

Effect of prone positioning on gas exchange according to lung morphology in patients with acute respiratory distress syndrome

Na Young Kim¹, Si Mong Yoon², Jimyung Park¹, Jinwoo Lee¹, Sang-Min Lee^{1,2}, Hong Yeul Lee²

¹Division of Pulmonary and Critical Care Medicine, Department of Internal Medicine and ²Department of Critical Care Medicine, Seoul National University Hospital, Seoul, Korea

Background: There are limited data on the clinical effects of prone positioning according to lung morphology. We aimed to determine whether the gas exchange response to prone positioning differs according to lung morphology.

Methods: This retrospective study included adult patients with moderate-to-severe acute respiratory distress syndrome (ARDS). The lung morphology of ARDS was assessed by chest computed tomography scan and classified as "diffuse" or "focal." The primary outcome was change in partial pressure of arterial oxygen to fraction of inspired oxygen (PaO₂/FiO₂) ratio after the first prone positioning session: first, using the entire cohort, and second, using subgroups of patients with diffuse ARDS matched 2 to 1 with patients with focal ARDS at baseline.

Results: Ninety-five patients were included (focal ARDS group, 23; diffuse ARDS group, 72). Before prone positioning, the focal ARDS group showed worse oxygenation than the diffuse ARDS group (median PaO₂/FiO₂ ratio, 79.9 mm Hg [interquartile range (IQR)], 67.7–112.6 vs. 104.0 mm Hg [IQR, 77.6–135.7]; P=0.042). During prone positioning, the focal ARDS group showed a greater improvement in the PaO₂/FiO₂ ratio than the diffuse ARDS group (median, 55.8 mm Hg [IQR, 11.1–109.2] vs. 42.8 mm Hg [IQR, 11.6–83.2]); however, the difference was not significant (P=0.705). Among the PaO₂/FiO₂-matched cohort, there was no significant difference in change in PaO₂/FiO₂ ratio after prone positioning between the groups (P=0.904).

Conclusions: In patients with moderate-to-severe ARDS, changes in PaO₂/FiO₂ ratio after prone positioning did not differ according to lung morphology. Therefore, prone positioning can be considered as soon as indicated, regardless of ARDS lung morphology.

Key Words: acute respiratory distress syndrome; dynamic compliance of lung; lung morphology; oxygenation; prone positioning

Original Article

Received: March 16, 2022

Revised: May 9, 2022

Accepted: May 13, 2022

Corresponding author

Hong Yeul Lee
Department of Critical Care
Medicine, Seoul National University
Hospital, 101 Daehak-ro, Jongno-gu,
Seoul 03080, Korea
Tel: +82-2-2072-2957
Fax: +82-2-762-9662
E-mail: takumama@naver.com

Copyright © 2022 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/licenses/by-nc/4.0/>) which permits
unrestricted noncommercial use, distribution,
and reproduction in any medium, provided the
original work is properly cited.

INTRODUCTION

Acute respiratory distress syndrome (ARDS) is a clinical syndrome of hypoxemic respiratory failure and is associated with a high mortality rate. Ventilator-induced lung injury can occur

in patients receiving invasive mechanical ventilation and can contribute to multiple organ dysfunction [1]. Several lung protective ventilation strategies and adjunctive management have been proposed to reduce the deleterious consequences of lung injury [2-5]. Prone positioning, one of several interventions, has been implemented widely in patients with moderate-to-severe ARDS to reduce mortality [6]. Additionally, a recent study from our institution found that an improvement in oxygenation after prone positioning is a useful predictor of survival [7]. Prone positioning minimizes regional differences in lung aeration, compliance, and shear strain, leading to clinically significant improvements in oxygenation [8].

The value of precision medicine has recently emerged, and attempts have been made to identify meaningful subgroups of critical illness syndrome considering the heterogeneity in the field of critical care medicine [9]. Calfee et al. [10] used inflammatory biomarkers, such as interleukin-6 and interferon-gamma, to identify two biologically distinct groups and found that the reactive phenotype was associated with worse clinical outcomes. Moreover, ARDS can be subdivided using clinical imaging as a surrogate marker of lung recruitment potential. The lung imaging morphology for ventilator settings in an ARDS study (LIVE study) conducted in France suggested that an approach based on the lung morphology of ARDS could reduce mortality. However, there was a limitation in that the misclassification rate of lung morphology was high due to the small proportion of patients with available computed tomography (CT) scans [11]. This study investigated whether the improvement in oxygenation after prone positioning differed between lung morphologies as assessed by CT scan.

MATERIALS AND METHODS

Study Design and Patients

This was a retrospective cohort study in the medical intensive care unit (ICU) at Seoul National University Hospital, a tertiary care referral hospital in Seoul, Korea. The requirement of written informed consent was waived due to the retrospective nature of the study.

We reviewed the medical records of adult patients aged >18 years who were diagnosed with moderate-to-severe ARDS and underwent prone positioning between January 1, 2014, and May 31, 2021. According to the Berlin definition [6,12], moderate-to-severe ARDS is defined as partial pressure of arterial oxygen to fraction of inspired oxygen ($\text{PaO}_2/\text{FiO}_2$) ratio <150 mm Hg with positive end-expiratory pressure (PEEP) ≥ 5 cm of

KEY MESSAGES

- Changes in the partial pressure of arterial oxygen to fraction of inspired oxygen ($\text{PaO}_2/\text{FiO}_2$) ratio and respiratory compliance after prone position did not differ according to the lung morphology of acute respiratory distress syndrome (ARDS).
- Prone position can be considered as soon as possible, regardless of the morphological phenotype in patients with moderate-to-severe ARDS.
- In a multivariable Cox proportional hazard regression analysis of patients with moderate-to-severe ARDS, the improvement in $\text{PaO}_2/\text{FiO}_2$ ratio after prone position was independently associated with mortality.

water. We also assessed lung morphology using chest CT scans performed within 1 week of the first prone session. Patients who were in prone position for less than 12 hours were excluded from this study [13].

Radiologic Findings

We assessed the lung morphology of ARDS by chest CT and classified it as “diffuse” or “focal.” The lung morphology characterization of ARDS was defined as (1) diffuse ARDS (widespread loss of lung aeration distribution throughout the lungs or uneven lung attenuation) and (2) focal ARDS (predominant loss of aeration in the lower lobar distribution or gravitationally-dependent areas) [14,15]. The chest CT findings were assessed visually by three independent readers, and two physicians (NYK and SMY) blindly reviewed the CT images. Any disagreement between the two physicians was reassessed under the supervision of a third blinded physician (HYL).

Prone Positioning Methods

We have several devices in place to reduce chest wall pressure during prone positioning, and this pressure can be minimized by padding all facial areas in contact with the bed. We placed foam dressings on the following areas before prone positioning: facial cheeks, shoulders, anterior iliac spine, and knees to protect from skin lesions. After changing to prone position, the chest and the ipsilateral leg were supported with a pillow. One arm was raised, with the head rotated toward the raised arm and the opposite arm positioned alongside the body, in the Swimmer’s position. We tilted the patient into reverse Trendelenburg position and continued lateral repositioning every 2 hours [16].

Study Outcomes and Data Collection

The primary outcome was change in $\text{PaO}_2/\text{FiO}_2$ ratio after the first prone positioning session. Secondary outcomes were 28-day mortality, ICU mortality, and changes in dynamic lung compliance (C_{dyn}) after the first prone positioning session. The $\text{PaO}_2/\text{FiO}_2$ ratio and C_{dyn} were evaluated at three time points for each patient by collecting the results of arterial blood gas analysis and ventilator setting at the time of blood tests at (1) baseline, before initiation of prone positioning; (2) time P8-12, approximately 8–12 hours after initiation of prone positioning; and (3) time S4-12, approximately 4–12 hours after resuming the supine position. The driving pressure was defined as the difference between plateau pressure and PEEP. If not recorded, peak inspiratory pressure was assumed to be equal to plateau pressure in pressure-controlled ventilation mode [17].

Statistical Analysis

Continuous variables were presented as means and standard deviations or medians and interquartile ranges (IQRs), while categorical variables were reported as numbers and percentages. All variables were compared between the diffuse and focal ARDS groups using the chi-square test or Student t-test, as appropriate. In primary and secondary outcomes, we performed a 2 to 1 matching procedure with the nearest-neighbor method without replacement to balance the two groups by $\text{PaO}_2/\text{FiO}_2$ ratio. Cox proportional hazard regression models were used to evaluate the effects of different variables on 28-day mortality in patients with ARDS who underwent prone positioning. All variables found to be significant ($P < 0.1$) in the univariable analysis were entered into a multivariable Cox regression model to avoid model overfitting and comply with the rule of thumb [18]. The results are presented as hazard ratios (HRs) with 95% confidence intervals (CIs). A two-way repeated-measures analysis of variance was applied to compare the extent of changes in $\text{PaO}_2/\text{FiO}_2$ from baseline to time S4-12. A two-tailed P-value less than 0.05 was considered statistically significant. All statistical analyses were performed using R (version 4.1.2) in R Studio (version 1.4.1717; R Foundation, Vienna, Austria).

RESULTS

Baseline Characteristics of the Patients

During the study period, we included 245 patients who underwent prone positioning to treat ARDS. Of these, 27 pa-

tients who were in prone position for less than 12 hours were excluded, and their median duration of prone positioning was 0.6 hours (IQR, 0.2–4.1 hours). A total of 95 patients had undergone chest CT within 1 week of the first prone session (Figure 1). In addition, 14 patients had undergone CT scans during mechanical ventilation, and 81 patients received CT scans before admission to the ICU. CT scan was used to assess lung morphology and classify 23 patients (24.2%) in the focal ARDS group and 72 patients (75.8%) in the diffuse ARDS group. Interobserver agreement was assessed by the kappa coefficient and was moderate (kappa coefficient, 0.58). The two groups had similar baseline characteristics (Table 1). The mean age was 65.1 ± 13.7 years, and 66 patients (69.5%) were males. The mean body mass index was $23.6 \pm 3.9 \text{ kg/m}^2$, and the main cause of ARDS was pulmonary (97.9%). Baseline vital signs and laboratory results before initiation of prone positioning did not differ between the two groups (Supplementary Table 1). The three scoring systems for prediction of mortality in critically ill patients did not differ between the two groups. The comorbidity was similar between the groups, but more than half of the included patients were diagnosed with malignancies due to the characteristics of the tertiary hospital.

The results of ventilator settings and arterial blood gas analysis at baseline did not differ significantly according to lung morphology, except for $\text{PaO}_2/\text{FiO}_2$ ratio (Table 2). Patients with

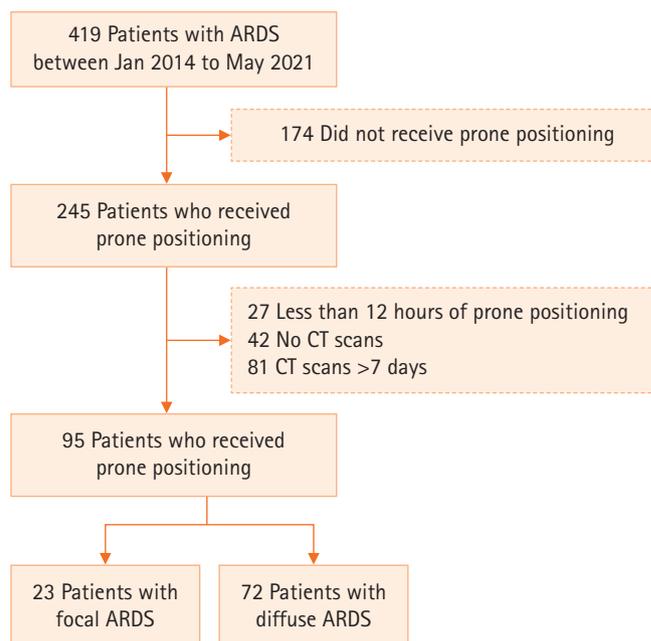


Figure 1. Flowchart of the study population. ARDS: acute respiratory distress syndrome; CT: computed tomography.

Table 1. Characteristics of the patients at baseline

Variable	Diffuse ARDS (n=72)	Focal ARDS (n=23)	P-value
Age (yr)	65.6±12.5	63.3±17.2	0.566
Male	48 (66.7)	18 (78.3)	0.429
Body mass index (kg/m ²)	23.7±3.9	23.2±4.2	0.589
Cause of ARDS			>0.999
Pulmonary	70 (97.2)	23 (100.0)	
Extrapulmonary	2 (2.8)	0	
Comorbidity			
Cardiovascular disease	11 (15.3)	4 (17.4)	>0.999
Diabetes mellitus	21 (29.2)	12 (52.2)	0.077
COPD	5 (6.9)	1 (4.3)	>0.999
Moderate to severe CKD ^a	7 (9.7)	1 (4.3)	0.706
Chronic liver disease	9 (12.5)	0	0.170
Solid tumor	32 (44.4)	8 (34.8)	0.566
Hematological malignancy	11 (15.3)	9 (39.1)	0.032
Connective tissue disease	10 (13.9)	1 (4.3)	0.384
Chronic neurologic disease	5 (6.9)	1 (4.3)	>0.999
Charlson comorbidity index	6.1±3.1	5.2±2.8	0.226
SOFA score	12.0±3.0	12.7±2.9	0.325
APACHE II score	26.3±6.6	28.1±7.8	0.282
SAPS II	58.7±15.5	63.1±14.4	0.226

Values are presented as mean±standard deviation or number (%).

ARDS: acute respiratory distress syndrome; COPD: chronic obstructive pulmonary disease; CKD: chronic kidney disease; SOFA: Sequential Organ Failure Assessment; APACHE: Acute Physiology and Chronic Health Evaluation; SAPS: Simplified Acute Physiology Score.

^aModerate CKD was defined as creatinine >3 mg/dL (0.27 mmol/L), and severe CKD was defined on dialysis, status post kidney transplant, and uremia.

Table 2. Pre-prone ventilator settings and arterial blood gas measurements

Variable	Diffuse ARDS (n=72)	Focal ARDS (n=23)	P-value
Ventilator setting			
Tidal volume (ml/kg PBW)	6.7±1.5	7.1±1.8	0.283
Respiratory rate (breaths/min)	24.4±5.0	24.4±4.9	0.974
PEEP (cm H ₂ O)	10.0 (8.0–12.0)	8.0 (7.0–10.5)	0.243
Driving pressure (cm H ₂ O)	16.9±4.9	17.0±4.8	0.954
FiO ₂	0.8 (0.6–1.0)	0.9 (0.7–1.0)	0.402
Total minute ventilation (L/min)	9.2 (8.0–10.0)	9.2 (8.4–11.2)	0.435
Dynamic compliance of lung (ml/cm H ₂ O)	21.3 (17.2–28.5)	26.0 (20.0–29.7)	0.195
Arterial-blood gas			
pH	7.3±0.1	7.3±0.1	0.796
PaO ₂ (mm Hg)	76.5 (64.9–91.0)	68.0 (62.2–76.5)	0.081
PaO ₂ /FiO ₂ (mm Hg)	104.0 (77.6–135.7)	79.9 (67.7–112.6)	0.042
PaCO ₂ (mm Hg)	47.0 (37.8–51.5)	44.7 (37.6–53.8)	0.924
Bicarbonate (mmol/L)	23.7 (21.0–27.0)	23.3 (22.1–25.9)	0.924
Lactate (mmol/L)	2.5 (1.7–3.3)	2.6 (2.0–4.4)	0.490

Values are presented as mean±standard deviation or median (interquartile range).

ARDS: acute respiratory distress syndrome; PBW: predicted body weight; PEEP: positive end expiratory pressure; FiO₂: fraction of inspired oxygen; PaO₂: partial pressure of oxygen; PaCO₂: partial pressure of carbon dioxide.

focal ARDS showed worse oxygenation than those with diffuse ARDS (median PaO₂/FiO₂ ratio, 79.9 mm Hg [IQR, 67.7–112.6] vs. 104.0 mm Hg [IQR, 77.6–135.7]; P=0.04). Before prone positioning, 98.9% of the patients received pressure-controlled ventilation. The mean tidal volume was 6.8±1.6 mL/kg predicted body weight, and the median PEEP was 10.0 cm H₂O (IQR, 8.0–11.5 cm H₂O). Adjunctive therapy during prone positioning sessions did not differ between the groups (Table 3). At the initiation of prone positioning, all patients received antibiotic therapy. Neuromuscular blockade, systemic steroids, vasopressors, inhaled nitric oxide, and renal replacement therapy were used in 93.7%, 90.5%, 74.7%, 21.1%, and 14.7% of the patients, respectively.

Oxygenation and Dynamic Compliance of Lung Response to First Prone Positioning

The median number of prone sessions was two per patient (IQR, 1–4), the median duration of the first prone session was 17.5 hours (IQR, 16.3–20.0 hours), and the median interval from intubation to initiation of the first prone positioning was 6.8 hours (IQR, 13.0–55.4 hours). These values did not differ significantly between the two groups (Table 3).

Changes in PaO₂/FiO₂ ratio after the first prone positioning session are shown in Figure 2. Measurement of the PaO₂/FiO₂ ratio at time P8–12 and time S4–12 was performed at a median of 10.2 hours (IQR, 9.0–11.4 hours) after the initiation

Table 3. Characteristics of adjunctive therapies and prone positioning sessions

Variable	Diffuse ARDS (n=72)	Focal ARDS (n=23)	P-value
Adjunctive therapy			
Neuromuscular blocker ^a	68 (94.4)	21 (91.3)	0.630
Glucocorticoids ^a	66 (91.7)	20 (87.0)	0.683
Vasopressors ^a	54 (75.0)	17 (73.9)	>0.999
Inhaled nitric oxide ^a	12 (16.7)	8 (34.8)	0.080
Renal replacement therapy ^a	12 (16.7)	2 (8.7)	0.506
Cross over to extracorporeal membrane oxygenation	1 (1.4)	0	>0.999
Prone positioning session			
Total number of sessions of prone positioning (day)	2.0 (1.0–4.0)	2.0 (1.0–2.0)	0.153
Median duration of prone positioning per session (hr)	17.5 (16.3–19.1)	19.0 (16.4–19.6)	0.192
Duration of the first prone positioning session (hr)	17.5 (16.1–20.0)	18.3 (16.4–19.9)	0.417
Interval between intubation and the first prone positioning session (day)	1.1 (0.6–2.4)	1.2 (0.5–2.1)	0.751

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

ARDS: acute respiratory distress syndrome.

^aDuring the first prone positioning session.

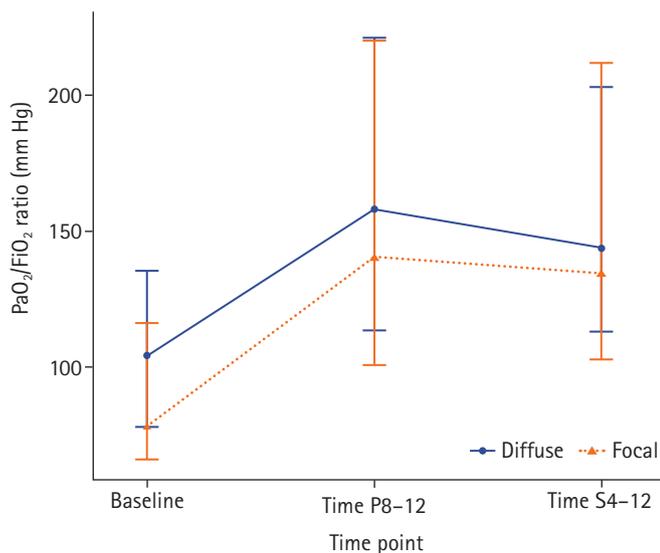


Figure 2. Median and interquartile range (error bars) of changes in the partial pressure of arterial oxygen to fraction of inspired oxygen ($\text{PaO}_2/\text{FiO}_2$) ratio during the first session of prone positioning. Time P8–12: approximately 8–12 hours after initiation of prone positioning; Time S4–12: approximately 4–12 hours after resuming the supine position.

of prone positioning and at 8.2 hours (IQR, 6.0–10.0 hours) after changing to supine position, respectively. The $\text{PaO}_2/\text{FiO}_2$ ratio was higher at time P8–12 in the diffuse ARDS group than in the focal ARDS group (median $\text{PaO}_2/\text{FiO}_2$ ratio, 157.9 mm Hg [IQR, 113.3–222.2] vs. 134.7 mm Hg [IQR, 99.2–217.5]; $P=0.39$). A detailed description of the ventilator settings and arterial blood gases during the first prone positioning session is provided according to lung morphology in [Supplementary](#)

Table 2. When comparing baseline and time P8–12, the absolute improvement in $\text{PaO}_2/\text{FiO}_2$ ratio was greater in patients with focal ARDS (median, 55.8 mm Hg [IQR, 11.1–109.2]) than in patients with diffuse ARDS (median, 42.8 mm Hg [IQR, 11.6–83.2]) ([Table 4](#)). However, the difference between the groups was not significant ($P=0.71$). After the patients returned to the supine position, the change in $\text{PaO}_2/\text{FiO}_2$ ratio from baseline was higher in patients with focal ARDS (median, 43.3 mm Hg [IQR, 24.0–98.0]) than in patients with diffuse ARDS (median, 41.4 mm Hg [IQR, 3.8–88.3]). The difference between the groups was not significant ($P=0.42$). Among the $\text{PaO}_2/\text{FiO}_2$ -matched cohort, there was no significant difference in change in $\text{PaO}_2/\text{FiO}_2$ ratio from baseline between the two groups at P8–12 and S4–12. Meanwhile, the change in dynamic compliance of the lung from baseline was not significantly different between the focal group and the diffuse group (median, -1.1 ml/cm H_2O [IQR, -4.6 to 4.6] vs. -1.0 ml/cm H_2O [IQR, -4.0 to 1.5]; $P=0.36$) at time P8–12. Additionally, at time S4–12, there were no significant differences in the improvement in dynamic compliance of the lung from baseline between the two groups (median, 2.3 ml/cm H_2O [IQR, -0.9 – 4.5] vs. 0.9 ml/cm H_2O [IQR, -1.9 – 5.3]; $P=0.48$). The results for the $\text{PaO}_2/\text{FiO}_2$ ratio-matched group were consistent with those for the entire group ([Table 4](#)).

Traditional PaO_2 responders, defined by Gattinoni et al. [19] as patients showing an increase in $\text{PaO}_2/\text{FiO}_2$ ratio ≥ 20 mm Hg from baseline to prone positioning, accounted for 69.4% of patients with diffuse ARDS and 65.2% with focal ARDS. According to the novel definition of prone responders using the per-

Table 4. Primary and secondary outcomes

Variable	Entire group			PaO ₂ /FiO ₂ -matched		
	Diffuse ARDS (n=72)	Focal ARDS (n=23)	P-value	Diffuse ARDS (n=46)	Focal ARDS (n=23)	P-value
Primary outcome						
PaO ₂ /FiO ₂ change from baseline to 8–12 hours after prone positioning (mm Hg)	42.8 (11.6–83.2)	55.8 (11.1–109.2)	0.705	48.2 (14.0–108.9)	55.8 (11.1–109.2)	0.904
PaO ₂ /FiO ₂ change from baseline to 4–12 hours after resuming the supine position (mm Hg)	41.4 (3.8–88.3)	43.3 (24.0–98.0)	0.419	54.1 (9.4–88.5)	43.3 (24.0–98.0)	0.800
Secondary outcome						
Change of C _{dyn} from baseline to 8–12 hours after prone positioning (ml/cm H ₂ O)	-1.0 (-4.0 to 1.5)	-1.1 (-4.6 to 4.6)	0.364	-1.2 (-4.0 to 1.7)	-1.1 (-4.6 to 4.6)	0.379
Change of C _{dyn} from baseline to 4–12 hours after resuming the supine position(ml/cm H ₂ O)	0.9 (-1.9 to 5.3)	2.3 (-0.9 to 4.5)	0.475	0.9 (-2.1 to 3.7)	2.3 (-0.9 to 4.5)	0.423
Mortality at 28 days	37 (51.4)	10 (43.5)	0.674	23 (50.0)	10 (43.5)	0.798
ICU mortality	35 (48.6)	9 (39.1)	0.580	20 (43.5)	9 (39.1)	0.931

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

ARDS: acute respiratory distress syndrome; PaO₂: partial pressure of oxygen; FiO₂: fraction of inspired oxygen; C_{dyn}: dynamic lung compliance; ICU: intensive care unit.

centage change in PaO₂/FiO₂ ratio from baseline to 8–12 hours after prone positioning, the proportion of prone responders was 41.7% for diffuse ARDS and 60.9% for focal ARDS (P=0.17) [7]. After PS matching, traditional prone responders represented 67.4% of patients with diffuse ARDS and 65.2% of those with focal ARDS (P>0.99). Prone responders based on the percentage change in PaO₂/FiO₂ ratio accounted for 56.5% of those with diffuse ARDS and 60.9% of those with focal ARDS (P=0.93). The change in PaO₂/FiO₂ ratio and dynamic compliance from baseline to the first prone session were not different between the two groups, although the results were limited to the responders (Supplementary Tables 3 and 4). Further details on study outcomes separately analyzing survivors and non-survivors within 28 days can be found in the supplementary information (Supplementary Tables 5–8).

Outcomes of Patients and Predictors of Mortality

Mortality was not significantly different between the two groups; 10 patients (43.5%) in the focal ARDS group and 37 patients (51.4%) in the diffuse ARDS group died within 28 days; 9 patients (39.1%) in the focal ARDS group and 35 patients (48.6%) in the diffuse ARDS group died in the ICU (Table 4). The main cause of 28-day mortality was ARDS (78.7%), while the others had causes such as septic shock, arrhythmia, and hypovolemic shock. ICU mortality was significantly higher in the excluded patients than in the included patients (61.3% vs. 46.3%, P=0.03); however, there was no difference in mortality at 28 days (58.0% vs. 49.5%, P=0.24). In the multivariable Cox regression analysis, baseline serum lactate level (HR, 1.25; 95%

CI, 1.05–1.48 per 1-mmol/L increase) and change in PaO₂/FiO₂ ratio within 8–12 hours after the prone positioning session (HR, 0.99; 95% CI, 0.98–1.00 per 1 mm Hg increase) were significantly associated with 28-day mortality (Table 5).

DISCUSSION

This study investigated changes in oxygenation and compliance after the first prone positioning session in patients with ARDS according to lung morphology. Overall, the improvement in oxygenation after prone positioning was greater in the focal ARDS group than in the diffuse ARDS group; however, there was no statistically significant difference between the two groups. Moreover, there were no differences in changes in respiratory system compliance after prone positioning between the two groups. Among the PaO₂/FiO₂-matched cohort, there were no significant differences in change in PaO₂/FiO₂ ratio and compliance of the respiratory system after prone positioning between the two groups.

The definition of ARDS has been controversial since it was first described in 1967 [1], but currently it follows the Berlin definition [12]. In the Berlin criteria, ARDS is defined by bilateral opacities on chest radiograph, although a CT scan can also visualize the disease. The interobserver reliability of the Berlin definition is moderate, driven primarily by variability in imaging interpretation [20]. In an international cohort study of patients with ARDS [21], the diagnosis of ARDS was missed in two-thirds of the patients, leading to failure of appropriate strategies to reduce mortality. Another prospective study at-

Table 5. Univariable and multivariable Cox regression analysis of the prognostic factors associated with 28-day mortality

Factor	Univariable analysis		Multivariable analysis	
	HR (95% CI)	P-value	HR (95% CI)	P-value
Focal ARDS	0.83 (0.41–1.67)	0.603		
Age (yr)	1.00 (0.98–1.02)	0.650		
Male	0.80 (0.44–1.46)	0.467		
Body mass index (kg/m ²)	1.02 (0.94–1.10)	0.684		
SOFA score	1.09 (0.99–1.21)	0.072	1.12 (0.97–1.29)	0.122
APACHE II score	1.04 (0.99–1.08)	0.124		
SAPS II	1.01 (0.99–1.03)	0.190		
Charlson comorbidity index	1.03 (0.94–1.13)	0.508		
Solid tumor	1.32 (0.75–2.35)	0.339		
Hematologic malignancy	1.49 (0.77–2.87)	0.236		
Lactate (mmol/L)	1.26 (1.06–1.49)	0.007	1.25 (1.05–1.48)	0.013
Baseline PaO ₂ /FiO ₂ ratio (mm Hg)	1.00 (0.99–1.01)	0.960		
Baseline driving pressure (mm Hg)	1.01 (0.95–1.08)	0.732		
Interval of intubation and prone positioning (day)	1.02 (0.96–1.09)	0.525	1.12 (0.99–1.27)	0.072
Change of PaO ₂ /FiO ₂ ratio from the baseline to 8–12 hours after prone positioning session (mm Hg)	0.99 (0.98–1.00)	<0.001	0.99 (0.98–1.00)	0.002
Change of Cdyn from the baseline to 8–12 hours after prone positioning session (ml/cm H ₂ O)	0.98 (0.95–1.02)	0.329		
Neuromuscular blocker	1.21 (0.38–3.92)	0.741		
Glucocorticoids	1.64 (0.51–5.30)	0.405		
Vasopressors	1.02 (0.53–1.96)	0.957		
Inhaled nitric oxide	1.24 (0.63–2.43)	0.538		
Renal replacement therapy	1.54 (0.72–3.29)	0.268		
Cross over to extracorporeal membrane oxygenation	0.37 (0.05–2.69)	0.326		

HR: hazard ratio; CI: confidence intervals; ARDS: acute respiratory distress syndrome; SOFA: Sequential Organ Failure Assessment; APACHE: Acute Physiology and Chronic Health Evaluation; SAPS: Simplified Acute Physiology Score; PaO₂: partial pressure of oxygen; FiO₂: fraction of inspired oxygen; Cdyn: dynamic lung compliance.

tempted to show the superiority of personalized treatment according to lung morphology but was unsuccessful due to misclassification of images [11]. To overcome this imaging limitation, chest CT can be used to increase the accuracy of diagnosis [22,23]. A retrospective observational study showed that CT scans led to changes in management in 26.5% of patients with ARDS [24]. To obtain a more accurate evaluation of ARDS patterns, we used chest CT scans to classify the subphenotypes in the current study.

In the LIVE study, personalized management of mechanical ventilation was applied, considering the advantages of being prone to focal ARDS and high PEEP and recruitment maneuvers to diffuse ARDS; however, the results did not show a significant decrease in mortality [11]. Here, we did not directly compare the tailored strategy according to lung morphology due to the retrospective nature of the study. However, we applied prone positioning to patients with ARDS, focusing on lung protective ventilation, and there were no differences in

gas exchange and lung compliance between the two groups. Moreover, in multivariable analysis, the interval between intubation and prone positioning tends to be associated with 28-day mortality (HR, 1.12; 95% CI, 0.99–1.27). Therefore, prone positioning can be considered as soon as indicated, regardless of ARDS lung morphology.

In a previous study, Gattinoni et al. [19] defined “prone responders” as patients with ARDS in whom the PaO₂/FiO₂ ratio increased to ≥20 mm Hg. The proportion of prone responders was 72.1% in the Gattinoni group and 68.4% in the present study. Our institution proposed classification of prone responders as those with an increase in the percentage change in PaO₂/FiO₂ ratio of 53.5% from baseline to 8–12 hours after prone positioning (time P8-12) [7]. Based on this new criterion, the proportion of prone responders was 41.7% in the diffuse group and 60.9% in the focal group, a non-significant difference. However, there is some possibility of underpowered results of primary and secondary outcomes. Although

not statistically significant, the proportion of prone responders according to this novel definition was relatively higher and the 28-day and ICU mortality rates were relatively lower in the focal ARDS group than in the diffuse ARDS group. Moreover, among the traditional prone responders, the percentage change in $\text{PaO}_2/\text{FiO}_2$ ratio from baseline to P8-12 was significantly higher in the focal ARDS group than in the diffuse ARDS group (median, 90.4% [IQR, 69.5–161.1] vs. 75.1% [IQR, 37.0–106.9]; $P=0.04$) (Supplementary Table 3). Additionally, focal ARDS is considered to have a low baseline $\text{PaO}_2/\text{FiO}_2$ ratio due to its large shunt fraction. Regional shunt fraction tends to be higher in a dependent area [25]. In the present study, $\text{PaO}_2/\text{FiO}_2$ ratio before prone positioning was significantly lower in focal ARDS patients, who can have a substantial true shunt in the dorsal lung [26]. As prone positioning improves the shunt fraction [25], the greater improvement of oxygenation in the focal ARDS group might be due to the higher shunt fraction. Nevertheless, in patients with moderate-to-severe ARDS, the proportion of prone responders was greater than 50% with both definitions. Therefore, if a prone position is indicated, it should be actively implemented regardless of lung morphology.

Known prognostic factors of ARDS are physiological and laboratory variables such as age, SOFA score, oxygenation index, and driving pressure [27–29]. Additionally, circulating plasma markers of inflammation such as IL-6, IL-8, sTNFR1, and PAI-1 are used to classify subphenotypes of ARDS, and the higher are concentrations of biological variables, the higher is the mortality rate [9]. Previous studies have also revealed the correlation between ground-glass opacity (GGO) extent and inflammatory cytokine concentrations [30], as well as the correlation between GGO extent and mortality [31]. Although it is difficult to compare in our study because we did not measure prognostic biomarkers, it is possible that mortality tends to be higher in the diffuse ARDS group due to the extent of pulmonary opacity. Recently, in our institution, we revealed that improvement in oxygenation after prone positioning can be a predictor of survival [7], and a recent Italian study in coronavirus disease 2019 (COVID-19) ARDS showed that the sustained improvement of $\text{PaO}_2/\text{FiO}_2$ ratio after the first prone positioning was related to shorter duration of mechanical ventilation and less ICU mortality [32]. Furthermore, we evaluated the relationships between patient variables and survival using multivariable Cox regression analysis. It was shown that the extent of improvement in $\text{PaO}_2/\text{FiO}_2$ ratio after initial prone positioning could be a predictor of mortality (HR, 0.99; 95% CI,

0.98–1.00; $P=0.002$), and it was confirmed that improvement of oxygenation after prone positioning was related to mortality.

This study had several limitations. First, this was a retrospective study conducted in a single center. Additionally, more than half of patients were diagnosed with malignancy. Therefore, our results are not necessarily generalizable to other hospital settings. Second, the number of patients with ARDS who underwent chest CT was relatively small. Patients who did not undergo CT scan had a higher ICU mortality rate than those who did undergo CT scan. Additionally, those who did not undergo CT scan might have had unstable vital signs and worse status, hindering the CT procedure. Furthermore, there is possibility of underpowered results of the $\text{PaO}_2/\text{FiO}_2$ -matched cohort analysis due to limited sample size. Therefore, the generalizability of the results is limited. Third, chest CT cannot usually be performed at the time of ARDS diagnosis due to the risk of in-hospital transfer and radiation. However, in this study, the median duration from CT scan to prone positioning was 2 days, which might be appropriate to differentiate between focal and diffuse ARDS. Future studies might use other modalities, such as lung ultrasound, dynamic lung tomography, and electrical impedance tomography, to identify subphenotypes of ARDS at the bedside to overcome these risks. Fourth, most patients were under pressure-controlled ventilation. As peak inspiratory pressure was assumed to be equal to plateau pressure in pressure-controlled ventilation mode, driving pressure might have been overestimated. Additionally, esophageal pressure could not be measured in the present study, and change in transpulmonary pressure could not be confirmed. Fifth, regardless of consolidation or GGO, the distribution of lung aeration loss was simply divided into focal and diffuse in this study. Moreover, radiological findings could not be quantified. In future studies, it is necessary to quantify the type and distribution of aeration loss.

In conclusion, the improvement in oxygenation after prone positioning did not differ according to lung morphology in patients with moderate-to-severe ARDS. The findings of our study suggest that prone positioning can be initiated as soon as indicated, regardless of ARDS lung morphology.

CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.

ORCID

Na Young Kim	https://orcid.org/0000-0002-4123-8619
Si Mong Yoon	https://orcid.org/0000-0001-8011-7986
Jimyung Park	https://orcid.org/0000-0003-2655-5517
Jinwoo Lee	https://orcid.org/0000-0003-0958-106X
Sang-Min Lee	https://orcid.org/0000-0002-1388-9318
Hong Yeul Lee	https://orcid.org/0000-0002-3638-8890

AUTHOR CONTRIBUTIONS

Conceptualization: NYK, HYL. Data Curation: NYK, SMY, JP, HYL. Formal analysis: NYK, HYL. Methodology: NYK, HYL. Drafting the manuscript: NYK. Supervision: HYL. Writing–original draft: all authors. Writing–review & editing: all authors.

SUPPLEMENTARY MATERIALS

Supplementary materials can be found via <https://doi.org/10.4266/acc.2022.00367>.

REFERENCES

- Ashbaugh DG, Bigelow DB, Petty TL, Levine BE. Acute respiratory distress in adults. *Lancet* 1967;2:319-23.
- Acute Respiratory Distress Syndrome Network; Brower RG, Matthay MA, Morris A, Schoenfeld D, Thompson BT, et al. Ventilation with lower tidal volumes as compared with traditional tidal volumes for acute lung injury and the acute respiratory distress syndrome. *N Engl J Med* 2000;342:1301-8.
- Hager DN, Krishnan JA, Hayden DL, Brower RG; ARDS Clinical Trials Network. Tidal volume reduction in patients with acute lung injury when plateau pressures are not high. *Am J Respir Crit Care Med* 2005;172:1241-5.
- Papazian L, Forel JM, Gacouin A, Penot-Ragon C, Perrin G, Loundou A, et al. Neuromuscular blockers in early acute respiratory distress syndrome. *N Engl J Med* 2010;363:1107-16.
- Briel M, Meade M, Mercat A, Brower RG, Talmor D, Walter SD, et al. Higher vs lower positive end-expiratory pressure in patients with acute lung injury and acute respiratory distress syndrome: systematic review and meta-analysis. *JAMA* 2010;303:865-73.
- Guérin C, Reignier J, Richard JC, Beuret P, Gacouin A, Boulain T, et al. Prone positioning in severe acute respiratory distress syndrome. *N Engl J Med* 2013;368:2159-68.
- Lee HY, Cho J, Kwak N, Choi SM, Lee J, Park YS, et al. Improved oxygenation after prone positioning may be a predictor of survival in patients with acute respiratory distress syndrome. *Crit Care Med* 2020;48:1729-36.
- Scholten EL, Beitler JR, Prisk GK, Malhotra A. Treatment of ARDS with prone positioning. *Chest* 2017;151:215-24.
- Reddy K, Sinha P, O’Kane CM, Gordon AC, Calfee CS, McAuley DF. Subphenotypes in critical care: translation into clinical practice. *Lancet Respir Med* 2020;8:631-43.
- Calfee CS, Delucchi K, Parsons PE, Thompson BT, Ware LB, Matthay MA, et al. Subphenotypes in acute respiratory distress syndrome: latent class analysis of data from two randomised controlled trials. *Lancet Respir Med* 2014;2:611-20.
- Constantin JM, Jabaudon M, Lefrant JY, Jaber S, Quenot JP, Langeron O, et al. Personalised mechanical ventilation tailored to lung morphology versus low positive end-expiratory pressure for patients with acute respiratory distress syndrome in France (the LIVE study): a multicentre, single-blind, randomised controlled trial. *Lancet Respir Med* 2019;7:870-80.
- ARDS Definition Task Force, Ranieri VM, Rubenfeld GD, Thompson BT, Ferguson ND, Caldwell E, et al. Acute respiratory distress syndrome: the Berlin Definition. *JAMA* 2012;307:2526-33.
- Hu SL, He HL, Pan C, Liu AR, Liu SQ, Liu L, et al. The effect of prone positioning on mortality in patients with acute respiratory distress syndrome: a meta-analysis of randomized controlled trials. *Crit Care* 2014;18:R109.
- Coppola S, Pozzi T, Gurgitano M, Liguori A, Duka E, Bichi F, et al. Radiological pattern in ARDS patients: partitioned respiratory mechanics, gas exchange and lung recruitability. *Ann Intensive Care* 2021;11:78.
- Costamagna A, Pivetta E, Goffi A, Steinberg I, Arina P, Mazzeo AT, et al. Clinical performance of lung ultrasound in predicting ARDS morphology. *Ann Intensive Care* 2021;11:51.
- Hao D, Low S, Di Fenza R, Shenoy ES, Ananian L, Prout LA, et al. Prone positioning of intubated patients with an elevated body-mass index. *N Engl J Med* 2022;386:e34.
- Schmidt M, Pham T, Arcadipane A, Agerstrand C, Ohshimo S, Pellegrino V, et al. Mechanical ventilation management during extracorporeal membrane oxygenation for acute respiratory distress syndrome. an international multicenter prospective cohort. *Am J Respir Crit Care Med* 2019;200:1002-12.
- Peduzzi P, Concato J, Kemper E, Holford TR, Feinstein AR. A simulation study of the number of events per variable in logistic regression analysis. *J Clin Epidemiol* 1996;49:1373-9.
- Gattinoni L, Vagginelli F, Carlesso E, Taccone P, Conte V, Chiumello D, et al. Decrease in PaCO₂ with prone position is

- predictive of improved outcome in acute respiratory distress syndrome. *Crit Care Med* 2003;31:2727-33.
20. Sjoding MW, Hofer TP, Co I, Courey A, Cooke CR, Iwashyna TJ. Interobserver reliability of the Berlin ARDS definition and strategies to improve the reliability of ARDS diagnosis. *Chest* 2018;153:361-7.
 21. Bellani G, Laffey JG, Pham T, Fan E, Brochard L, Esteban A, et al. Epidemiology, patterns of care, and mortality for patients with acute respiratory distress syndrome in intensive care units in 50 countries. *JAMA* 2016;315:788-800.
 22. Rouby JJ, Puybasset L, Nieszkowska A, Lu Q. Acute respiratory distress syndrome: lessons from computed tomography of the whole lung. *Crit Care Med* 2003;31(4 Suppl):S285-95.
 23. Chiumello D, Langer T, Vecchi V, Luoni S, Colombo A, Brioni M, et al. Low-dose chest computed tomography for quantitative and visual anatomical analysis in patients with acute respiratory distress syndrome. *Intensive Care Med* 2014;40:691-9.
 24. Simon M, Braune S, Laqmani A, Metschke M, Berliner C, Kalsow M, et al. Value of computed tomography of the chest in subjects with ARDS: a retrospective observational study. *Respir Care* 2016;61:316-23.
 25. Richter T, Bellani G, Scott Harris R, Vidal Melo MF, Winkler T, Venegas JG, et al. Effect of prone position on regional shunt, aeration, and perfusion in experimental acute lung injury. *Am J Respir Crit Care Med* 2005;172:480-7.
 26. Petersson J, Glenny RW. Gas exchange and ventilation-perfusion relationships in the lung. *Eur Respir J* 2014;44:1023-41.
 27. Amato MB, Meade MO, Slutsky AS, Brochard L, Costa EL, Schoenfeld DA, et al. Driving pressure and survival in the acute respiratory distress syndrome. *N Engl J Med* 2015;372:747-55.
 28. Balzer F, Menk M, Ziegler J, Pille C, Wernecke KD, Spies C, et al. Predictors of survival in critically ill patients with acute respiratory distress syndrome (ARDS): an observational study. *BMC Anesthesiol* 2016;16:108.
 29. Song M, Liu Y, Lu Z, Luo H, Peng H, Chen P. Prognostic factors for ARDS: clinical, physiological and atypical immunodeficiency. *BMC Pulm Med* 2020;20:102.
 30. Wu Z, Liu X, Liu J, Zhu F, Liu Y, Liu Y, et al. Correlation between ground-glass opacity on pulmonary CT and the levels of inflammatory cytokines in patients with moderate-to-severe COVID-19 pneumonia. *Int J Med Sci* 2021;18:2394-400.
 31. Chen YY, Kuo JS, Ruan SY, Chien YC, Ku SC, Yu CJ, et al. Prognostic value of computed tomographic findings in acute respiratory distress syndrome and the response to prone positioning. *BMC Pulm Med* 2022;22:71.
 32. Scaramuzza G, Gamberini L, Tonetti T, Zani G, Ottaviani I, Mazzoli CA, et al. Sustained oxygenation improvement after first prone positioning is associated with liberation from mechanical ventilation and mortality in critically ill COVID-19 patients: a cohort study. *Ann Intensive Care* 2021;11:63.