Adenocarcinoma arising from a heterotopic pancreas in the first portion of the duodenum: a case report

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Abstract

Background: Heterotopic pancreas (HP) is defined as pancreatic tissue in organs with no anatomical continuity with the orthotopic pancreas. Based on the number of cases reported in the literature between the year 2000 and 2020, HP is rarely found causing malignant transformation of the duodenum. We herein report a case of adenocarcinoma arising from the HP in the first portion of the duodenum.

Case presentation: A 77-year-old Japanese man presented to our hospital with epigastric pain. Despite having undergone laparoscopic surgery for early sigmoid colon cancer a month earlier, serum levels of tumor-specific antigens, such as CA19-9, were elevated. After undergoing a series of radiologic examinations, the first portion of the duodenum was found thickened. However, a biopsy of the lesion showed no malignancy. Four months later, follow-up computed tomography (CT) scans showed that the lesion was thicker and involved the gastroduodenal artery (GDA), suggesting tumor invasion. A new biopsy did not detect the malignancy. However, serum tumor-specific antigen levels increased, especially duke pancreatic monoclonal antigen type 2 (5287 U/mL), in the absence of tumor in the orthotopic pancreas. The follow-up CT imaging showed a malignant tumor in the first portion of the duodenum. Five months later, we performed a subtotal stomach-preserving pancreaticoduodenectomy (SSPPD) for duodenal or HP cancer in the first portion of the duodenum, finding a lesion from the pyloric bulbs to the first portion of the duodenum, which invaded the adjacent pancreas and GDA. The pathological examination of the specimens revealed adenocarcinoma arising from HP. Nine months after surgery, no recurrence was found by radiologic imaging or tumor-specific antigen laboratory testing.

Conclusions: HP adenocarcinoma is rare and difficult to diagnose preoperatively due to its submucosal location. Therefore, a careful follow-up with blood testing and radiologic imaging, as well as diagnostic surgery, is recommended.

Keywords: Heterotopic pancreas, Adenocarcinoma, SSPPD

Background

Heterotopic pancreas (HP), also called ectopic or aberrant pancreas, is defined as pancreatic tissue located in organs with no anatomical continuity with the orthotopic pancreas. HP commonly occurs at the submucosal layer of the gastrointestinal tract and is difficult to diagnose preoperatively. On the basis of abdominal surgery evidence, the incidence of HP is estimated to range from 0.25 to 1.2% [1, 2]. Notably, the incidence of malignant transformation in HP is extremely rare. To our knowledge, only 12 cases of HP cancer of the duodenum are found in the literature since the year 2000 after performing a PubMed search (keywords: heterotopic OR ectopic OR aberrant pancreas, carcinoma). Herein, we describe a case of adenocarcinoma arising from HP in the first portion of the duodenum and include a review of the literature.
Case presentation
A 77-year-old Japanese man presented to our hospital with epigastric pain and a medical history of type 2 diabetes and duodenal ulcer. One month earlier, the patient had undergone laparoscopic surgery for early stage sigmoid colon cancer, classified as pathological T1N0M0 stage I according to the 7th edition of the Union for International Cancer Control (UICC)’s TNM classification of malignant tumors. The results of the physical examination were unremarkable and his vital signs were within the reference range. Laboratory test results showed elevated serum levels of tumor-specific antigen: carbohydrate antigen 199 (CA19-9), 1130 U/mL (reference range, 0–37 U/mL). Enhanced multi-detector computed tomography images showed a significant gastric distension and slight thickening of the first portion of the duodenum (Fig. 1). Esophagogastroduodenoscopy results revealed an ulcer scar with slight stenosis and no evidence of malignancy after biopsy (Fig. 2). The patient was followed up for a short term on an outpatient basis. We evaluated the thickening lesion by laboratory testing, radiologic imaging, and endoscopic examination every 4 months. Four months later, his subjective symptoms remained unchanged, but follow-up CT showed thickening of the stenosis-surrounding tissue with the involvement of the gastroduodenal artery (GDA), suggesting tumor invasion (Fig. 3). Endoscopy examination was challenging due to the difficulty in passing through the thickening lesion (Fig. 4). CA19-9 serum levels remained elevated, and serum level of duke pancreatic monoclonal antigen type 2 (DUPAN-2), which is a pancreatic tumor-specific antigen, was especially elevated, 5287 U/mL (reference range, 0–37 U/mL). Because CT images showed no evidence of the tumor in the orthotopic pancreas, we suspected that the malignancy could be located in heterotopic pancreas tissue. After 5 months of diagnostic intervention, we performed a subtotal stomach-preserving pancreateodudodenectomy in the first portion of the duodenum to treat the duodenal and presumed heterotopic pancreatic cancer.

Intraoperative findings included the identification of an elastic hard tumor at the duodenal bulbs invading the adjacent pancreas and GDA (Fig. 5). The gastrointestinal reconstruction strategy involved the modified Child method with Kakita’s technique for pancreatojejunostomy. The resected specimen had severe stenosis from the pyloric bulb to the first portion of the duodenum, whereas the mucosa appeared normal (Fig. 6). The tumor was white-yellow in color on its cut surface and had a diameter of 55 mm. A moderately differentiated tubular adenocarcinoma continuous with the heterotopic pancreatic tissue expanded from the submucosal layer to

Fig. 1 Abdominal CT scan showing a thickened wall in the first portion of the duodenum (arrow)

Fig. 2 Gastroscopic image showing a slight pyloric stenosis with a small ulcer (arrow). The lesion was covered with normal mucosa (yellow dotted circles)

Fig. 3 Four months later, abdominal CT scan showing a more thickened wall than that observed in previous CT scans (arrow heads). The lesion involved the surrounding tissue of the gastroduodenal artery, suggesting invasion (arrow)
the muscularis propria of the first portion of the duodenum (Fig. 7a–c). The pancreatic tissue consisted of acini, ducts, and Langerhans islet cells (Fig. 7d) and was classified as Heinrich’s type I HP. Finally, pathological diagnosis was an adenocarcinoma arising from the HP in the first portion of the duodenum.

The postoperative course was uneventful, and the patient was discharged 25 days after surgery. One month after surgery, serum carcinoembryonic antigen level was 2.9 U/mL; CA19-9, 1.133 U/mL; and DUPAN-2, 1.613 U/mL. The patient underwent 5 courses of oral tegafur-gimeracil-dihydropyrimidine dehydrogenase (TS-1) as adjuvant chemotherapy and has remained free of disease 9 months after surgery, with no recurrence found by radiologic imaging.

Discussion

Heterotopic pancreas (HP), also called ectopic or aberrant pancreas, is defined as pancreatic tissue in organs with no anatomical continuity with the orthotopic pancreas. Shultz first reported the existence of HP in 1727. Almost 200 years later, in 1909, Von Heinrich classified HP tissue into four histopathologic types, by the presence or absence of pancreatic ducts, acini, and islet cells [3, 4]. Commonly, HP is a benign disease comprising submucosal tumors (SMT) in the gastrointestinal tract, in organs such as the stomach, duodenum, jejunum, mesocolon, and Meckel’s diverticulum. Thus, HP initially identified as SMT by radiologic or endoscopic imaging is rarely biopsied for a definitive diagnosis, and only follow-up imaging is performed because it rarely causes clinical symptoms. Conversely, when clinical symptoms occur, they include inflammation, bleeding, obstruction, and malignant transformation [5]; therefore, a biopsy for a definitive diagnosis as well as treatment is needed. Guillou et al. and Makhlof et al. independently reported that the incidence of malignancy due to HP was 0.7% and 1.8%, respectively [6, 7]. Malignant transformation of the HP needs to fulfill the following histopathologic criteria for diagnosis [6], as occurred in our case:

1. Tumor found within or close to the heterotopic pancreatic tissue
2. Direct transition observed between the pancreatic structures and the carcinoma (malignant transformation of the HP must be differentiated from a metastatic deposit or a neoplastic invasion from a neighboring digestive cancer, especially from the stomach, biliary tract, and heterotopic pancreas)
3. Non-neoplastic pancreatic tissue must comprise at least fully developed acini and ductal structures.
We found 12 other case reports of HP cancer on PubMed records from the year 2000 to 2020, as shown in Table 1 [3, 5, 7–15]. In those reports, the patients’ mean age was 69.7 years (range, 56–86 years). Eight patients were male and 5 were female. As far as the other reports have described, there were 7 cases of type I and one case each of type III and type IV as per the Heinrich classification of HP. All patients presented with clinical symptoms, such as abdominal pain or vomiting because of obstruction caused by the tumor. The mean tumor size was 30.4 mm (range, 12–55 mm). In few cases, serum levels of tumor-specific antigens were not elevated. Almost all patients had preoperative suspicion of malignancy, and only 2 could be confirmed by endoscopic ultrasonography (EUS). No cases of HP adenocarcinoma could be preoperatively diagnosed. Tumor location and the extent of invasion in the pancreas during the surgical procedure produced changes in the surgical plan. In addition, there were 2 cases of gastrectomy, 6 cases of pancreaticoduodenectomy (including SSPPD), and 5 cases of partial duodenectomy. Surgery with lymphadenectomy occurred in 9 cases, and 5 out of 9 cases had metastases to the lymph nodes. Although no standard chemotherapy for malignant HP has been established, 5 among the 13 cases received adjuvant chemotherapy. The mean time of progression-free survival (PFS) was 35 months.

In our case, because of the high levels of serum tumor-specific antigens, we suspected the existence of a malignant tumor. Remarkably, DUPAN-2 being a specific pancreatic tumor antigen, serves as a useful tool for detecting orthotopic pancreatic cancer as well as HP cancer. CT and endoscopy imaging examination initially showed the thickened wall in the first portion of the duodenum. However, several biopsies performed at different times could not reveal the location of the malignancy. Hence, the surgery decision was postponed during the 4 months of follow-up until a precise diagnosis could be achieved. Similar to the previous 12 reports, our case could not be preoperatively diagnosed by biopsy because the tumor existed in the submucosal layer, a difficult site to obtain a sample. Endo et al. [16] suggested the usefulness of preoperative EUS-guided fine-needle aspiration (EUS-FNA) to reveal submucosal malignancies, like HP cancer. Also, cytological examination with EUS revealed it in approximately 50% of the cases [17]. However, when tumor obstruction exists, it makes it difficult to pass through the lesion and perform a precise evaluation. In fact, only a few of the reported cases could be diagnosed using EUS-FNA. We believe that diagnostic surgery is still a good choice to evaluate the tumor location and malignancy and preferable to perform with a lymphadenectomy when considering malignant cases.
| Case | Year | Author | Age | Sex | Location | Size (mm) | Biopsy | Tumor-specific antigen | Diagnostic approach | Lymphadenectomy | Lymph node metastasis | Heinrich classification | Chemotherapy | PFS (months) |
|------|------|--------|-----|-----|----------|----------|--------|------------------------|-------------------|----------------|----------------------|------------------------|--------------|--------------|
| 1    | 2007 | Tison  | 72  | M   | 2nd      | N.D      | Negative | N.D                    | Ope (PD)          | +              | N.D                  | N.D                    |              | N.D          |
| 2    | 2007 | Kawakami| 65  | F   | 2nd      | 12       | Not performed | WNL                  | Ope (SSPPD)       | N.D            | N.D                  | N.D                    |              | 19           |
| 3    | 2008 | Rosok  | 59  | F   | Proximal | 50       | Negative | N.D                    | Ope (lap-D)       | –              | N.D                  | Not performed            |              | 36           |
| 4    | 2010 | Inoue  | 75  | M   | 2nd      | 30       | Negative | WNL                    | Ope (PPPD)        | +              | III                  | Not performed            |              | 72           |
| 5    | 2010 | Bini   | 56  | M   | 1st      | N.D      | Adenocarcinoma | WNL                  | EUS-FNA → Ope (PD) | –              | I                    | Performed (N.D)      |              | N.D          |
| 6    | 2011 | Stock  | 79  | F   | 4th      | 30       | Negative | Chromogranin-A, 53 mg/mL | Ope (D)          | +              | I                    | Performed (N.D)      |              | N.D          |
| 7    | 2012 | Kinoshita| 62  | F   | 1st      | 34       | Negative | CEA, 8.1 U/mL          | Ope (PD)          | –              | I                    | Not performed            |              | 12           |
| 8    | 2013 | Ginori | 86  | F   | 1st      | 30       | Not performed | N.D                    | Ope (subTG)       | –              | I                    | Not performed            |              | N.D          |
| 9    | 2013 | Alireza| 58  | M   | 1st      | 27       | Not performed | N.D                    | Ope (D)          | –              | N.D                  | Performed (XELOX)      |              | 18           |
| 10   | 2014 | Endo   | 75  | M   | 2nd      | 22       | Adenocarcinoma | CEA, 55.4 U/mL         | EUA-FNA → Ope (SSPPD) | N.D         | IV                   | Not performed            |              | 60           |
| 11   | 2015 | Fukino | 62  | M   | 4th      | 15       | Negative | CEA, 548 U/mL               | Ope (D)          | N.D            | N.D                  | Performed (SP)              |              | N.D          |
| 12   | 2019 | Kaneko | 81  | M   | 1st      | 30       | Negative | WNL                    | Ope (DG)          | +              | I                    | Not performed            |              | 18           |
| 13   | 2020 | Our case | 77  | M   | 1st      | 55       | Negative | CEA, 1130 U/mL            | Ope (SSPPD)       | +              | I                    | Performed (TS1)        |              | 9            |

CEA carcinoembryonic antigen, CA carbohydrate antigen, DUPAN duke pancreatic monoclonal antigen, EUS-FNA endoscopic ultrasonography-guided fine-needle aspiration, N.D not dated, PFS progression-free survival, D (partial) duodenectomy, DG distal gastrectomy, TG total gastrectomy, PD pancreatoduodenectomy, PPDP pylorus-preserving pancreatoduodenectomy, SSPPD subtotal stomach-preserving pancreatoduodenectomy, SMT submucosal tumors, TS1 tegafur-gimeracil-dihydropyrimidine dehydrogenase, SP TS1+cisplatin, XELOX capecitabine+oxaliplatin, WNL within normal limits.
The recurrence rate of HP cancer is low. Similar to what has been observed in previous reports, the PFS of the HP adenocarcinoma was longer than that of orthotopic pancreatic cancer: this is probably because the tumor enlargement might cause earlier and more diverse symptoms, thus allowing better control of the disease.

Conclusions
HP adenocarcinoma is rare and difficult to diagnose due to its submucosal location. EUS is a useful procedure to detect SMT, except for lesions in which the tumor causes obstruction. Therefore, in cases where the primary or malignant lesions are difficult to detect, a careful and short-term follow-up, including blood testing and radiologic imaging, is needed. Additionally, diagnostic surgery should be considered as a choice for evaluating the location of the malignant tissue.

Abbreviations
CA: Carbohydrate antigen; DUPAN: Duke pancreatic monoclonal antigen; EUS: Endoscopic ultrasonography; EUS-FNA: Endoscopic ultrasonography-guided fine-needle aspiration; HP: Heterotopic pancreas; GDA: Gastroduodenal artery; PFS: Progression-free survival; SSPPD: Subtotal stomach-preserving pancreatoduodenectomy; SMT: Submucosal tumors; TS-1: Tegafur-gimeracil-dihydropyrimidine dehydrogenase

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Authors’ contributions
TM wrote the manuscript and performed the literature search. TM and TT treated and observed the patient. YN performed the pathological examinations. TT and MT supervised the preparation of this case report. All authors have read and approved the final manuscript.

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Availability of data and materials
The datasets supporting the conclusions of this article are included in the article.

Ethics approval and consent to participate
Informed consent was obtained from the patient to publish the details of the case, and his identity has been protected.

Consent for publication
The institutional ethics committee approved the publication of this case report.

Competing interests
The authors declare that they have no competing interests.

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