Unilateral spontaneous adrenal hemorrhage in a young patient

Muhammet Ferhat Çelik, M.D., Cevher Akarsu, M.D., Ahmet Cem Dural, M.D., Murat Çikot, M.D., Mustafa Gökhan Ünsal, M.D., Halil Alış, M.D.

Department of General Surgery, Bakirkoy Dr. Sadi Konuk Training and Research Hospital, İstanbul

ABSTRACT

The objective of this study was to report an unusual case of unilateral adrenal hematoma in a 19-year-old young man who did not have a history of any specific systemic disease. The patient was admitted to hospital with chest pain that lasted for one day. Preoperative contrast-enhanced computerized tomography evaluated an adrenal mass (sized, 10.5x12.7 cm) adjacent to the anterior of the left kidney, and findings were indicative of adrenal hematoma. The final pathological diagnosis was adrenal adenoma.

Key words: Adrenal gland; adrenal hematoma; adrenal hemorrhage.

INTRODUCTION

Adrenal hemorrhage (AH) can result from a variety of reasons. The large majority of patients with unilateral adrenal hemorrhage do not have clinically obvious signs of adrenal insufficiency, and diagnosis is usually made incidentally by the imaging performed for another reason. When unilateral, it is often clinically silent.[1]

Adrenal hemorrhage is associated with not only meningococcal septicemia but also disseminated intravascular coagulation (Waterhouse-Friderichsen syndrome); however, trauma, complications of pregnancy, tumors, surgical stress or anticoagulation therapy may also cause AH.[2] Primary adrenal cortical neoplasm rarely presents with spontaneous retroperitoneal hemorrhage. Nonetheless, spontaneous or idiopathic, AH is extremely rare in adults.[2]

This article reported the case of a young patient with spontaneous unilateral adrenal hemorrhage.

CASE REPORT

A 19-year-old man was admitted to hospital for acute chest pain and generalized weakness that continued for one day. There was no history of abdominal trauma, fever, hematuria, urinary symptoms or any specific systemic disease. On examination, the patient had tachycardia (120 beats per minute), but his blood pressure was normal. The patient had normal abdominal examination except for mild abdominal distension. Hematological investigation revealed severe anemia (hemoglobin 6.8 g/dL) with decreased hematocrit (23.2%) and neutrophilic leukocytosis (19800/mm³). Routine urine examination revealed 2 red blood cells and 10 pus cells/high power field (HPF). The coagulation profile as well as the serum amylase and lipase levels were normal. Abdominal ultrasound revealed diffuse free liquid into the peritoneal cavity. A computerized tomography (CT) scan of the abdomen was performed, which showed a left-sided perirenal solid mass and hematoma suggestive of renal origin in size of 10.5x12.7 cm (Fig. 1). The endocrinological examinations of the patient revealed Aldosterone 23.1 g/dL and Cortisol 34.6 µg/dL.

Preoperatively, six bags of erythrocyte suspension and four bags of fresh frozen plasma was transfused to the patient. An emergency operation was performed due to the worsening of the patient’s general condition on day three. A laparotomic left adrenalectomy and drainage of the hematoma was performed. The excised specimen contained an adenoma of the adrenal gland (Fig. 2). Postoperative recuperation was uneventful. Therefore, the final pathological diagnosis was adrenal adenoma. This study reported a case of spontaneous AH in a young man.
DISCUSSION

Common etiologies of spontaneous retroperitoneal hemorrhage in adults include trauma, coagulopathy or anticoagulation therapy, ruptured aneurysm of the aorta, splenic or renal artery and an arteriovenous malformation.[3] Spontaneous or idiopathic adrenal hemorrhage is extremely rare in adults. The incidence of spontaneous AH has been reported between 0.14% and 1.1%, usually involving the right gland.[4] Spontaneous hemorrhage can also occur in the retroperitoneum.

Adrenal hematomas have numerous and highly dissimilar radiographic appearances in CT. Magnetic resonance imaging is an alternative imaging modality for characterization of adrenal masses including adrenal hemorrhage.[5]

Since the 1990s, open adrenalectomy has been replaced with laparoscopic adrenalectomy (LA), and the implementation of minimally invasive surgery in adrenal gland surgery has gained significant momentum. Currently, LA has become a standard surgical method in many centers.[6]

AH rarely results from ruptured adrenal neoplasm. AH from a malignant primary adrenal neoplasm is also rare.[3] Adrenocortical carcinoma is a rare malignancy with an annual incidence of 1-2/million.[7] It is common in the fifth decade of life and in children under the age of 5.[8]

Baksi et al.[3] have presented a similar case with spontaneous retroperitoneal hemorrhage in a 21-year-old woman. Histological examination of the resected specimen has revealed an adrenocortical neoplasm.

Patients with hemorrhage usually do not present with hemorrhagic shock. Initially, they can be managed conservatively and investigated prior to surgery.

Our patient had no fever, neck stiffness, findings of meningitis and meningococcal septicemia; and therefore, these diseases were excluded.

Endocrinological examinations could not be fully carried out as an emergency surgery had to be performed in the preoperative preparation stage. In addition, it was initially thought as having renal origin. Possible differential diagnoses included a ruptured splenic or renal vascular aneurysm, an arteriovenous malformation or an adrenal vascular lesion. In this regard, angio-embolization of the bleeding adrenal vessels was reported.[9] Interventional radiology concluded that embolization was not suitable for the patient as the bleeding came from the mass. In a retrospective study of one hundred and forty-one patients of spontaneous AH admitted in the Mayo Clinic, College of Medicine from 1972 to 1997, sixteen patients presented with sudden abdominal pain and unilateral AH, and only seven required surgery in order to control the
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The possibility of AH should be kept in mind for acute abdominal pain, especially in patients with a diagnosis of a perirenal hematoma upon imaging. It can be concluded that in clinically stable patients, preoperative investigation and diagnosis can be performed in detail with a conservative approach.

Conflict of interest: None declared.

REFERENCES