

Case Report

Uncommon Pancreatic Metastasis 10 Years After Nephrectomy in Clear Cell Renal Carcinoma

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Introduction

Adult kidney cancer represents approximately 3% of all malignancies and ranks third among urological cancers, following prostate and bladder cancers. While the most common sites for metastasis include the lungs, bones, liver, and brain, pancreatic metastases are relatively rare. This report presents a case of pancreatic metastasis from renal cell carcinoma (RCC), which is an uncommon occurrence [1].

Case Report

A 56-year-old female patient with a history of clear-cell Renal Carcinoma (RCC) treated 10 years earlier presented with progressive weight loss and epigastric pain over the past week. A contrast-enhanced CT scan was performed to further evaluate the symptoms.

Imaging revealed a large, rounded mass with lobulated contours located in the head of the pancreas. The mass demonstrated significant enhancement following contrast administration (**Figure 1, 2**), measuring $67 \times 51 \times 37$ mm (height x width x anterior-posterior). Notably, the mass caused dilation of both the extra- and intra-hepatic bile ducts, as well as the pancreatic duct (Wirsung's duct), with a corresponding atrophy of the downstream pancreas. Additionally, a necrotic, heterogeneous mass was observed in the left adrenal gland.

The right nephrectomy bed was unremarkable, with the contralateral kidney appearing normal (Figure 3). Ultrasound and biopsy were performed, confirming the metachronous nature of the pancreatic mass as originating from the prior renal carcinoma (Figure 4, 5). Subsequently, the patient underwent surgical resection of the pancreatic mass with curative intent.

Discussion

Clear cell renal carcinoma typically metastasizes to the lungs, bones, and liver. Pancreatic metastases from RCC are rare, with a latency period of several years following nephrectomy, typically ranging from 10 to 27 years [2-3]. The spread to the pancreas may occur through lymphatic pathways, connecting

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Figure 1: Axial CT scan showing a mass in the head of the pancreas (C-phase) with lobulated contours.

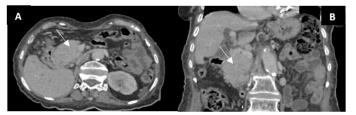


Figure 2: (A) Axial CT scan revealing a hypervascular mass in the pancreas; (B) Coronal CT section demonstrating the hypervascular mass in the pancreatic head.

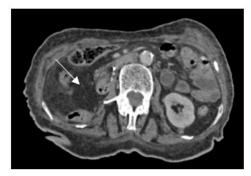


Figure 3: Axial CT scan illustrating the empty right nephrectomy bed.

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Figure 4: Ultrasound showing a heterogeneous, hypoechoic mass in the head of the pancreas.

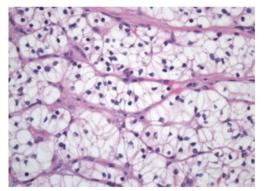


Figure 5: Histopathological image of the clear-cell adenocarcinoma in the pancreatic tissue.

the pancreas to the renal arteries [4], or via venous routes, particularly through portocaval shunts associated with renal tumors [5].

While many patients with pancreatic metastasis from RCC remain asymptomatic, common clinical presentations include abdominal pain (often epigastric), jaundice, weight loss, palpable masses, and, in some cases, duodenal invasion leading to gastrointestinal hemorrhage [6].

Radiologically, pancreatic metastases from RCC can present as hypodense lesions with well-defined borders and intense, heterogeneous enhancement after contrast administration [7]. These lesions may also contain hypodense areas, which is a characteristic finding on CT imaging. On ultrasound, such lesions are typically hypoechoic, although cystic forms can occasionally be observed [7]. The differential diagnosis includes pancreatic neuroendocrine tumors, which require careful consideration in imaging [8].

The definitive diagnosis is usually made through histopathological examination following surgical resection or biopsy, as was performed in this case [9,10]. Surgical management is the treatment of choice for solitary pancreatic metastases from RCC. Common procedures include cephalic duodenopancreatectomy or left-sided pancreatectomy. In cases of multiple pancreatic metastases, subtotal pancreatectomy may be necessary [11]. The prognosis following surgical resection of pancreatic metastases from RCC is generally favorable, with reported 5-year survival rates ranging from 34% to 88%.

Conclusion

Pancreatic metastases from clear cell renal carcinoma are rare but may manifest several years after nephrectomy. Surgical resection remains the most effective treatment modality, offering the potential for improved survival and quality of life.

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Informed consent: Written informed consent was obtained from a legally authorized representative(s) for anonymized patient information to be published in this article.

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