Case Report
Pituitary Apoplexy and Subdural Hematoma after Caesarean Section

Van Trung Hoang,1 The Huan Hoang,1 Thanh Tam Thi Nguyen,2 Vichit Chansomphou,3 and Duc Thanh Hoang4,5

1Department of Radiology, Thien Hanh Hospital, Buon Ma Thuot, Vietnam
2Department of Radiology, FV Hospital, Ho Chi Minh City, Vietnam
3Department of Radiology, Savannakhet Medical-Diagnostic Center, Kaysone Phomvihane, Laos
4Department of Endocrinology, Diabetes and Metabolism, Walter Reed National Military Medical Center, Bethesda, USA
5Department of Medicine, Uniformed Services University of the Health Sciences, Bethesda, USA

Correspondence should be addressed to Van Trung Hoang; dr.hoangtrungradiology@gmail.com, Vichit Chansomphou; dr.vichit1991@gmail.com, and Duc Thanh Hoang; tdhthanh@gmail.com

Received 10 May 2022; Revised 7 June 2022; Accepted 14 June 2022; Published 23 June 2022

Copyright © 2022 Van Trung Hoang et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Pituitary apoplexy can occur postpartum, and subdural hematoma following epidural anesthesia is a rare complication. Cooccurrence of these two complications is extremely rare and has not been previously reported in the literature. In this article, we present a case of pituitary apoplexy along with intracranial subdural hematoma happening two days after spinal anesthesia for cesarean section. The patient presented with peripheral facial nerve paralysis accompanied by headache, eye pain, and blurred vision and was diagnosed by imaging modalities. The patient made a good recovery with conservative treatment without serious health events.

1. Introduction
Epidural anesthesia and spinal anesthesia to assist in obstetric procedures have become the current standard techniques with proven safety and efficacy. However, anesthesia and caesarean section may cause some certain complications such as intracranial bleeding that can be potentially lethal. Few cases of intracranial subdural hematoma following the administration of epidural anesthesia have been reported in the literature [1–5]. Very few cases of pituitary apoplexy have been reported in postpartum patients [6–10]. To our knowledge, the cooccurrence of pituitary apoplexy and subdural hematoma after spinal anesthesia for cesarean section has not been reported before. We report a rare case where multiple intracranial complications occurred simultaneously after a cesarean section using spinal anesthesia with the presence of symptoms of peripheral facial paralysis. The patient was treated conservatively and recovered well thereafter.

2. Case Presentation
A 34-year-old primigravida at 38 weeks gestation starting labor was hospitalized. She had had a benign tumor in the left ovarian which had been operated on 2 years ago. Cardiotocography showed category II, and clinical examination manifested cephalopelvic disproportion, so cesarean section was indicated. The baby was delivered through cesarean section by the Pfannenstiel line under spinal anesthesia. After the surgery, the patient recovered within normal limits. After two days of surgery, she showed signs of right facial paralysis. Her chief complaints were headaches, mild eye pain, and blurred vision. She was normotensive with a blood pressure of 110/60 mmHg, pulse rate 100 bpm, respiratory rate 21 per minute, and temperature 37°C. Physical examination revealed mild right-sided facial and mouth distortion (drooping). Otherwise, no other focal neurological signs were found. Laboratory results performed before and after
cesarean section in this case including a complete blood count, coagulation, blood glucose, liver and kidney function, TSH, and free T4 and T3 were within normal ranges. She underwent a head computed tomography (CT) scan showing mildly hyperattenuating pituitary gland and thin layer subdural hematoma in the left frontal region (Figure 1). A subsequent head magnetic resonance imaging (MRI) revealed pituitary apoplexy and left frontal subdural hematoma (Figure 2). The patient was treated conservatively with the usual supportive measures such as vital sign monitoring, airway support, intravenous fluids, and nutritional support. She did not need hormone imbalance or surgery. Her symptoms were relieved, and she was discharged after 7 days. She continued doing well at one-year follow-up. Laboratory tests and imaging exams to check pituitary function did not reveal any abnormalities.

3. Discussion

Epidural anesthesia and spinal anesthesia are frequently used in obstetric procedures; however, they can cause complications such as back pain, headache, hemorrhage, infection, nerve injury, and intracranial hematoma. Intracranial subdural hematoma is a rare complication but can lead to life threatening [11]. The clinical appearance of a subdural hematoma includes persistent headache, vomiting, drowsiness, disorientation, blurring of vision, and other neurological symptoms [12]. Pituitary apoplexy is defined as acute nonhemorrhagic or hemorrhagic infarction of the pituitary gland. The risk of pituitary hemorrhage and infarction is increased during pregnancy and the postpartum period. Pituitary apoplexy occurring after spinal anesthesia for cesarean sections is an extremely rare acute clinical condition that presents with symptoms including sudden headache, nausea, vomiting, visual disturbances, meningeal signs, and altered consciousness [13, 14]. However, the symptoms are the combined results of subdural hematoma and pituitary apoplexy as such they may overlap each other [1, 2, 6, 10].

These situations can be explained by several pathogenesis. First, for subdural hematoma, the primary mechanism leading to it after spinal anesthesia is intracranial hypotension that causes a caudal shift of the brain. This leads to traction of broken dural vessels (such as a small cerebral cortical vein, the bridging dural vein, or dural venous sinus wall) that result in blood extravasation and the formation of subdural hematoma [2, 15].

Second, for pituitary apoplexy, a mechanism may be explained by the increased volume of the pituitary gland in pregnant women due to lactotroph cells undergoing massive hyperplasia. Estrogen receptors are expressed in the lactotroph cells, and estrogen levels become very high during pregnancy. Even though lactotroph cells become larger, both in the pituitary gland and the pituitary adenoma, the blood supply remains limited leading to pituitary stroke [10, 13, 16].

Another possible mechanism for pituitary apoplexy in this circumstance can be the subacute, excessive growth of the preexisting adenoma, which outgrows its blood supply with eventual ischemic necrosis followed by hemorrhage. Some causes of pregnancy-related pituitary apoplexy include intracranial hypertension, diabetes, dynamic pituitary testing, bromocriptine, and anticoagulants. Identified risk factors are tumor factors including histological type, nonfunctioning tumor or prolactinoma, and size (macroadenoma), along with patient factors such as pregnancy, systemic hypertension, dopamine agonist administration, dynamic pituitary function tests, and anticoagulant agents. Indeed, pathologic and dynamic imaging studies have shown that macroadenomas, as well as microadenomas, are less vascularized than the pituitary gland so that relatively fast and sizable growth can exceed this low blood supply. However, this theory does not explain the onset of pituitary apoplexy in patients with small adenomas or with a healthy pituitary [6, 13, 14, 16].

Another hypothesis is that tumor compression of the infundibulum and superior pituitary arteries may cause infarction of the normal pituitary gland; however, ischemia of the tumor mass itself is less probable in this case because the vessels supplying the adenoma are attributable to the inferior pituitary circulation. Therefore, pituitary tumors probably suffer from an intrinsic vasculopathy that can lead to spontaneous infarction and hemorrhage [13, 16].

Sheehan syndrome is a typical obstetric-related inclined factor and a rare cause of pituitary apoplexy as well as panhypopituitarism. It occurs only in women who suffered a severe postpartum hemorrhage with the critical hypovolemic shock resulting in ischemic pituitary necrosis. Sheehan syndrome should be suspected in the case of persistent hypotension and tachycardia after severe obstetric hemorrhage treatment. Other premature signs of Sheehan syndrome are related to hypopituitarism, such as hypoglycemia and breastfeeding difficulty or inability [17].

Typical imaging features of intracranial subdural hematoma are generally changed in the cases happening after spinal anesthesia for cesarean section from a small hematoma to a large hematoma leading to a mass effect. CT and MRI are easy to diagnose intracranial subdural hematoma [18]. In our case, the subdural hematoma is presented as a high-density thin layer in the left frontal region. Radiological manifestations of pituitary apoplexy vary depending on the duration of apoplexy. A large pituitary gland can be observed in most cases. CT images can show patchy or confluent areas of hyperdensity in a pituitary lesion if a hemorrhagic component is present. The density of the lesion is often mixed and varies over time due to hemolysis. MRI may identify the presence of an adenoma and hemorrhagic degeneration of pituitary apoplexy lesions. Early findings appear as sellar enlargement and abnormal signal intensity with poor enhancement or rim enhancement. Late findings appear with an empty sella of normal size [13, 14, 19, 20].

Immediate medical treatment of subdural hematoma and pituitary apoplexy should begin with careful assessment of the patient’s condition, management of the airway, and supportive measures to ensure hemodynamic stability, fluid, and electrolyte balance. After monitoring and stabilization of the patient, a secondary care plan should be implemented. Conservative nonsurgical management for subdural hematoma may be considered if the accumulation does not cause
impingement on the brain or brain stem. Surgical evacuation should be performed promptly if there are signs of increased intracranial pressure threatening vital functions [21, 22]. Hypopituitarism can resolve on its own in some cases; in others, it will lead to irreversible hormone deficiencies. Hormone replacement therapy must be individualized to the needs of each specific patient. Patients may need lifelong hormone replacement therapy [23–25].
4. Conclusion

The coexistence of multiple intracranial complications is uncommon after spinal anesthesia for cesarean section. However, in the specific case, no connection exists between subdural hematoma, connected with anesthesia, and pituitary apoplexy, which specifically has no diagnosed cause. In short, it is a casual association. This case helps to add knowledge to the current literature and may be useful to clinicians managing patients if similar situations arise. It is essential to diagnose these conditions early and manage appropriately.

Data Availability

There are no data to share. All the details have been mentioned in the manuscript.

Consent

Written informed consent was obtained from the patient in this case report.

Conflicts of Interest

The authors declare that there is no conflict of interest regarding the publication of this article.

Authors’ Contributions

VTH and VC contributed to the conception and design of the manuscript and data acquisition and wrote the initial draft. THH and TTTN critically revised the manuscript. DTH supported the revision and critically reviewed the paper. All authors participated in the approval of the final version.

Acknowledgments

The authors thank the patient for providing the data and for the permission to publish this article.

References


