



CASE REPORT

A Huge Chromophobe Renal Cell Carcinoma

Omar Jendouzi*, Melang Alexis Mvomo, Larbi Hamdoune, Mohamed Alami and Ahmed Ameur

Department of Urology, Military Teaching Hospital-Rabat, Morocco

*Corresponding author: Omar Jendouzi, Department of Urology, Military Teaching Hospital-Rabat, Morocco, Tel: +212669185144



Abstract

We report the clinical case of a 32-year-old female with no underlying disease who underwent in our structure a nephrectomy for a huge renal tumor. The histopathological finding revealed a chromophobe cell carcinoma and the evolution was favorable with no recurrence or metastasis.

Keywords

Huge tumor, Chromophobe carcinoma, Kidney

Chromophobe renal cell carcinoma accounts for 3.6 to 10.4% of renal cancers and corresponds to the third histological subtype in frequency, after clear cell carcinoma and papillary carcinoma. It has a good prognosis, it is most often limited to the kidney and has low grade nuclear [1,2].

We report the clinical case of a 32-year-old female with no underlying disease, admitted for management of isolated flank pain. Physical examination revealed right lumbar contact. An abdominal computed tomography (CT) showed a voluminous right renal tumor process (approximately 20 cm × 16 cm × 13.5 cm) with heterogeneous enhancement. There were no thrombi in the left renal vein. The patient underwent right radical nephrectomy through transperitoneal laparotomy and the exploration revealed a huge kidney tumor, which adherence was mild and it was removed surgically en bloc. The surgery took less than 2 hours, and the blood loss was 800 mL. The histopathological study was in favor of chromophobe carcinoma of the kidney, pT2 (TNM 1997), a grade 2 nuclear furhman grading (Figure 1).

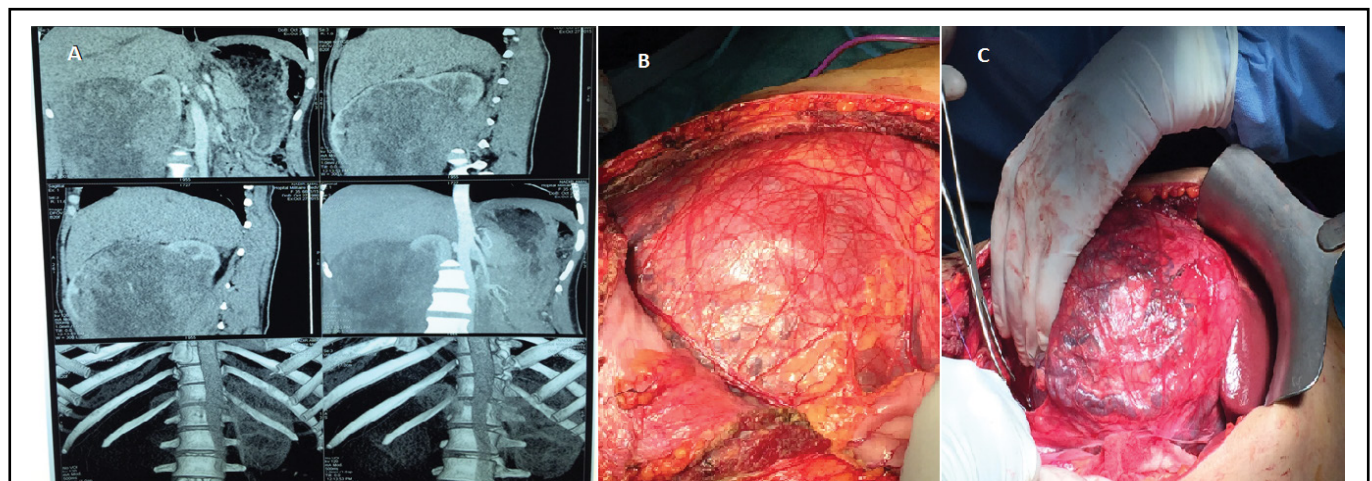


Figure 1: A) CT sections showing a large heterogeneous tumor mass depending on the right kidney. (B and C) Intraoperative images showing a huge encapsulated right renal tumor occupying the right hypochondrium.

There has been no recurrence or metastasis so far (12 months after surgery) and no adjuvant therapy was realized.

References

1. Capitanio U, Cloutier V, Zini L, Isbarn H, Jeldres C, et al. (2009) A critical assessment of the prognostic value of clear cell, papillary and chromophobe histological subtypes in renal cell carcinoma: A population-based study. *BJU Int* 5: 1-5.
2. Amin MB, Pauner GP, Alvarado-Caberero I, Young AN, Stricker HJ, et al. (2008) Chromophobe renal cell carcinoma: histopathologic characteristics and evaluation of conventional pathologic prognostic parameters in 145 cases. *Am J Surg Pathol* 32: 1822-1834.