Imaging autoimmune pancreatitis

Computed tomography and MRI findings of 40 cases of autoimmune pancreatitis (AIP) were reviewed, and AIP classified into three types: diffuse, segmental and multifocal. A capsule-like rim can play an important role in diagnosing AIP. A smooth contour of the lesion without circumference fat tissue change is also considered to be a characteristic of AIP. The important differential diagnosis of diffuse type AIP is lymphoma. Differentiating clues are main pancreatic duct change and enhancement pattern on dynamic study. Differentiation between segmental type AIP and pancreatic cancer is often difficult or may be almost impossible. Although rare, AIP may show multifocal delayed enhanced masses within the pancreas. Recognition of this type of AIP is mandatory in diagnosing pancreatic abnormalities.

Autoimmune pancreatitis (AIP) is characterized by diffuse or segmental enlargement of the pancreas with irregular narrowing of the main pancreatic duct, and fibrotic change with lymphocyte infiltration histopathologically. Diagnosis of AIP is very important, since it can be treated by steroid therapy.

Previously, diffuse pancreatic enlargement with diminished signal intensity on T1-weighted MR images (T1WI), delayed parenchymal enhancement on dynamic imaging and a capsule-like rim, which shows as low density on computed tomography (CT) and hypointense on T1-weighted MR images, have been considered to be characteristic of AIP [1]. However, as recognition of the disease concept of AIP has spread and many cases have been reported, it is now recognized that AIP displays various CT and MRI findings [2-5].

Difficulties in differentiating AIP-related atypical imaging findings, such as focal enlargement of the pancreas, from pancreatic carcinoma due to focal mass formation have been reported [6,7]. It is recognized that differentiating AIP from other pancreatic disorders may often be difficult.

We reviewed CT and MRI findings of 40 cases of AIP, and classified AIP into three types: diffuse, segmental and multifocal. We then attempted to differentiate AIP and other pancreatic disorders on CT and MRI according to those types of AIP.

**Review of CT & MRI findings of AIP**

**Patient population**

The patient population consisted of 40 patients (29 men, 11 women; age: 19–81 years, mean: 58) who fulfilled the diagnostic criteria for AIP according to the Japan Pancreas Society 2006 (Box 1) [8], and had received no treatment before.

**Image acquisition**

Dynamic CT was performed in all 40 patients (single-detector helical CT or four- or 64-detector row helical CT). MRI (Signa 1.5 T, Magnetom Vision or Symphony 1.5 T) was available in 37 patients (all include dynamic study).

All patients received 100 ml of intravenous nonionic contrast material containing 370 mg I/ml by means of an automated power injector. Slice thickness was 3–5 mm. The early phase was obtained 35–45 s after the initiation of contrast material injection at an injection rate of 3.0 cc/s. The portal venous and delayed phases were acquired at 70 and 240 s, respectively, after the initiation of contrast material injection.

Unenhanced MR images included axial T2-weighted turbo spin echo, axial T1-weighted gradient-echo images with and without fat saturation, and coronal magnetic resonance cholangiopancreatography. Contrast-enhanced, 3D dynamic MRI studies using gadopentetate dimeglumine were performed at an injection rate of 2.0 cm3/s, including early, portal and delayed phases. The early phase was obtained 30 s after the start of the injection, and the portal and delayed phases were obtained 60 and 240 s after initiation of the intravenous contrast medium injection, respectively. Postcontrast axial or coronal T1-weighted gradient-echo images with fat saturation were also obtained.
Box 1. Clinical diagnostic criteria.

- Diffuse or segmental narrowing of the main pancreatic duct with irregular wall, and diffuse or localized enlargement of the pancreas by imaging studies, such as abdominal ultrasonography, computed tomography and MRI.
- High serum γ-globulin, IgG or IgG4, or the presence of autoantibodies, such as antinuclear antibodies and rheumatoid factor.
- Marked interlobular fibrosis and prominent infiltration of lymphocytes and plasma cells in the periductal area, occasionally with lymphoid follicles in the pancreas.
- Diagnosis of autoimmune pancreatitis is established when criterion 1, together with 2 and/or 3, are fulfilled. However, it is necessary to exclude malignant disease such as pancreatic or biliary cancers.

Data taken from [8].

Box 2. Summary of computed tomography and MRI findings.

- Diffuse enlargement: 29
  - Abnormal parenchyma (delayed enhancement on dynamic study and hypointensity on T₁WI): 24
    - With capsule-like rim: 16 (Figure 1)
    - Without capsule-like rim: 8 (Figure 2)
  - Normal parenchyma (early enhancement on dynamic study and hyperintensity on T₁WI): 3
    (Figure 3)
  - Multifocal nodular lesions showing delayed enhancement on dynamic study and hypointensity on T₁WI: 2 (Figure 4)
- Segmental enlargement: 11
  - Segmental delayed enhancement of the same segmental area on dynamic study and hypointensity on T₁WI
    - With capsule-like rim: 3 (Figure 5)
    - Without capsule-like rim: 8 (Figure 6)
- Presence of delayed enhancement area on dynamic study and its extent
  - Present: 37
    - Diffuse: 24
    - Segmental: 11 (head: 2, head–body: 2, body–tail: 7)
    - Multifocal and nodular: 2
  - Absent: 3
- Presence of hypointense area on T₁WI and its extent (n = 37)
  - Present: 35
    - Diffuse: 22
    - Segmental: 11 (head: 2, head–body: 2, body–tail: 7)
    - Multifocal and nodular: 2
  - Absent: 2

The results of the hypointense area on T₁WI were the same as the results of the delayed enhancement area on dynamic study:

- Presence of capsule-like rim
  - Present: 19
    - Diffuse enlargement case: 16
    - Segmental enlargement case: 3
  - Absent: 21
    - Diffuse enlargement case: 13
    - Segmental enlargement case: 8

A summary of the CT and MRI findings are shown in Box 2 (Supplementary Figures 1–6) (see online www.futuremedicine.com/toc/iim/1/1):
Contour of the lesion and change of circumference fatty tissue
- Smooth contour without circumference fatty tissue abnormality: 37 (Supplementary figure 6)
- Irregular contour with obliteration of circumference fatty tissue: 3 (Supplementary figure 7)
- A smooth contour of the lesion without circumference fat tissue change is considered to be characteristic for AIP

Others (e.g., calcification and cysts)
- Calcification: 4 (Supplementary figure 8)
- Cysts: 3 (Supplementary figure 9)
- Main pancreatic duct (MPD) dilatation in the pancreatic body–tail: 4 (Supplementary figure 7)

It should be noted that these findings are not so rare and can sometimes be observed in AIP.

Three types of AIP based on CT & MRI findings
According to the results, we classified AIP into three types: diffuse, segmental and multifocal.

Diffuse type
Diffusely enlarged pancreas demonstrates delayed parenchymal enhancement on dynamic study and hypointensity on T1WI, with (subtype: 1) or without (subtype: 2) a capsule-like rim. Rarely, the pancreatic parenchyma may show normal enhancement pattern on dynamic study and normal hyperintensity on T1WI (subtype: 3).

The subtypes are as follows:
- Diffuse with capsule-like rim: 40% (Supplementary figure 1)
- Diffuse without capsule-like rim: 20% (Supplementary figure 2)
- Diffuse with normal parenchyma: 7.5% (Supplementary figure 3)

Segmental type
Segmental type AIP shows segmental pancreatic enlargement with delayed enhancement on dynamic study and hypointensity on T1WI of the same segmental area, with (subtype: 1) or without (subtype: 2) a capsule-like rim. Correct diagnosis of segmental type AIP may often be difficult in imaging findings alone.

The subtypes are as follows:
- Segmental with capsule-like rim: 7.5% (Supplementary figure 5)
- Segmental without capsule-like rim: 20% (Supplementary figure 6)

Multifocal type
Multifocal masses, which demonstrate delayed enhancement on dynamic study and hypointensity on T1WI, are found within the diffusely enlarged pancreas. Multifocal type AIP (Supplementary figure 4) is rare (5%), but there have been several case reports of this type [9,10]. We believe this to be a distinct type of AIP and should be recognized as such by radiologists.

Differentiation between AIP & other pancreatic disorders on CT & MRI according to the type of AIP
Diffuse type
**Diffuse with capsule-like rim**
This type of AIP is most common and should be diagnosed correctly. In acute pancreatitis, phlegmon surrounding the pancreas may resemble a capsule-like rim; however, it does not show enhancement on dynamic study [11] and differentiation is not difficult (Supplementary figure 10). There is no other differential diagnosis.

**Diffuse without capsule-like rim**
This type of AIP is somewhat difficult to diagnose compared with diffuse type with capsule-like rim. Smooth contour of the pancreas without circumference fatty tissue abnormality as well as narrowing or nonvisualization of MPD on magnetic resonance cholangiopancreatography are the key findings for obtaining a correct diagnosis (Supplementary figure 2). Differential diagnoses include mild acute pancreatitis, malignant lymphoma, leukemia, diffuse pancreatic cancer and diffuse pancreatic metastasis.

Mild acute pancreatitis often shows diffuse enlargement of the pancreas, however, early enhancement on dynamic study is usually preserved and it can be differentiated from this type of AIP (Supplementary figure 11).

There are several differential observations between malignant lymphoma (Supplementary figure 12) and diffuse type without capsule-like rim. The main pancreatic duct is usually well preserved in lymphoma [12]. The enhancement pattern on dynamic study of lymphoma may differ from AIP (e.g., delayed washout), since lymphoma is a hypercellular tumor and has a small amount of fibrous stroma. It is well known that malignant lymphoma shows hyperintensity on diffusion-weighted MRI with high b-factor due to the hypercellularity and a high nuclear-to-cytoplasmic ratio of the tumor [13]. Leukemia has a similar appearance to lymphoma [14], and is included in the differential diagnosis (Supplementary figure 13).
Diffuse pancreatic cancer should invade the peripancreatic region and major vessels, and differentiation from AIP is thought to be easy. However, diffuse pancreatic cancer is very rare, and we have not experienced this type of cancer.

Diffuse pancreatic metastasis is thought to be end-stage metastatic disease, usually accompanied with other organic metastases (Supplementary figure 14) [15,16].

**Diffuse with normal parenchyma**

This type of AIP may reflect early stages of AIP [4]. Since the size of a normal pancreas varies, diagnosis of pancreatic enlargement is usually made subjectively and is therefore difficult. Thus, diagnosis of this type of AIP is very difficult. Differential diagnosis is mild acute pancreatitis and differentiation between them requires clinical information such as serum amylase levels (Supplementary figure 11).

**Segmental type**

**Segmental with capsule-like rim**

A capsule-like rim may be a specific finding of AIP; therefore, diagnosis of this type AIP is not so difficult (Supplementary figure 5). Detailed evaluation to detect a capsule-like rim is essential for correct diagnosis. Differential diagnosis is segmental acute pancreatitis with phlegmon around, but differentiation is usually easy since phlegmon does not show enhancement on dynamic study.

**Segmental without capsule-like rim**

The most important and difficult differential diagnosis is pancreatic cancer. Contour of the affected part on AIP is usually smooth and AIP does not obliterate peripancreatic fat tissue in contrast to pancreatic cancer, a finding that should be considered when differentiating between them (Supplementary figure 6) [17]. Focal enlargement of the pancreatic head without upstream MPD dilatation may suggest AIP rather than pancreatic cancer.

In the case of upstream MPD dilatation, this type of AIP is very difficult to diagnose correctly (Supplementary figure 7). In this situation, differentiation between this type of AIP and pancreatic cancer, based on CT and MRI findings alone, may be almost impossible, and usually requires clinical information such as IgG4 levels [18]. A diagnostic trial of steroid therapy may be used for evaluation; however, it should be done very carefully by experts, since steroid therapy may elevate the risk of progression to pancreatic cancer [19].

**Multifocal type**

Although rare, AIP may show multifocal delayed-enhanced masses within the pancreas [9,10]. Recognition of this type of AIP is mandatory in diagnosing pancreatic abnormalities.

The most important differential diagnosis is malignant lymphoma, the same as diffuse type AIP without capsule-like rim. Different enhancement patterns and diffusion-weighted MRI may be useful in differentiating lymphoma from AIP (Supplementary figure 15).

Multiple pancreatic cancer can also be a differential diagnosis, but it is extremely rare. Multiple pancreatic metastases may be another differential diagnosis. History of hypovascular primary cancer is a key point in diagnosing multiple pancreatic metastases, and it often accompanies other organic metastases (Supplementary figure 16) [14].

**Conclusion**

AIP can be classified into three types: diffuse, segmental and multifocal. Differentiation between AIP and other pancreatic disorders on CT and MRI may be easier, considering those types of AIP.

Diffuse type AIP should be relatively easy to diagnose; however, an important differential diagnosis is lymphoma. Differentiating segmental type AIP and pancreatic cancer is difficult, especially in AIP without capsule-like rim and with upstream MPD dilation.

The presence of a third type of AIP, multifocal AIP, should be recognized, and important differential diagnoses include lymphoma and metastases.

**Future perspective**

In patients with AIP, serum IgG4 levels are frequently and significantly elevated, and various extrapancreatic lesions are present. Dense infiltration of IgG4-positive plasma cells as well as fibrosis have been found in the peripancreatic retroperitoneal tissue, bile duct, gallbladder wall, periportal area of the liver, salivary glands and the pancreas. Furthermore, all of the extrapancreatic lesions associated with AIP, such as sclerosing cholangitis, sclerosing sialadenitis and retroperitoneal fibrosis, show infiltration of abundant IgG4-positive plasma cells. Both the pancreatic and extrapancreatic lesions of AIP respond well to steroid therapy. Thus, as a novel clinicopathological entity, IgG4-related sclerosing disease has been proposed. Although the precise pathogenesis and clinicopathologic features have not been well recognized, they will be clarified in the near future.
Financial & competing interests disclosure
The authors have no relevant affiliations or financial involvement with any organization or entity with a financial interest in or financial conflict with the subject matter or materials discussed in the manuscript. This includes employment, consultancies, honoraria, stock ownership or options, expert testimony, grants or patents received or pending, or royalties. No writing assistance was utilized in the production of this manuscript.

Executive summary

Review of computed tomography & MRI findings of autoimmune pancreatitis

- Diffuse enlargement: 29
  - Abnormal parenchyma: 24
    - With capsule-like rim: 16
    - Without capsule-like rim: 8
  - Normal parenchyma: 3
  - Multifocal nodular lesions: 2
- Segmental enlargement: 11
  - With capsule-like rim: 3
  - Without capsule-like rim: 8

A smooth contour of the lesion without circumference fat tissue change is considered to be a characteristic of autoimmune pancreatitis (AIP). It should be recognized that calcification, cysts and main pancreatic duct dilatation in the pancreatic body/tail are not so rare and can sometimes be observed in AIP.

Three types of AIP based on computed tomography & MRI findings

- Diffuse: 67.5%
  - With capsule-like rim: 40%
  - Without capsule-like rim: 20%
  - With normal parenchyma: 7.5%
- Segmental: 27.5%
  - With capsule-like rim: 7.5%
  - Without capsule-like rim: 20%
- Multifocal: 5%

Differentiation between AIP & other pancreatic disorders on computed tomography & MRI according to the type of AIP

- Differential diagnoses of diffuse type without capsule-like rim include mild acute pancreatitis, malignant lymphoma, leukemia, diffuse pancreatic cancer and diffuse pancreatic metastasis. The main pancreatic duct is usually well preserved in lymphoma and the enhancement pattern on dynamic study of the lymphoma may differ from AIP.
- The most important and difficult differential diagnosis of segmental type without capsule-like rim is pancreatic cancer. Contour of the affected portion on AIP is usually smooth and AIP does not obliterate peripancreatic fat tissue in contrast to pancreatic cancer, which may be a useful observation when differentiating between them. Focal enlargement of the pancreatic head without upstream main pancreatic duct dilatation may suggest AIP rather than pancreatic cancer.

Bibliography

Papers of special note have been highlighted as:

5. Detailed imaging evaluation using various imaging modalities was performed.
7. Reports on cases of autoimmune pancreatitis (AIP) in Japan and proposed new diagnostic criteria.
9. Reports difficulties of differentiating segmental type AIP from mass-forming pancreatitis and pancreatic cancer.


* This is the first case report of multifocal type AIP.


