CASE REPORT

Osteoclast-like Giant Cell-type Pancreatic Anaplastic Carcinoma Presenting with a Duodenal Polypoid Lesion

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Abstract:
Osteoclast-like giant cell-type (OCGC) anaplastic carcinoma is a rare variant of pancreatic ductal adenocarcinoma, and its imaging characteristics and progression pattern have not been fully clarified. The patient was a 73-year-old man who had been incidentally found to have a pancreatic head tumor. Computed tomography demonstrated a 3-cm marginally enhanced mass at the pancreatic head, continuing toward the duodenum. Diffusion-weighted magnetic resonance imaging showed a retained diffusion capacity. Duodenoscopy revealed a 1.5-cm polypoid lesion, covered by a dirty coat, near the major papilla. Surgical material revealed OCGC pancreatic anaplastic carcinoma protruding to the duodenum, accompanied by multiple hemorrhagic foci and hemosiderin precipitations.

Key words: pancreas, duodenum, anaplastic carcinoma, osteoclast-like giant cell, hemorrhaging

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Introduction

Anaplastic carcinoma is a rare undifferentiated variant accounting for only 0.08% of total pancreatic cancers, 0.5% (1) of resected pancreatic cancers, and 1.4% (2) of resected pancreatic ductal adenocarcinoma (PDAC). Pancreatic anaplastic carcinoma (PAC) shows oncogenic molecular alterations similar to those of PDAC (i.e. activation of K-ras oncogene and inactivation of suppressor genes, including CDKN2A/p16, TP53, and DPC4) (3) and is thought to represent a subtype of PDAC. PAC is thought to show an aggressive biological behavior; however, the patient survival can be significantly improved by surgical resection (4) or depending on the presence of osteoclast-like giant cells (OCGCs) (5) and the purity of OCGCs (i.e. no epithelial neoplasm) (3). However, studies showing clinical images and the progression pattern of PAC have been lacking. We herein report a case of OCGC-type PAC that presented with a protruding duodenal mass and developing pancreatitis, which enabled us to make a preoperative diagnosis.

Case Report

A 73-year-old man was referred to our hospital for the investigation of a mass lesion located at the pancreatic head. He had a history of diabetes, hyperlipidemia, and lung cancer (bronchioloalveolar carcinoma, 24 mm, pT1bN0M0, stage IA) at 71 years of age, for which he had undergone right-upper lobectomy and subsequent adjuvant chemotherapy with tegafur/uracil. His blood tests showed slight anemia (hemoglobin: 12.9 g/dL, normal: 13.5-17.6 g/dL) and elevated levels of serum amylase (431 U/L, normal: 37-125 U/L), hemoglobin A1c (6.8%, normal: 4.3%-5.8%), and cancer antigen 19-9 (CA19-9)(294 U/ml, normal: <37 U/mL). The current pancreatic mass had been incidentally detected by computed tomography (CT) conducted to follow his post-operative lung lesion. His family history was not notable.
Abdominal ultrasonography (Fig. 1) revealed an irregular-margin, low-echoic pancreatic mass, 30 mm in size. Enhanced CT (Fig. 2) demonstrated a heterogeneously ill-attenuated mass located at the pancreatic uncinate process and continuously protruding into the duodenum. The enhancement recovered slightly in the late phase, especially at the tumor margin. Only the peripancreatic lymph nodes were slightly swollen, they can thus be interpreted as either “inflammatory or non-cancerous” or “cancer metastasis-positive”. Magnetic resonance imaging (MRI) (Fig. 3) showed a low-signal-intensity lesion in both T1- and T2-weighted images. Diffusion-weighted imaging (DWI) showed a retained diffusing capacity at the lesion when compared with the surrounding pancreas.

During these imaging examinations, the patient complained of intermittent abdominal pain; this was suggestive of pancreatitis, based on the elevated level of serum amy-
and neural invasion was recognized. The two nearest lymph
tations into the lymph duct and peripheral vein were positive,
type anaplastic carcinoma of the pancreas. Cancer permea-

Figure 4. Endoscopic retrograde pancreatography. A: An indigocarmine-sprayed duodenoscopic
view showed a protruding tumor, covered with a dirty coat, at the anal side of the major papilla. B: Pancreatography visualized faint stenosis of the main pancreatic duct (arrow) and a floating filling
defect nearby (small arrow). C: A forceps biopsy specimen showed undifferentiated carcinoma with
many osteoclast-like giant cells (Hematoxylin and Eosin staining, ×100).

lase. Duodenoscopy (Fig. 4A) revealed an approximately 15-
mm polypoid lesion, covered with a dirty coat, at the anal side of the major papilla. Endoscopic retrograde pancrea-
tography demonstrated the irregularly narrowed main pan-
creatic duct (MPD) and a crumbling filling defect inside the
MPD (Fig. 4B). Endoscopic naso-pancreatic duct drainage
(ENPD) was performed to relieve pancreatitis. The patient’s
serum amylase level had normalized by the next day (76 U/
L), and his abdominal pain had disappeared. Histology of
the forceps biopsy specimen from the duodenal polyp re-

Discussion

PAC is a rare variant of pancreatic cancer, accounting for
1.4% (2) of resected PDAC. OCGC-type PAC is even rarer, accounting for only 0.2% of pathology-confirmed
PDAC (5). Hoshimoto et al. analyzed 60 previously reported
cases of Japanese PAC (mostly written in the Japanese lan-
guage) and summarized the characteristics of these PAC
cases. They presented a mean age of 61.5 years (range: 32
to 85 years), a slight male predominance (63%), main com-
plaints of abdominal pain (48%) and back pain (17%), fre-
quent elevation of serum CA19-9 (55%), almost equal distri-

A macroscopic view of the surgical material showed a heterogeneously hemorrhagic whitish mass (3 cm) and the
continuing brownish tumor protruding into the duodenal lu-
men (2 cm) (Fig. 5A). The macroscopic brownish areas were microscopically hemorrhaging and hemosiderin pre-
cipitation (Fig. 5B). The tumor consisted mostly of undiffer-
entiated carcinoma containing many OCGCs (Fig. 5C) and
partially of well-differentiated adenocarcinoma (Fig. 5D).
The pathological diagnosis was osteoclast-like giant cell-
type anaplastic carcinoma of the pancreas. Cancer permea-
tions into the lymph duct and peripheral vein were positive,
and neural invasion was recognized. The two nearest lymph
nodes were positive for carcinoma metastasis. Immunostain-
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TP53 diffusely overexpressed in the tumor cells. Cytokeratin
AE1/AE3 and MUC1 were similarly positive in the differenti-
ted adenocarcinoma components but negative in the dedif-
erentiated cancer cells. Immunostaining of CA19-9 and
Vimentin was diffusely positive in the cancer cells. KP-1
was positive not only in the inflammatory cells but also in
the cancer cells, including OCGCs. The Ki-67 labeling in-
dex was 42% as the average of 3 randomly chosen areas
consisting of 500 cancer cells each (Fig. 5E, F, G and H).

The patient’s postoperative course was uneventful, and he
was discharged after two weeks. The patient was transferred
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with gemcitabine. However, five months after the surgery,
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Figure 5. Pathological view of the resected specimen. A macroscopic view of the resected materials (A) showed a pancreatic tumor (small arrow) and tumor continuously protruding into the duodenum (arrow). The cut surface of the tumor was mostly accompanied by brownish areas or hemorrhaging. The main pancreatic duct (black arrowhead) was involved with the tumor at the marginal site. The major papilla (yellow arrow) was located at the margin of the duodenal diverticulum (asterisk). A low-power view of the protruding duodenal tumor and duodenal wall (B) showed hemorrhagic lesions (arrow) with hemosiderin precipitates [inset, Hematoxylin and Eosin (H&E) staining, ×100] and adenocarcinoma invading the submucosa (small arrow) (H&E staining, ×12.5). A high-power view of the tumor demonstrated many osteoclast-like giant cells within the undifferentiated carcinomas and red blood cells (C) (H&E staining, ×200, a bar size: 50 μm) and a well-differentiated adenocarcinoma component (D) (H&E staining, ×100, a bar size: 100 μm). Immunostaining of CDKN2A/p16 was diffusely repressed (E) (×200), whereas that of TP53 was overexpressed in the cancer cells (F) (×200). The KP-1 protein expression was positive not only in the inflammatory cells but also in the cancer cells, including the osteoclast-like giant cell-type anaplastic carcinoma cells (G) (×200). The Ki-67 labeling index was 42% in the randomly selected area (H) (×200).
bution within the pancreas (head: 53%, body to tail: 42%, and entire pancreas: 5%), and a median size of 6 cm (range: 1.5 to 24 cm) (6).

Little is known about the imaging features and progression patterns of this rare tumor type (7, 8). Fukukura et al. analyzed the CT and MRI scans of seven cases of OCGC-type PAC and reported several findings: a smooth-margined low-attenuated mass on CT and a low signal intensity on T1-weighted, T2-weighted, and diffusion-weighted imaging, reflecting hemosiderin precipitates (7). Khashab et al. used endoscopic ultrasonography (EUS) to examine six cases of PAC; five cases were observed as heterogeneous low-echoic masses. Four of these six cases demonstrated PAC components in EUS-fine-needle aspiration (EUS-FNA) samples (9).

In the current case, the imaging findings for CT, MRI, and ultrasonography were quite similar to those of the cases reported by Fukukura et al (7). Multiple foci of hemosiderin precipitates were also recognized within the tumor, as reflected in the low signal intensity on all three MRI modalities (7).

Hoshimoto et al. reported that the macroscopic view of the resected materials of PAC often indicate intra-tumor hemorrhaging (77%) and sometimes cystic formation (33%) (6). These findings are suggestive of the natural course of tumor development causing hemorrhagic necrosis, as the hemorrhaging changes into a hemosiderin clot, while necrosis leads to cyst formation. In our case, we observed hemorrhaging and diffuse hemosiderin precipitates in both the pancreatic and duodenal lesions, but no prominent cystic changes were apparent, suggesting a relatively early stage of detection by postoperative screening. The viable interstitial and tumor cells predominantly seen at the marginal area were thought to be reflected as marginal enhancement on late-phase images (Fig. 2).

According to a report from the Mayo Clinic, the prognosis in cases of PAC is poor if the lesions cannot be treated with surgery (overall survival [OS]: 34.1 months in resected cases vs. 3.3 months in non-resected cases, P=0.001) (4). When surgically resected, in a condition-matched comparison, patients had a survival similar to ordinary PDAC (44.1 vs. 39.9 months, P=0.763) (4). This trend was the same when analyzing the American National Cancer Data Base (NCDB); the post-surgery 5-year survival rate was 22% in the PAC group and 17% in the PDAC group (P<0.032) (1). However, comparison of PACs with and without OCGCs indicated that tumors with OCGCs portended a significantly better prognosis (5-year survival: 50% vs. 15%, P < 0.001) (1). Another clinicopathological study by Luchini et al. also showed that the post-operative OS was significantly better in patients with pure OCGC-PAC (n=9) than in those with OCGC-PAC with a PDAC component (n=13) (median OS: 36 months vs. 15 months, P=0.04) (3). The current case of OCGC-PAC was accompanied by an adenocarcinoma component (Fig. 5D) and showed aggressive progression resistant to adjuvant chemotherapy with gemcitabine.

A recent European and American multicenter study demonstrated a significantly higher incidence of PD-L1 positivity in OCGC-type PACs (63%) than in ordinary PDACs (P=0.04) and a worse prognosis in OCGC-type PAC cases with PD-L1 expression than in those without (hazard ratio: 3.98, P=0.12) (10). However, in cases of non-small-cell lung cancers, immune-checkpoint inhibitors, such as pembrolizumab and atezolizumab, are more effective in PD-L1-positive cases (11, 12). In the future, we need to keep these data in mind when treating patients with OCGC-type PACs.

In conclusion, we encountered a case of osteoclast-like giant cell-type pancreatic anaplastic carcinoma forming a duodenal protruding mass and causing pancreatitis. A relatively smooth demarcation of the tumor margin, ill (or marginal) enhancement, intra-tumorous hemorrhaging, and hemosiderin precipitation were viewed as characteristic findings of this tumor.

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References


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