

Features of Recurrent Bowel Obstruction with Nonobstructive Duodenal Neuroendocrine Tumor

Authors: Ruchi Sharma, MD¹; Hammad Zafar, MD¹; Scott K. Sherman, MD²; Fadi Niyazi, MD³

Organizations/Affiliations: ¹ Department of Internal Medicine, University of Iowa Hospital and Carver College of Medicine

²Department of Surgical Oncology and Endocrine Surgery, University of Iowa Hospital and Carver College of Medicine

³Department of Gastroenterology and Hepatology, University of Iowa Hospital and Carver College of Medicine

Introduction

Neuroendocrine tumors (NET) are a rare cause of bowel obstruction which is usually mechanical in nature. We present an unusual case of bowel obstruction in a patient with NET where no mechanical obstruction was found despite extensive work up.

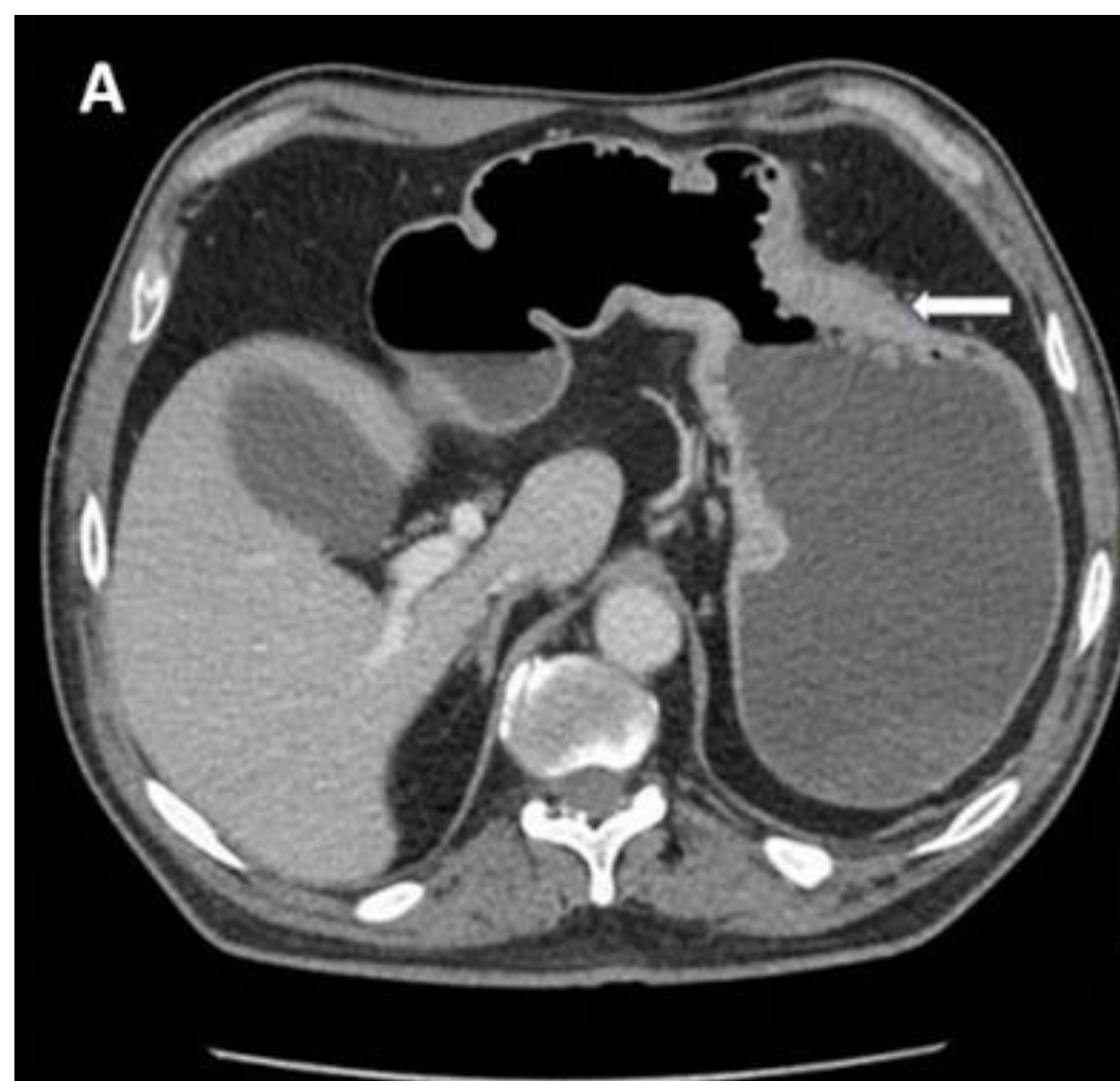


Figure 1 (A): CT abdomen with dilated stomach and duodenum with decompressed distal small intestine (arrow)

Case Description

A 56-year-old male was admitted after 9-10 admissions to another facility with bowel obstruction and 50-pound weight loss over 10 months. His symptoms included abdominal pain, distention, and vomiting. He had no prior abdominal surgeries. Previous CT abdomen revealed small bowel obstruction with transition point in the terminal ileum. Current CT showed gastric outlet obstruction (Figure 1A). Formerly he was managed nonoperatively with nasogastric tube (NGT) decompression. Prior esophagogastroduodenoscopy and colonoscopy did not reveal an obstruction. NGT output was 2.5L. Push enteroscopy, video capsule endoscopy did not reveal any mechanical obstruction. MR enterography showed two right mesenteric nodules, larger being 2.1 x 2.3 cm in size. These were moderately FDG avid on PET-CT. Endoscopic ultrasound biopsy revealed a well differentiated NET. The lesions were intensely somatostatin receptor positive on Gallium-68 DOTATOC PETCT (Figure 1B). Pancreatic polypeptide was 2145pg/mL(0-435), chromogranin A 1458ng/mL(0-103), serotonin 278ng/mL(50-200), and gastrin 208ng/mL(0-100). He underwent partial duodenal resection with primary repair and peripancreatic lymph node resection. No obstructive pathology was found intraoperatively. Histopathology confirmed well differentiated NET with metastatic peripancreatic nodes. He has had no admissions over the past 3 months and has gained 33 pounds.

Discussion

NET are rare tumors and use of appropriate diagnostic modalities are key to timely diagnosis. Despite improvement and availability of diagnostic modalities, most patients have metastatic disease at diagnosis. NET presenting with bowel obstruction usually have an obstructing tumor. We wish to bring to light the unusual presentation of NET with mechanical obstruction on imaging but extensive negative work up for an actual obstructing lesion. It is possible that mesenteric inflammation and edema secondary to NET may have caused recurrent obstruction in this case. NET can present with mechanical obstruction and Gallium-68 DOTATOC PETCT may be a reasonable investigation in a patient with unexplained recurrent gastrointestinal obstruction.

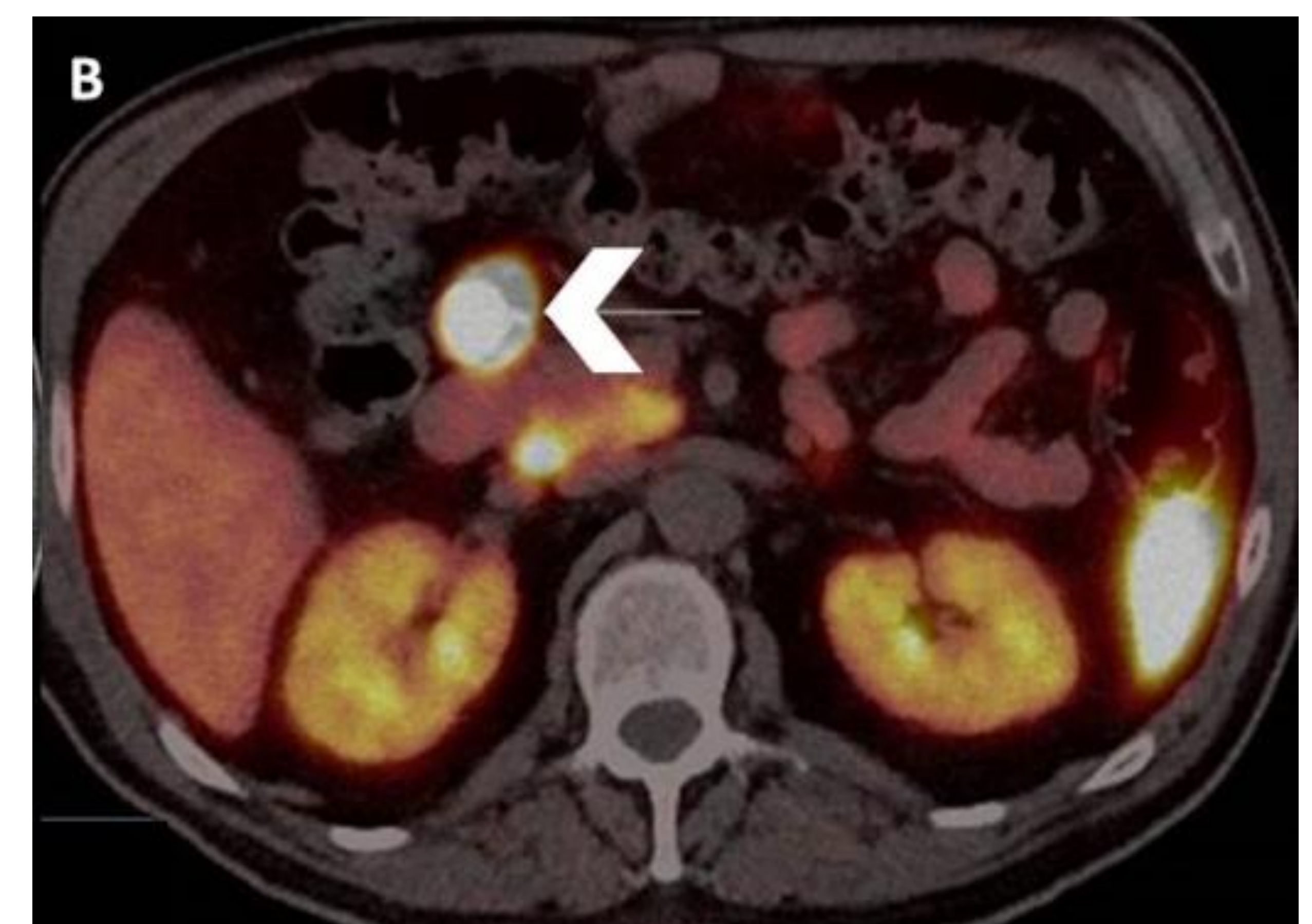


Figure 1(B): Ga-68 DOTATOC PET/CT scan showing somatostatin receptor-positive peripancreatic node (chevron)