Abstract. Acute lung injury (ALI) is a severe health issue with significant morbidity and mortality. Artemisinin is used for the treatment of fever and malaria in clinical practice. Dihydroartemisinin (dHA), the major active metabolite of artemisinin, plays a role in anti-organizational fibrosis and anti-neuronal cell death. However, whether dHA can attenuate ALI remains unclear. The current study thus examined the effects of dHA on ALI and primary macrophages. The results revealed that dHA attenuated lipopolysaccharide (LPS)-induced pulmonary pathological damage. dHA suppressed the LPS-induced infiltration of inflammatory cells, the elevation of myeloperoxidase activity, oxidative stress and the production of pro-inflammatory cytokines, including interleukin (IL)-1β, tumor necrosis factor-α, and IL-6. Furthermore, dHA reduced the LPS-induced inflammatory response by suppressing the degradation of I-κB and the nuclear translocation of nuclear factor-κ-light-chain-enhancer of activated B cells (NF-κB)/p65 in vivo and in vitro. DHA activated the nuclear factor-erythroid 2 related factor 2 (Nrf2) pathway, which was suppressed by LPS treatment. The Nrf2 inhibitor, ML385, diminished the protective effects of dHA against LPS-induced inflammation in macrophages. On the whole, the findings of this study demonstrate that DHA exhibits anti-inflammatory activities and may be a therapeutic candidate for the treatment of ALI.

Introduction

Acute lung injury (ALI) is a leading cause of acute respiratory failure (1). ALI is characterized by extreme inflammation, the release of pro-inflammatory cytokines, excessive neutrophil infiltration and lung endothelial/epithelial cell injury, resulting in edema and gas exchange deterioration (2). However, the clinical mortality rate of the severe form of ALI, acute respiratory distress syndrome (ARDS), remains >40.0% (1). Therefore, more effective therapeutic strategies for ARDS are urgently required.

Macrophages, the principal immune cells in the lungs, produce inflammatory molecules and carry out vital functions in the molecular mechanisms of ALI, such as boosting neutrophil infiltration and triggering inflammatory reactions (3). Neutrophils trigger the release of pro-inflammatory cytokines, such as tumor necrosis factor (TNF)-α, interleukin (IL)-1β and IL-6 (4). These pro-inflammatory cytokines induce the production of oxidants, which are associated with the activation of nuclear factor κ-light-chain-enhancer of activated B cells (NF-κB), eventually contributing to ALI (5,6). Accordingly, oxidative stress is increased in lipopolysaccharide (LPS)-induced ALI (7). The transcription factor, nuclear factor-erythroid 2 related factor 2 (Nrf2), plays a critical role in protection against ALI by inducing the expression of antioxidant and detoxifying enzymes and proteins (8). For example, it has been reported that Nrf2 attenuates ALI and inflammation by suppressing Toll-like receptor (TLR)4 and Akt signaling (9).

Artemisinin is isolated from Artemisia annua, a Chinese traditional medicinal herb. Artesunate is a water-soluble hemisuccinate derivative of artemisinin. Studies have reported that artemesunate inhibits ischemia-reperfusion-induced lung inflammation and LPS-induced ALI (10,11). Dihydroartemisinin (DHA), the major active metabolite of artemisinin or artemesunate, is an effective and widely distributed anti-malarial drug with good absorption (12,13). DHA is more stable and ten times more effective than artesunate (14). Recent studies have demonstrated that DHA not only exerts an anti-malarial activity but also shows anti-inflammatory effects in vivo and in vitro. For example, DHA attenuates lipopolysaccharide (LPS)-induced acute lung injury in mice by suppressing NF-κB signaling in an Nrf2-dependent manner.
effect, but also exerts anticancer, anti-organizational fibrosis and anti-neuronal cell death effects (15-17). However, whether DHA can attenuate ALI and affect NF-κB signaling activation in macrophages remains unclear.

The present study thus hypothesized that DHA may attenuate LPS-induced ALI and evaluated the effects of DHA on LPS-treated macrophages to elucidate the mechanisms through which DHA attenuates LPS-induced ALI.

Materials and methods

Ethics statement. The Ethics Committee of the Center for Scientific Research with Animal Models at Central South University (Changsha, China) approved the experiments, which were performed in accordance with the guidelines of the National Institutes of Health. Mice were anesthetized with pentobarbital sodium (80 mg/kg, intraperitoneal injection), and all necessary efforts were taken to minimize suffering prior to the experiments.

Animal experiments. Male adult C57bl/6 mice were kept in climate-controlled quarters with a 12-h light/dark cycle and a relative humidity of 40-60%, and were provided with food and water ad libitum at a temperature of 25°C. The mice were housed for 1 week for environmental adaptation prior to experimentation. The mice were randomly divided into 4 groups (weight, 20-25 g; age, 8 weeks; n=24 in each group) as follows: i) The control group; ii) ALI group; iii) DHA group; and iv) ALI + DHA group. The ALI model was induced by the intratracheal injection of LPS (E. coli O111:B4; Sigma-Aldrich; Merck KGaA) in 50 µl saline as described in our previous studies (18,19). Mice in the control group received saline. Mice in the DHA group received DHA (75 mg/kg; Sigma-Aldrich; Merck KGaA) only. Mice in the LPS + DHA group were treated with DHA (75 mg/kg) via intragastric administration 1 h prior to LPS administration. A total of 12 h following the LPS administration, the mice were anesthetized by an intraperitoneal (i.p.) injection of sodium pentobarbital (80 mg/kg) and blood was collected for further analysis. Following the collection of lung tissue, the mice were then sacrificed by an i.p. injection of 200 mg/kg sodium pentobarbital.

Survival experiment. For survival analysis, another 80 mice were randomly divided into 4 groups as follows: i) The control group; ii) ALI group; iii) DHA group; and iv) ALI + DHA group (n=20 in each group). The mice were treated with LPS at a lethal dose (25 mg/kg, intratracheal) (20-23). Mice in the DHA group received DHA (75 mg/kg; Sigma-Aldrich; Merck KGaA) only. Mice in the LPS + DHA group were treated with DHA (75 mg/kg) via intragastric administration 1 h prior to LPS administration. A total of 12 h following the LPS administration, the mice were anesthetized by an intraperitoneal (i.p.) injection of sodium pentobarbital (80 mg/kg) and blood was collected for further analysis. Following the collection of lung tissue, the mice were then sacrificed by an i.p. injection of 200 mg/kg sodium pentobarbital.

Histological analysis. Lung tissue was excised and immersed in 4% paraformaldehyde for 24 h at 4°C. Paraffin-embedded lung tissue was sectioned at a thickness of 4 µm and was then stained with haematoxylin and eosin (cat. no. G1120; Solarbio) for 5 min at room temperature for pathological analysis. According to a previous study, the severity of inflammation was graded between 0 and 4 as follows: 1, <25% lung involvement; 2, 25-49% lung involvement; 3, 50-75% lung involvement; and 4, >75% lung involvement (24). Lung injury was scored by 3 pathologists blinded to the treatments.

Bronchoalveolar lavage fluid (BALF) acquisition and analysis. BALF was collected by lavaging the lungs with 1 ml PBS 3 times and was centrifuged at 800 x g for 5 min at 4°C. Total cells, macrophages and neutrophils were counted with a hemocytometer following Wright-Giemsa staining for 10 min at room temperature. The cell-free supernatant of BALF was harvested, and total protein content was determined using a bicinchoninic acid protein assay kit (Thermo Fisher Scientific, Inc.).

Lung wet-to-dry weight ratio. The lungs were excised from the mice and blood was removed by blotting the tissue with filter papers until dry. After weighing (wet weight), the lungs were placed in an incubator at 60°C for 48 h and then weighed again (dry weight). The wet-to-dry ratio of the lungs was calculated to reflect edema.

Lactate dehydrogenase (LDH) activity assay. LDH activity in BALF was determined with an LDH Cytotoxicity assay kit (cat. no. A020; Nanjing Jiancheng Bioengineering Institute) according to the manufacturer's protocol. The absorbance at 490 nm was measured using a microplate reader (Thermo Fisher Scientific, Inc.).

Measurement of myeloperoxidase (MPO), malondialdehyde (MDA), superoxide dismutase (SOD) and glutathione (GSH) levels. The MPO, MDA, SOD and GSH levels were detected using the related kits (MPO: cat. no. A044; MDA: cat. no. A003; SOD: cat. no. A001; GSH: cat. no. A005; Nanjing Jiancheng Bioengineering Institute). All procedures were performed according to the manufacturer's protocols.

Primary peritoneal macrophages. Male C57bl/6 mice (8-week-old, n=20) were used to extract primary peritoneal macrophages. At 3 days following the intraperitoneal injection of 3 ml 3% thioglycolate (Sigma-Aldrich; Merck KGaA), the animals were euthanized by CO₂ inhalation (flow rate of CO₂, 20%) and peritoneal macrophages were harvested by peritoneal lavage with cooled RPMI-1640 (Gibco; Thermo Fisher Scientific, Inc.). Cells were collected by centrifugation (room temperature, 1,200 g, 10 min) and resuspended with culture medium. Cells were plated into 6- or 12-well plates (1x10⁶ cells/well) and the culture medium was discarded 2 h later. After being allowed to rest overnight, the cells were treated with DHA (20 µM) for 1 h followed by treatment with LPS (100 ng/ml) for a further 6 h at 37°C. To investigate the effects of ML385 (an Nrf2 inhibitor; MedChemExpress) on macrophages, primary macrophages were pre-treated with ML385 (20 µM) for 30 min, and the cells were then treated with DHA (20 µM) for 1 h followed by treatment with LPS (100 ng/ml) for a further 6 h.

Immunofluorescence. Primary peritoneal macrophages were plated on polylysine-coated coverslips. Following fixing with 4% paraformaldehyde at 4°C and permeabilization with Triton X-100, cells were incubated with antibodies against NF-κB p65, and the expression of NF-κB p65 was detected using a fluorescence microscopy.
X-100, macrophages were incubated overnight at 4°C with anti-CD68 antibody (cat. no. ab125212; 1:50; Abcam) and anti-F4/80 antibody (cat. no. ab6640; 1:50; Abcam), FITC (cat. no. ab6717; 1:200; Abcam) and Cy3 (cat. no. ab6953; 1:200; Abcam)-conjugated secondary antibody were then applied for 1 h at room temperature. Following nuclear staining with DAPI (cat. no. C0065; Solarbio), the macrophages were viewed under a fluorescence microscope (Thermo Fisher Scientific, Inc.).

**Cytokine measurements.** The levels of TNF-α, IL-1β and IL-6 in mouse sera and the culture supernatants of primary peritoneal macrophages were detected using enzyme-linked immunosorbent assay (ELISA) kits (TNF-α, cat. no. BMS607-3; IL-1β, cat. no. BMS6002; IL-6, cat. no. BMS603-2; Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions.

**Total RNA extraction and reverse transcription-quantitative polymerase chain reaction.** Total RNA was extracted from the lung tissue or macrophages using RNeasy reagent (cat. no. 5301100; Takara Bio, Inc.). Total RNA (1 μg) was used for the synthesis of cDNA using a PrimeScript RT Reagent kit with gDNA Eraser (cat. no. RR047A; Takara Bio, Inc.). qPCRs were run using SYBR-Green Real-time PCR Master mix (Thermo Fisher Scientific, Inc.) on a Bio-Rad real-time PCR system (CFX96 Touch™; Bio-Rad Laboratories, Inc.). The amplification conditions were as follows: Pre-degeneration at 95°C for 10 min, then 40 cycles of denaturing at 95°C for 15 sec, annealing at 60°C for 30 sec and extension at 72°C for 30 sec, and a final extension at 72°C for 5 min. Relative fold expression levels were normalized to GAPDH and calculated using the 2^-ΔΔcq method (25). The primer sequences were as follows: TNF-α forward, ACAGCAAGGACTAGCA GGAG and reverse, GGAGTGCTCTTCTGCGCAT; IL-1β forward, GGCGCTCAAAGGAAAGATC and reverse, TAC CAGTTGGGAAACTCTGC; IL-6 forward, CTGGGGGAT GTCTTGCAGCTC and reverse, CTGTGAGTCTCTCCTCC GG; and β-actin forward, TCTTGTGACGCTCTTCGTT and reverse, TCTTCTGACCCATTCACC.

**Western blot analysis.** The lung tissues or cell samples were lysed in RIPA buffer (cat. no. P0013K; Beyotime) at 4°C for 30 min, and the total proteins were quantified using a BCA kit (cat. no. P0010; Beyotime). Briefly, 30 μg proteins were separated by 12% SDS-PAGE gels and transferred onto polyvinylidene difluoride membranes (EM Millipore). The membranes were blocked with 5% fat-free milk for 2 h and then probed at 4°C overnight with the following primary antibodies: Anti-phosphorylated(p)-IκB (1:1,000; cat. no. 2859; Cell Signaling Technology, Inc.), anti-IκB (1:1,000; cat. no. 4812; Cell Signaling Technology, Inc.), anti-Nrf2 (1:1,000; cat. no. YTH3189; ImmunoWay), anti-heme oxygenase 1 (HO-1; 1:1,000; cat. no. 43966; Cell Signaling Technology, Inc.), anti-p65 (1:1,000; cat. no. 10745; Proteintech), anti-β-actin (1:1,000; cat. no. 4970; Cell Signaling Technology, Inc.) and anti-p-p65 (1:1,000; cat. no. 3033; Cell Signaling Technology, Inc.). After washing with TBST 3 times, the membranes were incubated with secondary antibodies (1:7,500; cat. no. ab6721; Abcam) at room temperature for 1 h. Enhanced chemiluminescence (EMD Millipore) was used to detect the protein content. Images were obtained using a ChemiDoc XRS system (Bio-Rad Laboratories, Inc.).

**Measurement of ROS production.** The lung tissues were homogenized and stained with 50 μM of DCFH-DA (cat. no. S0033; Beyotime) at 37°C in the dark for 30 min. DCF fluorescence intensities were detected by a multi-detection reader (Thermo Fisher Scientific, Inc.) at an excitation and emission wavelength of 485 and 535 nm.

**Statistical analysis.** Data were analyzed using SPSS 19.0 software (SPSS, Inc.). All data are expressed as the means ± standard error of the mean. Means were compared by two-way ANOVA followed by Tukey’s post-hoc test to assess significance. Survival analysis was carried out using the Kaplan-Meier log-rank test. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**DHA attenuates lung tissue injury in mice with LPS-induced ALI.** The lung tissues of the control mice were with intact alveoli, while the lung tissues from the mice with ALI exhibited obvious pulmonary edema, alveolar disarray and inflammatory cell infiltration in the alveolar cavity. Following treatment with DHA, the pathological changes induced by LPS in the lungs were attenuated (Fig. 1A). DHA treatment also significantly reduced the lung injury score and LDH activity in the BALF of mice with ALI (Fig. 1B and C). LPS significantly increased the lung wet-to-dry weight ratio and the total protein content in BALF, and these were reduced by DHA treatment (Fig. 1D and E), indicating that DHA attenuated LPS-induced edema. In addition, it was identified that DHA treatment significantly decreased the numbers of total cells, macrophages and neutrophils in the BALF of mice with LPS-induced ALI (Fig. 1F-H). It was also revealed that DHA treatment significantly reduced MPO activity in the lungs of mice with LPS-induced ALI (Fig. 1I). These results indicate that DHA attenuates the lung injury induced by LPS in mice.

**DHA reduces the inflammatory response and oxidative stress in the lungs of LPS-exposed mice.** IL-1β, TNF-α and IL-6 are pro-inflammatory cytokines that are critical to the development of ALI (26). The results of the present study demonstrated that mice with ALI exhibited a significant increase in the mRNA levels of IL-1β, TNF-α and IL-6 in the lungs following LPS administration; however, these effects were significantly suppressed by DHA treatment (Fig. 2A-C). Accordingly, DHA reduced the protein levels of IL-1β, TNF-α and IL-6 in the serum of mice with LPS-induced ALI (Fig. 2D-F). LPS induced a significant increase in ROS generation and MDA content in the lungs, which was significantly suppressed by DHA treatment (Fig. 2G and H). By contrast, SOD activity and GSH content were decreased in the ALI group and were partially restored by treatment with DHA (Fig. 2I and J). These results indicate that DHA protects against LPS-induced lung injury by reducing inflammation and oxidative stress.
DHA inhibits inflammatory cytokine release and oxidative stress induced by LPS in primary macrophages. Treatment with DHA (20 µM) suppressed the increase in the IL-1β, TNF-α and IL-6 mRNA expression levels in primary macrophages and protein levels in the culture supernatant, which were induced by LPS (100 ng/ml) (Fig. 3A-G). Furthermore,
ROS generation and MDA levels were increased in the macrophages treated with LPS, whereas these responses were significantly suppressed by DHA (Fig. 3H and I). In addition, DHA treatment partially restored the decreased SOD activity and GSH content in macrophages exposed to LPS (Fig. 3J and K). Collectively, these findings suggest that DHA reduces the LPS-induced inflammatory response and oxidative stress in primary macrophages.

Inhibition of the NF-κB pathway by DHA is dependent on Nrf2 in primary macrophages. DHA inhibited I-κB degradation and reduced the increase in p-65 expression induced by LPS in the lung tissue of mice and primary macrophages (Fig. 3J and K). Collectively, these findings suggest that DHA reduces the LPS-induced inflammatory response and oxidative stress in primary macrophages.

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Therapeutic effect of DHA on LPS-induced ALI mice. Clinically, pharmacotherapy does not commence until there is an approved diagnosis of ALI. In this study, the therapeutic effects of DHA on mice with ALI were subsequently examined. It was identified that treatment with DHA 2 h post-exposure to LPS significantly improved the survival rate of mice with
LPS-induced ALI (Fig. 5). This result suggests a therapeutic effect of DHA against ALI in mice.

Discussion

Artemisinin and its derivative, artesunate, have been reported to exert protective effects against lung inflammation (10,11), which indicates the potential use of artemisinin and its derivative in therapy. As a more effective agent than artesunate, DHA has been reported to exert anti-inflammatory and anti-fibrotic effects against bleomycin-induced pulmonary fibrosis in rats (27). The present study first reported that DHA attenuated lung tissue injury in a murine model of LPS-induced ALI and suppressed macrophage activation induced by LPS. Notably, it was identified that treatment with DHA 2 h post-exposure to LPS significantly improved the survival rate of LPS-exposed mice, indicating a therapeutic effect of DHA against ALI. Mechanistically, it was determined that the DHA-mediated suppression of inflammatory injury was dependent on Nrf2. The present findings suggest the use of DHA as a potential therapeutic agent for patients with ALI in the future.
Nrf2 plays a critical role in the regulation of oxidative stress, which is the key pathogenic mechanism of ALI (28-30). Generally, Nrf2 exists as a complex with Keap1 in the cytoplasm. When cells are sensitized to ROS, Nrf2 is released from the complex and translocates to the nucleus, promoting the expression of antioxidants, such as HO-1.
to upregulate Nrf2 and HO-1 expression in lung tissue, which indicated that dHA inhibits the NF-κB pathway by activating Nrf2. Additionally, inhibition of Nrf2 abolished the protective effects of DHA, indicating a role of Nrf2 in the therapeutic effects of DHA. The Nrf2/ARE signaling pathway also regulates the expression of anti-inflammatory genes and inhibits the progression of inflammation (33). Nrf2 negatively regulates LPS-induced NF-κB/p65 translocation induced by DHA in macrophages, which indicates that DHA inhibits the NF-κB pathway in a Nrf2-dependent manner in macrophages. The NF-κB pathway is a key target for the development of anti-inflammatory agents (34). Numerous natural products have been screened for anti-inflammatory activities by inhibiting NF-κB (35-37). Artemisinin significantly inhibits NF-κB activation by suppressing the phosphorylation and degradation of IκBα and p65 nuclear translocation (38). Artesunate has been reported to suppress LPS-induced TLR4 expression and NF-κB activation in lung tissue and to upregulate Nrf2 and HO-1 expression in lung tissue in vivo (11). However, whether DHA can affect the activation of NF-κB and the underlying mechanisms in macrophages remain unclear. The present study first reported that DHA significantly mitigated NF-κB pathway activation in the lungs of ALI mice and in primary macrophages exposed to LPS. It has also been reported that DHA inhibits the NF-κB pathway in rat chondrocytes (39) and tumor cell invasion (40). While the exact mechanisms remain unclear, the present study provides a novel mechanism through which DHA inhibits the NF-κB pathway by activating Nrf2. This indicates that DHA is a potential anti-inflammatory and anti-oxidative agent.

Macrophages are the principal immune cells of inflammatory molecules in pulmonary tissue and exert a vital function in the molecular mechanisms of ALI, triggering inflammation reactions and boosting the infiltration of neutrophils (3). There is increasing evidence to suggest that macrophages, which act as the first line of defense in the lungs, are key factors in the pathogenesis of ALI (41). The depletion of macrophages has been found to mitigate lung injury significantly at 4 h following the administration LPS in mice by attenuating neutrophilic alveolitis and reducing pro-inflammatory cytokines (42). Under the LPS challenge, the pro-inflammatory M1 alveolar macrophages are mainly derived from the bone marrow. Those alveolar macrophages are the triggers of the uncontrolled inflammatory response during ALI. However, it is hard to harvest a sufficient amount of alveolar macrophages from healthy mice to conduct an experiment. In this study, the primary peritoneal macrophages were recruited to the peritoneal cavity by 3% thioglycolate. Thus, these macrophages are also derived from bone marrow. Notably, the adoptive transfer of peritoneal macrophages into the lungs results in the expression of certain alveolar macrophage-specific genes (43). In some studies, primary peritoneal macrophages are used to investigate the role of macrophages in lungs (19,44-46). The present study focused on the role of DHA in the LPS-challenged inflammatory response in macrophages in vitro. It was found that DHA inhibited inflammatory cytokine release and oxidative stress induced by LPS in primary peritoneal macrophages. Collectively, we hypothesized that primary murine peritoneal macrophages share, at least partly, the response to LPS-challenge with alveolar macrophages.

The limitations of the present were the following: First, only the protective effects of DHA against LPS-induced ALI in mice were examined. To further clarify the effects of DHA on ALI, the protective effects of DHA in other models of ALI should also be investigated. Second, the mechanisms underlying the suppression of ALI by DHA are not completely clear. In addition to the NF-κB pathway, the effects of DHA on the activation of the NLRP3 inflammasome should be examined, which is a vital mechanism underlying the uncontrolled inflammation during ALI (47). Artesunate has been identified to alleviate renal ischemia-reperfusion-induced lung inflammation by attenuating the activation of NLRP3 inflammasome (10).

In conclusion, this study demonstrates that DHA exerts protective and therapeutic effects against LPS-induced ALI by inhibiting the NF-κB signaling pathway in a Nrf2-dependent manner. The present findings provide further evidence that DHA may be a valuable therapeutic candidate for use in the treatment of ALI.

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Availability of data and materials
The data used to support the findings of this study are presented in the present study or are available from the corresponding author upon request.

Authors’ contributions
XTH designed and performed most of the experiments, analyzed and interpreted the data, and wrote the manuscript. WL, CXH, YaZ, CYZ, and CCS assisted during the acquisition, analysis, and interpretation of data and revised the manuscript. ZQL and YoZ assisted with data acquisition and revision of the manuscript. SYT performed experiments and prepared the manuscript. All authors have reviewed and approved the final version of the manuscript.

Ethics approval and consent to participate
The Ethics Committee of the Center for Scientific Research with Animal Models at Central South University (Changsha, China) approved the experiments, which were performed in accordance with the guidelines of the National Institutes of Health.

Patient consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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