CASE REPORT

Isolated Pancreatic Tuberculosis Mimicking Focal Pancreatitis and Causing Segmental Portal Hypertension

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ABSTRACT

Context Isolated pancreatic tuberculosis is a common mimicker of pancreatic malignancy and a common presentation is abdominal pain. However, segmental portal hypertension is very uncommon. Case report A case of isolated pancreatic tuberculosis mimicking focal pancreatitis and causing segmental portal hypertension is presented. Conclusion A histological or microbiological diagnosis of the presence of pancreatic masses is necessary to detect potentially treatable diseases, such as tuberculosis, in endemic countries like India.

INTRODUCTION

Pancreatic tuberculosis is very rare and it closely resembles a pancreatic malignancy, both clinically and radiologically [1]. Therefore, most cases of pancreatic tuberculosis have been diagnosed after surgery carried out for a suspected malignancy [2]. However, improvement in imaging techniques and the resulting image-guided interventions have helped us in the preoperative diagnosis of pancreatic masses. We present a rare case of isolated pancreatic tuberculosis that mimicked focal pancreatitis and caused segmental portal hypertension.

CASE REPORT

A 40-year-old female presented with abdominal pain of three months duration. The pain was located in the upper abdomen with radiation to the back. There were no exacerbating factors and the pain subsided with the ingestion of oral opioids. The patient also complained of weight and appetite loss. There was no history of alcohol ingestion. Her laboratory studies revealed an elevated erythrocyte sedimentation rate (46 mm in first hour; reference range: 0-20 mm) and were otherwise unremarkable. Serum amylase was 320 IU/L (reference range: 0-160 IU/L). A chest radiograph did not reveal any abnormality. Ultrasound of the abdomen did not reveal any abnormality and the pancreas was obscured by the bowel gases. Contrast-enhanced computerized tomography (CECT) of the abdomen demonstrated the bulky head and body of the pancreas with heterogeneous areas of non-enhancement suggestive of necrosis and peripancreatic fat stranding (Figure 1). The splenic vein was not visualized which was suggestive of thrombosis. The CECT findings were suggestive of focal pancreatitis involving the head and body of the pancreas with splenic vein thrombosis. As the patient had no history of severe pain which required admission, had ongoing chronic pain with loss of weight and appetite and a bulky head and body of the pancreas, image-guided fine needle aspiration was carried out to exclude a pancreatic malignancy.

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Figure 1. Contrast-enhanced computerized tomography (CECT) showing the bulky head and body of the pancreas with heterogeneous areas of non-enhancement suggestive of necrosis and peripancreatic fat stranding.
cytological examination of the CT-guided fine needle aspiration revealed caseous necrosis. No acid-fast bacilli could be demonstrated. The Mantoux test was negative. However, a culture of the aspirated material demonstrated the growth of *Mycobacterium tuberculosis*. The patient was treated with antitubercular therapy for twelve months (four drugs for two months and two drugs for ten months). The patient improved with the therapy and was asymptomatic after completing two months of therapy. A repeat CECT of the abdomen performed after completion of the therapy revealed a normal pancreas with no focal lesions (Figure 2). The splenic vein was attenuated and collaterals were noted at perigastric, perisplenic and gastrohepatic locations (Figure 3). No varices were noted on upper gastrointestinal endoscopy. After four months of follow-up, the patient is asymptomatic.

**DISCUSSION**

Abdominal tuberculosis includes the infection of varying combinations of the intestinal tract, peritoneum, lymph nodes and solid organs, such as the liver, spleen, and pancreas. Involvement of solid abdominal organs is usually seen in association with miliary tuberculosis. Isolated abdominal organ involvement, especially of the pancreas, is unusual, even in the setting of miliary disease (ranging from 2.1 to 4.7%) [3, 4]. Postmortem analysis of 300 miliary tuberculosis patients over twelve years in India did not reveal even one with pancreatic involvement [3]. This is probably due to the antibacterial effect of the pancreatic enzymes [1].

Pancreatic tuberculosis usually affects young adults having a mean age of approximately 40 years and with equal sex distribution. Higher incidences have been observed in endemic areas and in the immunocompromised population. It has been speculated that tuberculous involvement of the pancreas might occur as a result of direct extension, lymphohematogenous dissemination, and reactivation of a previous abdominal focus or immune reaction to generalized tuberculosis [5, 6].

The clinical and radiological findings of pancreatic tuberculosis usually mimic pancreatic malignancy as a result of common features such as weight loss, anorexia, abdominal pain, jaundice and a pancreatic mass [2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15]. Both tend to occur in the head and uncinate process, probably due to the rich dual vascular supply [7]. Other rare presentations, such as acute pancreatitis, portal hypertension, gastrointestinal bleeding, diabetes mellitus and obstructive jaundice have also been described in the published literature [2, 8, 9, 10, 11, 12, 13, 14, 15]. A case report of pancreatic tuberculosis presenting as a head mass along with a pancreaticobiliary fistula has also been reported [14]. The Mantoux test may also be negative because of poor nutritional status leading to a weak immune response as happened in our case. The imaging findings may suggest the possibility of tuberculosis, but none of the findings are pathognomonic for pancreatic tuberculosis. The CT findings include hypodense lesions and irregular borders usually in the head of the pancreas, diffuse enlargement of the pancreas or enlarged peripancreatic lymph nodes. The presence of hypodense lymph nodes with rim enhancement in the peripancreatic region, ascites and/or mural thickening affecting the ileocecal region may suggest the possibility of tuberculosis [16]. MRI findings of focal pancreatic tuberculosis include a sharply delineated mass located in the pancreatic head, showing heterogeneous enhancement. These lesions usually are hypointense on fat-suppressed T1-weighted images and show a mixture of hypo- and hyper-intensity on T2-weighted images [17]. The common bile duct and the pancreatic duct have been reported to be normal in patients with pancreatic tuberculosis, even if the tuberculous mass is centrally positioned in the pancreatic head [17, 18]. This is in contrast to pancreatic adenocarcinoma where the pancreatic duct is dilated in centrally located tumors in the head region. The diffuse form of pancreatic tuberculosis is characterized by pancreatic enlargement with narrowing of the main pancreatic duct and...
heterogeneous enhancement [17]. On ERCP, bile cytology also shows poor results in diagnosing tuberculosis, although it may occasionally help in establishing the diagnosis [15, 19]. Since there are no clinical, laboratory or radiological features which are specific for pancreatic tuberculosis, histopathological or cytological as well as bacteriological confirmation is necessary for establishing the diagnosis of isolated pancreatic tuberculosis. Most of the cases in the literature have been diagnosed after laparotomy for a suspected pancreatic malignancy. Percutaneous imaging or endoscopic ultrasound-guided fine needle aspiration of the pancreatic lesion have been reported for establishing a diagnosis of pancreatic tuberculosis [20, 21, 22, 23]. Because of the rarity of this disease, there are no specific treatment guidelines. The majority of cases of pancreatic tuberculosis respond well to 6-12 months of anti-tubercular therapy and their prognosis is good [24].

In conclusion, isolated pancreatic tuberculosis is a rare entity and can be clinically elusive. Fine needle aspiration cytology and definitive biopsy are the primary techniques used for reaching a definitive diagnosis and avoiding surgery in people living in an endemic area who present with pancreatic mass lesions.

Conflict of interest The authors have no potential conflict of interest

References