

[CASE REPORT]

Acute Myeloid Leukemia Developing with Acute Pancreatitis Mimicking Autoimmune Pancreatitis

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Abstract:

A 33-year-old man was admitted to our hospital for fever and abdominal pain. A blood analysis revealed pancytopenia and increased serum pancreatic enzymes with disseminated intravascular coagulation. A detailed examination revealed acute pancreatitis, with diffuse swelling of the pancreas and diffuse beaded dilatation of the main pancreatic duct, which mimicked autoimmune pancreatitis complicated by acute myeloid leukemia. Systemic cytotoxic chemotherapy led to the remission of leukemia and pancreatitis. We hypothesized that the etiology of acute pancreatitis was invasion of leukemia cells. Acute pancreatitis is rare as a symptom of leukemia; however, we should consider the possibility of leukemia during the differential diagnosis of acute pancreatitis.

Key words: acute pancreatitis, autoimmune pancreatitis, acute myeloid leukemia

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Introduction

In general, acute leukemia develops with symptoms accompanying pancytopenia, such as anemia, bleeding tendency and febrile neutropenia (1). It is therefore possible to miss leukemia if other organ symptoms are dominant. We report a case of acute myeloid leukemia (AML) accompanied by abdominal pain due to acute pancreatitis.

Case Report

A 33-year-old man with a 5-year history of back pain due to lumbar disc herniation underwent microscopic herniotomy in April 2016. At one week after the surgery, he developed persistent fever and hypochondralgia. Abdominal computed tomography (CT) demonstrated swelling of the pancreatic head and dilatation of the main pancreatic duct. He was transferred to our hospital with acute pancreatitis. On physical examination, he had epigastric tenderness without the

Blumberg sign, left cervical lymphadenopathy, bilateral pretibial petechiae and splenomegaly with a two-finger breadth. Blood tests demonstrated pancytopenia with a hemoglobin level of 11.4 g/dL, a white blood cell count of 3,000/ μ L (blasts 12.5%, mono 28.5%) and a platelet count of 6.3×10^4 / μ L. Hemostatic tests revealed an elongated prothrombin time of 14.7 seconds and an activated partial thromboplastin time of 40.5 seconds, with a D-dimer level of 3.0 μ g/mL. Blood chemistry and immunological analyses revealed the following findings: amylase, 802 IU/L; pancreatic amylase, 762 IU/L; elastase 1, 5,587 IU/L; total bilirubin, 0.4 mg/dL; LDH, 245 IU/L; corrected calcium, 9.1 mg/dL; CRP, 4.8 mg/dL; IgG, 1,528 mg/dL and IgG4, 72.8 mg/dL. The patient was negative for autoantibodies. A bone marrow examination revealed increased monoblasts (38.6%) and eosinophils (8.1%), with 270,000 copies/ μ gRNA of the CBF β /MYH11 fusion gene. On flow cytometry, monoblastic cells were positive for CD 13, 33, 34, and HLA-DR and negative for CD 2, 3, 4, 5, 7, 8, 10, 19, 20, 16, 56, 14, and 41. A chromosomal analysis demonstrated 46,XY,inv(16)(p

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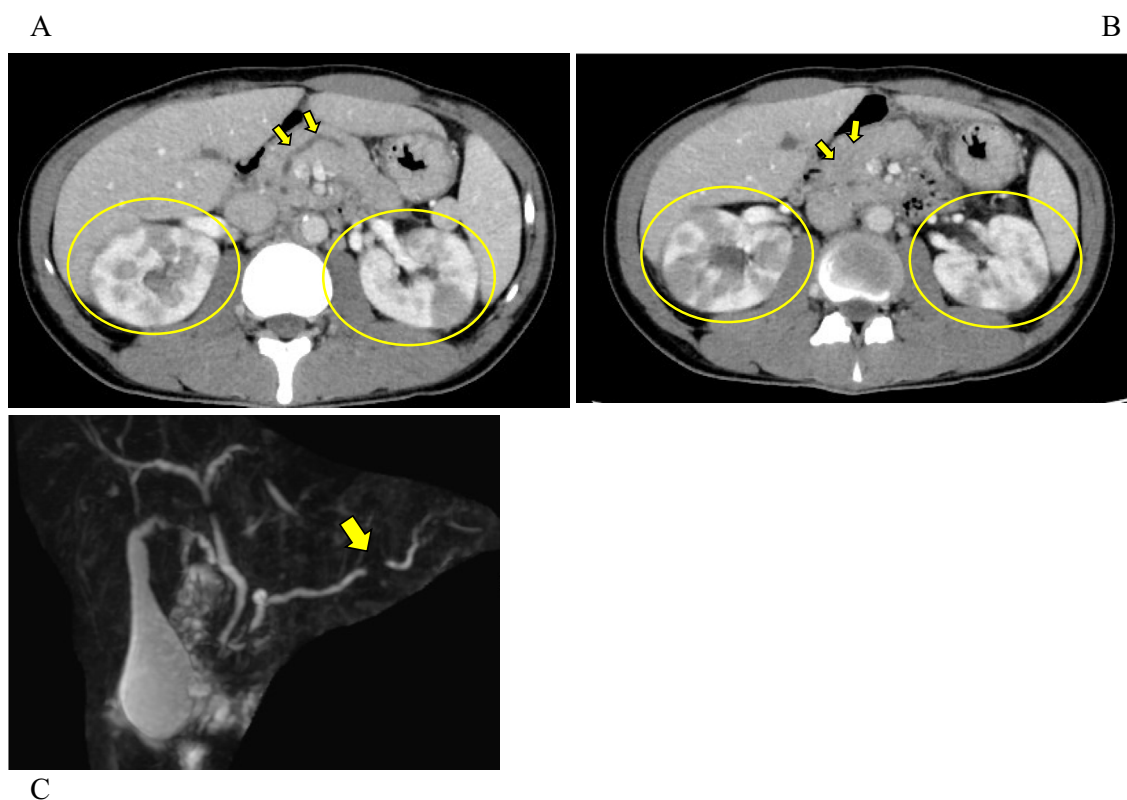


Figure 1. A, B: Computed tomography image of the abdomen on admission. Diffuse swelling of the pancreas, dilatation of the main pancreatic duct (arrows) and multiple enhancement defects in the bilateral kidneys (circles) were observed. C: Magnetic resonance cholangiopancreatography on admission showed beaded dilatation of the pancreatic duct and disruption of the main pancreatic duct (arrow).

13.1q22)[13]/47, idem, +22[5]. *C-kit* mutation was not investigated. Enhanced CT confirmed the dilatation of the main pancreatic duct, swelling of the pancreatic head, multiple cuneiform defects of the bilateral kidney and splenomegaly. Magnetic resonance cholangiopancreatography (MRCP) revealed disruption and dilation of the main pancreatic duct, which had a beaded appearance, without pancreatobiliary maljunction or biliary stones. We diagnosed the patient with AML with inv(16)(p13.1q22), and suspected invasion of leukemia cells to the kidneys and pancreas. Induction chemotherapy with idarubicin and cytarabine achieved hematological remission (Fig. 2). During consolidation therapy, MRCP and CT confirmed the disappearance of dilatation of the main pancreatic duct, swelling of the pancreas and multiple kidney defects. Disruption of the main pancreatic duct remained. No pancreatic pseudocyst developed in the healed pancreas after the treatment of pancreatitis. After eight months and three courses of high-dose cytarabine, the patient developed molecular relapse. Pancreatitis did not develop to the point of molecular relapse before hematological relapse. We performed re-induction therapy followed by unrelated bone marrow transplantation with a conditioning regimen consisting of fludarabine, busulfan and melphalan. The disruption of the main pancreatic duct was not improved at the time of bone marrow transplantation. The patient has been alive for 3 years since the initial development

of AML without relapse of AML or pancreatitis.

Discussion

We reported the case of a patient with AML who developed acute pancreatitis as an initial symptom. We diagnosed the cause of acute pancreatitis as invasion of AML. As fever with abdominal pain is a relatively atypical symptom for AML, systemic chemotherapy followed by bone marrow transplantation resolved the pancreatitis and leukemia without pancreatic sequelae.

Acute pancreatitis involves inflammation of the pancreas, and may be associated with a systemic inflammatory response that can impair the function of other organs or systems (2). Most cases develop with sudden abdominal pain and severe cases are associated with a high rate of mortality (3). The pathophysiology of acute pancreatitis by imaging is divided into interstitial edematous pancreatitis and necrotic pancreatitis. The etiologies of pancreatitis are alcohol-induced, gall bladder stone-induced and post endoscopic retrograde cholangiopancreatography, but drug-induced and mumps virus-induced pancreatitis have also been reported (4). L-asparaginase-associated pancreatitis is well known to occur in association with treatment in leukemia patients (5). Malignant tumors rarely cause acute pancreatitis; however, pancreatic cancer, intraduct papillary mucinous

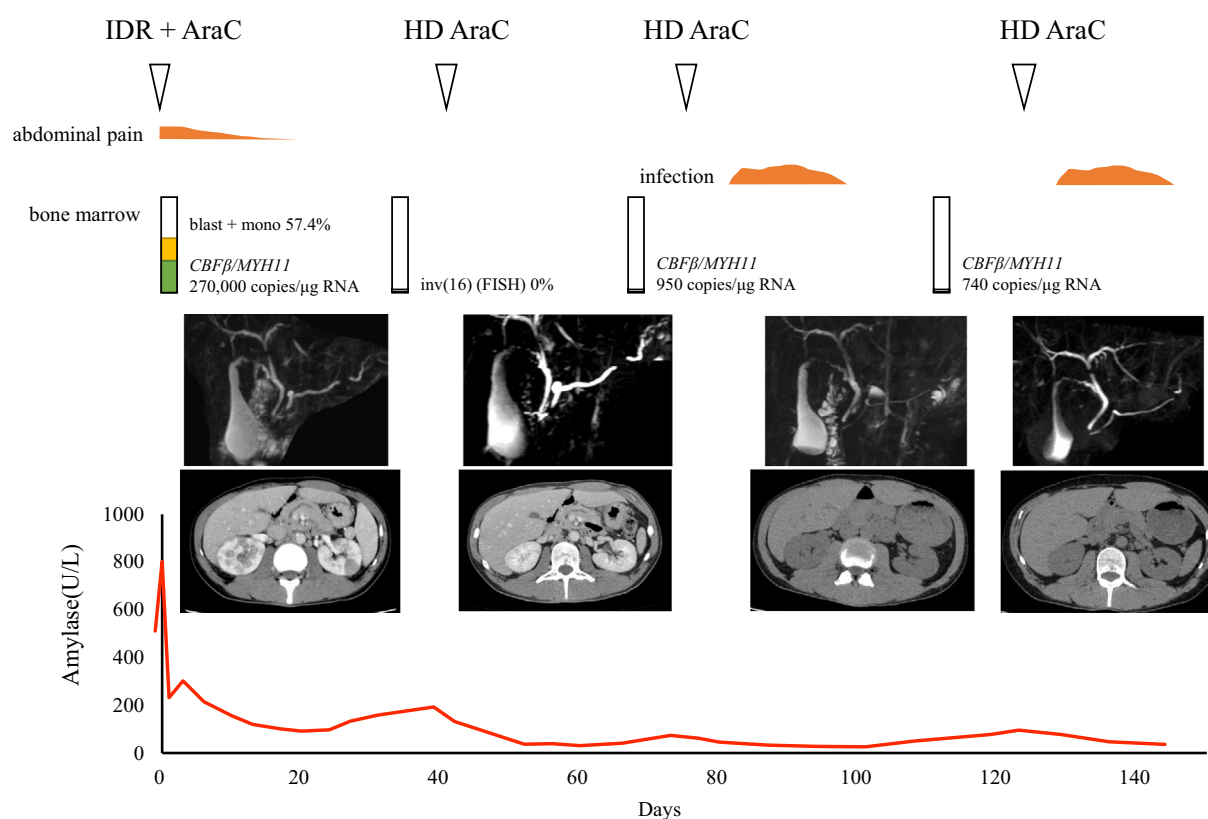


Figure 2. Clinical course. The patient received idarubicin and cytarabine as induction therapy and achieved a hematological remission. Leukemia and pancreatitis both improved after chemotherapy, and the abnormal pancreas and kidney findings disappeared after the first course of consolidation chemotherapy. IDR: idarubicin, AraC: cytarabine, HD AraC: high-dose cytarabine, FISH: fluorescence *in situ* hybridization, mono: monocyte, RNA: ribonucleic acid

neoplasm, metastatic pancreatic cancer, pancreatic neuroendocrine tumor, biliary cancer and papillary cancer of the duodenal papilla were previously reported to induce acute pancreatitis (6). We diagnosed our patient with acute pancreatitis based on abdominal pain, increased serum amylase and the swelling of the pancreas on CT. Furthermore, a bone marrow examination confirmed AML. As disseminated intravascular coagulation developed on admission, we were unable to perform a biopsy because of the high possibility of pancreatic hemorrhage after the procedure. However, the cause of pancreatitis was suggested to be AML invasion because induction chemotherapy improved both pancreatitis and leukemia.

Eleven cases of acute pancreatitis with leukemia, excluding cases of pancreatitis induced by chemotherapy, have been reported (7-16) (Table). The ages ranged from 11 years to 76 years, and six of the patients were male. The types of leukemia were as follows: adult T-cell leukemia/lymphoma (n=5), acute lymphoblastic leukemia (n=3), and acute myeloid leukemia, acute promyelocytic leukemia and chronic lymphocytic leukemia (CLL; n=1 each). Imaging of pancreatitis revealed diffuse swelling in 8 cases, and stenosis or dilatation was noted in the main pancreatic duct. The suspected etiologies of pancreatitis included hypercalcemia, invasion of leukemia cells, and obstruction of the pancreatic

duct by leukemia cells. Autoimmune pancreatitis was confirmed in the case of CLL. Needle biopsy was performed in two cases and revealed invasion of leukemia. Systemic chemotherapy was effective for both leukemia and pancreatitis in two of three cases.

One of the characteristics in our case was imaging findings mimicking autoimmune pancreatitis. In 1995, autoimmune pancreatitis was proposed as a subtype of pancreatitis with an autoimmune etiology for which steroid therapy is effective (17). Its imaging findings consist of diffuse swelling of the pancreas and stenosis of the main pancreatic duct (18, 19). High serum IgG4 was often described as a feature of autoimmune pancreatitis; however, normal serum IgG4 levels were observed in 24% of autoimmune pancreatitis cases (20). Our case was atypical as autoimmune pancreatitis because imaging of the main pancreatic duct revealed dilatation rather than stenosis. Furthermore, speckled/dotted enhancement, a capsule-like rim and duct-penetrating sign, characteristics of autoimmune pancreatitis, were not observed. We concluded that pancreatitis developed due to the invasion of leukemia cells because systemic cytotoxic chemotherapy induced the simultaneous remission of both pancreatitis and leukemia. We hypothesize that anti-tumor immune reactions and/or invasion of leukemia cells indirectly found in the pancreas by imaging modalities were due

Table. Cases of Acute Pancreatitis at the Onset of Leukemia.

Reference	Age and sex	Leukemia subtype	Imaging	Cause of pancreatitis	Treatment
7	38F	ATLL	swelling, patchy low density areas within pancreas	hypercalcemia	chemotherapy
7	39M	ATLL	swelling, macular echo pattern, smoothly dilated PD	hypercalcemia	supportive care
8	44F	ATLL	swelling, ascites, fat stranding	hypercalcemia	supportive care
9	50F	ATLL	swelling, irregular dilation of main PD	infiltration (diagnosed via FNB)	chemotherapy
10	42M	B precursor ALL	fat stranding	infiltration (presumed by other lesions)	chemotherapy
11	76M	CLL/SLL	fullness of the pancreas head, common bile duct stricture	AIP (resected specimen)	surgery
12	34F	ATLL	swelling, fat stranding	infiltration or hypercalcemia	N/A
13	30M	AML relapse after U-BMT	swelling, irregular narrowing of main PD	infiltration (diagnosed via FNB)	supportive care
14	11M	T-ALL	dilation of PD, duodenal wall thickening	PD obstruction	surgery
15	25F	T-ALL	swelling, low density areas	infiltration	supportive care
16	49M	APL	N/A	N/A	supportive care
Present case	33M	AML with inv(16)	swelling (head>body or tail), beaded dilation and obstruction of main PD	infiltration	chemotherapy

PD: pancreatic duct, N/A: not available, FNB: fine needle biopsy, ATLL: adult T-cell leukemia/lymphoma, CLL/SLL: chronic lymphocytic leukemia/small lymphocytic lymphoma, ALL: acute lymphoblastic leukemia, APL: acute promyelocytic leukemia, AML: acute myeloid leukemia, Ara-C: cytarabine, AIP: autoimmune pancreatitis

to an autoimmune pancreatitis-like diffuse swelling pattern, which is atypical of conventional autoimmune pancreatitis. In the relapse phase of AML, autoimmune pancreatitis-like findings on magnetic resonance imaging have been reported and paraneoplastic autoimmune pancreatitis in metastatic breast cancer patients was also reported recently (13, 21, 22). However, acute pancreatitis in leukemia is rare and the mechanisms underlying the development of pancreatitis in our patient remain unclear. More clinical cases are needed.

AML with inv(16) (p13.1q22) is considered a favorable subset of AML; however, our patient relapsed relatively soon after initial therapy and developed extramedullary infiltration of leukemia. Extramedullary leukemia in inv(16)(p13q22) is relatively frequent, developing at a rate of 19% in 27 cases, with the most common lesions being cervical lymphadenopathy with or without tonsillar enlargement; this cervico-tonsillar extramedullary involvement is associated with a shorter duration of first remission (23). In another report, the frequency of extramedullary tumors in leukemia with *RUNX1-RUNX1T1* and *CBFβ-MYH11* was 10%, and they were a negative prognostic factors in leukemia with *RUNX1-RUNX1T1*, but not *CBFβ-MYH11* (24). *C-kit* and *FLT3* mutations were reported to be negative prognostic factors in AML with inv16(p13.1q22) (24-26). We did not examine these mutations, but they may have played a role in the early relapse in our case. A chromosomal analysis at the diagnosis demonstrated trisomy 22, which was reported to

be a favorable factor in AML with inv(16) or t(16;16) (26); however, how this chromosomal abnormality affected the prognosis is unclear. The association between AML complicated by acute pancreatitis and the prognosis remains unknown. The accumulation of further cases is required.

In conclusion, we described a case of AML that initially developed with acute pancreatitis. We considered the etiology of acute pancreatitis to be the invasion of leukemia cells because systemic cytotoxic chemotherapy ameliorated both leukemia and pancreatitis. Acute pancreatitis is rare as a symptom of leukemia; however, we should consider the possibility of leukemia during the differential diagnosis of acute pancreatitis.

The authors state that they have no Conflict of Interest (COI).

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