Adult gastric duplication: an unknown condition within the spectrum of gastric submucosal lesions

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ABSTRACT

Background: Gastric duplication is an uncommon condition in adults representing only 4-8% of all gastrointestinal duplications. It develops as a cystic growth within the thick gastric wall that is surrounded by a layer of smooth muscle and lined with gastric mucosa. It is usually asymptomatic but may present with nonspecific abdominal symptoms as well as cyst-related complications such as gastrointestinal (GI) bleeding, ulceration and even pancreatitis. They are difficult to distinguish from other gastric submucosal lesions and may also mimic a gastrointestinal tumor (GIST) or mucinous cyst of the pancreas. Endoscopic ultrasound (EUS) is the most widely used diagnostic procedure and histology provides a definitive diagnosis.

Case report: We report the case of a 36-year-old female patient with a history of a "contained" perforated gastric ulcer that was managed conservatively. The patient was diagnosed with a gastric GIST which prompted a laparoscopic subtotal gastrectomy and Roux-en-Y reconstruction with a good postoperative outcome.

Discussion: The definitive histological diagnosis was gastric duplication. Thus, the previous diagnosis of a "contained" perforated gastric ulcer was in fact a complication of the gastric duplication itself.

Key words: Gastric duplication. Stomach. Submucosal tumors. Gastrointestinal stromal tumor. Mucinous pancreatic cyst.

INTRODUCTION

Gastric duplication is a rare congenital malformation representing only 4% of gastrointestinal (GI) tract duplications. It develops as a cystic tumor within the gastric wall surrounded by a smooth muscle layer and lined with gastric mucosa, although it may also contain ectopic duodenal or pancreatic mucosa (1,2). It is considered as a rare condition in adults. The lesion is usually asymptomatic and difficult to distinguish from other submucosal growths. Thus, it is often mistaken for a GI stromal tumor or a mucinous pancreatic tumor. We report the case of a 36-year-old female patient with a history of a gastric ulcer who was subsequently diagnosed with a gastrointestinal stromal tumor (GIST) after work-up for dyspepsia. After surgery the patient received a definitive histological diagnosis of a gastric duplication.
The pathology analysis described a subtotal gastrectomy specimen 14 x 6 x 3 cm in size with no gross changes either externally or after opening the specimen. Serial slices of the specimen revealed a cystic cavity 0.8 cm in diameter within the gastric wall, with mucinous-looking contents. Microscopically, the gastric mucosa exhibited normal characteristics and a cystic cavity lined with normal gastric mucosa was found within the wall with a well delimited muscular wall and focal pancreatic metaplasia. These findings were consistent with gastric duplication (Fig. 2).

DISCUSSION

Gastrointestinal tract duplications are defined as cystic structures adjacent to a GI segment (1,2). The ileum is the most commonly involved site (35%), followed by the jejunum, colon and stomach in only 2-9% of cases (2). It is considered as a congenital malformation that is identified in over 70% of cases in patients younger than 12 years of age. Diagnosis of this lesion is rare during adulthood.

Patients are usually asymptomatic or have vague abdominal symptoms, and this condition is often detected as an incidental finding. The presence of ectopic gastric, duodenal or pancreatic tissue is characteristic and clinically significant as it may result in complications such as upper GI bleeding, ulcer and episodes of pancreatitis (2). In rare cases it may become malignant and mainly develops into adenocarcinoma. Most are isolated from the lumen of the GI tract and 50% are associated with other...
congenital abnormalities (pancreatic duplications, ectopic or annular pancreas) or spinal disorders (spina bifida) (3). Pancreas related abnormalities reflect the common embryonic origin of the pancreas and stomach.

The etiopathogenesis remains unknown. The condition is thought to originate from the dorsal aspect of the primitive gut tube during development. Thus, most cases involve the greater curvature of the stomach (4) and only 5.5% are located at the lesser curvature. These duplications develop before the epithelium differentiates into the several types present in adults, and are thus designated according to the associated organ. Multiple theories exist, of which three are of interest: a) persistence of a vacuole or diverticulum during bowel development; b) failed fusion of longitudinal gastric folds resulting in an epithelial bridge; and c) traction diverticulum formation from anomalous endodermal and notochordal development (5).

Most cases reported in adults are preoperatively diagnosed as GISTs or mucinous pancreatic cysts (4) with a definitive diagnosis achieved by the histological examination of the surgical specimen. This suggests a lack of awareness with regard to this condition, particularly from the radiographic and echoendoscopic perspectives. Contrast-enhanced CT shows cystic growths with thick walls near the stomach with inner layer enhancement. These radiographic features are similar to those of mucinous pancreatic cysts, which makes an accurate diagnosis difficult. Magnetic resonance imaging does not improve identification and both imaging techniques are estimated to yield an incorrect diagnosis in up to 43-70% of cases.

EUS is highly advisable for the characterization of gastric submucosal lesions. This technique is key for the identification of these lesions as cyst type and location can be defined. FNAP can be performed for the differential diagnosis from other submucosal lesions and it informs about potential malignant sites (5-7). The lesion is observed as an anechoic round or ovoid image lined with a thin echogenic muscle layer, giving the impression of a “double wall” or a “muscular rim” (8,9).

Diagnosis requires histological confirmation according to the Rowling’s criteria: the cyst wall should be contiguous with the gastric wall and the lesion must be surrounded by smooth muscle in continuity with the gastric muscle and lined with gastric, epithelial or intestinal mucosa (5).

There is no standard treatment. Symptomatic or complicated cases should be treated surgically. However, no consensus exists with regard to asymptomatic patients. Some researchers consider that these patients only require follow-up while others advocate for surgery after diagnosis, regardless of symptoms, as there is a risk for malignancy or complications (5). Enucleation and endoscopic submucosal resection have been reported (10), but complete excision is recommended, as in this case.

This rare congenital malformation in an adult is a lesson to be learned as it is an uncommon condition initially diagnosed as a simple perforated ulcer and, subsequently, as a gastric submucosal mass. The contained perforated ulcer event the patient experienced in 2009 may have been a complication of the gastric duplication itself. In conclusion, we consider that gastric duplication should be included in the differential diagnosis of gastric submucosal growths, where EUS plays a key role for preoperative diagnosis. These lesions should be surgically removed once diagnosed due to the risk of complications or malignization. A minimally invasive surgical approach is the most appropriate approach in these cases.

REFERENCES