

Metachronic Pancreatic Metastasis of Renal Clear Cell Carcinoma

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Abstract

Case Report

Pancreatic metastases from kidney cancer are rare and generally late onset. They are most often detected by an abdominal computed tomography performed as part of kidney cancer surveillance. We report the observation of a 50-year-old diabetic patient who had a nephrectomy for a clear cell renal adenocarcinoma seven years ago, during follow-up, abdominal CT and nuclear magnetic resonance imaging (MRI) revealed two heterogeneous pancreatic nodules, the kinetics of enhancement resembles that of the primary tumor, and pathology has in fact revealed metachronic pancreatic metastases of his clear cell renal adenocarcinoma.

Keywords: Pancreatic metastases, kidney cancer, imaging.

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INTRODUCTION

The secondary locations of renal clear cell carcinomas are most often pulmonary, hepatic or bone. Pancreatic locations are exceptional and represent, depending on the series, 2 to 4% of pancreatic malignant lesions. We report the observation of a pancreatic tumor secondary to a renal clear cell adenocarcinoma which appeared seven years after nephrectomy.

OBSERVATION

This is a 50-year-old diabetic (type II) patient who had seen for gross macroscopic hematuria that had been progressing for a week. An emergency ultrasound followed by an abdominal computed tomography (CT) revealed a tissue mass of 50 mm at the expense of the lower pole of the right kidney. The extension assessment, including chest CT and bone scan, was negative.

An enlarged right nephrectomy was performed. Pathology has confirmed the presence of a clear cell renal adenocarcinoma with invasion of perirenal fat and neoplastic thrombus in the right renal vein performing a pT3N0 stage according to the TNM classification. Seven years later, during follow-up, abdominal CT and nuclear magnetic resonance imaging (MRI) revealed two heterogeneous pancreatic nodules. One was 14 mm and was in the isthmus position (Figure 1), the other was 20 mm and was located in the more caudal position (Figure 2). The hypervascular character

of these nodules immediately oriented towards metastases of his kidney cancer more than towards a primary pancreatic tumor. The patient then benefited from a splenopancreatectomy enlarged to the isthmus to perform the excision of these two nodules. Pathology has in fact revealed three pancreatic metastases of his clear cell adenocarcinoma. Three months later, an abdominal CT scan showed a 20 mm nodule in the incinatus process which had not been detected until now. The patient benefited from a totalization of his pancreatotomy to perform the removal of this nodule which was also a metastasis of his kidney cancer. Post-operative follow-ups were simple and insulin therapy was started. Three years after his pancreatotomy, the patient was in good general condition and free from any tumor recurrence.

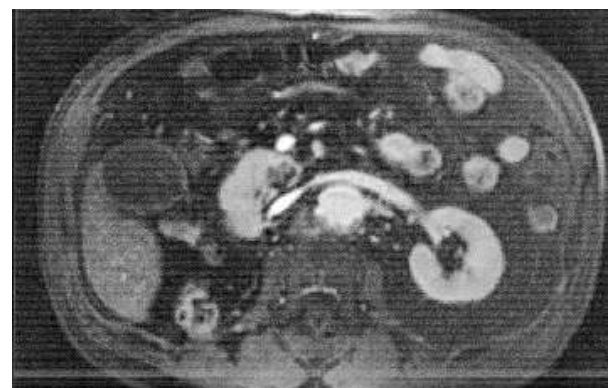


Fig- 1: Abdominal MRI. T1 acquisition after Gadolinium injection. Cross section through the head of the pancreas. The hypervascular metastasis is well visualized at the level of the incinatus process



Fig-2: Abdominal MRI. T1 acquisition after Gadolinium injection. Cross section through the tail of the pancreas. Metastasis is well visualized

DISCUSSION

Pancreatic metastases from kidney cancer are rare and appear late. Very few cases have been reported in the literature [1, 3, 4, 8, 11, 12]. The sex ratio is 1.7 with a male predilection, which can be superimposed on that of the epidemiology of kidney cancer [9]. At diagnosis, the average age is 61 years with extremes ranging from 39 to 76 years in the literature [9]. The average delay between nephrectomy and the appearance of metastases is long with an average of 10 years but they can appear earlier or even synchronous with the primary renal tumor.

The clinical manifestations of pancreatic metastases are not very specific [7]. They can reveal themselves by abdominal pain, a table of deterioration of the general state with slimming, an exocrine or endocrine pancreatic insufficiency, an acute pancreatitis even a compression of vicinity (stomach, vessels, and duodenum) responsible for digestive hemorrhage. However, due to their slow growth, metastases are often asymptomatic and therefore discovered incidentally in the context of surveillance [11].

Abdominal ultrasound hardly explores the pancreatic gland and has no specificity for the diagnosis of secondary lesions; the lesions appear in the form of hypoechogenic solid masses located in the pancreatic parenchyma [3]. Doppler ultrasound has a better sensitivity since it will highlight the hypervascular character of the lesion [2]. The abdominal CT shows a well circumscribed mass which is iso or hypodense compared to the normal pancreas on the sections without contrast. Enhancement is generally heterogeneous, but tends to be homogeneous in small lesions and peripheral in larger lesions, probably due to central necrosis, in general, the kinetics of enhancement resembles that of the primary tumor [2-4]. Calcifications are rare and pancreatic duct obstruction is common for lesions of the head and body, seen in about 40% of cases, and can be associated with obstruction of

the main bile duct and intrahepatic biliary dilation. The hypervascularized nature of these metastases manifests itself in the stereotypes by a lesion which takes the contrast unlike the primary tumors of the pancreas, which are hypovascularized [2, 3, 5].

However, the puncture is justified if no primary tumor is known, if the appearance of the pancreatic lesion suggests another histology (in the case of a hypervascular pancreatic mass, the main differential diagnosis is that of endocrine tumor) or if the lesion is not resectable in order to obtain histological evidence of the pancreatic lesion [7, 11, 13].

Secondary pancreatic locations for kidney cancer appear to be of lymphatic origin. Indeed, a study carried out by Nagakawa by injection of radiolabelled carbon in patients with cancer of the head of the pancreas shows that the lymph node extension extends to behind the renal arteries [10]. Thus, a lymph node invasion along the renal pedicle can explain the pancreatic involvement. Another possible mode of spread is the hematogenous pathway. The existence of arteriovenous shunts within the tumor of the kidney leads to a decrease in vascular resistance with as a consequence an ectasia of the peri-tumoral and perirenal drainage veins. The venous pressure in this exorenal system being higher than that of the adjacent structures, large anastomoses are created with the adrenal veins and the portal system. Thus and especially if there is a thrombosis of the renal vein, neoplastic emboli can migrate in a reno-portal shunt [9].

The treatment of pancreatic metastases is surgical [12]. Pancreatic resection is most often major, the type of which depends on the topography of the lesion [6, 11]. A cephalic duodenopancreatectomy or a splenopancreatectomy are most often necessary [13]. Hernandez reports the case of a patient who underwent a laparoscopic caudal pancreatectomy in this indication [5]. The multiple and diffuse nature of secondary pancreatic lesions can lead to total pancreatectomy as is the case in our observation. Some authors opt for a pancreatic resection limited to the tumor by performing atypical pancreatectomies in order to preserve as much as possible the healthy parenchyma under cover of an extemporaneous examination of the section sections [16]. This attitude does not prevent the occurrence of pancreatic fistula and does not seem to modify either the frequency of recurrences or the survival of patients [11,16].

Chemotherapy with vinblastine or immunotherapy with interferon alpha have not proven to be effective in the treatment of pancreatic metastases of kidney tumors, but they may nevertheless be useful after excisional surgery with lymph node involvement or minimal tumor residue [11, 15].

Survival after surgical resection of pancreatic metastases is satisfactory with a median survival of 30 months. For Tuech, the 5-year survival rate is 31% in the case of metastases localized only to the pancreas [14, 17].

The good prognostic factors for these pancreatic metastases appear to be the late onset of a pancreatic location, an asymptomatic form, a single pancreatic lesion or a radiological appearance of central necrosis of the tumor lesion [11].

CONCLUSION

Pancreatic metastases from kidney cancer are rare and generally late onset. They are most often detected by an abdominal computed tomography performed as part of kidney cancer surveillance. Early diagnosis makes it possible to attempt a curative surgical gesture (pancreatic resection adapted to tumor location) which seems to be the only treatment capable of prolonging the survival of patients.

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