

Spontaneous perirenal hematoma - foreshadowing of an impending renal cell carcinoma - does it need close surveillance? A case report and literature review

Hematoma perirrenal espontâneo - prenúncio de um carcinoma de células renais iminente - é necessária uma vigilância rigorosa? Relatório de caso e revisão de literatura

Ankit Gupta¹ , Abhishek G Singh¹, Ravindra B Sabnis¹

ABSTRACT

Renal cell carcinoma classically presents as a triad of hematuria, loin pain, and a palpable mass. Sometimes it may present as an incidental finding, whereas spontaneous renal hemorrhage or Wunderlich syndrome is a rare presentation of renal cell carcinoma. We present a clinical case of 51 years old gentleman who presented with right flank pain with no other significant history. Computed tomography scan showed right perirenal hemorrhage with no abnormality in the kidney. A drain was placed and the hematoma was evacuated. The patient was discharged. On follow-up at 6 months, he was asymptomatic but found to have a renal lesion on ultrasound, which was evaluated with computed tomography scan and partial nephrectomy was done and histopathology was suggestive of papillary renal cell carcinoma. Renal cell carcinoma being an aggressive tumor, any spontaneous perirenal hemorrhage with unknown etiology mandates a strict follow-up with prompt imaging for concerns of impending renal cell carcinoma.

Keywords: Hemorrhage; Renal cell carcinoma; Angiomyolipoma.

1. Muljibhai Patel urological hospital Nadiad Gujarat, Urology - Nadiad - Gujarat - India.

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Correspondence author: Ankit Gupta.

E-mail: ankitsurg@gmail.com

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RESUMO

O carcinoma de células renais apresenta-se classicamente como uma tríade de hematuria, dor lombar e massa palpável. Às vezes, pode se apresentar como um achado incidental, enquanto hemorragia renal espontânea ou síndrome de Wunderlich é uma apresentação rara de carcinoma de células renais. Apresentamos o caso clínico de um senhor de 51 anos que apresentava dor no flanco direito sem outros antecedentes significativos. A tomografia computadorizada mostrou hemorragia perirrenal direita sem anormalidades no rim. Um dreno foi colocado e o hematoma foi evacuado. A paciente recebeu alta. No acompanhamento de 6 meses, ele era assintomático, mas apresentava lesão renal na ultrassonografia, que foi avaliada com tomografia computadorizada e nefrectomia parcial e a histopatologia foi sugestiva de carcinoma papilar de células renais. Sendo o carcinoma de células renais um tumor agressivo, qualquer hemorragia perirrenal espontânea com etiologia desconhecida exige um acompanhamento rigoroso com exames de imagem imediatos para preocupações do carcinoma de células renais iminente.

Descritores: Hemorragia; Carcinoma de células renais; Angiomiolipoma.

INTRODUCTION

Spontaneous retroperitoneal hemorrhage (SPH) is a rare clinical entity that can occasionally be life-threatening. It poses a dilemma in diagnosis and management due to its varied clinical presentation. The common causes of non-traumatic causes of perirenal hemorrhage include renal cell carcinoma (RCC), renal angiomyolipoma (AML) vascular diseases, blood dyscrasia, and infections^[1-3]. Imaging plays a significant role in the diagnosis and management of this potentially lethal entity.

Herein, we present a clinical case of a patient with spontaneous perirenal hematoma with unclear etiology, who was treated with drainage of the hematoma; subsequently, he was diagnosed to have a renal mass at 6-month follow-up.

CASE REPORT

A 51-year-old male patient presented to outpatient department with complaints of right flank pain, since one day, which was sudden in onset, colicky, moderate in intensity and non-radiating in nature. Pain was constant over the last 6 hours and was associated with one episode of vomiting. There was no history of trauma, any bleeding tendency, hematuria, or stone disease. He did not have any previous history of a medical disorder, and presently, was not on any medication. He was a nonsmoker. On examination, his vitals were stable. Abdominal examination showed mild tenderness in the right flank. Bowel sounds were present; the abdomen was soft with no palpable mass. Ultrasonography of the abdomen was done which was suggestive of 106mm x 61mm x 71mm right perirenal heterogenous soft tissue lesion standing order hematoma (Figure 1).

Contrast-enhanced computed tomography scan (CT scan) abdomen and pelvis was done which was

suggestive of 15.2 x 10.6x 6.8cm right perinephric collection standing order hematoma (30-35HU [Hounsfield unit]) extending into the right pelvis. The post-contrast study does not show significant enhancement in the collection (Figures 2A and 2B). As the patient was symptomatic so ultrasound-guided drain was placed into the perinephric collection. The

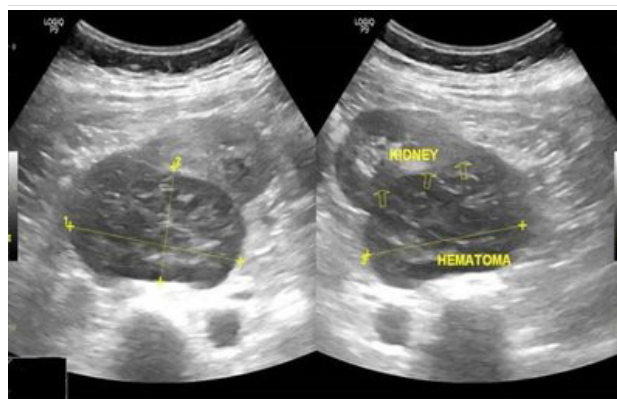


Figure 1. Ultrasound pictures showing the right kidney with perirenal hematoma (106mm x 61mm x 71mm).

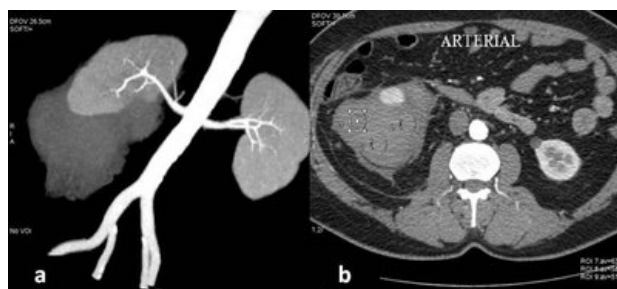


Figure 2. A. Computed tomography (CT) scan reconstructed image of right kidney with perirenal hematoma; B. Cross section CT image during arterial phase showing non-enhancing perirenal hematoma with no other obvious abnormality in the kidney.

drain output was 50-60ml per day, which comprised old clotted blood. Over a period of 7 days, the drain output decreased to 5-10ml/day, and the collection was also seen regressed on ultrasound, so the drain was removed and the patient was discharged.

The patient was followed up after one month and an ultrasound was done which showed minimal perinephric collection and the patient was asymptomatic. Thus, patient was advised to follow up after 6 months.

At 6 months follow-up, the patient was asymptomatic, on screening ultrasound, there was a suspicious lesion at the lower pole of the right kidney, so a plan was made to do a CT scan. CT scan was suggestive of a well-defined, iso to hypodense lesion seen at the posteromedial aspect of the mid and lower poles of the right kidney (Avg-36HU). It measured 7.5 x 6.2 x 6.1cm. The lesion showed an exophytic component with post-contrast enhancement (85HU) in some parts of the lesion (Figures 3A and 3B).

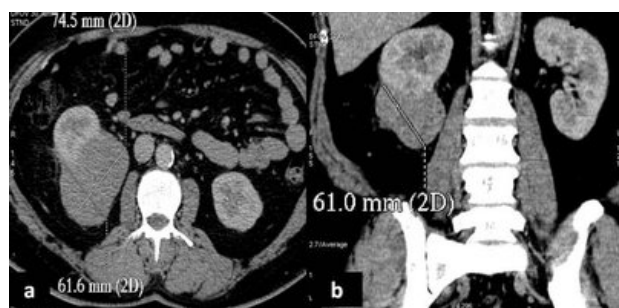


Figure 3. CT Scan image showing heterogeneously enhancing lesion in the right kidney measuring 74.5mm x 61.6mm x 61mm. A. Image showing transverse section; B. Image showing coronal section.

Trucut biopsy was taken from the lesion, histopathology was suggestive of fibro-adipocytic tissue with one tiny bit of renal tissue and malignant changes were not detected.

In view of the CT scan findings and inconclusive biopsy report, it was decided for surgical removal of the renal lesion. In view of the history of perirenal hematoma and previous intervention, the patient was taken up for right open partial nephrectomy.

Postoperatively the patient recovered well. Histopathological examination of the surgical specimen revealed papillary renal cell carcinoma (type II), nuclear grade "4, which was 7.4cm in largest dimension, and the lesion was assigned pathologic staging of pT2a Nx (Figures 4A and 4B). The patient was discharged in stable condition. The follow-up CT scan done 6 months post-surgery showed no signs of any recurrence.

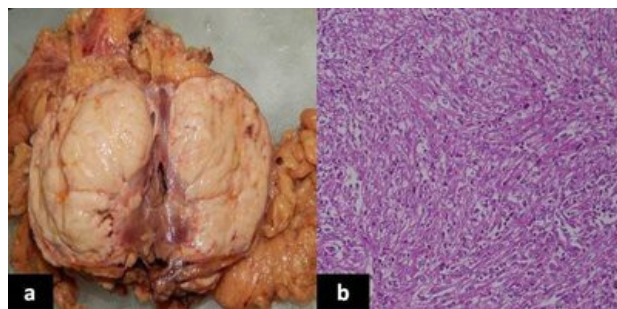


Figure 4. A. Cut section of the partial nephrectomy specimen; B. Low power microscopic image showing papillary fronds suggestive of papillary renal cell carcinoma.

DISCUSSION

In the present era, RCCs are more commonly discovered as incidentalomas on radiological imaging done for other abdominal ailments. This is due to the advancement and invention of newer imaging techniques. The usual presentation of RCC as flank pain, abdominal mass, and total hematuria are less commonly seen now a days. Dougal et al. (1975) discovered 78 documented cases of SPH since 1975, out of which 58% of cases were due to tumors, and the rest were due to benign causes. Similar findings were reported by Cinman et al. (1985) in 27 additional cases of SPH.^[2-5]

In 1956, Carl Reinhold August Wunderlich described spontaneous hemorrhage of the kidney with seepage of blood into the perinephric spaces.^[6] Mostly, Wunderlich syndrome is due to a benign cause.^[7] Wunderlich syndrome (WS) is a rare condition characterized by acute onset of spontaneous, nontraumatic renal hemorrhage into the subcapsular and perirenal spaces. Wunderlich syndrome is classically characterized by the Lenk's triad: acute flank pain, flank mass, and hypovolemic shock.^[8]

In a meta-analysis by Zhang et al. (2002), it was seen that more than 60% of cases of SPH were due to neoplasms.^[8,9]

The treatment of Wunderlich syndrome ranges from conservative management to immediate exploratory surgery. This depends upon the presentation of the patient and the choice of the operating surgeon. Few urologists are inclined towards early surgery in cases of SPH.^[9] There are various other ways of management which include interventional radiology-based embolization procedures to stop hemorrhage and stabilize the patient.^[10] These procedures may make further surgical resection difficult.

CT scan remains one of the best modalities to diagnose retroperitoneal hemorrhage and RCCs but there is a huge concern regarding the efficiency of CT to diagnose renal tumors when there is bleeding around and in the kidney. As reported by Kendall et al. (1988), an initial CT scan showed no signs of

malignancy in about 60% of the patients with SPH. This observation was also seen in our present case and is also supported by a case report by Aldughiman et al., in 2020.^[11-14]

If there is a case of SPH, it can be dealt with either by immediate radical nephrectomy as proposed by Kendall et al. (1988), if the opposite kidney is normal or the nephrectomy to be done in patients with non-fatty lesions except hematomas and rest patients to be followed up with imaging.^[12,13]

As a rule, if the etiology cannot be determined at the primary examination, we should perform a follow-up CT scan at 3-month intervals so that the hematoma can resolve and we can clinch a definitive diagnosis. Malignant renal masses diagnosed on CT scan can be managed by radical nephrectomy or partial nephrectomy. Whereas a benign renal bleed demands an angioembolization. Moreover, if these cases are diagnosed with a renal malignancy in the future, this will affect the staging and prognosis.^[13,14]

For patients having contrast allergy or renal sufficiency, magnetic resonance imaging (MRI) can be used for follow-up. Hashimoto et al. (2007) reported a case in their series who had a renal hemorrhage and the imaging both before and after the drainage of the hematoma showed no malignancy. The patient presented with hematuria after 1 year and was diagnosed with RCC in the same kidney.^[15] This case presentation was quite similar to our case.

Initially, it was advised to perform radical nephrectomy in all patients with spontaneous renal hemorrhage after excluding benign causes.^[12] But in the present era, newer advanced imaging modalities like multiphasic helical CT and MRIs have reduced the need for surgical exploration of patients with spontaneous renal hemorrhage (SRH). These modalities have also helped in better diagnosis and follow-up. In our case the tumor was a papillary RCC which is a comparatively slow-growing tumor as compared to other types, this is particularly important in tumor progression.

CARE reporting guidelines were used to prepare this manuscript.^[16]

CONCLUSIONS

Patients with spontaneous perirenal hemorrhage with unexplained causes should be followed up regularly with imaging in the form of CT scans so that an impending renal tumor can be identified at the earliest. In case of undiagnosed etiology at first presentation, imaging in the form of CT scan/MRI should be done early in follow-up at least at 2-3 months to clinch a definitive diagnosis.

CONFLICT OF INTEREST

There is no conflict of interest.

AUTHORS' CONTRIBUTIONS

AG	Collection and assembly of data, Conception and design, Manuscript writing
AGS	Final approval of manuscript, Provision of study materials or patient
RBS	Final approval of manuscript

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