Ligustilide modulates oxidative stress, apoptosis, and immunity to avoid pathological damages in bleomycin induced pulmonary fibrosis rats via inactivating TLR4/MyD88/NF-KB P65

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Background: Pulmonary fibrosis (PF) is a fatal disease with increasing incidence. Ligustilide (LIG) has been shown to inhibit oxidative stress, apoptosis, and inflammation. Here we investigated the possible effect of LIG on bleomycin-induced PF in Sprague-Dawley rats.

Methods: PF rats were set up through a single endotracheal injection of bleomycin (5 mg/kg). Then rats were treated with 20, 40, and 80 mg/kg LIG for four weeks, and the effects were estimated.

Results: Overall, LIG significantly improved ventilation and reduced hyperplasia, and treatment of LIG reduced fibrosis as indicated by Masson staining and reduced expression of transforming growth factor-beta (TGF-β), Fibronectin, and alpha-smooth muscle actin (α-SMA). Oxidative stress was induced with bleomycin while inhibited with LIG, as showed with rebalanced serum lactate dehydrogenase (LDH), and tissue superoxide dismutase (SOD), glutathione peroxidase (GSH) and malondialdehyde (MDA). Apoptosis was further inhibited with LIG, as shown with Terminal dUTP nick-end labeling (TUNEL) staining and expression of Caspase-3, Caspase-9, Bax, and Bcl-2. Th1/Th2 balance was also rebuilt as evaluated with CD4 and IFNγ/IL-4 labeled flow cytometry of peripheral blood mononuclear cells (PBMCs) and expression of inducible nitric oxide synthase (iNOS) and IL-10 in the serum and lung. Protein expression of Toll-like receptor 4 (TLR4), HSP60-TLR4-myeloid differentiation factor 88 (Myd88) and nuclear factor-kappa B (NF-κB) p-P65/P65 was significantly reduced with LIG treatment. All the effects of LIG exhibited in a dose-dependent way.

Conclusions: LIG improved bleomycin-induced PF with improved ventilation, reduced fibroblast, reduced oxidative stress and apoptosis, and rebalanced Th1/Th2 immunity, through TLR4/MyD88/NF-κB P65 signaling.

Keywords: bleomycin; idiopathic pulmonary fibrosis (IPF); ligustilide (LIG); rat

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Introduction

Pulmonary fibrosis (PF), mainly idiopathic pulmonary fibrosis (IPF) is the most common interstitial lung diseases and is a progressive, irreversible, and usually fatal disease (1). With aging and environmental pollution appearing, the incidence of PF increased. However, as a result of limited early diagnosis and intervention, the average life expectancy of IPF patients is 3–5 years after diagnosis (2). As multiple driven forces promoted PF with different characters, PF or IPF are not correct concepts, and pathogenesis of PF...
possibly could be far more than fibrosis. Generally, in PF, aberrantly activated secretion of lung epithelial cells leads to fibroblast proliferation and differentiation into highly active myofibroblasts, which deposit excessive amounts of extracellular matrix (ECM) and irreversibly destroy the lung architecture (3).

Oxidative stress and apoptosis are potential therapeutic targets of PF. Simply elicitation of oxidative stress is sufficient to induce PF in Atp8b1 mutant mice (4). Bleomycin (bleomycin) induced lung injury is the most commonly used model of PF (5). Oxidative stress promoted apoptosis was the target of several potential therapeutics of PF (6). Reduction of oxidative stress, for example, with Sulforaphane significantly reduced PF in C57/BL6 mice (7), while enhancing of oxidative stress promoted toxicity of bleomycin (8). However, the exact influence of oxidative stress and apoptosis in PF patients still needs to be characterized.

Immunne responses are essential players in the progression of PF. Dysregulated release of fibroblast cells in PF lung induces a series of pro-inflammatory immune responses, which in turn promotes the progression of PF (9). The overall inflammatory response in PF closely resembles a Th2 immune response, while lung fibroblasts tend to induce a Th1 immune response (9). Th1/Th2 balance could have essential affects in the regulation of inflammation and apoptosis, thus determining cell fate and progression of PF. Currently, the balance of Th1/Th2 was considered to be regulated with NF-κB, Nrf-2, and MyD88 (7,10,11).

Ligustilide (LIG) is the main lipophilic component of the Umbelliferae family in Traditional Chinese Medicine, including Radix angelicae sinensis and Ligusticum chuanxiong (12). LIG has been directly and indirectly shown to inhibit fibrosis in multiple ways. The antioxidant and anti-apoptotic properties of LIG were characterized as neuroprotective (13). In lung tissues, LIG promotes apoptosis of cancer-associated fibroblasts via TLR4 (14) and inhibits pro-inflammatory damage-associated molecular patterns (15). In fibrosis of other organs, LIG inhibition of fibrosis, apoptosis, and inflammation was reported to be related to TLR4, NF-κB, and MyD88 (16,17). Sprague-Dawley (SD) rats were used in the present study show the possible role of LIG in lung fibrosis by referring to previous researches (18). This research highlighted the possible role of LIG in bleomycin-induced fibrosis of rats.

We present the following article in accordance with the ARRIVE reporting checklist (available at http://dx.doi.org/10.21037/atm-20-4233).

Methods

Animals and treatments

All animal experiments complied with the guidelines developed by the Health Department of Local Government and Experimentation Ethics Committee of the Universities and approved by the Department of Health of the Hong Kong Government and the Animal Ethics Committee of The Chinese University of Hong Kong (Ref: 13/085/GRF-5). A total of 30 SD rats (adult, male, 180–220 g) were bought from Shanghai SIPPR-Bk Laboratory Animal Co. Ltd. The rats were raised at 25 °C with 12-hour light/dark cycles with relative humidity between 40–70%, fed with standard pellet chow, and had free access to food and water. After acclimated for a week, rats were randomly assigned to control, bleomycin (marked as bleomycin), bleomycin plus 20 mg/kg of LIG (marked as LIG 20 mg/kg), Bleomycin plus 40 mg/kg of LIG (marked as LIG 40 mg/kg), and Bleomycin plus 80 mg/kg of LIG (marked as LIG 80 mg/kg), groups (n=6 per group, 2 rats per cage). PF was established through a single endotracheal injection of bleomycin (5 mg/kg; Zhejiang Hisun Pharmaceutical Co., Ltd., China) in all groups except the control (5,18). After 24 hours, control and bleomycin groups were treated with 2 mL/kg/day of saline and other groups with 20, 40, and 80 mg/kg/day LIG (Sichuan Weikeqi Biotechnology Co., Ltd., China) treatments through intraperitoneal injection for four weeks. Then Lung functions were measured, and rats were sacrificed, and tissues, serum, and blood were collected.

Lung function measurements and sample collection

After treated for four weeks, Lung function measurements were carried out as described (19). Briefly, a whole-body flow plethysmograph (Buxco Electronics Inc., Wilmington, NC, USA) was used to measure minute ventilation volume directly, and then airway resistance was measured with a FinePointe-series all-in-one Resistance/Compliance (RC) system (Buxco Electronics). After calibration of the equipment with airflow, each SD rat was placed in a small plexiglass container within the plethysmograph chamber and was measured over 5 minutes at room temperature. Then rats were anesthetized with ketamine/xylazine,
intubated with a sterile cannula that was held in place with surgical thread and then connected to the ventilator with a fixed breathing rate of 145 bpm. Airway resistance of each rat was recorded over 3 minutes using Biosystem XA. After measurements, blood was withdrawn by a retro-orbital puncture. Then peripheral blood mononuclear cells (PBMCs) were collected from peritoneum with PBS and cultured in DMEM (HyClone, USA) (supplemented with 10% FBS (PAN, Germany) and 1% penicillin/streptomycin) at 37 °C and 5% CO₂. Finally, the anesthetized rat was euthanized following an IACUC approved protocol after the experiment. The blood was centrifuged at 500 g for 15 minutes, and sera were collected. Left lungs (about 200 mg) in each group were collected for making tissue homogenate, and right lungs in each group were fixed for making paraffin sections. Samples were temporarily stored at −80 °C before the test.

**Histology evaluation**

The diced lung tissues were marked, then fixed in 4% Paraformaldehyde Fix Solution (Beyotime Biotechnology Co., Ltd., China) at 4 °C overnight. Tissues were then dehydrated in a series of ethanol and xylene solutions and embedded in paraffin. Sections of 4-μm thickness were cut then deparaffinized. Hematoxylin and eosin (H&E Staining Kit, Beyotime Biotechnology Co., Ltd., China) staining and Masson staining were performed according to protocol. Then the slides were observed in 200× scope.

**Test of serum lactate dehydrogenase (LDH) and tissue superoxide dismutase (SOD), glutathione peroxidase (GSH) and malondialdehyde (MDA)**

The determinations of LDH, SOD, GSH, and MDA in serum or tissues were conducted to evaluate the changes in the oxidant-antioxidant systems. The levels of serum LDH were measured using collected serum and related assay kits (Jiancheng Institute of Biotechnology, Nanjing, China). Lung tissues were sonicated in a 50 mM phosphate buffer, and the supernatant was gained after centrifuged at 3,000 g for 10 minutes. The supernatant was used to test SOD (tested with SOD assay kit, Jiancheng Institute of Biotechnology, Nanjing, China), GSH (tested with GSH-PX assay kit, Jiancheng Institute of Biotechnology, Nanjing, China) and MDA (tested with MDA assay kit, Jiancheng Institute of Biotechnology, Nanjing, China) according to instructions.

**Terminal dUTP nick-end labeling (TUNEL)**

The paraffin section of lung tissue was taken to receive conventional dewaxing, antigen repairing, and blocking. Sections were incubated in the working solution (20 µg/mL proteinase K dissolving in 10 mmol/L Tris-HCl, PH: 7.4–8; Sigma-Aldrich) at 37 °C for 15 minutes. Then, 50 µL TUNEL reaction solutions (Roche Applied Science, Indianapolis, Inc., IN, USA) were added for incubation at 37 °C for 1 hour. Next, the sections were washed with PBST, incubated with 100 µL diaminobenzidine (DAB) for 30 minutes in converter-peroxidase (POD) (50 µL, at 37 °C) and wash with PBST. Finally, the slides were observed in 200x scope, four fields of each tissue were randomly harvested, and the average apoptotic cells per 100 cells were analyzed with Image J.

**Western blotting**

Tissues were then homogenized in RIPA lysis buffer (Beyotime Biotechnology Co., Ltd., China), tested with a BCA kit (Beyotime Biotechnology Co., Ltd., China) and balanced with 1X PBS. Then about 30 µg of protein for each sample was added with a loading buffer, separated on an SDS-PAGE gel, transferred to a PVDF membrane, and incubated with primary and secondary antibodies. The antibodies used were: anti-Bax (182733), anti-Bcl-2 (196495), anti-Caspase-3 (179517), anti-Caspase-9 (ab184786), anti-MyD88 (ab2064), anti-P65 (ab32536) and anti-p-P65 (ab86299) purchased from Abcam; anti-TGF-beta (#3709), anti-Fibronection (#4705), anti-alpha-SMA (#68463), anti-TLR4 (#14358), anti-beta-Actin (#12748) and all secondary antibodies bought from Cell Signaling Technology.

**Flow cytometry**

Flow cytometry was applied to determine CD4+IFN-γ⁺ and CD4+IL-4⁺ lymphocytes balance. The collected PBMCs were cultured in a 5% CO₂ incubator at 37 °C for 24 hours so that unattached cells were discarded. Then cells were collected in an EP tube. FITC-anti-CD4 was incubated at room temperature in the dark for 30 minutes. Then the cells were washed with cell staining buffer, mixed with 500 µL fixation/permeabilization solution (BD, 555028), and incubated in the dark for 45 minutes. Next, 1X BD perm/wash buffer washed cells were incubated with PE-anti-IL-4 and APC-anti-IFN-γ in the dark for 45 minutes, and washed again with 1X BD perm/wash buffer, and resuspended
with 300 μL flow washing liquid. Th1 cells (defined as CD4^+IFN-γ^+) and Th2 cells (defined as CD4^+IL-4^+) were detected by flow cytometry (FACSAria; BD Biosciences), and data were analyzed using CellQuest software.

**Enzyme-linked immunosorbent assay (ELISA)**

Tissue samples were prepared and balanced as mentioned in western blotting, serum samples were from centrifuged blood. Elisa kits (Cloud-Clone Corp., Wuhan, China) were used to measure the changes in iNOS and IL-10 in rat serum and lung tissue in each group following the instruction. Standard preparation (50 μL) was added in the first and second wells, followed by multiple proportion dilution. Sample diluent (40 μL) and sample (10 μL) were added in each well at 37 °C for 30 minutes. Washing liquid was then used to wash the wells for five times, and a secondary antibody (50 μL) was added in each well at 37 °C for 30 min. After washed out again, chromogenic agents A and B (50 μL for each) were added in each well at 37 °C in the dark for 15 minutes. A termination solution (50 μL) was added in each well. A microplate reader (Bio-Rad, USA) was used for detection. Sample concentrations were calculated according to the curve of the standard.

**Statistical analysis**

All the data were analyzed by Image J and Graphpad Prism 7.0 and presented as the mean ± standard error of the mean (SEM). Differences were determined with P<0.05 of the t-test for at least three independent experiments.

**Results**

**LIG improved lung function of bleomycin treated rats**

Bleomycin treated rats were administered with 20, 40, and 80 mg/kg of LIG, respectively, to estimate the overall effects of LIG on PF. HE stains of lung tissues indicated bleomycin-induced rats developed lung injury compared to normal individuals, while LIG dose-dependently reduced hyperplasia (Figure 1A). Similar results were seen in minute pulmonary ventilation (Figure 1B) and airway resistance (Figure 1C), as bleomycin significantly reduced minute ventilation and airway resistance, and this was dose-dependently inhibited with LIG (*P<0.05 versus control and #P<0.05 versus bleomycin). These revealed that the inhibited lung function with bleomycin could be improved with LIG.

**LIG reduced fibrosis of bleomycin treated rats**

PF of the rats was showed with Masson staining and related expression of TGF-β, Fibronectin, and α-SMA. As expected, bleomycin-induced fibrosis, obviously, and treatment of LIG alleviated the burden of fibrosis (Figure 2A). Similarly, TGF-β, Fibronectin, and α-SMA all were induced with bleomycin while dose-dependently reduced with LIG (Figure 2B, *P<0.05 versus control and #P<0.05 versus bleomycin). These results revealed that LIG
could reduce bleomycin-induced fibrosis of SD rats.

**LIG reduced the oxidative stress of bleomycin treated rats**

The relative content of LDH (Figure 3A) in the serum and SOD (Figure 3B), MDA (Figure 3C) and GSH (Figure 3D) in the tissue were then determined to show the involvement of oxidative stress in LIG improved lung function. Serum LDH and lung MDA were significantly induced with bleomycin, which was dose-dependently reduced with LIG (*P<0.05 versus control and ^P<0.05 versus bleomycin). The activity of SOD and GSH were significantly inhibited with bleomycin and dose-dependently recovered with LIG (*P<0.05 versus control and ^P<0.05 versus bleomycin). The reduced LDH and MDA, together with recovered SOD and GSH, showed that LIG dose-dependently reduced lung oxidative stress-induced with bleomycin in rats.

**LIG reduced apoptosis of bleomycin treated rats**

Apoptosis in rat lung was then evaluated with TUNEL staining and expression of apoptosis-related proteins. As indicated in Figure 4A, a single dose of bleomycin markedly induced apoptosis of the rat lung in day 28, and treatment LIG significantly reduced apoptotic cells (*P<0.05 versus control and ^P<0.05 versus bleomycin). The bleomycin-induced and LIG reduced apoptosis was further confirmed with western blotting. The ratio of Bax/Bcl-2, cleaved Caspase-3/Caspase-3 and cleaved Caspase-9/Caspase-9 were all elevated with bleomycin while dose-dependently reduced with LIG (Figure 4B, *P<0.05 versus control and ^P<0.05 versus bleomycin).

**LIG rebuilt Th1/Th2 balance of bleomycin treated rats**

Th1/Th2 Balance was further evaluated with CD4 and IFNγ/IL-4 labeled flow cytometry of PBMCs. CD4^IFNγ^ cells were markedly induced with bleomycin and dose-dependently reduced with LIG (Figure 5A, B, *P<0.05 versus control and ^P<0.05 versus bleomycin). CD4^IL4^ cells were not significantly changed with bleomycin, while 40 and 80 mg/kg of LIG promoted the amount of CD4^IL4^ cells (Figure 5A, C, *P<0.05 versus bleomycin). The bleomycin increased, and LIG inhibited Th1 immunity, and the LIG increased Th2 immunity, which was further validated with an expression of iNOS and IL-1 in the blood (Figure 5D).
and lung tissue. Bleomycin obviously induced iNOS both in the serum and lung tissue, and LIG dose-dependently reduced iNOS; bleomycin had no significant effects on IL-10, and LIG dose-dependently reduced IL-10 both in the serum and lung tissue. These showed the rebalancing of Th1/Th2 with LIG.

**LIG reduced fibrosis of bleomycin treated rats through TLR4/MyD88/P65**

The involvement of TLR4 (Figure 6A,B), MyD88 (Figure 6A,C), and NF-κB P65 (Figure 6A,D) in LIG reduced fibrosis of bleomycin was then investigated with western blotting. The relative expression of TLR-4, MyD88, and the ratio of p-NF-κB P65/ NF-κB P65 were all markedly induced with bleomycin, and dose-dependently decreased with LIG (*P<0.05 versus control and ^P<0.05 versus bleomycin). Results above show that TLR4/MyD88/P65 was involved in LIG reduced fibrosis of bleomycin in SD rats.

**Discussion**

Currently, PF remains an irreversible and deadly disease, in which acute exacerbations remain a significant cause of morbidity and mortality (2). Inhibition of fibrosis and maintenance of ventilation was the main issue in treating PF, including IPF. In this study, we showed that LIG dose-dependently reduced the progression of fibrosis and improved ventilation. The results supported LIG as a potential prodrug for PF.

Oxidative stress, pro-inflammatory cytokines, and apoptosis were accumulated, and under PF, inhibition of oxidative stress effectively reduced bleomycin toxicity and PF (7,20). Accumulated oxidative stress would induce mitochondrial dysfunction and apoptosis, which further promoted PF (6). Activation of inflammatory responses is also a key process in PF; for example, the NLRP3 inflammasome is reported to be activated to the promoted progression of bleomycin-induced PF (21), and targeting of NLRP3 inflammasome effectively reduced PF (22). The activation of oxidative stress, inflammation, and apoptosis

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**Figure 3** Reduced oxidative stress with LIG in bleomycin treated rat lung. Rats were pretreated with bleomycin once and treated with an indicated dosage of LIG for 28 days and (A) LDH activity in serum, (B) SOD activity in the lung, (C) MDA activity in the lung, and (D) GSH activity in the lung were tested with related kits. n=6 for each group, ^P<0.05 versus control and *P<0.05 versus bleomycin with t-test. Data were shown as mean ± SEM. LIG, ligustilide; LDH, lactate dehydrogenase; SOD, superoxide dismutase; MDA, malondialdehyde; GSH, glutathione peroxidase.
Reduced apoptosis with LIG in bleomycin treated rat lung. Rats were pretreated with bleomycin once and treated with an indicated dosage of LIG for 28 days and (A) TUNEL staining of the lung tissue observed under 200× scope and average apoptotic cells (Yellow-stained) per 100 cells counted. (B) Apoptosis related proteins Bax, Bcl-2, Caspase-3, and Caspase-9, were evaluated with western blotting, and the relative expression of proteins was evaluated with Image J software. n=6 for each group, *P<0.05 versus control and #P<0.05 versus bleomycin with t-test. Data were shown as mean ± SEM. LIG, ligustilide.

could mutually interact to promote PF, and co-targeting of these processes, especially targeting of TLR4, could be a potentially effective way of therapy (21). LIG was shown to inhibited oxidative stress in multiple ways, including Nrf-2 and ferroportin (23,24). Current studies highlighted the anti-inflammation effects of LIG, primarily through TLR4 and NF-κB (25). In this study, we showed that the anti-oxidative, anti-apoptotic and anti-inflammation effects of LIG also exist in bleomycin-induced PF. As expected, TLR4, MyD88, and NF-κB P65 were inactivated with LIG dose-dependently.

Inflammatory immune response was activated in PF, in which the Th2-type immune response was overactivated, while lung fibroblasts possibly tend to induce a Th-type immune response (9). Regulation of Th1/Th2 immunity was an essential target of treating PF. For example, Valsartan showed a significant protective effect against bleomycin-induced PF, possible through NF-κB expression and regulation of Th1/Th2 cytokines (10). During the treatment of LIG, the Th1-type immune response was inhibited, and the Th2-type immune response was activated. These results were possible as a result of decreased fibrosis, as indicated with Masson staining.

The pathogenesis of PF remained mostly unclear, and
Figure 5 Rebalanced Th1/Th2 with LIG in bleomycin treated rat lung. Rats were pretreated with bleomycin once and treated with an indicated dosage of LIG for 28 days. (A) CD4+IFNγ+ and CD4+IL-4+ PBMCs were evaluated with flow cytometry and (B) CD4+IFNγ+ and (C) CD4+IL-4+ cells were shown in bar graphs. Expression of iNOS and IL-10 in serum (D) and Lung tissue (E) were evaluated with ELISA. n=3 for each group for flow cytometry and n=6 for Elisa, *P<0.05 versus control and #P<0.05 versus bleomycin with t-test. Data were shown as mean ± SEM. LIG, ligustilide.
the treatment of PF is still limited. Current research and this paper indicated the benefits of TLR4/MyD88/NF-κB P65 targeting in PF. However, there may be another mechanism of action for LIG in treating PF, such as referring myofibroblast differentiation, matrix deposition, fibroblast proliferation, senescence-associated secretory phenotype, or regulation of microRNAs (26). So the effects and mechanisms characterized here could merely result in form ameliorated fibrosis but not the reason by which fibrosis ameliorated.

In conclusion, LIG ameliorated bleomycin-induced PF with improved ventilation, reduced fibroblast, reduced oxidative stress and apoptosis, and rebalanced Th1/Th2 immunity, possibly through TLR4/MyD88/NF-κB P65 signaling.

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Footnote

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Figure 6 Expression of TLR4, MyD88, and P65 in LIG and bleomycin treated rat lung. Rats were pretreated with bleomycin once and treated with an indicated dosage of LIG for 28 days. (A) Protein expression of TLR-4, MyD88, P65, and p-P65 were tested with western blotting and relative expression levels of TLR4 (B), MyD88 (C), and the ratio of p-P65 to P65 (D) was evaluated with Image J software. n=6 for each group, *P<0.05 versus control and #P<0.05 versus bleomycin with t-test. Data were shown as mean ± SEM. LIG, ligustilide.
The authors have no conflicts of interest to declare.

**Ethical Statement:** The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. All animal experiments complied with the guidelines developed by the Health Department of Local Government and Experimentation Ethics Committee of the Universities and approved by the Department of Health of the Hong Kong Government and the Animal Ethics Committee of The Chinese University of Hong Kong (Ref: 13/085/GRF-5).

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