

Review began 03/12/2025 Review ended 03/20/2025 Published 03/20/2025

© Copyright 2025

Sadek Azar et al. This is an open access article distributed under the terms of the Creative Commons Attribution License CC-BY 4.0., which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

DOI: 10.7759/cureus.80911

Subcapsular Hepatic Hematoma Secondary to Preeclampsia With a Successful Maternal Outcome: A Case Report

Rashid Sadek Azar ¹, Jocelyn M. Leon Turriza ¹, Roberto Velazquez Tass ¹, Roxana Guillemin Sosa Montejo ¹

1. Obstetrics and Gynecology, Hospital General de Especialidades, Campeche, MEX

Corresponding author: Rashid Sadek Azar, rashid_sadek92@hotmail.com

Abstract

Subcapsular hepatic hematoma (SHH) is a rare and life-threatening condition, often leading to severe maternal complications. Its diagnosis is frequently challenging due to nonspecific symptoms and a reliance on ultrasound findings. Case reports for this type of pregnancy complication are limited due to its low incidence.

A 30-year-old pregnant woman at 32.3 weeks with preeclampsia with severe features presented with epigastric pain and high blood pressure, requiring an emergency C-section. She developed a peri-hepatic subcapsular hematoma, leading to hypovolemic shock and multiple surgeries, including liver packing. After stabilization and follow-up, she was discharged without complications.

We reported a rare case of a patient with hepatic subcapsular hematoma secondary to preeclampsia with severe features. SHH is a rare but serious pregnancy complication, often linked to hemolysis, elevated liver enzymes, and low platelets (HELLP) syndrome and preeclampsia, with a high maternal mortality rate. Diagnosis is challenging due to nonspecific symptoms, requiring imaging studies such as ultrasound or CT. Treatment varies from conservative management to surgical intervention, depending on hemodynamic stability.

Categories: Family/General Practice, Obstetrics/Gynecology, Radiology

 $\textbf{Keywords:} \ damage \ control \ laparotomy, \ liver \ lesion, \ preeclamps ia \ with \ hellp \ syndrome, \ preeclamps ia \ with \ severe \ features, \ subcapsular \ hepatic \ hematoma$

Introduction

A subcapsular hepatic hematoma (SHH) is an accumulation of blood located between Glisson's capsule and the liver parenchyma. Its incidence during pregnancy is between 2% and 8%, with a high maternal mortality risk ranging from 17% to 59%, particularly in cases associated with hemolysis, elevated liver enzymes, and low platelets (HELLP) syndrome and preeclampsia with severe features [1]. The clinical presentation of a SHH is nonspecific, making diagnosis challenging. Diagnosis is confirmed through imaging studies, with ultrasound as the first-line option, followed by CT and MRI [2-5]. The treatment of SHHs can be conservative or non-conservative, depending on the patient's hemodynamic stability and the severity of the liver injury [1,2,4].

Case Presentation

A 30-year-old female, currently in her second pregnancy with a previous miscarriage and no other significant medical history, was at 32.3 weeks of gestation with a diagnosis of gestational hypertension, previously managed with methyldopa 500 mg every 8 hours. She presented to the obstetric ED with epigastric pain radiating to the right shoulder for one week, accompanied by elevated blood pressure readings of up to 180/105 mmHg at the time of evaluation. This required the administration of two boluses of hydralazine 5 mg and the initiation of neuroprotection with magnesium sulfate using a modified Zuspan protocol.

Upon admission to the obstetric surgery unit, her blood pressure remained above the target range (160/110 mmHg), accompanied by epigastric pain radiating to the shoulder and other signs of vasospasm. Laboratory tests were ordered, revealing the following results: hemoglobin 13.8 g/dL, hematocrit 39%, platelets 145,900/mm³, aspartate aminotransferase (AST) 83 U/L, alanine aminotransferase (ALT) 111 U/L, and a urinalysis showing protein levels of 30 mg/dL (Table 1). Due to the persistence of symptoms, an urgent cesarean section was performed to improve maternal and fetal outcomes.



Laboratory Parameter	Initial Admission	Postoperative (After First Surgery)	Postoperative (After Second Surgery)	Normal Range
Hemoglobin (g/dL)	13.8	9.52	7.21	12-15
Hematocrit (%)	39	26	21	36-48
Platelets (/mm³)	145,900	157,500	131,900	150-450
AST (U/L)	83	87	500	15-38
ALT (U/L)	111	1000	907	13-40
Proteinuria (mg/dL)	30	Not reported	Not reported	Not detected

TABLE 1: Patient's laboratory results from initial admission to postoperative period after the second surgery.

AST: Aspartate aminotransferase; ALT: Alanine Aminotransferase.

A male newborn was delivered at 05:30 AM, weighing 1800 g, with an Apgar score of 3-5, gestational age of 33 weeks, and a length of 45 cm. The infant had no muscle tone and required two cycles of positive pressure ventilation, orotracheal intubation, and admission to the Neonatal Intensive Care Unit (NICU). Additional findings included scant, clear amniotic fluid, a normal umbilical cord, an anterior placenta with 20% abruption, and two fibroids on the anterior uterine wall measuring approximately 1×1 cm. The total blood loss was 800 cc.

At the end of the procedure, during monitoring, signs of hypovolemic shock were detected, with a blood pressure of 70/40 mmHg and a heart rate of 130 bpm. A maternal code was activated, and fluid resuscitation along with vasopressor support was initiated. An intraoperative abdominal ultrasound revealed a perihepatic subcapsular hematoma predominantly affecting the right lobe, measuring approximately $16.5 \times 4.3 \times 11.4$ cm (Figure 1), with an estimated volume of 430 ml. Additionally, there was non-quantifiable free fluid in the perisplenic region, approximately 60 ml in the left paracolic gutter, 50 ml in the right paracolic gutter, and 80-100 ml in the periuterine pelvic cavity. The uterus was involuting, with post-surgical changes in the myometrium and no evidence of retained placental tissue. Laboratory results showed hemoglobin 9.52 g/dL, hematocrit 26%, platelets 157,500/mm³, AST 87 U/L, and ALT 1000 U/L (Table 1).



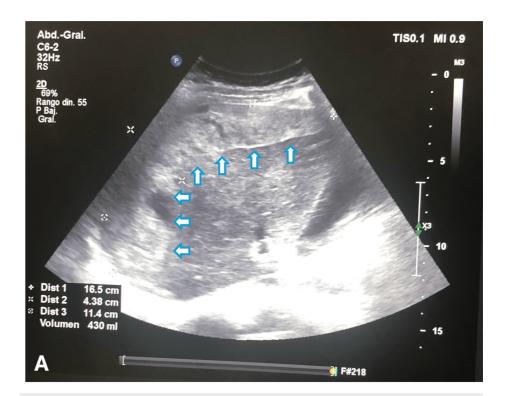


FIGURE 1: Abdominal ultrasound showing a subcapsular hepatic hematoma in the right lobe.

An emergency exploratory laparotomy was performed by the General Surgery team, revealing $800 \, \text{ml}$ of hemoperitoneum and a ruptured SHH measuring $15 \, \text{cm}$ in segments VI and VII (Figure 2). The liver was packed with five gauze pads wrapped in a latex glove. A $\frac{34}{100}$ -inch Penrose drain was placed in the foramen of Winslow and exteriorized through a counter-incision in the right flank.



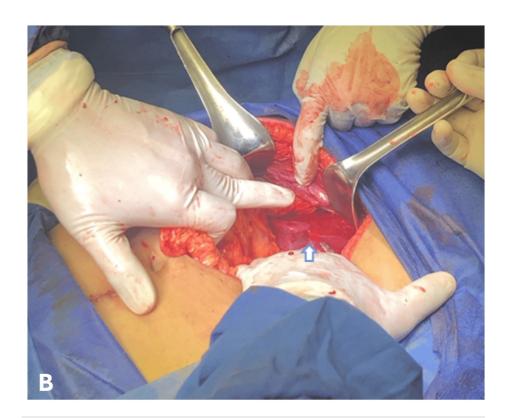


FIGURE 2: Hepatic laceration is observed following the drainage of a subcapsular hepatic hematoma.

Following surgery, the patient was admitted to the adult ICU. Four hours later, a second maternal code was activated due to active, non-quantifiable bleeding from the Penrose drain. A reoperation was performed by the General Surgery team, revealing an extension of the hematoma toward the posterolateral liver with 850 cc of hemoperitoneum. A second packing procedure was performed using six gauze pads after dissecting the round and cardinal ligaments of the liver, with intact hepatic veins observed.

Forty-eight hours after the second surgery, unpacking and abdominal closure were performed. Six gauze pads were removed from the hepatic bed without active bleeding; however, a hematoma in segments VI and VII with 150 ml of blood was noted. This was managed conservatively by the General Surgery team.

Postoperative laboratory tests showed hemoglobin 7.21 g/dL, hematocrit 21%, platelets 131,900/mm 3 , AST 500 U/L, and ALT 907 U/L (Table 1). The patient was discharged without complications 48 hours after the last surgical event, with a follow-up scheduled at six weeks, during which no further complications were reported.

Discussion

Subcapsular hepatic hematoma is a rare but severe complication of hypertensive disorders of pregnancy, particularly preeclampsia with severe features and HELLP syndrome. It occurs due to hepatic vasospasm, endothelial dysfunction, and microvascular injury, leading to hemorrhage beneath the liver capsule [3]. If left undiagnosed or untreated, SHH can rupture, resulting in massive hemoperitoneum, hemorrhagic shock, and high maternal-fetal mortality [2]. In this case, the patient presented with epigastric and right shoulder pain, a hallmark symptom of hepatic involvement in preeclampsia. Despite antihypertensive management with methyldopa, her blood pressure remained severely elevated, necessitating emergency cesarean delivery. The presence of persistent pain and signs of hemorrhagic shock postoperatively prompted urgent imaging, leading to the diagnosis of a ruptured hepatic hematoma. Immediate surgical intervention, including liver packing and damage control resuscitation, was essential for hemodynamic stabilization.

The management depends on its size, stability, and presence of rupture [2,5]. In stable, unruptured cases, conservative management with close monitoring may suffice. However, in cases like this one, where a rupture occurs, surgical intervention is life-saving [2,5]. Treatment options include hepatic packing, embolization, or even partial hepatectomy in severe cases [5]. This case demonstrates the importance of a high index of suspicion in hypertensive pregnant patients with unexplained epigastric pain, as timely intervention can prevent maternal and fetal morbidity and mortality.



Despite a challenging clinical course, the patient had a favorable outcome due to early recognition, multidisciplinary surgical management, intensive postoperative care, and the patient made a full recovery without long-term complications. This case underscores the critical role of prompt diagnosis, aggressive surgical intervention, and vigilant hemodynamic monitoring in managing hepatic complications of severe preeclampsia.

Conclusions

This case highlights the rare but life-threatening complication of SHH in the context of preeclampsia with severe features. The patient's initial presentation with epigastric pain, right shoulder pain, and severe hypertension was indicative of hepatic involvement, which ultimately led to the rupture of a subcapsular hematoma following delivery. Early recognition and multidisciplinary management, including urgent abdominal delivery, intraoperative ultrasound, exploratory laparotomy, and sequential surgical interventions, were critical in preventing maternal mortality. This case underscores the importance of early diagnosis, close monitoring, and timely surgical intervention in managing hepatic complications of hypertensive disorders in pregnancy.

Additional Information

Author Contributions

All authors have reviewed the final version to be published and agreed to be accountable for all aspects of the work

Concept and design: Rashid Sadek Azar, Jocelyn M. Leon Turriza, Roberto Velazquez Tass, Roxana Guillemin Sosa Montejo

Acquisition, analysis, or interpretation of data: Rashid Sadek Azar, Jocelyn M. Leon Turriza, Roberto Velazquez Tass, Roxana Guillemin Sosa Montejo

Drafting of the manuscript: Rashid Sadek Azar, Jocelyn M. Leon Turriza, Roberto Velazquez Tass, Roxana Guillemin Sosa Montejo

Critical review of the manuscript for important intellectual content: Rashid Sadek Azar, Jocelyn M. Leon Turriza, Roberto Velazquez Tass, Roxana Guillemin Sosa Montejo

Supervision: Rashid Sadek Azar

Disclosures

Human subjects: Consent for treatment and open access publication was obtained or waived by all participants in this study. **Conflicts of interest:** In compliance with the ICMJE uniform disclosure form, all authors declare the following: **Payment/services info:** All authors have declared that no financial support was received from any organization for the submitted work. **Financial relationships:** All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. **Other relationships:** All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

- Chen H, Li Y, Yang S, et al.: Spontaneously ruptured huge hepatic subcapsular hematoma associated with preeclampsia: a case report and literature review. J Int Med Res. 2024, 52:3000605241264220. 10.1177/03000605241264220
- Bradke D, Tran A, Ambarus T, Nazir M, Markowski M, Juusela A: Grade III subcapsular liver hematoma secondary to HELLP syndrome: a case report of conservative management. Case Rep Womens Health. 2020, 25:e00169. 10.1016/j.crwh.2019.e00169
- Luhning K, MacCormick H, Macaulay B, Saunders M, Craig C: Subcapsular hepatic hematoma as a complication of severe preeclampsia: a case report. J Med Case Rep. 2021, 15:625. 10.1186/s13256-021-07166 pt.
- Moore EE, Cogbill TH, Jurkovich GJ, Shackford SR, Malangoni MA, Champion HR: Organ injury scaling: spleen and liver (1994 revision). J Trauma. 1995, 38:323-324. 10.1097/00005373-199503000-00001
- Kapan M, Evsen MS, Gumus M, Onder A, Tekbas G: Subcapsular liver hematoma in HELLP syndrome: case report. Gastroenterology Res. 2010, 3:144-146. 10.4021/gr2010.04.178e