CASE REPORT

Pancreas-Preserving Duodenal Resections with Bile and Pancreatic Duct Replantation for Duodenal Dystrophy. Two Case Reports

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ABSTRACT

Context Duodenal dystrophy is a rare disease, characterized by the chronic inflammation of the aberrant pancreatic tissue in the duodenal wall. Case reports Two middle-aged men were admitted with upper abdominal pain of several months duration, periodic nausea and vomiting after meals, intermittent jaundice and weight loss. A diagnosis of cystic dystrophy of the vertical part of the duodenum without chronic inflammation of the orthotopic pancreas was established in both cases by multi-detector computed tomography, magnetic resonance imaging and endosonography. Both patients were successfully treated by two modifications of pancreas-preserving duodenal resections with reimplantation of the bile and pancreatic ducts into the neoduodenum. Conclusion These cases are a good example of a pancreas-preserving approach to duodenal dystrophy treatment and can be an alternative to the Whipple procedure in cases of mild changes of the orthotopic gland.

INTRODUCTION

Duodenal dystrophy, which is the chronic inflammation of the aberrant pancreatic tissue in the duodenal wall, is a relatively rare disease [1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13] despite the fact that heterotopic pancreatic tissue is found in the duodenum in 25-35% of all cases of pancreatic ectopia in the digestive tract [11, 12, 13, 14, 15]. The heterotopic pancreas is usually functioning and acute or chronic pancreatitis is more probable in the ectopic tissue than in the orthotopic gland as a result of an underdeveloped duct system [8, 11, 12]. The progression of ectopic pancreatitis associated with increasing cystic formation could result in a blockade of the major or minor duodenal papilla and consequent chronic pancreatitis in the (orthotopic) pancreas proper [10, 11, 12, 13, 14, 15]. Furthermore, malignant transformation of the aberrant pancreas is not a rare occurrence [16, 17]. It makes a timely diagnosis of this condition vital as it may determine the choice of the surgical procedure [3, 4, 5, 6, 9, 10, 12, 13]. Duodenal dystrophy management remains a topic of discussion and pancreaticoduodenectomy is currently the method of choice when conservative treatment fails [3, 4, 5, 6, 9, 10, 11, 12, 13]. The purpose of this report was to show the possibility of preserving the orthotopic pancreas in cases of cystic dystrophy in the heterotopic gland located in the region of the papilla of Vater.

Case #1

A 32-year-old man was admitted to the 4th Clinical Hospital, Moscow, Russia, after two months of upper abdominal pain, periodic nausea and vomiting after meals and weight loss. The patient had a short history of alcohol and tobacco abuse and, a few months previously, he had been treated for bleeding from an ulcer in the vertical duodenal branch. A physical examination revealed no cardiac, respiratory or urinary disorders. His pulse was 72 beats/min, sinus rhythm, and his blood pressure was 120/80 mmHg. There was slight tenderness in the right upper abdomen quadrant without a distinct palpable mass. Hematological and biochemical blood tests did not show any considerable shifts, including bilirubin, alanine and aspartate aminotransferase, gamma GT levels and oncological markers. Serum amylase was mildly elevated to 350 U/mL (reference range: 60-100 U/mL) and biochemical signs of hepatitis C minimal activity were revealed. The pH of the urine was 7 and relative density was 1.016. The sediment contained 1-2 white cells per high-power field.
An upper GI endoscopy revealed deformation and infiltration of the second portion of the duodenum, mainly at the expense of an ulcerated medial wall. The main papilla was involved in the infiltrated tissue, but the bile flow was preserved. A biopsy of the duodenal mucosa detected moderate inflammation. A barium meal revealed mild evacuation disorders due to a narrowing of the second part of the duodenum. Abdominal ultrasonography did not reveal any significant changes. Contrast multi-detector computed tomography and magnetic resonance imaging showed a fibrous parietal thickening and a large ovoid mass (4.2x2.5 cm) of fluid density, divided by the septa, in the medial wall of the second portion of the duodenum. The pancreatic head, body and tail were unchanged in size and structure. The gastroduodenal artery was shifted notably forward and to the left (Figure 1). The diagnosis of cystic duodenal dystrophy was confirmed by an endosonography examination which revealed an ovoid septated cystic structure of the same size (4.2x2.5 cm) located in the submucosa and muscularis propria, associated with infiltration and sclerosis of the second portion of the duodenal wall (Figure 2). There were no signs of tumor growth in the duodenum, common bile duct, pancreas or the peripancreatic lymph nodes.

The patient was operated on seven days later, after preoperative treatment. The hepatic flexure of the colon was deflected through a midline incision, and the duodenum and the head of the pancreas were extensively mobilized from the retroperitoneum well to the left of the midline. The stomach was dilated due to the stenosis of the second part of the duodenum. A cholecystectomy was performed and the papilla was stented by a Dogliotti probe through the cystic duct. The proximal jejunum was transected 15 cm below the Treitz ligament, detached from its mesentry and transferred to the right behind the superior mesenteric vessels. The duodenum was transected 1.5 cm distal to the pylorus in the unaffected zone. The second, third and fourth portions of the duodenum were detached from the pancreas by the division and ligature of the short vessels and removed with the main papilla after transection of the common bile and the main pancreatic ducts; a pancreas-preserving subtotal duodenectomy was performed (Figure 3). The proximal jejunal was passed through the mesocolon, and the bile and pancreatic ducts were implanted in the shifted jejunum 4 cm below the future duodenojejunal anastomosis using a single layer of interrupted 5/0 absorbable sutures. An end-to-end duodenoenterostomy was performed with a single layer of 4/0 absorbable continuous suture (Figure 4) and a microdrain was placed through the cystic duct stump into the common bile duct.
The postoperative course was uneventful. Cholangiography and a barium meal (Figure 5) on the 6th postoperative day revealed no leaks and good functioning of the neoduodenum and choledocho-pancreaticojejunostomy. The bile drain was removed 3 weeks later. Throughout the ensuing six-month follow-up, the patient had no complaints and gained 4 kg.

**Case #2**

A 43-year-old man with abdominal pain of a one year duration, nausea, vomiting, weight loss and intermittent jaundice was referred to the Hospital of the Russian Federal Security Service, Moscow, Russia, with a clinical diagnosis of chronic pancreatitis. The patient had a history of previous extensive small bowel resection; one year previously he had been operated on
for acute pancreatitis with abdominal draining and a cholecystectomy. The epigastrium was mildly sensitive on physical examination. The laboratory findings were normal, except for a slight increase in the bilirubin level (40 µmol/L; reference range: 3.5-19.0 µmol/L). Abdominal US showed diffuse changes of the liver and pancreas. On computed tomography (Figure 6) and magnetic resonance imaging (Figure 7), a thick-walled cystic lesion 4x4.5 cm which had infiltrated the medial wall of the second part of the duodenum was observed against the background of the minimally changed pancreatic structure with pancreatic and common bile duct dilation. Endosonography revealed a solid and septated cystic lesion mainly located in the submucosa of the narrowed second portion of the duodenal wall.

The patient was operated on with a diagnosis of “chronic inflammation of the aberrant pancreatic tissue in the duodenal wall (duodenal dystrophy), bile and pancreatic hypertension”. Exploration and an extensive Kocher’s maneuver were performed through a midline incision. The stomach and the first portion of the duodenum were dilated due to the intraluminal mass of the second part of the duodenum connected to its medial wall. The gallbladder (14x7 cm) and common bile duct (13 mm) were tense and dilated. After the cholecystectomy, the papilla was stented using a Dogliotti probe, the tip of which was found exactly at the low pole of the intraluminal mass. Taking into
consideration the shortness of the small bowel, we decided to save as much of its length as possible. The duodenum was transected 2 cm below the pylorus and 3 cm below the main papilla. Following the transection of the common bile and the main pancreatic ducts, and detachment of the duodenum from the pancreatic head, the whole of its second part, including the main papilla, was removed (Figure 8). Fifty centimeters below the Treitz ligament, a 10 cm segment of the proximal jejunum, supplied by the artery and vein, was cut out and passed through the mesocolon (Figure 9). The shifted segment was interposed between the first and the third parts of the duodenum and jeuno-jejuno- and distal duodeno-jejuno-anastomoses were performed. The bile and pancreatic ducts were sutured together and implanted in the neoduodenum 4 cm below the final proximal duodeno-jejuno-anastomosis. All the bowel anastomoses were end-to-end and carried out using a single layer continuous 4/0 absorbable suture. The choledocho-pancreaticojejunostomy was carried out with a single layer of interrupted 5/0 absorbable sutures (Figure 10). The procedure was completed by drainage of the common bile duct through the cystic duct stump and drainage of the right abdominal cavity. The postoperative course was uneventful. Cholangiography and a barium meal (Figure 11) on the 7th postoperative day revealed no leaks and good functioning of the neoduodenum and choledocho-pancreatico-jejunostomy. The bile drain was removed 4 weeks

Figure 10. Scheme of the pancreas-preserving resection of the second portion of the duodenum (Case #2). The shifted segment is interposed between the 1st and the 3rd parts of the duodenum. Jejunoo-jejuno- and duodeno-jejuno-anastomoses are performed. The bile and the pancreatic ducts were implanted in the neoduodenum 4 cm below the proximal duodeno-jejuno-anastomosis. The common bile duct stent has been omitted.

Figure 11. X-ray barium meal examination seven days after the pancreas-preserving resection of the second portion of the duodenum (Case #2). No leaks and good functioning of the interposed jejunal segment (arrow).

Figure 12. Microscopy (Case #1; H&E, x50). The fragment of pancreatic tissue heterotopia presented mainly by ductal structures in the submucosa of the duodenum.

Figure 13. Microscopy (Case #2; H&E, x400). The fragment of the dilated ectopic pancreatic duct with PanIN-2 changes (arrows) in the submucosa and muscularis propria of the duodenal wall.
later. During the subsequent five-month follow-up, the patient had no symptoms, returned to work and gained 6 kg.

In both cases, the diagnosis was confirmed macroscopically (Figure 8) and histopathologically (Figures 12 and 13) after detecting pancreatic tissue in the duodenal wall which was fully isolated from the pancreas proper.

DISCUSSION

Both cases were characterized by cystic dystrophy surrounded by inflammation and fibrosis in the heterotopic pancreas located in the duodenal wall; in both cases, the diagnosis was established by computed tomography and confirmed by magnetic resonance imaging and endosonography. Although similar lesions are frequently associated with chronic inflammation of the orthotopic pancreas [8, 9, 10, 11], in these cases, the changes of the pancreas proper were minimal. The main symptoms in both cases were abdominal pain, nausea and vomiting, attributed to inflammatory changes as well as to duodenal, bile and pancreatic duct stenoses.

The choice of a pancreas-sparing procedure instead of the conventional Whipple procedure [3, 4, 5, 6, 9, 10, 11, 12, 13] was determined by the absence of heavy scarring changes in the pancreatic head and duodenum, which usually present in the event of associated “orthotopic” chronic pancreatitis. These conditions usually lead to an unnecessary excessive procedure, such as a pancreaticoduodenectomy.

A pancreas-preserving procedure is theoretically the optimal modality for the treatment of duodenal dystrophy due to the removal of the pathologic tissue and the preservation of the exocrine and endocrine pancreatic functions. The aim of this report was to describe the possibility of such surgery for duodenal dystrophy which, in our opinion, clearly demonstrates the essence of the desirable therapy. We did not come across any reports in the literature regarding a pancreas-preserving procedure for duodenal dystrophy [18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37] nor did we find any information about the resection of the second duodenal portion with the main papilla followed by the reconstruction with bowel interposition [19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39]. We believe that both types of pancreas-preserving procedures are better choices than a pancreaticoduodenectomy for duodenal dystrophy or for any other non-malignant condition of the duodenum; the former is easier to perform and the latter is preferable in selected cases of concomitant short bowel when direct duodeno-duodenostomy [38, 39] is risky because of excessive tension.

In our opinion, the risk of such a procedure is the same as that of a pancreaticoduodenectomy and experience with a total duodenectomy for familial polyposis has shown that mortality and morbidity in specialized departments after pancreas-preserving procedures were comparable with those after a pancreaticoduodenectomy which is still considered the primary therapeutic choice in duodenal dystrophy [34, 35, 37]. The Whipple procedure is a major operation with a high risk of diabetes, exocrine insufficiency and significant mortality (1-5%) and we feel that both pancreaticoduodenectomies and pancreas-preserving procedures should be performed only in specialized departments. Comparison of the efficacy of these procedures will only be possible after the assessment of long-term results of a sufficient number of cases.

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Conflict of interest

The authors have no potential conflict of interest

Reference


