Giant Brunner’s Gland Hamartoma of the Duodenal Bulb Presenting with Upper Gastrointestinal Bleeding and Obstruction

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Brunner’s gland hamartomas are small benign lesions that are most commonly found in the bulb of the duodenum. They are very uncommon, and most are found incidentally during upper gastrointestinal series or esophagogastroduodenoscopy. The lesions tend to be asymptomatic, but patients may present with symptoms of duodenal obstruction or hemorrhage secondary to ulceration. Histologically, a Brunner’s gland hamartoma consists of the components of Brunner’s gland cells, as well as glandular, adipose and muscle cells. In this study, we report the case of a 30-year-old man who presented with upper gastrointestinal bleeding and obstructive symptoms due to a giant Brunner’s gland hamartoma in the duodenal bulb. The hamartoma was successfully removed by endoscopic resection. No significant complications were observed. Microscopically, the lesion was found to be entirely composed of variable Brunner’s glands and adipocytes.

Key Words: Brunner glands; Hamartoma; Gastrointestinal bleeding; Gastrointestinal obstruction

INTRODUCTION

Brunner’s gland hamartoma, also known as Brunner’s gland adenoma or brunneroma, is a rare benign tumor arising from the Brunner’s glands of the duodenum.1 Brunner’s gland hamartomas account for 10.6% of all benign tumors of the duodenum, and have been found in 0.008% of a single series of 215,000 autopsies.1,2 These lesions typically form polypoid pedunculated masses as they grow. The majority of Brunner’s gland hamartomas are asymptomatic; however, in symptomatic patients, they can be treated by endoscopic or surgical excision. We report a case of the successful endoscopic removal of a large Brunner’s gland hamartoma in a patient who presented with upper gastrointestinal bleeding and obstruction.

CASE REPORT

A 30-year-old man presented with a 3-month history of nausea, bloating, dizziness, dyspnea on exertion, and self-limiting episodes of melena. His past medical history included obesity, hypertension, diabetes mellitus, hypercholesterolemia, and metabolic syndrome. He was initially seen at a primary clinic. A subepithelial duodenal lesion was detected on esophagogastroduodenoscopy, and this was diagnosed as a lipoma on endoscopic ultrasonography. He was managed conservatively, and received two 8-unit blood transfusions for his iron deficiency anemia. Subsequent abdominal ultrasound and colonoscopy findings were normal. However, the patient did not improve over the next 3 months and complained of persistent dizziness and nausea.

He was then referred to our hospital for further investigation. On questioning, he denied weight loss or the use of nonsteroidal anti-inflammatory drugs. His height and weight were 180.9 cm and 145.8 kg, and he had a body mass index of
44.5 kg/m². Physical examination did not reveal any abdominal tenderness, rebound tenderness, or palpable masses. His complete blood count was white blood cells 9,770/μL, hemoglobin 9.7 g/dL, hematocrit 31.6% (mean corpuscular volume 71.8 fl, mean corpuscular hemoglobin 22 pg), and platelets 281,000/μL. Serum total bilirubin was 1.2 mg/dL, serum as-
partate aminotransferase was 37 IU/L, and serum alanine transaminase was 67 IU/L. Abdominal computed tomography showed a large polypoid mass in the duodenum (Fig. 1). It was a hypodense tubular intraluminal mass measuring about 9.3×2 cm in the first and second parts of the duodenum. Esophagogastroduodenoscopy showed a large pedunculated and eroded mass with a thick stalk arising from the duodenal bulb. The tumor was located on the anterior surface of the duodenal bulb and had a wide base, causing near-total obstruction (Fig. 2A-C). The tip of the tumor also was covered with exposed vessels and hematin, but was not actively bleeding (Fig. 2D). Endoscopic polypectomy was performed. However, the lesion was too large to fit into the conventional snare. We grasped a portion of the mass and pulled it up into the stomach with the snare, and then performed two partial snare polypectomies (Fig. 3A). Next, we performed endoscopic submucosal dissection on the bulb of the duodenum after the duodenal U-turn (Fig. 3B, C). Hemoclips were applied to prevent bleeding and perforation. The gross endoscopic resection specimen showed a large duodenal lesion measuring 9.3×2 cm (Fig. 4A). Histology confirmed a Brunner's gland hamartoma (Fig. 4B, C).

The patient did not experience any serious complications from this procedure. He was started on proton pump inhibitor therapy and discharged after 7 days. His dizziness and melena resolved after the endoscopic procedure, and his anemia resolved after 1 month. At 4 months after discharge, his complete blood count had normalized. At 6 months, a repeat esophagogastroduodenoscopy did not find any evidence of recurrence or tumor remnants (Fig. 3D).

DISCUSSION

Brunner's gland hamartoma, also known as Brunner's gland adenoma or brunnneroma, is a rare benign tumor arising from the Brunner's glands of the duodenum.1 Brunner's glands were
first described by Brunner in 1688. These glands are branching actinotubular glands that arise in the duodenal submucosa. They secrete urogastrone, pepsinogen, and mucus to inhibit acid secretion and protect the duodenum from stomach acid. As such, they are crucial for preventing duodenal ulcers. The majority of these glands are seen in the first part of the duodenum, and they become smaller and less common towards the distal part of the duodenum. Their proportion in the duodenum decreases from 55% at birth to 35% by 50 years of age.2

Brunner's gland hamartomas typically develop into polypoid pedunculated masses as they enlarge. The exact pathogenesis is unknown. One hypothesis suggests that they may be related to irritation of the local lining, chronic pancreatitis, *Helicobacter pylori* infection, or parasympathetic activity. It is often difficult to differentiate between Brunner's gland hyperplasia and a Brunner's gland hamartoma.3 However, lesions smaller than 5 mm, whether single or multiple, are defined as Brunner's gland hyperplasia, whereas lesions larger than 5 mm are termed hamartomas.3

Brunner's gland hamartomas are most commonly observed between the ages of 50 and 60 years, and the incidence is similar in both genders.3 Most patients are asymptomatic, but some may present with non-specific symptoms such as abdominal bloating, abdominal pain, and nausea. Brunner's gland hamartomas can be found incidentally during esophagastroduodenoscopy or on imaging.6 Although symptomatic lesions are uncommon, patients can present with hemorrhage or obstruction. Gastrointestinal bleeding can occur because of ulcers or tumor erosion, causing in melena, iron deficiency anemia, fatigue, and in rare cases, hematemesis.5 They are mostly found in the duodenal bulb (57%), which may be due to the vascular damage and stress from gastrointestinal motility present here. However, they can also be seen in the second (27%) and third (7%) parts of the duodenum.6 In this case, the
patient was referred to our hospital after two episodes of large amounts of melena requiring blood transfusions. He was then found to have a large, elongated and pedunculated mass on the surface of the duodenal bulb during esophagogastroduodenoscopy.

Patients may experience obstruction when a mass becomes too large, or when there is diffuse hyperplasia. These patients often present with persistent nausea, vomiting, weight loss, epigastric pain, abdominal pain, and early satiety. Other uncommon symptoms include obstructive jaundice and intussusception due to the location and size of the tumor. These patients tend to have hamartomas exceeding 2 cm in size. In our patient, the Brunner's gland hamartoma had a thick stalk, which caused near-complete obstruction, resulting in the obstructive symptoms of nausea and vomiting. Several diagnostic tools including esophagogastroduodenoscopy and endoscopic ultrasound were used to identify the submucosal polypoid mass.

Brunner's gland hamartomas have characteristic histological features. They consist of an unusual mix of normal tissue and Brunner's glands, adipose cells, lymphoid cells, as well as ductal tissue. Dysplasia is not observed.

The differential diagnosis for duodenal mass lesions include leiomyomas, adenomas, lipomas, adenocarcinomas, carcinoid tumors, lymphomas, leiomyosarcomas, pancreatic or ampullary tumors, and the lesions of Peutz-Jeghers syndrome. Treatment depends on tumor size, symptoms, and the possibility of malignancy. Asymptomatic Brunner's gland hamartomas do not usually require treatment because their neoplastic potential is low. However, large or symptomatic lesions that cause obstruction or hemorrhage need to be removed. This can be done either endoscopically or surgically. Endoscopic treatment is preferred for pedunculated lesions; however, surgical resection is necessary if endoscopic methods fail, particularly if there is a suspicion of malignancy.

In this present case, endoscopic removal proceeded successfully despite the size of the tumor. The mass was too large to fit into the conventional snare, and had to be pulled into the stomach and removed with two partial snare polypectomies. Following this, endoscopic submucosal dissection was performed after the U-turn in the duodenal bulb. This case shows that a non-malignant lesion can have significant clinical ramifications, and that Brunner's gland hamartomas can be treated successfully by careful use of endoscopic procedures.

Conflicts of Interest

The authors have no financial conflicts of interest.

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