Grade III subcapsular liver hematoma secondary to HELLP syndrome: A case report of conservative management

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Abstract

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Subcapsular liver hematoma (SLH) is a rare condition that is associated with preeclampsia and HELLP (hemolysis, elevated liver enzymes, and low platelets) syndrome. A high level of suspicion, early diagnosis, and coordinated, intensive multidisciplinary management are necessary to monitor for serious complications and prevent death. Options include conservative management, hepatic resection, hepatic artery ligation and liver transplantation. This paper describes a 34-year-old woman with HELLP syndrome who developed a large grade III SLH that was managed conservatively.

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1. Introduction

Subcapsular liver hematoma (SLH) is spontaneous bleeding between the Glisson’s capsule and the liver parenchyma. It occurs in approximately 2% of pregnancies complicated by HELLP (hemolysis, elevated liver enzymes, and low platelets) syndrome. Unrecognized SLH can have devastating consequences, one of the worst complications being hepatic rupture, which has a significant mortality rate. Due to the acute and critical nature of SLH, even with aggressive medical and/or surgical interventions the outcome can be devastating.

2. Case Report

A 34-year-old woman, gravida 2, para 0, at 37 weeks of gestation presented to the labor and delivery department for evaluation of new-onset hypertension in pregnancy with a blood pressure (BP) of 140/92 mmHg. On initial presentation, the patient had no symptoms of preeclampsia but was noted to have hyperreflexia (3+ deep-tendon reflexes) on physical exam. Laboratory findings were significant for 2+ proteinuria, a platelet count (PLT) of 122 × 103/mcL, an aspartate aminotransaminase (AST) level of 116 U/L, alanine aminotransaminase (ALT) level of 162 U/L and lactate dehydrogenase (LDH) level of 300 U/L. Based on the laboratory findings, in conjunction with the physical exam, and elevated BP, the diagnosis of HELLP syndrome was made. HELLP syndrome is associated with adverse pregnancy outcomes; therefore the decision was made to induce labor. The patient was started on magnesium sulfate for seizure prophylaxis and induction was started using vaginal misoprostol for cervical ripening.

During the induction process, serial samples were drawn for laboratory analysis. AST trended upwards to a maximum of 3110 U/L, ALT to 2607 U/L, LDH to 2991 U/L, and platelets fell to 58 × 103/mcL (Table 1). The patient progressed appropriately, leading to an uncomplicated vaginal delivery of a viable male infant with APGAR scores of 8 and 9. Within hours of delivery, the patient had an abrupt onset of pain in the right upper quadrant, which proved to be persistent. Physical exam revealed a tender right upper quadrant with rebound and a liver edge beyond the costal margin. A subcapsular liver hematoma was suspected and confirmed on abdominal ultrasound (Fig. 1). A computerized tomography (CT) angiogram was performed with liver mass protocol, which demonstrated a 16 × 6 × 22 cm subcapsular liver hematoma without evidence of active extravasation (Fig. 2). A multidisciplinary consultation was rapidly convened that included obstetrics, maternal-fetal medicine, general surgery, radiology, intensive care, and hematology. The decision was made to transfer the patient to the intensive-care unit (ICU) for conservative management, blood product repletion, and monitoring.

Once in the ICU, the patient underwent close continuous monitoring and serial abdominal exams were performed. The patient was transfused a total of 5 units of packed red blood cells (PRBC) and 2 units of platelets. Laboratory results remained stable. A repeat CT scan on postpartum day 1 showed a stable hematoma (Fig. 3). Pain was managed initially with patient-controlled intravenous hydromorphone, and...
then switched to oral hydromorphone and cyclobenzaprine. The patient improved clinically and was eventually downgraded to the postpartum unit. A repeat CT scan on postpartum day 8 showed that the hematoma had decreased in size (Fig. 4) and laboratory values were noted as ALT 207 U/L, AST 81 U/L, LDH 1132 U/L, and PLT 464 × 10³/mcL. Marked improvement continued and the patient was discharged home on postpartum day 10. The patient then proceeded to a full recovery with resolved transaminitis. Long-term follow-up was with gastroenterology and a CT scan 2 months after delivery showed interval organization of the hematoma; otherwise, the CT scan of the liver was grossly normal (Fig. 5).

3. Discussion

Subcapsular liver hematoma (SLH) is a rare but potentially fatal complication of HELLP syndrome, occurring in about 1–2% of all patients with preeclampsia and HELLP syndrome [1,2]. The pathogenesis of SLH in HELLP syndrome is not well understood but may involve fibrin deposition leading to platelet activation, thrombus formation, and occlusion of capillaries, which ultimately progresses to hepatic necrosis and hemorrhage [2,3]. The most concerning potential complication of SLH is hepatic rupture, which may occur in the setting of trauma, such as abdominal palpation, manual removal of the placenta, and uterine contractions [1]. Hepatic rupture due to SLH most commonly involves the right lobe of the liver [2]. Maternal mortality of SLH ranges from 17% to 59%, depending on rupture of the hematoma, timing of diagnosis, and availability of therapeutic interventions [2].

The signs and symptoms of SLH are nonspecific. They include hemodynamic instability and sudden-onset pain in the right upper quadrant or epigastric pain, severe pain in the right shoulder, and/or nausea and vomiting due to distention of the hepatic parenchyma and Glisson’s capsule [2,4]. In the case of hepatic rupture, abdominal distension due to hemoperitoneum and hypovolemic shock may be seen [2]. The clinical presentation of SLH may be confused with the usual pain in the right upper quadrant seen in preeclampsia, other intra-abdominal pathology such as cholecystitis, or even pulmonary embolism [2].

As with this case, early suspicion for SLH should lead to prompt evaluation and diagnosis. The choice of imaging studies should be based on availability and how quickly they can be performed. Therefore, portable ultrasound is a good screening tool as it is the quickest and can be

| Table 1 |
| Laboratory values. |
| Prior to induction | T+ 7.5 h | T+ 12 h | Delivery +2 h | Delivery +6 h | Delivery +14 h | Postpartum day 1 | Postpartum day 2 | Postpartum day 3 |
| Leukocytes (4.5–11.0 × 10³/mcL) | 11.4 | 15.1 | 11.5 | 12.3 | 12.4 | 14.4 | 15.4 | 13.5 | 13.6 |
| Hemoglobin (12.0–15.0 g/dL) | 12.2 | 8.2 | 9.4 | 8.9 | 9.1 | 8.9 | 8.5 | 7.8 | 8.7 |
| Hematocrit (35.0–45.0%) | 35.3 | 23.1 | 26.9 | 25.8 | 25.8 | 25.8 | 24.7 | 22.1 | 24.9 |
| Platelet (150–450 × 10³/mcL) | 79 | 70 | 58 | 85 | 85 | 98 | 117 | 113 | 203 |
| PT (9.8–11.7 Seconds) | 9.4 | 9.9 | 10.3 | 9.8 | 9.7 | 9.5 | 9.6 | 9.5 | 10.5 |
| INR (0.9–1.1) | 0.9 | 0.9 | 1 | 0.9 | 0.9 | 0.9 | 0.9 | 0.9 | 1 |
| PTT (24.3–30.4 Seconds) | 23.2 | 21.3 | 22.9 | 23.3 | 23.3 | 23.3 | 23.7 | 25 | 24.4 |
| Fibrinogen (210–400 mg/dL) | 520 | 399 | 341 | 367 | 314 | 385 | 424 | 546 | – |
| Glucose (90–110 mg/dL) | 110 | 96 | 91 | 91 | 84 | 78 | 87 | 90 | 122 |
| BUN (6.0–23.0 mg/dL) | 12 | 23 | 22 | 18 | 16 | 12 | 11 | 11 | 8 |
| Creatinine (0.60–1.30 mg/dL) | 0.48 | 0.63 | 0.52 | 0.46 | 0.43 | 0.4 | 0.35 | 0.38 | 0.38 |
| Uric Acid (2.6–7.4 mg/dL) | 6.3 | 8.8 | 8.4 | 7.7 | 6.9 | 6.4 | – | 4.5 | – |
| Urine Protein (0–10.0 mg/dL) | 117.1 | 75.4 | – | 47.8 | – | – | – | – | – |
| Urine Creatinine (mg/dL) | 151.2 | 184.6 | – | 109.2 | – | – | – | – | – |
| ALT (6–52 Unit/L) | 332 | 2444 | 2607 | 2445 | 2186 | 1962 | 1464 | 932 | 580 |
| AST (15–40 Unit/L) | 327 | 2615 | 3110 | 2606 | 2262 | 1739 | 762 | 289 | 110 |
| LDH (95–230 Unit/L) | 513 | 3028 | 3223 | 2438 | 2055 | 1399 | 551 | 356 | – |
| ALP (46–116 Unit/L) | 172 | 159 | 162 | 155 | 158 | 160 | 152 | 147 | 142 |

Fig. 1. Ultrasound scan of the right upper quadrant.

Fig. 2. Computerized tomography angiogram, postpartum day 0.
performed at the bedside [2,5] and a FAST (focused abdominal ultrasound for trauma) exam should be the main imaging modality in unstable patients. CT and magnetic resonance imaging (MRI) are both highly sensitive for the detection and evaluation of SLH [3]. However, as CT is more readily available and takes less time to complete, it is more appropriate in emergency cases where patients are becoming hemodynamically unstable [6]. In the case presented, abdominal ultrasound was used to establish the diagnosis of SLH while CT angiogram was used to confirm the diagnosis and further describe the extent of the hematoma. MRI should be reserved for non-emergency case or for pregnant patients [2].

In order to classify injuries and help guide management, the American Association for the Surgery of Trauma (AAST) created a classification system that grades the extent of injury from grade I to grade VI (Table 2). As hepatic hematomas from HELLP syndrome are rare, critical care management can be guided using the AAST classification. In data published by the National Trauma Data Bank (NTDB), approximately 67% of hepatic injuries were between grade I and grade III [8]. Hemodynamically stable patients with low-grade injuries (I, II, III) were able to be successfully managed without surgical intervention [8]. For patients with grade IV and V injuries, non-surgical management was less successful, and patients with grade VI trauma (hepatic avulsion) were universally unstable and required surgical intervention [8]. Our patient had a hematoma that was >10 cm intraparenchymal and therefore categorized as grade III. Similar to the published trauma literature, our patient's grade III hematoma responded to non-surgical management.

Multiple case reports stress that the management of patients with SLH should be at tertiary centers, in order for timely recognition and for treatment to have an optimal outcome [7]. Traditionally, first-line treatment for SLH has been surgical management. Patients with hepatic rupture and unstable patients are managed with immediate exploratory laparotomy and transfusion of blood products [5]. Surgical options for stabilization of patients with a ruptured hematoma include perihepatic packing and drainage of the surgical site, ligation of the appropriate branch of the portal vein or hepatic artery, patching of the omentum and partial liver resection [2,12]. Liver transplant should be considered in patients with refractory hepatic hemorrhage or with rapidly progressing fulminant liver failure [12]. However, in non-bleeding, hemodynamically stable patients with SLH, as was the case with our patient, non-operative, conservative management is the preferred approach [2,9].

While no specific guidelines exist to aid in clinical decision-making, the AAST scoring system is a useful tool in deciding between surgical and non-surgical management in hemodynamically stable patients. Conservative management has been shown to be successful in treating non-ruptured hematomas and should be advocated in these cases to

| Grade | Type of injury | Description of injury |
|-------|----------------|-----------------------|
| I     | Hematoma       | Subcapsular, <10% Surface Area |
|       | Laceration     | Capsular tear, <1 cm, Parenchymal Depth |
| II    | Hematoma       | Subcapsular, 10% to 50% of Surface Area |
|       | Laceration     | Intraparenchymal <10 cm in Diameter |
| III   | Hematoma       | Subcapsular, >50% Surface Area of Ruptured Subcapsular or Parenchymal Hematoma |
|       | Laceration     | Intraparenchymal Hematoma >10 cm or Expanding |
| IV    | Laceration     | >3 cm Parenchymal Depth |
| V     | Laceration     | Parenchymal Disruption Involving >75% of Hepatic Lobe or >3 Couinaud’s Segments |
|       | Vascular       | Juxtahepatic Venous Injuries. Ex: Retrohepatic Vena Cava or Central Major Hepatic Veins |
| VI    | Vascular       | Hepatic Avulsion |

* For Multiple Injuries, Advance One Grade up to Grade III.
avoid unnecessary liver trauma, which could lead to rupture and catastrophic bleeding [2,9,13]. This involves intensive care with infusion of fluid, transfusion of blood products when necessary, correction of coagulopathy, and serial imaging with ultrasound or CT to monitor the size of the SLH [10]. Percutaneous trans catheter hepatic artery embolization by interventional radiology has also been shown to be a successful alternative to control hemorrhage [2,11]. Surgical evaluation is necessary if the hematoma increases in size or the condition of the patient deteriorates. While complete resolution of the hematoma can take several months, if the patient’s condition and the size of the hematoma remain stable, the patient can be discharged and followed up as an outpatient [2]. As SLH due to HELLP is a rare condition, long-term outcome data are limited to case reports or are extrapolated from trauma surgery and critical care medicine. At this time, long-term outcomes appear to depend on the liver injury grade, amount of or lack of hepatic necrosis and/or fibrosis, and the existence of patient co-morbidities.

In conclusion, SLH is a rare complication of HELLP syndrome but should be considered due to the risk of hepatic rupture. Early recognition and treatment are crucial to optimizing patient outcomes. While surgical management has traditionally been the first-line treatment for SLH, conservative management with a multidisciplinary team that includes critical care, surgery, hematology, and gastroenterology should be considered in hemodynamically stable patients with AAST grade I, II and III hematomas, such as the patient presented in this report.

**Contributors**

Daniel Bradke contributed to the conception and design of the study, acquisition of data, analysis and interpretation of data, and drafting or revision of the article.

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Alexander Jusela contributed to the conception and design of the study, acquisition of data, analysis and interpretation of data, and drafting or revision of the article.

**Conflict of Interest**

The authors declare that they have no conflict of interest regarding the publication of this case report.

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