Solid Pseudopapillary Neoplasm of the Pancreas Presenting With Gastric Outlet Obstruction

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Abstract

Solid pseudopapillary neoplasm (SPN) of the pancreas are rare tumors accounting for a minor portion of all exocrine pancreatic tumors. It usually occurs in young women. It has a very low malignant potential with a relatively indolent clinical course. A small subset of patients exhibits pathological features of malignancy. Herein, we present a rare case of pancreatic SPN that presented with gastric outlet obstruction. Despite the characteristic computed tomography (CT) findings, due to its rarity, it was missed in more common conditions such as gastrointestinal stromal tumors. In our case, we found that the tumor was causing extrinsic duodenal compression leading to gastric outlet obstruction, creating a diagnostic dilemma.

Categories: General Surgery

Keywords: outcome, surgery, small intestinal obstruction, cystic neoplasm of the pancreas, solid pseudopapillary neoplasm

Introduction

Pancreatic solid pseudopapillary tumor (SPT) is a rare neoplasm, usually characterized by a well-encapsulated mass, with very low malignant potential. It is predominantly seen in young females, with a female/male ratio of 9.5:1 [1]. It was first described by Frantz in 1959 [2]. The first surgical resection of this neoplasm was performed by Grosfeld, and the electron microscopic appearance was first described by Hamoudi et al. in 1970 [3].

These are often clinically asymptomatic. Due to its indolent course and low malignant potential, mostly, it presents with gradually enlarging abdominal mass or nonspecific abdominal discomfort [4,5]. On palpation, the abdomen is usually non-tender but may become tender if obstructive symptoms occur in cases of large lesions compressing the adjacent viscera. SPT can be visualized in many imaging modalities, such as ultrasonography (USG), computed tomography (CT), and magnetic resonance imaging (MRI), which can be used to differentiate it from other pancreatic lesions [6]. Herein, we present a case of SPN in a male patient where the tumor was causing extrinsic duodenal compression, leading to gastric outlet obstruction.

Case Presentation

A 22-year-old male presented with intermittent, dull aching type of abdominal pain for the last four months. It was localized to the epigastrium and reached a peak within 45 minutes of food intake, persisting for 3-4 hours. It was non-radiating and not related to bowel habits but associated with multiple episodes of non-bilious vomiting that contained food particles. He denied any history of abdominal distension, jaundice, fever, weight loss, appetite loss, blood in stool, and altered bowel and bladder habits. There was no history of passage of clay-colored stool or biliary instrumentation. He also gave a history of alcohol abuse for the past three years.

Physical examination was noncontributory. Provisional diagnoses considered at this point were pyloric stricture, gastric tumor, cystic pancreatic tumor, and duodenal mass.

Biochemical and serological investigations are shown in Table 1. Blood cell counts and renal function tests were within normal limits. Inflammatory markers such as C-reactive protein and pro-calcitonin were within the normal range.
| Laboratory test                  | Laboratory value (normal range) |
|---------------------------------|---------------------------------|
| Total bilirubin                 | 0.7 mg/dL (0.2-1.2 mg/dL)       |
| Aspartate transaminase          | 65 U/L (8-44 U/L)               |
| Alanine transaminase            | 75 U/L (7-55 U/L)               |
| Alkaline phosphatase            | 230 IU/L (44-147 IU/L)          |
| Albumin                         | 2.2 g/dL (3.5-5.5 g/dL)         |
| Amylase                         | 52 U/L (40-140 U/L)             |
| Lipase                          | 68 U/L (10-160 U/L)             |
| Hepatitis B surface antigen     | NR                              |
| Anti-hepatitis C virus antibody  | NR                              |
| HIV serology                    | NR                              |
| Autoimmune hepatitis markers    | NR                              |

**TABLE 1: Details of the initial laboratory investigations**

NR: nonreactive

Transabdominal ultrasound evaluation of the abdomen showed a homogeneous hypoechoic soft tissue texture lesion adjacent to the gallbladder fossa, duodenum, and pancreas measuring 4.4 × 4.8 × 4 cm with no internal calcifications. Contrast-enhanced computed tomography of the abdomen showed a well-defined mild heterogeneously enhancing soft tissue lesion measuring about 5.8 × 4 cm noted in the right subhepatic space around the pyloroduodenal junction, suggestive of gastrointestinal stromal tumor (Figure 1). Upper gastrointestinal endoscopy showed extrinsic compression at the D1-D2 junction. Carbohydrate antigen 19-9 was 15 U/mL (normal range: 0-37 U/mL).
The differential diagnoses considered were exophytic gastrointestinal stromal tumor and exophytic pancreatic tumor. In view of the above findings, the patient was taken up for Whipple’s pancreaticoduodenectomy and feeding jejunostomy under general anesthesia. Intraoperatively, a 6 × 4 cm space-occupying lesion of the pancreatic head occupying the pancreaticoduodenal groove was observed (Figure 2). The resected specimens were sent for histopathological examination, which revealed a cellular neoplasm consisting of sheets of small, uniform tumor cells surrounding delicate hyalinized fibrovascular stroma forming pseudopapillae. Intracellular and extracellular hyaline globules along with foamy cells were also seen. The overall histological features were suggestive of a solid pseudopapillary tumor (Figure 3). The tumor margin was negative for malignant cells, and there was no breach in the capsule of the lesion. The resected lymph nodes were free from malignant cells. The patient had an uneventful postoperative recovery and was discharged on postoperative day 7. A lifelong need for insulin and pancreatic enzyme supplementation was explained. The patient was advised six-monthly follow-up with ultrasonographic imaging of the abdomen and was doing well on the 12-month follow-up. There was no radiological evidence of recurrence of the disease on check ultrasonographic imaging.

FIGURE 1: Contrast-enhanced computed tomography of the abdomen showing a well-defined mild heterogeneously enhancing soft tissue density measuring about 5.8 × 4 cm noted in the right subhepatic space around the pyloroduodenal junction (blue arrow)
Discussion

Solid pseudopapillary neoplasm (SPN) is described as a rare pancreatic epithelial tumor that accounts for 1%-2% of all non-endocrine pancreatic tumors. There has been a steady increase in the number of diagnosed cases of cystic neoplasm of the pancreas including SPN. This is mostly attributed to the increased use of various radiological investigations [7]. It is frequently observed in young people and women, with a female/male ratio of 9.5:1 [1,6]. Although the cause of marked female predominance is still unknown, Kosmahl et al. have suggested that the migration of primordial ovarian cells to the developing pancreatic tail may give rise to SPN [8]. Recent studies have identified various molecular events involved in the pathogenesis of SPN. There is a clear distinction in the genetic profile of SPN from that of pancreatic adenocarcinoma. SPNs show β-catenin gene mutations, which is a downstream regulator of Wnt signaling [9,10].

The World Health Organization has classified SPN as a low-grade malignant tumor of the exocrine pancreas [11]. It is usually noninvasive in nature and hence asymptomatic and only detected after growing as large as 2.5-10 cm in diameter. Clinical manifestations are therefore due to the compression of the adjacent structures. These are equally distributed in the head, neck, body, and tail of the pancreas. A few cases of SPN of the pancreas associated with sinistral portal hypertension were previously reported [12]. There is also a case of SPN presenting with obstructive jaundice [3]. Gastric outlet obstruction as a presenting feature of SPN has been rarely reported before [13]. The majority of tumors are diagnosed through radiological modalities such as ultrasound or computed tomography of the abdomen, but magnetic resonance imaging helps in defining the hypervascular, well-encapsulated, round tumors with mixed cystic and solid components. Endoscopic ultrasonography-guided fine-needle aspiration biopsy is beneficial for
preoperative pathological diagnosis [14]. Certain histopathological features, such as extensive necrosis, nuclear atypia, high mitotic rate, sarcomatoid areas, and immunohistochemistry evaluation showing high expression of Ki-67, suggest the aggressive behavior of the tumor [15-17].

Despite reports suggesting locally aggressive features, the tumor has a very low-grade malignant potential and tends to have a favorable prognosis, even in the presence of metastatic disease. The overall five-year survival is as high as 97% in patients undergoing surgical resection [18].

Conclusions

In this case, the solid pseudopapillary neoplasm (SPN) of the pancreas was affecting a male patient. Herein, we report that SPN can cause complications such as gastric outlet obstruction despite its indolent course. Moreover, in a background of alcohol abuse with a pancreatic head mass, clinicians may face a diagnostic dilemma to rule out malignancy versus groove pancreatitis. Timely resection on diagnosis provides long-term survival. Histopathological evaluation of the sample stays as the investigation of choice as every pancreatic neoplasm behaves uniquely.

Additional Information

Disclosures

Human subjects: Consent was obtained or waived by all participants in this study. The Institute of Postgraduate Medical Education and Research (PGMER&R) Research Advisory Committee issued approval NA. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

1. Klimstra DS, Wenig BM, Helfless CS: Solid-pseudopapillary tumor of the pancreas: a typically cystic carcinoma of low malignant potential. Semin Diagn Pathol. 2000, 17:66-80.
2. Frantz VR: Tumors of the pancreas. Atlas of tumor pathology. Armed Forces Institute of Pathology, Washington, DC; 1959. 32-5.
3. Hamoudi AB, Misugi K, Grosfeld JL, Reiner CB: Papillary epithelial neoplasm of pancreas in a child. Report of a case with electron microscopy. Cancer. 1970, 26:1126-34. 10.1002/1097-0142(197011)26:5<1126::aid-cncr2820260245>3.0.co;2-k
4. Yu PF, Hu ZH, Wang XB, Guo JM, Cheng XD, Zhang YL, Xu Q: Solid pseudopapillary tumor of the pancreas: a review of 553 cases in Chinese literature. World J Gastroenterol. 2010, 16:1209-14. 10.3748/wjg.v16.i10.1209
5. Chen S, Zou S, Dai Q, Huang P, Cai W, Li H: The clinicopathological and immunochemical analysis of solid-pseudopapillary tumor of the pancreas: report of 9 cases. J Nanjing Med Univ. 2007, 21:598-401. 10.1007/s10367-007-0087-2
6. Choi JY, Kim MJ, Kim JH, et al.: Solid pseudopapillary tumor of the pancreas: typical and atypical manifestations. AJR Am J Roentgenol. 2006, 187:W178-86. 10.2214/AJR.05.0569
7. Zhan H, Cheng Y, Wang L, et al.: Clinicopathological features and treatment outcomes of solid pseudopapillary neoplasms of the pancreas: a 10-year case series from a single center. J Laparoendosc Adv Surg Tech A. 2019, 29:600-7. 10.1089/lap.2018.0704
8. Kosmahl M, Seada LS, Jänig U, Harms D, Klipple G: Solid-pseudopapillary tumor of the pancreas: its origin revisited. Virchows Arch. 2000, 456:473-80. 10.1007/s004280050475
9. Abraham SC, Klimstra DS, Wilentz RE, et al.: Solid-pseudopapillary tumors of the pancreas are genetically distinct from pancreatic ductal adenocarcinomas and almost always harbor beta-catenin mutations. Am J Pathol. 2002, 160:1361-9. 10.1016/s0002-9440(10)62563-1
10. Tanaka Y, Kato K, Notohara K, et al.: Frequent beta-catenin mutation and cytoplasmic/nuclear accumulation in pancreatic solid-pseudopapillary neoplasm. Cancer Res. 2001, 61:8401-4.
11. Hamilton SR, Aaltonen LA: Pathology and genetics of tumours of the digestive system. JARC Press, Lyon, France; 2000.
12. Dhali A, Pasangha E, D’Souza C, Hazra A, Dhali GK: Papillary pseudopapillary tumor of the pancreas: a rare cause of extrahepatic portal hypertension. Cureus. 2021, 15:e15707. 10.7759/cureus.15707
13. McFarlane ME, Plummer JM, Patterson J, Pencle FK: Solid-pseudopapillary tumour of the pancreas as a rare cause of gastric outlet obstruction: a case report. Cases J. 2008, 1:574. 10.1186/1757-1626-1-574
14. Zuurraizen A, Nir I, Brocklage T, Rajput A: Pseudopapillary tumor of the pancreas in a 17-year-old girl. J Clin Oncol. 2011, 29:e395-6. 10.1200/JCO.2010.33.5364
15. Tang LH, Aydin H, Brennan MF, Klimstra DS: Clinically aggressive solid pseudopapillary tumors of the pancreas: a report of two cases with components of undifferentiated carcinoma and a comparative clinicopathologic analysis of 34 cases. Am J Surg Pathol. 2005, 29:512-9. 10.1097/01.sap.0000155159.28530.88
16. Shimizu M, Matsumoto T, Hirokawa M, Morobe Y, Iwamoto S, Tsunoda T, Manabe T: Solid-pseudopapillary carcinoma of the pancreas. Pathol Int. 1999, 49:231-4. 10.1046/j.1440-1827.1999.00852.x
17. Kang CM, Kim KS, Choi J, Kim H, Lee WJ, Kim BR: Solid pseudopapillary tumor of the pancreas suggesting...
malignant potential. Pancreas. 2006, 32:276-80. 10.1097/01.mpa.0000202956.41106.8a
18. Eder F, Schulz HU, Röcken C, Lippert H: Solid-pseudopapillary tumor of the pancreatic tail. World J Gastroenterol. 2005, 11:4117-9. 10.3748/wjg.v11.i26.4117