

## **POSTER PRESENTATION**

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# Effects of pressure control and pressure support ventilation on ventilator induced lung injury in experimental acute respiratory distress syndrome with intra-abdominal hypertension

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#### Introduction

In acute respiratory distress syndrome (ARDS), intraabdominal hypertension (IAH) increases intra-thoracic pressures, leading atelectasis and deterioration of respiratory mechanics and gas-exchange. The optimal setting of mechanical ventilation (MV) and its impact on respiratory function and ventilator-induced lung injury (VILI) in ARDS associated with IAH needs to be better clarified. Lung-protective MV with low tidal volume  $(V_T)$  and positive end-expiratory pressure (PEEP) has been recommended; however, assisted MV may be a favorable alternative to controlled MV at the early phase of ARDS, since it requires less sedation, no paralysis and is associated with better lung protection, reducing the risk of VILI. We hypothesized that pressure-support ventilation (PSV) improve pulmonary morphofunction and minimize lung injury in ARDS with IAH.

## **Objectives**

To compare the effects of PSV with protective MV (PCV) on arterial blood gases, lung mechanics and histology, as well as to identify biological markers of inflammation and fibrogenesis in a model of ARDS with IAH.

### Methods

24 Wistar rats (250-300 g) were submitted to the a sequence of events: 1) receive *Escherichia coli* lipopoly-saccharide (LPS) intraperitoneally (1,000 µg); 2) waiting

period of 24 hours for development of ARDS; 3) anesthesia and mechanical ventilation; 4) induction of IAH (15 mmHg) or not; 5) random assignment to PCV ( $V_T = 6 \text{ mL/kg}$ , respiratory rate (RR) = 80 breaths/min, fraction of inspired oxygen (FIO<sub>2</sub>) = 0.4 and PEEP = 5 cmH<sub>2</sub>O) or PSV. During PCV, animals were paralyzed with pancuronium bromide. In PCV and PSV, the driving pressure was adjusted to achieve  $V_T = 6 \text{ ml/kg}$ . In addition, in PCV, the RR was controlled to keep minute ventilation constant (160 ml/min). Peak (Ppeak,RS), and mean (Pmean,RS) airway pressures and arterial blood gases were analyzed at baseline and at the end of 1 h ventilation. Lungs were removed for lung histology and molecular biology analysis [mRNA expression of interleukin (IL)-6, and pro-collagen type III (PCIII)].

### Results

PSV improved oxygenation regardless of IAH. In ARDS with IAH, PSV, compared to PCV group, was associated with greater reduction in Ppeak, (PSV: 11.4  $\pm$  2.4 cmH<sub>2</sub>O, PCV: 16.9  $\pm$  0.5 cmH<sub>2</sub>O, p < 0.05) and Pmean, (PSV: 5.8  $\pm$  1.9 cmH<sub>2</sub>O, PCV: 9.6  $\pm$  0.2 cmH<sub>2</sub>O, p < 0.05). Furthermore, PSV reduced the amount of alveolar collapse, and the mRNA expression of interleukin (IL)-6 and type III procollagen compared to PCV.

## **Conclusions**

In this model of ARDS with IAH, PSV, compared to PCV, promoted functional and lung morphological benefit thus mitigating VILI.

Full list of author information is available at the end of the article



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