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## Parthenolide attenuates LPS-induced activation of NF- $\kappa$ B in a time-dependent manner in rat myocardium<sup>☆</sup>

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### Abstract

Parthenolide (PTN), a selective nuclear factor kappa B (NF- $\kappa$ B) inhibitor, has been used extensively to inhibit NF- $\kappa$ B activation. The duration of the inhibitory effect of PTN on NF- $\kappa$ B *in vivo* remains unclear. This study was to determine whether a lipopolysaccharide (LPS) challenge 6, 12 and 24 h after the administration of PTN could activate NF- $\kappa$ B. Rats were divided into five groups. The rats in the PTN, PTN+LPS and DMSO groups were injected intraperitoneally with PTN or DMSO. After 6, 12 or 24 h, LPS was administered in LPS and PTN+LPS groups. The expressions of NF- $\kappa$ B p50, I $\kappa$ B $\alpha$  and p-I $\kappa$ B $\alpha$  were inhibited in both PTN and PTN+LPS group at end of 6 and 12 h and no effects at 24 h. In summary, myocardial NF- $\kappa$ B expression occurs 1 h after the administration of LPS. PTN blocks this effect given at 6 h and no inhibitory effect 24 h after administration *in vivo*.

**Keywords:** parthenolide, nuclear-factor- $\kappa$ B, lipopolysaccharide, myocardium

### INTRODUCTION

Nuclear factor-kappa B (NF- $\kappa$ B) participates in the regulation of multiple immediate early gene products involved in immune, acute phase and inflammatory

responses, in addition to programmed cell death and cellular proliferation<sup>[1]</sup>. At present, NF- $\kappa$ B as a central event leading to the activation of sepsis and septic shock can be activated by a variety of pathogens known to cause septic shock syndrome<sup>[2]</sup>.

Parthenolide (PTN), an NF- $\kappa$ B inhibitor, is responsible for many anti-inflammatory effects and has been used extensively in previous studies<sup>[3-8]</sup>. *In vitro* study has suggested that PTN can inhibit IKK and also directly inactivate NF- $\kappa$ B<sup>[4]</sup>. In addition, other studies have reported that PTN inhibits NF- $\kappa$ B activation in cultured cells and experimental models<sup>[5-8]</sup>. Evidence supports the concept that PTN can inhibit NF- $\kappa$ B activation by selective inhibition of IKK activation and I $\kappa$ B $\alpha$  degradation; however, data regarding effectiveness of PTN duration *in vivo* are currently unavailable.

Treating cells with lipopolysaccharide (LPS) results in the dissociation of cytoplasmic complexes containing NF- $\kappa$ B and the translocation of free NF-

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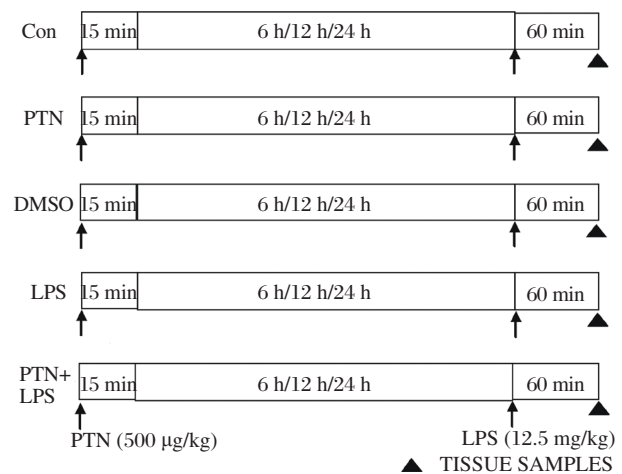
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$\kappa$ B to the nucleus<sup>[9]</sup>. LPS is a primary inducer of inflammatory responses through the biosynthesis and release of a variety of inflammatory mediators via the mononuclear phagocyte system, which is involved in the immune inflammatory response, leading to sepsis and septic shock. NF- $\kappa$ B is a central mediator leading to sepsis and septic shock and can be activated by a variety of pathogens known to cause septic shock syndrome. The role of NF- $\kappa$ B activation in the pathophysiology of sepsis and the signal transduction pathways leading to NF- $\kappa$ B activation during sepsis has been intensively investigated<sup>[2]</sup>. NF- $\kappa$ B activity has been found to be markedly increased in every organ studied to date in both animal models of septic shock and humans with sepsis. The inhibition of NF- $\kappa$ B activation prevents multiple organ injury and improves survival in a rodent model of septic shock. Thus, NF- $\kappa$ B activation plays a central role in the pathophysiology of septic shock<sup>[2,10]</sup>. PTN is a selective NF- $\kappa$ B inhibitor and protects against septic shock syndrome by selective inhibition of IKK activation and I $\kappa$ B $\alpha$  degradation<sup>[8]</sup>. In the present study, we investigated the specific role of NF- $\kappa$ B translocation in rat myocardium after an LPS challenge and determined whether the administration of PTN as a NF- $\kappa$ B inhibitor 6, 12 and 24 h prior could inhibit this effect.

## MATERIALS AND METHODS

### Grouping and experimental protocol

This study was approved by the Animal Care Committee of the Medical College of Soochow University, and all experiments were conducted in accordance with the guidelines for animal care of the National Institutes of Health. A total of 90 adult male Sprague-Dawley rats (10–12 w, 250–350 g) were used in this study. **Fig. 1** illustrates the treatment groups in our study. We randomly divided the rats into 5 groups ( $n = 6$  in each group) at 6, 12 and 24 h after receiving intraperitoneally (i.p.) PTN, respectively: control (CON), PTN, DMSO, LPS and PTN+LPS. The rats in the PTN, PTN+LPS and DMSO groups were injected i.p. with PTN (500  $\mu$ g/kg) or DMSO as indicated, while the rats in the other groups were injected i.p. with an equivalent amount of saline<sup>[11]</sup>. At 6, 12 and 24 h after PTN injection, LPS (12.5 mg/kg) was administered subcutaneously to the LPS and PTN+LPS groups<sup>[12]</sup>, while the rats in the other groups were injected subcutaneously with an equivalent volume of saline. Hearts were collected 1 h after LPS injection. The rats were anesthetized with an i.p. injection of sodium pentobarbital (50 mg/kg) and anticoagulated with an i.p. injection of heparin (1,000 U/kg). Left ventricular samples were quickly excised, cut into five or six cross-sections



**Fig. 1 Schematic illustration of the experimental protocol used in Western immunoblotting experiments.**

Rats were divided into five groups at 6, 12 and 24 h after receiving intraperitoneally PTN, respectively: control CON, PTN, DMSO, LPS and PTN+LPS. LPS was administered subcutaneously at 6, 12 and 24 h after PTN injection. For Western immunoblotting measurements, the tissue samples were obtained 1 h after receiving LPS. PTN: parthenolide; DMSO: dimethylsulfoxide; LPS: lipopolysaccharide; NS: physiological normal saline.

tional slices of 2 mm thickness, snap frozen in liquid nitrogen and stored at  $-80^{\circ}\text{C}$  for subsequent analysis.

### Western blotting analysis of NF- $\kappa$ B p50, I $\kappa$ B $\alpha$ and p-I $\kappa$ B $\alpha$ expression levels

Western immunoblotting was performed as described previously<sup>[13,14]</sup>. Briefly, frozen tissue samples were homogenized using a polytron homogenizer (Beyotime Institute of Biotechnology, Haimen, China) in ice-cold lysis buffer with the complete protease inhibitor phenylmethylsulfonyl fluoride. After centrifugation at 15,000  $g$  for 15 min at  $4^{\circ}\text{C}$ , total protein was isolated. The clarified supernatant was collected to quantify total protein expression. Protein concentrations were determined using an enhanced BCA protein assay kit (Beyotime Institute of Biotechnology). Equivalent amounts (100  $\mu$ g) of protein samples were mixed with Laemmli buffer and heated at  $100^{\circ}\text{C}$  for 5 min, then separated by 5% to 10% sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) and transferred to nitrocellulose membranes (Pall Corporation, East Hills, USA) at 100 V for 1.5 h. The membranes were blocked with 5% nonfat dry milk in TBST (10 mmol/L Tris-HCl, pH 7.5, 150 mmol/L NaCl, 0.05% Tween-20) for 1 h at room temperature and immunoblotted overnight at  $4^{\circ}\text{C}$  with a rabbit polyclonal anti-NF- $\kappa$ B p50 antibody (Santa Cruz Biotechnology, Santa Cruz, USA), a rabbit polyclonal anti-phospho-I $\kappa$ B $\alpha$  antibody (Cell Signaling Co., Danvers, USA) or a mouse monoclonal

anti-phospho-I $\kappa$ B $\alpha$  antibody (Santa Cruz Biotechnology) diluted 1:500 in 5% nonfat dry milk buffer. Membranes were then washed three times in TBST and incubated with a peroxidase-labeled anti-rabbit or anti-mouse IgG secondary antibody diluted 1:5,000 in 5% nonfat dry milk buffer (Cell Signaling Co.) for 1 h at room temperature. Membranes were washed three times in TBST before the antigen-antibody complexes were detected. Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) was used as a reference for quantitative analysis. Specific antigen-antibody complexes were detected using an enhanced chemiluminescence system (ECL, Merck, Darmstadt, Germany). Protein densitometry was performed using Sigma Pro5.0 (Sigma, Saint Louis, USA).

### Statistical analysis

Data were shown as mean  $\pm$  SD. Statistical analyses were performed with SAS 8.1 software (SAS Institute Inc, Cary, USA). Differences between groups were evaluated using two-way analysis of variance, when appropriate, and the post hoc test used was the Newman-Keuls test. All *P* values were two-tailed and a *P* value  $< 0.05$  was considered significant.

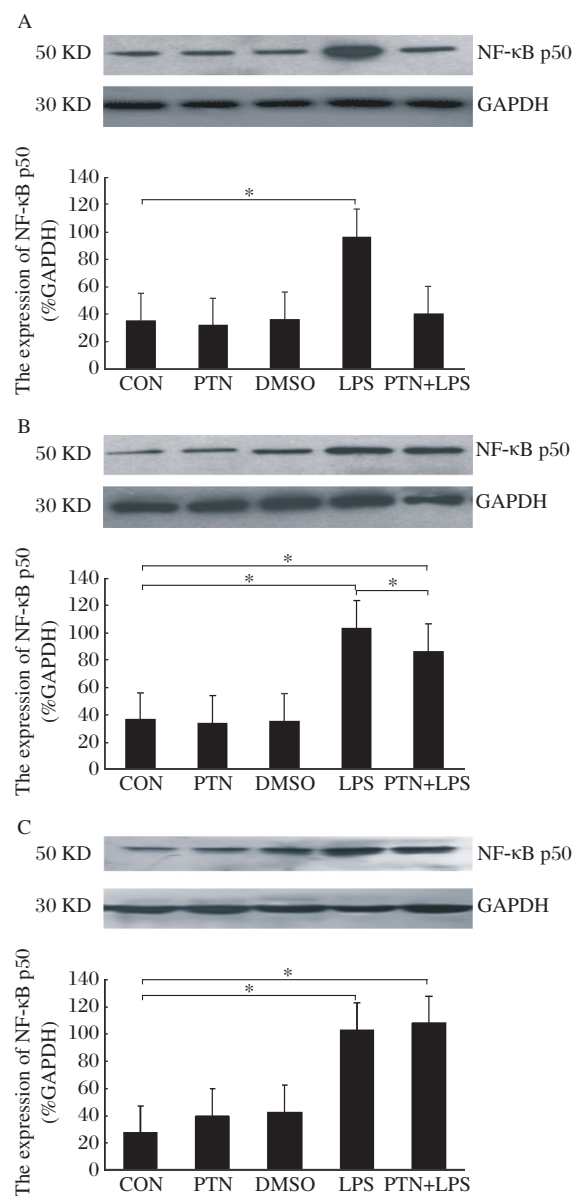
## RESULTS

### Expression of NF- $\kappa$ B p50 protein

The NF- $\kappa$ B p50 protein at 6 h was up-regulated in the LPS group [(97  $\pm$  10)%, *P*  $< 0.05$ ] in comparison with the CON group [(36  $\pm$  11)%], but there was no significant difference in the expression of NF- $\kappa$ B p50 in the PTN, DMSO and PTN+LPS groups [(32  $\pm$  9)%, (37  $\pm$  12)% and (41  $\pm$  13)%, respectively] compared to the CON group (**Fig. 2A**).

The NF- $\kappa$ B p50 protein at 12 h had no significant difference in the PTN and DMSO groups [(34  $\pm$  10)% and (36  $\pm$  19)%, respectively] compared to the CON group [(37  $\pm$  14)%], but the NF- $\kappa$ B p50 protein was up-regulated in the LPS and PTN+LPS groups [(104  $\pm$  4)% and (87  $\pm$  17)% respectively, *P*  $< 0.05$ ] compared to the CON group. However, the up-regulated amplitude of the PTN+LPS group was smaller than that of the LPS group (*P*  $< 0.05$ , **Fig. 2B**).

Western blot analysis showed that the NF- $\kappa$ B p50 protein at 24 h was up-regulated in the LPS and PTN+LPS groups [(103  $\pm$  17)% and (108  $\pm$  20)%, respectively, *P*  $< 0.05$ ] 1 h after subcutaneous injection of LPS compared to the CON group [(27  $\pm$  10)%], but there was no significant difference in the expression of NF- $\kappa$ B p50 protein in the PTN and DMSO groups [(40  $\pm$  15)% and (43  $\pm$  14)%, respectively] compared to the CON group (**Fig. 2C**).



**Fig. 2** Representative effect of PTN on NF- $\kappa$ B p50 in the presence or absence of LPS. A, B and C stand for the expression of NF- $\kappa$ B p50 at 6, 12 and 24 h after receiving intraperitoneally PTN, respectively. Data are shown as the mean  $\pm$  SD, with *n* = 6 in each group. GAPDH: glyceraldehyde-3-phosphate dehydrogenase; PTN: parthenolide; DMSO: dimethylsulfoxide; LPS: lipopolysaccharide.

### Expression of I $\kappa$ B $\alpha$ protein

At 6 h, there were no changes in I $\kappa$ B $\alpha$  expression in the PTN, DMSO and PTN+LPS groups [(68  $\pm$  10)%, (71  $\pm$  12)%, and (65  $\pm$  14)% respectively] compared to the CON group [(69  $\pm$  9)%], but the I $\kappa$ B $\alpha$  protein expression of the LPS group [(39  $\pm$  7)%, *P*  $< 0.05$ ] was down-regulated compared to the CON group (**Fig. 3A**).

At 12 h, the Western blotting analysis showed that the I $\kappa$ B $\alpha$  protein expression in the PTN and DMSO groups had no significant difference [(66  $\pm$  8)% and

(63±10)%, respectively] compared to the CON group [(69±12)%], but in the LPS and PTN+LPS groups IκBα was decreased [(38±11)% and (52±7)%, respectively,  $P < 0.05$ ] compared to the CON group. However, the amplitude in the PTN+LPS group was smaller than that of the LPS group ( $P < 0.05$ , **Fig. 3B**).

At 24 h, Western blotting analysis revealed that there was no significant difference in IκBα expression in the PTN and DMSO groups [(85±9)% and (82±5)%, respectively,  $P > 0.05$ ] compared to the CON group (80±11)% 1 h after injecting LPS subcutaneously. IκBα levels in the cytoplasmic fraction of the heart, however, decreased in the LPS and PTN+LPS groups [(38±5)% and (37±5)%, respectively,  $P < 0.05$ , **Fig. 3C**].

### Expression of p-IκBα protein

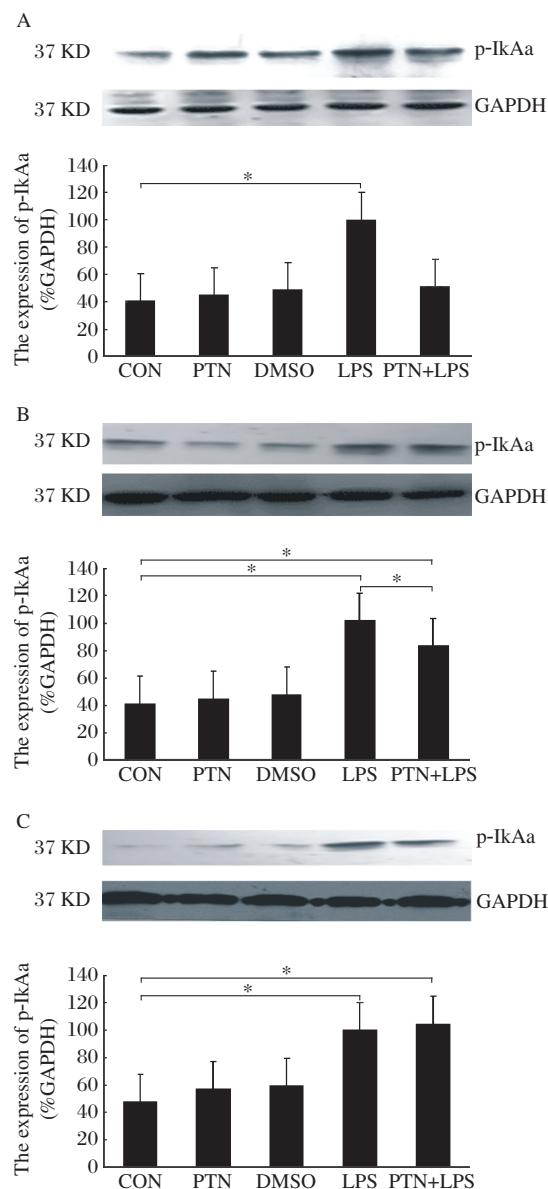
At 6 h, the Western blotting analysis showed that p-IκBα protein was up-regulated in the LPS group [(100±28)%,  $P < 0.05$ ] compared to the CON group [(41±16)%], but there was no evident change of the expression of p-IκBα in the PTN, DMSO and PTN+LPS groups [(45±21)%, (49±24)% and (51±16)%, respectively] compared to the CON group (**Fig. 4A**).

At 12 h, the expression of p-IκBα had no significant difference in the CON, PTN and DMSO groups [(41±15)%, (45±19)% and (48±20)%, respectively], but the p-IκBα protein was up-regulated in the LPS and PTN+LPS groups [(102±16)% and (83±11)%, respectively,  $P < 0.05$ ] compared to the CON group. However, the up-regulated amplitude in the p-IκBα protein of the PTN+LPS group was smaller than that of the LPS group ( $P < 0.05$ , **Fig. 4B**).

At 24 h, p-IκBα expression was also determined by Western blot analysis. Compared to the CON group [(48±23)%], there was no significant difference in the expression of cytoplasmic p-IκBα in the PTN and DMSO groups [(57±23)% and (60±22)%, respectively]. Compared to the CON group [(48±23)%], however, the expression of p-IκBα was significantly increased in the LPS and PTN+LPS groups [(100±35)% and (105±36)%, respectively,  $P < 0.05$ , **Fig. 4C**].

## DISCUSSION

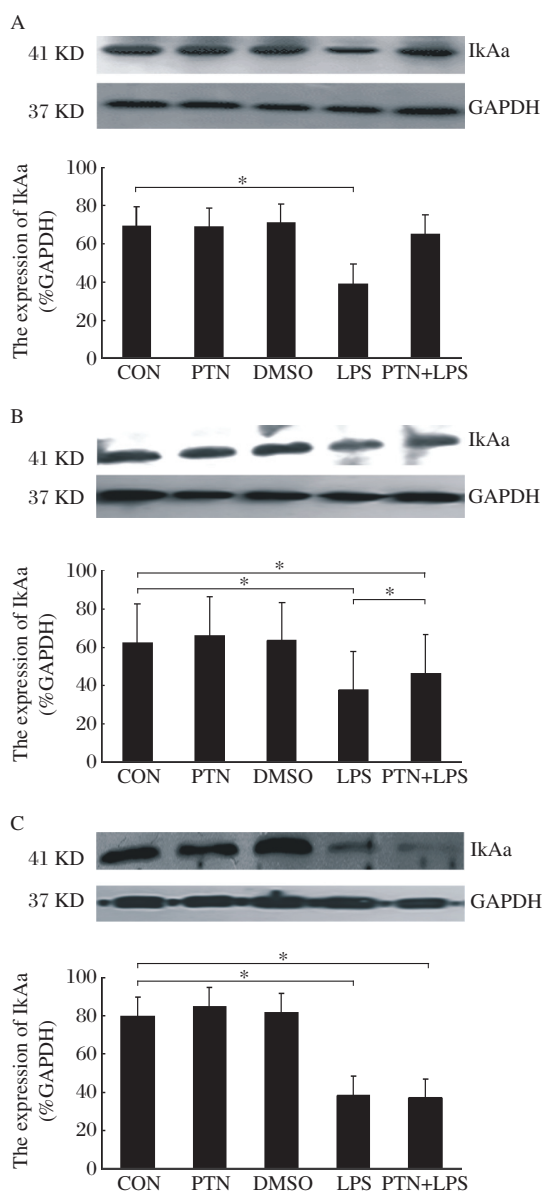
It is well known that NF-κB transcription factors are critical regulators of inflammation, innate and adaptive immunity, and the suppression of apoptosis<sup>[15]</sup>. In resting cells, NF-κB is present in the cytosol as an inactive heterodimer, mainly consisting of the p65/p50 subunits bound to the IκB inhibitor, which is rapidly degraded in response to stimuli such as bacte-



**Fig. 3** The effect of PTN on myocardial IκBα levels in the presence or absence of LPS. A, B and C stand for the expression of IκBα at 6, 12 and 24 h after receiving intraperitoneally PTN, respectively. Data are shown as the mean±SD, with  $n = 6$  in each group. GAPDH: glyceraldehyde-3-phosphate dehydrogenase; PTN: parthenolide; DMSO: dimethylsulfoxide; LPS: lipopolysaccharide.

rial LPS, resulting in NF-κB nuclear entry<sup>[16]</sup>. Therefore, we assessed cardiac NF-κB activation after an LPS challenge and determined whether the administration of PTN, a selective NF-κB inhibitor, 6, 12 and 24 h prior to this challenge could inhibit the effects. Our results indicate that myocardial NF-κB can be activated following LPS administration, but PTN could block this effect completely when given 6 h and partly 12 h prior to the LPS challenge, and did not block NF-κB activation at 24 h. That means that PTN can block





**Fig. 4 Representative effect of PTN on myocardial p-I $\kappa$ B $\alpha$  levels in the presence or absence of LPS.** A, B and C stand for the expression of p-I $\kappa$ B $\alpha$  at 6, 12 and 24 h after receiving intraperitoneally PTN, respectively. Data are shown as the mean  $\pm$  SD, with  $n = 6$  in each group. GAPDH: glyceraldehyde-3-phosphate dehydrogenase, PTN: parthenolide; DMSO: dimethylsulfoxide; LPS: lipopolysaccharide.

this activation when given within 24 h.

The activation of NF- $\kappa$ B is a key factor in excessive inflammatory responses and inflammatory injury. LPS is a primary inducer of inflammatory responses through the biosynthesis and release of a variety of inflammatory mediators, which is involved in the immune inflammatory response, leading to sepsis and septic shock<sup>[17]</sup>. Consistent with the theory that septic shock is a clinical syndrome with diverse etiologies, and NF- $\kappa$ B can be activated by a variety of bacteria, bacterial products and proinflammatory cytokines that

are released during sepsis. NF- $\kappa$ B is the final target of these septic shock inducers<sup>[18]</sup>.

NF- $\kappa$ B can be activated by numerous bacteria, bacterial toxins and proinflammatory mediators known to cause septic shock. LPS-induced activation of NF- $\kappa$ B is biphasic: an early phase occurs at 0.5-2 h post stimulation, and a late phase occurs at 8-12 h post stimulation. LPS and other early inflammatory mediators cause the early phase of NF- $\kappa$ B activation, while TNF and IL-1 $\beta$  mediate the late phase<sup>[19]</sup>. The effects of bacterial toxins on NF- $\kappa$ B activity are dramatic and widespread, leading to massive elevation in NF- $\kappa$ B activity in all organs studied to date<sup>[20-23]</sup>. This is consistent with the idea that multiple organs are involved in septic shock. In the heart, cardiac-specific overexpression of I $\kappa$ B $\alpha$  protects against cardiac injury during trauma<sup>[24]</sup>, and the inhibition of NF- $\kappa$ B activation prevents LPS-induced increase in microvascular permeability<sup>[23]</sup>. Cultured cells or animals developing a tolerance to endotoxin exhibited down-regulated NF- $\kappa$ B activity and reduced expression of NF- $\kappa$ B-dependent genes when subsequently exposed to endotoxin<sup>[25]</sup>. This blunted NF- $\kappa$ B response is believed to be the result of increased I $\kappa$ B $\alpha$ <sup>[26]</sup>, p50 or p52 expression or a switch from the formation of the p50/p65 heterodimer to the p50/p50 homodimer<sup>[27]</sup>. Studies using promoter deletion mutagenesis and reporter gene analysis have demonstrated that NF- $\kappa$ B plays a crucial role in the LPS-activated promoter activity of over 200 genes, many of which play important roles in the development of septic shock<sup>[28-30]</sup>. A major feature of the pathophysiology of septic shock is cardiovascular dysfunction; however, the inhibition of NF- $\kappa$ B activation corrects the cardiovascular functional abnormalities seen in septic shock. Studies have demonstrated that cardiac-specific overexpression of I $\kappa$ B $\alpha$  prevents LPS-induced repression of cardiac systolic and diastolic function<sup>[31]</sup>. Furthermore, inhibition of the NF- $\kappa$ B pathway can ameliorate the vascular derangement in both LPS and polymicrobial models of septic shock<sup>[32,33]</sup>.

The results of an endotoxemia study of jejunal mucosa induced by LPS suggest that activated NF- $\kappa$ B consists primarily of p50 subunits. As an inhibitor of NF- $\kappa$ B, I $\kappa$ B has a variety of forms, mainly consisting of I $\kappa$ B $\alpha$  and I $\kappa$ B $\beta$ . The main function of I $\kappa$ B $\alpha$  is to regulate and control the activation of NF- $\kappa$ B. I $\kappa$ B $\alpha$  demonstrates rapid degradation and resynthesis in response to stimuli such as LPS and TNF- $\alpha$  and is responsible for the acute-phase activation of NF- $\kappa$ B. In contrast, I $\kappa$ B $\beta$  demonstrates delayed degradation and resynthesis and is responsible for the persistence of NF- $\kappa$ B activation<sup>[12]</sup>. This process requires the phos-

phorylation of I $\kappa$ B by the IKK complex and the activation of NF- $\kappa$ B in response to proinflammatory stimuli and pathogen-associated molecular patterns<sup>[15,16]</sup>. In this study, therefore, we determined whether the model of septic shock was successful by measuring the changes of NF- $\kappa$ B p50 and I $\kappa$ B $\alpha$  expression 1 h after the administration of LPS. Our results show that the majority of NF- $\kappa$ B p50 is activated 1 h after the administration of LPS, inducing corresponding changes in the expression of I $\kappa$ B $\alpha$  and p-I $\kappa$ B $\alpha$ .

PTN has attracted considerable interest, and PTN-containing products are commonly used to treat a variety of inflammatory conditions such as migraine, rheumatoid arthritis and asthma<sup>[34]</sup>. Several studies have reported that PTN inhibits NF- $\kappa$ B activation in cultured cells and experimental models<sup>[5-8,35]</sup>. Furthermore, *in vitro* experiments with LPS-stimulated vascular smooth muscle cells and monocytes have shown that PTN at noncytotoxic doses inhibits I $\kappa$ B $\alpha$  degradation, preventing NF- $\kappa$ B activation and the subsequent gene expression. PTN also has beneficial effects during myocardial ischemia, endotoxic shock and renal disease through NF- $\kappa$ B inhibition<sup>[8]</sup>. Studies have found that PTN prevents NF- $\kappa$ B activation and dependent gene expression in both human and murine cells<sup>[36]</sup>, and evidence indicates that PTN may inhibit the phosphorylation and degradation of I $\kappa$ B $\alpha$ <sup>[8]</sup>, directly affect the DNA-binding ability of NF- $\kappa$ B without changing I $\kappa$ B $\alpha$  degradation or block a signaling pathway other than NF- $\kappa$ B. Both the *in vitro* and *in vivo* anti-inflammatory effects of PTN have been associated with the inhibition of I $\kappa$ B $\alpha$  depletion, which in turn results in the inhibition of excessive activation of NF- $\kappa$ B<sup>[37]</sup>. Some studies have reported that PTN protects against myocardial ischemia and reperfusion injury in rats by selective inhibition of IKK activation and I $\kappa$ B $\alpha$  degradation<sup>[8]</sup>. In this study, the expression levels of NF- $\kappa$ B, I $\kappa$ B $\alpha$  and p-I $\kappa$ B $\alpha$  in the LPS group at 6 h were different from those in the PTN+LPS group, which means that PTN blocked the activation of NF- $\kappa$ B induced by LPS within 6 h. The expression levels of NF- $\kappa$ B, I $\kappa$ B $\alpha$  and p-I $\kappa$ B $\alpha$  in the LPS and PTN+LPS groups had a similar trend at 12 h, but amplitude was different between them, showing that PTN has a partial effect in blocking NF- $\kappa$ B activation. Inversely, the expression levels of NF- $\kappa$ B, I $\kappa$ B $\alpha$  and p-I $\kappa$ B $\alpha$  in the PTN+LPS group were similar to those in the LPS group and different from those in the PTN group when given at 24 h prior to LPS administration, which means that PTN did not block the activation of NF- $\kappa$ B induced by LPS at 24 h. From what has been discussed above, it would be reasonable to believe that PTN blocked the activation of NF- $\kappa$ B induced by LPS

within 6 and 12 h, not at 24 h.

Our study has several potential limitations. We did not specifically investigate different doses of PTN in the pharmacological inhibition of NF- $\kappa$ B (500  $\mu$ g/kg) for the second window of protection (SWOP)<sup>[8]</sup>. In addition, we focused on a single organ, the heart, at only a 1-h interval of LPS challenge. In future studies, we will design and execute a comprehensive approach to accomplish our original goal of determining the duration of PTN following *in vivo* administration.

In conclusion, myocardial NF- $\kappa$ B expression occurs 1 h after the administration of LPS, and PTN, a selective inhibitor of NF- $\kappa$ B, could block this effect completely when given at 6 h, and partly block the activity 12 h prior to the LPS challenge. There was no block after 24 h. This further indicates that PTN has no inhibitory effect 24 h after administration.

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