

# World Journal of *Clinical Cases*

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## Primary duodenal dedifferentiated liposarcoma: A case report and literature review

Nah Ihm Kim, Ji Shin Lee, Chan Choi, Jong Hee Nam, Yoo Duk Choi, Hee Joon Kim, Sung Sun Kim

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### Abstract

#### BACKGROUND

Dedifferentiated liposarcoma (DDLPS) is an extremely rare neoplasm that exhibits various morphologies. The tumor is characterized by immunoreactivity to MDM2 and CDK4 and can be confirmed by detecting *MDM2* amplification *via* fluorescence *in situ* hybridization (FISH). Herein, we report an unusual case of DDLPS arising from the duodenum.

#### CASE SUMMARY

A 64-year-old man presented with repeated abdominal pain and weight loss. Radiologic studies revealed a mass of the duodenum involving the pancreas. The patient was treated with pylorus-preserving pancreaticoduodenectomy. Histologically, the tumor showed a high-grade sarcoma. Immunohistochemistry demonstrated that the tumor cells were positive for MDM2 and CDK4 expression. *MDM2* amplification was detected *via* FISH, leading to the final diagnosis of DDLPS. Following surgery, the patient was treated in the intensive care unit due to peritonitis, and died 60 d after surgery.

#### CONCLUSION

To the best of the authors' knowledge, this is the first case of primary duodenal DDLPS in Korea and the third case in the English-language literature. Care must be taken not to misdiagnose DDLPS as another high-grade tumor. Liposarcoma should be in the differential diagnosis list.

**Key Words:** Liposarcoma; Primary; Small bowel; Duodenum; Immunohistochemistry;

Case report

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**Core Tip:** Primary dedifferentiated liposarcoma (DDLPS) originating from the duodenum are rare and can be diagnosed based on histology, immunohistochemistry and *MDM2* amplification *via* fluorescence *in situ* hybridization. Differential diagnoses are required along with consideration of DDLPS.

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## INTRODUCTION

Liposarcoma is the most common form of malignant soft tissue tumor and tends to occur in the retroperitoneum, deep soft tissues of the trunk, and extremities[1]. Primary liposarcoma of the gastrointestinal tract is extremely rare, with only a few cases reported in the literature[2-10]. In this report, we describe a unique case of dedifferentiated liposarcoma (DDLPS) originating from the duodenum and review previously reported cases. Pathologists should keep liposarcoma in the differential diagnosis list.

## CASE PRESENTATION

### Chief complaints

A 64-year-old male presented with a 7-d duration of repeated abdominal pain.

### History of present illness

The patient also complained of 8 kg weight loss in 3 mo.

### History of past illness

The patient had been on antihypertensive medication for 20 years.

### Personal and family history

He had no other significant personal or family medical history.

### Physical examination

Physical examination revealed diffuse abdominal tenderness, but a palpable mass was not detected.

### Laboratory examinations

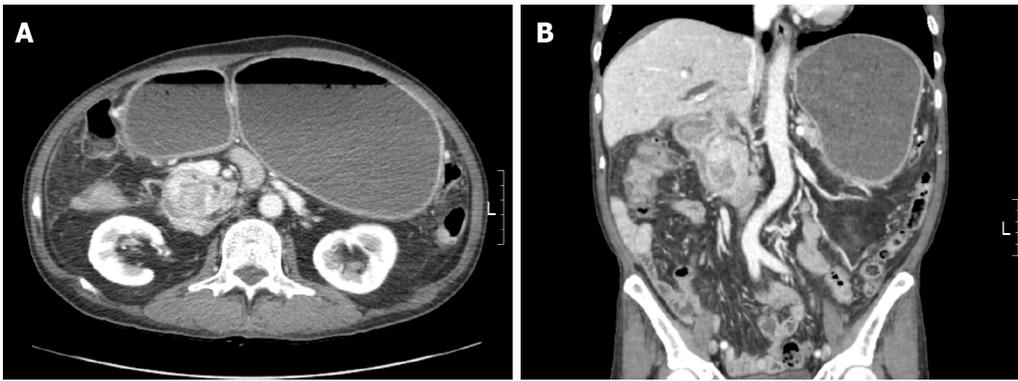
Laboratory findings showed slight elevation of aspartate aminotransferase (55 U/L). Tumor markers including carcinoembryonic antigen (2.13 mg/mL), were within the normal range.

### Imaging examinations

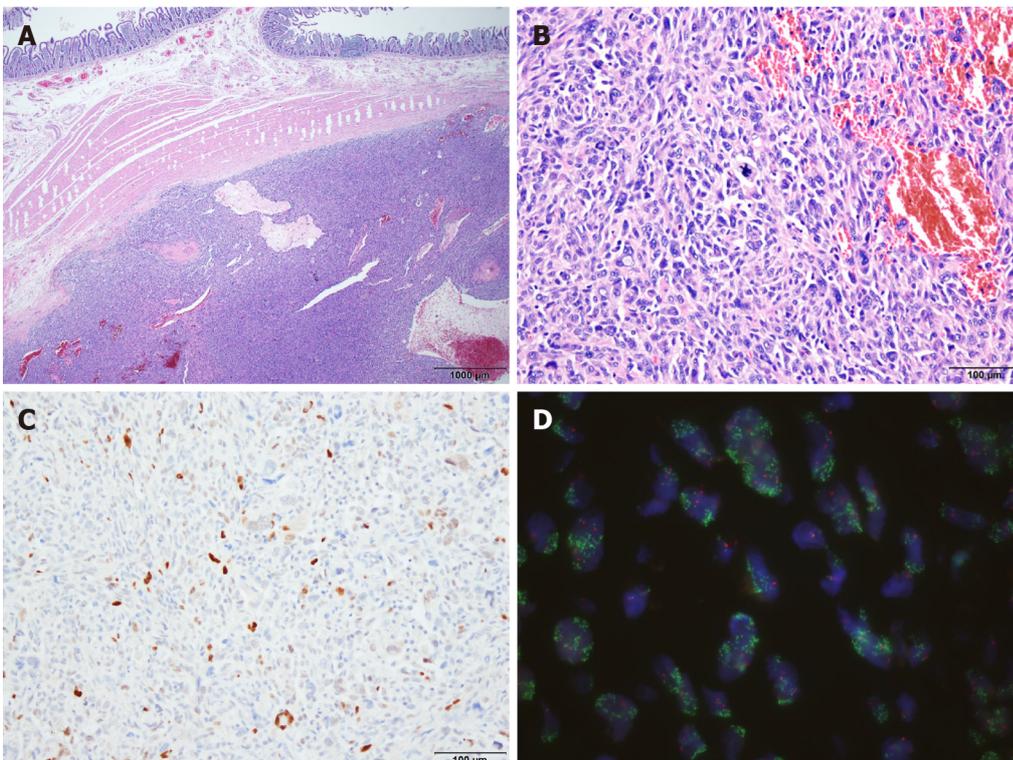
Abdominal computed tomography (CT) revealed a 3 cm-sized heterogeneously enhancing mass in the pancreaticoduodenal groove, causing the obstruction of the second portion of the duodenum.

## FINAL DIAGNOSIS

The preoperative differential diagnosis was duodenal adenocarcinoma, gastro-intestinal stromal tumor (GIST), or leiomyosarcoma (LMS). An accurate diagnosis *via* preoperative upper endoscopy was not possible because only the mucosal surface was collected due to duodenal stenosis. The duodenal tumor was diagnosed as DDLPS by combining all clinical, radiologic, and intraoperative findings and histologic data (Figure 1 and Figure 2).



**Figure 1 Radiologic findings.** A, B: Abdominal computed tomography scan demonstrated a heterogeneously enhanced mass in the pancreaticoduodenal groove with duodenal obstruction.



**Figure 2 Features of tumor cells.** A: The tumor was located in the submucosal layer of the duodenum (Hematoxylin-and-eosin stain,  $\times 20$ ); B: At a higher magnification, undifferentiated tumor cells were shown to have marked nuclear atypia with brisk mitotic activity (Hematoxylin-and-eosin stain,  $\times 200$ ); C: Immunohistochemistry revealed positivity for MDM2 in the tumor cells (Immunohistochemistry,  $\times 200$ ); D: *MDM2* amplification was detected by *MDM2/CEN12* fluorescence *in situ* hybridization assay (*MDM2*-green signals, *CEN12*-red signals,  $\times 1000$ ).

## TREATMENT

The patient underwent pylorus-preserving pancreaticoduodenectomy with *en bloc* right hemicolectomy, superior mesenteric vein segmental resection, and inferior vena cava wedge resection. Intraoperatively, the duodenal mass invaded the pancreas and hepatic flexure of the colon, resulting in gastric and duodenal distension. The tumor also appeared to invade adjacent large blood vessels. Although the duodenal mass was surgically removed, the entire tumor could not be completely removed as the tumor had already spread throughout the body.

Upon macroscopic examination, the surgical resection specimen appeared to have originated from the submucosal layer of the duodenum, measuring 3x3 cm at its greatest dimension. The nodular mass showed a white-to-tan colored cut surface with a focal area of hemorrhage. Pathologic evaluation revealed a tumor arising from the duodenum and extending to the pancreas, colon, and omentum. Histology showed the epicenter of the tumor was the submucosal layer with normal-looking mucosa. The tumor was relatively well circumscribed and composed of high-grade pleomorphic cells. The tumor

was arranged in a haphazard, fascicular growth pattern with telangiectatic-like feature. The majority of the tumor cells exhibited marked nuclear atypia with brisk mitotic activity. Undifferentiated tumor cells displayed large vesicular or hyperchromatic nuclei and prominent macronucleoli. Discohesive polygonal giant cells with abundant eosinophilic cytoplasm were also observed. Immunohistochemistry was positive for MDM2 and CDK4 but was negative for CK, Actin, Desmin, CD117, CD34, ERG, S100, TFE3, Melan A, and HMB45. The tumor was also found to harbor *MDM2* amplification *via* fluorescence *in situ* hybridization (FISH).

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## OUTCOME AND FOLLOW-UP

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Subsequently, chest CT and whole-body positron emission tomography/CT scans were performed to check for preexisting and unidentified intraabdominal liposarcoma outside the gastrointestinal tract and secondary duodenal involvement. No specific abnormalities or metastases were found.

However, due to the vessel invasion of the tumor, cancer seeding contributed to anastomotic dehiscence. Seven days after initial surgery, a 5 mm-sized leakage of the hepaticojejunal anastomosis occurred, leading to generalized peritonitis. The patient underwent a second operation. Large amounts of necrotic fluid had filled the abdominal cavity, and severe adhesions were observed. Hepaticojejunal anastomosis repair with superior mesenteric vein thrombectomy was performed. Fourteen days later, the patient developed sepsis due to the perforation of the gastrojejunal-anastomosis. A life saving emergency operation was performed. Intraoperatively, massive hematoma and bowel adhesion were observed in the upper abdomen. Overall, the tissue was very friable and edematous due to severe inflammation. An external stent was inserted and repaired using a T-tube in the gastrojejunal perforation site. Since the possibility of perforation was very high in the case of primary repair of the jejunal limb, an external stent was inserted using a hemovac drain. After massive irrigation, the operation was terminated.

Unfortunately, the patient did not recover from disseminated intravascular coagulation and peritonitis, and he died 60 d after surgery.

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## DISCUSSION

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Liposarcomas are subclassified as atypical lipomatous tumor/well-differentiated liposarcoma (ALT/WDLPS), DDLPS, myxoid liposarcoma, pleomorphic liposarcoma, or myxoid pleomorphic liposarcoma according to the World Health Organization classification[11]. WDLPS is a typically indolent histologic subtype that presents as slowly growing masses but can be locally aggressive with minimal to no distant metastatic potential, while DDLPS is a higher grade histology with the potential for rapid growth and distant metastatic potential[12,13].

The term DDLPS was first introduced by Evans in 1979[14] and is defined as a combination of ALT/WDLPS and a high-grade non-lipogenic sarcoma-like component of variable histologic grade. DDLPS can occur *de novo* (90%), with about 10% occurring from a pre-existing WDLPS[15]. The histologic hallmark of DDLPS is the transition from ALT/WDLPS to non-lipogenic sarcoma, although a well-differentiated component may not be identifiable.

The tumor usually presents as a large painless mass in late adult life with an equal distribution between males and females. The retroperitoneum is the most common location and occurrence in the extremities and subcutaneous tissue is very rare[16]. Since DDLPS primarily involving the intestine is extremely unusual, less than 10 cases have been reported to date[1-3,5-10,17]. Of those tumors arising in the small bowel, four originated in the jejunum, five in the ileum, and two in the duodenum (Table 1). Clinical information on the precise locations of the other three small bowel tumors is not available. The literature reveals that occurrences of DDLPS in the small intestine can cause various symptoms, such as intussusception, bleeding, obstruction, and abdominal discomfort. Since Okabayashi *et al*[6] reported the first case in 2013, there has only been one additional report of DDLPS in the duodenum[10].

Microscopically, the tumor exhibits variable histologic features but mostly undifferentiated pleomorphic cells with striking nuclear atypia. Such tumors with unusual locations and histopathological features may pose a diagnostic challenge. Thus, sarcomatoid carcinoma, GIST, LMS, malignant melanoma and other high-grade sarcomas should be among the list of differential diagnoses.

Sarcomatoid carcinomas should be considered as the primary differential diagnosis. These tumors are predominantly composed of poorly differentiated spindle cells and/or undifferentiated bizarre anaplastic cells resembling fibrosarcoma or LMS. Sarcomatoid carcinomas can be diagnosed by immunohistochemistry using cytokeratin to demonstrate epithelial derivation.

GIST may occur anywhere in the gastrointestinal tract with 30% arising in the small bowel, including the duodenum. The tumor also exhibits a broad morphologic spectrum. While most instances of GIST are spindle or epithelioid cell tumors, progression to high-grade sarcomatous morphology can be seen rarely. The majority of GISTs show expression of CD117, DOG1 and CD34, which may be helpful for diagnosis.

Table 1 Clinicopathologic features of liposarcoma from small bowel described previous and in present reports

No.	Ref.	Age (yr)	Sex	Clinical presentation	Location	Histology	Treatment
1	Atik <i>et al</i> [2]	58	F	Intussusception	Jejunum	LPS	Not stated
2	Papadopoulos <i>et al</i> [7]	52	M	Abdominal discomfort, vomiting	Jejunum	WDLPS	Small bowel resection
3	Rivkind <i>et al</i> [9]	Unknown	Unknown	Mimicking appendicitis	Small intestine	Myxoid LPS	Unknown
4	Benaragama <i>et al</i> [3]	76	M	Small bowel perforation	Iliac fossa	WDLPS	Segmental resection
5	Patel <i>et al</i> [8]	59	M	Palpable lump	Iliac fossa	DDLPS with WD component	Resection
6	Jeong <i>et al</i> [30]	45	M	Abdominal discomfort	Jejunum	DDLPS with WD component	Excision
7	Okabayashi <i>et al</i> [6]	55	M	Melena	Duodenum	DDLPS with WD component	Pancreaticoduodenectomy with partial colon resection
8	Nennstiel <i>et al</i> [17]	60	M	Gastrointestinal bleeding	Ileocecal valve	Pleomorphic LPS	Ileum segmental resection
9	Matsuo <i>et al</i> [5]	84	M	Intussusception	Ileum	DDLPS	Ileocecal resection
10	Gajzer <i>et al</i> [1]	51	M	Not provided	Small intestine	DDLPS with myofibroblastic differentiation	Excision
11	Gajzer <i>et al</i> [1]	75	M	Intestinal obstruction	Ileum	DDLPS with WD component	Segmental ileectomy
12	Gajzer <i>et al</i> [1]	53	M	Intestinal obstruction	Jejunum	DDLPS	Segmental jejunectomy
13	Gajzer <i>et al</i> [1]	68	M	Not provided	Small intestine	DDLPS with WD component	Excision
14	Whitham <i>et al</i> [10]	59	F	Fatigue, palpitation, shortness of breath	Duodenum	DDLPS	Segmental duodenal resection and distal gastrectomy
15	Present case	64	M	Abdominal pain, weight loss	Duodenum	DDLPS	Pylorus-preserving pancreaticoduodenectomy

F: Female; M: Male; LPS: Liposarcoma; DDLPS: Dedifferentiated liposarcoma; WDLPS: Well differentiated liposarcoma.

Gastrointestinal LMS is also very rare, with fewer than 100 cases reported in the English-language literature since 2000[18-21]. Typical LMS shows spindle cells with blunt-ended nuclei and eosinophilic fibrillary cytoplasm. The tumor cells are arranged in intersecting fascicles with varying degree of nuclear atypia, necrosis, and brisk mitotic activity. LMS can exhibit a poorly differentiated, pleomorphic appearance in addition to typical areas; this is known as pleomorphic LMS or dedifferentiated LMS[22, 23]. For this diagnosis to be established, morphological features characteristic of classic LMS must be present, and are usually positive for at least one myogenic marker, although staining is often weaker and more focal than in typical leiomyosarcomatous areas[22,23]. The diagnosis of LMS should be made on the basis of immunohistochemical stains along with the appropriate morphological features.

Most melanomas of the gastrointestinal tract are metastases from the skin, and primary small bowel melanoma of duodenal origin is extremely rare[24,25]. Because malignant melanomas display various histologic appearance, strong clinical suspicion and precise evaluation are needed to diagnose primary duodenal melanoma.

Based on the morphology of tumor cells in DDLPS, other types of high-grade tumors such as undifferentiated pleomorphic sarcoma, malignant peripheral nerve sheath tumor, and angiosarcoma should be included among the differential diagnoses. The tumor in the present case showed positive immunoreactivity for MDM2 and CDK4. DDLPS was diagnosed based on the histological and immunohistological findings combined with *MDM2* amplification *via* FISH.

Dual staining with MDM2 and CDK4 has been shown to be both sensitive and specific to DDLPS[26]. This is a result of the overexpression of the protein product from chromosomal amplification in the 12q13-15 region of the *MDM2* and *CDK4* oncogenes. Amplification of these genes can then be confirmed with FISH if diagnostic uncertainty remains[27,28].

However, MDM2 positivity by immunohistochemistry is not a specific indicator of *MDM2* amplification because MDM2 positivity is observed in many other sarcomas, including WDLPS, DDLPS, intimal sarcoma, LMS, angiosarcoma, and myxofibro-sarcoma[29]. Validation of *MDM2* amplification by FISH, which is currently a gold standard, is mandatory to confirm the diagnosis of DDLPS.

While the biologic behavior of DDLPS appears to be unfavorable, the most effective treatment modality is surgical resection. There have been no published studies of adjuvant therapy due to the paucity of intestinal DDLPS. Because there are only a limited number of cases, we cannot predict the outcomes for DDLPS arising in the small bowel, but we expect similar prognoses compared with soft tissue DDLPS.

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## CONCLUSION

We presented a unique case of DDLPS originating from the duodenum, one of the rarest locations for gastrointestinal sarcomas. DDLPS should be thoroughly distinguished from its morphological mimickers, as the tumor may be more highly aggressive and extensive than clinically and radiologically expected. While histopathologic features and immunohistochemistry offer evidence of DDLPS, *MDM2* FISH is essential to confirm the diagnosis. Pathologists should keep DDLPS among the initial histologic differential diagnoses for high-grade tumors of the gastrointestinal tract.

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## FOOTNOTES

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