

World Journal of *Clinical Cases*

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WJCC mainly publishes articles reporting research results and findings obtained in the field of clinical medicine and covering a wide range of topics, including case control studies, retrospective cohort studies, retrospective studies, clinical trials studies, observational studies, prospective studies, randomized controlled trials, randomized clinical trials, systematic reviews, meta-analysis, and case reports.

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Spontaneous remission of autoimmune pancreatitis: Four case reports

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Abstract

BACKGROUND

Autoimmune pancreatitis (AIP) is a particular type of chronic pancreatitis, and steroid treatment of AIP is effective. Spontaneous remission (SR) of AIP without steroids is relatively rare. The international consensus for the treatment of autoimmune pancreatitis suggests that patients with AIP with obstructive jaundice, abdominal pain, and back pain related to the pancreas or the bile duct should be treated with steroids; most asymptomatic patients with AIP may improve without steroids. However, in our clinical work, we found that the clinical characteristics of AIP patients with SR vary. Four of these cases are described here. In addition, to our knowledge, there is no previously published report of dynamic imaging before and after SR of AIP at present.

CASE SUMMARY

We present the cases of four patients with AIP (two females and two males) in which the AIP improved spontaneously without steroid treatment. Two patients were asymptomatic, one patient had abdominal pain with obstructive jaundice, and one patient had intermittent right upper abdominal pain. Three patients presented with localized pancreatic enlargement and one with diffuse pancreatic enlargement. In addition to the pancreatic lesions, bile duct involvement was seen in two patients, and no extra-pancreatic organ involvement was found in the other two patients. The serum IgG4 level of all patients was more than twice the normal level. After SR in the four patients, the affected pancreases exhibited three types of image features: Return to normal, progressive fibrosis, and atrophy and calcification.

CONCLUSION

The clinical features of SR in our four patients with AIP differ, but the imaging findings share some characteristics. After SR, in some cases the affected pancreas

could return to normal, although some patients suffer from progressive fibrosis and atrophy as well as calcification.

Key Words: Autoimmune pancreatitis; Spontaneous remission; Computed tomography; Magnetic resonance imaging; Case report

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Core Tip: We focus on spontaneous remission (SR) in patients with autoimmune pancreatitis due to the high risks and contraindications of steroid treatment. As with asymptomatic patients, some symptomatic patients may also have SR. Patients with localized or diffuse enlargement of the pancreas (the former may be more prone to SR), with or without involvement of the bile ducts, may have SR. After SR, the affected pancreases may return to normal, or exhibit progressive fibrosis and atrophy as well as calcification.

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INTRODUCTION

Autoimmune pancreatitis (AIP) is a type of chronic pancreatitis with a hypothesized autoimmune mechanism. It is characterized by: (1) Dense infiltration of plasma cells and lymphocytes; (2) Peculiar storiform fibrosis; and (3) Occlusive phlebitis[1]. The concept of AIP was proposed by Yoshida *et al*[2] in 1995. Subsequent studies showed that AIP is a disease with special clinical, radiological, and serological characteristics. It is a manifestation of pancreatic involvement in IgG4-related systemic diseases that often involve the bile ducts, salivary and lacrimal glands, kidneys, and lymph nodes[3]. The international consensus for the treatment of autoimmune pancreatitis suggests that AIP patients with obstructive jaundice, abdominal pain, and back pain related to the pancreas or the bile duct should be treated with steroids. "Watchful waiting" may be appropriate in most of the asymptomatic patients, because they may improve spontaneously without steroid treatment[4]. However, in our clinical experience, we found that the clinical characteristics of AIP patients with spontaneous remission (SR) are different, and the factors that lead to SR are not clear. In addition, as far as we know, there is no published report of dynamic imaging before and after SR of AIP up to now. Therefore, we report here the clinical and imaging findings of four AIP patients with SR in order to promote more appropriate individualized treatment strategies for patients with AIP.

CASE PRESENTATION

Chief complaints

Case 1: A 67-year-old woman had a pancreatic lesion found accidentally by yearly ultrasound for 2 mo.

Case 2: A 55-year-old woman complained of having a pancreatic lesion detected accidentally by computed tomography (CT) for 23 d.

Case 3: A 76-year-old man suffered from right upper abdominal pain for 1 mo, and his skin and sclera were yellow for more than 10 d.

Case 4: A 78-year-old man suffered from intermittent right upper abdominal pain for 6 d without obvious cause.

History of present illness

Case 1: Two months preceding our examination, an abnormal pancreatic lesion was found by routine ultrasound examination in this patient. She had no discomfort. For further diagnosis, plain and enhanced MR scanning of the abdomen was performed in the outpatient department of our hospital. The diagnosis was focal AIP of the pancreatic body and tail, which required follow-up closely, to rule out a malignant tumor. In order to further clarify the diagnosis and treatment, she was admitted to our hospital.

Case 2: This patient underwent physical examination 23 d before entering our care, and abdominal plain CT showed a mass at the body and tail of the pancreas. She went to a local hospital and underwent an abdominal enhanced CT scan. The result suggested that pancreatic cancer could not be excluded. Subsequently, she visited our hospital for surgical treatment. She had no discomfort, and her weight had not changed.

Case 3: One month prior to our examination, this patient complained of right upper abdominal pain of unknown cause, accompanied by abdominal distension. He went to a primary hospital and was diagnosed with gastroenteritis, and was treated with traditional Chinese medicine. However, his symptoms did not significantly improve. Ten days ago, he had yellow skin and sclera, and dark-colored urine. He went to a local hospital where abdominal ultrasound showed that his common bile duct was dilated and his pancreatic head was enlarged. The blood test indicated that his serum total bilirubin level (TBIL) was 11.59 mg/dL, direct bilirubin level (DBIL) was 9.23 mg/dL, alanine aminotransferase level (ALT) was 164.5 U/L, and aspartate aminotransferase level (AST) was 106.7 U/L. Then he visited our hospital for further treatment. Endoscopic cholangiopancreatography (ERCP) was performed after preoperative examination, and a severe common bile duct stenosis was seen. Later, he underwent duodenal papillotomy (EST), endoscopic nasobiliary drainage (ENBD), and endoscopic pancreatic stent drainage (ERPD). After the operation, he was sent to our department for further evaluation and treatment.

Case 4: Six days prior to our examination, this patient had right upper abdominal pain without obvious cause, accompanied by intermittent vomiting. He went to a local hospital where abdominal ultrasound showed mild dilatation of the common bile duct and a solid mass in the pancreatic head. The blood test indicated that his TBIL was 9.64 mg/dL, DBIL was 3.55 mg/dL, ALT was 84 U/L, and AST was 70 U/L. Subsequently, he came to our department for further diagnosis and treatment.

History of past illness

Case 1: This patient had a history of allergic asthma for 6 years; she was treated with intermittent Salmeterol Xinafoate and Fluticasone Propionate aerosol. She had a 1 year history of diabetes, and used 6-9 U subcutaneous insulin and 0.5 g oral metformin three times a day. Her fasting blood glucose (FBS) was controlled at 6.20-7.20 mmol/L.

Case 2: This patient's blood sugar level increased during the course of the last year; her FBS was controlled by diet at 7.40-8.90 mmol/L.

Case 3: This patient had a history of hypertension for more than 10 years; he regularly took 10 mg nifedipine sustained-release tablets twice a day, and his blood pressure was controlled at 130-140/70 mmHg. He had a history of a prior cerebral infarction more than 10 years ago without sequelae.

Case 4: This patient had a history of hypertension for more than 7 years; he took 40 mg nimodipine three times a day, and his blood pressure was controlled at 120-130/90 mmHg. He also had abnormal liver function in the last 2 years as well as chronic gastritis in the past year without treatment.

Personal and family history

Cases 1 and 2: These patients had no history of smoking or alcohol abuse.

Case 3: This patient smoked 20 cigarettes and drank about 20 g of alcohol per day for more than 50 years.

Case 4: This patient drank about 10 g of alcohol per day for more than 40 years. All patients had no unusual family history of disease.

Physical examination

Cases 1 and 2: These patients had no symptoms and no abnormalities were identified during the physical examination. All vital signs were within normal ranges.

Case 3: This patient had yellow skin and sclera, tenderness in the right upper abdomen, and no rebound pain and muscle tension, and no mass could be touched. The drainage tube of the bile duct was properly fixed on the skin and discharged dark green liquid. Vital signs were normal.

Case 4: This patient had right upper abdomen tenderness. Vital signs were normal.

Laboratory examinations

Case 1: Laboratory data were as follows: Red blood cell (RBC) count, 3.69×10^{12} /L; hemoglobin (HGB), 115 g/L; white blood cell (WBC) count, 6.52×10^9 /L; C-reactive protein (CRP), 23.0 mg/L; glycosylated hemoglobin A1c (HbA1c), 6.50%; blood amylase, 33 IU/L. The indices of liver function were normal. Serum IgG4 was elevated to 530 mg/dL; antinuclear antibody and antineutrophil antibody were

negative. Carbohydrate antigen 19-9 (CA 19-9), carcinoembryonic antigen (CEA), and alpha-fetoprotein (AFP) were normal.

Case 2: Laboratory data were as follows: Routine blood analysis, normal; HbA1c, 8.40%; blood amylase, 26 U/L. The indices of liver function were normal. Serum IgG4 was elevated to 407 mg/dL. CA 19-9, CEA, and AFP were normal.

Case 3: Laboratory data were as follows: RBC count, $3.23 \times 10^{12}/L$; HGB, 121 g/L; WBC count, $5.51 \times 10^9/L$; CRP, 68 mg/L; blood amylase, 62 IU/L; TBIL, 9.41 mg/dL; DBIL, 4.94 mg/dL; ALT, 84 U/L; AST, 58.5 U/L; alkaline phosphatase (ALP), 356 U/L; γ -glutamyltranspeptidase (γ -GT), 168 U/L. Serum IgG4 was 1310 mg/dL. CA 19-9 was 88.00 U/mL; CEA and AFP were normal.

Case 4: Laboratory data were as follows: RBC count, $2.59 \times 10^{12}/L$; HGB, 100 g/L; WBC count, $3.70 \times 10^9/L$; TBIL, 2.85 mg/dL; DBIL, 1.75 mg/dL; ALP, 185 U/L; ALT, 43 U/L; AST, 41 U/L. Serum IgG4 was 590 mg/dL. CA 19-9, CEA, and AFP were normal.

Imaging examinations

Case 1: Abdominal magnetic resonance imaging (MRI) showed obvious enlargement of the pancreatic body and tail. Compared with the pancreatic head, the lesion showed low signal intensity on T1WI, and high signal intensity on T2WI. Multiple enlarged lymph nodes were seen around the portal vein (Figure 1A and B). The lesion demonstrated decreased enhancement in the arterial phase, and progressive enhancement in the delayed phase on gadolinium-enhanced T1WI. A pseudocapsule was around the lesion and no expansion of the pancreatic duct was observed (Figure 1C and D). ECHO-3-22 needles were used to puncture the lesion under monitoring by endoscopic ultrasound. The pathology demonstrated well differentiated squamous epithelial and glandular epithelial cells, and a small number of lymphocytes and neutrophils, with no definite malignant tumor cells.

Case 2: Abdominal MRI showed obvious enlargement of the pancreatic body and tail. Compared with the pancreatic head, the lesion showed low signal intensity on T1WI, and slightly high signal intensity on T2WI (Figure 2A and B). There was gradual enhancement of the lesion, splenic vein stenosis posterior to the lesion, and dilation of the large curved side vein of the gastric body on gadolinium-enhanced T1WI. There was no dilation of the pancreatic duct (Figure 2C-E). Three ECHO-3-22 puncture needles were monitored by endoscopic ultrasound. Pathology examination demonstrated that there were a large number of lymphocytes, a small number of neutrophils, a small number of well differentiated epithelial cells, and proliferation of fibrous tissue in the enlarged pancreas. No malignant tumor cells were found.

Case 3: Abdominal MRI showed diffuse enlargement of the pancreas. Compared with adjacent liver parenchyma, the pancreas showed low signal intensity on T1WI, and high signal intensity on T2WI (Figure 3A and B). On gadolinium-enhanced T1WI, a sign of progressive enhancement of the pancreatic lesion and the pseudocapsule around the pancreas was observed (Figure 3C and D); furthermore, the hepatic hilar and extra-hepatic bile duct wall was diffusely thickened and abnormally enhanced (Figure 3E).

Case 4: Abdominal contrast enhanced CT showed that the pancreatic head was slightly enlarged with delayed enhancement (Figure 4A-C). There was no abnormality of the pancreatic body and tail. The bile duct wall in the pancreatic head was thickened and enhanced abnormally (Figure 4B). Magnetic resonance cholangiopancreatography showed that the main pancreatic duct was narrowed in a longer segment, and the upstream pancreatic duct was slightly dilated, with a diameter of 4 mm. Moreover, the common bile duct of the pancreatic segment was severely narrowed, and the proximal intra- and extra-hepatic bile ducts were dilated (Figure 4D). Three disposable puncture needles with COOK22G were introduced at the pancreatic head lesion with ultrasonic monitoring. Degenerative small round cells were seen, but no malignant tumor cells were found.

FINAL DIAGNOSIS

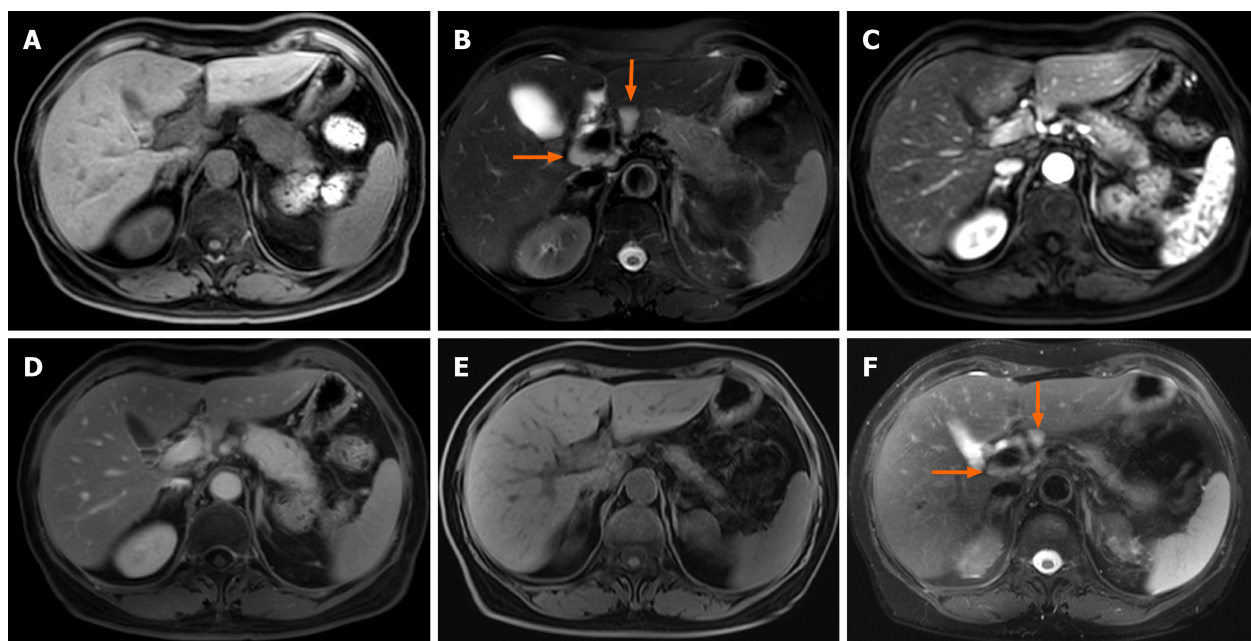
Cases 1 and 2: Focal AIP of the pancreatic body and tail.

Case 3: Diffuse AIP.

Case 4: Focal AIP of the pancreatic head.

TREATMENT

Cases 1 and 2: These patients refused steroid treatment, and were followed regularly in our outpatient



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Figure 1 Focal autoimmune pancreatitis with atrophy after spontaneous remission in case 1. A and B: The pancreatic body and tail enlarge, and multiple enlarged lymph nodes (arrows) are seen around the portal vein; C and D: Gradual enhancement of the lesion on enhanced scan; E and F: Magnetic resonance imaging follow-up at the 13th month. Pancreatic swelling dissipates and the volume of the pancreas decreases significantly, and the enlarged lymph nodes (arrows) are significantly reduced as well.

department.

Case 3: After the operation, this patient was treated by fasting and water prohibition, acid inhibition, and nutritional support, but he refused steroid treatment. Ten days after hospital admission, ERCP and EST and endoscopic biliary stent drainage were performed. After discharge, he took 50 mg ursodeoxycholic acid orally three times a day.

Case 4: After admission, this patient refused to use steroid and was given symptomatic treatment such as liquid food, acid suppression, and liver protection. After discharge, he received ursodeoxycholic acid (50 mg) and pancreatin enteric-coated capsules (50 mg) three times a day.

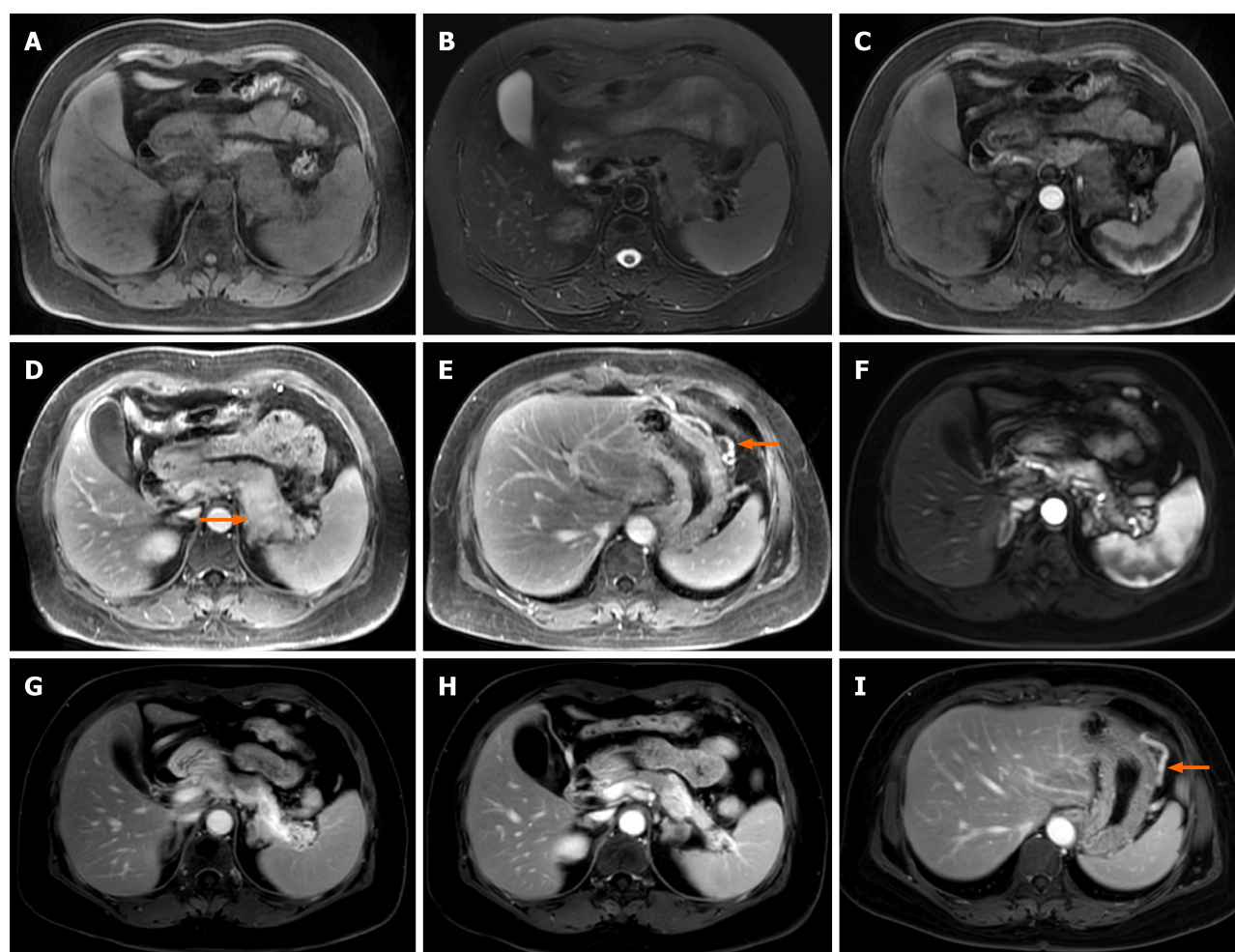
OUTCOME AND FOLLOW-UP

Case 1: During 15 mo following treatment, this patient had no symptoms. The HbA1c ranged from 7.20% to 8.30%. Her serum IgG4 had decreased to 162 mg/dL by the 11th month. Pancreatic swelling dissipated and the volume of the pancreas decreased significantly, and the enlarged lymph nodes were reduced obviously as observed on MRI at the 13-mo follow-up (Figure 1E and F).

Case 2: During 22 mo following treatment, this patient had no symptoms. The patient injected 6-12 U insulin subcutaneously once every night; FBS was in the range of 6.28-7.12 mmol/L. MRI follow-up at 22 mo showed that the body and tail of the pancreas were significantly reduced and more obviously gradually enhanced, which suggested that the affected pancreatic parenchyma suffered from progressive fibrosis. The stenosis of the splenic vein and the dilation of the large curvature vein of the gastric body were not improved (Figure 2F-I).

Case 3: This patient's clinical symptoms improved and his bilirubin levels decreased gradually. Twenty months after treatment, abdominal CT showed that the volume of the pancreas had decreased and the thickening of the bile duct wall had disappeared (Figure 3F).

Case 4: This patient's symptoms gradually improved. At the 12-mo follow-up, the patient's serum IgG4 was 570 mg/dL, the pancreatic head swelling had dissipated and it appeared to contain multiple punctate calcifications (Figure 4E). There was no expansion of the bile and pancreatic ducts (Figure 4F).



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Figure 2 Focal autoimmune pancreatitis with progressive fibrosis after spontaneous remission in case 2. A and B: Enlargement of the pancreatic body and tail; C, D, and E: In addition to the gradual enhancement of the lesion in the body and tail of the pancreas, stenosis of the splenic vein (arrow) and dilation of the large curved side vein of the gastric body (arrow) are observed; F and G: Magnetic resonance imaging (MRI) follow-up at 22 mo showing that the body and tail of the pancreas is significantly reduced and more obviously gradually enhanced; H: Lower level MRI showing that the volume of the pancreatic body and tail is significantly atrophied and obviously enhanced; I: The dilation of the large curvature side vein of the gastric body (arrow) is not improved.

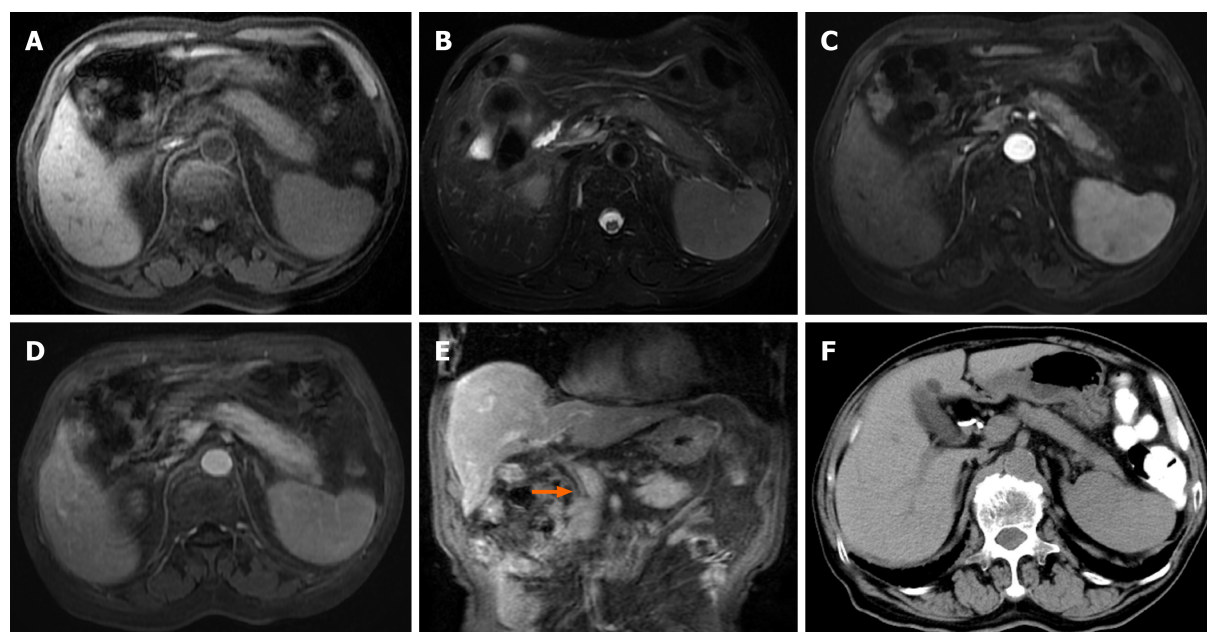
DISCUSSION

AIP remission without steroid treatment has been reported occasionally, but its mechanism is not clear. In order to deepen our understanding of the SR of AIP, we searched the database through MEDLINE using keywords "autoimmune pancreatitis" or "IgG4-related disease" and "spontaneous remission" or "spontaneous resolution" or "spontaneous release" or "natural course". After excluding cases with incomplete clinical data or cases with no evidence of remission of pancreatic lesions, nine publications were ultimately examined, including four original research papers[5-8] and five case reports[9-13] (Table 1). Kamisawa *et al*[5] reported 11 AIP patients who were followed during the natural course of the disease. Ultimately, six patients needed steroid treatment due to disease progression, while five patients were stable or had SR of the disease. They found that there were no significant differences in the clinical characteristics between the two groups. The other three studies on SR of AIP were reported by Kubota *et al*[6-8] in 2007, 2009, and 2011. Multivariate analysis of factors related to the SR of AIP showed no consistent pattern. The independent predictors of SR were negative staining for IgG4 in the duodenal papilla, early manifestations of endoscopic ultrasonography (lobularity and hyperechoic pancreatic duct margin), and IgG4 seronegativity. In five case reports of AIP patients with SR, the clinical symptoms were different in each one. In these patients, the pancreas could exhibit either diffuse swelling or contain a localized mass, and serum IgG4 could be slightly or significantly increased, with or without the involvement of extra-pancreatic organs. Our four cases and the literature reports suggest that the factors responsible for the SR of AIP are not clear, and that the clinical characteristics of the AIP patients with SR are quite different. Of the four patients that we describe, two had no symptoms, one had intermittent abdominal pain, and one had abdominal pain with obstructive jaundice. Among them, three had localized enlargement, and one had diffuse enlargement of the pancreas, accompanied with

Table 1 Literature review of autoimmune pancreatitis cases with spontaneous remission

Ref.	Age/sex	Clinical presentation	Serum IgG4 mg/dL	Type of pancreatic enlargement	Associated diseases	Treatment	Outcome
Ohtsubo <i>et al</i> [9], 2021	67/F	Dark urine	674 (< 135)	Pancreatic head mass	Nothing	Biliary plastic stent	Pancreatic head mass diminished after 1 month; multiple malignant lymphomas of the bile duct after 3 mo
Kusano <i>et al</i> [10], 2019	79/M	Dry mouth, weight loss, blood glucose control deteriorated	1830 (4.8-105)	Diffuse pancreatic enlargement	Lymphadenopathy	Intensive insulin treatment	Pancreatic enlargement disappeared but lymph nodes persisted after 4 years; IgG4 was 623
Uchida <i>et al</i> [11], 2007	54/M	Jaundice	213 (6-140)	Pancreatic head mass	Hepatic masses	Endoscopic biliary drainage Biopsy of hepatic mass	Pancreatic masses disappeared after 3 mo; hepatic masses disappeared after 12 mo
Araki <i>et al</i> [12], 2006	73/F	No symptoms	196 (4.8-105)	Pancreatic uncinata and tail mass	Nothing	Follow-up	Uncinate mass decreased but mass in the tail increased after 3.6.9 mo; IgG4 were 163-203
Ozden <i>et al</i> [13], 2005	58/F	Painless jaundice	Not mentioned	Pancreatic head body and tail mass	Gallbladder	Percutaneous cholangiography and external drainage; Laparotomy and Cholecystectomy	No evidence of focal lesion in pancreas after 6 mo; IgG4 was 1040

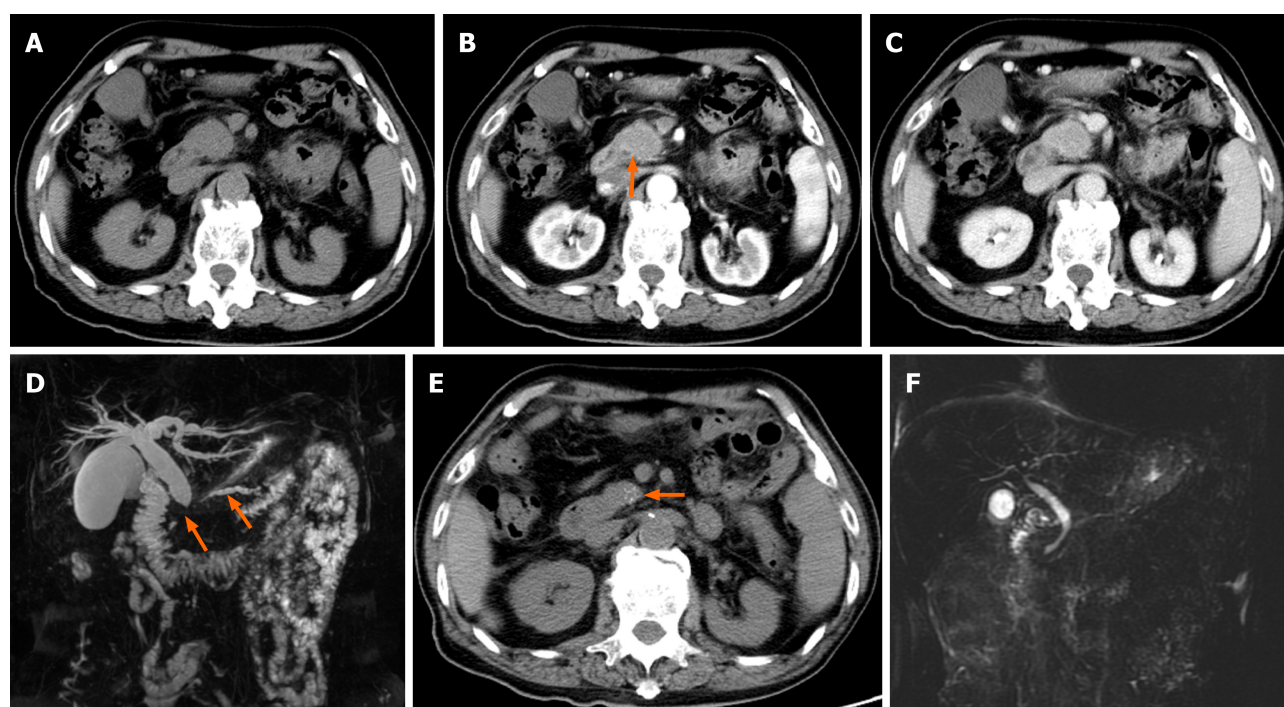
F: Female; M: Male.



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Figure 3 Diffuse autoimmune pancreatitis with spontaneous remission in case 3. A and B: Diffuse enlargement of the pancreas; C and D: Progressive enhancement of the pancreatic lesion and the pseudocapsule on enhanced scan; E: Extra-hepatic bile duct wall thickening and enhancement (arrow); F: Computed tomography follow-up at 20 mo showing that the pancreatic swelling has disappeared.

bile duct involvement in two, and no extra-pancreatic organ involvement in two cases. In all the presented cases, the level of serum IgG4 was more than twice the normal level. Our four cases suggest that there are three types of image features after SR of AIP: (1) The size and density (signal) of the pancreas return to normal; (2) There may be progressive fibrosis of the pancreas; and (3) There may be atrophy and calcification of the pancreas. Previous studies had shown that the atrophy and calcification of pancreatic parenchyma accompanying AIP were often related to fibrosis and recrudescence, and may be accompanied by irreversible internal and external pancreatic secretion dysfunction[14,15]. During



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Figure 4 Focal autoimmune pancreatitis with calcification after spontaneous remission in case 4. A, B, and C: The pancreatic head is enlarged with progressive enhancement, the wall of the common bile duct is thickened and abnormally enhanced (arrow); D: Magnetic resonance cholangiopancreatography (MRCP) showed a longer stenotic segment of the pancreatic duct (arrow) and bile duct, and dilatation of the upstream pancreatic duct (arrow) and proximal bile duct; E: Computed tomography follow-up after 12 mo showing that the swelling of the pancreatic head has dissipated, and multiple punctate calcifications (arrow) occur; F: The diseased pancreatic duct and bile ducts have nearly returned to normal on MRCP follow-up.

long-term follow-up, case 2 showed progressive fibrosis, and case 1 and case 4 showed atrophy and calcification of the affected pancreas without steroid treatment, respectively. Such imaging findings after SR of AIP have not previously been reported. With atrophy and progressive fibrosis, pancreatic endocrine dysfunction in case 1 and case 2 was aggravated and blood glucose was not well controlled. At the same time, case 2 had left portal hypertension secondary to splenic vein involvement. An MRI scan in this case indicated that the lumen of the splenic vein was narrowed and the large curved side vein of the gastric body was dilated. Juarez *et al*[16] reported that when AIP involved the splenic vein, blood flow was limited, resulting in left portal hypertension, which may lead to congestive splenomegaly and variceal bleeding. Therefore, these authors suggested treatment with steroids early to reduce the risk of irreversible complications. In fact, although we recommended case 2 to use steroids as soon as possible to prevent the progression of the disease, she refused. Therefore, we continued follow-up observations to determine her long-term prognosis. Of the four patients with SR of AIP, serum IgG4 was rechecked in two patients. Among these, serum IgG4 returned to normal levels in case 1, and it decreased slightly in case 4. We speculate that the change in serum IgG4 Level is not consistent with or synchronous with the imaging appearances of pancreatic lesion remissions, which is consistent with the reports of Kusano[10] and Araki[12]. It should be noted that the SR of AIP in this study is just the result of patients' rejection of steroids. Actually, a steroid therapy had been suggested for the four patients at the beginning when the final AIP diagnosis was established.

CONCLUSION

In conclusion, the clinical features of the SR of AIP in our four cases are different, but the imaging findings share some characteristics. For example, all cases had localized or diffuse enlargement of the pancreas (the former is more common), with or without involvement of the bile ducts. Follow-up CT and MRI showed that after the SR of AIP, the affected pancreas could return to normal, although some patients suffered from progressive fibrosis and atrophy as well as calcification.

FOOTNOTES

Author contributions: Zhang BB and Jin EH conceived the idea of the study; Zhang BB and Huo JW collected and analysed the data; Yang ZH and Wang ZC interpreted the results; Zhang BB wrote the paper; all authors discussed the results and revised the manuscript.

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