Experimental study of “Tong Xia” purgative method in ameliorating lung injury in acute necrotizing pancreatitis

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Abstract

AIM To investigate the role of tumor necrosis factor (TNF) in lung injury during acute necrotizing pancreatitis (ANP), and the therapeutic effect of “Tong Xia” purgative method in minimizing the severity of lung injury. METHODS Fourteen canines were randomly divided into 3 groups: the “Tong Xia” treatment group (n = 5) using Dachengqitang; saline control group (n = 5), and the sham operation group (n = 4). TNF activity in serum and in bronchoalveolar lavage fluid (BALF), the serum endotoxin levels were measured, and the severity of lung injury evaluated. RESULTS Elevation of TNF activity was more prominent in BALF than in serum. TNF activity in serum at 6 and 12 hours and in BALF was significantly decreased in the “Tong Xia” treatment group than in the saline control one (q = 21.11, q = 12.07, q = 9.03, respectively, \( P < 0.01 \)) and the lung injury was significantly alleviated at 12 hours as compared with that in the saline group, manifested as amelioration of the lung wet/dry weight ratio, decrease in protein concentration and neutrophils count in BALF, and improvement of pulmonary inflammatory changes. A positive correlation was demonstrated between serum TNF activity and endotoxin level. CONCLUSION Hypersecretion of TNF is shown to be one of the major causes of lung injury during ANP; “Tong Xia” purgative method could alleviate the degree of lung injury mediated by TNF.

INTRODUCTION

In the process of the multiple organ dysfunction syndrome (MODS) caused by severe acute pancreatitis, acute lung injury (ALI) occurs most frequently. A number of cytokines are involved in ALI caused by severe acute pancreatitis due to excessive stimulation of monocyte macrophages provoked by endotoxin. Dachengqitang can inhibit the absorption of enterogenous endotoxin and relieve endotoxemia so as to prevent and ameliorate the MODS including the lungs. The purpose of this study is to investigate the effects of TNF on ALI in acute necrotizing pancreatitis (ANP) and to determine whether the herb mixture could relieve the ALI, and investigate its mechanism of action.

MATERIALS AND METHODS

Grouping of experimental animals

Fourteen healthy adult canines, male and female, weighing 7.7 kg-9.9 kg (8.8±1.1), were used. The ANP model canines were divided randomly into 3 groups, i.e. saline control group (n = 5), Dachengqitang treatment group (n = 5), and sham operation group (n = 4).

Preparation of animal models

ANP models were made by the retrograde injection of 5% sodium taurocholate into the pancreatic duct. The canines were only accessible to drinking water. 24 hours later were anesthetized with 3% pentobarbital sodium (1 mL/kg) intraperitoneally...
in normal saline (10 mL·kg⁻¹·h⁻¹) dripping continuously. A median incision was made in the upper abdomen to expose the duodenum, then a transverse incision (2 cm) was made at the level of the orifice of the main pancreatic duct. A catheter was inserted and 5% sodium taurocholate (1 mL/kg) was injected into the pancreatic duct at a pressure of 16 kPa, then pulled out, the duodenal dwelling catheter was placed, and Chinese herbal medicine or normal saline could be given via the duodenum. In sham operation group, the catheter was pulled out and no medicine or saline was given.

**Preparation of Chinese herbal medicine**

Dachengqitang consisted of Radix Rhizoma Rhei 30 g, Natrii Sulphas 20 g, Cortex Magnoliae Officinalis 30 g, and Fructus Aurantii Immaturus 15 g. The latter two were decocted for 25 minutes with 500 mL water, then added with Radix Rhizoma Rhei, decocted for another 3 minutes, and then filtered to yield 150 mL medicinal fluid. Finally, the Natrii Sulphas was added. In treatment group, 50 mL of the medicinal fluid and saline were given through the duodenal fistula tube at 0.5, 4 and 8 hours, respectively.

**Observation indices**

**Serum amylase** It was measured before the model was made and at 2, 6 and 12 hours after.

**Serum endotoxin** It was measured with the o·azo coloration before the model preparation and at 2, 6 and 12 hours after, 2 mL blood was drawn from the femoral vein, and the serum was isolated for measurement. The standard curve correlation coefficient was $r = 0.999$.

**Serum TNF activity** It was measured with the method of the mouse fibroblast L929 biological activity quantitative analysis, the same procedure was carried out before and after.

**Broncho-alveolar lavage fluid (BALF)** The canines were exsanguinated to death 12 hours after the model preparation. After thoracotomy, the whole lungs together with the trachea were taken out, the left bronchus was ligated and the left lung was removed and weighed; the right lung together with the trachea was used for lavage purpose. The surface blood was washed away with normal saline. A total of 100 mL of normal saline was repeatedly lavaged for 5 times, and 80 mL of the lavage fluid was collected. Ten mL was centrifuged, and 2 mL of supernatant was obtained for protein content measurement; another 2 mL was used for the TNF activity measurement. The sediment was stained for the counting of all kinds of nucleated cells.

**Pathological examination**

The lung and the pancreatic tissues were sectioned and preserved in 10% formalin fluid.

**The lung wet/dry weight ratio** The wet left lung was weighed and placed in the oven for 24 hours at 60°C, and weighed when it was dry. The wet/dry weight ratio was calculated.

**Statistical analysis**

All data were placed in the computer for the statistical analysis with the STATA statistics software.

**RESULTS**

**Autopsy findings**

**Abdominal cavity** In both saline control group and treatment group, as much as 300 mL–500 mL of bloody ascitic fluid were present. There was no statistical significance between them. In sham operation group, only a small amount of clear ascitic fluid was noted. In saline control group and treatment group, the pancreas appeared violet black and enlarged, subserous hemorrhage and a little saponification were observed.

The lungs In saline control group, there were various degrees of pulmonary edema and extensive patchy or petechial hemorrhage. In treatment group, the pulmonary edema was less severe without any patchy hemorrhage. In sham operation group, the lung tissue was normal.

**Blood amylase (Figure 1).**

![Figure 1 Serum amylase.](image)

$^aP<0.01$ vs preoperation; $^bP<0.01$ vs sham operation.
Serum endotoxin (Figure 2).

Figure 2 Serum endotoxin.

Figure 3 Serum TNF activity.

Cell changes in BALF (Table 1).

Table 1 Cell changes in BALF (×10⁹/L)

| Group           | Total cell count | Differential count | Other |
|-----------------|------------------|--------------------|-------|
| Saline          | 36.4±8.2             | 3.1±1.2          | 11.2±3.5 | 22.1±7.6 |
| “Tong Xia”      | 20.7±10.3            | 0.8±0.2         | 5.3±2.3   | 14.6±6.3 |
| Sham operation  | 7.2±4.1             | 0.2±0.07        | 4.2±2.3   | 2.8±0.18 |

Protein content and TNF activity in BALF (Table 2).

Table 2 Protein content and TNF activity in BALF (×10⁹/L)

| Group           | TNF activity (IU/mL) | Protein content |
|-----------------|----------------------|-----------------|
| Saline          | 58.75±4.65           | 2.53±0.83       |
| “Tong Xia”      | 49.00±1.83           | 1.42±0.31       |
| Sham operation  | 17.00±0.82           | 0.93±0.21       |

The correlation analysis between serum TNF and endotoxin

The linear correlation analysis showed that there was a positive correlation between serum TNF activity and endotoxin level in saline control group ($r = 0.9706, P < 0.05$).

Microscopic changes

The pancreatic and pulmonary lesions in the three groups (Table 4) The pancreatic lesion was graded according to Meng’s criteria[2] whereas the pulmonary lesion was graded on basis of Lei’s criteria[3].

Table 4 The pancreatic and pulmonary lesions

| Group           | Pancreatic lesion | Pulmonary lesion |
|-----------------|------------------|-----------------|
| Saline          | Negative Mild Moderate Severe | 0 | 1 | 2 | 3 | 4 | 5 |
| “Tong Xia”      | 5 | 2 | 3 | 1 | 3 | 1 |
| Sham operation  | 4 | 4 | 4 | 4 | 4 |

DISCUSSION

MODS is frequently seen in severe acute pancreatitis. It is probably due to the cytokines produced by excessive activation of the monocyte-macrophage system, and the influx and infiltration of polymorphonuclear leukocyte cytokines as TNF, IL, inflammatory mediators, phospholipase A2, oxygen free radicals, cathepsin may all contribute[4]. Among them TNF plays an important role in the processes. Many authors reported that TNF activity was increased in the serum and BALF from the patients with severe acute pancreatitis and adult respiratory distress syndrome[5]. Majority of the authors believed that the increase in TNF activity was due to excessive stimulation of mononuclear-macrophage by endotoxin. In this study, serum TNF activity was remarkably increased, which indicated and was involved in the pathophysiological processes of ANP. TNF activity was also increased more markedly in the BALF than that in the serum, which suggested that TNF also came from pulmonary alveolar macrophages. There was a positive correlation between serum TNF activity and endotoxin level, which indicated that...
the increase in TNF activity was caused by the stimulation of endotoxin. In BALF, polymorphonuclears and macrophages were both increased more remarkably than those in sham operation group, indicating the involvement of polymorphonuclears in the lung injury. In treatment group, at 6 and 12 hours after the model preparation, TNF activities in both serum and BALF were lower than those in saline control group, and the lung injury was also much milder, indicating that Dachengqitang could ameliorate the lung injury mediated by TNF.

“Tong Xia” therapeutic method, is an important part of traditional Chinese medicine in treating acute pancreatitis, and has been studied extensively by Chinese scholars. As regards to the therapeutic mechanism, the herb mixture by purgation could decrease the absorption of the gut endotoxin, ameliorate the endotoxemia and translocation of gut bacteria, and protect the gut barrier function. In this study, at 6 hours after ANP, the serum endotoxin level was a little lower in the treatment group than that in saline control one, and the serum TNF activity was lowered very much, suggesting that Dachengqitang inhibited TNF activity via inhibiting the absorption of endotoxin.

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