Expression of TREM-1 mRNA in acute pancreatitis

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Received: 2003-09-06 Accepted: 2003-10-07

Abstract

AIM: To explore the expression of triggering receptor expressed on myeloid cells (TREM-1) mRNA in acute pancreatitis (AP).

METHODS: Using the reverse transcription polymerase chain reaction (RT-PCR), we examined the expression of TREM-1 mRNA in 10 cases of mild acute pancreatitis (MAP), 8 cases of severe acute pancreatitis (SAP), and 10 cases of healthy control subjects. And we also examined the expression of TREM-1 mRNA in 14 cases of AP (including 10 MAP and 4 SAP) before treatment, after successful therapy and clinically cured.

RESULTS: The expression of TREM-1 mRNA in the groups of MAP, SAP patients and healthy control subjects was 0.771±0.274, 1.092±0.331 and 0.459±0.175, respectively; there was a significant difference among the three groups (P<0.05). And there was also a significant difference between the AP patients (0.914±0.341) and healthy control subjects (0.459±0.175) (P<0.05). Moreover, in the 14 cases of AP, before treatment, after successful therapy and clinically cured, the expression of TREM-1 mRNA was 0.905±0.226, 0.739±0.169 and 0.633±0.140, respectively, and there was a significant difference among the three stages (P<0.05).

CONCLUSION: The expression of TREM-1 mRNA in the patients with AP increases obviously, and correlates with the degree of AP. Furthermore, the expression of TREM-1 mRNA is distinctly different at the different stages of AP. It indicates TREM-1 may play an important role in the occurrence and development of AP.

Wang DY, Qin RY, Liu ZR, Gupta MK, Chang Q. Expression of TREM-1 mRNA in acute pancreatitis. World J Gastroenterol 2004; 10(18): 2744-2746

http://www.wjgnet.com/1007-9327/10/2744.asp

INTRODUCTION

TREM-1 is a new activating receptor of the Ig superfamily expressed on human myeloid cells. And it is selectively expressed on blood neutrophils and a subset of monocytes and up-regulated by bacterial lipopolysaccharide (LPS)[1]. Researchers have demonstrated that TREM-1 mediates activation of neutrophil and monocytes, and may have a predominant role in inflammatory responses such as septic shock and systemic inflammatory response syndrome (SIRS)[2,3]. In this study, we aimed to explore the expression of TREM-1 mRNA in AP and tried to reveal etiology of AP and underlying mechanism.

MATERIALS AND METHODS

Patients and specimens

Data obtained from 18 AP patients without any treatment during 48 h of the onset of the disease and 10 healthy control subjects were explored. According to the standards of diagnosis and classification of AP proposed at the Sixth National Conference on Pancreatic Surgery of China in 1996[4], we divided the AP patients into two groups: 10 cases of MAP, and 8 of SAP. Among the AP patients, we examined the expression of TREM-1 in 14 patients at different stages of the disease (before treatment, after successful therapy and clinically cured). We collected 2 mL of blood from the patients and controls before breakfast in the morning and stored them in the EDTA tube. The total leucocytes in the blood were separated by centrifugation at 1 200 r/min for 15 min.

Reagents and primers

Trizol reagents, Oligo (dt) 15, M-MLV reverse transcriptase and DEPC were the products of Promega Co. (USA). Taq DNA polymerase were purchased from Sangon Co. (Shanghai). The RNase inhibitor was from Takara Co. (Japan). According to the human TREM-1 and β-actin cDNA sequence reported in GeneBank, we designed two sets of primers: TREM-1 sense 5’-GTG TGT GAT CAT CCA GCC T-3’, antisense 5’-TTC AGA GTC AGG AGT GGA G-3’ (amplified fragment length: 232 bp); β-actin sense 5’-GTG CGT GAG ATT AAG GAG-3’, antisense 5’-CTA AGT CAT AGT TCC CCT-3’ (amplified fragment length: 520 bp).

Total RNA extraction

The total RNA was extracted from the leucocytes by Trizol reagent. The extracted RNA concentrations and purity were determined at A260 and A280 spectrophotometrically (Beckman DU650, USA). Then 750 mL/L ethanol was used to preserve total RNA at -20 °C.

Reverse transcription polymerase chain reaction

The total RNA was denatured by incubation for 5 min at 70 °C and rapidly cooled on ice for about 2-3 min. Single-stranded cDNA was synthesized by mixing 1-2 µg denatured RNA with 0.5 µL (40 u/µL) RNase inhibitor, 1 µL (200 u/µL) M-MLV reverse transcriptase, 4 µL reaction buffer, 1 µL OligodT (15), 1 µL (10 mmol/L) dNTP, and adding DEPC-treated water up to 20 µL. The RT reaction mixture was incubated at 37 °C for 60 min, followed by 95 °C for 5 min to inactivate M-MLV reverse transcriptase, and stored at 4 °C for use. A 5 µL of the cDNA was used as template in PCR amplification for TREM-1. The mixture (50 µL) consisting of 1 µL of primers (p1/p2 of TREM-1 and β-actin) each, Taq DNA polymerase (0.5 µL (5 U/µL),10×PCR buffer 5 µL, MgCl2 3 µL (25 mmol/L), ), by addition of deionised water up to 50 µL, was reacted on a programmable thermocycler (Perkinelmer 9600). The DNA was prenatured at 95 °C for 5 min, followed by 35 cycles of denaturation (45 s at 95 °C), annealing (45 s at 65 °C), and extension (45 s at 72 °C). Then the mixture was finally extended for 10 min at 72 °C, and stored at -20 °C. β-actin was served as internal control in each experiment under the same conditions.

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Electrophoresis identification and quantitative scanning

PCR products (7-9 µL) with DL2000 marker were electrophoresed through 10 g/L agarose gel in 1×TBE at 70 V for 30 min. The samples were stained with ethidium bromide (0.5 mg/L) and observed by ultraviolet illumination and photographed. Each lane was scanned longitudinally. The A values of TREM-1 and β-actin bands were measured by a gel imaging analysis system (GELWORKS 1D advanced v4.01). The A ratios of TREM-1/β-actin representing the relative levels of the expression of TREM-1 mRNA were calculated for semi-quantitative analysis.

Statistical analysis

Results were presented as mean±SD. Statistical comparisons were analyzed through the SAS 8.1 software. Analysis of variance was used. P<0.05 was considered statistically significant.

RESULTS

Expression of TREM-1 mRNA and semi-quantitative analysis

The expression of TREM-1 mRNA in the groups of MAP, SAP patients and healthy control subjects were 0.771±0.274, 1.092±0.331 and 0.459±0.175, respectively. There was a significant difference among the three groups (P<0.05). The difference was also significant (P<0.05) (Figures 1, 3) between the AP patients (0.914±0.341) and healthy control subjects (0.459±0.175). In AP patients before treatment, after successful therapy and clinically cured, the expression of TREM-1 mRNA was 0.905±0.226, 0.739±0.169 and 0.633±0.140, respectively. There was a significant difference among the three stages (P<0.01 or P<0.05). And the expression of TREM-1 mRNA was higher in clinically cured group than in healthy control subjects (P<0.05) (Figures 2, 4).

DISCUSSION

AP is a disease of self-digestive chemical inflammation with unclear etiology. In most cases, they present a self-limited course. Twenty percent of patients with AP have a severe form of the disease. SAP is such a severe disease that it may be associated with both systemic and local complications, such as respiratory insufficiency with adult respiratory distress syndrome (ARDS), renal failure, the development of pancreatic and peripancreatic infection. The mortality of SAP is higher than 40%[4,5]. Even now, there are no efficient and sensitive methods to treat AP successfully because the pathogenesis of AP still remains obscure[6]. Recent studies[7] show that factors like the pancreatic enzymes activated abnormally lead the body to secrete and amplify large quantities of cytokines. Finally, the cytokines increase vascular permeability, activate complement components, resulting in microcirculatory impairment and imbalance of thrombo-fibrinolytic system. Many of these events occur not only in the pancreas itself, but also in the other vital organs and tissues, leading to SAP and complications[8,9]. Generally, there are three phases of AP progression: local acinar injury, systemic response, and general sepsis[7]. Moreover, some data suggest that the changes in concentration of proinflammatory cytokines can be an early diagnostic criterion for AP[10,11]. Most authors have realized that the pathogenesis of SAP is very complicated. It is a multifactorial as well as multifaceted disease. First of all, the etiologic agents initiate the pancreatic acinar injury by release of pancreatic enzymes and over-stimulation of...
In 2000, Bouchon et al. had identified a new activating receptor of the Ig superfamily expressed on human myeloid cells, called triggering receptor expressed on myeloid cells (TREM). TREM-1 is selectively expressed on blood neutrophils and a subset of monocytes and up-regulated by bacterial LPS. Engagement of TREM-1 triggers secretion of IL-8, monocyte chemotactic protein-1, and TNF-alpha and induces neutrophil degranulation. Intracellularly, TREM-1 induces Ca^{2+} mobilization and tyrosine phosphorylation of extracellular signal-related kinase 1 (ERK1), ERK2 and phospholipase C-gamma. To mediate activation, TREM-1 associates with the transmembrane adapter molecule DAP12. Furthermore, Bleharski et al. suggested that activation of TREM-1 on monocytes participated in the early induced and adaptive immune responses involved in host defense against microbial challenges. Thus, TREM-1 was considered mediating the activation of neutrophils and monocytes, and might have a predominant role in inflammatory responses. Wilson et al. discovered that SAP had many similarities to sepsis syndrome and septic shock. The haemodynamic features of cardiovascular instability, reduced ejection fraction and decreased systemic vascular resistance were indistinguishable in each of these conditions. In addition, there are many striking similarities in the cytokine and inflammatory mediator profiles, suggesting that the haemodynamic abnormalities may result from the same pathogenic mechanisms, albeit as a result of different inflammatory stimuli.

In 2001 through their experiments, Bouchon et al. believed that inflammatory responses to microbial products were amplified by a pathway mediated by TREM-1. TREM-1 was an activating receptor expressed at high levels in neutrophils and monocytes that infiltrate human tissues infected with bacteria. Furthermore, it was up-regulated in peritoneal neutrophils of patients with microbial sepsis and mice with experimental LPS-induced shock. Notably, blockade of TREM-1 protected mice against LPS-induced shock, as well as microbial sepsis caused by live Escherichia coli or caecal ligation and puncture. Their results demonstrate a critical function of TREM-1 in acute inflammatory responses to bacteria and implicate TREM-1 as a potential therapeutic target for septic shock. A recent study shows that TREM-1 amplifies Toll-like receptor-initiated responses against microbial challenges and potentiates the secretion of proinflammatory chemokines and cytokines in response to bacterial and fungal infections. Blockade of TREM-1 reduced inflammation and increased survival in animal models of bacterial infections that caused systemic hyper-inflammatory syndromes. But nobody knows how TREM-1 expressed in AP as yet. According to the theory mentioned above, we carried out a study of the expression of TREM-1 in AP. We demonstrated that the expression of TREM-1 in leukocytes of patients with AP was much higher than that in healthy control subjects and correlated with the degree of AP. Furthermore, significant differences in the expression of TREM-1 mRNA were found at the different stages of AP. The expression of TREM-1 decreased gradually with the improvement of severity of the AP patients. This experiment indicates that the expression of TREM-1 is relevant to progression of AP. Because of lack of relevant data about the patients after they were discharged from the hospital, we only compared the expression of TREM-1 between the cured group and healthy control subjects and found that there was still a significant difference between them. Therefore, we suggest that the pathological changes of the pancreas continued even after the patients were cured clinically. This study may help reveal the etiology of AP and the mechanism of its progression. It may provide a new standard for the diagnosis and classification of AP as well as an efficient method to cure AP by blocking the expression of TREM-1.

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