

FAKULTNÍ NEMOCNICE BRNO
A LÉKAŘSKÁ FAKULTA
MASARYKOVY UNIVERZITY



**KLINIKA DĚTSKÉ
ANESTEZIOLOGIE
A RESUSCITACE**

Data, které zásadně ovlivnili praxi při nastavení parametrů UPV

Milan Kratochvíl

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**M U N I
M E D**

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**VENTILATION WITH LOWER TIDAL VOLUMES AS COMPARED WITH
TRADITIONAL TIDAL VOLUMES FOR ACUTE LUNG INJURY
AND THE ACUTE RESPIRATORY DISTRESS SYNDROME**

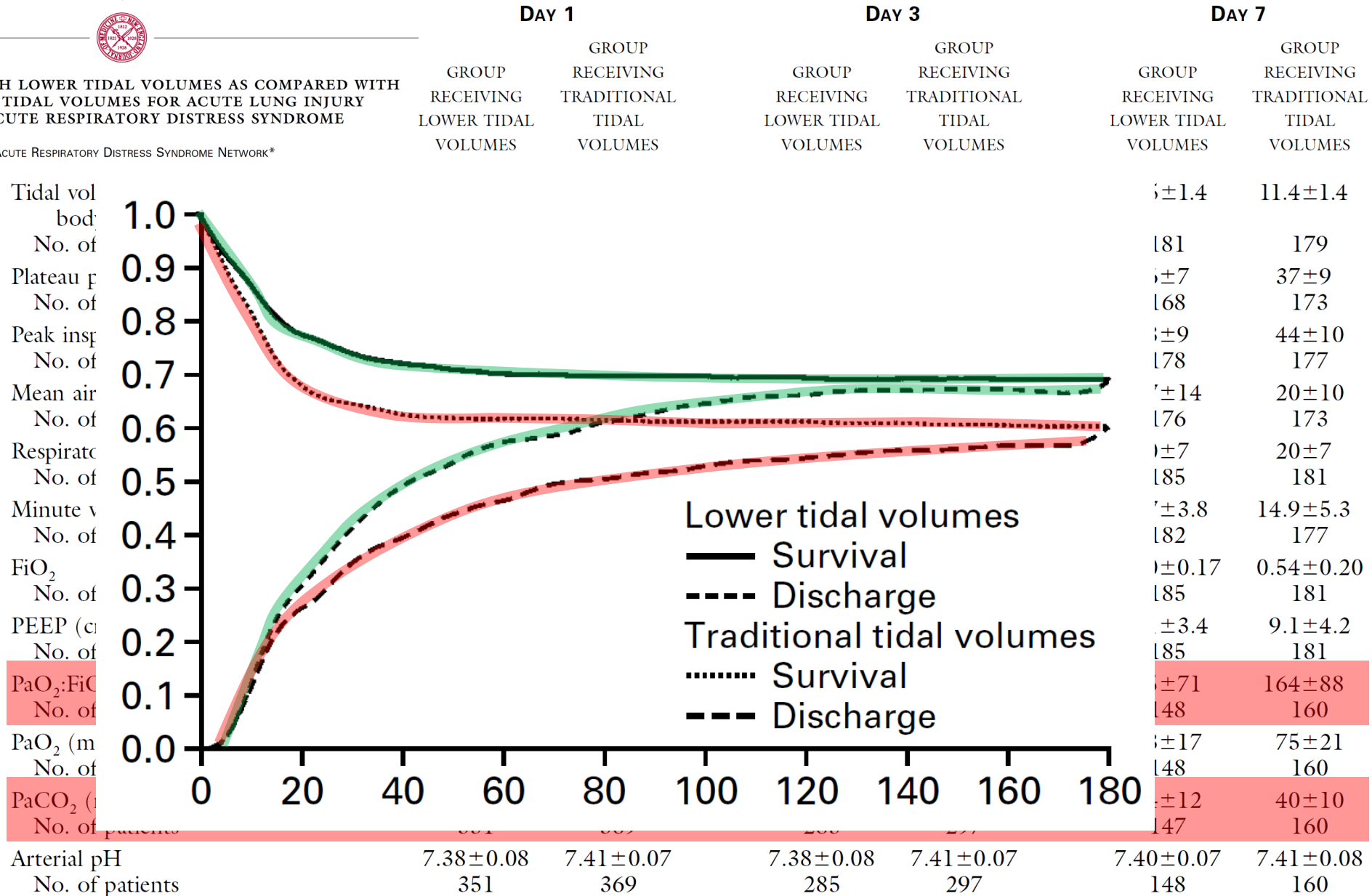
THE ACUTE RESPIRATORY DISTRESS SYNDROME NETWORK*



VENTILATION WITH LOWER TIDAL VOLUMES AS COMPARED WITH TRADITIONAL TIDAL VOLUMES FOR ACUTE LUNG INJURY AND THE ACUTE RESPIRATORY DISTRESS SYNDROME

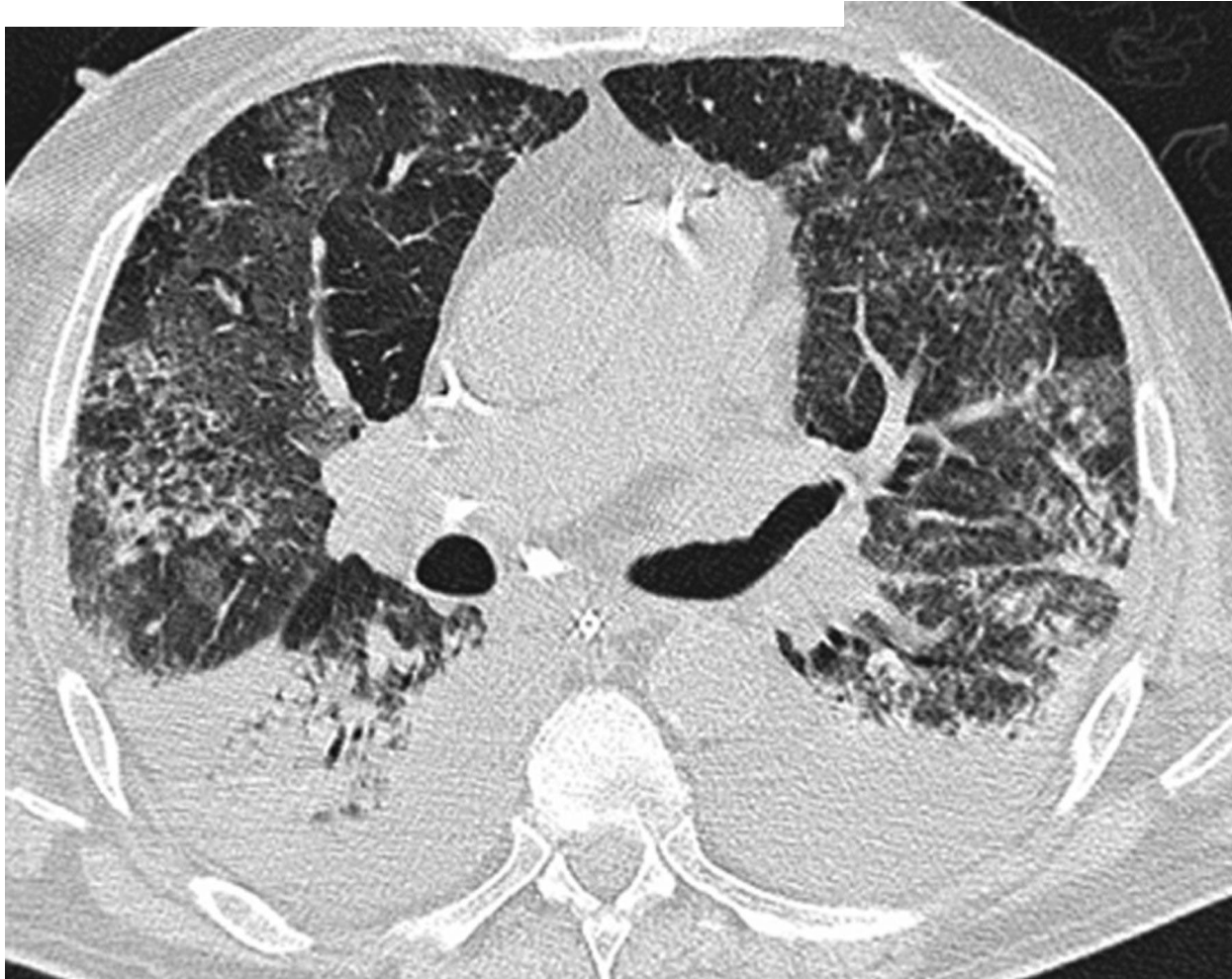
THE ACUTE RESPIRATORY DISTRESS SYNDROME NETWORK*

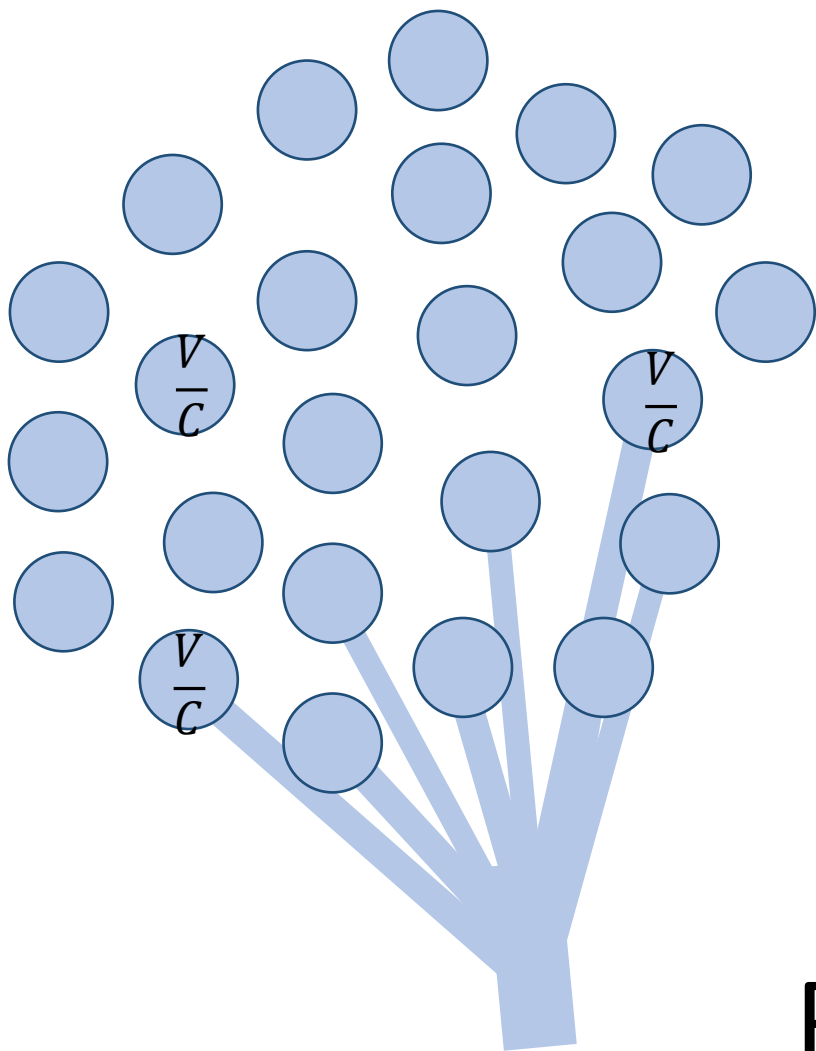
URING THE FIRST SEVEN DAYS OF TREATMENT IN PATIENTS WITH ACUTE LUNG INJURY AND THE ACUTE RESPIRATORY DISTRESS SYNDROME.*



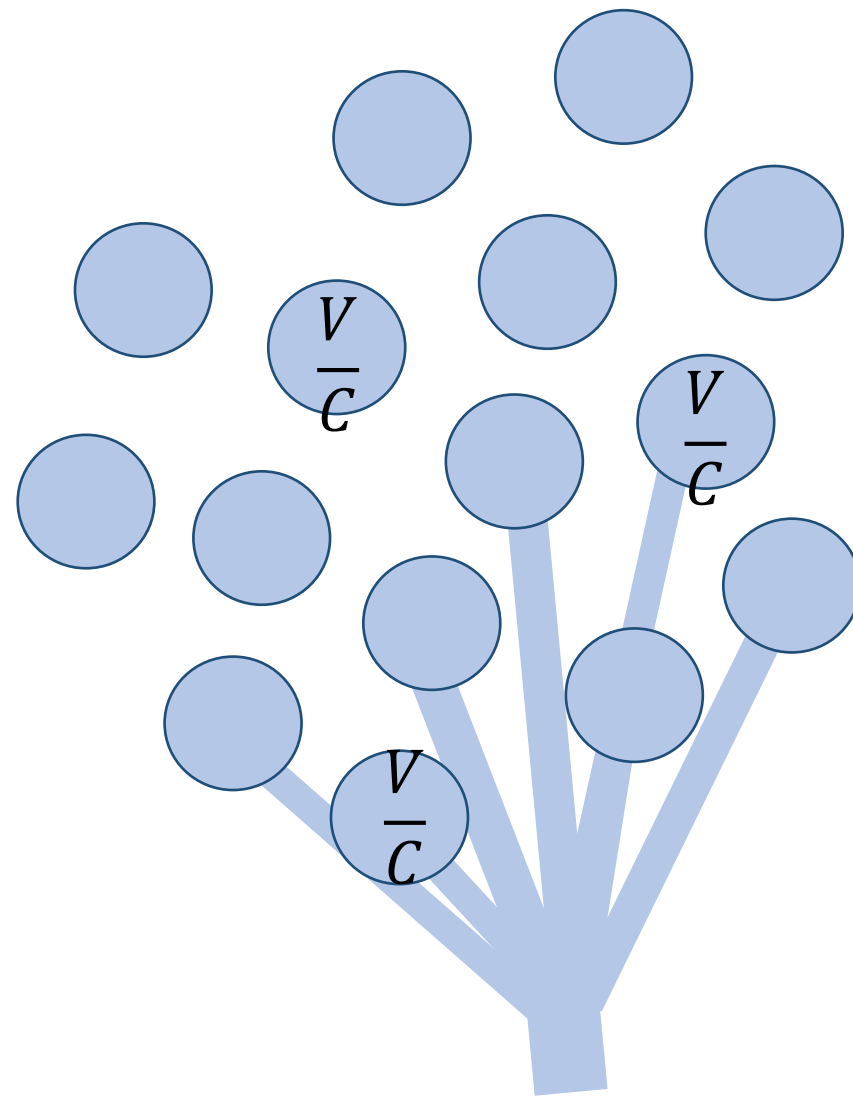
Luciano Gattinoni
Antonio Pesenti

The concept of “baby lung”





$$P = \frac{V}{C}$$



Luciano Gattinoni
Antonio Pesenti

The concept of “baby lung”

- \int JOURNAL OF APPLIED PHYSIOLOGY
Vol. 28, No. 5, May 1970. *Printed in U.S.A.*
- \int Stress distribution in lungs: a model of
- \int pulmonary elasticity

JERE MEAD, TAMOTSU TAKISHIMA, AND DAVID LEITH
Department of Physiology, Harvard University School of Public Health, Boston, Massachusetts 02115

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Ventilation Strategy Using Low Tidal Volumes, Recruitment Maneuvers, and High Positive End-Expiratory Pressure for Acute Lung Injury and Acute Respiratory Distress Syndrome

A Randomized Controlled Trial

Higher versus Lower Positive End-Expiratory Pressure in Patients with the Acute Respiratory Distress Syndrome

The National Heart, Lung, and Blood Institute

Maureen O. Meade, MD, MSc
Deborah J. Cook, MD, MSc
Gordon H. Guyatt, MD, MSc
Arthur S. Slutsky, MD
Yaseen M. Arabi, MD

Context Low-*tidal-volume* ventilation reduces mortality in critically ill patients with acute lung injury and acute respiratory distress syndrome. Instituting additional strategies to open collapsed lung tissue may further reduce mortality.

Objective To compare an established low-*tidal-volume* ventilation strategy with an experimental strategy based on the original "open-lung approach," combining low tidal volume, lung recruitment maneuvers, and high positive-end-expiratory pressure.

Positive End-Expiratory Pressure Setting in Adults With Acute Lung Injury and Acute Respiratory Distress Syndrome

A Randomized Controlled Trial

Alain Mercat, MD

Context The need for lung protection is universally accepted, but the optimal positive end-expiratory pressure is controversial. The need for lung protection is universally accepted, but the optimal positive end-expiratory pressure is controversial. The need for lung protection is universally accepted, but the optimal positive end-expiratory pressure is controversial.

Thomas E. Stewart, MD
for the Lung Open Ventilation Study Investigators

concealed allocation and blinded outcome assessment. The study was conducted in 30 intensive care units from March 2006 in 30 intensive care units.

Patients with acute lung injury and acute respiratory distress syndrome and fraction of inspired oxygen not exceeding 250.

Low tidal volumes of 6 mL/kg of predicted body weight, 5 cm H₂O, and conventional levels of positive end-expiratory pressure. The experimental strategy included target plateau pressures not exceeding 40 cm H₂O and positive end-expiratory pressures (n=475).

met criteria for acute respiratory distress syndrome. Mortality was similar in the 2 groups, and mean positive end-expiratory pressures were 14.6 (SD, 3.4) cm H₂O in the experimental group vs 9.8 (SD, 2.7) cm H₂O among controls during the first 72 hours (*P* < .001). All-cause hospital mortality rates were 36.4% and 40.4%, respectively (relative risk [RR], 0.90; 95% confidence interval [CI], 0.77-1.05; *P* = .19). Barotrauma rates were 11.2% and 9.1% (RR, 1.21; 95% CI, 0.83-1.75; *P* = .33). The experimental group had lower rates of refractory hypoxemia.

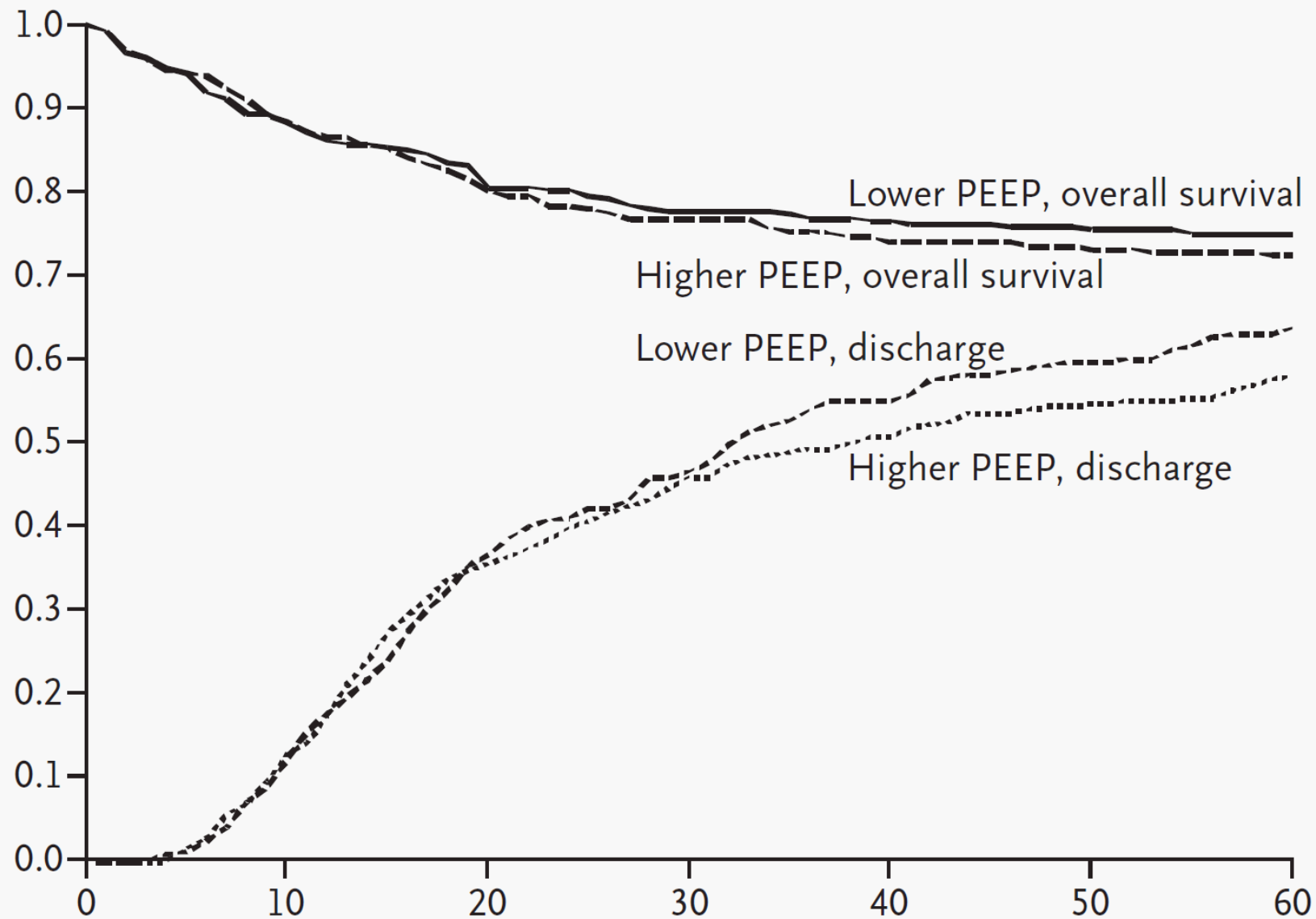
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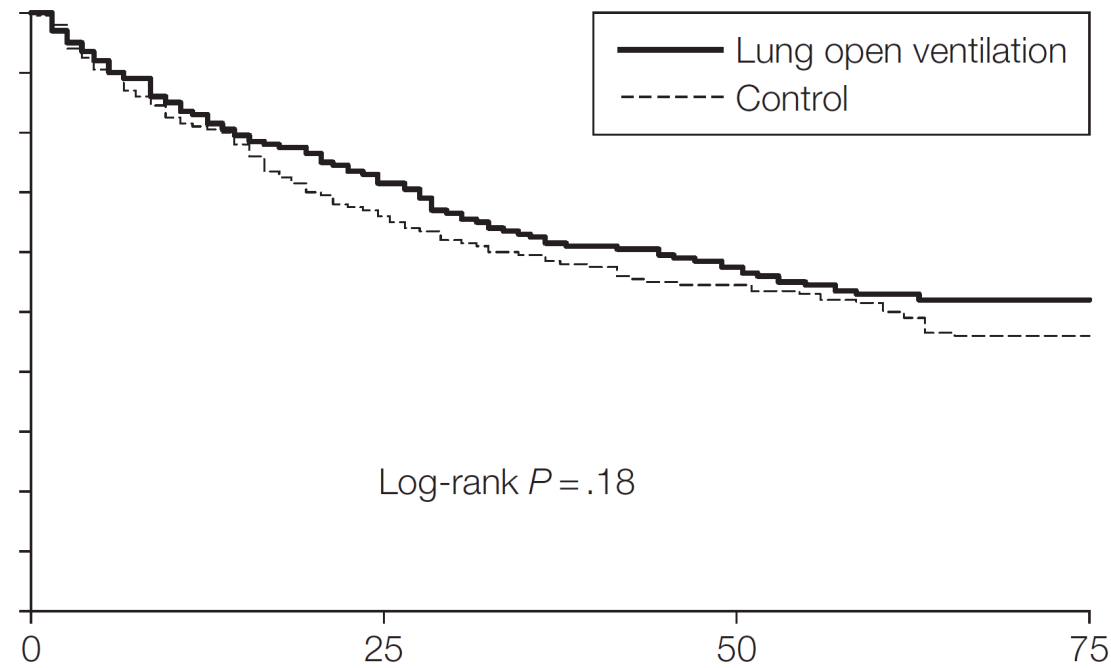
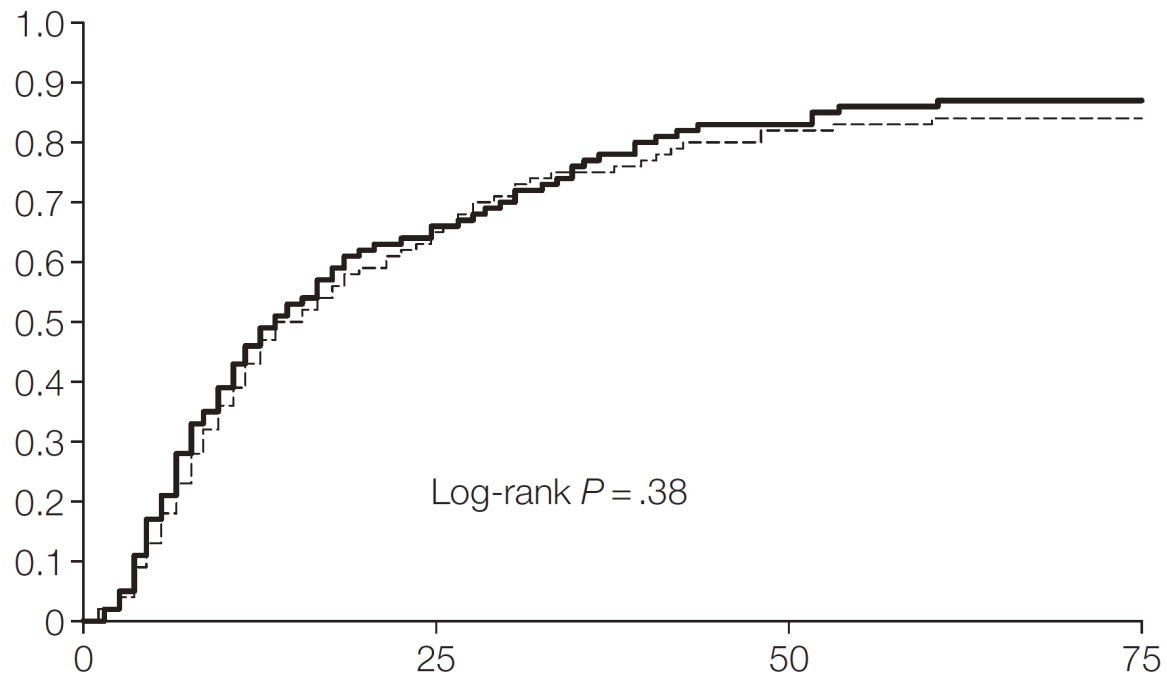
Higher versus Lower Positive End-E in Patients with the Acute Respirator

The National Heart, Lung, and Blood Institute ARDS



Ventilation Strategy Using Low Tidal Volumes, Recruitment Maneuvers, and High Positive End-Expiratory Pressure for Acute Lung Injury and Acute Respiratory Distress Syndrome

A Randomized Controlled Trial



Positive End-Expiratory Pressure Setting in Adults With Acute Lung Injury and Acute Respiratory Distress Syndrome

A Randomized Controlled Trial

Table 4. Main Outcome Variables

Outcome	Minimal Distension (n = 382)	Increased Recruitment (n = 385)	<i>P</i> Value
	No. (%)		
Death in the first 28 d ^a	119 (31.2)	107 (27.8)	.31
Death before hospital discharge	149 (39.0)	136 (35.4)	.30
Death in the first 60 d	151 (39.5)	138 (35.9)	.31
Pneumothorax between day 1 and day 28 ^b	22 (5.8)	26 (6.8)	.57
	Median (IQR)		
No. of days between day 1 and day 28			
Ventilator-free ^c	3 (0-17)	7 (0-19)	.04
Organ failure-free ^d	2 (0-16)	6 (0-18)	.04
Cardiovascular failure-free ^d	21 (4-26)	23 (10-26)	.09
Renal failure-free ^d	27.5 (8.0-28.0)	28.0 (11.0-28.0)	.23

Refining Ventilatory Treatment for Acute Lung Injury and Acute Respiratory Distress Syndrome

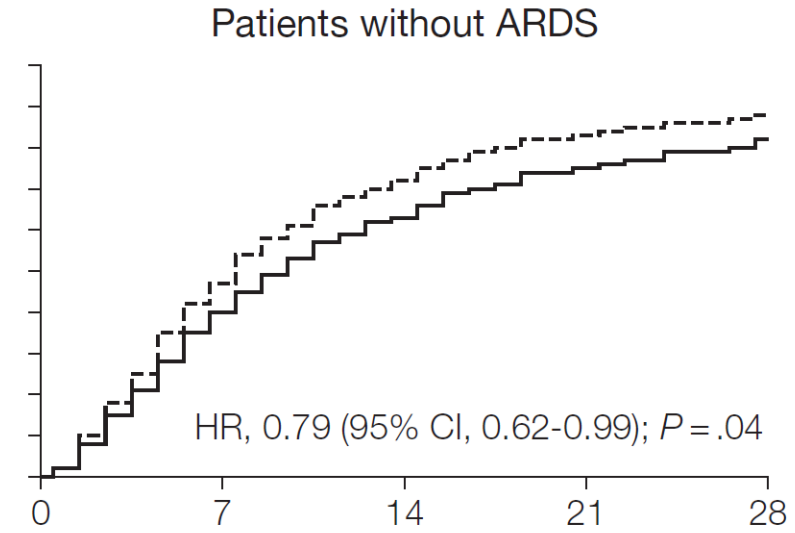
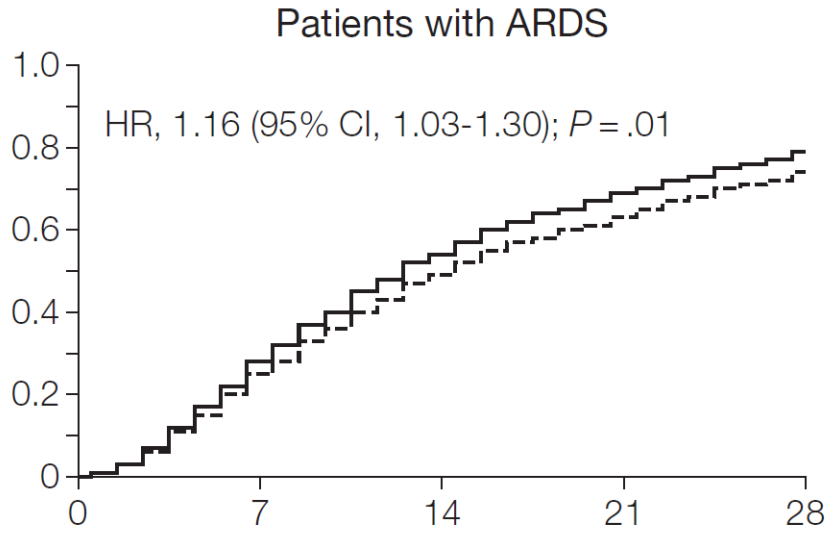
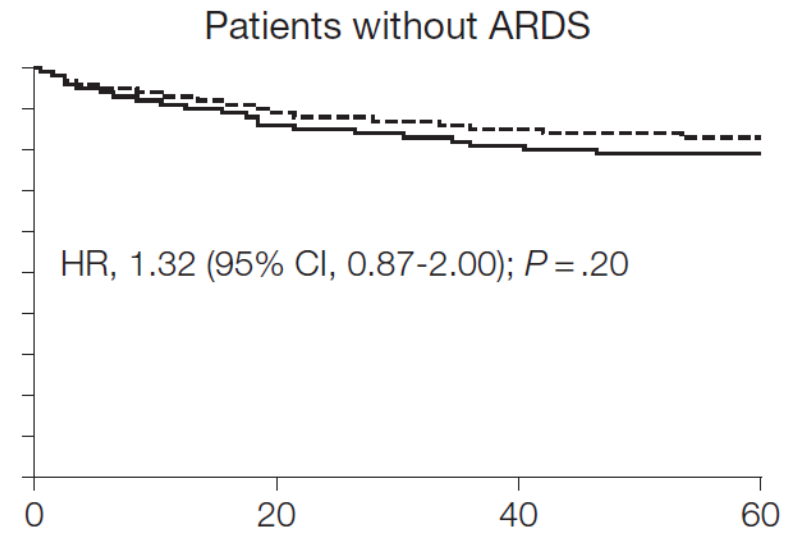
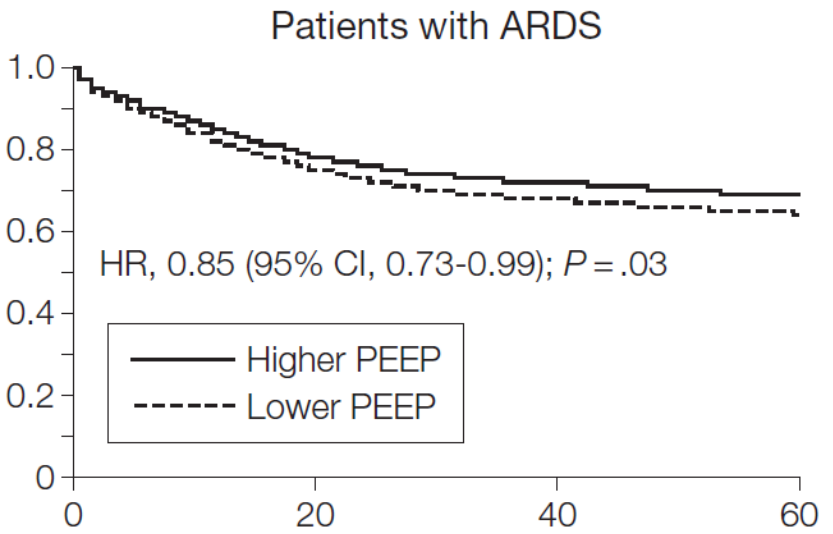
Luciano Gattinoni, MD, FRCP

Pietro Caironi, MD

at a FiO_2 of 0.80 or higher in the Express study.¹⁶ Taking the 2 studies together, the numbers of patients with severe hypoxemia requiring rescue therapy were 94 (10.9%) in the higher level of PEEP group and 184 (20.7%) in the lower level of PEEP group. Although mortality was similar (60.6% and 58.2% in the higher and lower PEEP groups, respectively), the difference in incidence suggests that **the rate of these pulmonary deaths was much lower in the group receiving a higher level of PEEP (6.6% vs 12.0%)**. One possible inference is that a higher level of PEEP may prevent a large number of pulmonary deaths, which may be the reason for the 3% to 4% mortality difference favoring the group receiving a higher level of PEEP observed both in the LOV study and in the Express study.

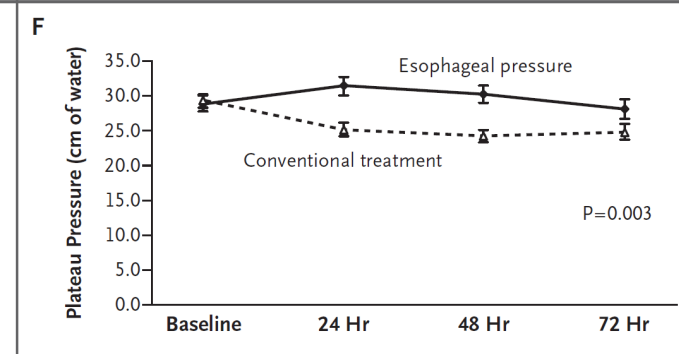
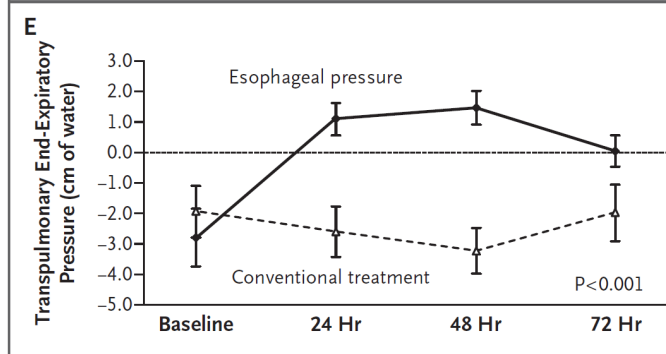
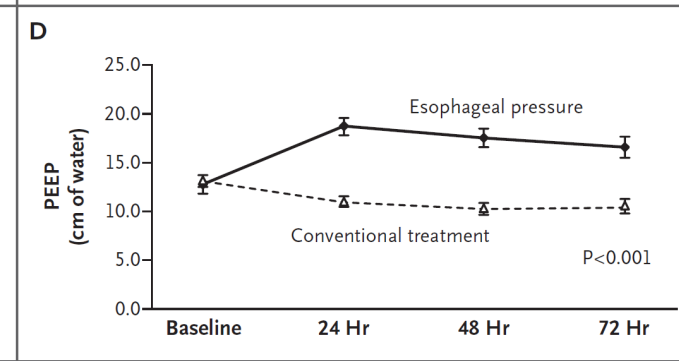
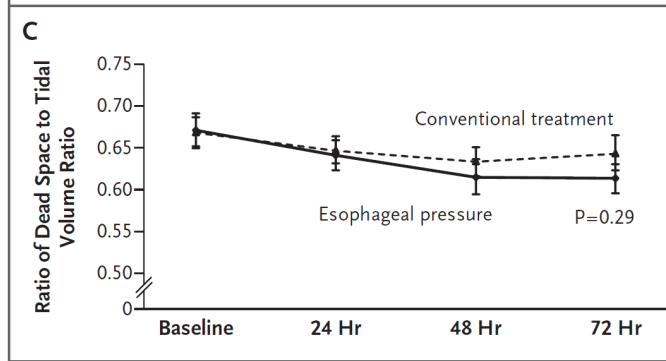
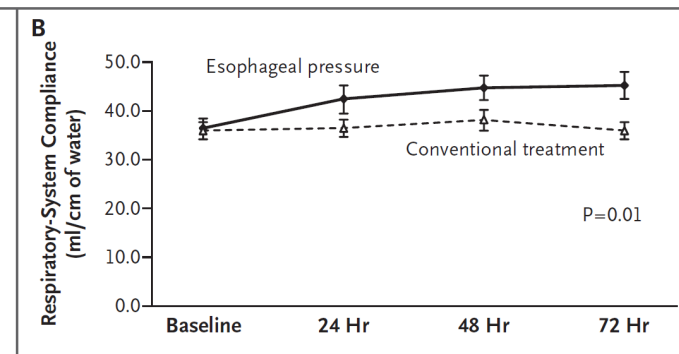
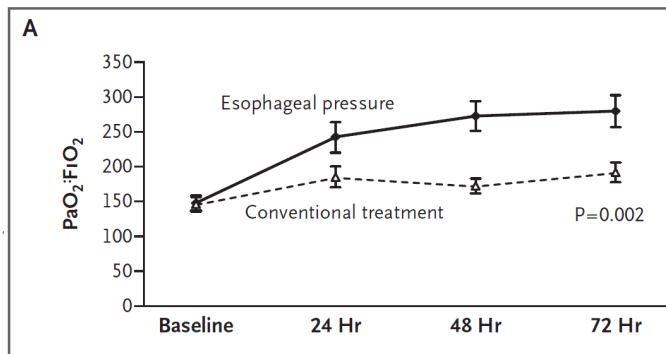
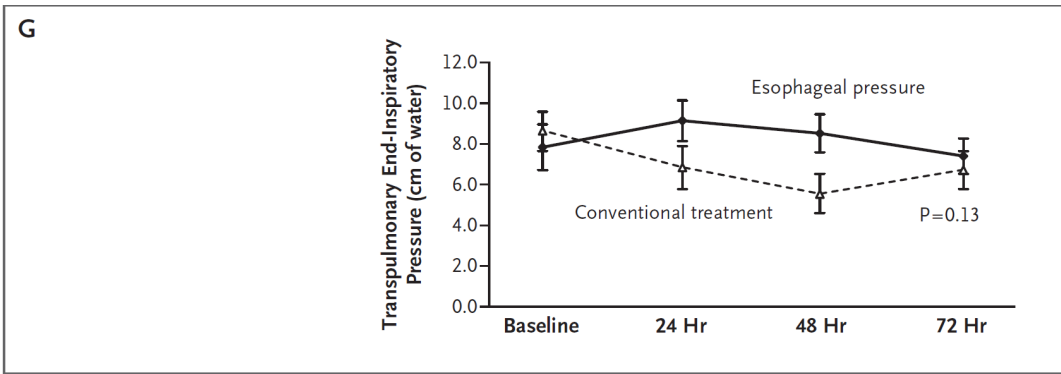
Higher vs Lower Positive End-Expiratory Pressure in Patients With Acute Lung Injury and F_{iO_2} System

In-hospital time to death



Mechanical Ventilation Guided by Esophageal Pressure in Acute Lung Injury

Daniel Talmor, M.D., M.P.H., Todd Sarge, M.D., Atul Malhotra, M.D., Carl R. O'Donnell, Sc.D., M.P.H., Ray Ritz, R.R.T., Alan Lisbon, M.D., Victor Novack, M.D., Ph.D., and Stephen H. Loring, M.D.



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Prone Positioning in Severe Acute Respiratory Distress
Syndrome

Claude Guérin, M.D., Ph.D., Jean Reignier, M.D., Ph.D., Jean-Christophe Richard, M.D., Ph.D., Pascal Beuret, M.D.,
Arnaud Gacouin, M.D., Thierry Boulain, M.D., Emmanuelle Mercier, M.D., Michel Badet, M.D.,
Alain Mercat, M.D., Ph.D., Olivier Baudin, M.D., Marc Clavel, M.D., Delphine Chatellier, M.D., Samir Jaber, M.D., Ph.D.,
Sylvène Rosselli, M.D., Jordi Mancebo, M.D., Ph.D., Michel Sirodot, M.D., Gilles Hilbert, M.D., Ph.D.,
Christian Bengler, M.D., Jack Richecoeur, M.D., Marc Gainnier, M.D., Ph.D., Frédérique Bayle, M.D.,
Gael Bourdin, M.D., Véronique Leray, M.D., Raphaele Girard, M.D., Loredana Baboi, Ph.D., and Louis Ayzac, M.D.,
for the PROSEVA Study Group*

Prone Positioning in Severe Acute Respiratory Distress Syndrome

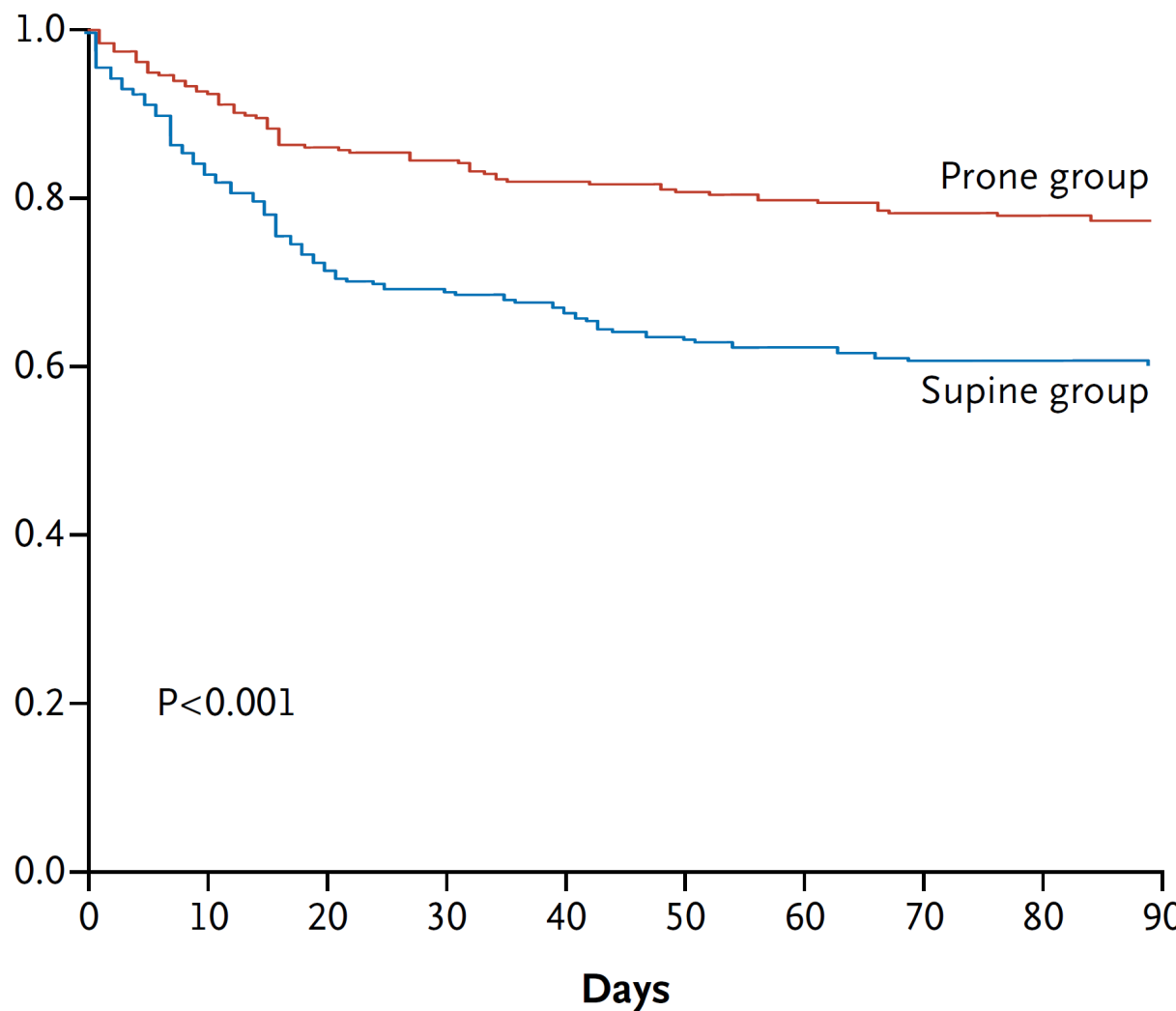


Table 3. Primary and Secondary Outcomes According to Study Group.*

Outcome	Supine Group (N=229)	Prone Group (N=237)	Hazard Ratio or Odds Ratio with the Prone Position (95% CI)	P Value
Mortality — no. (% [95% CI])				
At day 28				
Not adjusted	75 (32.8 [26.4–38.6])	38 (16.0 [11.3–20.7])	0.39 (0.25–0.63)	<0.001
Adjusted for SOFA score†			0.42 (0.26–0.66)	<0.001
At day 90				
Not adjusted	94 (41.0 [34.6–47.4])	56 (23.6 [18.2–29.0])	0.44 (0.29–0.67)	<0.001
Adjusted for SOFA score†			0.48 (0.32–0.72)	<0.001
Successful extubation at day 90 — no./total no. (% [95% CI])	145/223 (65.0 [58.7–71.3])	186/231 (80.5 [75.4–85.6])	0.45 (0.29–0.70)	<0.001
Time to successful extubation, assessed at day 90 — days				
Survivors	19±21	17±16		0.87
Nonsurvivors	16±11	18±14		
Length of ICU stay, assessed at day 90 — days				
Survivors	26±27	24±22		0.05
Nonsurvivors	18±15	21±20		
Ventilation-free days				
At day 28	10±10	14±9		<0.001
At day 90	43±38	57±34		<0.001
Pneumothorax — no. (% [95% CI])	13 (5.7 [3.9–7.5])	15 (6.3 [4.9–7.7])	0.89 (0.39–2.02)	0.85
Noninvasive ventilation — no./total no. (% [95% CI])				
At day 28	10/212 (4.7 [1.9–7.5])	4/228 (1.8 [0.1–3.5])	0.36 (0.07–3.50)	0.11
At day 90	3/206 (1.5 [0.2–3.2])	4/225 (1.8 [0.1–3.5])	1.22 (0.23–6.97)	1.00
Tracheotomy — no./total no. (% [95% CI])				
At day 28	12/229 (5.2 [2.3–8.1])	9/237 (3.8 [1.4–6.0])	0.71 (0.27–1.86)	0.37
At day 90	18/223 (8.1 [4.5–11.7])	15/235 (6.4 [3.3–9.5])	0.78 (0.36–1.67)	0.59

Závěr

- Nízké dechové objemy
- Optimálně titrovaný PEEP
- Prone positioning
- RM pouze jako rescue postup

Děkuji za pozornost