examination of the patient was performed to evaluate her motor and sensory functions, which were identical to the preoperative findings. Postoperative analgesia consisted of intravenous paracetamol 1 g every 8 hours and intravenous nefopam 100 mg per 24 hours. Fetal ultrasound revealed a HR of 150 beats per minute. After discussion with the obstetrician, there was no indication for tocolysis. The patient was continuously monitored for another 24 hours in the intensive care unit, and the postoperative period remained uneventful. She was transferred to the rehabilitation department where she underwent physical therapy. On postoperative day 2, she had slight sensation in both legs, and her muscle power showed a gradual improvement in the upper limbs (3/5 and 4/5, respectively, in the right and left upper limbs). After 1 month, power had improved to grade 4 in her left leg and to grade 3 in her right leg, and she was able to move her arms freely.

Discussion
Spontaneous spinal epidural hematoma (SSEH) is a very uncommon medical condition in obstetrical patients [1, 2]. It is spontaneous in the absence of trauma, arteriovenous malformation, blood disorders, and epidural puncture [1, 3]. It has been suggested that raised intra-abdominal pressure forces venous return from the pelvis and abdomen to the vertebral venous plexus. Rupture of these veins within the negative pressure compartment of the epidural space explains the hemorrhage [1, 3, 4]. It has also been suggested that structural vessel changes induced by estrogen and progesterone excess [5], along with various anatomical factors [6-8], may favor SSEH occurrence.

SSEH is most often located in the thoracic region because of the rich epidural plexus present in this region. Only ten cases of pregnant women with SCEH have been described in the literature to date (Table 1). The SSEH is usually posterior or posterolateral to the thecal sac [9-12]. Urgent surgical decompression and clot removal is the gold standard [13, 14], but some patients may be managed conservatively [15-18]. Incomplete preoperative sensorimotor loss and surgery within 36 hours of onset correlate with a favorable outcome [19].

The anesthetic management of SCEH in pregnancy is not well described. General anesthesia is necessary for cervical localizations, and the difficulty concerns airway management of these patients. Masski et al. reported a case of SCEH in a pregnant woman who was under general anesthesia using a rapid sequence technique with cricoid pressure [20]. The patient’s head was maintained.

| Author [reference] | Age, years | GW | Time from ND to decompression | Symptoms | Level | Timing of decompression related to delivery | Surgical positioning | Outcome (recovery) |
|--------------------|------------|----|-----------------------------|----------|-------|---------------------------------------------|----------------------|-------------------|
| Mahieu et al. [1]  | 26         | 30 | 3 hours                     | Neck pain Brown-Séquard syndrome | C3–T1  | After CS                                    | ?                    | Complete          |
| Singh et al. [2]   | 25         | 31 | 30 hours                    | Interscapular pain Quadriplegia | C3–C7  | Before VD                                   | ?                    | Complete          |
| Yonekawa et al. [6] | 20        | 37 | 14 hours                    | Neck pain Paraplegia            | C4–C6  | Before VD                                   | Sitting position     | No recovery        |
| Wang et al. [13]   | 29         | 40 | 6 hours                     | Neck pain Quadriplegia          | C5–C7  | After CS                                    | Prone position       | Mild (impaired sensation in fingers) |
| Iwatsuki et al. [15]| 27        | 37 | Spontaneously resolved      | Left shoulder pain              | C4–T1  | Abstention                                  | Not done             | Spontaneous recovery |
| Masski et al. [20] | 27         | 41 | 12 hours                    | Neck and arm pain Quadriplegia  | C7–T2  | After CS                                    | Prone position       | No recovery        |
| Binnert et al. [28]| 28         | 37 | 120 hours                   | Interscapular pain Paraplegia   | C7–T1  | After VD                                    | Sitting position     | No recovery        |
| Matsubara et al. [29] | 36    | 16 | 9 hours                     | Interscapular pain Paraplegia   | C3–C7  | Before CS                                   | ?                    | No recovery        |
| Tada et al. [30]   | 26         | 31 | 30 hours                    | Paraparesis Quadripareisis      | C4–T2  | After CS                                    | Prone position       | Complete           |
| Our patient        | 35         | 21 | 28 hours                    | Neck and interscapular pain Quadriplegia | C3–C6  | Before CS                                   | Prone position       | Mild               |

Abbreviations: CS Cesarean section, GW Gestational weeks, ND Neurological deficit, VD Vaginal delivery
in neutral position without a collar by a third person. Manual in-line stabilization (MILS) was used during direct laryngoscopy and intubation to minimize the risk of further cord compression. However, the MILS technique does not eliminate cervical spine movement [21], and it is known that MILS reduces mouth opening and leads to a poor laryngoscopic view [22].

In our patient, we used a different approach. We thought that conventional endotracheal intubation with direct laryngoscopy could have aggravated the injury and exposed the patient to a marked sympathetic response. Awake fiberoptic bronchoscopy was deemed most appropriate.

The management of acute spinal cord decompression should also include measures to preserve spinal cord perfusion [23] and adequate uterine blood flow [24] by ensuring volume expansion, using anesthetics that respect the hemodynamic status and administering vasopressors if needed (Table 2). Invasive BP monitoring with an indwelling arterial catheter is useful. Cardiac dysfunction secondary to loss of cardiac sympathetic tone is another reason for using invasive monitoring.

Lateral positioning was considered for our patient, but the neurosurgeons expressed their view that this position would have made the surgery technically difficult with an increase of the surgery duration and bleeding. Prone position is limited by the difficulty in monitoring the fetal HR and uterine activity. According to the American Congress of Obstetricians and Gynecologists, the decision whether to use intraoperative fetal monitoring should be determined by a multidisciplinary team and based on each patient’s unique circumstances and the surgery to be performed [25]. After a multidisciplinary team discussion, the prone position was selected for our patient to allow optimal surgical access. Careful positioning for laminectomy was done with help and guidance of the neurosurgical team. For this position, an armored tracheal tube is necessary.

Finally, at the end of the surgery, airway edema is one of the major concerns in patients undergoing surgery in prone position [26, 27], combined with effects of the gravid state and parenteral fluid administration. The precaution of direct laryngoscopy prior to extubation is justified. During the postoperative period, patients need expensive health care. Postoperative documentation of motor and sensory function is essential for these patients, and they should be monitored for any hemodynamic alterations in a high-dependency unit.

Conclusions
This case report highlights the management of an uncommon medical condition in obstetrical patients. The potential for difficult airway management with the patient under general anesthesia is one of the major concerns. Anesthetic management should also include measures to maintain the mean arterial pressure to improve spinal cord perfusion, as well as special considerations regarding positioning and postoperative management.

Abbreviations
BP: Blood pressure; CS: Cesarean section; GW: Gestational weeks; HR: Heart rate; MAC: Minimum alveolar concentration; MILS: Manual in-line stabilization; ND: Neurological deficit; SCEH: Spontaneous cervical epidural hematoma; SSEH: Spontaneous spinal epidural hematoma; VD: Vaginal delivery

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Authors’ contributions
MS and AE drafted the manuscript under the supervision of MB. All authors participated in patient care. All authors critically revised the manuscript, and all authors read and approved the final manuscript.

Competing interests
The authors declare that they have no competing interests.

Consent for publication
Written informed consent was obtained from the patient for publication of this case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

Table 2 Anesthetic management of published cases of spontaneous spinal epidural hematoma

| Author(s) | Localization | Induction of anesthesia | Intubation | Maintenance of anesthesia | Hemodynamic support | Outcome |
|-----------|--------------|-------------------------|------------|--------------------------|-------------------|---------|
| Jo et al. [25] | T1–T5 | Propofol 100 mg and rocuronium 50 mg | Direct laryngoscopy | Sevoflurane (1–1.5%) in 60% oxygen and air | Dopamine 5–10 μg/kg/minute (5 days) | Complete recovery |
| Doblar and Schumacher [4] | T6–T9 | Etomidate 12 mg and succinylcholine 120 mg | Direct laryngoscopy | Isoflurane at 0.25–0.5 MAC in oxygen and nitrous oxide | Phenylephrine infusion (7 days) | Mild recovery |
| Masski et al. [20] | C7–T2 | Thiopental 5 mg/kg and rocuronium 0.8 mg/kg | MILS + direct laryngoscopy | Not precise | Not used | No recovery |
| Our patient | C3–C6 | nebulization of lidocaine 2% and superior laryngeal block propofol 2.5 mg/kg, fentanyl 4 μg/kg, and rocuronium 0.6 mg/kg | Awake fiberoptic intubation | Sevoflurane (1–1.5%) in 50% oxygen and air | Not used | Mild recovery |

MAC Minimum alveolar concentration, MILS Manual in-line stabilization
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