Fatal hemorrhage from a gastroaortic fistula secondary to gastric ulceration associated with Nissen fundoplication and nonsteroidal anti-inflammatory drug use

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Acute gastrointestinal hemorrhage from a gastroaortic fistula in the gastric fundoplication pouch is a rare complication of Nissen fundoplication. The present case reports a gastroaortic fistula secondary to gastric ulceration associated with prior Nissen fundoplication and nonsteroidal anti-inflammatory drug use. A 55-year-old man presented with massive hematemesis and died of exsanguination during emergency laparotomy. Recognition of factors that predispose a patient to gastric ulceration after fundoplication, including nonsteroidal anti-inflammatory drug use, is critical to arouse the high index of suspicion required to diagnose and manage this life-threatening complication.

Key Words: Gastric ulceration; Gastroaortic fistula; Nissen fundoplication

Nissen fundoplication is a common and effective surgical procedure for gastroesophageal reflux disease refractory to medical therapy. Rare complications of this procedure include gastric ulceration leading to formation of a gastroaortic fistula (1-4). This may result in severe, potentially life-threatening, upper gastrointestinal hemorrhage. The present report documents the formation of a gastroaortic fistula likely secondary to nonsteroidal anti-inflammatory drug (NSAID)-induced gastric ulceration four years after Nissen fundoplication for gastroesophageal reflux disease.

CASE PRESENTATION

The patient was a 55-year-old man who presented to the emergency department at Kingston General Hospital (Kingston, Ontario) with unstable gastrointestinal bleeding. He had been found obtunded on the floor of his home with approximately 2 L of blood in the surrounding area. His wife gave an account of his medical history, which included complaints of epigastric pain and cramping for two days before the event. His medical history was significant for severe reflux esophagitis that had failed antireflux medical therapy. He had undergone a Nissen fundoplication four years earlier for treatment of esophagitis and gastroesophageal reflux disease. His symptoms had resolved after surgery. He was taking indomethacin and morphine sulfate for chronic back and pleuritic chest pain for 10 months before admission. The indomethacin dosage had been recently increased to 75 mg three times per day. On arrival at the hospital, he was in severe hypovolemic shock, with a systolic blood pressure of 60 mmHg. His initial laboratory investigations showed a hemoglobin level of 83 g/L, leukocytes $15.2 \times 10^9/L$, platelets $199 \times 10^9/L$, and an international normalized ratio of 1.5. An NSAID-induced gastric ulceration was suspected initially. The patient was aggressively resuscitated with nine units of packed red blood cells, five units of platelets and five units of fresh frozen plasma. He was intubated and an urgent upper endoscopy was performed, which demonstrated a large amount of fresh and clotted blood in the stomach. A specific site of hemorrhage was not identified. However, the proximal one-half of the stomach could not be adequately seen because of a large obstructing blood clot. A subsequent visceral angiogram including the celiac and superior mesenteric arteries also failed to show any evidence of a bleeding source. Due to ongoing bleeding, an emergency laparotomy was performed in an attempt to control the active hemorrhage. It was found intraoperatively that the stomach had ruptured posteriorly. An anterior gastrotomy was performed, which revealed massive bleeding originating in the proximal one-half of the stomach. A left lower anterior thoracotomy was performed, and the patient's hemorrhage was stopped. He underwent an emergent Nissen reversal. He was found obtunded on the floor of his home with approximately 2 L of blood in the surrounding area. His wife gave an account of his medical history, which included complaints of epigastric pain and cramping for two days before the event. His medical history was significant for severe reflux esophagitis that had failed antireflux medical therapy. He had undergone a Nissen fundoplication four years earlier for treatment of esophagitis and gastroesophageal reflux disease. His symptoms had resolved after surgery. 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and the aorta was occluded using digital palpation manipulation with good proximal control of the bleeding. However, the patient’s ventricular rhythm decompensated into ventricular fibrillation and he expired on the operating table.

The postmortem examination showed evidence of recent emergency laparotomy with gastrotomy. There was a 9.5 cm linear incision along the anterior wall of the body of the stomach. The gastric antrum contained only a few millilitres of blood, whereas the duodenum, jejunum and ileum were filled with a large amount of clotted blood. Evidence of remote Nissen fundoplication surgery was noted. A pouch-like component of the fundoplication wrap, measuring 5 cm × 3.5 cm × 3.5 cm in greatest dimensions, was identified situated on the left side of the esophagus. The posterior surface of this pouch was densely adhered to the anterior surface of the lower thoracic aorta. A 5 cm × 2 cm zone of mucosal ulceration was present on the posterolateral wall of the fundoplication pouch. A 2 mm perforation overlying a gastroaortic fistula was noted, involving the anterior wall of the lower thoracic portion of the aorta, secondary to the chronic gastric ulcer in the fundoplication (Figures 1 and 2). Aside from the Nissen fundoplication, the esophagus was unremarkable with no evidence of esophagitis. The aorta showed only minimal atherosclerosis.

Histological sections from the area of the fistulous tract showed erosion of the gastric mucosa with transmural ulceration. The posterior surface of the stomach was directly apposed to the aorta by fibrosis and chronic inflammation. Giemsa-stained sections of the gastric antrum and fundoplication were negative for Helicobacter pylori organisms.

**DISCUSSION**

Aortoenteric fistulas develop uncommonly as a result of either aortic or gastrointestinal pathology. Aortic abnormalities that may predispose a patient to fistula formation include aortic aneurysms, atherosclerotic disease, infections or postaortic vascular graft placement (5-7). Gastrointestinal pathology includes irradiation for malignancy, peptic ulcer disease, foreign bodies and ingestion of corrosive materials (8). A gastroaortic fistula can occasionally form secondary to gastric ulceration after Nissen fundoplication for gastroesophageal reflux disease. This complication is very rare and has been described in only a handful of cases (1-4,9).

The most common causes of gastric ulceration include H pylori and NSAIDs (10,11). However, these etiologies have not been directly implicated in the post-Nissen fundoplication gastroaortic fistula patient population. Instead, the cause of
ulceration after Nissen fundoplication has been hypothesized to be mechanical in nature. Possible mechanisms include surgical trauma, delayed gastric emptying due to vagal injury, hypergastrinemia, suture irritation from the wrap, and gastric wall ischemia due to ligation of short gastric vessels. Over time, these factors may result in gastric ischemia leading to ulceration, which given the close proximity of the aorta and the stomach due to the surgical fundoplication procedure, gives rise to the subsequent development of a fistula (12). In addition to the above potential mechanical factors, we believe the prolonged use of NSAIDs in the present case likely contributed to gastric ulceration of the fundoplication, resulting in fistula formation.

Due to the rarity of aortogastroenteric fistulas, the diagnosis is frequently missed or made too late for surgical correction, often resulting in a fatal outcome (4). A great proportion of the few cases of primary aortoenteric fistulas first presented with ‘herald bleeding,’ attributed to spasms of the intestinal musculature that temporarily close the fistula and stop the initial bleeding. A review of relevant literature by Wasvary et al (4) identified 10 of 14 cases that presented with a herald bleed before a massive upper gastrointestinal hemorrhage. The diagnostic triad described by Lewis and Allan (13) may be helpful in establishing the clinical diagnosis. It includes a history of hemorrhage with no obvious source, a pulsatile abdominal mass and recurrent cardiovascular collapse. Any previous gastroesophageal surgery or distortion of the gastrointestinal anatomy is also highly suggestive of a potential primary aortogastroenteric fistula in these circumstances. As demonstrated by our case, a history of Nissen fundoplication, especially with a background of NSAID use, should serve to bring this rare complication to the forefront of the differential diagnosis.

Regrettably, the diagnostic sensitivity of computed tomography, and even arteriography or gastroduodenoscopy is low (14). Definite diagnosis is often made only during operative exploration or at autopsy (4). In the present case, both the upper gastrointestinal endoscopy and the selective visceral angiogram did not identify the source of the hemorrhage. Specifically, the gastroaortic fistula was not seen endoscopically, likely due to its location within the fundoplication wrap itself. In this case, angiographic examination of the aorta proximal to the celiac artery was not undertaken. Accordingly, the fistula site was not included in the study. Whether aortography proximal to the celiac artery would have identified the fistula is speculative, but it is worth considering in similar circumstances.

**SUMMARY**

Gastroaortic fistula is a rare, but potentially life-threatening, complication after Nissen fundoplication. Recognition of ulcerogenic factors, including NSAID use, that predispose a postfundoplication patient to gastric ulceration is critical to arouse the high index of suspicion required to diagnose and manage this rare complication.
REFERENCES
