CASE REPORT

Inflammatory myofibroblastic tumors of the duodenum

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Summary Inflammatory myofibroblastic tumors (IMTs) are rare soft-tissue tumors that can occur at virtually any anatomical site. We report the case of a 58-year-old male with an IMT of the fourth part of the duodenum who presented with signs and symptoms of high intestinal obstruction and bilious vomiting. The patient underwent a surgical resection of the fourth part of the duodenum with end-to-end duodenojejunal anastomosis. The follow-up period of 6 months was uneventful with no evidence of recurrence. According to our knowledge, only six cases of duodenal IMTs have been reported in the literature thus far, and this is the first report of a duodenal IMT sited at the fourth part of the duodenum. The duodenum is among the rarest sites of IMTs. Signs and symptoms resulting from diagnostic imaging investigations are nonspecific and inadequate to obtain diagnosis accurately. In most cases, surgical treatment is considered a cure for IMTs. There is no evidence of deaths caused by duodenal IMT. IMT of the duodenum is a possible diagnosis in differential diagnosis of tumor-like lesions of the duodenum.

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1. Introduction

Inflammatory myofibroblastic tumors (IMTs) are rare tumors histologically composed of spindle myofibroblasts and an inflammatory infiltrate dominated by plasma cells, lymphocytes, and eosinophils.1 According to the World Health Organization, IMTs belong to a group of soft-tissue tumors, a subset of fibroblastic/myofibroblastic tumors.2 These principally occur in soft tissues and visceral organs at possibly any anatomical location. Nevertheless, duodenal IMTs remain an extremely rare condition. We report a case of IMT of the fourth part of the duodenum. Only six cases of...
2. Case report

A 58-year-old male who presented with a 1-week history of an intermittent epigastric pain, bilious vomiting, and unintentional weight loss of 10 kg in 4 weeks was admitted to the Department of Surgery of Clinical University Hospital Center. His medical history was significant for gastroesophageal reflux disease (GERD), Helicobacter pylori gastritis, Gilbert’s syndrome, and acute pancreatitis of unknown exact cause 2 months ago, which was successfully treated conservatively. Multiple abdominal ultrasonographies (USs) and multislice computed tomography (MSCT) scans excluded cholelithiasis and there was no evidence of alcohol abuse in the patient’s history (gamma glutamyl transferase: 17 U/L). Other causes of acute pancreatitis could neither be proven with certainty nor excluded.

A physical examination demonstrated abdominal bloating and visible distension with pain on palpation of the epigastric area of the abdomen. There was no jaundice, fever, or anemia. Laboratory tests revealed slightly elevated levels of aspartate transaminase (46 U/L), alanine transaminase (72 U/L), total bilirubin (58 µmol/L), and direct bilirubin (10 µmol/L). Other laboratory examinations analyzing the levels of C-reactive protein, erythrocyte sedimentation rate, complete blood count, coagulation factors, urea, creatinine, lipidogram, serum amylase, electrolytes including calcium, phosphorus, and magnesium were all within normal limits. Tumor markers such as cancer antigen 19-9, alpha-fetoprotein, and cancer antigen 125 were all negative.

Abdominal X-ray, abdominal US, and esophagogastroduodenoscopy showed no significant findings. Contrast imaging of the small intestine displayed a severe stenosis of the ascending duodenal portion (D4) measuring 0.8 cm in diameter with normal duodenal mucosal folds and normal morphology of the horizontal and descending portions of the duodenum (Fig. 1).

The MSCT scan of the abdomen revealed a filling defect in the fourth part of the duodenum measuring 4.3 × 3.6 cm² and dilatation of the second and third part of the duodenum. Triangle-shaped infiltration area of the adipose tissue measuring 2.1 × 1.9 cm² was identified cranial to the fourth part of the duodenum (Fig. 2). No regional lymphadenopathy or focal lesions of parenchymatous organs of the abdomen suggestive of metastatic lesions were identified.

Because imaging findings could not exclude malignancy and the mass appeared to be resectable, a surgical exploration was performed through an upper midline laparotomy. A firm, elastic tumor measuring 4 cm was found in the fourth part of the duodenum, adjacent to the paraduodenal adipose tissue and peritoneum surrounding the duodenojejunal flexure. The head of the pancreas and the hepatoduodenal ligament were intact. After the right colon and the mesenteric root mobilization, superior mesenteric artery 5 cm in length was exposed, and resection of the duodenojejunal flexure measuring 15 cm in length with 2-cm surgical resection margin and clearance of loco-regional lymph nodes were performed (Fig. 3). End-to-end duodenoojejunal anastomosis with single-layer continuous suture was made for reconstruction. Macroscopic pathology demonstrated a segment of the duodenum 15.5 cm in length and 3.2 cm in diameter with exophytic polyloid tumor measuring 2.1 × 1.6 cm² arising from the duodenal wall (Fig. 4). Histologically, the tumor was composed of spindle-shaped interspersed myofibroblasts separated with fibrous stroma infiltrated by predominantly mononuclear inflammatory cells (Figs. 5 and 6). The tumor cells were immunohistochemically positive for actin, vimentin, AE1/AE3, and negative for nonspecific esterase, CD34, CD117, S-100, and DOG1. The resection margins were clear of malignancy. In the surrounding adipose tissue, seven lymph nodes were found, which had a diameter from 0.4 to 1.2 cm, uninvolved with tumor tissue. A postoperative
course was uneventful with complete resolution of patient’s symptoms. Six months after the surgery the patient remained asymptomatic with no evidence of recurrence detected by the MSCT scans.

3. Discussion

Different terms for IMT have been used— inflammatory pseudotumor,9 fibrous xanthoma, plasma cell granuloma,10 pseudosarcoma, lymphoid hamartoma, myxoid hamartoma,11 inflammatory myofibrohistiocytic proliferation, pseudosarcomatous myofibroblastic proliferation.12 The term “inflammatory pseudotumor” has for many years been used for any clinically, macroscopic, or microscopic tumor-like lesion caused by inflammatory or reactive process.13 Later, the term has been applied only to neoplastic lesions microscopically characterized by the proliferation of spindle cells of mesenchymal origin with morphological characteristics of myofibroblasts and large inflammatory infiltration of different types of cells, usually with predominance of mature lymphocytes and plasma cells, regardless of etiology. However, during the last two decades, due to a greater understanding of the importance of myofibroblastic component in relation to inflammatory component1,14 and additional electron microscopic and immunohistochemical findings, this term has entirely replaced other terms.15

IMTs occur mainly in children and young adults, with more reports published on IMTs in adults recently.16,17 However, their etiology is unknown. Two basic hypotheses, reactive and neoplastic, are complemented with presumably infectious and autoimmune hypotheses. However, the exact infectious agents are not known. Some relate IMTs to Epstein–Barr virus, human herpesvirus-8, the bacterium Eikenella corrodens, and schistosomiasis.18–21 Others claim that previous surgical manipulation and

Figure 3  Surgical exploration of the tumor and the demonstration of the superior mesenteric artery.

Figure 4  Segment of the duodenum 15.5 cm in length and 3.2 cm in diameter with exophytic polypoid tumor measuring 2.1 × 1.6 cm arising from the duodenal wall.

Figure 5  Photomicrograph demonstrating an admixture of spindle-shaped and ovoid cells with a prominent inflammatory infiltrate (hematoxylin–eosin; low-power view; magnification 10×).

Figure 6  Photomicrograph showing a conspicuous admixture of lymphocytes and plasma cells (high-power view; magnification 40×).
chemotherapy or radiotherapy are potential causes.\textsuperscript{22–24} Some studies suggest the possible autoimmune origin of IMTs.\textsuperscript{25} According to these hypotheses, a medical history of acute pancreatitis, GERD, and H. pylori gastritis could be contributing factors in our case.

Based on the anatomical site, IMTs can be pulmonary and extrapulmonary.\textsuperscript{17} Extrapulmonary IMTs are most frequent in the abdomen including the mesentery and omentum.\textsuperscript{16} Other sites of the gastrointestinal tract are all visceral organs such as the appendix, Meckel’s diverticulum, and Vater’s papilla.\textsuperscript{2,23,24,26,27} Duodenal IMTs are among the rarest. To our knowledge, only six cases of duodenal IMTs have been reported, five of which had anatomical sites at the first and second parts of the duodenum, and one, particularly interesting, multiple IMT arising from the first part of the duodenum and extending up to the first part of the jejunum.\textsuperscript{3–8} According to the available literature, our patient is the first with IMT localization in the fourth part of the duodenum (Table 1). Specific risk factors are not defined and potential risk factors of duodenal IMT are provided in Table 1. Considering these cases, there seems to be no gender or age predilection (Table 1). Signs and symptoms of abdominal IMTs often mimic malignant tumors.\textsuperscript{22,28} Duodenal IMTs mostly present with upper gastrointestinal-tract obstructive symptoms, nausea, vomiting, pain in the epigastric area, anorexia, weight loss, malaise and fatigue, while in one of the cases night sweats were referred to as one of the significant symptoms. None of the cases presented with jaundice.\textsuperscript{3–8} Bilious vomiting, present in our patient, was not described previously (Table 1).

Diagnostic imaging techniques demonstrate tumor mass with its extent without specific features necessary for obtaining accurate diagnosis. Stricture of the duodenum, impossible passage, and irregular masses arising from different parts of the duodenum are most usual findings on radiological examination. It can be difficult to distinguish duodenal IMT from other groups of soft-tissue tumors or even pure (post)inflammatory strictures due to longstanding ulcer or diverticulitis. Stenosis due to duodenal ulcer can be divided into acute/active and chronic ulceration. Acute ulceration results in inflammatory edema and

<table>
<thead>
<tr>
<th>Author</th>
<th>Segment of duodenum</th>
<th>Age (y)/sex</th>
<th>Symptoms and signs</th>
<th>Treatment</th>
<th>Recurrence (treatment)</th>
<th>Potential risk factors</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fong et al\textsuperscript{3}</td>
<td>Second/third</td>
<td>57/F</td>
<td>Gastric outlet obstruction</td>
<td>Whipple operation</td>
<td>None</td>
<td>No report</td>
</tr>
<tr>
<td>Stringer et al\textsuperscript{4}</td>
<td>Second</td>
<td>5/F</td>
<td>Nonbilious vomiting; significant weight loss</td>
<td>Whipple operation with standard reconstruction</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>Mattei and Barnaby\textsuperscript{5}</td>
<td>First/second</td>
<td>13/M</td>
<td>Asymptomatic (incidentally diagnosed IMT)</td>
<td>Intravenous NSAID application (ketorolac); laparotomy (tumor mass regression)—Roux-en-Y gastrojejunostomy</td>
<td>None</td>
<td>Paratesticular rhabdomyosarcoma</td>
</tr>
<tr>
<td>Wynn et al\textsuperscript{6}</td>
<td>First</td>
<td>16/M</td>
<td>Abdominal pain; night sweats; malaise</td>
<td>En-bloc excision with primary closure of the duodenum</td>
<td>Yes (perioral corticosteroids; azathioprine)</td>
<td>No report</td>
</tr>
<tr>
<td>Mirshemirani et al\textsuperscript{7}</td>
<td>First</td>
<td>13/M</td>
<td>Abdominal pain; weight loss; epigastric mass</td>
<td>Surgical resection of the tumor; duodeno-duodenostomy</td>
<td>None</td>
<td>No report</td>
</tr>
<tr>
<td>Xiang et al\textsuperscript{8}</td>
<td>Multiple IMT (from the first part of the duodenum to starting part of the jejunum)</td>
<td>20/M</td>
<td>Intermittent right epigastric pain; nausea; vomiting</td>
<td>Whipple operation</td>
<td>None</td>
<td>Calculous cholecystitis</td>
</tr>
<tr>
<td>Current study</td>
<td>Fourth</td>
<td>58/M</td>
<td>Bilious vomiting; epigastric pain; significant weight loss</td>
<td>Resection of the duodenoejejunal flexure with termino-terminal anastomosis</td>
<td>None</td>
<td>Acute pancreatitis; GERD; H. pylori gastritis</td>
</tr>
</tbody>
</table>

GERD = gastroesophageal reflux disease; IMT = inflammatory myofibroblastic tumor; NSAID = nonsteroidal anti-inflammatory drug.
pylorospasm. Signs and symptoms of stenosis may regress if the ulcer heals under medical treatment, which is not noticed in IMT cases. Long-standing ulceration can present with radiological demonstration of scarring and deformities of the duodenum. In most cases, there are no signs of tumorous mass. Past medical history of peptic ulcer-type pain for 10 or more years strongly suggests the precise diagnosis contrary to duodenal diverticula, which are the uncommon site of inflammation due to their large size and mostly sterile content of the duodenum. The incidence of duodenal diverticula is approximately 22% with the incidence of complicated duodenal diverticulum estimated from 0.03% to 5%. Inflammation of the diverticulum can be distinguished before surgery by meticulous CT scan analysis. The CT findings vary between saccular and tubular configuration, diverticular content (homogenous liquid, liquid debris, and hydroaeric levels), wall (thickened, enhanced, and thin), retroperitoneal infiltration, free air, and signs of common bile duct dilatation. 

Definitive diagnosis of IMTs is confirmed by histopathological analysis and immunohistochemical tests of surgically removed specimen. Some histopathological IMT patterns such as compact fascicular spindle cells with myxoid or collagenized regions intermingled with inflammatory cells may resemble fibromatoses. Fibromatoses colli, juvenile hyaline fibromatosis, superficial fibromatosis, desmoid-type fibromatosis, and lipofibromatosis are well-defined soft-tissue tumors histopathologically differentiated from IMT. These benign neoplasms can extend into adjacent tissues, and sometimes have a tendency toward local recurrence, but do not metastasize. Except superficial fibromatosis and desmoid-type fibromatosis, other entities of this group occur mostly in infants or early childhood with no sex predilection. Superficial and desmoid-type fibromatosis occur mostly in adults with slight predominance in men of first entity and in women of second entity. To the best of our knowledge and according to available literature, no cases of fibromatoses of duodenum were reported so far. 

The basic principle for IMTs treatment is complete surgical resection. In nonreseetable tumors, a palliative treatment regimen with radiotherapy and nonsteroidal anti-inflammatory drugs (NSAIDs) can be applied. However, guidelines for this treatment approach do not exist. Some studies recommend the use of NSAIDs, corticosteroids, and immunosuppressive drugs after the surgery. One case of the duodenal IMT recurrence was successfully treated by immunosuppressive drugs (Table 1). In conclusion, the duodenum is among the rarest sites of IMTs. Signs and symptoms as results of diagnostic imaging investigations are nonspecific and inadequate to accurately obtain diagnosis. The diagnosis can be confirmed only by a histopathological analysis and immunohistochemical tests of the surgically removed specimens. In most cases of duodenal IMTs, surgical treatment is considered to cure the disease. There is no evidence of deaths caused by duodenal IMTs. All reported patients with duodenal IMTs had an uneventful recovery. Duodenal IMT should be considered in differential diagnosis of tumor-like lesions of the duodenum.

References


