Risk Factors for Recurrent Pancreatitis After First Episode of Acute Pancreatitis

Yingying Sun, Jie Jin, Aying Zhu, Hong Hu, Yingying Lu, Yue Zeng, Dadao Jing

1Department of Gastroenterology, Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine, Shanghai, 201600, People’s Republic of China; 2Department of Geriatrics, Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine, Shanghai, 200080, People’s Republic of China

Correspondence: Dadao Jing, Department of Gastroenterology, Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine, NO. 650 Xinsongjiang Road, Shanghai, 201600, People’s Republic of China. Tel +86-13816958050, Email dadaojing@126.com

Background: In recent years, the cure rate of acute pancreatitis (AP) has increased gradually, but the morbidity of recurrent acute pancreatitis (RAP) has not decreased. Patients with RAP have a poor quality of life and are more likely to develop into chronic pancreatitis. To investigate the risk factors of RAP after first pancreatitis attack is very necessary.

Methods: Patients with first episode AP admitted to Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine from January 2018 to May 2021 were included in this retrospective study, and follow-up was 3–45 months after discharge. The patients information were collected from medical records including laboratory tests and auxiliary inspection of their hospitalization. Univariate and multivariate Cox regression analysis were used to explore the risk factors of RAP. Cumulative risk of RAP was plotted using Kaplan–Meier curves.

Results: A total of 592 patients were enrolled in the study and 81 (13.7%) of the patients developed RAP. Among those RAP patients, the majority (67.9%) were men, with a median age of 43 years. The most common etiology of RAP was hypertriglyceridemia (38.3%). Multivariate Cox Analysis showed that smoking history (p < 0.001), infected pancreatic necrosis occurred during first admission (p = 0.005), and high low-density lipoprotein cholesterol (LDLc) level (p < 0.001) were significant independent risk factors for RAP. Patients with the above independent risk factors had increased 3-year cumulative risk of recurrence (32.2%, 45.5%, 28.9%, respectively).

Conclusion: Smoking history, infected pancreatic necrosis, and high LDLc level were the most decisive risk factors for RAP. Attention should be paid to the patients with the above factors.

Keywords: first episode pancreatitis, recurrent acute pancreatitis, risk factors, low-density lipoprotein cholesterol, infected pancreatic necrosis

Introduction

Acute pancreatitis (AP) is one of the most common critical diseases of the gastrointestinal tract. It is estimated that between 5 and 35 people per 100,000 are treated for AP annually, and this number has increased over the past decade.1 Biliary, hypertriglyceridemia and alcohol are the common causes of AP. Etiological composition varies in different countries and regions.2,3 Most AP can be completely recovered after standard treatment, but a few patients may have recurrent episodes. The recurrence rate of AP reported in previous studies is around 10–30%.2,4–6 It is important to note that patients with recurrent acute pancreatitis (RAP) are at high risk of progressing to chronic pancreatitis and even may be related to pancreatic cancer.7 And even in the absence of chronic pancreatitis, patients with RAP have lower physical and mental quality of life.8

It is necessary to find the risk factors of pancreatitis recurrence accurately and early. Previous studies have shown that factors that can lead to AP are more likely to become causes of RAP.9 Continued drinking after the first attack, delayed cholecystectomy, and poor control of triglyceride level are all well-known risk factors for RAP.9,10 The correlation between smoking and recurrence of AP is also clear, and it has been agreed that smokers have a significantly increased
The purpose of this study was to find the risk factors for recurrence of AP by analyzing the laboratory test indicators and clinical characteristics of patients with the first episode AP.

**Materials and Methods**

**Patients and Study Design**

Patients admitted to Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine with a first episode of AP between January 2018 and May 2021 were included in our study. The exclusion criteria included patients younger than 18 years of age, pregnancy, prior malignancy, chronic pancreatitis, and poor compliance. The inclusion and exclusion criteria of this study are shown in Figure 1. Patient characteristics, past medical history, tobacco and alcohol consumption, local complication, etiology and severity of AP, laboratory data and modified computed tomography severity index (MCTSI) were collected from medical records of their first hospitalization. All patients received standardized management according to guidelines and consensus. Follow-up of each patient began on the day of discharge and ended with a readmission for AP or the earliest of the following dates: death, loss of contact, or time to end date of our study: August 2021. The follow-up evaluation was based on medical records and telephone visit. This study

**Figure 1** Flowchart of the patients included in the study.

**Abbreviations:** AP, acute pancreatitis; OAP, once episode of acute pancreatitis; RAP, recurrent acute pancreatitis.
was reviewed and approved by the Medical Ethical Committee of Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine (No. 021KY049).

Definition
The diagnosis and classification of AP were based on the 2012 revised Atlanta classification criteria. The diagnosis of AP required two of the following three criteria: 1) typical abdominal pain consistent with AP; 2) serum amylase (or lipase) activity at least three times greater than the standard upper limit; and 3) characteristic findings consistent with AP on computed tomography or abdominal ultrasonography. RAP was defined as follows: 1) interval between attacks greater than 3 months; 2) first episode AP entirely or near wholly resolved; and 3) no conclusive evidence of underlying chronic pancreatitis.

Acute biliary pancreatitis (ABP) was defined as AP caused by biliary stones or biliary tract obstruction. Hypertriglyceridemic pancreatitis (HTGP) was described as a serum triglyceride level at admission greater than 11.3 mmol/L or greater than 5.65 mmol/L without other causes, such as biliary stones or obstruction, and alcohol excluded. Alcohol-induced AP was diagnosed as the consumption of more than 80 g of alcohol per day for at least 10 years or heavy alcohol consumption prior to onset of AP. The remaining cases, such as autoimmune pancreatitis, idiopathic pancreatitis, pancreas divisum, intraductal papillary mucinous neoplasm were defined as others etiology of AP.

Laboratory Detection
Whole blood and serum samples were collected at the first time of admission. Whole blood samples were used to detect C-reactive protein and hemoglobin (BC-1050, Mindray, Guangzhou, China). Serum samples were used to detect the concentrations of blood calcium, fasting blood glucose, glycated hemoglobin A1c, albumin, total cholesterol, triglyceride, high-density lipoprotein cholesterol, low-density lipoprotein cholesterol (LDLc) (ADIVA2400, Siemens, Munich, Germany) and D-dimer (CN6000, Sysmex, Kobe, Japan). Their concentrations were obtained by standard operating methods.

All laboratory indicators were classified according to the normal reference range of our hospital. D-dimer level was classified as high or normal based on whether they were above the upper limit of normal (0.55mg/L). Glycated hemoglobin A1c was divided into high and normal groups based on whether it was greater than 6%. Triglyceride level higher than 1.7mmol/L was classified as high-level. LDLc level was divided into the high-level group and the normal or low-level group according to whether they were greater than 4.14mmol/L. Serum calcium was divided into low (<2.10mmol/L) and normal or high (≥2.10mmol/L) groups. Fasting blood glucose was divided into high (>6.11mmol/L) and normal or low (≤6.11mmol/L) groups. Albumin was divided into low (<40g/L) and normal or high (≥40g/L) groups. Hemoglobin was divided into low (<131g/L) and normal or high (≥131g/L) groups. Total cholesterol was divided into high (>5.18mmol/L) and normal (≤5.18mmol/L) groups. High-density lipoprotein cholesterol was divided into low (<1.16mmol/L) and normal or high (≥1.16mmol/L) groups. C-reactive protein was divided into high and normal groups based on whether it was greater than 10mg/L.

Statistical Analysis
All statistical tests were conducted using SPSS version 26.0 and GraphPad Prism version 8.0. We expressed continuous variables as the mean ± standard deviation (Ma ± SD) or as the median ± interquartile range (Md ± IQR). Normally distributed data were analyzed by a 2-tailed t-test, whereas non-normally distributed data tested by the Mann–Whitney U-test. Categorical variables were indicated as proportions and analyzed using the χ² test. If more than 20% of the expected value was less than 5, Fisher’s exact test was used. Variables with a $p < 0.05$ on univariate analysis were entered into a multivariable Cox regression to identify independent predictors. Results were presented as hazard ratios (HRs) with 95% confidence intervals (CIs). Cumulative incidences of RAP were plotted using Kaplan–Meier curves. Statistical significance was defined as $p < 0.05$.

Results
Patient Characteristics
A total of 644 patients with first episode AP were admitted to the Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine from January 2018 to May 2021. Data were excluded from the exclusion criteria (Age...
<18 years, pregnancy, chronic pancreatitis, pancreatic cancer, other malignant tumor and poor cooperate), and a total of 52 people were excluded. Eventually, 592 patients were enrolled in this study. According to whether patients have recurrence, we divided them into RAP group (n = 81) and once episode of AP (OAP) group (n = 511). Baseline characteristics are presented in Table 1. The median age was 49±26 years, median body mass index (BMI) was 25.40 ±4.60 kg/m², and 383 (64.7%) were males. Seventy-two (12.2%) had a history of diabetes mellitus and 236 (39.9%) had a previous diagnosis of fatty liver. The etiology of AP was biliary in 289 patients (48.8%), hypertriglyceridemia (HTG) in 157 patients (26.5%), alcohol in 68 patients (11.5%), and others in 78 patients (13.2%). In total, 133 (22.5%) people were exposed to cigarette smoke and 150 (25.3%) people have a history of drinking. Of all local complications, acute peripancreatic fluid collection was the most common occurring in 257 (43.4%) patients during the first hospitalization.

Comparison of RAP and OAP
The patients were followed up for 3–45 months with a median follow-up time of 25 months. And, 13.7% (81/592) of patients with first episode AP developed RAP. The median age of the RAP patients was 43 ± 29.5 years, which was younger than that of OAP patients (50 ± 26 years, p = 0.123). The proportion of males was higher in the RAP group (55/81, 67.9%) than in the OAP group (328/511, 64.2%), but the difference was not statistically significant (p = 0.516). The median BMI for two groups were 25.8kg/m² and 25.4 kg/m², respectively (p = 0.459). There were significant differences in tobacco smoke exposure (p < 0.001) and alcohol consumption (p < 0.001) between the two groups. There was no significant difference in previous diagnosis of diabetes mellitus (p = 0.674) between the two groups while the RAP group had a higher rate of fatty liver history (p = 0.004).

The composition of etiology of AP was significantly different (p = 0.010). The most common etiology in the OAP group was biliary (51.5%), whereas the etiology was HTG (38.3%) in the RAP group. Of the 12 patients (14.8%) with others etiology in RAP group, all patients had completed endoscopic ultrasonography during the follow-up. Six of them were congenital malformation of the pancreas (pancreas divisum), four were idiopathic pancreatitis, and the rest of them were autoimmune pancreatitis and intraductal papillary mucinous neoplasm.

The incidence of acute peripancreatic fluid collection, pancreatic pseudocyst, and infected pancreatic necrosis in RAP group were significantly higher than that in OAP group during the first hospitalization (p = 0.000, p = 0.021, p = 0.010, respectively). There were significant differences in serum calcium, fasting blood glucose, and LDLc level between the two groups (p = 0.040, p = 0.015, p = 0.000, respectively). Differences in clinical feature profiles between RAP and OAP groups are presented in Table 1.

Risk of RAP
Regression analysis of risk factors for RAP is presented in Table 2. After univariate Cox regression analysis, the cigarette smoke exposure, alcohol consumption, fatty liver history, etiology and severity of AP, MCTSI, acute peripancreatic fluid collection, pancreatic pseudocyst, infected pancreatic necrosis, high fasting blood glucose level, high triglyceride level, and high LDLc level were significantly associated with RAP (p < 0.05). There is no correlation between the respective variables. Multivariate Cox regression analysis of these variables showed that cigarette smoke exposure (p < 0.001), infected pancreatic necrosis (p = 0.005) and high LDLc level (p < 0.001) were significantly independent risk factors for RAP.

Cumulative risk analysis of development of RAP stratified by cigarette smoke exposure, infected pancreatic necrosis and LDLc level is presented in Figure 2. The 3-year cumulative risk for RAP was highest in patients with infected pancreatic necrosis during the first episode pancreatitis (45.5%). Smokers or patients with high LDLc level have significantly higher cumulative risk for RAP over 3 years than non-smokers or patients with normal LDLc level (32.2% vs 9.5%, 28.9% vs 5%, respectively).

Discussion
Recurrent episodes of pancreatitis seriously affect the quality of life. In addition, some studies consider that AP, RAP, and chronic pancreatitis are the continuum of the same disease. It has even been suggested that RAP may be an
The occurrence of RAP was found to be 13.7% during follow-up, which was consistent with the conclusion of other studies. The etiology composition of AP varies between countries and regions. It has been reported that HTG has become the second most common cause after biliary in China. In our study, HTG and alcohol etiology caused more

Table 1 Basic Characteristics of Patients with First Episode Acute Pancreatitis

|                      | All Patients (n = 592) | OAP (n = 511) | RAP (n = 81) | p       |
|----------------------|------------------------|--------------|--------------|---------|
| **Sex**              |                        |              |              |         |
| Male                 | 383 (64.7%)            | 328 (64.2%)  | 55 (67.9%)   | 0.516a  |
| Female               | 209 (35.3%)            | 183 (35.8%)  | 26 (32.1%)   |         |
| **Age (years)**      | 49.0 ± 26.0            | 50.0 ± 26.0  | 43.0 ± 29.5  | 0.123b  |
| **BMI (kg/m^2)**     | 25.40 ± 4.60           | 25.40 ± 4.47 | 25.80 ± 4.95 | 0.459b  |
| **Habits**           |                        |              |              |         |
| Smoking              | 133 (22.5%)            | 93 (18.2%)   | 40 (49.4%)   | <0.001a |
| Drinking             | 150 (25.3%)            | 116 (22.7%)  | 34 (42.0%)   | <0.001a |
| **Diabetes mellitus**|                        |              |              |         |
|                      | 72 (12.2%)             | 61 (11.9%)   | 11 (13.6%)   | 0.674a  |
| **Fatty liver**      | 236 (39.9%)            | 192 (37.6%)  | 44 (54.3%)   | 0.004a  |
| **Etiology of AP**   |                        |              |              |         |
| Biliary              | 289 (48.8%)            | 263 (51.5%)  | 26 (32.1%)   | 0.010a  |
| HTG                  | 157 (26.5%)            | 126 (24.7%)  | 31 (38.3%)   |         |
| Alcohol              | 68 (11.5%)             | 56 (11.0%)   | 12 (14.8%)   |         |
| Others               | 78 (13.2%)             | 66 (12.9%)   | 12 (14.8%)   |         |
| **Local complication**|                       |              |              |         |
| Acute peripancreatic fluid collection | 257 (43.4%) | 206 (40.3%) | 51 (63.0%) | <0.001a |
| Acute necrotic collection | 58 (9.8%)  | 48 (9.4%)   | 10 (12.3%)   | 0.406a  |
| Pancreatic pseudocyst | 27 (4.6%)             | 19 (3.7%)    | 8 (9.9%)     | 0.021a  |
| Walled-off necrosis  | 14 (2.4%)              | 10 (2.0%)    | 4 (4.9%)     | 0.111a  |
| Infected pancreatic necrosis | 11 (1.9%) | 6 (1.2%)    | 5 (6.2%)     | 0.010a  |
| **MCTSI**            |                        |              |              |         |
| Mild                 | 212 (35.8%)            | 194 (38.0%)  | 18 (22.2%)   | 0.023a  |
| Moderate             | 328 (55.4%)            | 274 (53.6%)  | 54 (66.7%)   |         |
| Severe               | 52 (8.8%)              | 43 (8.4%)    | 9 (11.1%)    |         |
| **Severity of AP**   |                        |              |              |         |
| MAP                  | 368 (62.2%)            | 325 (63.6%)  | 43 (53.1%)   | 0.107a  |
| MSAP                 | 151 (25.5%)            | 128 (25.0%)  | 23 (28.4%)   |         |
| SAP                  | 73 (12.3%)             | 58 (11.4%)   | 15 (18.5%)   |         |
| **Laboratory test**  |                        |              |              |         |
| High D-dimer         | 466 (78.7%)            | 403 (78.9%)  | 63 (77.8%)   | 0.824a  |
| Low-Ca²⁺             | 231 (39.0%)            | 191 (37.4%)  | 40 (49.4%)   | 0.040a  |
| High-FBG             | 343 (57.9%)            | 286 (56.0%)  | 57 (70.4%)   | 0.015a  |
| High-HbA1c           | 188 (31.8%)            | 154 (30.1%)  | 34 (42.0%)   | 0.033a  |
| Low-ALB              | 314 (53.0%)            | 267 (52.3%)  | 47 (58.0%)   | 0.333a  |
| Low-Hb               | 184 (31.1%)            | 159 (31.1%)  | 25 (30.9%)   | 0.964a  |
| High-TC              | 229 (38.7%)            | 192 (37.6%)  | 37 (45.7%)   | 0.164a  |
| High-TG              | 234 (39.5%)            | 190 (37.2%)  | 44 (54.3%)   | 0.003a  |
| Low-HDLc             | 434 (73.3%)            | 370 (72.4%)  | 64 (79.0%)   | 0.212a  |
| High-LDLc            | 64 (10.8%)             | 32 (6.3%)    | 32 (39.5%)   | <0.001a |
| High-CRP             | 477 (80.6%)            | 409 (80.0%)  | 68 (84.0%)   | 0.408a  |

Notes: *χ² test, categorical variables were indicated as proportions. Mann-Whitney U-test, continuous variables as the median with the interquartile range. Abbreviations: OAP, once episode of acute pancreatitis; RAP, recurrent acute pancreatitis; BMI, body mass index; HTG, hypertriglyceridemia; MCTSI, modified computed tomography severity index; MAP, mild acute pancreatitis; MSAP, moderately severe acute pancreatitis; SAP, severe acute pancreatitis; Ca²⁺, serum calcium; FBG, fasting blood glucose; HbA1c, glycated hemoglobin A1c; ALB, albumin; Hb, hemoglobin; TC, total cholesterol; TG, triglyceride; HDLc, high-density lipoprotein cholesterol; LDLc, low-density lipoprotein cholesterol; CRP, C-reactive protein.

independent risk factor for pancreatic cancer. The occurrence of RAP was found to be 13.7% during follow-up, which was consistent with the conclusion of other studies. The etiology composition of AP varies between countries and regions. It has been reported that HTG has become the second most common cause after biliary in China. In our study, HTG and alcohol etiology caused more
AP recurrence than biliary, and HTG is the most frequent etiology factor of recurrence. This result may be related to poor lipid control and continued alcohol consumption after discharge. Tai et al also found higher recurrence rate of HTGP than ABP. HTGP is often associated with multiple factors (metabolic abnormalities, lack of exercise, alcohol

| Table 2 Cox Regression Analyses of the Risk Factors Associated with Recurrent Acute Pancreatitis |
|-----------------------------------------------|------------------|------------------|
|                                              | Univariate Analysis | Multivariate Analysis |
|                                              | HR (95% CI)        | HR (95% CI)        |
| Sex                                           |                   |                   |
| Male                                          | 0.883 (0.554–1.408) | 3.512 (2.010–6.139) |
| Female                                        |                   |                   |
| Age (years)                                   | 0.990 (0.977–1.004) | 1.326 (0.728–2.412) |
| BMI (kg/m²)                                   | 1.014 (0.958–1.074) | 0.631             |
| Habits                                        |                   |                   |
| Smoking                                       | 3.621 (2.342–5.598) | 3.512 (2.010–6.139) |
| Drinking                                      | 2.178 (1.401–3.387) | 1.326 (0.728–2.412) |
| Diabetes mellitus                             | 1.164 (0.616–2.197) | 1.078 (0.488–2.378) |
| Fatty liver                                   | 1.825 (1.179–2.826) | 1.026 (0.593–1.775) |
| Etiology of AP                                |                   |                   |
| Biliary                                       | Ref (1.00)         | Ref (1.00)         |
| HTG                                           | 2.201 (1.307–3.707) | 1.078 (0.488–2.378) |
| Alcohol                                       | 1.998 (1.008–3.959) | 0.671 (0.283–1.591) |
| Others                                        | 1.723 (0.869–3.414) | 1.709 (0.812–3.595) |
| Local complication                            |                   |                   |
| Acute peripancreatic fluid collection         | 2.243 (1.429–3.521) | 1.608 (0.760–3.401) |
| Acute necrotic collection                     | 1.372 (0.707–2.659) | 1.952 (0.818–4.659) |
| Pancreatic pseudocyst                         | 2.628 (1.266–5.455) | 0.010             |
| Walled-off necrosis                           | 2.460 (0.900–6.723) | 0.079             |
| Infected pancreatic necrosis                 | 4.908 (1.982–12.151) | 0.001             |
| MCTSI                                         |                   |                   |
| Mild                                          | Ref (1.00)         | Ref (1.00)         |
| Moderate                                      | 1.957 (1.148–3.337) | 1.209 (0.567–2.578) |
| Severe                                        | 2.169 (0.974–4.829) | 0.721 (0.211–2.461) |
| Severity of AP                                |                   |                   |
| MAP                                           | Ref (1.00)         | Ref (1.00)         |
| MSAP                                          | 1.272 (0.767–2.111) | 0.647 (0.348–1.204) |
| SAP                                           | 1.885 (1.047–3.393) | 1.101 (0.488–2.483) |
| Laboratory test                               |                   |                   |
| High D-dimer                                  | 0.933 (0.533–1.576) | 0.797             |
| Low-Ca²⁺                                      | 1.262 (0.783–2.034) | 0.339             |
| High-FBG                                      | 1.331 (1.048–1.689) | 1.356 (0.785–2.342) |
| High-HbAlc                                    | 1.553 (0.999–2.416) | 0.351             |
| Low-ALB                                       | 1.148 (0.738–1.787) | 0.647 (0.348–1.204) |
| Low-Hb                                        | 0.991 (0.618–1.588) | 0.616             |
| High-TC                                       | 1.367 (0.883–2.118) | 0.019             |
| High-TG                                       | 1.842 (1.189–2.851) | 0.006             |
| Low-HDLc                                      | 1.382 (0.810–2.360) | 0.006             |
| High-LDLc                                     | 2.548 (2.038–3.186) | 0.006             |
| High-CRP                                      | 1.251 (0.691–2.264) | 0.006             |

Note: Risk factors showing potential association (p < 0.05) in univariate analysis were subsequently entered into a multivariable regression.
Abbreviations: BMI, body mass index; HTG, hypertriglyceridemia; MCTSI, modified computed tomography severity index; MAP, mild acute pancreatitis; MSAP, moderately severe acute pancreatitis; SAP, severe acute pancreatitis; Ca²⁺, serum calcium; FBG, fasting blood glucose; HbAlc, glycated hemoglobin Alc; ALB, albumin; Hb, hemoglobin; TC, total cholesterol; TG, triglyceride; HDLc, high-density lipoprotein cholesterol; LDLc, low-density lipoprotein cholesterol; CRP, C-reactive protein.
intake, microcirculation disorders) and is more severe and has a worse prognosis than other causes, which may also contribute to its higher recurrence rate. We also found a proportion of patients with pancreas congenital malformation in the RAP group. Earlier studies had shown that pancreas divisum is associated with RAP, which may be related to sustained ductal hypertension due to narrower minor papilla and minor duct. Bertin et al suggested that pancreas divisum itself was not a cause of pancreatitis and that pancreatitis occurred as a cumulative effect of CFTR mutations and pancreas divisum. The different etiologies of acute pancreatitis and the mechanisms associated with the recurrence of acute pancreatitis remain to be further explored. As is known to all that factors that can lead to AP are more likely to become causes of RAP. It is essential to determine the etiology of the disease in time to apply the most appropriate therapy.

There seems to be a consensus that cigarette consumption is a risk factor for RAP. We found that smokers had a significantly higher cumulative risk for RAP than non-smokers. However, the mechanism by which cigarette consumption increases the recurrence rate of pancreatitis remains unclear. Wittel et al suggested that tobacco smoke exposure causes pancreatic damage, in part, by altering the balance between trypsinogen and pancreas-specific trypsin inhibitor, which creates a microenvironment that is conducive to pancreatic autodigestion and promotes acinar cell injury. Chowdhury et al showed that nicotine can activate multiple pathways in acinar cells and induce oxidative stress. These changes cause calcium influx leading to cell injury. In addition, we suspect that chronic vascular damage caused by smoking may also be a factor of recurrence. Previous studies have shown that exposure to tobacco smoke impairs the function of vascular endothelial cells, but most of the research focused on coronary arteries. Further research is needed on the pathogenesis of RAP caused by smoking and whether smoking cessation can reduce the risk of recurrence. Anyway, physicians should be advising patients to quit smoking at all times.

In this study, we found that high LDLc level is associated with pancreatitis recurrence. As far as we know, this is the first time that the conclusion has been put forward. It is well known that high LDLc level triggers a vicious cycle of oxidative stress that leads to vascular endothelial dysfunction. We suspect this may be related to recurrent episodes of pancreatitis. We know that one consequence of pancreatitis is the vascular damage within the pancreas. However, whether chronic vascular endothelial dysfunction leads to pancreatic vulnerable is unknown. Due to the effects of cigarette smoke exposure and high LDLc level on vascular injury are clear, we hypothesized that these two factors may contribute to recurrence of pancreatitis by causing chronic vascular damage.

The link between RAP and pancreatic cancer was not fully elaborated. A population-based cohort study found that RAP may be an independent risk factor for pancreatic cancer. Although RAP can be a sign of pancreatic cancer, it is also plausible that inflammatory process of the pancreas triggers the development of pancreatic cancer. Identifying the risk factors for recurrence of AP can not only prevent RAP, but also reduce the incidence of pancreatic cancer possibly. Previous studies have shown that the risk of AP in diabetic patients is 1.86–2.89 times higher than that in the general population. However, there are few studies on whether first episode AP with diabetes mellitus increases the risk of pancreatitis recurrence. Our research found that patients with high-HbA1c at
admission for the first attack of AP were more likely to have RAP. Unfortunately, this is not an independent risk factor for recurrence of AP.

The severity of the first episode of AP does not seem to be significantly related to the recurrence in this study. This is consistent with previous research. Cho et al found that a local complication at index admission in patients with first episode pancreatitis was the strongest risk factor for RAP. The occurrence of local complications may be related to structural changes in the pancreas after recovery from pancreatitis. Our regression analysis found that patients who developed infected pancreatic necrosis during their first hospitalization had a fivefold higher risk of recurrence than those who without. A cross-sectional study in the Netherlands found that pancreatic necrosis was an independent risk factor for recurrent pancreatitis. The scar formed during the healing process of pancreatic necrosis, especially the scar of pancreatic duct, may be an important cause of pancreatitis repeated attacks.

There are a number of limitations in our study. First, this is a observation study conducted in a single center. Second, our cases were followed for a short time that the incidence of RAP may be underestimated. A longer follow-up duration may have provided further insights. Third, the effect of smoking on recurrence of pancreatitis was not quantified. Fourth, our strict definition of alcohol-induced AP may understate the incidence of this etiology. In the future, we will verify our results in pathophysiology and design large-scale prospective studies to quantify risk factors.

Conclusions
In this study, 13.7% of the patients developed RAP after a first-attack of AP. Smokers, patients with infected pancreatic necrosis during first admission and patients with high LDLc level had significantly higher cumulative risk for RAP over 3 years. Cigarette consumption, infected pancreatic necrosis and high LDLc level were independent risk factors of RAP. In the future, we should help patients improve their awareness of risk factors for RAP, improve compliance behavior and advocate a healthy lifestyle, so as to prevent recurrence of pancreatitis.

Abbreviations
OAP, once episode of acute pancreatitis; RAP, recurrent acute pancreatitis; BMI, body mass index; HTG, hypertriglyceridemia; MCTSI, modified computed tomography severity index; MAP, mild acute pancreatitis; MSAP, moderately severe acute pancreatitis; SAP, severe acute pancreatitis; Ca²⁺, serum calcium; FBG, fasting blood glucose; HbA1c, glycated hemoglobin A1c; ALB, albumin; Hb, hemoglobin; TC, total cholesterol; TG, triglyceride; HDLc, high-density lipoprotein cholesterol; LDLc, low-density lipoprotein cholesterol; CRP, C-reactive protein; IPN, infected pancreatic necrosis.

Data Sharing Statement
The data used and analyzed in the current study can be made available upon request to the corresponding author.

Ethics Approval and Consent to Participate
The study has been ethically approved by the ethics committee of Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine (IRB No. 021KY049). The respondents were informed about the aim of the study. The respondents provided informed consent for the survey. This study was conducted in accordance with the declaration of Helsinki. The confidentiality and anonymity of the data was also ensured.

Author Contributions
All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Disclosure

The authors report no conflicts of interest in this work.

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