LETTER

A Large Gastric Duplication (Choristoma) in an Adult Mimicking a Mucinous Cystic Tumor of the Pancreas

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Summary

Gastric duplication represents a very rare entity in the adult population. Most of the symptoms are due to the presence of ectopic gastric mucosa (30-35% of cases), gastrointestinal bleeding or perforation or neoplasm formation. We report a case of gastric duplication in an adult mimicking a mucinous cystic neoplasm of the pancreas.

Dear Sir:

Gastric duplications represent between 3 and 4% of foregut duplications and are a rare entity in adults [1]. They are predominantly diagnosed in infancy (before age two). Gastric duplication may be tubular or cystic, the cystic form being more frequent. They are usually located in the greater curvature of the stomach, without communication with the gastric lumen and often mimic other disease entities [1]. Ectopic gastric mucosa is observed in 30-35% of the cases, intestinal epithelium in 24% and heterotopic pancreatic tissue in 1% [1, 2]. They sometimes present as pancreatic complications: relapsing pancreatitis, pancreatic pseudocysts, endoluminal bleeding and neoplasia formation [3, 4, 5, 6, 7, 8, 9]. We present a case of gastric duplication (foregut choristoma) in an adult mimicking a mucinous cystic neoplasm of the pancreas. To our knowledge, only one similar case has been reported worldwide.

Case report

A 38-year-old woman with a previous diagnosis of ileal Crohn’s disease, presented to our institution complaining of epigastric discomfort, bloating, postprandial nausea and 2-3 bowel movements per day. She had no history of pancreatitis and an unremarkable gynecologic history. The patient was further investigated with abdominal ultrasonography and computerized scan tomography (CT) which revealed a large cystic mass (8.5 cm diameter) in the pancreatic tail (displacing the stomach wall) (Figure 1a). Fine needle-aspiration showed a mucinous content with detritus and acellular fluid. The patient underwent a distal pancreatectomy with splenectomy. No communication between the pancreatic mass and the lumen of the stomach was assessed. The postoperative...
course was uneventful. The patient remains free of symptoms 10 years after the procedure. Pathological study revealed a 9.5 cm diameter cystic mass with a homogeneous wall (Figure 1b). Part of the cyst was lined with gastric mucosa surrounded by a wall of muscularis propria well delineated from the normal pancreas tissue (Figure 2ab). A focal hyperplasia of simple cuboid epithelium and ciliated pseudostratified epithelium was observed (Figure 2cdef). The diagnosis of a gastric duplication cyst (foregut choristoma) of the pancreas was made.

Discussion

Gastric duplications represent between 3 and 4% of foregut duplications which include the esophagus, stomach, 2nd and 3rd portion of the duodenum and the pancreas [1, 2]. They usually present in infancy as cystic formations with occasional compression symptoms, such as intestinal sub-occlusion, palpable mass, poor weight gain or symptoms secondary to the heterotopic mucosa, such as gastrointestinal bleeding, perforation into the peritoneal cavity or neoplasm formation which is very rare in adults. Some authors have coined the term “choristoma” derived from an ancient Greek word (χωριστϖμα), the result of separation in order to recall the embryonic origin of this entity. The case reported showed the clinical and radiological pattern of a mucinous cystic tumor of the pancreas. Due to its size (9.5 cm diameter) and the presence of symptoms, a distal pancreatectomy with splenectomy was carried out. The patient is free of symptoms 10 years after the procedure. We have found four cases of gastric duplications presenting as pancreatic complications in adults: one case of relapsing pancreatitis, two cases of pancreatic pseudocyst and a case of mucinous cystic tumor [3, 5, 6, 7]. Due to the fact that gastric duplication is located in the greater curvature of the stomach, it can present as a pseudocyst or as a cystic tumor of the pancreas. Complete surgical resection is warranted as a tendency of the heterotopic gastric mucosa to become malignant has been described [10]. Table 1 shows the cases reported which present pancreatic complications in children and adults.

Figure 2. Microscopic aspect of the cyst wall. Some areas are lined by gastric-type mucosa, either oxyntic-type (a; b: detail of a), or antral mucosecretory-type epithelium with mild hyperplasia (c; d: detail of c). Flat ciliated pseudostratified epithelium covering other areas of the inner face of the cyst (e; f: detail of e). (H&E; a., c., e.: 15x; b., d.: 200x; f: 400x).
Conflict of interest

The authors have no potential conflict of interest.

References