

| pISSN 2586-6052 | eISSN 2586-6060

## Association between mechanical power and intensive care unit mortality in Korean patients under pressure-controlled ventilation

Jae Kyeom Sim<sup>1</sup>, Sang-Min Lee<sup>2</sup>, Hyung Koo Kang<sup>3</sup>, Kyung Chan Kim<sup>4</sup>, Young Sam Kim<sup>5</sup>, Yun Seong Kim<sup>6</sup>, Won-Yeon Lee<sup>7</sup>, Sunghoon Park<sup>8</sup>, So Young Park<sup>9</sup>, Ju-Hee Park<sup>10</sup>, Yun Su Sim<sup>11</sup>, Kwangha Lee<sup>12</sup>, Yeon Joo Lee<sup>13</sup>, Jin Hwa Lee<sup>14</sup>, Heung Bum Lee<sup>15</sup>, Chae-Man Lim<sup>16</sup>, Won-II Choi<sup>17</sup>, Ji Young Hong<sup>18</sup>, Won Jun Song<sup>19</sup>, Gee Young Suh<sup>20</sup>

For further information on the authors' affiliations, see Additional information.

**Background:** Mechanical power (MP) has been reported to be associated with clinical outcomes. Because the original MP equation is derived from paralyzed patients under volume-controlled ventilation, its application in practice could be limited in patients receiving pressure-controlled ventilation (PCV). Recently, a simplified equation for patients under PCV was developed. We investigated the association between MP and intensive care unit (ICU) mortality.

**Methods:** We conducted a retrospective analysis of Korean data from the Fourth International Study of Mechanical Ventilation. We extracted data of patients under PCV on day 1 and calculated MP using the following simplified equation:  $MP_{PCV} = 0.098 \cdot respiratory rate \cdot tidal volume \cdot (\Delta P_{insp} + positive end-expiratory pressure), where <math>\Delta P_{insp}$  is the change in airway pressure during inspiration. Patients were divided into survivors and non-survivors and then compared. Multivariable logistic regression was performed to determine association between  $MP_{PCV}$  and ICU mortality. The interaction of  $MP_{PCV}$  and use of neuromuscular blocking agent (NMBA) was also analyzed.

**Results:** A total of 125 patients was eligible for final analysis, of whom 38 died in the ICU.  $MP_{PCV}$  was higher in non-survivors (17.6 vs. 26.3 J/min, P<0.001). In logistic regression analysis, only MP-PCV was significantly associated with ICU mortality (odds ratio, 1.090; 95% confidence interval, 1.029–1.155; P=0.003). There was no significant effect of the interaction between  $MP_{PCV}$  and use of NMBA on ICU mortality (P=0.579).

**Conclusions:** MP<sub>PCV</sub> is associated with ICU mortality in patients mechanically ventilated with PCV mode, regardless of NMBA use.

Key Words: artificial respiration; intensive care unit; mechanical ventilators; mortality

## INTRODUCTION

Mechanical ventilation is an essential component of critical care, but it can damage the lungs, an event called ventilator-induced lung injury (VILI). Therefore, the primary goal of mechanical ventilation is to maintain adequate gas exchange and to reduce the work of breathing while minimizing VILI [1]. To achieve this goal, lung protective strategies, in which

## **Original Article**

Received: July 3, 2023 Revised: December 11, 2023 Accepted: December 13, 2023

#### Corresponding author

Gee Young Suh Division of Pulmonary and Critical Care Medicine, Department of Medicine, Samsung Medical Center, Sungkyunkwan University School of Medicine, 81 Irwon-ro, Gangnam-gu, Seoul 06351, Korea Tel: +82-2-3410-3429 Fax: +82-2-3410-6956 Email: smccritcare@gmail.com

© 2024 The Korean Society of Critical Care Medicine

This is an Open Access article distributed under the terms of Creative Attributions Non-Commercial License (https://creativecommons.org/ li-censes/by-nc/4.0/) which permits unrestricted noncommercial use, distribution, and reproduction in any medium, provided the original work is properly cited.



tidal volume and plateau pressure are limited, have been widely adopted [2]. However, other ventilator variables such as respiratory rate and driving pressure have also been shown to be associated with the development of VILI [3,4]. Because of the interdependence of the variables and the requirement for adequate gas exchange, adjustment of one variable results in changes in the other variables. Thus, it is difficult to predict how the adjustment of one variable will affect VILI.

Gattinoni et al. [5] proposed the mechanical power (MP) concept, which refers to the amount of energy transferred to the lungs as the result of mechanical ventilation and integrates various ventilator variables affecting VILI; these authors proposed a calculation of MP based on the equation of motion. Experimental studies have found correlations between MP and lung injury [6-8]. In a large observational study, MP was associated with higher mortality, longer intensive care unit (ICU) and hospital lengths of stay, and fewer ventilator-free days [9]. Gattinoni and colleagues' original equation for MP is based on volume-controlled ventilation with a linear increase in airway pressure and was validated in paralyzed patients [5]. Thus, this equation may not be useful in a significant number of mechanically ventilated patients because the use of pressure-regulated ventilation has been increasing [10], and restricted use of neuromuscular blocking agents (NMBAs) is advocated due to their detrimental effects [11,12].

Recently, Becher et al. [13] developed an equation to calculate MP for patients under pressure-controlled ventilation (PCV) mode. This equation easily can be calculated with readily available parameters and may serve as a useful monitoring index in patients on PCV but has not been studied extensively, especially in patients undergoing ventilation with or without NMBA. We aimed to examine the association between ICU mortality and MP calculated using Becher's equation ( $MP_{PCV}$ ) in patients undergoing PCV and to investigate whether the use of NMBA affects this relationship.

## MATERIALS AND METHODS

#### **Design and Population**

This study was a retrospective analysis of a prospective Korean cohort that formed part of an international study [14]. The study protocol was approved by the Institutional Review Boards of all participating hospitals, and the need for informed consent was waived due to the non-interventional nature of the protocol. In 2016, 226 patients from 18 Korean ICUs participated in the Fourth International Study of Mechanical

#### KEY MESSAGES

- Mechanical power calculated using Becher's simplified equation (MP<sub>PCV</sub>) was significantly associated with intensive care unit (ICU) mortality in Korean patients on pressure-controlled ventilation.
- The association between MP<sub>PCV</sub> and ICU mortality was not affected by the use of neuromuscular blocking agent.

Ventilation of the VENTILA group [15]. That was a prospective, international, multicenter, non-interventional cohort study that enrolled adult patients who received invasive mechanical ventilation for at least 12 hours or non-invasive ventilation for more than 1 hour (https://clinicaltrials.gov/ct2/show/record/NCT02731898). Patients who were ventilated with PCV on day 1 were included in this study.

#### Data Collection Time

According to the parent study protocol, the day of initiation of mechanical ventilation was considered day 0, and the next day was considered day 1. Data were collected for the duration of mechanical ventilation or until day 28. For patients with invasive mechanical ventilation, blood gas analysis, ventilator mode and settings, and co-adjuvant therapies (sedatives, analgesics, NMBAs) were recorded daily at 8 am from day 1 (on day 0, data were collected within the first hour of starting mechanical ventilation). Documented ventilator settings were as follows: total and ventilator respiratory rates (per minute), tidal volume (mL), peak pressure (cm H<sub>2</sub>O), plateau pressure (cm H<sub>2</sub>O), and applied positive end-expiratory pressure (PEEP; cm H<sub>2</sub>O). We extracted and analyzed the day 1 data. Basal demographics, primary reasons for invasive mechanical ventilation, and discharge status were also collected.

#### Calculation of MP

We calculated the MP of patients under PCV using Becher's simplified equation:

$$MP_{PCV} = 0.098 \cdot RR \cdot V_{T} \cdot (\Delta P_{insp} + PEEP)$$

where  $\Delta P_{insp}$  is the change in airway pressure during inspiration (cm H<sub>2</sub>O), RR is respiratory rate (per minute), V<sub>T</sub> is tidal volume (L), and 0.0998 is a correction factor to convert the units to J/min [13]. If the total respiratory rate and ventilator



respiratory rate were different, the total respiratory rate value was entered in the RR term of the equation. Peak pressure was substituted for the last term of the equation.

#### **Statistical Analysis**

Patients were divided into survivors and non-survivors. Categorical variables are reported as number and percentage and were compared using Fisher's exact test or the chi-square test. Continuous variables are reported as median with interquartile range and were compared using the t-test or Mann-Whitney U-test, as appropriate. The normality of distributions was examined by the Kolmogorov-Smirnov test. Multivariable logistic regression analysis was performed to identify factors associated with ICU mortality. Variables with P-value less than 0.2 in univariable analysis and clinical variables shown to be important in previous studies (age and sex) were included in the multivariable analysis to investigate prognostic factors for ICU mortality. We also analyzed the interaction between  $MP_{PCV}$  and the use of NMBA to investigate an effect on the association between MP<sub>PCV</sub> and ICU mortality. Multivariable logistic regression analysis for ICU mortality was conducted and included the interaction term of  $\mathrm{MP}_{\mathrm{PCV}}$  and NMBA use. In addition, we conducted subgroup analysis by categorizing patients based on the presence or absence of spontaneous breathing effort, using respiratory rate as the criterion. When the ventilator respiratory rate was equal to the total respiratory rate, this was considered absence of spontaneous breathing effort (controlled ventilation group). Conversely, if the total respiratory rate was higher than the ventilator respiratory rate, this was considered presence of spontaneous breathing effort (spontaneous breathing group). The logistic regression results are reported as odds ratio (OR) of each variable with 95% confidence interval (CI). All tests were two-sided, and a P-value less than 0.05 was considered statistically significant. All statistical analyses were performed using IBM SPSS ver. 20.0 (IBM Corp.) and SAS ver. 9.4 (SAS Institute Inc.).

#### RESULTS

Two hundred twenty-three patients were treated with invasive mechanical ventilation on day 1, and PCV mode was applied to 160 of them.  $MP_{PCV}$  could be obtained for 125 patients. Of these, 87 survived and 38 died in the ICU (Figure 1). Overall, baseline characteristics were comparable between survivors and non-survivors (Table 1). Median age was 68 years, and two-thirds of patients were male. The majority of patients re-

ceived mechanical ventilation owing to acute respiratory failure. The use of analgesics was more frequent in non-survivors (72.4 % vs 92.1%, P=0.014). NMBA was used twice as frequently in non-survivors than survivors, but this difference was not significant (11.5% vs. 23.7%, P=0.081).

 $MP_{PCV}$  in the whole study population was 21.7 J/min and was significantly higher in non-survivors than survivors (17.6 J/min vs. 26.3 J/min, P<0.001). Overall tidal volume per predicted body weight (V<sub>T</sub>/PBW) was 7.2 mL/kg and was higher in non-survivors than survivors (7.0 ml/kg vs. 8.2 ml/kg, P=0.025). PEEP was similar in the two groups (Table 2).

Table 3 shows the results of logistic regression analysis. Analgesic use,  $MP_{PCV}$  and  $V_T/PBW$  were significantly associated with ICU mortality in univariable analysis and were included in the multivariable analysis. Age, sex, and other variables with p-value less than 0.2 in the univariable analysis were also included in multivariable analysis. Among them, only  $MP_{PCV}$  was significantly associated with ICU mortality (OR, 1.090; 95% CI, 1.029–1.155; P=0.003).

Regarding ICU mortality, there was no significant interaction



Figure 1. Study flowchart. PCV: pressure-controlled ventilation; ICU: intensive care unit.

Λ	

Table 1.	Baseline	characteristics
----------	----------	-----------------

Variable	All (n=125)	Survivor (n=87)	Non-survivor (n=38)	P-value
Age (yr)	68 (57–78)	68 (58–78)	66 (55–75)	0.425
Male	83 (66.4)	58 (66.7)	25 (65.8)	0.924
Weight (kg)	60.0 (50.0-68.0)	60.0 (49.0-68.0)	60.0 (53.0-68.5)	0.531
Height (cm)	165.0 (158.0–170.0)	164.0 (157.0–170.0)	165.5 (159.8–170.0)	0.503
BMI (kg/m <sup>2</sup> )	22.0 (19.0-24.0)	22.0 (19.0-25.0)	22.0 (20.0-24.0)	0.851
SAPS II	50 (40–61)	50 (42–63)	48.5 (36–60)	0.205
Primary reason for mechanical ventilation <sup>a)</sup>				0.435
Acute on chronic respiratory failure	8 (7.3)	5 (6.7)	3 (8.6)	
COPD	4 (3.6)	3 (4)	1 (2.9)	
Asthma	2 (1.8)	2 (2.7)	0	
Other chronic respiratory disease	2 (1.8)	0	2 (5.7)	
Acute respiratory failure	91 (82.7)	60 (80)	31 (88.6)	
ARDS	6 (5.5)	2 (2.7)	4 (11.4)	
Postoperative	0	0	0	
Congestive heart failure	11 (10.0)	9 (12.0)	2 (5.7)	
Aspiration	14 (12.7)	12 (16)	2 (5.7)	
Pneumonia	30 (27.3)	16 (21.3)	14 (40)	
Sepsis	20 (18.2)	12 (16)	8 (22.9)	
Trauma	1 (0.9)	1 (1.3)	0	
Cardiac arrest	5 (4.5)	5 (6.7)	0	
Other acute respiratory failure	4 (3.6)	3 (4)	1 (2.9)	
Coma	9 (8.2)	8 (10.7)	1 (2.9)	
Neuromuscular disease	2 (1.8)	2 (2.7)	0	
pH <sup>b)</sup>	7.36 (7.29–7.45)	7.38 (7.29–7.46)	7.34 (7.27–7.41)	0.230
$PaCO_2 (mm Hg)^{b}$	39.0 (31.0-46.0)	38.0 (31.0-51.0)	40.0 (31.0-43.3)	0.642
$PaO_2 (mm Hg)^{b)}$	87.0 (68.5–116.5)	85.0 (67.0-120.0)	89.5 (77.8–114.5)	0.239
PaO <sub>2</sub> /FiO <sub>2</sub> ratio <sup>b)</sup>	157 (106–230)	165 (114–256)	137 (95–211)	0.105
Analgesic <sup>b)</sup>	98 (78.4)	63 (72.4)	35 (92.1)	0.014
Sedative <sup>b)</sup>	86 (68.8)	57 (65.5)	29 (76.3)	0.231
NMBA <sup>b)</sup>	19 (15.2)	10 (11.5)	9 (23.7)	0.081

Values are presented as median (interquartile range) or number (%).

BMI: body mass index; SAPS: Simplified Acute Physiology Score; COPD: chronic obstructive pulmonary disease; ARDS: acute respiratory distress syndrome; PaCO<sub>2</sub>: partial pressure of carbon dioxide; PaO<sub>2</sub>: partial pressure of oxygen; FiO<sub>2</sub>: fraction of inspired oxygen; NMBA: neuromuscular blocking agent.

a) Data were not available in 12 patients among survivors and 3 patients among non-survivors; b) Data were collected at 8 am on day 1.

between the  $MP_{PCV}$  and the use of NMBA (P=0.579). In patients who were not treated with NMBA,  $MP_{PCV}$  was significantly associated with ICU mortality (OR, 1.081; 95% CI, 1.017–1.149; P=0.013). In patients who received NMBA, ICU mortality also tended to increase with higher  $MP_{PCV}$ , although significance was not achieved (OR, 1.125; 95% CI, 0.987–1.283; P=0.078) (Table 4).

In subgroup analysis, 51 patients (42.5%) were assigned to the controlled ventilation group. The  $MP_{PCV}$  was higher in non-survivors than survivors (21.6 J/min vs. 26.7 J/min, P=0.045) in the controlled ventilation group (Supplementary Table 1). In multivariable analysis, we observed a significant association between high  $MP_{PCV}$  and an increase in ICU mortality rate (OR 1.177, 95% CI, 1.030–1.344; P=0.016) (Supplementary Table 2). Similar results were found in the spontaneous breathing group (Supplementary Tables 3 and 4).

## DISCUSSION

In the present study, we found that  $\rm MP_{PCV}$  which can be calculated with universally monitored parameters at the bedside, was significantly associated with ICU mortality. In fact,  $\rm MP_{PCV}$ 



Table 2. Comparison of day 1 ventilator variables between survivors and non-survivors

Variable	All (n=125)	Survivor (n=87)	Non-survivor (n=38)	P-value
Mechanical power (J/min)	21.7 (14.3–26.6)	17.6 (12.7–23.7)	26.3 (20.7–33.9)	<0.001
V <sub>T</sub> /PBW (ml/kg)	7.2 (6.2–8.7)	7.0 (6.0–8.1)	8.2 (6.6–9.2)	0.025
PEEP (cm $H_2O)^{a}$	5.0 (5.0–8.0)	5.0 (5.0-8.0)	7.0 (5.0–8.0)	0.176

Values are presented as median (interquartile range).

V<sub>T</sub>/PBW: tidal volume per predicted body weight; PEEP: positive end-expiratory pressure.

a) Data were not available in one patient in survivors.

Table	3. Lo	aistic	rearession	analvsi	s for	intensive	care	unit	mortality

Verieble	Univariable and	alysis	Multivariable and	Multivariable analysis		
Variable	OR (95% CI)	P-value	OR (95% CI)	P-value		
Age (yr)	0.989 (0.963–1.016)	0.423	0.992 (0.961–1.026)	0.652		
Male	0.962 (0.430–2.150)	0.924	1.109 (0.441–2.789)	0.826		
BMI (kg/m <sup>2</sup> )	1.009 (0.915–1.113)	0.850	-	-		
SAPS II	0.985 (0.963-1.008)	0.205	-	-		
Primary reason for mechanical ventilation, acute respiratory ailure	1.937 (0.592–6.337)	0.274	-	-		
pH <sup>a)</sup>	0.127 (0.004-3.683)	0.230				
$PaCO_2 (mm Hg)^{a)}$	0.978 (0.948-1.009)	0.158	0.975 (0.938–1.013)	0.191		
$PaO_2 (mm Hg)^{a}$	1.003 (0.996–1.010)	0.426	-	-		
PaO <sub>2</sub> /FiO <sub>2</sub> ratio <sup>a)</sup>	0.997 (0.993-1.001)	0.129	1.000 (0.995-1.004)	0.844		
Analgesic <sup>a)</sup>	4.444 (1.249–15.816)	0.021	3.568 (0.834–15.256)	0.086		
Sedative <sup>a)</sup>	1.696 (0.711-4.043)	0.233	-	-		
NMBA <sup>a)</sup>	2.390 (0.882-6.474)	0.087	1.935 (0.606–6.178)	0.265		
Mechanical power (J/min) <sup>a)</sup>	1.105 (1.053–1.159)	<0.001	1.090 (1.029–1.155)	0.003		
V <sub>T</sub> /PBW <sup>a)</sup>	1.239 (1.022-1.502)	0.029	1.079 (0.835–1.395)	0.561		
PEEP (cm $H_2O)^{a)}$	1.086 (0.920-1.283)	0.329	-	-		

Variables with P-value less than 0.2 in univariable analysis and clinical variables with important meanings (age, sex) were included in the multivariable analysis. OR: odds ratio; CI: confidence interval; BMI: body mass index; SAPS: Simplified Acute Physiology Score;  $PaCO_2$ : partial pressure of carbon dioxide;  $PaO_2$ : partial pressure of oxygen; FiO<sub>2</sub>: fraction of inspired oxygen; NMBA: neuromuscular blocking agent;  $V_T/PBW$ : tidal volume per predicted body weight; PEEP: positive end-expiratory pressure.

a) Data were collected at 8 am on day 1.

Table 4. Association between  $MP_{PCV}$  and intensive care unit mortality according to the use of neuromuscular blocking agents

Variable	OR (95% CI)	P-value
Non-use of NMBA	1.081 (1.017–1.149)	0.013
Use of NMBA	1.125 (0.987–1.283)	0.078
Interaction of $MP_{PCV}$ and NMBA	-	0.579

A high OR (>1) indicates that as the  $MP_{PCV}$  increases, intensive care unit mortality also increases.

MP<sub>PCV</sub>: mechanical power obtained by Becher's equation; OR: odds ratio; CI: confidence interval; NMBA: neuromuscular blocking agent.

was the only independent predictor of ICU mortality in the multivariable logistic regression analysis. In addition, the association between  $MP_{PCV}$  and ICU mortality was not affected by

the use of NMBA, which suggests  $\mathrm{MP}_{\mathrm{PCV}}$  as a predictor of mortality regardless of the use of NMBA.

One of the most important findings in this study is that  $MP_{PCV}$  was associated with poor outcomes in Korean patients undergoing mechanical ventilation in PCV mode for respiratory failure. Although several studies have investigated the associations between MP and clinical outcomes, most of the studies have enrolled patients undergoing ventilation in volume-controlled mode and calculated MP using Gattinoni's equation. For example, in one study that reported an association between poor outcome and MP calculated with Gattinoni's equation, 90% of patients were treated with volume-controlled ventilation [16]. Other large-scale studies either did not provide information on ventilation modes [9,17,18] or included some patients who

# <mark>∧CC</mark>√

were ventilated in PCV mode [19] but used Gattinoni's equation to calculate MP. Since the use of pressure-regulated ventilation is increasing globally, especially in Korea, the results of our study can be clinically helpful [10,14,20]. Moreover, we used Becher's simplified equation, which is easy to calculate at the bedside because it incorporates only variables that are readily obtained from the ventilator.

Another notable aspect of this study is that we analyzed the effect of NMBA on the relationship between MP and ICU mortality. If we want to use MP as a surrogate for risk for VILI, it should be calculated with patients under passive conditions because published equations for MP cannot account for changes in values induced by spontaneous effort of the patient. In Becher's equation, the effect of spontaneous breathing efforts on calculated  $MP_{PCV}$  can be variable. Tidal volume should increase if there is spontaneous breathing effort at the same peak inspiratory pressure; at the same time, lower peak inspiratory pressure is needed to achieve the same tidal volume when the patient has spontaneous breathing efforts. However, only a subset of patients treated with mechanical ventilation receives NMBA, and this proportion is gradually decreasing [10,21], which would lessen the clinical usefulness of MP as a bedside monitoring tool or ventilator-adjustment target if it has value only in passive patients. Indeed, only 15% of participants received NMBA in our study. Previous studies on the association between MP and clinical outcomes have either tried to exclude patients with spontaneous efforts by excluding patients with a higher measured respiratory rate than the set respiratory rate [18,19] or included patients with spontaneous breathing in their analysis as well [17]. Our study is meaningful in that we assessed spontaneous breathing (or paralysis) based on NMBA use and subsequently analyzed the impact of NMBA use on the association between MP and clinical outcomes. We demonstrated that MP<sub>PCV</sub> was independently associated with ICU mortality irrespective of the use of NMBA. Serpa Neto et al. [9] also found no interaction between NMBA use and clinical outcomes, although they used the MP equation, which is based on volume-controlled ventilation.

Interestingly,  $MP_{PCV}$  was the only variable associated with ICU mortality in multivariable analysis.  $V_T/PBW$  was significantly higher in non-survivors than survivors and was significantly associated with ICU mortality in univariable analysis, but this significance did not persist in multivariable analysis. This should not be interpreted to mean that  $V_T/PBW$  does not affect clinical outcomes. It is well-known that different variables used to calculate MP have different impacts on MP,

and tidal volume is one of the variables with the strongest influence. For example, tidal volume, PEEP, and respiratory rate can all contribute to MP; however, when each respective value is doubled, MP increases by 4 times, 2 times, and 1.4 times [5]. Moreover, the effect of  $V_T$ /PBW would be diluted in studies involving a relatively small number of patients if a low tidal volume ventilation strategy is being universally applied [21].

There are several limitations to this study. First and foremost, MP<sub>PCV</sub> as calculated by Becher's equation may not represent the true MP delivered to the patient by the ventilator. Since Becher's equation calculates MP under the assumption of an ideal "square wave," this assumption may not hold true in a spontaneously breathing patient. Moreover, as spontaneous efforts increase, the airway pressure will tend to deviate increasingly from this assumption. Also when there is spontaneous effort, tidal volume will increase at the fixed inspiratory pressure, which might result in overestimation of the MP delivered by the ventilator. However, distending pressure and tidal volumes created by spontaneous effort could also be damaging [22], and MP<sub>PCV</sub> calculated by Becher's equation may prove to have prognostic significance even in spontaneously breathing patients, as was shown in this study. Further studies are needed on this subject. Second, since this was a retrospective study, we are not able to exclude the possibility of residual confounding factors. Third, only  $MP_{PCV}$  on day 1 was used for analysis. Thus, we did not evaluate  $MP_{PCV}$  on days other than day 1 or changes in  $MP_{PCV}$  and their potential impact on patient outcomes. Fourth, due to the relatively small size of the study population, caution should be used when generalizing the results, as there may be other potentially important differences that were not found due to lack of statistical power. However, all patients starting mechanical ventilation were prospectively included from 18 Korean ICUs during the study period according to an established protocol. Fifth, the presence or absence of NMBA use may not completely distinguish passive ventilation and spontaneous breathing. Finally, we did not compare the equation to calculate MP in this study with other equations proposed to calculate MP in PCV mode [13,23,24]. However, other equations are more complex, and we wanted to evaluate the usefulness of Becher's simplified equation because it can be readily applied at the bedside.

In conclusion,  $MP_{PCV}$  was associated with ICU mortality in patients mechanically ventilated with PCV mode regardless of NMBA use. High  $MP_{PCV}$  during the initial phase of mechanical ventilation could be predictive of a poor prognosis. Further prospective studies are needed to establish a specific cut-off



## ADDITIONAL INFORMATION

<sup>1</sup>Division of Pulmonary, Allergy, and Critical Care Medicine, Department of Internal Medicine, Korea University Guro Hospital, Korea University College of Medicine, Seoul, Korea

- <sup>2</sup>Division of Pulmonary and Critical Care Medicine, Department of Internal Medicine, Seoul National University Hospital, Seoul National University College of Medicine, Seoul, Korea
- <sup>3</sup>Division of Pulmonary and Critical Care Medicine, Department of Internal Medicine, Inje University Ilsan Paik Hospital, Inje University College of Medicine, Goyang, Korea
- <sup>4</sup>Department of Internal Medicine, Daegu Catholic University Medical Center, Daegu Catholic University School of Medicine, Daegu, Korea
- <sup>b</sup>Division of Pulmonology, Department of Internal Medicine, Institute of Chest Disease, Severance Hospital, Yonsei University College of Medicine, Seoul, Korea
- <sup>6</sup>Division of Pulmonary, Allergy, and Critical Care Medicine, Department of Internal Medicine, Pusan National University Yangsan Hospital, Pusan National University School of Medicine, Yangsan, Korea
- <sup>7</sup>Divison of Pulmonary, Allergy, and Critical Care Medicine, Department of Internal Medicine, Yonsei University Wonju Severance Christian Hospital, Yonsei University Wonju College of Medicine, Wonju, Korea
- <sup>8</sup>Division of Pulmonary, Allergy, and Critical Care Medicine, Department of Internal Medicine, Hallym University Sacred Heart Hospital, Hallym University College of Medicine, Anyang, Korea
- <sup>9</sup>Division of Pulmonary, Allergy, and Critical Care Medicine, Department of Internal Medicine, Hallym University Kangdong Sacred Heart Hospital, Seoul, Korea
- <sup>10</sup> Division of Pulmonary and Critical Care Medicine, Department of Internal Medicine, Dongguk University Ilsan Hospital, Goyang, Korea
- <sup>11</sup>Division of Pulmonary, Allergy, and Critical Care Medicine, Department of Internal Medicine, Hallym University Kangnam Sacred Heart Hospital, Seoul, Korea
- <sup>12</sup>Division of Pulmonary, Allergy, and Critical Care Medicine, Department of Internal Medicine, Pusan National University School of Medicine, Busan, Korea
- <sup>13</sup>Division of Pulmonary and Critical Care Medicine, Department of Internal Medicine, Seoul National University Bundang Hospital, Seongnam, Korea
- <sup>14</sup> Division of Pulmonary and Critical Care Medicine, Department of Internal Medicine, Ewha Womans University College of Medicine, Seoul, Korea
- <sup>15</sup>Division of Respiratory Disease and Critical Care Medicine, Department of Internal Medicine, Jeonbuk National University Medical School and Hospital, Jeonju, Korea
- <sup>16</sup>Division of Pulmonary and Critical Care Medicine, Department of Internal Medicine, Asan Medical Center, University of Ulsan College of Medicine, Seoul, Korea
- <sup>17</sup>Department of Internal Medicine, Myongji Hospital, Hanyang University College of Medicine, Goyang, Korea
- <sup>18</sup>Division of Pulmonary, Allergy, and Critical Care Medicine, Department of Internal Medicine, Chuncheon Sacred Heart Hospital, Hallym University Medical Center, Chuncheon, Korea
- <sup>19</sup>Department of Critical Care Medicine, Sungkyunkwan University School of Medicine, Kangbuk Samsung Hospital, Seoul, Korea
- <sup>20</sup> Division of Pulmonary and Critical Care Medicine, Department of Medicine, Samsung Medical Center, Sungkyunkwan University School of Medicine, Seoul, Korea

## CONFLICT OF INTEREST

Kwangha Lee is an editorial board member of the journal but was not involved in the peer reviewer selection, evaluation, or decision process of this article. No other potential conflict of interest relevant to this article was reported.

#### FUNDING

None.

#### ACKNOWLEDGMENTS

Part of this work was presented at the 26th Congress of the Asian Pacific Society of Respirology (APSR 2022). The authors would like to thank all members of the Korean Study Group on Respiratory Failure (KOSREF). They would also like to thank Soon-Young Hwang from the Department of Biostatistics at Korea University College of Medicine for providing valuable advice on statistical analysis.

## ORCID

Jae Kyeom Sim Sang-Min Lee Hyung Koo Kang Kyung Chan Kim Young Sam Kim Yun Seong Kim Won-Yeon Lee Sunghoon Park So Young Park Ju-Hee Park Yun Su Sim Kwangha Lee Yeon loo Lee Jin Hwa Lee Heung Bum Lee Chae-Man Lim Won-Il Choi Ji Young Hong Gee Young Suh

https://orcid.org/0000-0003-0873-2807 https://orcid.org/0000-0002-1388-9318 https://orcid.org/0000-0001-9671-0944 https://orcid.org/0000-0001-5697-9674 https://orcid.org/0000-0001-9656-8482 https://orcid.org/0000-0003-4328-0818 https://orcid.org/0000-0002-5461-6770 https://orcid.org/0000-0001-7004-6985 https://orcid.org/0000-0003-2718-0518 https://orcid.org/0000-0002-0932-121X https://orcid.org/0000-0002-3746-4947 https://orcid.org/0000-0001-9878-201X https://orcid.org/0000-0001-7697-4272 https://orcid.org/0000-0003-0843-9862 https://orcid.org/0000-0002-8267-8434 https://orcid.org/0000-0001-5400-6588 https://orcid.org/0000-0001-7705-0098 https://orcid.org/0000-0002-3132-7706 https://orcid.org/0000-0001-5473-1712

## AUTHOR CONTRIBUTIONS

Conceptualization: JKS, GYS. Data curation: all authors. For-

mal analysis: JKS, GYS. Visualization: JKS. Supervision: GYS. Writing–original draft: JKS. Writing–review & editing: SML, HKK, KCK, YSK, YSK, WYL, SP, SYP, JHP, YSS, KL, YJL, JHL, HBL, CML, WIC, JYH, WJS, GYS. All authors read and agreed to the published version of the manuscript.

## SUPPLEMENTARY MATERIALS

Supplementary materials can be found via https://doi. org/10.4266/acc.2023.00871.

#### REFERENCES

- Slutsky AS, Ranieri VM. Ventilator-induced lung injury. N Engl J Med 2013;369:2126-36.
- 2. Cho YJ, Moon JY, Shin ES, Kim JH, Jung H, Park SY, et al. Clinical practice guideline of acute respiratory distress syndrome. Korean J Crit Care Med 2016;31:76-100.
- **3.** Hotchkiss JR, Blanch L, Murias G, Adams AB, Olson DA, Wangensteen OD, et al. Effects of decreased respiratory frequency on ventilator-induced lung injury. Am J Respir Crit Care Med 2000;161(2 Pt 1):463-8.
- 4. Chiumello D, Carlesso E, Brioni M, Cressoni M. Airway driving pressure and lung stress in ARDS patients. Crit Care 2016;20:276.
- 5. Gattinoni L, Tonetti T, Cressoni M, Cadringher P, Herrmann P, Moerer O, et al. Ventilator-related causes of lung injury: the mechanical power. Intensive Care Med 2016;42:1567-75.
- 6. Cressoni M, Gotti M, Chiurazzi C, Massari D, Algieri I, Amini M, et al. Mechanical power and development of ventilator-induced lung injury. Anesthesiology 2016;124:1100-8.
- 7. Vassalli F, Pasticci I, Romitti F, Duscio E, Aßmann DJ, Grünhagen H, et al. Does iso-mechanical power lead to iso-lung damage?: an experimental study in a porcine model. Anesthesiology 2020;132:1126-37.
- **8.** Scharffenberg M, Wittenstein J, Ran X, Zhang Y, Braune A, Theilen R, et al. Mechanical power correlates with lung inflammation assessed by positron-emission tomography in experimental acute lung injury in pigs. Front Physiol 2021;12:717266.
- **9.** Serpa Neto A, Deliberato RO, Johnson AE, Bos LD, Amorim P, Pereira SM, et al. Mechanical power of ventilation is associated with mortality in critically ill patients: an analysis of patients in two observational cohorts. Intensive Care Med 2018;44:1914-22.
- **10.** Esteban A, Frutos-Vivar F, Muriel A, Ferguson ND, Peñuelas O, Abraira V, et al. Evolution of mortality over time in patients receiving mechanical ventilation. Am J Respir Crit Care Med

2013;188:220-30.

 Oh Y, Kang Y, Lee K. Development of a prognostic scoring system in patients with pneumonia requiring ventilator care for more than 4 days: a single-center observational study. Acute Crit Care 2021;36:46-53.

- Murray MJ, DeBlock H, Erstad B, Gray A, Jacobi J, Jordan C, et al. Clinical practice guidelines for sustained neuromuscular blockade in the adult critically ill patient. Crit Care Med 2016;44:2079-103.
- Becher T, van der Staay M, Schädler D, Frerichs I, Weiler N. Calculation of mechanical power for pressure-controlled ventilation. Intensive Care Med 2019;45:1321-3.
- 14. Sim JK, Lee SM, Kang HK, Kim KC, Kim YS, Kim YS, et al. Change in management and outcome of mechanical ventilation in Korea: a prospective observational study. Korean J Intern Med 2022;37:618-30.
- Esteban A, Anzueto A, Frutos F, Alía I, Brochard L, Stewart TE, et al. Characteristics and outcomes in adult patients receiving mechanical ventilation: a 28-day international study. JAMA 2002;287:345-55.
- 16. Chi Y, Zhang Q, Yuan S, Zhao Z, Long Y, He H. Twenty-fourhour mechanical power variation rate is associated with mortality among critically ill patients with acute respiratory failure: a retrospective cohort study. BMC Pulm Med 2021;21:331.
- 17. Urner M, Jüni P, Hansen B, Wettstein MS, Ferguson ND, Fan E. Time-varying intensity of mechanical ventilation and mortality in patients with acute respiratory failure: a registry-based, prospective cohort study. Lancet Respir Med 2020;8:905-13.
- 18. Tonna JE, Peltan ID, Brown SM, Grissom CK, Presson AP, Herrick JS, et al. Positive end-expiratory pressure and respiratory rate modify the association of mechanical power and driving pressure with mortality among patients with acute respiratory distress syndrome. Crit Care Explor 2021;3:e0583.
- 19. van Meenen DM, Serpa Neto A, Paulus F, Merkies C, Schouten LR, Bos LD, et al. The predictive validity for mortality of the driving pressure and the mechanical power of ventilation. Intensive Care Med Exp 2020;8(Suppl 1):60.
- 20. Jeong BH, Suh GY, An JY, Park MS, Lee JH, Lee MG, et al. Clinical demographics and outcomes in mechanically ventilated patients in Korean intensive care units. J Korean Med Sci 2014;29:864-70.
- **21.** Peñuelas O, Muriel A, Abraira V, Frutos-Vivar F, Mancebo J, Raymondos K, et al. Inter-country variability over time in the mortality of mechanically ventilated patients. Intensive Care Med 2020;46:444-53.
- 22. Brochard L, Slutsky A, Pesenti A. Mechanical ventilation to



minimize progression of lung injury in acute respiratory failure. Am J Respir Crit Care Med 2017;195:438-42.

- 23. van der Meijden S, Molenaar M, Somhorst P, Schoe A. Calculating mechanical power for pressure-controlled ventilation. Intensive Care Med 2019;45:1495-7.
- 24. Trinkle CA, Broaddus RN, Sturgill JL, Waters CM, Morris PE. Simple, accurate calculation of mechanical power in pressure controlled ventilation (PCV). Intensive Care Med Exp 2022;10:22.