Pancreatic Arteriovenous Malformation

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Key Words
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Abstract
An unusual case of pancreatic arteriovenous malformation (P-AVM) combined with esophageal cancer is reported. A 59-year-old man was admitted with upper abdominal pain. Contrast-enhanced computed tomography showed numerous strongly enhanced abnormal vessels and a hypovascular lesion in the area of the pancreatic tail. Angiographic study of the celiac artery confirmed racemose vascular networks in the tail of the pancreas. Endoscopic retrograde pancreatography revealed narrowing and displacement of the main pancreatic duct in the tail of the pancreas. Screening esophagoscopy showed a 0-IIa+IIc type tumor in the lower thoracic esophagus. Histological examination of esophagogscopic biopsies showed squamous cell carcinoma. Based on these findings, P-AVM or pancreatic cancer and esophageal cancer were diagnosed. Video-assisted thoracoscopic esophagectomy and distal pancreatectomy were performed. Histological examination of the resected pancreas revealed abundant abnormal vessels with intravascular thrombi. In addition, rupture of a dilated pancreatic duct with pancreatic stones and both severe atrophy and fibrosis of the pancreatic parenchyma were observed. The final diagnoses were P-AVM consequent to severe chronic pancreatitis and esophageal carcinoma. The patient’s postoperative course was relatively good.
Introduction

An arteriovenous malformation (AVM) is a complex tangle of abnormal arteries and veins linked by one or more direct connections called fistulas or shunt. AVM can occur anywhere in the body. An increasing number of patients with AVM in the digestive organs are being diagnosed due to widespread use of imaging techniques, but pancreatic AVM (P-AVM) is a very rare disease. To the best of our knowledge, fewer than 70 cases with this disease have been reported in the English language literature.

Many cases of P-AVM are associated with Rendu-Osler-Weber syndrome and are known to be a part of the visceral angiodysplasia of hereditary hemorrhagic telangiectasia. The majority of patients with P-AVM remain asymptomatic, but some present with abdominal pain or gastrointestinal bleeding. The most frequently involved portion of the pancreas has been reported to be the head (59.4%), followed by the body and tail (33.3%) and the entire pancreas (7.2%) [1]. The definitive treatment for P-AVM is surgical resection. A case of P-AVM with esophageal cancer who underwent distal pancreatectomy and esophagectomy is presented, and the relevant literature is reviewed.

Case Report

A 59-year-old man was admitted to our hospital with intermittent upper abdominal pain. He had a history of cervical disc herniation, lumbar compression fracture and abdominal aortic aneurysm. Physical examination revealed slight tenderness in the left upper quadrant of the abdomen, and no masses were palpable. Laboratory data on admission showed: erythrocyte count 411 × 10^4/mm^3 (normal 410–530), hemoglobin 13.7 g/dl (normal 14–18), leukocyte count 9,800/mm^3 (normal 4,000–8,000), platelet count 27.1 × 10^4/mm^3 (normal 15–40), serum total protein 6.8 g/dl (normal 6.5–8.0), total bilirubin 0.67 mg/dl (normal 0.2–1.0), aspartate aminotransferase 16 IU/l (normal 10–40), alanine aminotransferase 12 IU/l (normal 5–40), alkaline phosphatase 339 IU/l (normal 115–360), lactate dehydrogenase 256 IU/l (normal 119–229), γ-glutamyltranspeptidase 56 IU/l (normal 10–47), serum amylase 100 U/l (normal 33–116), blood urea nitrogen 14.0 mg/dl (normal 8–22), creatinine 0.8 mg/dl (normal 0.6–1.1), and C-reactive protein 1.71 mg/dl (normal 0–0.3). The serum level of the tumor marker carcinoembryonic antigen (CEA) was slightly elevated to 6.5 ng/ml (normal 0–5), and the carbohydrate antigen (CA19-9) level was elevated to 159.3 U/ml (normal 0–37).

Contrast-enhanced computed tomography (CT) showed numerous strongly enhanced abnormal vessels and a hypovascular lesion in the area of the pancreatic tail (fig. 1a). Angiographic study of the celiac artery confirmed racemose vascular networks in the tail of the pancreas and early venous return to the portal system (fig. 1b). Endoscopic retrograde pancreateography (ERP) revealed narrowing and displacement of the main pancreatic duct in the tail of the pancreas (fig. 1c). Screening esophagoscopy showed a 0-IIa+IIc type tumor in the lower thoracic esophagus and multiple Lugol-voiding lesions. Histological examination of esophagogastroduodenoscopy biopsies showed squamous cell carcinoma.

Based on these findings, P-AVM and chronic pancreatitis or pancreatic cancer with esophageal cancer were diagnosed. Video-assisted thoracoscopic esophagectomy and distal pancreatectomy with partial transverse colectomy were performed. The operative time was 402 min, and the amount of blood loss was 956 ml. There was a honeycomb-like structure with collected vessels in the pancreas. The splenic vein was occupied by thrombi (fig. 2).
Histological examination of the resected pancreas revealed rupture of the dilated pancreatic duct with bleeding (fig. 3a), pancreatic stones (fig. 3b) and severe atrophy with fibrosis of the pancreatic parenchyma (fig. 3c). These findings were consistent with chronic pancreatitis. In addition, many abnormal vessels, some containing thrombi, were observed in the pancreas (fig. 3d). The final diagnosis was P-AVM consequent to severe chronic pancreatitis and esophageal carcinoma. The patient had an uneventful postoperative course, and CA19-9 levels normalized to 27.6 U/ml.

Discussion

P-AVM is a very rare condition. Fewer than 70 cases with this disease have been reported in the English language literature since the first report of P-AVM by Halpern et al. in 1968 [2]. A review of AVM by Meyer et al. [3] revealed that 78% of AVMs were located in the cecum and right colon, whereas only 0.9% were located in the pancreas.

The etiology of P-AVM is classified into congenital and acquired. About 90% of patients with P-AVM are thought to be the congenital type, including patients with Rendu-Osler-Weber syndrome. The acquired type is usually secondary to inflammation, tumor or trauma [4, 5]. In the present case, there was no family history of Rendu-Osler-Weber syndrome and no telangiectasia; however, histological evidence of chronic pancreatitis was found. Therefore, it was thought to be an acquired case secondary to chronic inflammation.

P-AVM is frequently associated with symptoms of abdominal pain and gastrointestinal bleeding. Aida et al. [6] classified the mechanism of gastrointestinal bleeding into the following five types: (1) bleeding from esophagogastric varices secondary to portal hypertension; (2) bleeding from the AVM to the pancreatic duct; (3) bleeding from the AVM to the bile duct; (4) bleeding from intestinal mucosa in contact with the AVM; (5) bleeding from a duodenal ulcer associated with the AVM. So-called steal syndrome may cause abdominal pain, with shunting of blood away from the mesenteric circulation through the AVM [7].

Song et al. [1] showed the efficacy of contrast-enhanced dynamic CT as the first-line method for the diagnosis of P-AVM. The CT findings suggested that the characteristic features of P-AVM included strong enhancement or conglomeration of small hypervascular spots in the lesion and early contrast filling of the portal vein. Angiographic study is useful for the diagnosis of P-AVM and subsequent interventional therapy. The findings of P-AVM are characterized by dilated and tortuous feeding arteries, with a racemose vascular network, followed by a transient dense pancreatic stain and early venous filling. Kanno et al. [8] suggested that the diagnosis of this disease be confirmed by angiography. However, these findings can be seen in pancreatitis or hypervascular neoplasms such as cystadenoma, cystadenocarcinoma, angiosarcoma and islet cell tumor [5]. In addition, the procedure is relatively invasive. Koito et al. [9] discussed the usefulness of color Doppler ultrasonography for the detection of P-AVM, which was reported to be almost equal to that of angiography; however, it is difficult to identify the feeding arteries with ultrasonography; therefore, the procedure is relatively invasive. Koito et al. [9] discussed the usefulness of color Doppler ultrasonography for the detection of P-AVM, which was reported to be almost equal to that of angiography; however, it is difficult to identify the feeding arteries with ultrasonography; therefore, it is difficult to identify the feeding arteries with ultrasonography [10]. In the present case, contrast-enhanced CT showed numerous strongly enhanced abnormal vessels, but a hypovascular region was present in the area of the pancreatic tail. ERP revealed narrowing and displacement of the main pancreatic duct in the tail of the pancreas. The serum level of the tumor marker CA19-9 was elevated to 159.3 U/ml. Based on these findings, pancreatic cancer was considered in the differential diagnosis.

According to a recent review of 69 P-AMVs, 46.4% underwent pancreatic resection, 10.1% were treated by extended devascularization, and 15.9% were treated by transcatheter arterial embolization (TAE). Despite a small number of cases, irradiation therapy after
TAE due to recurrent bleeding and a transjugular intrahepatic portosystemic shunt (TIPS) has been reported [1]. P-AVM has multiple feeding arteries; therefore, it is very difficult to ligate or embolize them all. If no treatment is given, a P-AVM will grow and cause portal hypertension, gastrointestinal bleeding or rupture of esophageal varices [1, 5]. Furthermore, it has been reported to be impossible to reduce portal hypertension, once established, even if the AVM is surgically removed [4]. Therefore, surgical resection of the affected pancreatic lesion at an early stage is needed for complete cure of P-AVM. TAE, irradiation therapy and TIPS are options for high-risk surgical patients.

In summary, to the best of our knowledge, we have described for the first time a case of P-AVM combined with esophageal cancer. P-AVM is a very rare disease, but should be considered in the differential diagnosis of a hypervascular pancreatic lesion.

Disclosure Statement

The authors have no funding or conflicts of interest to disclose.

References

Fig. 1. **a** Contrast-enhanced CT showed numerous strongly enhanced abnormal vessels (arrowheads) and a hypovascular lesion in the area of the pancreatic tail (arrow). **b** Angiographic study of the celiac artery confirmed racemose vascular networks in the tail of the pancreas (arrowhead) and early venous return to the portal system (arrow). **c** ERP showed narrowing and displacement of the main pancreatic duct in the tail of the pancreas (arrow).
Fig. 2. A honeycomb-like structure with collected vessels was observed in the pancreas. The splenic vein was occupied by thrombi (asterisks).

Fig. 3. a Histological examination of the resected pancreas showed rupture of the dilated pancreatic duct with bleeding (HE, ×10). b Pancreatic stones (HE, ×10). c Severe atrophy with fibrosis of the pancreatic parenchyma (HE, ×10). d Abundant abnormal vessels were observed in the pancreas, some containing thrombi (Elastica van Gieson, ×10).