6-2007

Massive Retroperitoneal Hematoma Caused by Retroperitoneal Ectopic Pregnancy

Jay Goldberg
Thomas Jefferson University, Jay.Goldberg@jefferson.edu

Michael Weinstein
Thomas Jefferson University, Michael.Weinstein@jefferson.edu

Kris Kaulback
Thomas Jefferson University, Kris.Kaulback@jefferson.edu

Aileen Gariepy
Thomas Jefferson University

George Bega
Thomas Jefferson University, Gjergji.Bega@jefferson.edu

Let us know how access to this document benefits you

Follow this and additional works at: https://jdc.jefferson.edu/surgeryfp

Part of the Obstetrics and Gynecology Commons, and the Surgery Commons

Recommended Citation
Goldberg, Jay; Weinstein, Michael; Kaulback, Kris; Gariepy, Aileen; and Bega, George, "Massive Retroperitoneal Hematoma Caused by Retroperitoneal Ectopic Pregnancy" (2007). Department of Surgery Faculty Papers. Paper 61.
https://jdc.jefferson.edu/surgeryfp/61

This Article is brought to you for free and open access by the Jefferson Digital Commons. The Jefferson Digital Commons is a service of Thomas Jefferson University's Center for Teaching and Learning (CTL). The Commons is a showcase for Jefferson books and journals, peer-reviewed scholarly publications, unique historical collections from the University archives, and teaching tools. The Jefferson Digital Commons allows researchers and interested readers anywhere in the world to learn about and keep up to date with Jefferson scholarship. This article has been accepted for inclusion in Department of Surgery Faculty Papers by an authorized administrator of the Jefferson Digital Commons. For more information, please contact: JeffersonDigitalCommons@jefferson.edu.
CASE REPORT

Massive Retroperitoneal Hematoma Caused by Retroperitoneal Ectopic Pregnancy

Jay Goldberg, MD, MSCP; Michael Weinstein, MD; Kris Kaulback, MD; Aileen Gariepy, MD; George Bega, MD

A massive retroperitoneal hematoma caused by a retroperitoneal ectopic pregnancy is managed successfully utilizing multidisciplinary cooperation and transfusion of blood products. An ectopic pregnancy is defined as a gestation implanted in a location other than within the uterine cavity. While these most commonly occur in the fallopian tube, ectopic pregnancies may rarely occur in the cervix, ovary, or abdominal cavity. The authors report an unusual case of a retroperitoneal ectopic pregnancy presenting as a massive retroperitoneal hematoma in a hemodynamically unstable patient.

CASE REPORT

A 24-year-old woman (gravida 4, para 1021) presented to the emergency department at 8 weeks’ gestation by last menstrual period. She complained of increasing abdominal pain of 1 day’s duration. She was unaware that she was pregnant. She had no history of prior surgeries or sexually transmitted infections, and denied any recent trauma.

On presentation, the patient’s vital signs were initially stable. She was noted on examination to have significant abdominal pain with rebound and guarding but no palpable masses. Pelvic examination revealed a nongravid-sized uterus with no adnexal tenderness, adnexal enlargement, or cervical motion tenderness.

The patient’s serum human chorionic gonadotropin (hCG) level was 24,872 mIU/mL, and the hemoglobin value was 11.4 g/dL. Transabdominal and transvaginal ultrasonography showed a nongravid-sized uterus, with no evidence of an intrauterine gestation.

FIGURE 1.

Axial computed tomographic arteriogram showing a 20-×-15-cm hemorrhagic retroperitoneal mass below the left anterior perirenal space extending down into the upper pelvis.

Courtesy of Jay Goldberg, MD, MSCP.
and not bleeding into the peritoneal cavity. Due to continued hypotension and an intraoperative hemoglobin level of 5.5 g/dL, 4 units of packed red blood cells (RBCs) and 3 units of fresh frozen plasma (FFP) were transfused. At that point, it was decided that further exploration of the hematoma was neither necessary nor advisable, and the patient’s abdomen was closed.

Immediate postoperative computed tomographic arteriography of the abdomen and pelvis was performed, demonstrating a 20×15-cm hemorrhagic retroperitoneal mass below the left anterior perirenal space extending down into the upper pelvis (Figures 1 and 2). There was no evidence of pseudoaneurysm or active bleeding. An interventional radiologist was consulted, but the hematoma was not amenable to embolization.

Given the suspicion of an ectopic pregnancy, possibly retroperitoneal in location, a 90-mg intramuscular dose of methotrexate was administered based on her calculated body surface area. The plan was to closely monitor the patient, transfuse additional blood products as needed, and return to the operating room if she became unstable.

Approximately 20 hours later, the patient’s hemoglobin values had dropped from 9.6 to 6.8 g/dL and she again became hypotensive, so she was returned to the operating room for exploration of the retroperitoneal hematoma. A midline skin incision was made extending above the umbilicus. There was a significant amount of intra-abdominal blood arising from a small hole in the very tense retroperitoneal hematoma. The hematoma was opened with cautery, and approximately 2 L of clotted blood were removed. Bright red blood was oozing from a quarter-sized vascular area in the most superior portion of the hematoma cavity. Clips were applied to the vascular structures, resulting in hemostasis. No products of conception were grossly visible at the time of surgery. In total, from admission to completion of the second surgery, the patient received 12 units of packed RBCs and 8 units of FFP. Pathologic evaluation of the evacuated clotted blood revealed chorionic villi (Figure 3), confirming the suspected diagnosis of retroperitoneal ectopic pregnancy leading to a hematoma.

Pathologic evaluation of the evacuated clotted blood revealed chorionic villi (Figure 3), confirming the suspected diagnosis of retroperitoneal ectopic pregnancy leading to a hematoma.

Two days after evacuation of the retroperitoneal hematoma, the patient’s serum hCG level had decreased from 24,872 to 6,000 mIU/mL. Two days later, it had declined to 3,200 mIU/mL. Her postoperative course was complicated by intermittent fever and a mild ileus. These resolved without specific interventions. The patient was discharged 8 days after her initial
Microscopic image (200x) of degenerating chorionic villi identified in the evacuated retroperitoneal blood clot.

Courtesy of Jay Goldberg, MD, MSCP.

DISCUSSION
The management of retroperitoneal hemorrhage is based on hemodynamic parameters and secondary clinical findings. The majority of retroperitoneal hemorrhage is from venous sources and thus can be managed with expeditious correction of coagulopathy, volume replacement, and close monitoring. Patients who continue to bleed despite these efforts often have an arterial etiology and may benefit from angiography with embolization or stenting. Surgery to control bleeding in the retroperitoneum can be quite challenging, and is usually reserved for patients who do not respond to initial volume replacement or who have failed angiography. Rarely, a decompressive operation may be indicated when retroperitoneal bleeding causes femoral or lumbar plexus neuropathies or abdominal compartment syndrome.

Spontaneous retroperitoneal hematomas are uncommon. The majority of retroperitoneal hematomas are secondary to anticoagulation therapy. Other etiologies include neoplasm, arterial aneurysm, pseudoaneurysm rupture, and trauma.

Very few cases of retroperitoneal ectopic pregnancy have been reported. Following in vitro fertilization-embryo transfer (IVF-ET), a retroperitoneal ectopic pregnancy was hypothesized to have occurred via uterine perforation, with retroperitoneal placement of the transferred embryo. Reports in the absence of IVF-ET due to retroperitoneal implantation of an ectopic pregnancy. While rare, successful outcome of this life-threatening condition is predicated on multidisciplinary cooperation, clinical vigilance, and potentially massive transfusion of blood products.

REFERENCES