Images — Isolated caval recurrence of renal cell carcinoma

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Introduction

Renal cell carcinoma (RCC) is unique in that it has a biological propensity for vascular invasion. Approximately 10% of RCC cases present with venous tumour thrombus involving the renal vein or inferior vena cava (IVC)^{1,2} and 1% with a thrombus extending into the right atrium.³ Isolated caval recurrence of RCC following radical nephrectomy, however, is a rare event. Although systemic treatment may have a role in managing patients with distant metastases, in the case of local recurrence, surgical intervention remains the only effective treatment.³

Resection and reconstruction of the IVC for primary or recurrent malignancy is performed infrequently due to the prevalence of concurrent metastasis and mortality risks. Despite this, aggressive management may be associated with longer disease-free survival.4 Recurrence in the vena cava is important because surgical intervention is technically challenging, requiring cardiopulomary bypass (CPB) and deep hypothermic circulatory arrest (DHCA) if the thrombus extends above the hepatic veins and into the right atrium.⁵ Beyond this, the presence of a venous thrombus is associated with risks such as emboli and venous congestion.3 There is little information regarding operative management of recurrent tumour thrombi due to the rareness of its occurrence. We present a case of RCC with isolated caval recurrence 15 years following a radical nephrectomy. Surgical resection of the thrombus with CPB and DHCA was successfully performed.

Case report

A 52-year-old male presented initially with gross hematuria and right lower quadrant pain radiating to his testicles. A computed tomography (CT) scan of the abdomen and pelvis described a 12 cm right renal mass with tumour invasion into the renal vein. There was no evidence of metastatic

disease. The patient subsequently underwent a right radical nephrectomy with a renal vein tumor thrombectomy. The tumour invaded the right renal vein and venous wall. The tumor was also seen grossly invading into Gerota's fascia. The IVC appeared patent. Pathology described a 14 x 11 x 11 cm right renal mass, which was well- to moderately differentiated RCC, clear-cell, with some papillary architecture. Following five years of followup without recurrence of disease, the patient was discharged back to his family physician.

Ten years later, the patient presented to the emergency department complaining of a few weeks of right lower quadrant pain. He was otherwise well. An abdominal ultrasound identified a large thrombus in the upper IVC, approximately 11.5 cm in length (arising around the location of the right renal vein). The thrombus appeared to change in position with the cardiac cycle. Doppler studies demonstrated flow within and around the thrombus. A CT scan confirmed presence of the IVC mass extending to the intrahepatic IVC. Extension into the right atrium could not be determined. It also demonstrated a 4.5 x 2.0 cm mass posterior to the IVC in the renal fossa. A magnetic resonance imaging (MRI) angiogram (Fig. 1) and 3D reconstruction demonstrated thrombus to the level of the right atrium (Fig. 2).

Taking into account the new caval recurrence, the patient was taken for a laparotomy, median sternotomy, and CPB with DHCA. The IVC thrombus, renal vein stump, and paracaval mass were removed en bloc (Fig. 3). A 20 mm polytetrafluoroethylene (PTFE) graft was placed in the caval defect because of the circumferential caval wall invasion.

Frozen section taken during the operation showed clear-cell variant RCC. Final pathology of the posterior caval mass and caval thrombus demonstrated: clear-cell, Fuhrman grade II/IV, invaded skeletal muscle fibers, indicating likely invasion of the psoas and/or paraspinal muscles posterior to the IVC. Paracaval lymph nodes were negative for RCC. The procedure was well-tolerated and he recovered appropriately. Subsequent followup CT scans of the abdomen and pelvis showed no evidence of recurrence at 60-month followup. Metastatic workup was also negative at 60 months. Ultrasound demonstrated patency of his graft, without thrombus.

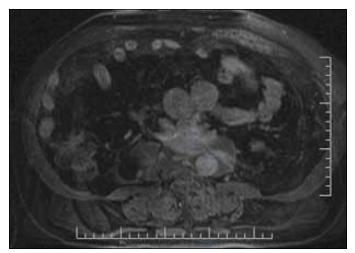


Fig. 1. A magnetic resonance imaging angiography demonstrating tumour thrombus to the level of the right atrium.

Discussion

There are very few documented cases in the literature surrounding isolated caval recurrence of RCC following nephrectomy. The results of our literature review demonstrated that surgical management of the recurrent tumour thrombus is complex and often multidisciplinary. Other services involved included cardiothoracic surgery, general surgery, hepatobiliary surgery, and vascular surgery. Compared with prior reports, our case demonstrated different findings with respect to the timing of recurrence, where the majority of previous experiences have shown a range of 6–48 months from nephrectomy to the development of recurrent tumour thrombus.

In RCC patients with isolated caval recurrence, several surgical methods have been discussed. Regardless of the surgical intervention, most reports emphasized technical difficulty. Given the involvement of tumour thrombus invading up to the level of the right atrium, we felt a combined approach — including open midline incision, median sternotomy, and CPB with DHCA — would most likely allow successful removal of the tumour and, importantly, minimize intraoperative complications.

Conclusion

While RCC is the sixth and eleventh most common cancer diagnosed in men and women, respectively,⁶ concomitant RCC with isolated caval recurrence following nephrectomy is incredibly rare. Only 26 cases have been reported. Surgery is challenging and usually requires caval replacement and CPB with DHCA if thrombus is extending into the right atrium. Although our experience may justify close long-term followup surveillance for RCC patients with T3 disease and venous wall invasion, this may be more a consequence of the biology of



Fig. 2. A magnetic resonance imaging with 3D reconstruction demonstrating tumor thrombus extending to the intrahepatic inferior vena cava.

this specific cancer and it is, therefore, difficult to make any inferences about management for cases going forward.

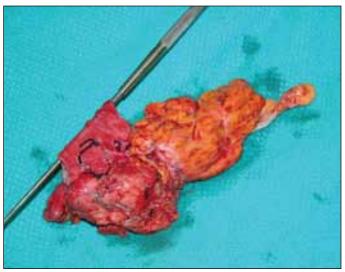


Fig. 3. Inferior vena cava tumour thrombus, renal vein stump, and paracaval mass. Probe transverses the vena cava.

Competing interests: The authors report no competing personal or financial interest related to this work.

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