Regulatory effect of cytokine-induced neutrophil chemoattractant, epithelial neutrophil-activating peptide 78 and pyrrolidine dithiocarbamate on pulmonary neutrophil aggregation mediated by nuclear factor-κB in lipopolysaccharide-induced acute respiratory distress syndrome mice

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Abstract. In the present study, the regulatory effect of cytokine-induced neutrophil chemoattractant (CINC) and epithelial neutrophil-activating peptide 78 (ENA-78) on pulmonary neutrophil (PMN) accumulation in lipopolysaccharide (LPS)-induced acute respiratory distress syndrome (ARDS) mice, and the therapeutic effect of pyrrolidine dithiocarbamate (PDTC), was investigated. BALB/c mice were divided into control, LPS and PDTC + LPS groups using a random number table. The phosphorylation of nuclear factor-κB (NF-κB) was detected using a western blot, and the mRNA expression levels of CINC were evaluated using reverse transcription-quantitative polymerase chain reaction. The expression of NF-κB, CINC and ENA-78 was detected using immunohistochemistry. The production of interleukin (IL)-8 and IL-10 in serum and broncho-alveolar lavage fluid (BALF) was analyzed using an enzyme-linked immunosorbent assay. The total number of leukocytes and proportion of PMNs in BALF was also determined. Following injection with LPS (20 mg/kg), the expression levels of p-NF-κB, CINC and ENA-78 were increased in lung tissue, and the expression levels of IL-8, IL-10 and the number of PMNs increased in serum and BALF. However, in comparison with the LPS group, the degree of lung injury was reduced in ARDS mice that were treated with PDTC. In addition, the expression level of p-NF-κB and the production of chemokines in lung tissue decreased in ARDS mice that were treated with PDTC, and the number of PMNs in BALF also decreased. In conclusion, the results of the present study suggest that the LPS-induced phosphorylation of NF-κB may result in the synthesis and release of CINC and ENA-78, which induce the accumulation of PMNs in the lung. Therefore, PDTC may be used to reduce the production of chemokines and cytokines, thereby decreasing the activation of PMNs in lung tissue and reducing the damage of lung tissue in ARDS.

Introduction

Acute respiratory distress syndrome (ARDS) is a common and severe lung disease that is associated with high rates of mortality and morbidity, and leads to reduced levels of oxygen in the blood (1,2). The characteristic features of ARDS include excessively uncontrolled inflammation, hypoxemia and non-cardiogenic pulmonary edema formation (3). These are caused by various increased inflammatory cytokines and pulmonary microvascular permeability (4).

The main sites of cell injury in ARDS are the vascular endothelium and alveolar epithelium (4). Polymorphonuclear neutrophils (PMNs) contribute towards lung inflammation and serve important roles in the pathogenesis and progression of ARDS (3). Lung injuries may result in the infiltration and activation of PMNs, involving a complex process including the recruitment, adhesion and chemotaxis of PMNs (3,4). Bhatia and Moochhala (5) have demonstrated that a large number of PMNs can accumulate in lung tissue and release inflammatory cytokines, such as interleukin (IL)-8, IL-10 and tumor necrosis factor-alpha (TNF-α),...
which serve important roles in activating and maintaining the inflammatory response (6). IL-8 and the epithelial neutrophil activating peptide (ENA-78) are critical for the accumulation of PMNs in lung tissue (7); the expression of adhesion molecules on the surface of PMNs increases with appearance of these chemokines (5). Adhesion molecules and their ligands provide a strong adhesion between PMNs and endothelial cells, allowing PMNs to migrate to the vessel wall (8,9). Cytokine-induced neutrophil chemoattractant (CINC), a member of the IL-8 family, is a specific PMN chemokine that serves a crucial function in the aggregation of PMNs in lung tissue (10,11).

Nuclear factor-κB (NF-κB) is a critical nuclear transcription factor, able to activate the transcription of a number of inflammatory cytokine genes and regulate the inflammatory response and immunoreaction (12). Typically, NF-κB exists as an inactive dimer in the cytoplasm and directly combines with an inhibitor of nuclear factor-κB (IκB) to produce a trimeric complex (13). The P50/P65 heterodimer serves an important physiological function during inflammation, and NF-κB P65 is the principal subunit (14).

Lipopolysaccharide (LPS) is the main component of gram-negative bacteria outer membranes, which is a common trigger of sepsis, and is the initiation factor that activates the NF-κB signaling pathway (15). Phosphorylated (p)-NF-κB can enter into the nucleus and bind to specific DNA sequences when stimulated by LPS (15). The activation of NF-κB is closely associated with the overexpression of adhesion molecules, chemokines and other cytokines involved in the migration of PMNs, and therefore serves an important role in the regulation of inflammation and ARDS (16). However, IL-10 is a principal anti-inflammatory cytokine and can inhibit the activation of NF-κB (17).

Pyrrolidine dithiocarbamate (PDTC), an inhibitor of NF-κB, has been reported to inhibit the expression of inflammatory cytokines, such as ILs and TNF-α, at the stage of transcription (18). In addition, PDTC reduces the expression of chemokines, such as CINC and ENA-78, and the accumulation of inflammatory cells in lung tissue in order to alleviate the pathological changes in the lung tissue of ARDS (11,19). It is therefore important to reduce the expression of CINC and ENA-78, and the accumulation of PMNs in lung tissue, by regulating NF-κB phosphorylation and hindering NF-κB activation. The present study aimed to investigate the regulatory effect of CINC and ENA-78 on PMN aggregation mediated by NF-κB, and the direct protective effects of PDTC on lung tissue in LPS-induced ARDS mice.

Materials and methods

Animals. A total of 90 BALB/c mice (age, 8-10 weeks; weight, 20±2 g) were purchased from the Experimental Animal Center of Shandong University (Jinan, China) and housed at room temperature (24°C) with a 12-h light/dark cycle. The mice were allowed free access to water and standard laboratory chow. The experimental procedures were approved by the Ethics Review Committee for Animal Studies at Qilu Hospital, Shandong University (Jinan, China) and performed in accordance with animal welfare and animal experimental guidelines.

Mice received an intraperitoneal (i.p.) injection of 20 mg/kg LPS (Escherichia coli O55:B5; Sigma-Aldrich, St. Louis, MO, USA) and an i.p. injection of PDTC (0, 40, 120 or 160 mg/kg; L04358; USA Alikesi International Group (China), Ltd.). PDTC (Beyotime Institute of Biotechnology, Haimen, China) was administered 30 min prior the injection of LPS.

To further investigate the protective effect of PDTC on LPS-induced ARDS mice, 90 mice were randomly divided into three groups (n=30/group), as previously described (20): Control (20 ml/kg normal saline, i.p.); LPS (20 mg/kg, i.p.); and PDTC (120 mg/kg, i.p.) + LPS (20 mg/kg, i.p.).

Specimen collection. Blood, lung tissue and bronchoalveolar lavage fluid (BALF) samples from each group of mice were collected simultaneously after modeling for 2, 6, 12 or 24 h. The mice were anesthetized by intraperitoneal injection with 10% chloral hydrate (3.5 ml/kg; Sigma-Aldrich), prior to sacrifice via aortic phlebotomy at the indicated time points. Subsequently, the lungs were extracted and the left lung was prepared for hematoxylin and eosin (HE) staining (Beyotime Institute of Biotechnology). Subsequently, the lungs were extracted and the left lung was prepared for hematoxylin and eosin (HE) staining (Beyotime Institute of Biotechnology) and immunohistochemistry, while the right lung was prepared for western blot analysis. PMNs were isolated from BALF using Wright-Giemsa staining (Beijing Leagene Biotech, Co., Ltd., Beijing, China). After centrifugation at 1,200 x g for 10 min at 4°C, the supernatant was collected and the expression of IL-8 and IL-10 was detected using enzyme-linked immunosorbent assay (ELISA) kits. Specifically, the expression of IL-8 was detected using the Quantikine ELISA kit from R&D Systems Europe, Ltd. (Abingdon, UK), whereas the expression of IL-10 was detected using the LEGEND MAX™ Mouse IL-10 ELISA kit from BioLegend, Inc. (San Diego, CA, USA).

Histopathological analysis. The left lung was fixed with 4% paraformaldehyde (Beijing CellChip Biotechnology, Co., Ltd., Beijing, China) for 24 h, embedded in paraffin and cut into 4 μm sections. Once stained with hematoxylin and eosin, an evaluation was performed to characterize the degree of lung injury. Briefly, the lung injury score was calculated by assessing the degree of inflammatory cell infiltration, hemorrhage, interstitial and alveolar edema and the thickness of the alveolar septum in five random fields in a blind manner using a light microscope (Olympus BX43; Olympus Corporation, Tokyo, Japan).

Determination of the difference between alveolar and arterial oxygen partial pressure [Pa(A-a)O2]. PaO2 and PaCO2 were analyzed in 150-μl arterial blood samples and the oxygen partial pressure (alveolar oxygen partial pressure) was calculated according to the results of a blood gas analysis: PaO2 = (atmospheric pressure - 47) x FiO2 - PaCO2 / R (R, the exchange rate; R=0.8). Alveolar - arterial oxygen partial pressure difference (A-a) O2 = (atmospheric pressure - 47) x FiO2 - PaCO2 / R - arterial blood oxygen partial pressure. The linear correlation coefficient was calculated to study the efficacy of gas exchange.

Extraction of cytoplasm and nuclear proteins. Lung tissue samples weighing ~100 mg were washed with 0.01 M
phosphate-buffered saline (PBS) supplemented with 1.5 ml nuclear protein extract lysis buffer A (BioTeke Corporation, Beijing, China). Subsequently, the samples were placed on ice for 15-30 min and homogenized using an electric homogenizer following the addition of 0.5 ml ice-cold NP-40 (10%; BioVision, Inc., Milpitas, CA, USA). Then, the samples were vortexed for 10 sec and centrifuged at 4°C and 12,000 x g for 30 sec; the supernatant produced was the cytoplasmic protein extract. The precipitate was washed once with cold PBS and then centrifuged at 4°C and 12,000 x g for 30 sec, after which the supernatant was discarded. Subsequently, 1.5 ml join nucleoprotein extract lysis buffer B (BioTeke Corporation) was added and the samples were placed on ice for 30 min, prior to centrifugation at 4°C and 12,000 x g for 2 min in order to produce the aspirate supernatant (nucleoprotein). Using the Bradford Protein Assay, the nuclear protein concentration was adjusted to 0.5-1.0 µg/µL and the samples were placed and stored at -70°C, according to a previous study (20).

**Measurement of NF-κB protein expression by western blot analysis.** The total protein quantity was analyzed using a Pierce BCA Protein Assay kit (#23250; Thermo Fisher Scientific, Inc., Waltham, MA, USA). Equal quantities of total protein (30 µg) were separated on a 10% Bis-Tris gel in NuPAGE MOPS SDS Running Buffer (all Invitrogen; Thermo Fisher Scientific, Inc.) and transferred to a polyvinylidene difluoride (PVDF) membrane (Immobilon; EMD Millipore, Billerica, MA, USA). The PVDF membrane was then blocked using 5% skimmed milk in Tris-buffered saline (TBS; Tris-Cl, 50 mM; NaCl, 150 mM; pH 7.5; Thermo Fisher Scientific, Inc.) for 60 min at room temperature. Next, the membrane was incubated for 15 h at 4°C with mouse anti-pNF-κB monoclonal antibody (1:200; cat. no. sc-166748; Santa Cruz Biotechnology, Inc., Dallas, TX, USA), rabbit anti-pNF-κB p65 polyclonal antibody (1:200; cat. no. ab16502; Abcam, Cambridge, UK) and mouse anti-β-actin monoclonal antibody (1:500; #BA2305; Boster Systems, Inc., Pleasanton, CA, USA). The membrane was washed three times for 5 min each with 1X TBS containing 0.15 Tween-20, and then incubated for 1 h with horse-radish peroxidase (HRP)-conjugated secondary antibodies (1:1,000; sc-2370; Santa Cruz Biotechnology, Inc.) at room temperature. The membrane was exposed to high performance autoradiography film (#87897; Fuji XR film; Fujifilm Corporation, Tokyo, Japan) and visualized using enhanced chemiluminescence reagents and the ChemiScope 2850 Fluorescence/Chemiluminescence Imaging system (Clinx Science Instruments Co., Ltd., Shanghai, China). The integrated density value of the band intensities on the film was analyzed using ImageQuant version 5.2 software (Molecular Devices, Sunnyvale, CA, USA).

**Evaluation of CINC mRNA expression level in lung tissue using reverse transcription-quantitative polymerase chain reaction (RT-qPCR).** RNA was isolated from lung tissue using TRIzol (Invitrogen; Thermo Fisher Scientific, Inc.) and quantified using a NanoDrop 2000/2000c (Thermo Fisher Scientific, Inc.). Subsequently, 1 µg total RNA from each sample was denatured at 70°C for 10 min and chilled on ice for 10 min. RT reactions were performed in a volume of 20 µl containing 4 µl 5X RT buffer (Toyobo Co., Ltd., Osaka, Japan), 1 µl RT Enzyme Mix (Toyobo Co., Ltd.) and 1.0 µl (5 pmol) of the sense and antisense primers (BGI, Shenzhen, China) in the presence of PCR buffer (Toyobo, Co., Ltd.). qPCR was performed in a volume of 20 µl containing 2 µl cDNA, 8 µl forward and reverse primers (10 pmol/µl, 10 µM) and 10 µl QuantiTect SYBRs Green PCR kit (QIAGEN, Inc., Toronto, Canada), which consisted of DNA polymerase, deoxyxynucleotide mix, buffer, MgCl₂, and fluorescent dyes. The gene-specific primer sequences were as follows: CINC forward, 5'-ATTGGAAGACCATTGGT-3' and reverse, 5'-CATAAAATGTCCAAGGGAAG-3'; and GAPDH forward, 5'-GAACTCCAGGTGTTAAGG-3' and reverse, 5'-TCGGTGCAATCTATCTTCTCTT-3'. The PCR protocol consisted of three stages: Denaturation, amplification and melting curve analysis for product identification. The denaturation and amplification conditions were 95°C for 20 min, followed by 40 cycles of 30 sec denaturation at 95°C, 10 sec annealing at 60°C and 15 sec elongation at 72°C. The temperature transition rate was 20°C/sec, except when heating at 72°C, when it was 5°C/sec. Fluorescence was measured at the end of every cycle to allow quantification of cDNA. Relative mRNA expression levels were calculated using the 2^[-ΔΔCq] method (21). This value was then used to determine the relative amount of amplification in each sample by interpolating from a standard curve. The mRNA expression level of CINC was normalized to that of GAPDH. Nuclease-free water was used as a RT-minus control.

**Immunohistochemical analysis NF-κB, CINC and ENA-78 expression in lung tissue.** The lobes of lungs from mice were dissected, fixed in 10% formaldehyde and processed in preparation for immunohistochemistry. Slides were dewaxed and rehydrated, and antigen retrieval was performed using 10 mM sodium citrate (pH 6.1; Beyotime Institute of Biotechnology) and blocked using 5% bovine serum albumin (Sigma-Aldrich) for 60 min at room temperature. Sections of lung tissue (4 µm) were incubated with primary antibodies anti-NF-κB (1:300; Santa Cruz Biotechnology, Inc.), anti-CINC (1:300; Santa Cruz Biotechnology, Inc.) and anti-ENA-78 (1:200; Santa Cruz Biotechnology, Inc.) overnight at 4°C and then with polyclonal immunoglobulins/HRP (1:100; Beyotime Institute of Biotechnology) for 1 h at room temperature. Nuclei were counterstained with HE. Control samples were incubated with the same antibodies. Cover slips were mounted with 80% glycerol (Beijing Zhongshnan Golden bridge Biotechnology, Co., Ltd., Beijing, China). Samples were examined under a microscope equipped with a digital camera (BX51 TRF; Olympus Corporation). ENA-78 positive areas were quantified by densitometry using Image-Pro Plus software 6.0 (Media Cybernetics, Inc., Rockville, MD, USA).

**Measurement of leukocytes and proportion of PMNs in BALF.** A total of 0.5 ml BALF was collected and total cells and neutrophils were counted using a hemocytometer in a double-blind manner for the measurement of leukocytes and the proportion of PMNs.

**Statistical analysis.** The results are expressed as the mean ± standard deviation. Statistical analyses consisted
of a one-way analysis of variance and a Student's t-test was used to compare groups. P<0.05 was considered to indicate a statistically significant difference. The statistical analysis was conducted using SPSS 19.0 software (IBM SPSS, Armonk, NY, USA).

Results

Animal phenotype. The mice in the control group were breathing easily and their lungs appeared pink upon dissection. In the ARDS mice, after 6 h induction with LPS, shortness of breath with oral cyanosis and blood-like liquid in the nose was observed. In addition, the lung volume was increased, the lungs were deep purple in appearance and flake bleeding was observed under the visceral pleura. In the PDTC + LPS group, the shortness of breath was attenuated and the lung was less inflamed and reddened. The scattered bleeding on the surface of the lung and the capsular tension was reduced on the lungs of mice in the PDTC + LPS group in comparison with the LPS group (Fig. 1).

Pathology of lung tissue determined by HE staining. In the control group, the structure of the lung tissue was normal and no inflammatory cells were detected. However, in the LPS group after 12 and 24 h, the alveoli septum was thickened, the pulmonary interstitial was highly congested, the alveolar wall was fractured and a large number of infiltrative inflammatory cells were detected. In addition, as time progressed there was an increase in inflammatory cell infiltration and aggravated lung tissue damage. Similar symptoms were detected in the PDTC + LPS group; however, the lung tissue injury was milder compared with LPS group, as determined by HE staining (Fig. 2).

Effect of PDTC on P(A-a)O₂ in LPS-induced ARDS mice. P(A-a)O₂ was significantly higher in the LPS group in comparison with the control group at each time point (P<0.01; Fig. 3). Following treatment with PDTC, the level of oxygenation improved in lung tissue as time progressed, and the P(A-a)O₂ gradually decreased in comparison with the LPS group (P<0.05). However, P(A-a)O₂ in the PDTC + LPS group was significantly higher compared with the control group (Fig. 3).

Differences in p-NF-κB and NF-κB P65 protein expression in the cytoplasm and nucleus in lung tissue. The expression levels of p-NF-κB were significantly higher in the LPS group in comparison with the control group (P<0.01). However, p-NF-κB protein expression in the lung tissue of the PDTC + LPS group was significantly reduced, as compared with the LPS group (P<0.05), although it remained increased, as compared with the control group. This suggests that PDTC can effectively inhibit the phosphorylation of NF-κB (Fig. 4A and B). NF-κB P65 cytoplasmic protein expression was significantly lower in the LPS group in comparison with the control group (P<0.01); however, the production of P65 was significantly higher in the pulmonary nucleus of the LPS group in comparison with the control group (P<0.01; Fig. 4C and D). P65 protein expression in the cytoplasm of lung tissue was significantly increased in the PDTC + LPS group in comparison with the LPS group (P<0.05), and the expression of P65 was significantly decreased in the nuclei of the PDTC+LPS group in comparison with the LPS group (P<0.01; Fig. 4E and F).

CINC mRNA expression in lung tissue. The expression of CINC mRNA in lung tissue was significantly increased in the mice that received an i.p. injection of LPS in comparison with the control group (P<0.01), and continued to increase over time. In the PDTC + LPS group, CINC mRNA expression in lung tissue was significantly decreased in comparison with the LPS group (P<0.05; Fig. 5).

Expression of NF-κB, CINC and ENA-78 in lung tissue. The immunohistochemistry results demonstrated that the expression levels of NF-κB, CINC and ENA-78 were markedly increased in the lung tissue of LPS-induced ARDS mice. Cell morphological analysis revealed that the majority of cells in the LPS lung tissue were infiltrated PMNs and endothelial cells. However, in the control group, the expression of NF-κB, CINC and ENA-78 was not detected in the lung tissue of mice. The number of positive cells stained with NF-κB, CINC and ENA-78 in PDTC + LPS group was markedly decreased in comparison with the LPS group (Figs. 6-8).

Expression of IL-8 and IL-10 in serum and BALF. The expression levels of IL-8 and IL-10 in the serum and BALF from the LPS group was significantly increased, as compared with the control group (P<0.01). In the PDTC + LPS group, IL-8 expression was significantly decreased, as compared with the LPS group (P<0.05), whereas it was increased in comparison with the control group. Over the time period, however, the production of IL-10 in the serum and BALF from the LPS group was increased. In the PDTC + LPS group, the expression level of IL-10 was substantially higher than that in the control group, whereas it was significantly reduced in comparison with the LPS group (P<0.05; Fig. 9).

Number of leukocytes and proportion of PMNs in BALF. The total number of leukocytes was significantly increased in the LPS group, as compared with the control group (P<0.01). In the PDTC + LPS group, the number of leukocytes was significantly decreased, as compared with the LPS group (P<0.05) at each time point, but was significantly increased, as compared with the control group (P<0.05). The proportion of PMNs in BALF was significantly higher in the LPS and PDTC + LPS groups in comparison with the control group (P<0.01), but the proportion of PMNs was significantly reduced in the PDTC + LPS group in comparison with the LPS group (P<0.05; Fig. 10).

Discussion

NF-κB is a target for anti-inflammatory ARDS treatment and serves an important role in initiating and developing inflammatory reactions (16,22). The association between NF-κB, cytokines and chemokines has become a topic of interest with regards to the pathogenesis of ARDS (12). The expression of inflammatory mediators can be downregulated by delaying the activation of the NF-κB signal pathway,
Figure 1. General condition (left panel) and lungs (right panel) of mice in the (A) normal saline control, (B) lipopolysaccharide (LPS) and (C) LPS + pyrrolidine dithiocarbamate (PDTC) group (n=12). Mice were treated with 20 mg/kg LPS with PDTC (0, 40, 120 or 160 mg/kg, intraperitoneally).

Figure 2. Effect of pyrrolidine dithiocarbamate (PDTC) on lung histopathological changes of mice at 12 and 24 h following intraperitoneal injection with saline, and lipopolysaccharide (LPS) with or without PDTC (stain, hematoxylin and eosin; magnification, x400). (A) Normal saline group with integrated lung tissue and no inflammatory cells; (B) LPS group at 12 h; (C) LPS group at 24 h with widened lung intervals, highly congested pulmonary interstitia, fractured alveolar walls and numerous infiltrative inflammatory cells; and (D) PDTC + LPS group with milder lung tissue injury in comparison with the LPS group.
thereby suppressing PMN accumulation in the lungs, effectively repressing the excessive activation of the inflammatory response and reducing the damage inflicted on lung tissue (23).

A previous study observed that the activation of NF-κB in ARDS could increase the production of inflammatory cytokines, including adhesion molecules (such as CD11b/CD18, ICAM-1), chemokines (such as IL-8), TNF-α and IL-1β, involving a number of downstream gene transcriptions (5). NF-κB was demonstrated to be a central transcription factor that promotes the production of cytokines, adhesion molecules and chemokines by specifically binding to κB promoters in the nucleus (24,25). The P50/P65 NF-κB heterodimer serves a primary physiological function during inflammation (12,13). As the principal subunit, NF-κB P65 exists in an inactive dimer form in the cytoplasm and directly combines with the inhibitory protein IκB to form a trimeric complex (26,27). Previous studies have demonstrated that when NF-κB is excessively activated, IL-8 and TNF-α are overexpressed, whereas the expression of IL-10 is decreased, in ARDS (20). An imbalance of the inflammatory response is the principal feature in ARDS (28,29). Therefore, inhibiting the activation of the NF-κB signal pathway and reducing the expression of inflammatory cytokines and chemokines may inhibit the inflammatory response at the source, leading to novel strategies in treating ARDS.

In the present study, the widened lung interval, highly congested pulmonary interstitial, fractured alveolar wall and increased infiltrative inflammatory cell in the LPS group aggravated lung tissue damage, gradually increased P(A-a)O₂ and resulted in oxygenation obstacles over time. In addition, the results of the NF-κB western blot demonstrated that p-NF-κB protein expression was significantly increased in the lung cells of the ARDS mice at each time point in comparison with the control group. The expression of the P65 protein in the nucleus was significantly enhanced, suggesting that NF-κB may be activated by phosphorylation and degradation. These results are consistent with previous studies (20).

IL-8, a proinflammatory cytokine, is a promoter for ARDS and serves a crucial role in the course of PMN migration from circulating blood to sites of inflammation (30). In patients with ARDS, the body can prevent excessive activation of the inflammatory response by activating an anti-inflammatory response and releasing anti-inflammatory cytokines, such as IL-10, in order to attenuate the extent of tissue damage (31). IL-10 is a crucial anti-inflammatory cytokine in the immune response and can inhibit the production of IL-8 and TNF-α in monocytes and macrophages, and repair injured tissue by regulating the immune response; however, overexpression can aggravate the immune response (17,32). Previous studies demonstrated that the genes of cytokines such as IL-8 and IL-10 encode the binding site for NF-κB (20), and that LPS-activated NF-κB binds to these genes, resulting in their expression (5,33). In the present study, it was observed that the production of IL-8 was rapidly increased after 2 h in the serum and BALF of ARDS mice; however, the expression of IL-10 was not significantly increased during the first 6 h following injection of LPS. This indicates that there is an imbalance of proinflammatory and anti-inflammatory molecules during the early stage of ARDS. Although the level of IL-8 and IL-10 were increased simultaneously over time, the pro- and anti-inflammatory cytokines were imbalanced, and a large number of proinflammatory cytokines released from inflammatory cells further aggravated lung tissue damage. Thus, a cascade of inflammatory responses commenced.

The results from the current study demonstrated that chemokines (in particular IL-8, ENA-78 and CINC) released by damaged cells in ARDS stimulated PMN extravasation and their migration to sites of inflammation. CINC, a member of the IL-8 family, is a specific PMN chemokine (11). CINC-1 and -3 serve important roles in PMN recruitment to the lung in ARDS induced by LPS (10). The enhanced expression of CINC mRNA and protein inhibits PMN apoptosis and activates a large number of PMNs, causing them to accumulate in the lung and result in lung injury (11). Endogenous TNF-α is associated with the increased expression of ENA-78. ENA-78, a member of CXC chemokine superfamily, can activate PMNs, induce secretion of numerous cytokines in PMNs and aggravate inflammation (19). In the present study, the results demonstrated that the expression of CINC mRNA was higher than that in the control group, and the immunohistochemical results demonstrated that there were a large number of positive immunoreactive cells with CINC and ENA-78 in the cytoplasm of macrophages and pulmonary interstitial cells. The number of leukocytes and PMNs in BALF were increased in comparison to the control group, which indicated that the high expression of chemokines (CINC and ENA-78) is closely related to the accumulation of PMNs.

As a dithiocarbamate of the pyrrole derivatives, PDTC can inhibit the activation of NF-κB by hindering the dissociation of IκB from the NF-κB complex, decrease the production of CINC and ENA-78, and prevent the aggregation of PMNs in the lung tissue of ARDS (18,34). The results from the present study suggest that PDTC can attenuate the mRNA expression of p-NF-κB, CINC and ENA-78 by inhibiting the activation of NF-κB, reducing the expression of IL-8 and IL-10, and inhibiting the activation of PMNs in BALF. Pretreatment with PDTC may have partially reduced lethality in LPS-induced mice and attenuated lung tissue
edema, damage, the production of inflammatory cytokines and chemokines, the infiltration of PMNs in the lung and pulmonary capillary permeability. This suggests that the NF-κB signaling pathway may become an important target in regulating ARDS in the lung, and that it may be useful in investigating the pathogenesis of ARDS and exploring more effective and targeted clinical therapies.

In conclusion, in the lung tissue of mice with ARDS induced by LPS, the degradation of NF-κB was activated by phosphorylation and NF-κB p65 migration from the cytoplasm to the nucleus. This was followed by NF-κB activation that commenced the synthesis and release of CINC and ENA-78, and resulted in the imbalanced expression of proinflammatory and anti-inflammatory cytokines, such as IL-8 and IL-10. These cytokines then stimulated the extravasation of PMNs and their direct migration to sites of inflammation, and resulted in a widened lung interval, highly congested pulmonary interstitial, fractured alveolar wall, increased inflammatory cell infiltration, the P(A-a)O₂ and oxygenation obstacles. Therefore, it can be hypothesized that

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Figure 4. Effect of PDTC on p-NF-κB and NF-κB P65 protein expression in lung tissue and the cytoplasm. (A and B) The expression of p-NF-κB was significantly increased in the LPS group (P<0.01) and significantly decreased in the PDTC + LPS group (P<0.05), demonstrating that PDTC can inhibit the activation of NF-κB. (C and D) NF-κB P65 protein expression was reduced in the cytoplasm of the LPS group (P<0.01); the production of P65 was significantly increased in nuclei of the LPS group in comparison with the NS group (P<0.01). (E and F) P65 expression was significantly increased in the cytoplasm (P<0.05), and significantly decreased in the nucleus, of the PDTC + LPS group in comparison with the LPS group (P<0.01). PDTC, pyrrolidine dithiocarbamate; NF-κB, nuclear factor-κB, p-NF-κB, phosphorylated-NF-κB; LPS, lipopolysaccharide; NS, normal saline; h, hours.

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Figure 5. Change of CINC mRNA expression in lung tissue detected by qPCR. (A) CINC expression was detected by RT-PCR. (B) mRNA expression of CINC was detected by qPCR. **P<0.01 vs. NS group; ***P<0.01 vs. L group. CINC, cytokine-induced neutrophil chemoattractant; NS, normal saline; L, lipopolysaccharide; P, pyrrolidine dithiocarbamate; qPCR, quantitative polymerase chain reaction.
PDTC, a specific inhibitor of NF-κB, can reduce the release of chemokines and cytokines via the NF-κB signal pathway, thereby decreasing PMN accumulation in lung tissue, lung tissue damage and improved oxygenation. These results indicate that strategies to regulate the expression of chemokines, cytokines and the NF-κB signal pathway in ARDS should focus on restricting the duration of PMN infiltration and the subsequent effects at inflammatory sites in ARDS.
Figure 8. Expression of epithelial-derived neutrophil-activating protein 78 (ENA-78) in lung tissue, detected using immunohistochemistry (stain, hematoxylin and eosin; magnification, x400). (A) Normal saline (NS), (B) 2 h lipopolysaccharide (LPS), (C) 12 h LPS and (D) 12 h pyrrolidine dithiocarbamate + LPS group. No ENA-78 positive cells were detected in the NS group. ENA-78 expression was increased in the PDTC + LPS group at 12 h, as compared with the NS group, and decreased in comparison with the LPS group.

Figure 9. Expression of IL-8 in (A) serum and (B) BALF, and the expression of IL-10 in (C) serum and (D) BALF. *P<0.05, **P<0.01 vs. the NS group; #P<0.05, ##P<0.01, ###P>0.05 vs. the L group. IL‑8, interleukin‑8; BALF, bronchoalveolar lavage fluid; NS, normal saline; L, lipopolysaccharide; P, pyrrolidine dithiocarbamate.

Figure 10. Number of (A) leukocytes and the proportion of (B) PMNs in the BALF of different groups. *P<0.01 vs. the NS group; #P<0.05, **P<0.01 vs. the L group. PMN, polymorphonuclear; BALF, bronchoalveolar lavage fluid; NS, normal saline; L, lipopolysaccharide; P, pyrrolidine dithiocarbamate; WBC, white blood cell.
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