Hemorrhagic ascites from spontaneous ectopic mesenteric varices rupture in NASH induced cirrhosis and successful outcome: A case report

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Core tip: Nonalcoholic steatohepatitis (NASH) related cirrhosis of the liver is an emerging disease. With the advent of new therapies for hepatitis C and the potential for cure, NASH will most likely be the leading cause of decompensated liver disease in the future. We present a rare case of hemorrhagic ascites from ectopic variceal rupture as the initial decompensating event in a young patient with a recent diagnosis of cirrhosis from NASH. A multidisciplinary, methodical treatment plan was undertaken, culminating in orthotopic liver transplantation and successful outcome. We briefly discuss presentation, diagnosis and management of ectopic variceal bleeding, which is not so commonly encountered in routine clinical practice in this case report.

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INTRODUCTION

Ectopic variceal bleeding in cirrhotic patients with significant portal hypertension is a well-known although rare complication, accounting for about 5% of all variceal bleeding[1]. It can often present as obscure gastrointestinal bleed depending on location of varices. Unlike esophageal variceal bleeding, for which definite guidelines exist in terms of management and surveillance, ectopic varices due to their sporadic occurrence and varying presentations depending on location, do not...
Hemorrhagic ascites in decompensated cirrhosis have specific guidelines in terms of management. Recognizing the various presentations of ectopic variceal bleed is vital to the survival of these patients and a high degree of suspicion is necessary for timely diagnosis and effective management. Hemorrhagic ascites presenting as a complication of intraperitoneal ectopic variceal rupture is a dreaded and potentially fatal complication, management of which requires a multi-disciplinary approach along with timely recognition and diagnosis.

CASE REPORT

A 37 years old Caucasian male presented to the emergency room with a 3 day history of right upper and lower quadrant abdominal pain with distension. Physical examination revealed palmar erythema, icterus and moderate ascites with tenderness in the right lower quadrant. He was tachycardic but hemodynamically stable at presentation with a low grade fever. Review of his recent medical records revealed that he was newly diagnosed with NASH cirrhosis at an outside facility, about 3 mo prior to this presentation, as a part of work up of abdominal pain which demonstrated abnormal appearing liver. Subsequent workup and liver biopsy (Figure 1) suggested the diagnosis of NASH related cirrhosis after ruling out all other etiologies. He did not have any abdominal surgeries in the past and was not on any medications.

Investigations

Initial laboratory data at the time of this presentation showed; Hb: 8.7 gm/dL, Hct: 25.2, WCC: 149 and INR: 1.81. Complete metabolic panel revealed Cr: 0.97 mg/dL, protein: 6.7 gm/dL, albumin: 2.9 gm/dL, total bilirubin: 18.2 mg/dL, direct bilirubin: 13.2 mg/dL, Alkp: 164U/L, AST: 140U/L, ALT: 44U/L and a MELD score of 24. Computerized tomography of the abdomen with contrast (Figure 2) revealed lobulated contour of the liver consistent with cirrhosis (arrow), severe porto-systemic collateral circulation with esophageal varices, mesenteric stranding and small to moderate volume ascites with dependent hyper dense areas within the fluid suggestive of hemorrhagic component. Patient underwent diagnostic paracentesis with removal of 6 liters of grossly bloody fluid consistent with intra-peritoneal hemorrhage.

Clinical course

Upon admission patient was initially resuscitated with blood products, fluids, correction of coagulopathy and empirical antibiotics were initiated. His clinical condition continued to worsen with drop in blood pressure and hemoglobin. A CTA and CTV of the abdomen were performed (Figure 3) to identify the source of bleeding. This revealed right abdominal mesenteric varices with a focal area of more consolidated abnormal enhancement.

Figure 1 Macrovesicular steatosis (A) and steatosis and cirrhotic nodules (B).

Figure 2 Contrast computed tomography abdomen (arrow). The liver consistent with cirrhosis (arrow).
in the right lower quadrant (arrow). A possible nidus of venous malformation between superior mesenteric vein and right common iliac vein varices was suggested with more hyper dense ascites adjacent to this suggestive of possible origin of hemoperitoneum.

The patient underwent an angiogram by intervention radiology with trans-hepatic access of the portal vein, portal venogram, superior mesenteric venogram and coil embolization of colonic branch of superior mesenteric vein and cecal branch of the superior mesenteric vein (Figure 4). There were no immediate complications. The procedure controlled his intra-abdominal bleeding and he remained hemodynamically stable. Over the next 48 h he developed first episode of hematochezia with hypotension. Upper gastrointestinal (GI) endoscopy revealed large esophageal varices with evidence of recent bleeding. He underwent an emergent Trans jugular Intrahepatic Portosystemic shunt (TIPS) procedure with successful decompression of portal system. The hepatic venous pressure gradient was reduced from 17 to 9 mmHg post procedure. He eventually underwent successful orthotopic liver transplantation after appropriate evaluation of decompensated end stage liver disease during the same presentation and was discharged home in a stable condition.

**DISCUSSION**

Ectopic varices are defined as large porto-systemic venous collaterals occurring anywhere in the abdomen other than the cardio-esophageal region. These are uncommon and account for less than 5% of all cases of variceal bleeding. Most patients show presence of esophageal varices simultaneously and may have a history of treatment for them in the form of endoscopic intervention or primary prophylaxis. They have been reported to occur at numerous sites which include 17% in the jejunum or ileum, 17% in the duodenum, 14% in the colon, 8% in the rectum and 9% in the peritoneum in a retrospective study. Ectopic varices may bleed even when portal venous pressure is low. They have a 4 fold increased risk of bleeding when compared to esophageal varices and can have a mortality rate as high as 40%.

Clinical presentation of ectopic variceal bleed is based on location of the varices. Luminal varices are fortunately more common, usually easier to detect and manifest earlier than the non-luminal ectopic varices. Clinical manifestations include overt GI bleeding of obscure origin, occult GI bleeding, accidental finding, iron deficiency anemia, hematemia, hematochezia, hemoperitoneum, hypovolemic shock, hemorrhagic pleural effusion and sometimes diagnosis is made only at autopsy. Ectopic varices should be considered in all patients with portal hypertension and GI bleeding, if both upper and lower endoscopies failed to show obvious source. Awareness of the condition is a necessity for any physician dealing with GI bleeding, especially in the setting of cirrhosis and portal hypertension.

Diagnosis of ectopic variceal bleed in a timely manner is important due to the high mortality rate associated with the condition. Endoscopic and angiographic techniques may be required as they can appear as filling defects in barium studies of the bowel and may be misdiagnosed as polyps or tumors. Abdominal wall varices usually rupture externally and can be easily diagnosed. Mesenteric, diaphragmatic, faliform ligament, splenic ligament and rectovesical varices may rupture into the peritoneal cavity causing internal bleeding and fatal outcomes. This requires high degree of clinical suspicion when presentations include rapid accumulation of ascites, reduction in hematocrit and signs of hypovolemic shock. CT scan of the abdomen and pelvis aids in the diagnosis and confirmed by detection of bloody ascites at paracentesis. Angiographic techniques which include percutaneous transhepatic portography, transjugular transhepatic portography, splenic portography and umbilical vein catheterization can be used to identify the location of bleeding, extent of portosystemic collaterals, direction of flow and simultaneously measure the pressure in the portal venous system.

There are no large randomized controlled trials that have previously addressed the therapeutic modalities for ectopic varices. Most of the available knowledge is obtained from small case series, case reports and mini reviews. The management of ectopic varices requires a multidisciplinary team of hepatologists, gastroenterologists, surgeons and interventional radiologists. Due to the diversity of their location and clinical presentation it is very difficult to draw treatment guidelines. Optimal treatment depends on location of varices, patient’s condition and the availability of local expertise and resources. Initial management includes appropriate resuscitation, emergent evaluation to localize site and source of bleeding followed by suitable treatment. The use of vasoactive medications like somatostatin (octreotide) and terlipressin to reduce splanchic blood flow and variceal pressure may be of benefit in patients with bleeding from gastroesophageal varices. No solid data exists on the role of beta blockers in the long term management of ectopic varices. Case reports have been contradictory as to their
Endoscopically accessible varices can be treated by band ligation or sclerotherapy. Those which are not accessible by endoscopic techniques can be treated by interventional radiology and may include coil embolization, balloon occluded retrograde transvenous obliteration and if unable to access the site a TIPS shunt to decompress the portal pressure could be a lifesaving intervention. Ultimately the long term survival of these patients is dependent on their liver function and liver transplantation should be considered as a definitive therapy after appropriate evaluation as in our case report.

In conclusion, Ectopic variceal bleeding is a rare event unlike esophageal variceal bleeding, for which specific treatment guidelines do not exist. A high degree of suspicion is essential for the entity as a delay in diagnosis could lead to fatal consequences. Due to the lack of specific guidelines in management, a multi-disciplinary approach utilizing local expertise and resources has been recommended. Early diagnosis, prompt resuscitation followed by appropriate intervention is crucial in the management of these patients, who have severe portal hypertension and are already coagulopathic from end stage liver disease. Appropriate referral to a tertiary care center should be initiated when resources are not available locally and liver transplantation should be considered as a definitive treatment to give them the best chance of long
term survival.

**COMMENTS**

**Case characteristics**
A 37-year-old male with recently diagnosed nonalcoholic steatohepatitis cirrhosis presented with abdominal pain and distension.

**Clinical diagnosis**
Palmar erythema, scleral icterus, shifting dullness on abdominal percussion and right lower quadrant tenderness.

**Differential diagnosis**
Aspects from portal hypertension, spontaneous bacterial peritonitis, intra-abdominal hemorrhage from variceal rupture, hepatocellular carcinoma rupture and perforated viscus.

**Laboratory diagnosis**
Hemoglobin: 8.7 g/dL, WCC: 17.6, platelets: 149, INR: 1.8, bilirubin: 18.2 mg/dL, alkaline phosphatase: 164, AST: 140, ALT: 44, creatinine: 0.97 mg/dL, and MELD score: 24.

**Imaging diagnosis**
CT/CTA/CVT of abdomen: Cirrhosis, ascites, right abdominal mesenteric varices with focal area of enhancement in the right lower quadrant indicating hemoperitoneum.

**Pathological diagnosis**
Paracentesis revealed hemorrhagic ascites. Trans-hepatic portal vein angiogram revealed mesenteric varices with bleeding from colonic branch and cecal branch.

**Treatment**
Coil embolization of bleeding mesenteric varices by intervention radiologist. Trans jugular intrahepatic Porto-systemic shunt for esophageal variceal bleed and eventually orthotopic liver transplantation for end stage liver disease.

**Related reports**
Ectopic variceal bleed accounts for about 5% of variceal hemorrhage and high degree of suspicion is necessary in cirrhosis with obscure bleeding.

**Term explanation**
Ectopic varices are porto-systemic collaterals occurring anywhere in the abdomen other than cardio-esophageal region. Orthotopic liver transplant is replacement of complete liver by donor liver in the normal anatomical position.

**Experiences and lessons**
This case report not only describes a very rare presentation of decompensated cirrhosis, but also managed successfully in a multi-disciplinary fashion utilizing multiple medical and surgical specialists.

**Peer review**
This article reported successful management of ectopic mesenteric variceal rupture and is an important diagnosis to consider in massive obscure bleeding in cirrhotic patients.

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