

# Transplantation of Viable Allogeneic Mitochondria Improves Gas Exchange and Alveolar-Capillary Permeability in Rats with Endotoxin-Induced Acute Lung Injury

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## Research Article

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## Abstract

**Background:** Acute lung injuries (ALI) cause disruption of the alveolar-capillary barrier and remains as the leading cause of death in critically ill patients. This study tested the hypothesis that the administration of freshly isolated viable allogeneic mitochondria can prevent injury of the alveolar-capillary barrier at the endothelial level, as mitochondrial dysfunction of the pulmonary endothelium is a critical aspect of ALI progression.

**Methods:** ALI was induced by intratracheal lipopolysaccharide instillation (LPS, 1mg/kg) in anesthetized rats. Mitochondria (100 µg) were isolated from the freshly harvested soleus muscles of naïve rats and stained with a green fluorescence MitoTracker™ dye. A mitochondria or placebo solution was randomly administered into the jugular veins of the rats at 2 h and 4 h after ALI induction. An arterial blood gas analysis was done 20 h later. The animals were then sacrificed, and lung tissues were harvested for analysis.

**Results:** An IVIS Spectrum imaging system was used to obtain *ex vivo* heart-lung block images and demonstrated the enhancement of MitoTracker™ fluorescence in the lungs. Mitochondria transplantation significantly improved arterial oxygen contents ( $\text{PaO}_2$  and  $\text{SaO}_2$ ), while reduced  $\text{CO}_2$  tension in rats with ALI. Animals with mitochondrial transplants had significantly increased ATP concentration in lung tissues. Allogeneic mitochondria transplantation preserved alveolar-capillary barrier function, as shown by a reduction in protein levels in the bronchoalveolar lavage fluid, and decrease of extravasated Evans blue dye and hemoglobin content in lung tissues. Relaxation responses to acetylcholine and eNOS expression were potentiated in injured pulmonary arteries, and inflammatory cells infiltration into lung tissue was also reduced following mitochondrial transplantation.

**Conclusions:** Transplantation of viable mitochondria protects the integrity of endothelial lining of the alveolar-capillary barrier, thereby improving gas exchange during the acute stages of endotoxin-induced ALI. However, the longer-term effects of mitochondrial transplantation on pulmonary function recovery after ALI requires further investigation.

## Introduction

Acute lung injury (ALI) and acute respiratory distress syndrome (ARDS) represent the most common causes of intensive care unit (ICU) admission, need for ventilatory support, and death in critically ill patients with a high mortality rate of up to 35-45%<sup>1</sup>. In severe ALI, intense pro-inflammatory responses and leukocyte infiltration into lung tissue destroy the integrity of the alveolar epithelial and pulmonary artery endothelial layers, leading to the breakdown of the alveolar-epithelial barrier. This in turn, promotes the formation of protein-rich pulmonary edema and impairs blood-gas exchange<sup>2-4</sup>. Since pulmonary vascular endothelium is the key modulator of ALI progression, pharmacologic- and cell-based endothelial-targeting therapies have become emerging candidate therapies for ALI/ARDS in the past two decades<sup>5, 6</sup>.

Under normal conditions, the pulmonary vascular endothelium inhibits the majority of inflammatory and coagulation responses in the lung; but in ALI, these endothelial cells are activated and contribute to the pro-inflammatory and pro-coagulant responses<sup>7</sup>. Mitochondria serve as the most important sensors in regulating intra- and inter-cellular communications for the maintenance of healthy cellular homeostasis. In order to maintain homeostasis, mitochondria are also extremely sensitive to environmental stressors<sup>8</sup>. When the excessive physical stress is generated and the cellular environment becomes toxic or allostatic, defects in mitochondrial electron transport, intracellular Ca<sup>2+</sup> homeostasis, ion transport, mitochondrial biogenesis, cellular metabolism, redox regulation, and mitochondrial quality control result in mitochondrial dysfunction<sup>8, 9</sup>. Recent evidence suggests that impaired mitochondrial homeostasis in pulmonary epithelial and vascular endothelial cells contributes to the progress of ALI<sup>10, 11</sup>, and the loss of mitochondrial genomic integrity is the crucial initial step that drives lung injury propagation during ALI<sup>12</sup>. Since viable exogenous mitochondria transplantation has been shown therapeutic potential in various pre-clinical ischemia-reperfusion injury models<sup>13–15</sup> and in pediatric patients with myocardial dysfunction<sup>16</sup>, we conducted the first pre-clinical study that tested the hypothesis that the administration of freshly-isolated viable allogeneic mitochondria can reduce endothelial injury at the alveolar-capillary barrier and improve gas exchange in an experimental model of endotoxin-induced ALI.

## Methods

### Rat model of acute lung injury

All experimental procedure were performed in accordance with the ARRIVE Guideline and the guidelines for animal research issued by the E-Da Hospital and I-Shou University. The study was approved by the Institutional Animal Care and Use Committee of the E-Da Hospital, Kaohsiung, Taiwan (IACUC approval number 109024). Adult male Sprague-Dawley rats were anesthetized using intraperitoneal Zoletil cocktail anesthetic (30 mg/kg) injections. Following mini-open incision, a placebo solution (0.9% saline, 0.5 ml) or Escherichia coli lipopolysaccharide (LPS, 1 mg/kg in 0.5 ml of saline; Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) was instilled through glottis with a catheter for sham operations and ALI induction, respectively<sup>17</sup>.

### Isolation and administration of mitochondria

Mitochondria were freshly isolated from health naïve rats within 1h before delivery. Rats were asphyxiated with CO<sub>2</sub> and their bilateral soleus muscles were harvested. A commercially available mitochondria isolation kit (Thermo Scientific, Waltham, MA) was used to isolate viable mitochondria<sup>18</sup>. The freshly isolated soleus muscles were briefly washed with a phosphate buffer solution (PBS, containing 0.1 M sodium phosphate and 0.15 M sodium chloride at pH 7.2) then incubated with a trypsin solution (0.3 mg/ml) for 3 minutes. The muscles were homogenized using a grinder and the resulting tissue samples were centrifuged at 11,752 g for 3 minutes at 4°C. After three cycles of

suspension and centrifugation, the mitochondrial pellets were collected, and the cytosolic components were discarded using mitochondria isolation reagents (A to C solutions).

## **Allocation of Experimental Groups and Study Protocol**

Animals were randomly allocated to sham (0.9% saline) or ALI group (LPS). Rats in the ALI group were then randomly re-assigned to either the control or mitochondrial group. The control group received intravenous PBS (200 µL) and the mitochondria group received skeletal muscle-derived mitochondria in PBS (100 µg in 200 µL PBS). Solutions in both groups were delivered via the jugular veins. Each animal in the control or mitochondrial group received the treatment solution at 2 h and 4 h after intratracheal instillation of LPS and they recovered from anesthesia under a warm blanket. A total of 11 rats received the sham operation, 23 rats received LPS treatment (including 10 controls and 13 mitochondrial transplantation), and 6 rats were served as naïve animals for tissue harvesting and *ex vivo* image scanning. The experimental animals were sacrificed under deep anesthesia 24 h after instillation of saline or LPS to collect blood samples and lung tissues for analysis.

## **Identification of the Distribution of Exogenous Mitochondria following systemic delivery**

In some experiments, the isolated mitochondria were stained with a green MitoTracker™ fluorescence dye (catalog #M7514, Invitrogen™, Carlsbad, CA) for the *ex vivo* imaging under an *in vivo* imaging system (IVIS, Caliper IVIS Spectrum System, PerkinElmer, Waltham, MA). Heart and lungs were harvested *en bloc* at 30 min after intravenous delivery of MitoTracker fluorescence-labelled mitochondria and scanned under the 3-dimensional tomography imaging system.

## **Artery blood gas analysis (ABG)**

Arterial blood was drawn from the femoral artery of anesthetized rats under room air condition. Oxygen saturation ( $\text{SaO}_2$ ), partial pressure of oxygen ( $\text{PaO}_2$ ) and  $\text{CO}_2$  ( $\text{PaCO}_2$ ), and other ABG parameters in the arterial blood were analyzed using a blood gas analyzer (ABL 520, Radiometer, Copenhagen, Denmark).

## **Measurement of vascular reactivity**

Pulmonary artery rings (approximately 2 mm long) were freshly isolated and mounted in organ chambers containing 25 ml of Krebs solution. The chambers were maintained at 37°C and aerated continuously with 94%  $\text{O}_2$ /6%  $\text{CO}_2$ . Changes in isometric force were recorded continuously using an isometric force-displacement transducer (Grass IT30; Grass Instrument, West Warwick, RI). Each ring was gradually stretched to 2.0 g. After a 45-min equilibration period, the rings were induced to contract using KCl (40 mM) and slow phenylephrine titration (PE,  $10^{-9}$  to  $10^{-5}$  M). Concentration-response curves were charted by recording acetylcholine ( $10^{-9}$  to  $10^{-5}$  M; Sigma-Aldrich, Burlington, MA) during contraction to a median PE effective concentration (EC50).

## **Assessment of pulmonary capillary permeability and lung wet-to-dry ratio**

The permeability of alveolar-capillary barrier was assessed by measuring protein content in the bronchoalveolar lavage fluid (BALF), the Evans blue assay and lung wet-to-dry ratio (LWDR)<sup>19</sup>. The trachea in the lab animals were dissected and lungs were lavaged with 6 ml/kg chilled PBS. The recovered lavage fluid was analyzed for total protein content using a protein assay kit (Bio-Rad Laboratories, Hercules, CA). Evans blue dye (2% v/v, 20 mg/kg, Sigma Aldrich) was administered via the jugular vein at 30 min before euthanization and lung tissue was perfused free of blood with normal saline. Evans blue was extracted from the freshly isolated lung tissues by incubating in 99.5% formamide (4 ml/g tissue; Sigma Aldrich) for 48h and the tissue concentrations were determined by a spectrophotometry at the wavelength of 620 nm. To assess lung water content, the left lung was excised and weighed immediately. Lung tissues were dried in an oven at 60°C for 24 h and reweighed. LWDR was obtained by dividing the mass of the initial specimen by the mass of the dried specimen.

## **Measurement of lung hemoglobin content and myeloperoxidase activity assay**

The hemoglobin colorimetric assay was performed using a hemoglobin colorimetric assay kit (catalog #700540, Cayman Chemical, Ann Arbor, MI). Lungs were excised and homogenized in 5 ml PBS solutions (pH 7.4, with 0.16 mg/ml heparin, per gram weight of tissue). Homogenized lung tissues were centrifuged at 10,000 g for 15 minutes at 4°C. After the supernatant was removed, the tissue samples were added to assay wells and incubated at room temperature for 15 minutes. The absorbance at 560-590 nm was read and analyzed. The enzymatic activity of myeloperoxidase (MPO) in the homogenized lungs was measured by a commercially available MPO assay kit (Cell Biolabs Inc., San Diego, CA).

## **Measurement of Tissue adenosine triphosphate (ATP)**

Tissue ATP concentrations were analyzed using an ATP assay kit (catalog #ab83355, Abcam, Cambridge, MA). The right lungs of lab animals were excised and homogenized in the ATP assay buffer. The homogenized tissue was centrifuged at 13,000 g for 5 minutes at 4°C. Insoluble components were removed and the solution was deproteinized using a sample preparation kit. The supernatant was then collected for ATP colorimetric assay.

## **Western Blot**

Pulmonary artery trunk and lung tissues were minced and homogenized in lysis buffer. Equal amount of proteins (100 mg) were loaded into polyacrylamide gels and transferred to nitrocellulose membranes using the wet transferring method. The membranes were incubated overnight at 4°C with primary antibodies (endothelial nitric oxide synthase (eNOS), phosphorylated eNOS-S1177, CD11b and iNOS) in 1:1000 dilutions. All primary antibodies were purchased from BD Biosciences (New Jersey, USA), except CD11b was obtained from Abcam (Cambridge, MA). After washing with PBS, the membranes were incubated with horseradish peroxidase-linked secondary antibodies for 1 hour at room temperature. Bands were visualized using enhanced chemiluminescence and quantified with scanning densitometry (the ImageJ; 1.48v, National Institutes of Health, Bethesda, Md).

# Histologic and immunostaining examinations

Lung tissues were immersed in 10% buffered formal saline and fixed for 24 h. Biopsies were then processed through increasing grades of alcohol and embedded in paraffin wax. Hematoxylin and eosin (H&E)-stained lung sections were examined under a light microscope and photographed. Lung injury was assessed by an investigator blinded to the treatment groups according to a scale of 0-2 for each of the following criteria: (1) infiltration of neutrophils in the alveolar and interstitial space, (2) amount of hyaline membranes and proteinaceous debris, (3) degree of lung hemorrhage and (4) extent of parenchymal consolidation<sup>20</sup>.

## Statistical analysis

All data sets were tested for the normality assumption using the Shapiro-Wilk test before statistical procedures. The values of continuous variables were compared using the Mann-Whitney U test was used. Two-way repeated measures (RM) ANOVA was used to compare differences in vasomotor reactivity between the control and mitochondrial transplantation groups at the different concentrations of vasoactive drugs. A Dunn's post hoc procedure was used for multiple comparisons. Results are presented as the median and interquartile range (IQR). Statistical significance was accepted at a level of  $P < 0.05$ . All of the statistical analyses were performed using the SigmaPlot 14.0 (Systat Software Inc., San Jose, CA).

## Results

### Uptake of allogenic mitochondria following intravenous administration

Compared with naïve animals, MitoTracker™ Green fluorescence densities were more significantly enhanced in the *ex vivo* lung lobes of rats with ALI that received skeletal muscle-derived allogeneic mitochondria transplantation (Figure 1A), indicating that exogenous mitochondria in the pulmonary circulation are retained following intravenous administration. Tissue ATP concentration in the lung homogenates of animals with ALI was restored following allogeneic mitochondrial transplantation (Figure 1B).

### Transplantation of mitochondria improves gas exchange and preserves function of alveolar-capillary barrier

Intratracheal LPS instillation successfully impaired pulmonary gas exchange, as shown by the significant reduction of  $\text{PaO}_2$  ( $95.9 \pm 5.0$  mmHg vs  $78.9 \pm 3.8$  mmHg, sham vs ALI, respectively;  $P < 0.001$ ) and increase in  $\text{PaCO}_2$  ( $51.6 \pm 3.7$  mmHg vs  $61.1 \pm 3.8$  mmHg, sham vs ALI, respectively;  $P < 0.001$ ) (Table 1). Level of  $\text{HCO}_3^-$  was increased in ALI group, but there was no difference in arterial pH (Table 1). Following two doses of mitochondrial transplants, arterial  $\text{O}_2$  and  $\text{CO}_2$  partial pressures were improved in rats with ALI (Table 1). Saturation of oxygen content ( $\text{SaO}_2$ ) in arterial blood was also significantly improved in

mitochondrial group (Table 1). Three different means were used to measure the integrity of the alveolar-capillary barrier in rats with ALI. Compared to shams, the LWDR was reduced in the lungs of ALI animals (Figure 2A), and transplantation of mitochondria did not change the lung weight ratio (Figure 2A). The BALF protein content and extravasated Evans blue dye content were significantly reduced in the lung tissues of mitochondrial-transplanted animals to levels similar to the sham-operated group (Figures 2B and 2C), suggesting improvement in pulmonary endothelial permeability following mitochondrial transplantation.

Table 1  
Measurements of arterial blood gases (ABG)

Parameters	Sham (n=10)	ALI (n=8)	ALI+Mito (n=8)
PaO <sub>2</sub> (mmHg)	95.96±5.03	78.88±3.82*,†	93.44±5.29
PaCO <sub>2</sub> (mmHg)	51.59±3.71	61.16±3.82*,†	56.11±4.54
SaO <sub>2</sub> (%)	96.80±0.52	94.95±1.30*,†	96.50±0.93
HCO <sub>3</sub> <sup>-</sup> (mmol/L)	28.38±2.58	32.84±3.07*	30.59±2.24
pH	7.363±0.03	7.342±0.03	7.345±0.03

Blood samples were collected via femoral artery and analyzed for ABG at 24 h after induction of acute lung injury (ALI); ALI+mito: rats with ALI received allogeneic mitochondrial transplantation; PaO<sub>2</sub> and PaCO<sub>2</sub>: partial pressures of oxygen and carbon dioxide in artery blood; SaO<sub>2</sub>: oxygen saturation in the artery blood. Data were analyzed using the Mann-Whitney U test and are presented as mean±SD. \*P<0.005 compared with sham group and †P<0.05 compared with ALI+mito group.

## Transplantation of mitochondria potentiates pulmonary vascular endothelial function

Transplantation of mitochondria did not affect the contraction responses to KCl depolarization and α1-adrenergic stimulation in the isolated pulmonary artery of rats with ALI (Figures 3A and 3B). However, the maximal tension of relaxation induced by 10<sup>-5</sup> M acetylcholine was significantly restored in the mitochondrial transplanted group (55.9±12.0 vs 73.6±5.6%, ALI vs mitochondrial groups, respectively; P=0.006; Figure 3C). However, the concentration-response curves of endothelial-dependent relaxation of pulmonary arteries were superimposed in the two ALI groups (Figure 3C). eNOS is one of the most important regulator of maintaining normal cellular function in the vascular endothelium.<sup>21</sup> Protein expressions of eNOS and its phosphorylated form (p-eNOS-S1177) in the pulmonary artery were suppressed in rats with ALI, but they were normalized following mitochondrial administration (Figure 3D).

# Transplantation of mitochondria attenuates damage of lung parenchyma

The severity of lung damage was assessed using a lung injury score, pulmonary hemorrhage, and tissue inflammation. Lung injury was precipitated by increased neutrophil infiltration and red blood cell accumulation in the alveolar and interstitial spaces at 24 h after intratracheal lipopolysaccharide instillation (Figure 4A). Consolidation, alveolar ectasia and hyaline membrane formation were noted in the lung parenchyma of rats with ALI (Figure 4A). Overall lung injury scores were significantly reduced in ALI animals received mitochondrial transplantation ( $1.57 \pm 0.11$  vs  $1.11 \pm 0.17$ , ALI vs ALI+mito, respectively;  $P=0.004$ ) (Figure 4B). Tissue hemoglobin content was reduced in animals with mitochondria transplants compared to placebo rats with ALI, which indicates pulmonary hemorrhage improvement in ALI (Figure 4C). Inflammatory cell infiltration was determined by the protein levels of iNOS and CD11b in the lung homogenates<sup>17</sup>. Mitochondrial transplantation significantly suppressed expressions of iNOS and CD11b in the lungs of endotoxin-induced ALI (Figure 5A). However, no differences in the lung myeloperoxidase activity were found between ALI animals that received placebo or mitochondrial treatment (Figure 5B).

## Discussion

Dr. McCully and his colleagues reported the first clinical application of mitochondrial transplantation in pediatric patients with ischemia-reperfusion-associated myocardial dysfunction<sup>16</sup>. Mitochondria transplantation have since been considered as a revolutionary approach for regenerative medicine<sup>22</sup>. Mitochondrial transplantation has the advantage of rapid isolation and purification to meet the clinical needs<sup>23</sup>. Mitochondria are unlikely to induce alloreactivity and damage-associated molecular pattern molecules reaction. Therefore, mitochondria have potential for both syngeneic and allogeneic transplants<sup>24</sup>. Furthermore, exogenous mitochondria can be conveniently administrated through intravenous route<sup>14, 18</sup>.

Intratracheal lipopolysaccharide instillation destroys lung parenchyma through the generation of proteases and reactive oxygen and nitrogen species produced by activated polymorphonuclear cells in the interstitial and alveolar compartments that is comparable to the pathophysiological changes in patients with ALI/ARDS<sup>25</sup>. Previous studies found that the intra-alveolar inflammatory reaction composed of a neutrophilic exudate at the initial 6 to 12 h, a monocytic exudate peaking at 24 h, and then followed by a lymphocytic exudate<sup>26</sup>. After lipopolysaccharide instillation, we detected hypoxemia and hypercapnia, indicating the impaired gas exchange due to development of alveolar dead space and ventilation-to-perfusion mismatch in these animals<sup>27</sup>. The alveolar-capillary barrier is organized by a network of collagen and laminin that separates the epithelium and endothelium<sup>28</sup>. The capillary endothelium is a semipermeable barrier to fluid exchange, whereas the alveolar epithelium is a tight layer that restricts the passage of water, electrolytes and hydrophilic solutes to the air space<sup>29</sup>. Increased

protein contents in the BALF, pulmonary hemorrhage, inflammatory cell infiltration, and parenchymal consolidation further confirmed alveolar-capillary disruption.

Under IVIS, we detected the engraftment of transplanted mitochondria into the pulmonary circulation was significantly more enhanced in lungs of rats with ALI. The uptake of these exogenous mitochondria was further confirmed by the increased ATP concentrations in the lungs. Consistent with our previous report<sup>18</sup>, these findings suggest that intravenous delivery of exogenous mitochondria can selectively locate and be taken up by areas of the pulmonary arterial system with mitochondrial dysfunction. After two mitochondrial transplants, the protein content recovered in BALF and extravasated Evans blue dye was significantly reduced, suggesting improved integrity of the monolayer capillary endothelium in preventing leakage of plasma proteins and albumin into the air-space<sup>19, 30</sup>. However, lung water content measured by LWDR was not differ significantly among the three treatment groups. We speculate that the development of extensive alveolar ectasia, lung consolidation and predominant infiltration of lymphocytes at 24 h after ALI<sup>31</sup>, might obscure the formation of pulmonary edema.

Pulmonary endothelial function was also assessed using vasomotor function tests and through the expression of eNOS in the pulmonary artery. The isometric tension analysis found no differences in the concentration response curves of phenylephrine and acetylcholine of ALI animals treated with PBS or mitochondria. However, the relaxation response to maximal concentration of acetylcholine, endothelial-dependent relaxation induced by stimulating the muscarinic receptors, was significantly potentiated in animals received mitochondrial treatment. The increased isometric tension in the pulmonary artery of ALI rats that received PBS at high acetylcholine concentrations might be due to vasoconstriction reaction mediated by direct stimulation of the muscarinic receptors on the vascular smooth muscle cells where the endothelial layer is damaged or denuded<sup>5, 32</sup>. Since the lung injury primarily originated from the bronchoalveolar site in this model, it was therefore reasonable that fewer changes in the vasoreactivity tests were detected in comparison to experimental models of direct pulmonary endothelial injury, such as intravenous administration of oleic acid<sup>5</sup>. Endothelial NOS is an important enzyme in the endothelial cells that synthesizes optimal amount of NO in order to maintain a normal endothelial homeostasis<sup>21, 33</sup>. Phosphorylation of eNOS at Ser1177 through Akt/Protein kinase B or AMP-activated protein kinase is a critical requirement for eNOS activation<sup>34</sup>. The Western blot analysis confirmed that mitochondria transplantation restored eNOS and p-eNOS-S1177 levels in the pulmonary artery, further supporting the engraftment of exogenous mitochondria into the pulmonary circulation can improve endothelial function in rats with ALI. In addition, the increase in capillary endothelium integrity also reduces trans-endothelial migration of immune cells towards the inflammatory cascade in lung tissues<sup>35</sup>. iNOS and CD11b are considered as the cell markers of activated M1 macrophages during the pro-inflammatory phase of ALI<sup>36</sup>. The significant suppression of iNOS and CD11b expressions in the injured lung following mitochondrial transplantation might imply the reduced pro-inflammatory cell infiltration in rats with ALI.

The exact mechanisms underlying mitochondrial transplantation in tissue regeneration remains undetermined, but three potential mechanisms have been proposed<sup>37</sup>. The Ca<sup>2+</sup> buffering capacity of

mitochondria may attenuate the environmental  $\text{Ca}^{2+}$  overload during cellular stress through the opening of voltage-dependent anion channels<sup>38</sup>. The second theory proposes that the internalization of exogenous mitochondria in recipient cells improves the mitochondrial function of target cells by increasing ATP generation and mitochondrial oxygen consumption<sup>39</sup>. Finally, the viable exogenous mitochondria may release ATP into the extracellular environment and salvage the dysfunctional cells<sup>37, 40</sup>. However, there is currently insufficient evidence to support the direct anti-inflammatory potential of transplanted mitochondria during tissue injury. It is then expected that we did not find any significant effects of mitochondrial transplant on the myeloperoxidase activity assay in the lung tissue exposed to endotoxin instillation.

There are a number of limitations in our study. First, the intracellular localization of these exogenous mitochondria was not determined in this study. Secondly, this study recorded very few ALI-related mortality throughout the study period. The effects of mitochondrial transplantation on other outcome measurements (e.g. overall mortality rate and lung parenchymal repair) were not studied. Thirdly, the optimal dose of mitochondria and the frequency of treatment were rather arbitrary, as there is still no general consensus on the standard dosing for mitochondrial transplantation<sup>41</sup>. Fourthly, this was a pre-clinical, proof-of-concept research for the potential application of mitochondrial transplantation in ALI/ARDS and more mechanistic analysis are still under investigation in our laboratory.

In conclusion, this is the first report demonstrating that intravenous transplantation of viable allogeneic mitochondria in rats with endotoxin-induced ALI significantly improves gas exchange by reducing alveolar-capillary barrier endothelial disruption. However, the effects on survival outcomes and long term recovery of pulmonary function in subjects with ALI after mitochondrial transplants require further investigation.

## **Declarations**

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## **DATA AVAILABILITY**

The datasets used and analyzed during the current study are available from the corresponding author on reasonable request.

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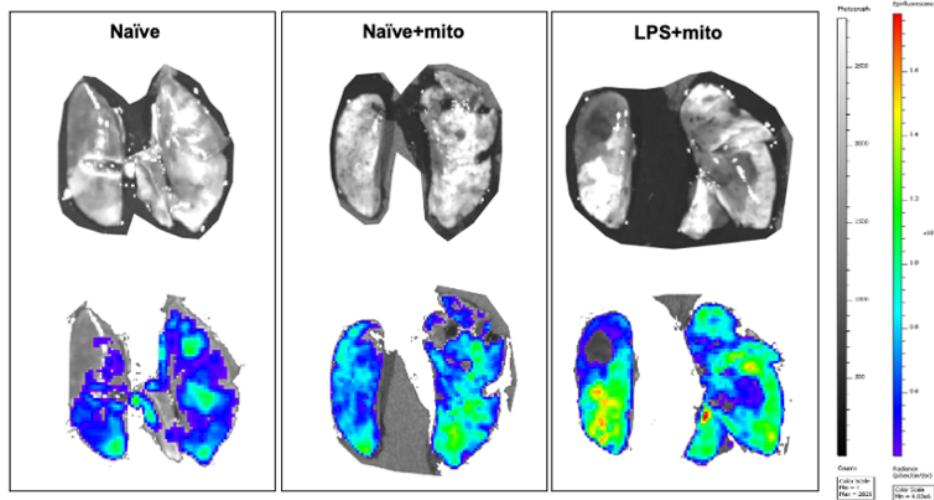
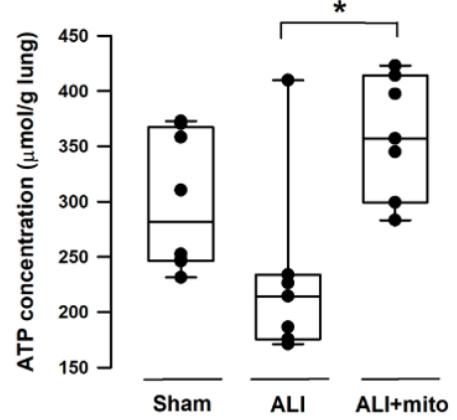
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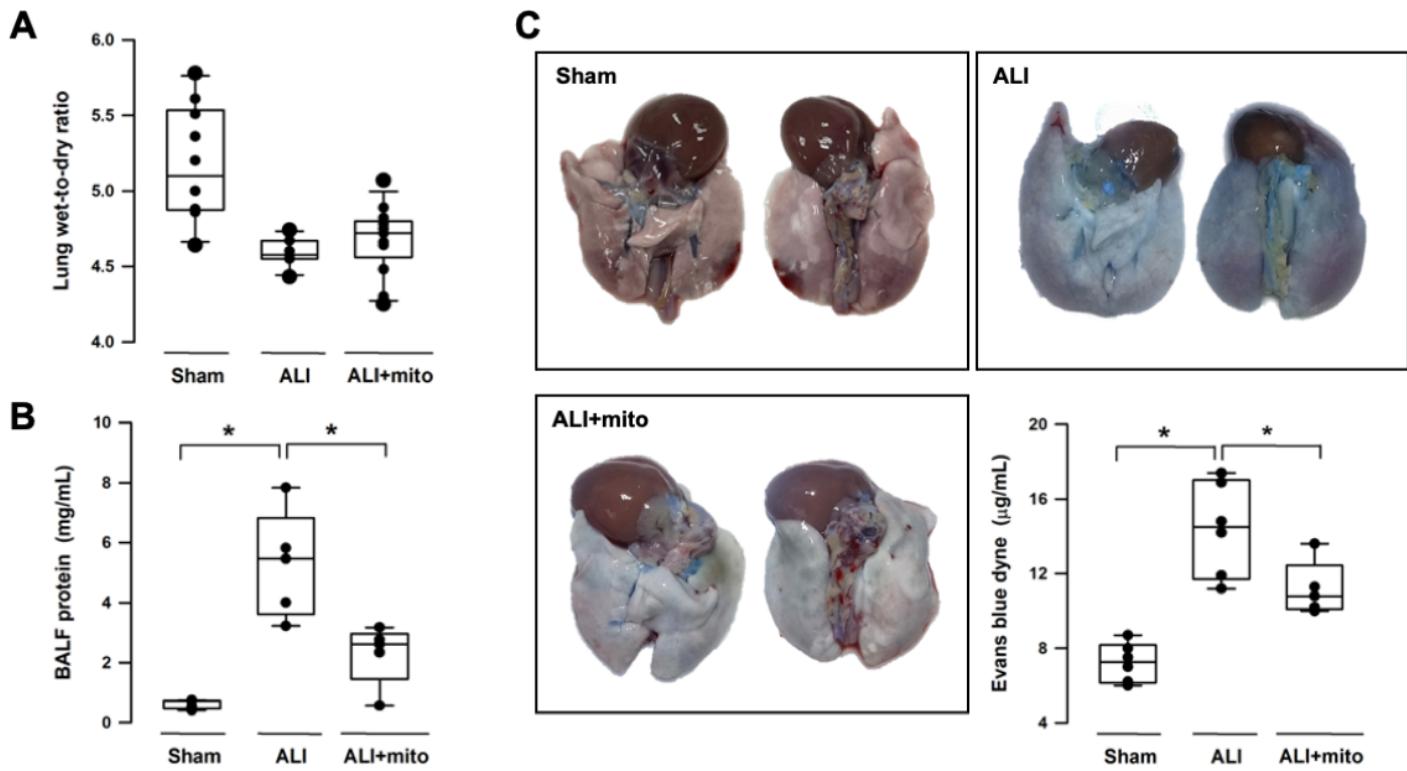
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## Figures

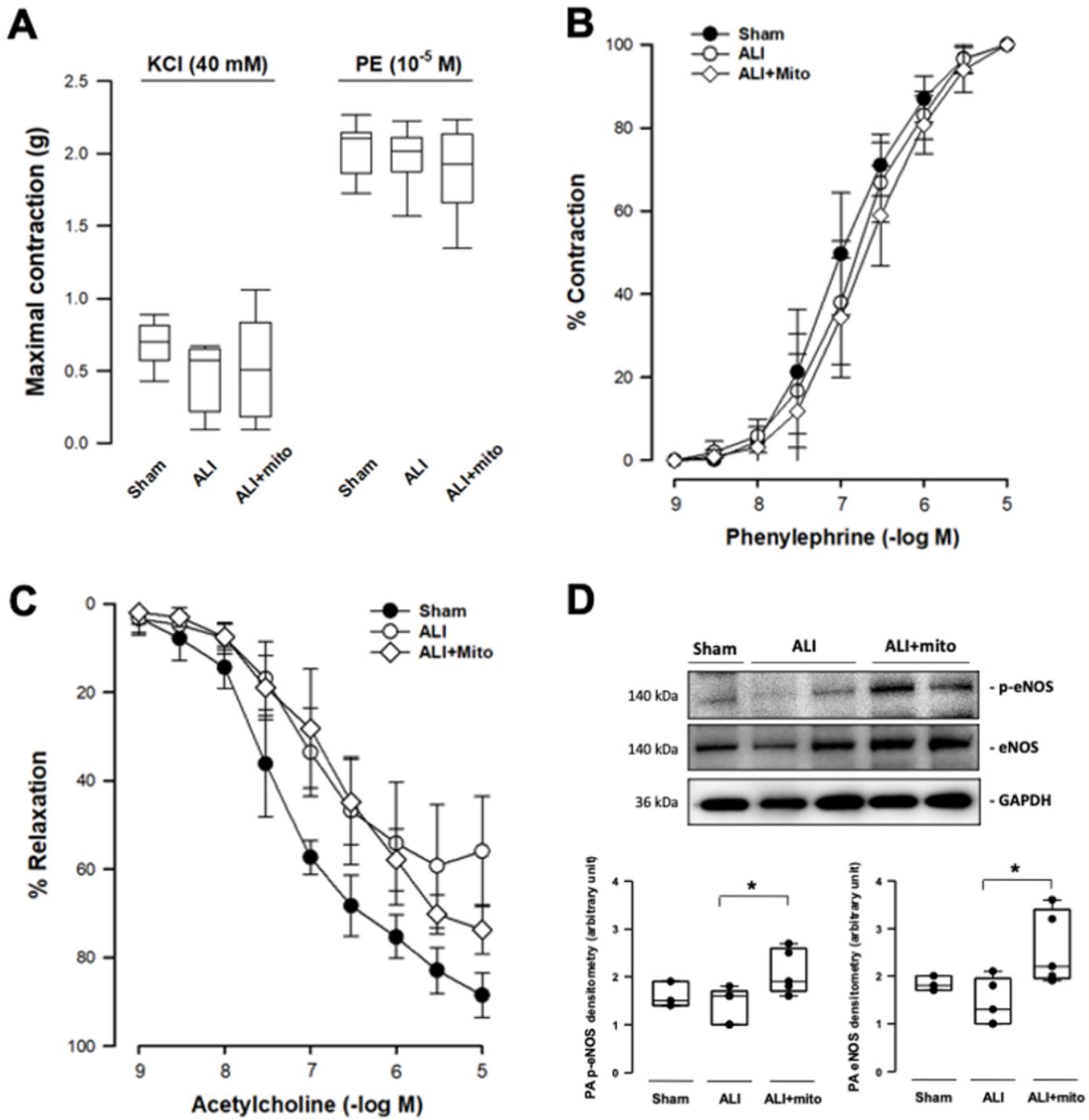
**A****B****Figure 1**

**(A).** Representative images acquired by an *in vivo* imaging system (IVIS) (Caliper IVIS Spectrum System). Mitochondria were freshly isolated from the soleus muscles of naïve rats and were labeled with a green fluorescence dye (MitoTracker™ Green, Invitrogen™) at 30 min before intravenous administration. Upper panels were photographs taken under normal light and the lower panels were the fluorescence photographs taken under IVIS. Lower MitoTracker™ fluorescence densities (blue to green) were found in the lungs of naïve rats with or without mitochondria. Fluorescence densities were significantly enhanced in the lungs of rats treated with lipopolysaccharide (LPS) (yellow to red). Images were taken from 3 different animals in each group. **(B).** Tissue ATP concentration in the lungs of animals with ALI was restored following transplantation of freshly-isolated allogeneic mitochondria. Results are presented as box-and-whisker plots, in which the horizontal lines of color boxes indicate the 75<sup>th</sup> percentile, median and 25<sup>th</sup> percentile of the distribution, and the upper and lower whiskers indicate the maximal and minimal values. Data were analyzed using the Rank Sum Test, \*P=0.009, n=8 different animals in each group.



**Figure 2**

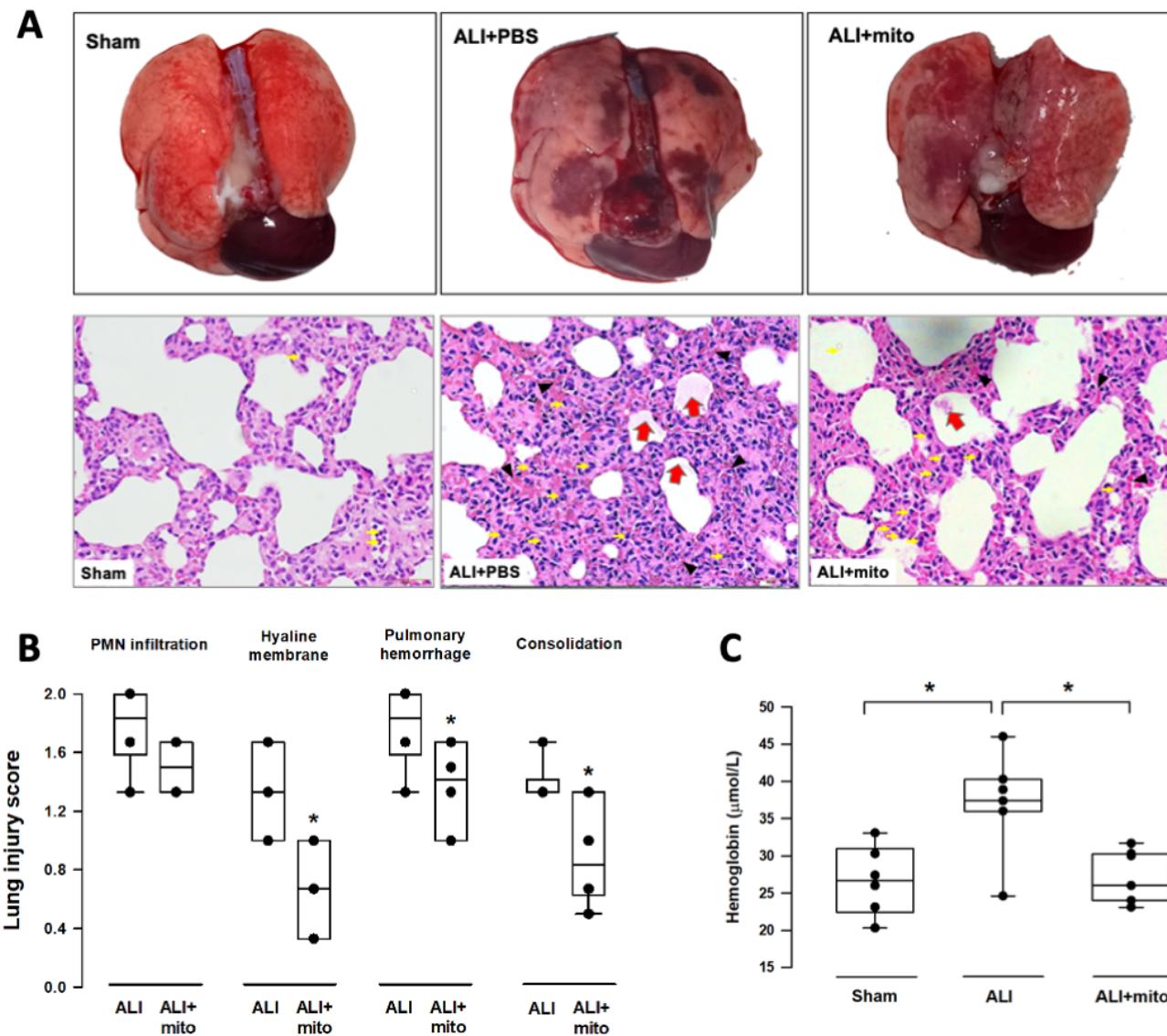
Changes in the alveolar-capillary permeability were assessed by degree of lung edema (lung wet-to-dry ratio, LWDR), protein content in the broncho-alveolar lavage fluid (BALF) and extravasated Evan blue dyne in the lung tissue. There were not differences in the LWDR among the three treatment groups (**A**), but total protein discovered from the BALF (**B**) and Evan blue dyne (**C**) in the lung homogenates was significantly reduced in endotoxin-induced acute lung injury (ALI) rats treated with allogeneic mitochondria. Results are presented as box-and-whisker plots, in which the horizontal lines of color boxes indicate the 75<sup>th</sup> percentile, median and 25<sup>th</sup> percentile of the distribution, and the upper and lower whiskers indicate the maximal and minimal values. Data were analyzed using the Rank Sum Test, \*P<0.05, n=5-12 different animals in each group.



**Figure 3**

Measurements of isometric force of isolated pulmonary artery. **(A)**. Contraction responses to KCl (40 mM) and maximal concentration of phenylephrine (PE,  $10^{-5}$  M). **(B)** Contraction responses to cumulative addition of phenylephrine (PE,  $10^{-9}$  to  $10^{-5}$  M). There were no differences in the contraction responses to KCl depolarization and  $\alpha$ 1-adrenergic stimulation in the pulmonary artery segments of acute lung injury (ALI) rats with or without mitochondrial transplantation. **(C)**. Endothelium-dependent relaxation response to cumulative addition of acetylcholine ( $10^{-9}$  to  $10^{-5}$  M). The concentration response curves were

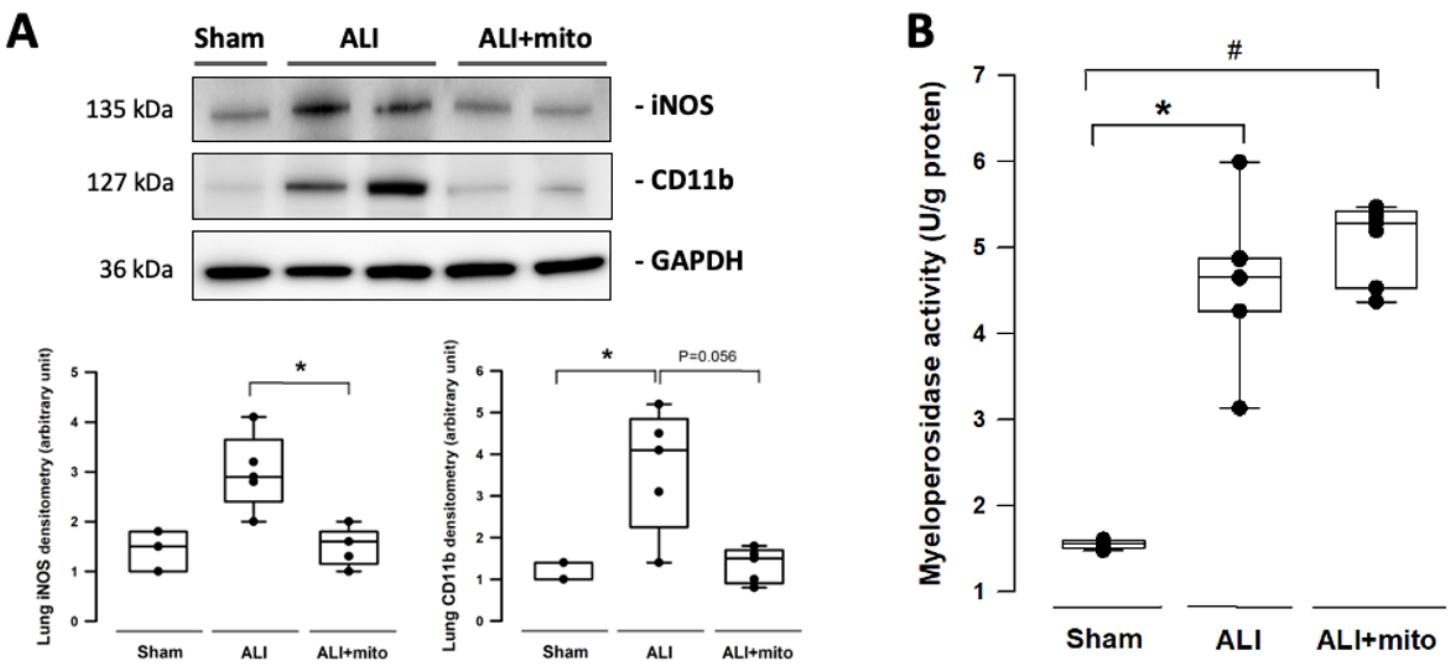
significantly impaired in ALI groups, as compared with sham-operated animals. The maximal tension of relaxation induced by  $10^{-5}$  M acetylcholine was significantly restored in the mitochondrial transplanted group ( $55.9\pm12.0$  vs  $73.6\pm5.6\%$ , ALI vs mitochondrial groups, respectively;  $P=0.006$  analyzed using the Rank Sum Test).  $n=6-8$  different animals in each group for isometric force measurement. **(C)**. Protein expressions of eNOS and phosphorylated eNOS (p-eNOS) in the rat pulmonary arteries. Results are presented as box-and-whisker plots, in which the horizontal lines of color boxes indicate the 75<sup>th</sup> percentile, median and 25<sup>th</sup> percentile of the distribution, and the upper and lower whiskers indicate the maximal and minimal values. Data were analyzed using the Rank Sum Test, \* $P<0.05$ ,  $n=3-5$  different animals in each group for Western blot analysis.



**Figure 4**

**(A).** Gross findings in lungs with acute lung injury (ALI) were heterogeneous, consisted of patchy consolidation and hemorrhage on the pleural surface of the lungs. The injured lungs were in unevenly

bluish-red color, as compared with the normal pink aerated appearance in the lungs of sham-operated rats. The gross findings of lung lobes are presented as dorsal views. Representative lung tissue sections stained with hematoxylin and eosin show alveolar wall thickening and consolidation, neutrophilic infiltration in the alveolar and the interstitial spaces (yellow arrows), extravasation of erythrocytes in the lung interstitium (black arrowheads) and filling of proteinaceous debris or formation of hyaline membrane in the airspaces (red arrows) in the rats with ALI. **(B)**. Severity of lung injury was scored using a semiquantitative histopathology score system<sup>19</sup>, which evaluates lung injury by degree of neutrophil infiltration, amount of hyaline membranes or proteinaceous debris, level of lung hemorrhage and extent of parenchymal consolidation. **(C)**. Quantification of lung hemorrhage by measuring hemoglobin content in the lungs. Results are presented as box-and-whisker plots, in which the horizontal lines of color boxes indicate the 75<sup>th</sup> percentile, median and 25<sup>th</sup> percentile of the distribution, and the upper and lower whiskers indicate the maximal and minimal values. Data were analyzed using the Rank Sum Test, \*P<0.05, n=6-7 different animals in each group.



**Figure 5**

**(A)**. Infiltration of leukocytes was assessed by expression of iNOS and CD11b in the lungs. **(B)**. Measurement of myeloperoxidase activity in the lungs of rats with or without endotoxin-induced acute lung injury (ALI). Results are presented as box-and-whisker plots, in which the horizontal lines of color boxes indicate the 75<sup>th</sup> percentile, median and 25<sup>th</sup> percentile of the distribution, and the upper and lower whiskers indicate the maximal and minimal values. Data were analyzed using the Rank Sum Test, \*P<0.05, #P=0.056.

whiskers indicate the maximal and minimal values. Data were analyzed using the Rank Sum Test,  
\*P<0.05, n=3-7 different animals in each group.