**Dexmedetomidine attenuates ventilator-induced lung injury in rats by up-regulating NLRC3**

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**Contributions:** (I) Conception and design: L Sun, M Wang; (II) Administrative support: B Zhang, X Zhang; (III) Provision of study material or patients: B Zhang, X Zhang; (IV) Collection and assembly of data: B Zhang, X Zhang; (V) Data analysis and interpretation: Q Li, F Ma; (VI) Manuscript writing: All authors; (VII) Final approval of manuscript: All authors.

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**Background:** Mechanical ventilation is a dispensable work in clinical treatment and rescue, and always caused of ventilator-induced lung injury (VILI). Dexmedetomidine is a clinical drug to prevent lung injury, but its mechanism still unclear.

**Methods:** Thirty-six SD rats were randomly divided into three groups: self-breathing control group (Group C), high tidal volume (V_T 20 mL/kg) group (Group H) and high V_T + dexmedetomidine group (Group DEX). Serum, lung tissue, bronchoalveolar lavage fluid (BALF) were collected after rats were sacrificed by anesthetic drug of pentobarbital sodium. The pathological changes of lung tissue were observed by hematoxylin and eosin stain (HE staining), and the lung injury score and wet/dry (W/D) ratio were tested to assess lung injury. The total protein level in BALF and contents of the interleukin-1β (IL-1β), IL-18 in serum and BALF were detected by enzyme-linked immunosorbent assay (ELISA), the mRNA and protein expression level of NLR Family CARD Domain Containing 3 (NLRC3), NLR Family Pyrin Domain Containing 3 (NLRP3), Apoptosis associated speck-like protein containing a CARD domain (ASC) and caspase-1 were measured by qRT-PCR and Western Blotting respectively.

**Results:** Compared with Group C, VILI mode of Group H were success established because of lung injury score and W/D value increased. when compared with Group H, which were decreased significantly in Group DEX (P<0.05), and the total protein level in BALF and the contents of IL-1β, IL-18 in serum and BALF of Group DEX were reduced markedly (P<0.05). Besides the mRNA and protein expression of NLRP3, ASC and caspase-1 in lung tissue of Group DEX were lowered dramatically (P<0.05). However, mRNA and protein expression of NLRC3 in lung tissue of Group DEX were up-regulated observably (P<0.05).

**Conclusions:** This study demonstrates that NLRC3 is involved in the VILI of rats, and dexmedetomidine can attenuate the VILI in rats by up-regulating the expression level of NLRC3.

**Keywords:** Dexmedetomidine; ventilator-induced lung injury (VILI); NLRC3; NLRP3

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**Introduction**

As an important means of treatment and rescue, mechanical ventilation plays an indispensable role in clinical work and was used widely (1,2), while positive pressure ventilation or anti-physiological ventilation was used in the application of mechanical ventilation, which may induce or aggravate lung injury, named ventilator-induced lung injury (VILI) (3). VILI includes of direct mechanical damage and indirect biological damage, and current researches divide mechanical...
damage into three parts: barotrauma, volutrauma and atelectrauma. Mechanical damage can mediate biotrauma through many pathways (4,5), during which the lung cell wall was damaged directly by external mechanical force, which results in a large release of cytokines in the alveolars and circulatory system to lead local or systemic inflammatory reactions subsequently (6). The mainly pathophysiological changes of VILI include the alveolar structural destruction, the increased pulmonary vascular permeability, the activation of inflammatory cells and inflammatory factors and the imbalance of oxidation and antioxidation, etc. (7). VILI has become one of the most serious complications of mechanical ventilation (8). Therefore, the prevention and treatment of VILI have become importantly and necessarily.

During mechanical ventilation, the external mechanical force will activate NLRP3 inflammasome, which is a member of the nucleotide oligomerization domain-like receptor (NLR) family. NLR family composed of NLRP3, an apoptosis-associated speck-like protein containing a CARD domain (ASC), and pro-caspase-1 (9). The activation of NLRP3 inflammasome releases the proinflammatory cytokines of interleukin (IL)-1β, IL-18 and finally induces lung and body inflammatory response (10). NLRC3, like NLRP3, also belongs to the NLR family. Its main structures include a nucleotide-binding oligomerization domain (NACHT) in the middle, a C-terminal leucine-rich repeat domain (LRR) on one side and an N-terminal cysteine aspartate recruitment domain (CARD) on the other side (11). Unlike NLRP3, NLRC3 does not participate in the composition of the inflammasome and, contrary to NLRP3, plays a negative regulatory role in inflammatory responses (12,13). However, the relationship between NLRC3 and VILI in rats still not clarified.

Dexmedetomidine, a fat-soluble alpha2 adrenergic receptor agonist, was first used in clinical work as a sedative and an analgesic drug in 1999 (14). Because dexmedetomidine has a weak inhibitory effect on the respiratory system and circulatory system, it is mainly used for sedation and analgesia in the intensive care unit, and as an anesthesia adjuvant for a perioperative period (15). The study found that dexmedetomidine has protective effects on a variety of organs, including lung. Dexmedetomidine can down-regulate high mobility protein 1 (HMGB1), reduce the expression of TNF-α, IL-1 and IL-6 to attenuate inflammatory response. Kim et al. (16) found that dexmedetomidine can inhibit the inflammatory response by blocking the TLR-4/NF-κB pathway. The research by Meng et al. (17) showed that infusion of dexmedetomidine can improve lung oxygenation in patients with restrictive lung disease and morbid obesity. At the same time, some studies have pointed out that dexmedetomidine can alleviate lipopolysaccharide-induced acute lung injury by inhibiting the activation of NLRP3 inflammasome (18).

Although previous studies have shown that dexmedetomidine can alleviate lung injury, its specific mechanism for lung protection is not yet perfect clarified. The purpose of this study is to discuss whether NLRC3 is involved in the process of VILI, and whether the lung protection mechanism of dexmedetomidine is related with NLRC3. We present the following article in accordance with the ARRIVE reporting checklist (available at http://dx.doi.org/10.21037/apm-19-375).

Methods

This study was approved by The Ethics Committee of Qingdao Municipal Hospital affiliated to Qingdao University in compliance with guidelines for the care and use of animals. Animals were treated in accordance with Guide for the Care and Use of Laboratory Animals (8th edition, National Academies Press). Experiments were performed under a project license (NO. 2018033).

Experimental animal and grouping

Thirty-six clean male SD rats, 240–260 g, 8 weeks old, were provided by Pengyue Experimental Unit (Jinan, China), with the animal certificate number SCXK (Shandong) 20140007. At temperature of 23–26 °C, all rats were free to diet for one week in SPF environment, and divided which into 3 groups by random number labeling method: self-breathing control group (Group C), high tidal volume (Vt) group (Group H), high Vt+ dexmedetomidine group (Group DEX), 12 rats per group.

Instruments and reagents

Small animal ventilator (R415, Ruiwode Life Technology Co., Ltd, China), BCA protein concentration kit ( P0010, Biyuntian Biotechnology Co., Ltd., China), IL-1β and IL-18 ELISA kit (JW-E14034, Jiwei Biological Technology Co., Ltd, China), PCR instrument (A600, Mio Instrument Co., Ltd., China), NanoDrop spectrophotometer (Nanodrop one, Thermo fisher, USA), Ultraviolet analyzer (JY0S-3D, Junyi Oriental Electrophoresis Equipment Co., Ltd., China), Real-time PCR instrument (Applied Biosystems, USA), Trizol RNA Extraction Reagent...
(B511311, Sangon Biochemical Technology Co., Ltd, China), PrimeScript™ RT reagent Kit with gDNA Eraser (RR047, TaKaRa, Japan), SYBR® Premix Ex Taq™ II (RR820A, TaKaRa, Japan), the primer of NLRC3, NLRP3, ASC, caspase-1 and internal reference β-actin (GENEWIZ, USA), NLRC3, NLRP3, ASC, caspase-1 and internal reference β-actin Primary antibody (ARG65683, Arigo Biotechnology Co., Ltd, China), goat anti-rabbit secondary antibody (ab182016, Abcam company., Ltd, USA).

Animal models
All rats were fasted for 8 hours before the experiment, and then were intraperitoneally injected with 3% pentobarbital sodium (50 mg/kg). The rats were fixed on the test bench in a supine position after anesthesia, and tracheotomy intubation was performed after neck skin preparation and disinfection. Arterial blood pressure was monitored by a right carotid artery, and venous access was established by a right jugular vein. The anal temperature of rats was maintained at 37–38 ℃ by an animal insulation blanket.

The method of reference (19) was used to establish the VILI model of rats. After successful tracheotomy and intubation, the rats in the Group C maintained spontaneous respiration; rats in the Group H and Group DEX were injected with rocuronium 0.1 mg/kg via the right jugular vein, and then connected to the small animal ventilator for mechanical ventilation for 4 hours after the spontaneous respiration disappeared. Rats in the Group DEX were intraperitoneally injected with dexmedetomidine 50 μg/kg 30 minutes before mechanical ventilation, and rats in the Group C and Group H were intraperitoneally injected with the same amount of normal saline 30 minutes before mechanical ventilation. Mechanical ventilation parameters were set as follows: V̇ T was set to 20 mL/kg, the suction and respiration ratio (I:E) was 1:1, the respiratory rate (f) was 80 times/min, the oxygen concentration (FiO₂) was 21%.

Specimen collection
After 4 hours of mechanical ventilation, 5 mL blood was extracted from the right carotid artery, centrifuged at 4 ℃ and 3,000 r/min for 10 minutes, and then the supernatant was collected and stored at −80 ℃. Then, it was dried in an electric drying box at 80 ℃ to a constant weight. The dry weight was weighed and the W/D of lung tissue was calculated.

Determination of wet/dry (W/D) ratio of lung tissue
The upper lobe of the right lung was taken and the surface moisture was absorbed by filter paper to weigh wet mass. Then it was dried in an electric drying box at 80 ℃ to a constant weight. The dry weight was weighed and the W/D of lung tissue was calculated.

Pathological evaluation of lung tissue and lung injury score
The middle lobe of the right lung was taken, fixed with 4% paraformaldehyde for 24 hours, embedded in paraffin, sectioned, stained with hematoxylin and eosin (HE), and observed under a light microscope. Lung injury score was performed by the method of reference (20). The scoring criteria included pulmonary edema, bleeding, neutrophil infiltration and small airway injury, each of which was rated as 0–4 points according to the severity of the lesion (0: no lesion or very mild disease; 1: mild lesion; 2: moderate lesions; 3: severe lesions; 4: extremely severe lesions). The total score of 4 items is the lung injury score.

Determination of total protein in BALF
The total protein content in BALF of rats was determined by BCA method according to the procedure described in the instructions of BCA Protein Concentration Kit.

Determination of cytokines in serum and BALF
The IL-1β and IL-18 levels in serum and BALF of rats were determined by ELISA according to the procedure described
in the instructions of ELISA Kit.

qRT-PCR

Partial tissues of the lower lobe of the right lung were taken, and the total RNA of lung tissues was extracted by Trizol one-step method after liquid nitrogen grinding according to the procedure described in the instructions. The concentration and purity of RNA were determined by ultraviolet absorption assay. The mRNA expressions of NLRC3, NLRP3, ASC and caspase-1 were determined by qRT-PCR, cDNA was synthesized by cDNA synthesis kit, and the mRNA of NLRC3, NLRP3, ASC and caspase-1 were amplified by ABI 7500 fluorescence quantitative PCR instrument. The primer sequences of NLRC3, NLRP3, ASC, caspase-1 and internal reference β-actin (Table 1). The reaction system of qRT-PCR was as follows: pre-denaturation at 95 °C, 30 seconds, denaturation at 95 °C, 5 seconds, annealing at 60 °C, 34 seconds, a total of 40 cycles. The differences between groups were compared by 2−ΔΔCt (ΔCt = target gene Ct value – internal reference Ct value), using the instrument's own software to do data analysis.

Western blotting

Partial tissue of the lower lobe of the right lung was cut into the size of rice grains and ground in a pre-cooled mortar. Tissue cell lysate was added. After 30 minutes on ice, the supernatant was centrifuged at 4 °C and 12,000 r/min for 15 minutes and the protein concentration was determined by BCA method. The total protein samples were taken out from the refrigerator at −80 °C. After dissolution, 5× loading buffer was added and reacted at 95 °C for 10 minutes. The membrane was transferred by SDS-PAGE and PVDF membrane wet transfer method under the conditions of 130 mA current, 60 V voltage and 60 minutes membrane transfer time. After sealing in the sealant for 1 hour, NLRC3 primary antibody (1:1,000), NLRP3 primary antibody (1:1,000), ASC primary antibody (1:1,000), caspase-1 primary antibody (1:1,000) and internal reference β-actin primary antibody (1:5,000) were added and incubated overnight at 4 °C. Then the solution was washed 3 times with 1× TBST, 10 minutes each time. The primary antibody membrane was placed into goat anti-rabbit secondary antibody (1:5,000) and incubated at room temperature for 1 hour. Then the solution was washed 3 times with 1x TBST, 10 minutes each time. Using IMAGE J software (NIH, USA) to scan the gray value of protein bands, the content of target protein was expressed by the ratio of the gray value of protein bands to the gray value of internal reference β-actin bands.

Statistical analysis

SPSS19.0 was used for statistical analysis. The measurement data of normal distribution were expressed as mean ± standard deviation ( x ± S ), and one-way ANOVA analysis was conducted for comparison between groups. P<0.05 was considered statistically significant.

Results

Dexmedetomidine alleviates VILI

Firstly, under light microscope, we observed that the alveolar structure of rats in Group C was normal, the alveolar wall was continuous, and the boundary was clear. In Group H, alveolar cavity fusion, alveolar wall thickening and rupture, pulmonary interstitial edema and a large number of inflammatory cell infiltration were observed. In Group DEX, alveolar injury was milder than that in Group H, slight edema was observed in the pulmonary interstitium, and a small amount of inflammatory cells

Table 1 The primer sequence of NLRC3, NLRP3, ASC, caspase-1 and internal reference β-actin

<table>
<thead>
<tr>
<th>Gene name</th>
<th>Forward primer (5'-3')</th>
<th>Reverse primer (5'-3')</th>
</tr>
</thead>
<tbody>
<tr>
<td>NLRC3</td>
<td>CAGATTGGAACAAAGGAGCCA</td>
<td>CGTTCCGTGTTATCCTCAGAGCA</td>
</tr>
<tr>
<td>NLRP3</td>
<td>CAGGGCGATCAAAGCGAGAC</td>
<td>AGAGATATCCAGCAACCTATCCA</td>
</tr>
<tr>
<td>ASC</td>
<td>TGTGGTTAGAGACATGGGCACTACAG</td>
<td>GCCATACAGGACATCCAGCAA</td>
</tr>
<tr>
<td>Caspase-1</td>
<td>ACTCGTACACGTCTTGCCCTCA</td>
<td>CTGGGCAGGCGAGCAGATAATC</td>
</tr>
<tr>
<td>β-actin</td>
<td>TCCTGTGGGATCAATGAAAACT</td>
<td>GAAGCATTGCGGTCACGAAGAT</td>
</tr>
</tbody>
</table>

exuded from alveolar cavity (Figure 1).

**Dexmedetomidine decreases lung injury score to prevent VILI**

Next, lung injury score was measured in different groups. We discovered that compared with Group C, the lung injury scores of Group H and Group DEX were increased nearly 6 folds (P<0.01), and compared with Group H, the lung injury score of Group DEX was largely reduced (P<0.05, Figure 2).

**Dexmedetomidine reduces W/D of lung tissue to alleviate VILI**

Latterly, the effect of dexmedetomidine on W/D of lung tissue was checked. Compared with Group C, the W/D of Group H and Group DEX were significantly increased (P<0.05). Compared with Group H, the W/D of Group DEX was decreased 1.4 folds obviously (P<0.05, Figure 3).

**Dexmedetomidine down-regulates the content of total protein in BALF to prevent VILI**

And compared with Group C, the total protein content in BALF of Group H and Group DEX were increased (P<0.05). Compared with Group H, the total protein content in BALF of Group DEX was also significantly reduced (P<0.05, Figure 4).

**Dexmedetomidine decreases cytokines in serum and BALF to inhibit VILI**

Besides, total protein in BALF were measured by ELISA kits, we found that compared with Group C, the levels of IL-1β and IL-18 in serum and BALF of Group H and Group DEX were obviously increased (P<0.05). Compared
with Group H, the levels of IL-1β and IL-18 in serum and BALF of Group DEX were significantly decreased (P<0.05, Figure 5).

**Dexmedetomidine reduces NLRP3, ASC, caspase-1 mRNA level, and increases NLRC3 to prevent VILI**

Moreover, qRT-PCR results showed that Compared with Group C, the mRNA expression of NLRC3 in lung tissues of Group H was increased (P<0.01), while the mRNA expression of NLRC3 in lung tissues of Group DEX was increased (P<0.01), and the mRNA expression of NLRP3, ASC and caspase-1 in lung tissue of Group H and Group DEX were significantly increased (P<0.05). Compared with Group H, the mRNA expression of NLRC3 in lung tissues of Group DEX was increased, the mRNA expression of NLRP3, ASC and caspase-1 in lung tissue of Group DEX were decreased (P<0.05, Figure 6).

**Dexmedetomidine reduces NLRP3, ASC, caspase-1 mRNA protein level, and increases NLRC3 to alleviate VILI**

At last, Western blotting results in consistence results with qRT-PCR. Compared with Group C, the protein expression of NLRC3 in lung tissues of Group H was decreased (P<0.01), while the protein expression of NLRC3 in lung tissues of Group DEX was increased (P<0.01), and the protein expression of NLRP3, ASC and caspase-1 in lung tissue of Group H and Group DEX were increased (P<0.05). Compared with Group H, the protein expression of NLRC3 in lung tissues of Group DEX was increased, the protein expression of NLRP3, ASC and caspase-1 in lung tissue of Group DEX were decreased (P<0.05, Figure 7).

**Discussion**

Mechanical ventilation plays an irreplaceable role in clinical work and functions as an effective means for life support and treatment, but its application in VILI will cause pulmonary complications. VILI is wildly used in people with lung disease. VILI deteriorates into acute respiratory distress syndrome (ARDS) or even result in death of patients was due to alveolar collapse or gas exchange disorder (21). In addition, with the aggravation of pulmonary inflammation, some inflammatory cytokines can spread to other organs of the body through the circulatory system, which leading to the outbreak of whole-body cascades of inflammation to endanger the health and safety of patients. Therefore, it has great clinical significance to explore the mechanism and preventive measures of VILI. VILI model of rats were established by referring to Wu reported methods (19). After mechanical ventilation with VT of 20 mL/kg for 4 hours in Group H, the phenomenon of alveolar cavity fused, alveolar wall thickened and fractured, pulmonary interstitial edema, and a large number of inflammatory cells infiltrated in the alveolar cavity were observed. Lung injury score, W/D of lung tissue and total protein content of BALF were significantly higher than those in Group C, indicating that VILI model of rats has been success established.

NLRP3 inflammasome belongs to NLR family, which is a type of pattern recognition receptor (PRR) in cytoplasm (22). Studies have shown that NLR, includes
NLRP3, apoptosis-associated speck-like protein containing a CARD domain (ASC) and pro-caspase-1, is mainly expressed in macrophages, lymphocytes (23). During mechanical ventilation treatment, mechanical force acts on the lungs, causing direct mechanical damage to alveolar cells due to the accumulation of reactive oxygen species (ROS) in the mitochondria of alveolar macrophages (24). NLRP3 exposes its effect domain through oligomerization of NOD domain to collect ASC after sensing ROS and other stimulation. ASC is a kind of adaptor protein containing CARD and pyrin domain (PYD). When ASC is recruited, its PYD combines to the N-terminus of NLRP3 and its CARD binds pro-caspase-1 to form the NLRP3 inflammasome (25-27). After the activation of NLRP3 inflammasome, pro-caspase-1 cleaved itself into caspase-1 initiates a cascade inflammatory response mediated by the maturation and release of IL-1β and IL-18 protein expressions of NLRP3. ASC and caspase-1 in the lung tissues of rats in Group H were significantly increased compared with those in Group C, and the contents of IL-1β and IL-18 in serum and BALF of rats in Group H had the similar trend with above. These results suggest that NLRP3 inflammasome is involved in the process of VILI activated by it and the downstream pro-inflammatory cytokines of IL-1β and IL-18 are matured and released to induce the pulmonary inflammatory response eventually. NLRC3 also belongs to the NLR family, which exists in the cytoplasm and wildly discovered in macrophages, lymphocytes, epithelial cells and other immune cell (3,28-30). Unlike NLRP3, NLRC3 plays a negative regulatory role in inflammatory response. Previous studies (31) have shown that NLRC3 can monitor the production of inflammatory cytokines in macrophages. NLRC3 inhibits TLR4/NF-κB pathway activation, by reducing the level of TRAF6, and affects the transmission of inflammatory signals in the downstream of TLR4/NF-κB pathway by inhibited the
phosphorylation of NF-κB. Besides, further studies (32,33) showed that NLRC3 binds to Stimulator of Interferon Inducible Genes (STING) to affect the interaction between STING and TANK-binding kinase 1 (TBK1) and the production of downstream type I interferon (IFN), and then inhibits the innate immune response of STING to viral DNA, cytosolic DNA or c-di-GMP. In addition, NLRC3 can bind to pro-caspase-1 in NLRP3 inflammasome, which affects the interaction between ASC and pro-caspase-1 and then inhibits the activation of NLRP3 inflammasome (34). There is still no relevant study on whether NLRC3 is involved in the process of VILI in rats. Our results showed that the mRNA and protein expressions of NLRC3 in the lung tissues in rats of Group H were significantly down-regulated compared with Group C. These results suggest that NLRC3 is involved in the process of VILI and the down-regulated expression of NLRC3 can induce or aggravate VILI occurrence in rats.

Dexmedetomidine, a safe alpha-2 adrenergic receptor agonist, is widely used as a sedative in clinical practice. In recent years, studies have found that dexmedetomidine plays a certain protective effect on the lungs and it can reduce acute lung injury caused by sepsis in rats by down-regulating the RAGE pathway signal (35); Dexmedetomidine inhibits the expression of IL-17, and decreases its effect on NF-κB signaling pathway activation, to inhibit inflammatory response (36). Moreover, studies showed that it can negatively regulate the inflammatory response through the TLR4/NF-κB signaling pathway mediated by HMGB1 and PI3K/Akt/mTOR signaling pathway (37,38). Besides, Dexmedetomidine may also reduce LPS-induced acute lung injury by inhibiting the activation of NLRP3 inflammasome (18). However, it inhibits the activation of NLRP3 inflammasome through which pathway remains unclear. Our results showed that the pathological injury of lung tissue and the W/D of lung

Figure 6 The mRNA expression of NLRC3, NLRP3, ASC and caspase-1 in lung tissue in rats of different groups. (A) The mRNA expression level of NLRC3 in three groups of rats; (B) the mRNA expression level of NLRP3; (C) the mRNA expression level of ASC; (D) the mRNA expression level of caspase-1. Group C: self-breathing control group, Group H: high tidal volume (V_T) group (V_T is 20 mL/kg), Group DEX: high V_T + dexmedetomidine group (V_T is 20 mL/kg). *, P<0.05 vs. Group C; #, P<0.05 vs. Group H. NLRC3, NLR Family CARD Domain Containing 3; NLRP3, NLR Family Pyrin Domain Containing 3; ASC, apoptosis associated speck-like protein containing a CARD domain.
tissue or total protein content in BALF in Group DEX were obviously reduced in contrast with Group H, these results demonstrate that dexmedetomidine can reduce VILI in rats. Meanwhile, the mRNA and protein expressions of NLRC3 in Group DEX was up-regulated compared with that in Group H. Compared with Group H, the mRNA and protein expressions of NLRP3, ASC and caspase-1 in Group DEX were down-regulated, and the level of pro-inflammatory cytokines such as IL-1β and IL-18 in serum and BALF were largely reduced also. These results suggest that dexmedetomidine can reduce inflammation and inhibit VILI in rats by up-regulating the expression of NLRC3, and the mechanism may be related to NLRC3 inhibiting the activation of NLRP3 inflammasome to block the maturation and release of pro-inflammatory cytokines such as IL-1β and IL-18. However, the study still not clarified the specific mechanism that dexmedetomidine regulates NLRC3, nor the necessary association between NLRC3 and NLRP3 inflammasome, which present the limitations of the study. Therefore, in future studies, we hope that we can in-depth study of the role of NLRC3 in VILI with cell experiments, and explore the specific mechanism of dexmedetomidine on NLRC3.

**Conclusions**

The study demonstrates that NLRC3 is involved in the process of VILI in rats. Dexmedetomidine can alleviate VILI occurrence by up-regulating the expression of NLRC3, and its mechanism may be result of NLRC3
inhibits the activation of NLRP3 inflammasome and blocks the maturation and release of pro-inflammatory cytokines of IL-1β and IL-18.

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**Footnote**

**Reporting Checklist:** The authors have completed the ARRIVE Reporting Checklist. Available at [http://dx.doi.org/10.21037/apm-19-375](http://dx.doi.org/10.21037/apm-19-375)

**Data Sharing Statement:** Available at [http://dx.doi.org/10.21037/apm-19-375](http://dx.doi.org/10.21037/apm-19-375)

**Conflicts of Interest:** All authors have completed the ICMJE uniform disclosure form (available at [http://dx.doi.org/10.21037/apm-19-375](http://dx.doi.org/10.21037/apm-19-375)). 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. Experiments were performed under a project license (NO. 2018033) granted by The Ethics Committee of Qingdao Municipal Hospital affiliated to Qingdao University in compliance with guidelines for the care and use of animals. Animals were treated in accordance with Guide for the Care and Use of Laboratory Animals (8th edition, National Academies Press).

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