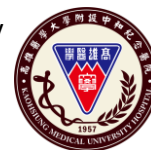




Electric impedance tomography guided-PEEP titration for ARDS

2025-11-8

高雄醫學大學附設中和紀念醫院
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History of ARDS

- Acute respiratory distress syndrome (ARDS) was first described by Ashbaugh in 1967 writing in The Lancet.
- His study was based on a case series of 12 patients treated in a civilian environment in the USA.

ARDS would later become closely associated with the Vietnam war, in which the American military deployed unprecedented medical resources to revive its wounded service men. Patient survival was further enhanced by the introduction of helicopter evacuation, which enabled casualties to reach an intensive care environment less than an hour after injury.

During this conflict, doctors from the American Army

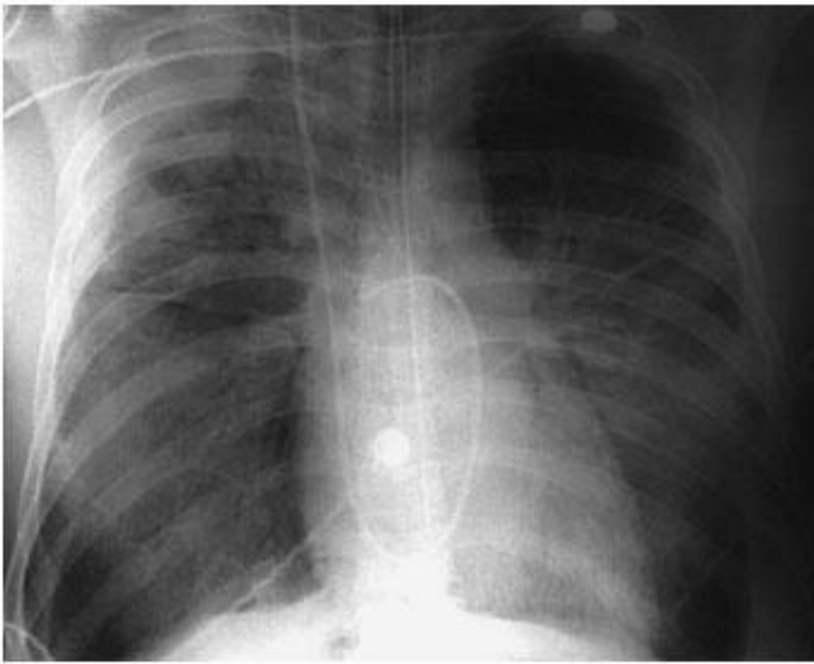
Medical Corps identified a subgroup of patients who appeared to respond to initial resuscitation, but later succumbed to respiratory distress. Previously fit young servicemen developed a persistent hypoxia that did not respond to an oxygen mask. More intriguing still, many of these patients actually lacked an obvious chest wound, with some suffering from burns, isolated head injury, or trauma to the limbs.

non-cardiogenic pulmonary oedema

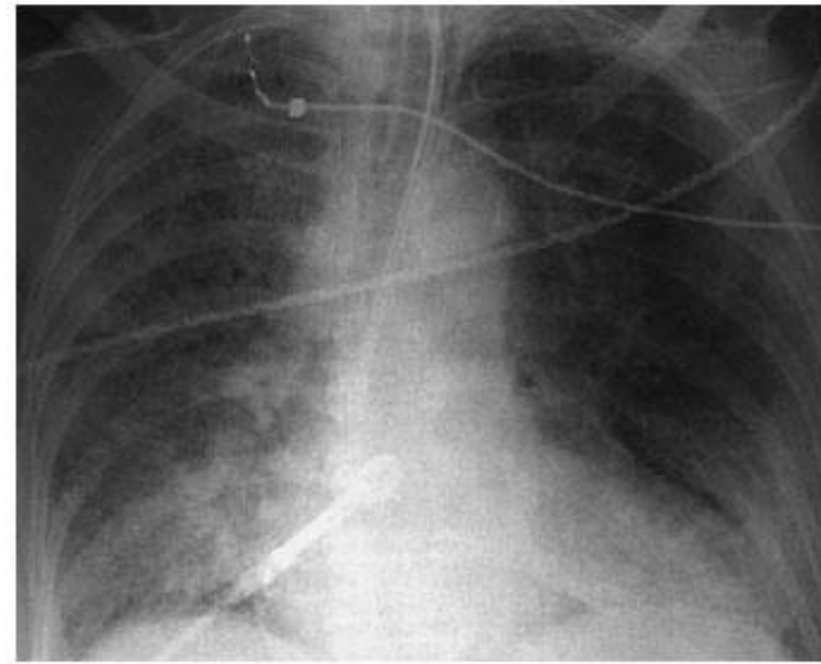
The

From 1 November 1955 to 30 April 1975

VIENTNAM WAR



A

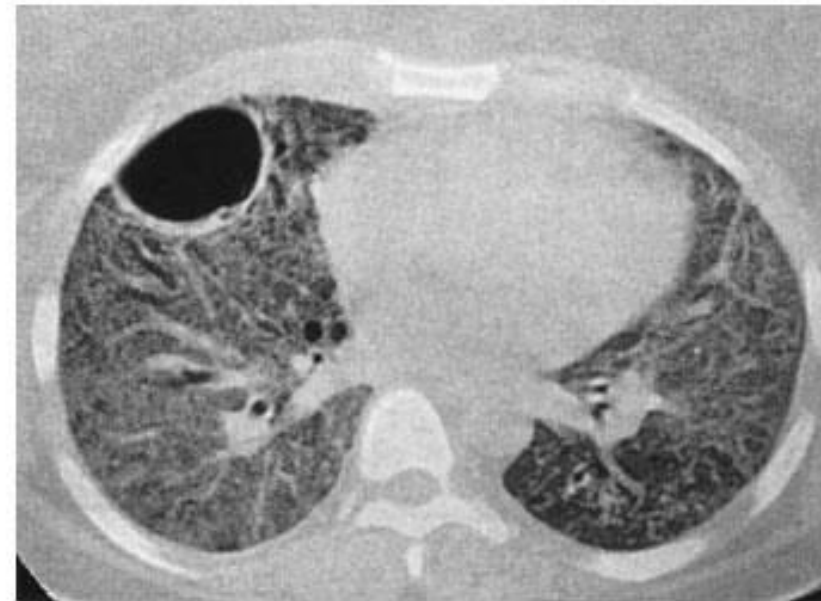


B

Acute, or Exudative, Phase (Panels A and C) Fibrosing-Alveolitis Phase (Panels B and D)

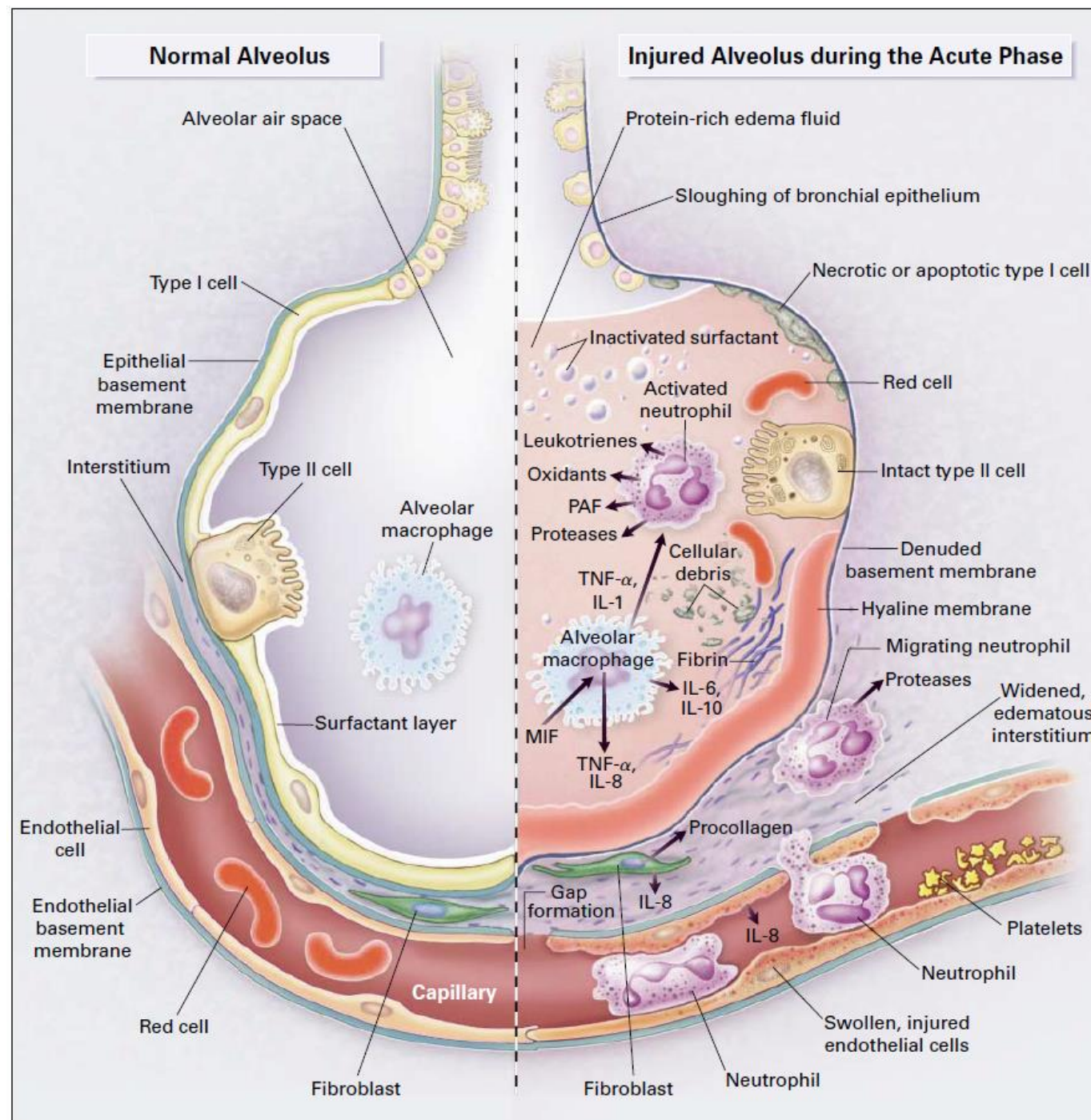


C

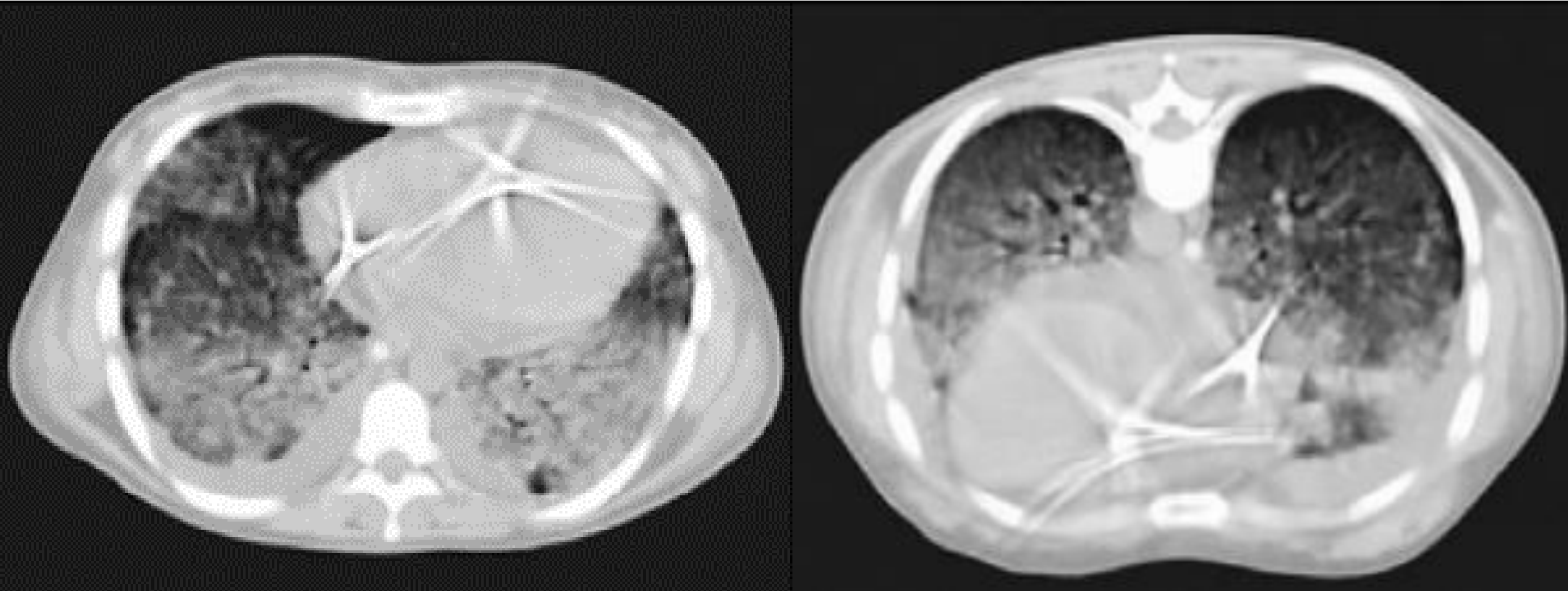


D

Ware LB, Matthay MA. N Engl J Med 2000;342:1334-1349.

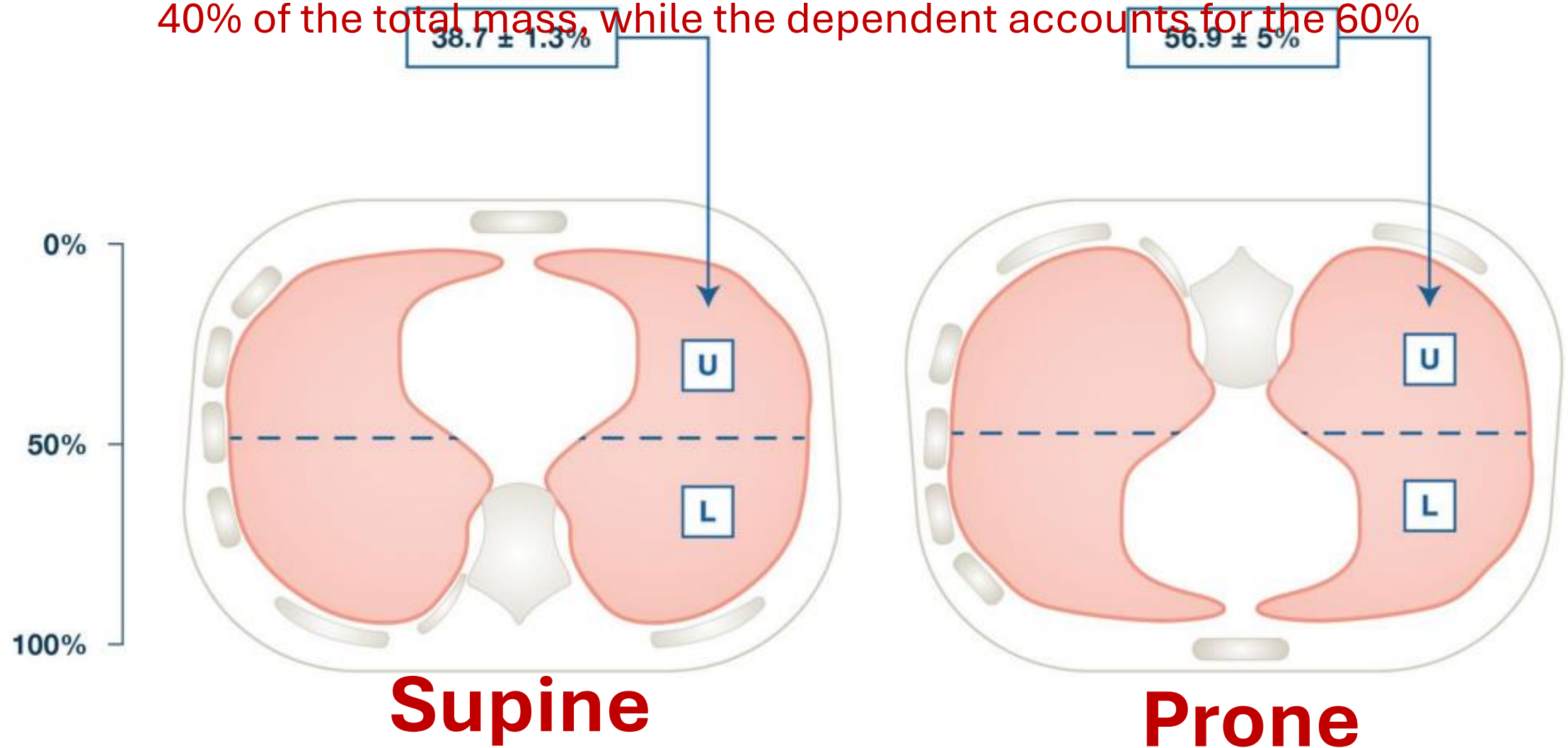


Prone Position Therapy



The open, non-dependent lung mass (at 50% of the sternum-vertebra distance) is about

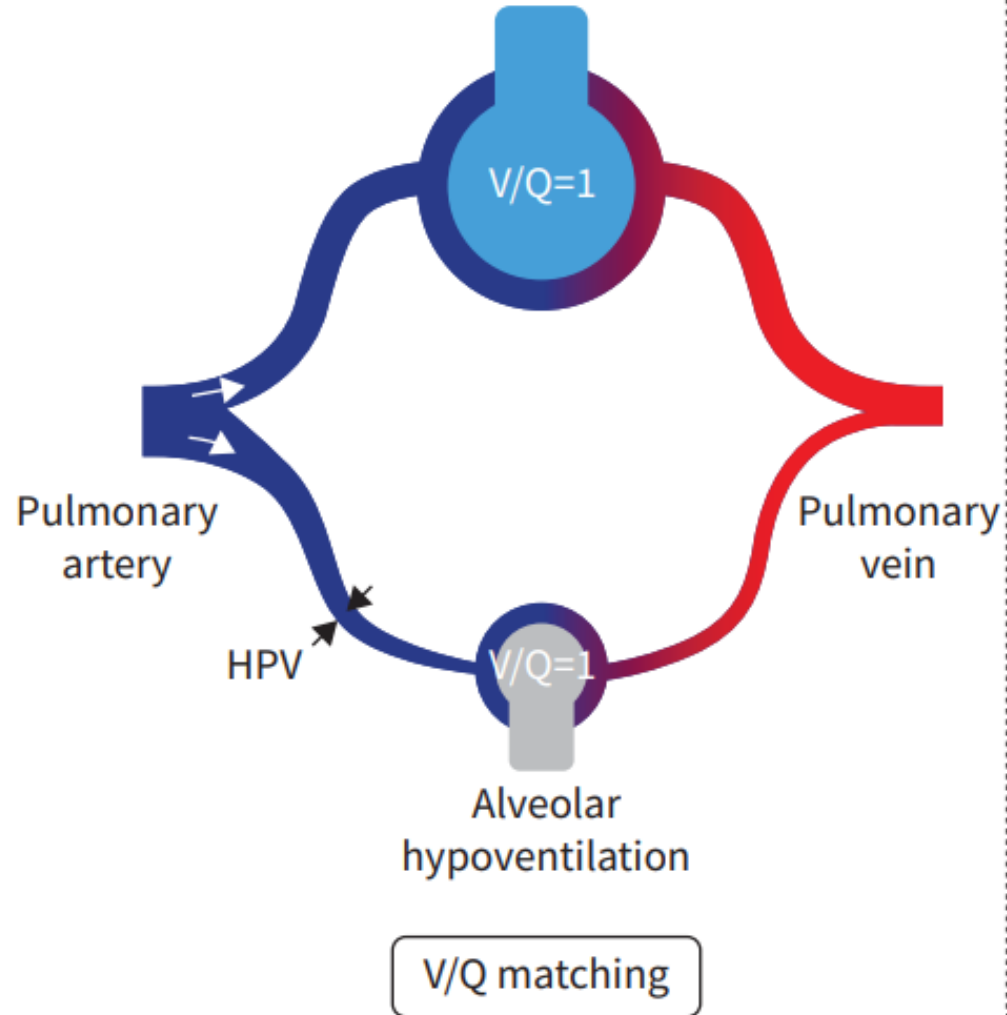
40% of the total mass, while the dependent accounts for the 60%



b)

Healthy

Normal ventilation



c)

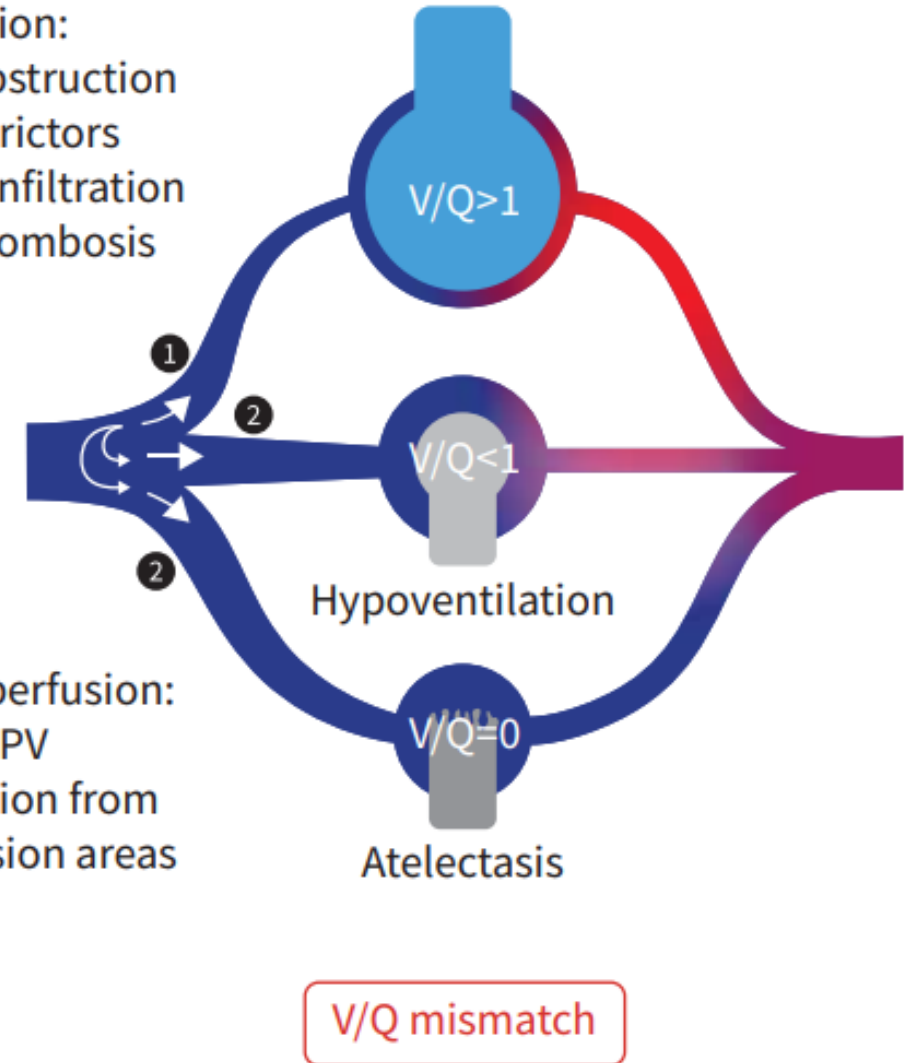
ARDS

Normal ventilation

- 1 Low perfusion:
Vascular obstruction
Vasoconstrictors
Oedema/infiltration
(Micro)thrombosis

- 2 Increased perfusion:
Impaired HPV
Redistribution from low perfusion areas

$P_{VO_2} \downarrow$
(if $CO \downarrow$)



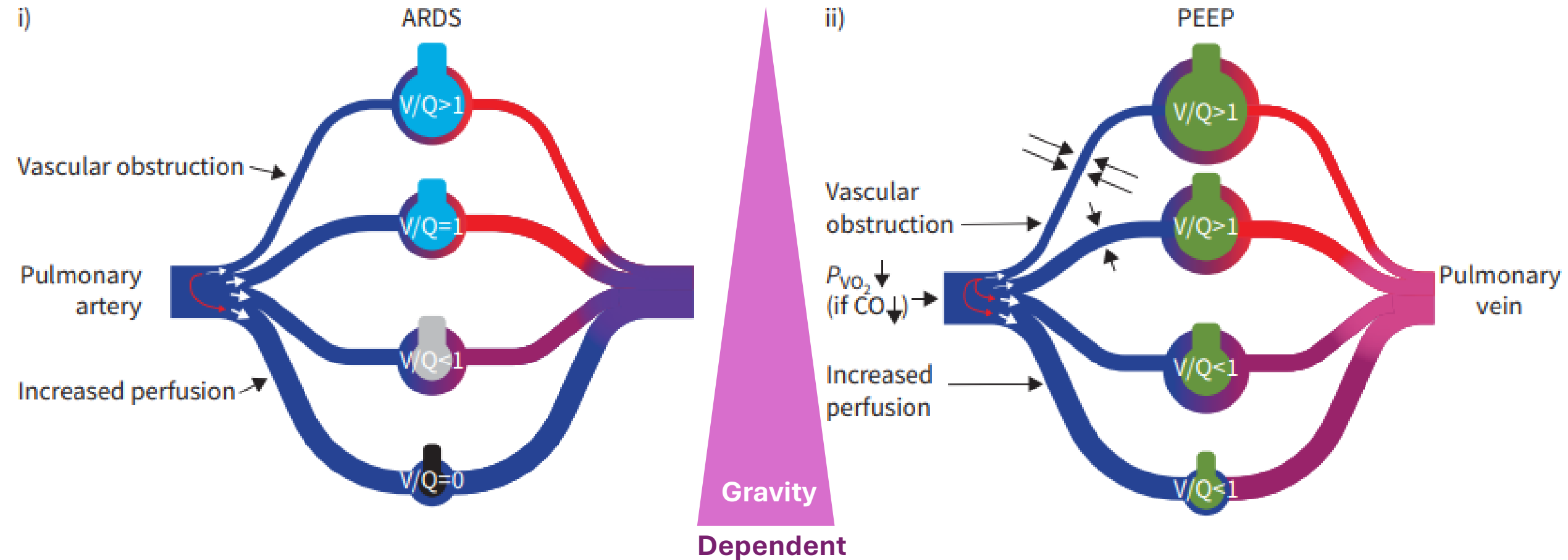
Supine

Prone

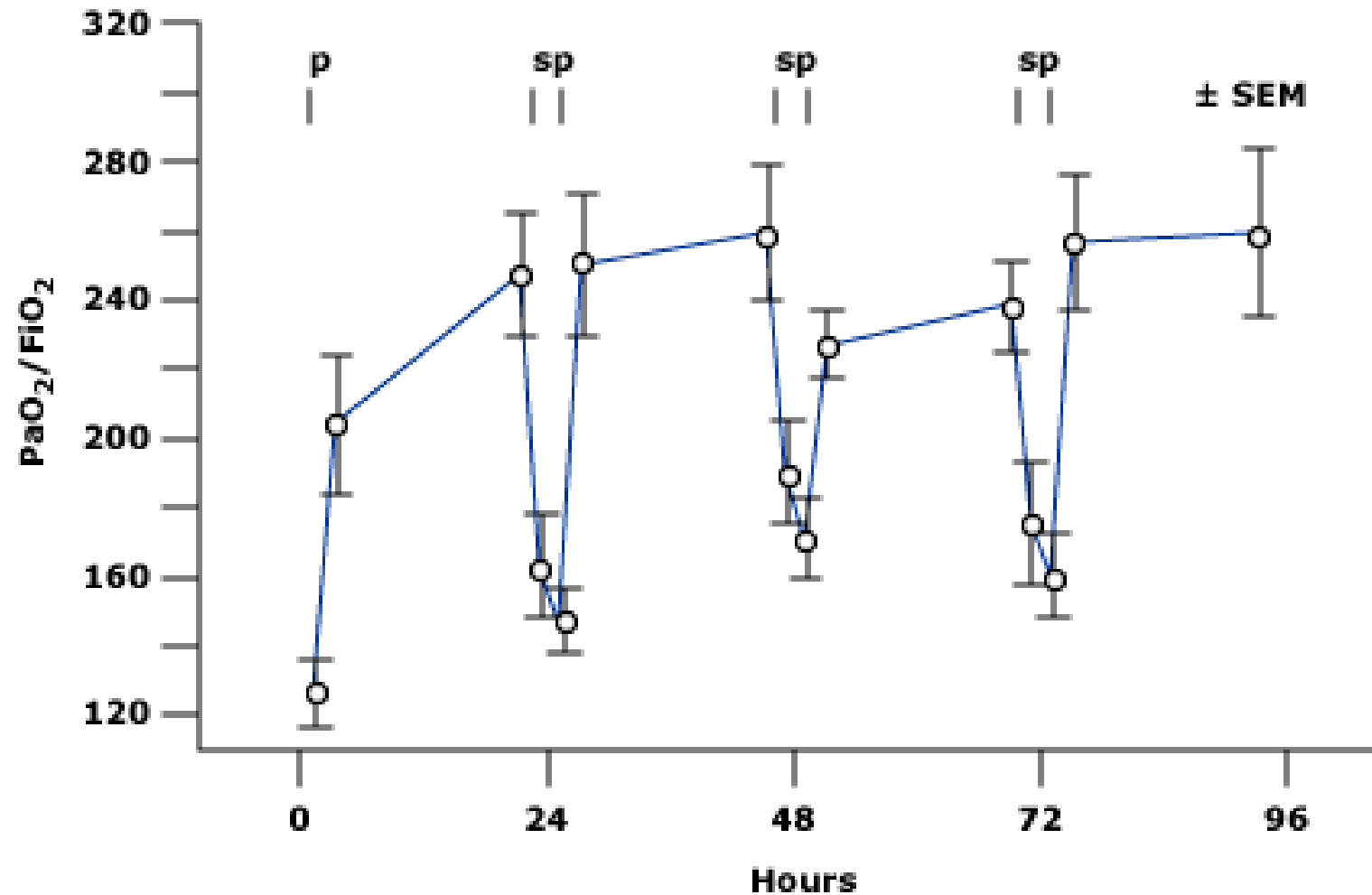
Non-Dependent

Gravity

Dependent



Course of $\text{PaO}_2/\text{FiO}_2$ during four consecutive 24-hour periods of prone positioning

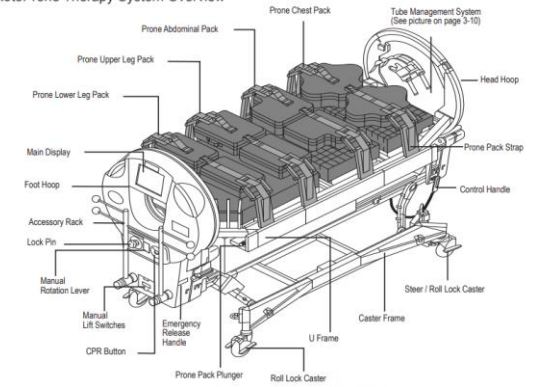


RoToProne

ARJO



RoToProne Therapy System Overview

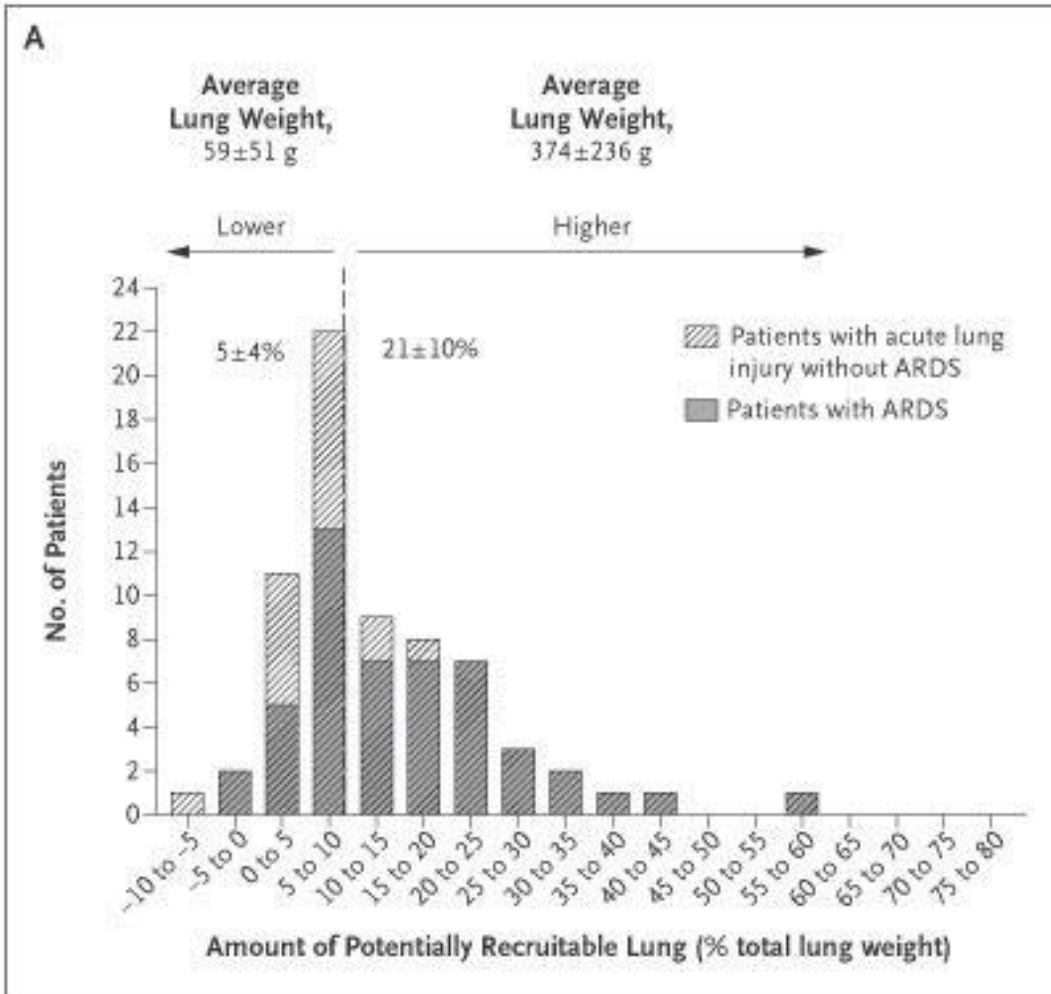


<https://www.youtube.com/watch?v=x9AD3d6zBW8>

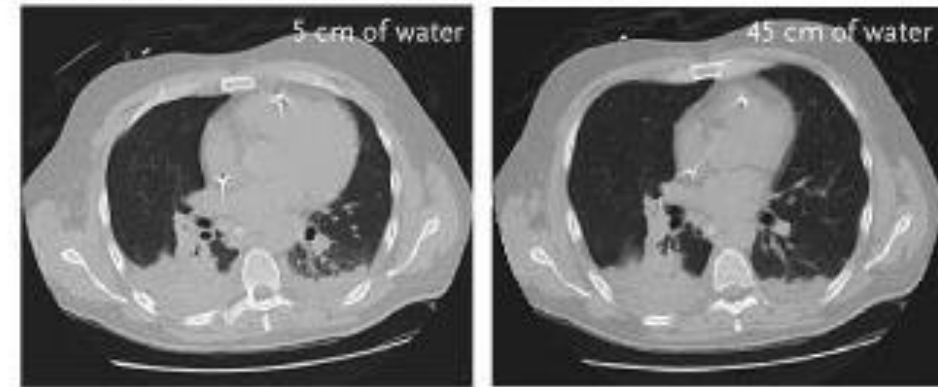
<https://www.arjo.com/en-us/products/medical-beds/critical-care>



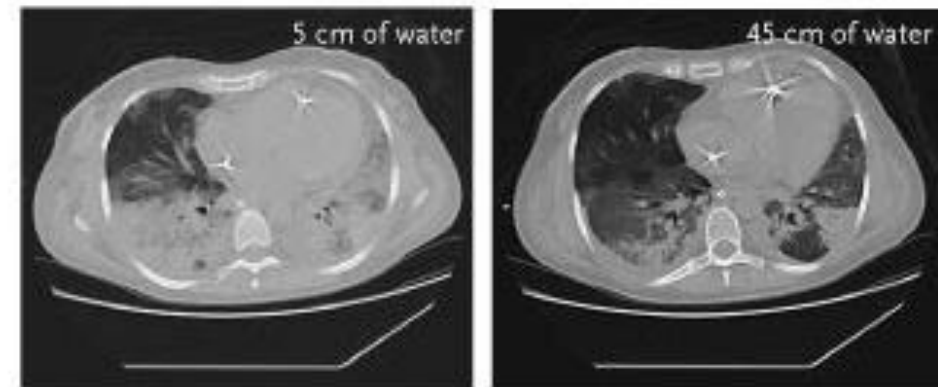
KMUH

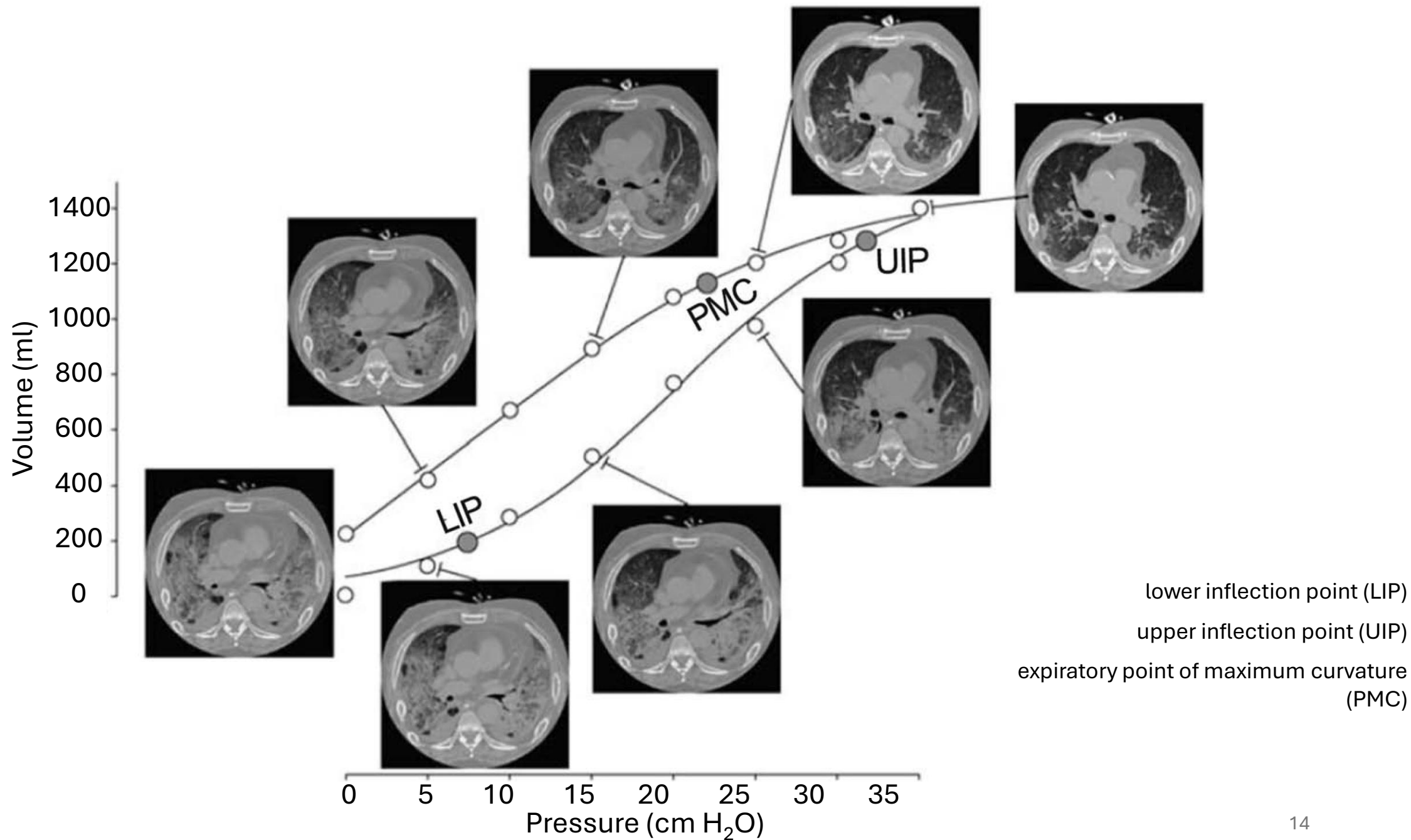


B Lower Percentage of Potentially Recrutable Lung

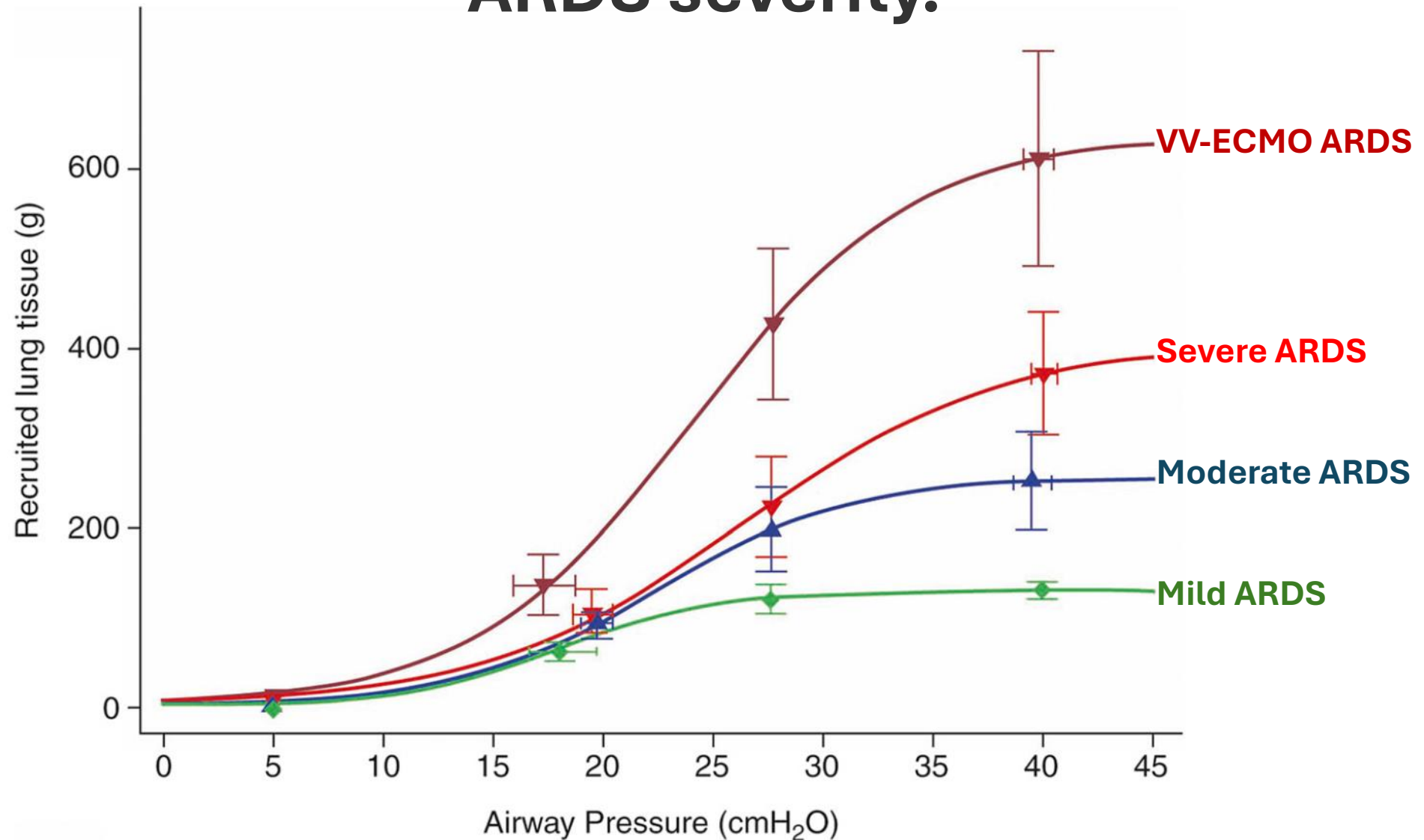


C Higher Percentage of Potentially Recrutable Lung





The amount of recruitable tissue increases with ARDS severity.





Ventilator Recruitment Maneuvers (RMs)

- **Definition:**

Temporary increase in airway and transpulmonary pressure above tidal ventilation levels.

Goal: Promote re-aeration of previously collapsed lung regions (lung recruitment).

- **Mechanism:**

Pressure during RMs typically exceeds closing pressure, aiding in the reopening of collapsed lung units.

- **Benefits:**

Increased End-Expiratory Lung Volume: Achieved after RM, potentially durable.

Improved Gas Exchange: Enhances oxygenation and CO₂ removal.

Homogenization of Alveolar Distension: Promotes uniform lung expansion.

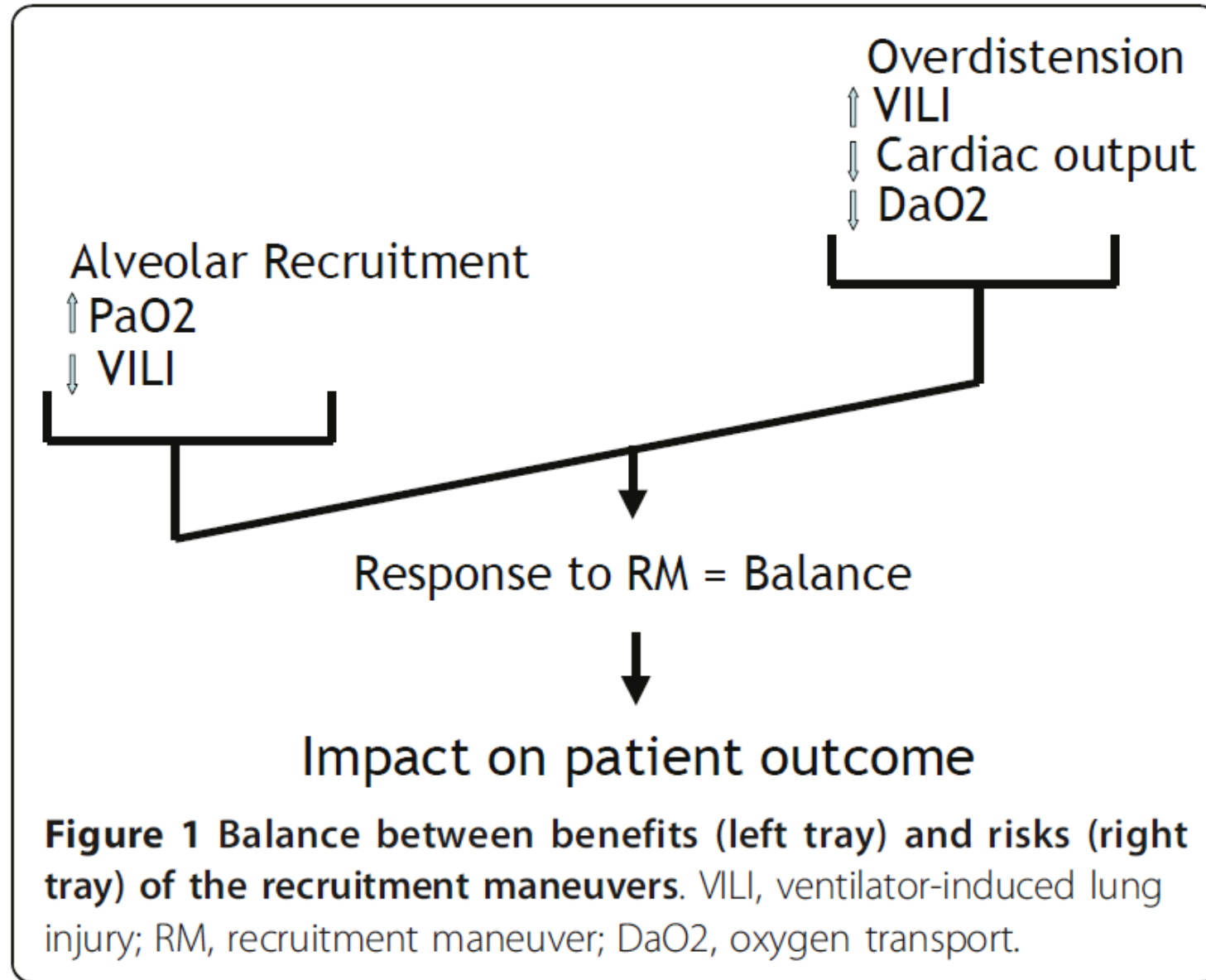
Decreased Lung Stress and Strain: Reduces risk of ventilator-induced lung injury.

- **Considerations:**

Variability in occurrence and durability of effects.

Risks of High-Pressure Maneuvers:

- **Over-Distension Complications:**
 - Barotrauma
 - Reduced venous return
 - Increased pulmonary vascular resistance
 - Right ventricular failure
 - Potential for hemodynamic collapse
- **Strategies for Performing RMs:**
- Variations in:
 - **Duration:** Length of the maneuver
 - **Pressure Target(s):** Levels of pressure applied
 - **Frequency:** How often RMs are performed
 - **Ventilator Maneuver:** Specific techniques used



Electrical Impedance tomography

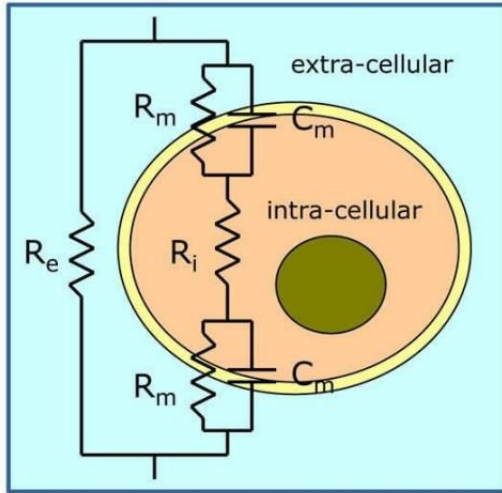


Fig.1: The equivalent electrical circuit model for tissues.

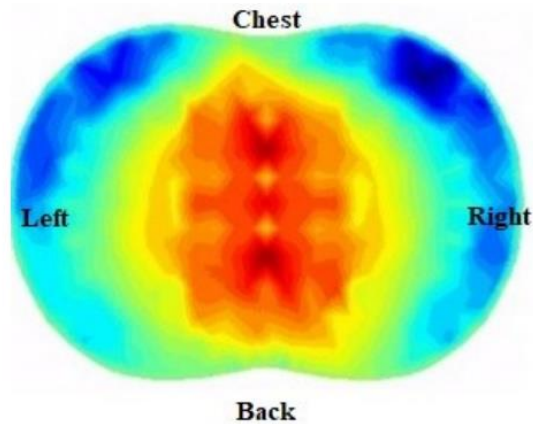


Fig. 3: Chest image reconstruction by EIT [21].

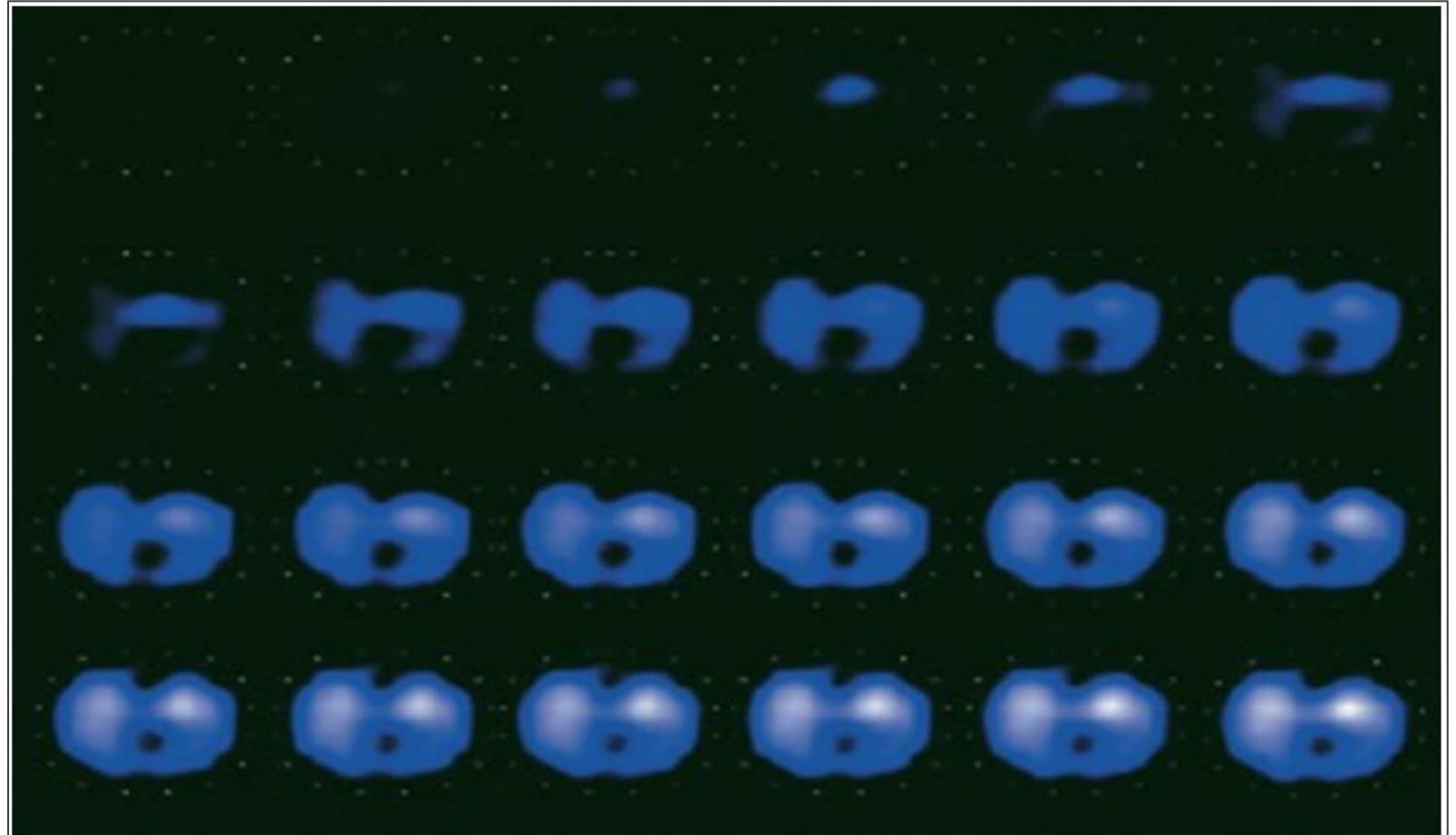


Fig. 6: Series of dynamic images showing air filling during inspiration by the PulmoVista500 system [81].

History of EIT - Applied potential tomography

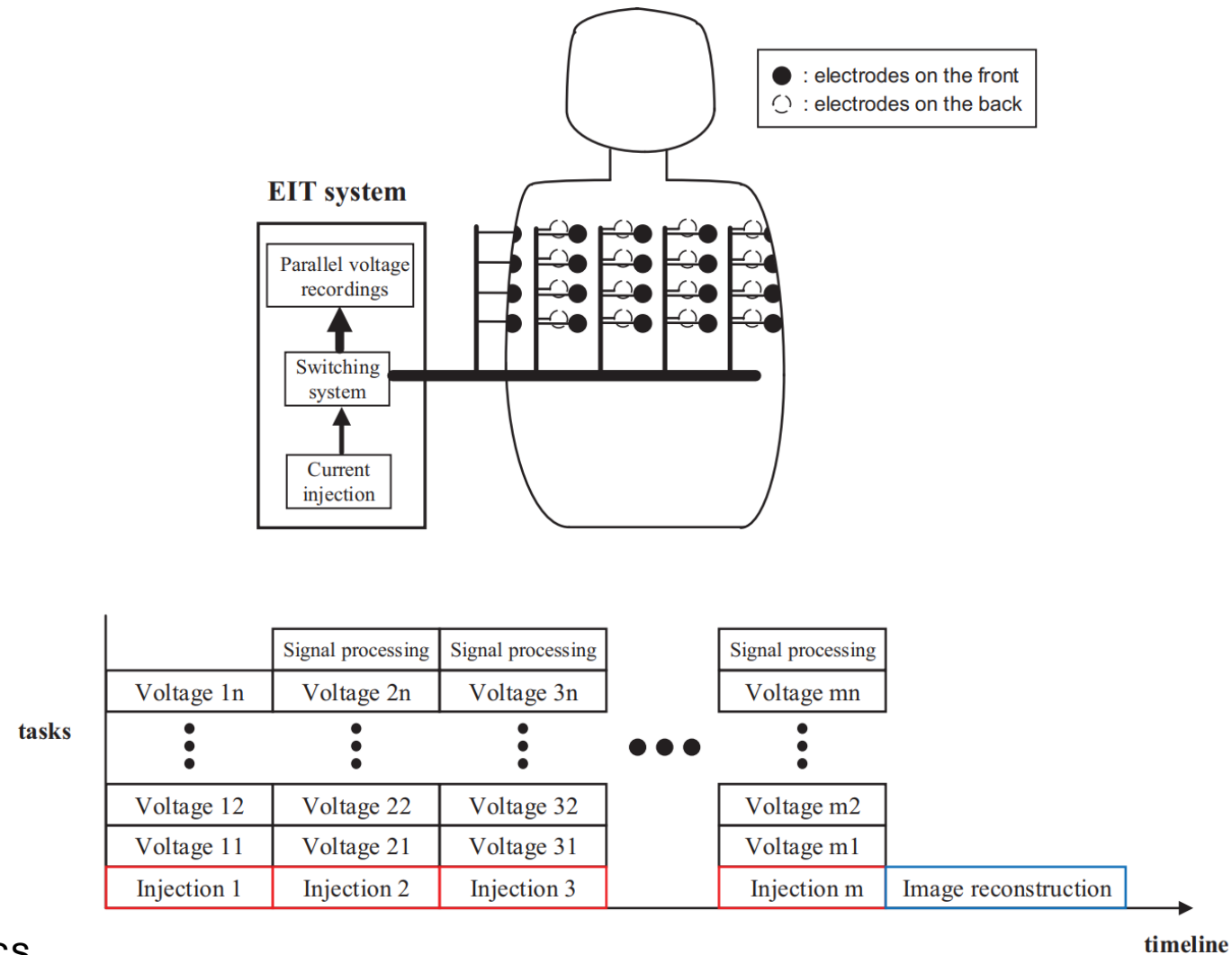
Definition and Principle

- APT is a novel imaging technique designed to detect **changes in the electrical resistivity distribution** within the human body.
- These changes reflect physiological activities such as **respiration, cardiac cycles, or fluid redistribution**.

Physiological Applications

- **Respiration:** Variations in lung air volume cause measurable resistivity changes.
- **Cardiac Cycle:** Blood flow within the thorax alters electrical conductivity during each heartbeat.
- **Fluid Redistribution:** Under simulated weightlessness, body fluid shifts produce observable impedance variations.

EIT system recording voltage in parallel



Non-invasive real time lung monitoring—the upcoming future?

基本概念

- EIT 是一種 非侵入性、無輻射、可即時影像化的肺通氣監測技術。
- 利用電流注入胸壁電極測量電壓變化，重建胸腔內的電導（或阻抗）分布。

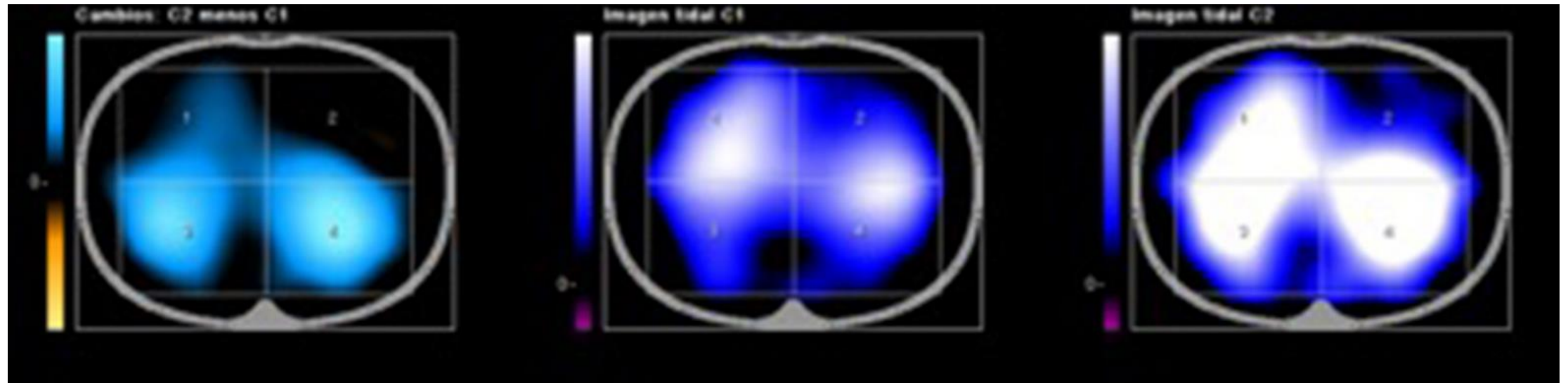
電極與測量方式

- 通常使用 **16 或 32 個電極** 的皮帶環繞胸廓（約第 4–5 肋間）。
- 小電流以兩電極注入，其餘電極量測電壓；逐對切換，共形成 **208 組量測值（frames）**。
- 每秒可重建約 **50 幅影像**，顯示胸腔阻抗變化。

生理意義

- 空氣量增加 → 阻抗上升（導電性下降）。
- 血液或液體增加 → 阻抗下降（導電性上升）。
- 因此能即時偵測肺內氣體含量、液體堆積或血流變化。

Non-invasive real time lung monitoring—the upcoming future?



Basal EIT

PEEP 5-10

PEEP 10

Real-time effects of PEEP and tidal volume on regional ventilation and perfusion in experimental lung injury

Experimental Subjects

- A total of **11 piglets** were studied: **4 healthy controls** and **7 with lung injury** induced by repeated saline lung lavages. Due to one cardiovascular event, **10 animals completed the experiment** and were included in the final analysis.

Ventilation Settings

- Mechanical ventilation was applied using **three tidal volumes (VT) of 7, 10, and 15 mL/kg**, and **four levels of positive end-expiratory pressure (PEEP): 5, 8, 10, and 12 cmH₂O**. All **12 combinations** of VT and PEEP were tested in **randomized order** for each animal.

Electrical Impedance Tomography (EIT) Monitoring

- Each pig had **32 electrodes** evenly placed around the thoracic circumference to record impedance changes within the lung. The lungs were divided into **three gravity-dependent regions of interest (ROIs): nondependent, middle, and dependent** regions.
- Ventilation maps were obtained from **changes in air content**, while perfusion maps were derived by injecting **10 mL of 10% NaCl** as a conductive contrast agent during an **end-expiratory breath-hold**. The resulting impedance changes were analyzed to **estimate regional pulmonary blood flow**.

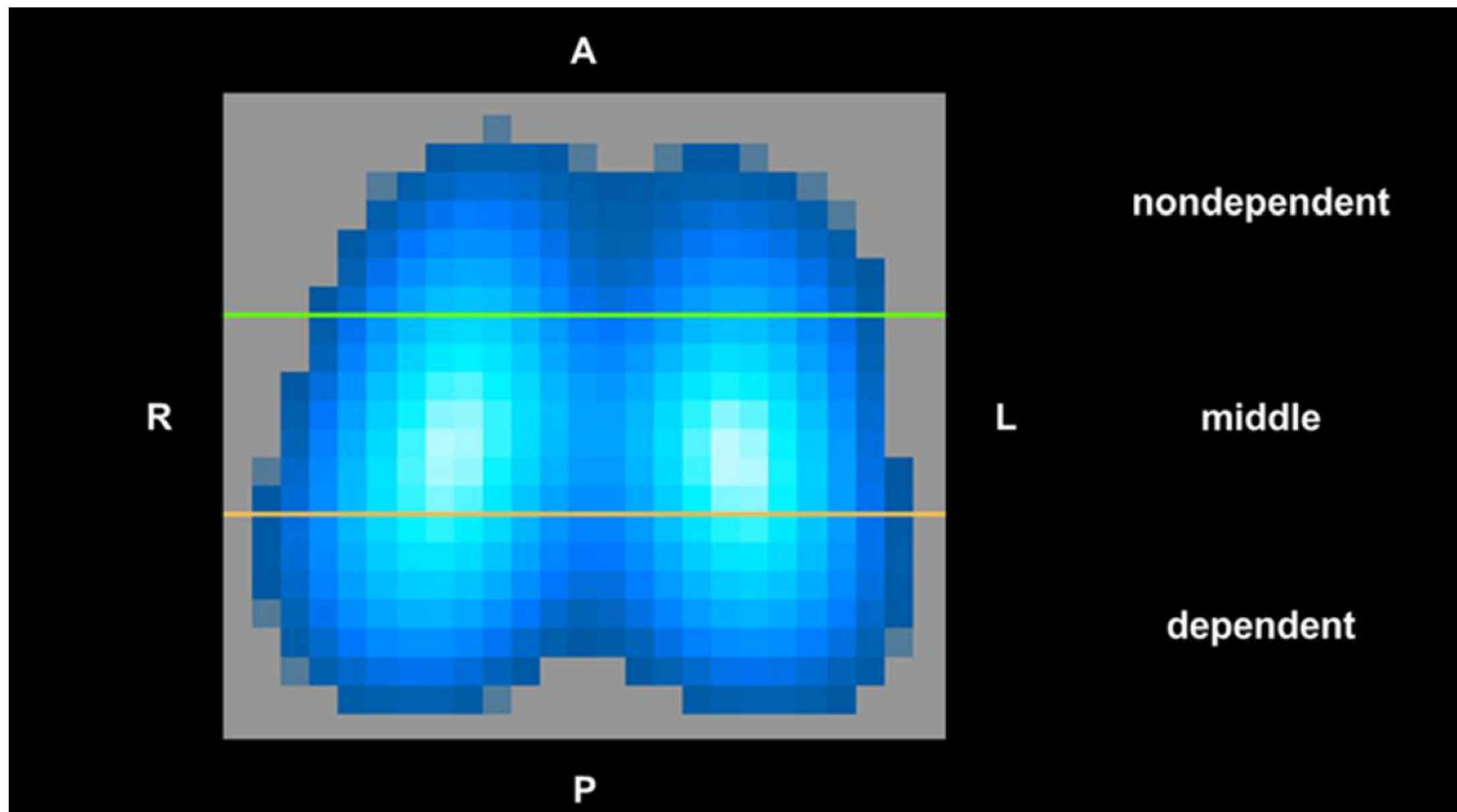
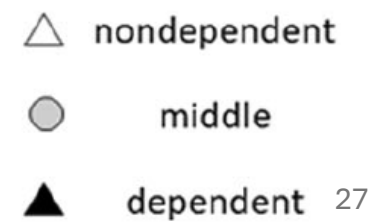
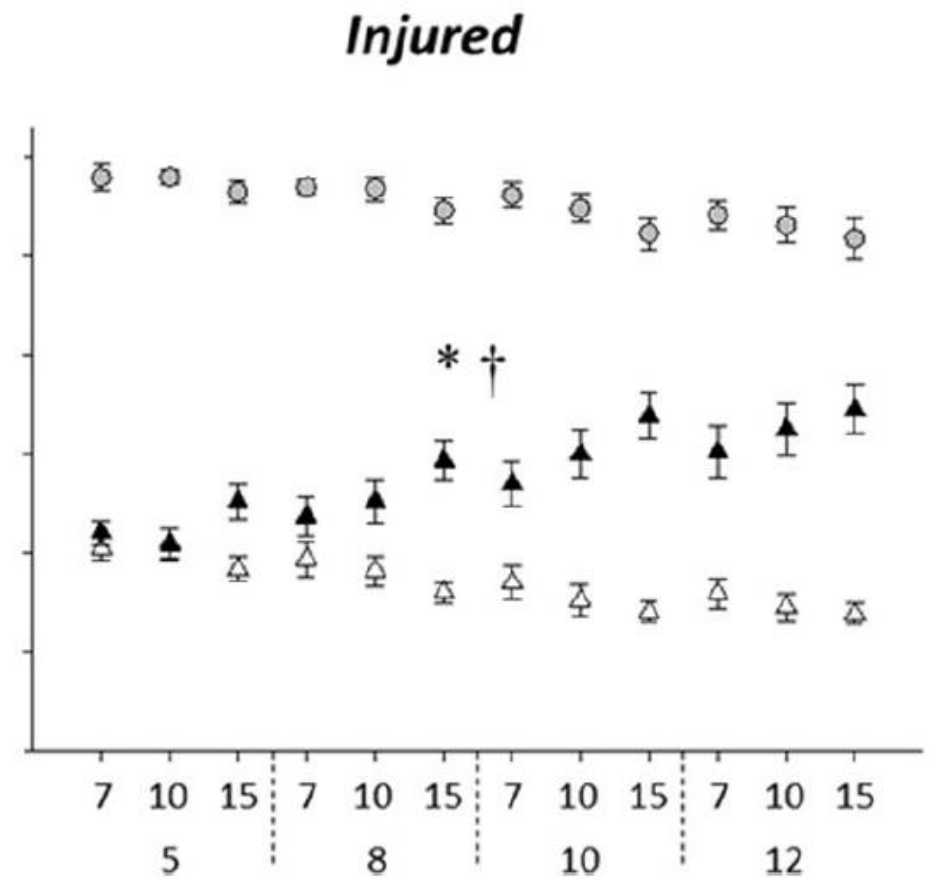
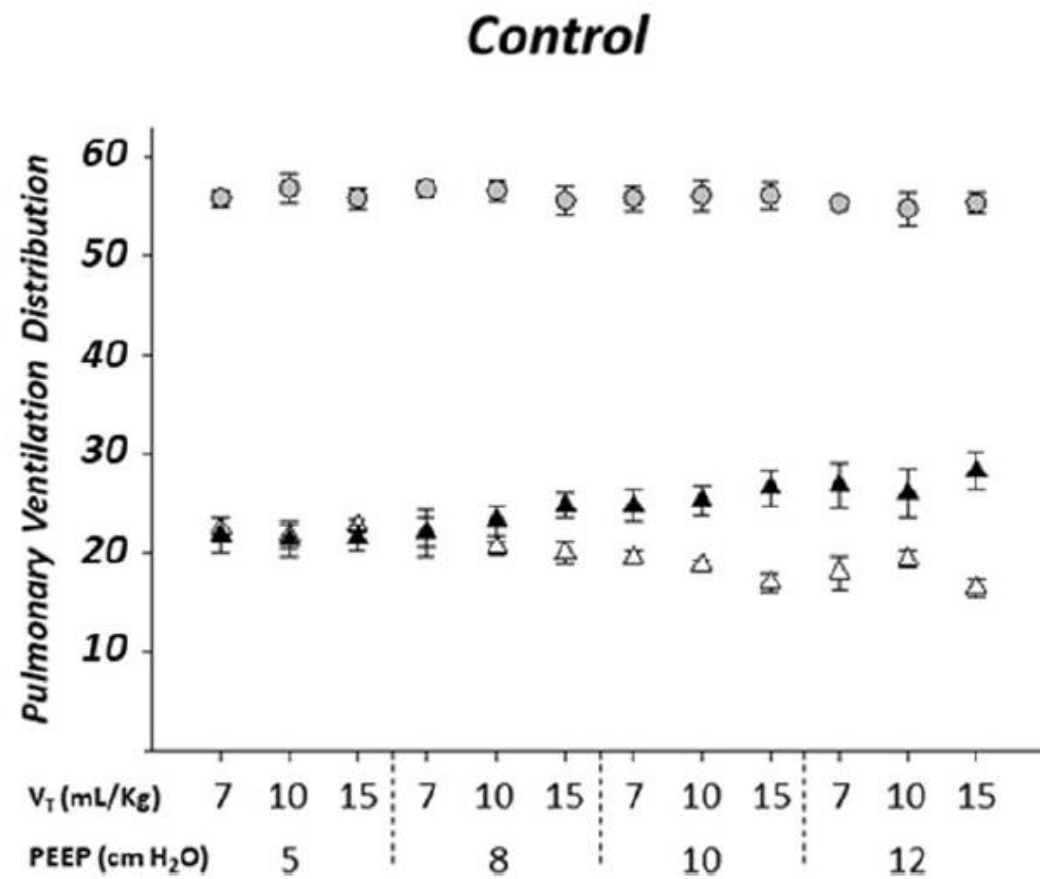
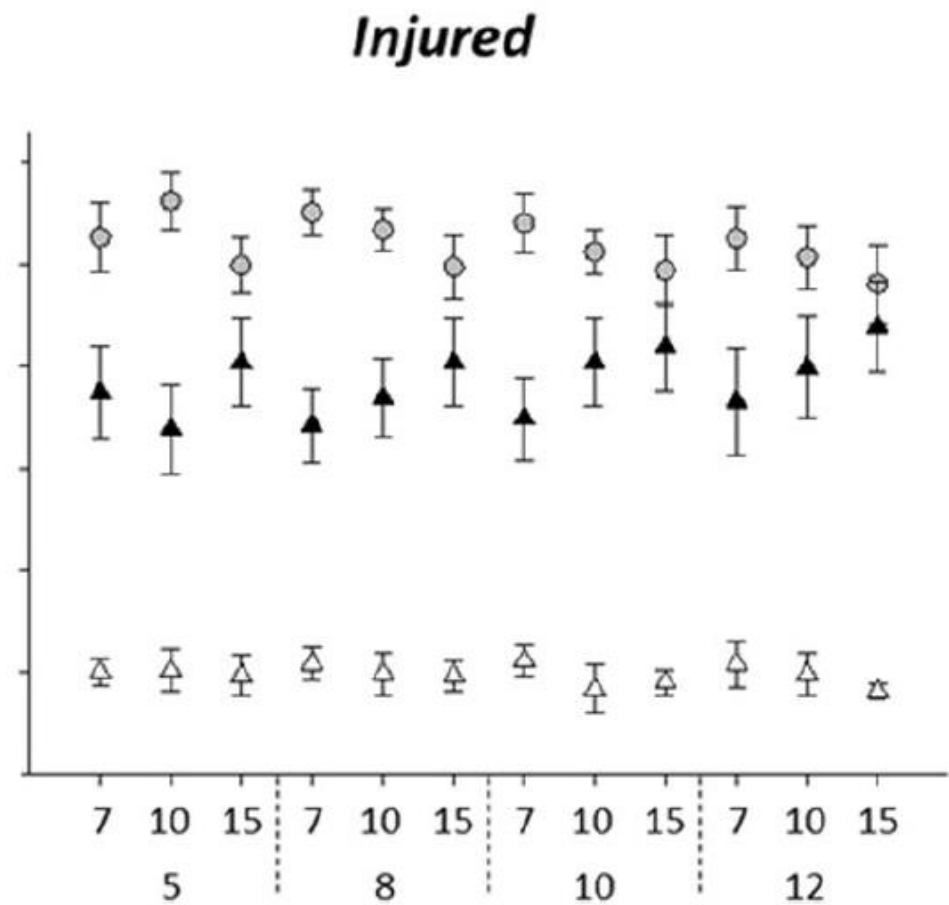
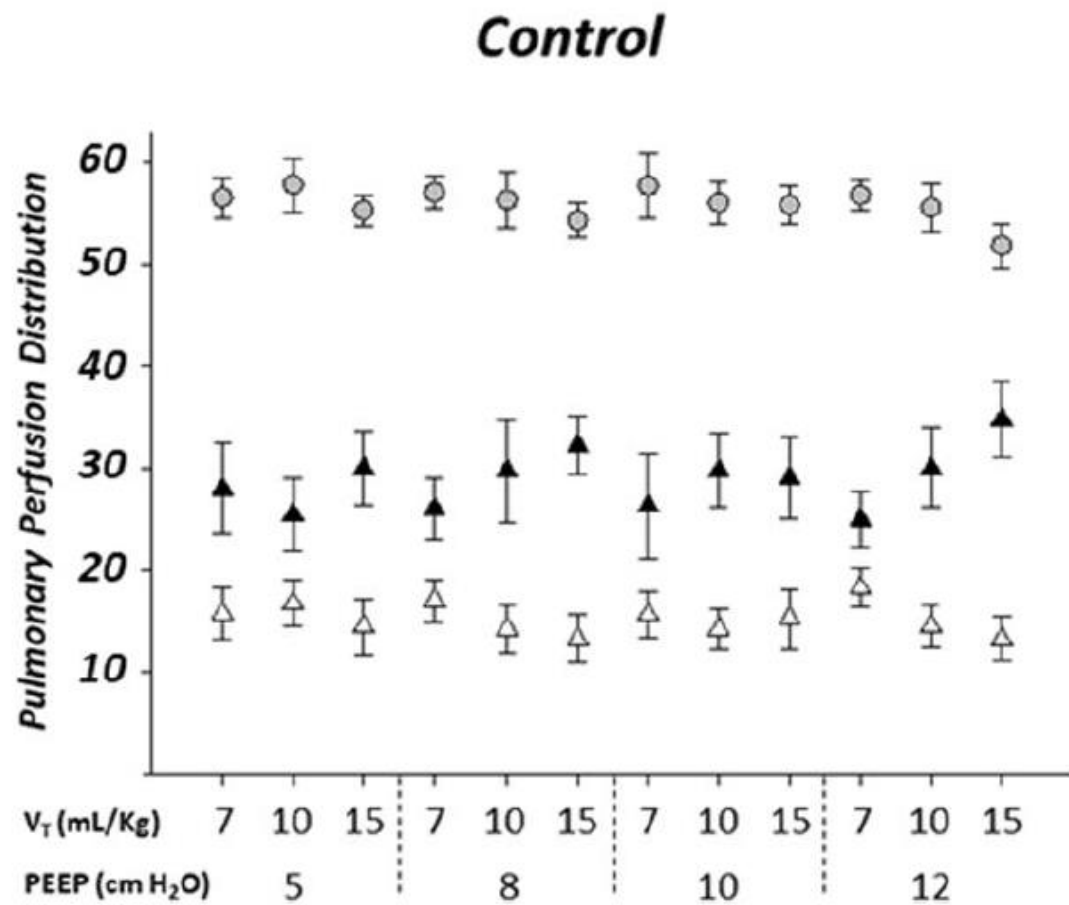


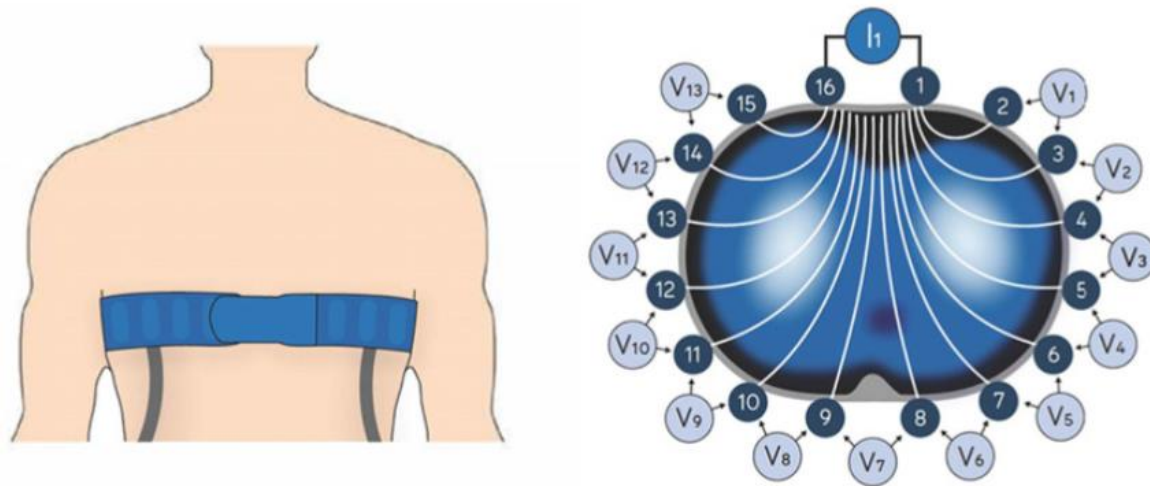
Table 1 Cardiopulmonary parameters

Parameter	Control				Injured						Control	Injured
	C1	C2	C3	C4	I1	I2	I3	I4	I5	I6	Mean (SD)	Mean (SD)
Weight (kg)	30	33	30	33	30	28	29	33	31	31	31 (2)	30 (2)
Lavage volume (L)	0	0	0	0	4	5	2	4	9	2	0 (0)	4 (3)
P/F ratio (mmHg)	287	384	441	373	158	206	153	104	164	143	371 (64)	154 (33)*
ABP (mmHg)	65	78	89	79	68	95	97	97	75	62	78 (10)	82 (16)
PAP (mmHg)	20	27	14	20	34	27	27	20	32	34	20 (5)	29 (6) #
CVP (mmHg)	6	11	11	6	13	8	11	6	8	12	9 (3)	10 (3)
HR (bpm)	75	87	82	143	97	105	98	107	107	93	97 (31)	101 (6)
CO (L/min)	1.9	4.0	3.0	6.1	3.6	3.9	3.2	4.0	4.1	3.4	3.8 (1.8)	3.7 (0.4)



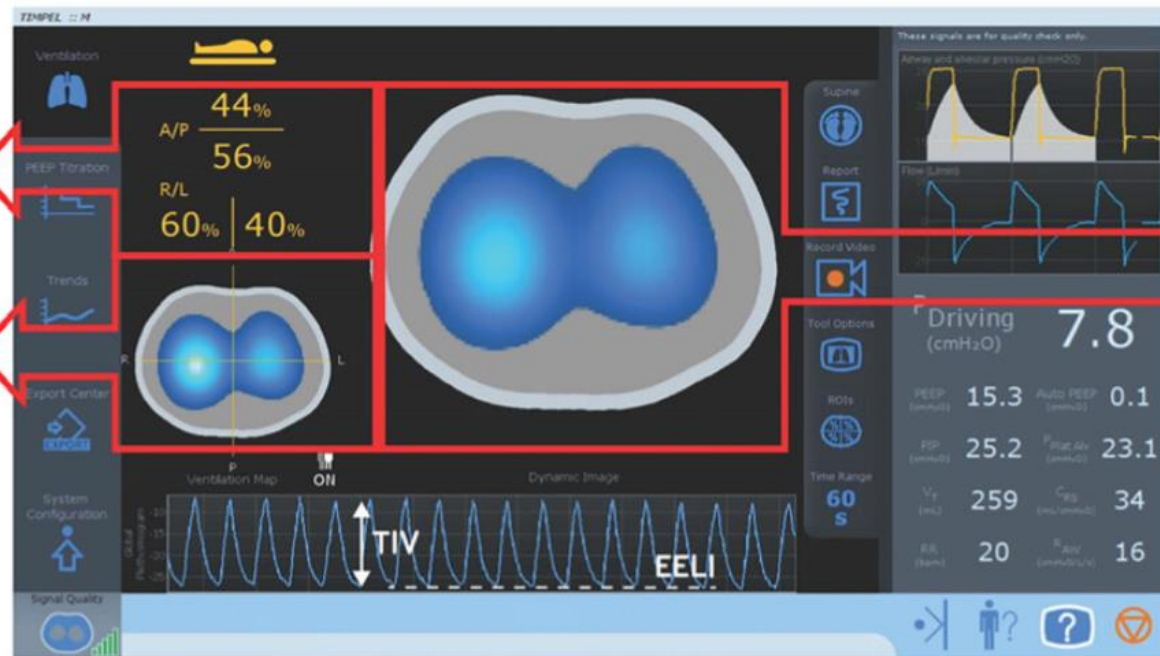


- △ nondependent
- middle
- ▲ dependent



Distribution of ventilation depending on prespecified ROI

Tidal image: distribution of tidal volume over the last breath, with selected region of interest



Dynamic image: real time distribution of tidal volume over time

Table 1. Definition, Clinical Application, and Limitations of the Most Commonly Used EIT Indices in Patients with Hypoxemic Respiratory Failure

EIT Indices	Definition	Potential Applications	Limits
TIV (tidal impedance variation)	Variation of impedance values between end of inspiration and end of expiration (amplitude of plethysmogram signal)	• Reflects global and regional relative distribution of V_T in lungs during ventilation • Contributes to identification of pleural effusion, pneumothorax, endotracheal tube malposition, pendelluft	• Cannot be used for detailed anatomic diagnosis • Only relative distribution of V_T
EELI (end-expiratory lung volume)	Regional lung impedance at end of expiration	• EELI changes are correlated with EELV variation • Reflects lung recruitment/derecruitment • Contributes to identification of pneumothorax	• Affected by factors other than lung volume (inflatable mattress, fluid infusion)
Global inhomogeneity index	Sum of absolute differences between median TIV and each pixel TIV normalized to sum of each pixel TIV	• Reflects heterogeneity of ventilation	• Does not reflect local distribution of TIV • Does not take into account overdistension, collapse, or any other pathological situation

Table 1. Definition, Clinical Application, and Limitations of the Most Commonly Used EIT Indices in Patients with Hypoxemic Respiratory Failure

EIT Indices	Definition	Potential Applications	Limits
Overdistension and collapse estimation	Estimation of relative local compliance loss presented as percentage of collapse and overdistension during decremental PEEP trial	• Selection of PEEP level that jointly minimizes collapse and overdistension • Applicable even on venovenous ECMO when V_T is very low	• Assumes that ΔP on ventilator is reliable surrogate of regional ΔP • Depends on highest and lowest values of PEEP applied during PEEP trial
Regional ventilation delay	Time between start of inspiration and aeration of lung regions	• Identify recruitable lung regions and cyclic opening/closing phenomenon • Could be used to identify best PEEP level with most homogenized tidal inflation	• Requires a slow inflation maneuver with constant flow • Depends on defined threshold • Does not detect not-recruitable or overdistended regions
Center of ventilation	Describes geometrical center of ventilation	• Reflects global ventilation homogeneity to guide and monitor impact of mechanical ventilation management or adjunct therapies (i.e., prone positioning) • Identification of adverse events (e.g., malpositioning of endotracheal tube or development of pneumothorax)	• Reflects ventilation shifts but not their origin • Imprecise interpretation of ventilation distribution (ventilation on one side can be compensated by other side). • Setting PEEP aiming for >50% dorsal distribution could imply overdistension of whole lung
Dorsal fraction of ventilation	Describes percentage of V_T distributed in dorsal region		

Table 2. Summary of Recent Clinical Trials Evaluating the Outcomes of an EIT-based PEEP in Patients with ARDS

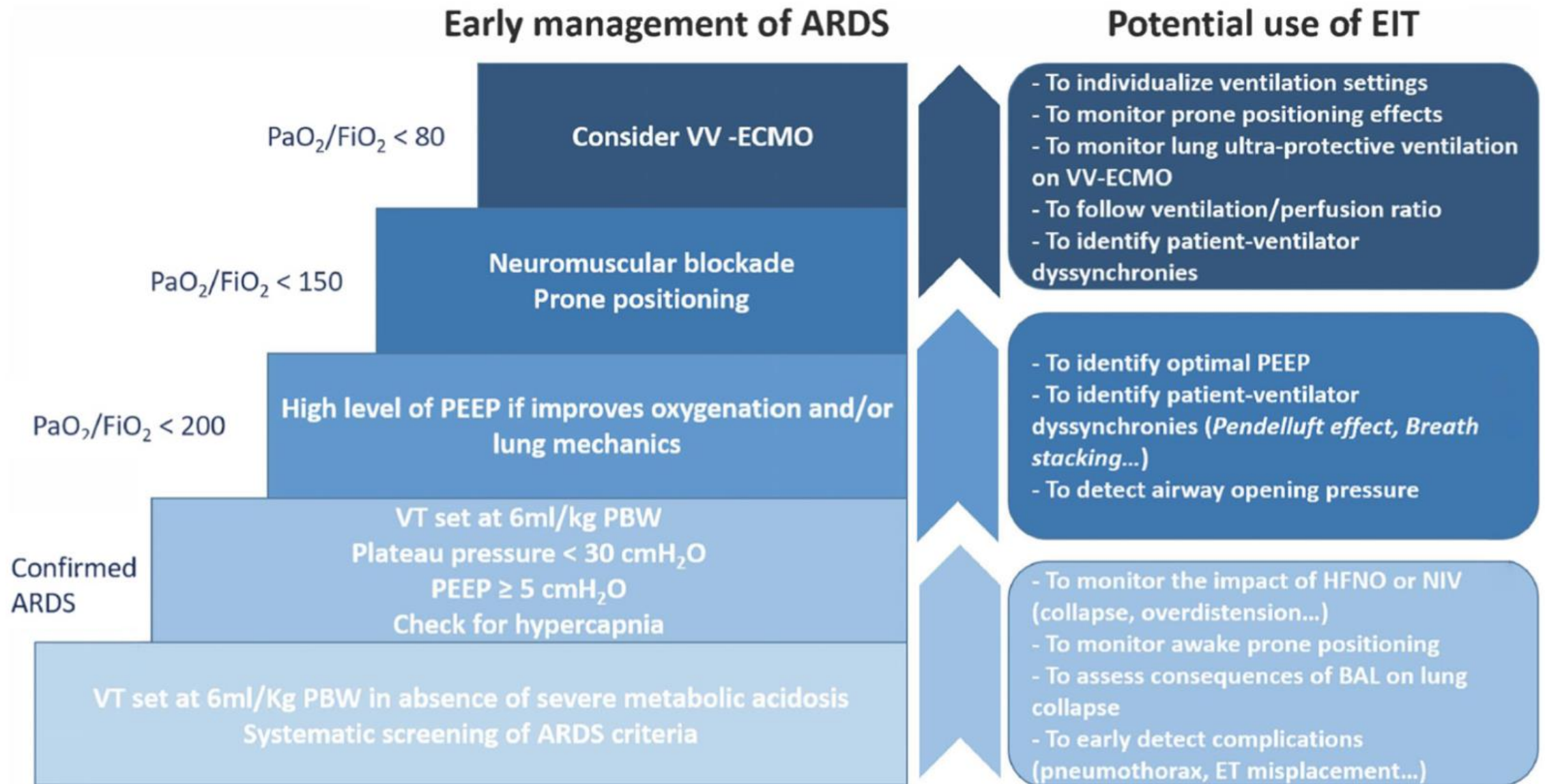
Population	Severity of the Population	Mean PEEP in the EIT group	Mean PEEP in the control group	Results
24 patients with ARDS vs. 31 historical patients (26)	APACHE II score 23.5	Based on overdistension and collapse: 18 cm H ₂ O	2 cm H ₂ O above inflection point of pressure–volume curve: 14 cm H ₂ O	- Higher compliance in EIT-based PEEP group: 26 vs. 20 ml/cm H₂O - No difference in clinical outcomes
117 patients with ARDS (83)	19.7% severe ARDS, 45.3% moderate ARDS, APACHE II score 19	Based on overdistension and collapse: 8 cm H ₂ O	Based on PEEP/FiO ₂ table: 8 cm H ₂ O	- Nonsignificant 28-d mortality in EIT group: 21% vs. 27% - ICU length of stay: 13 vs. 10 d - Ventilator-free days: 14 vs 23 d
87 patients with ARDS (81)	75.9% severe ARDS, inclusion ≤24 h after intubation	Based on overdistension and collapse: 16.2 cm H ₂ O	Based on pressure–volume curve: 17.4 cm H ₂ O	- Lower hospital mortality in EIT group: 44.4% vs 69.0% - Lower ΔP in EIT group: 10.9 vs. 12.4 cm H₂O
12 patients with ARDS with a cross-over protocol (84)	PaO ₂ /FiO ₂ , 130; MV duration at inclusion, 0.6 d; SAPS II, 38	EIT-selected PEEP was lower: mean difference of change, –2 cm H ₂ O		- Lower mechanical power in EIT group: 11.42 J/min, with lower ΔP and higher static respiratory system compliance

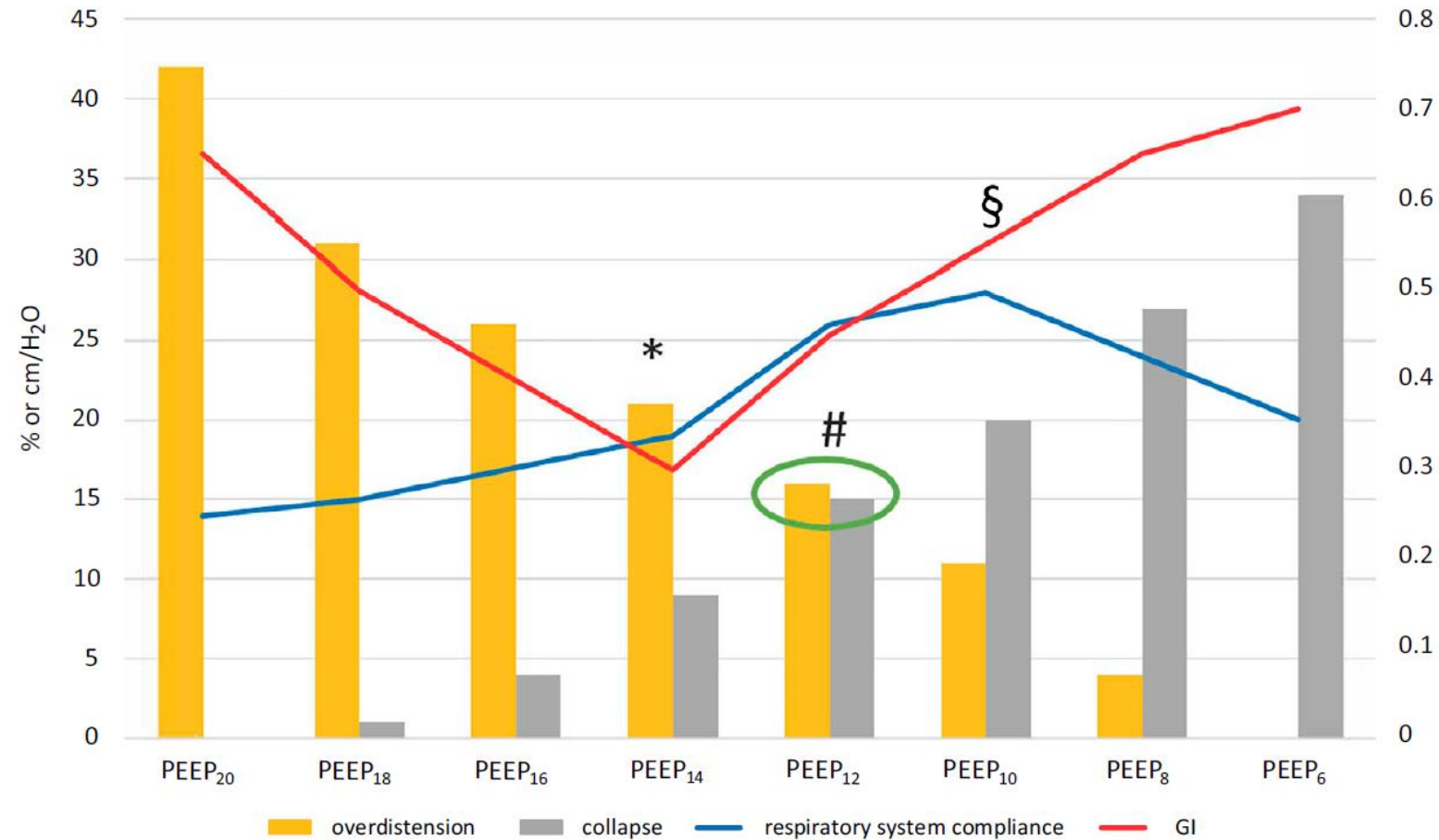
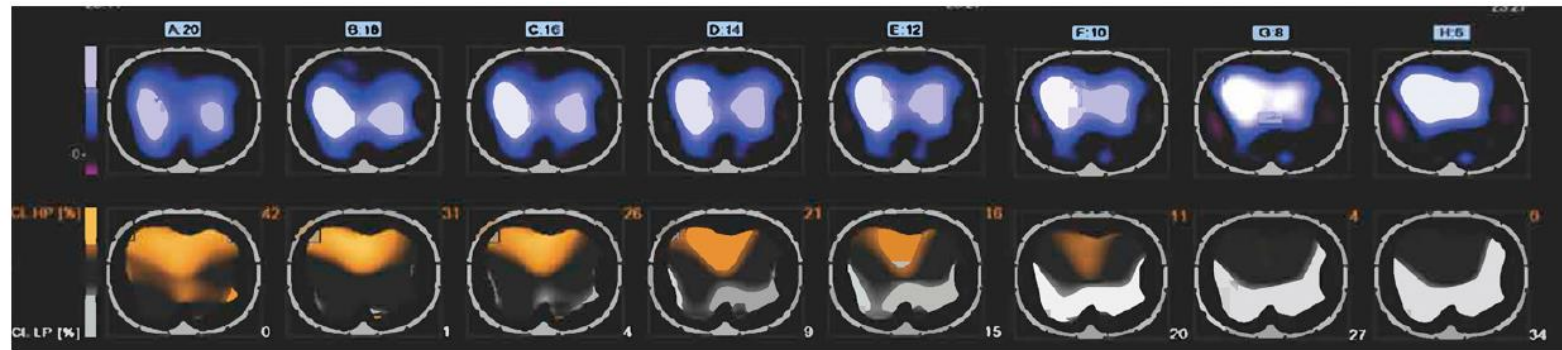
Table 3. Main Advantages and Limitations of EIT and Future Improvements to Address Limitations

Main Advantages	Main Limitations	Direct Impact on Clinical Practice	Potential Future Improvements
• Real-time monitoring of lung ventilation • Evaluation of regional compliance before and after therapeutic procedures (e.g., recruitment) • Visualization and quantification of overdistension and collapse • Identification of pathological situations (inhomogeneity, Pendelluft effect)	• Low spatial resolution • Most sensitive to detect electrical impedance changes occurring in plane of electrode belt • Performance, relevance, and reference values of EIT indices still lacking	• EIT cannot be used for precise anatomic diagnosis • EIT analysis cannot be extrapolated to lung part outside visualized area • EELV cannot be precisely inferred from EELI • Performance/relevance of available EIT indices to assess optimal PEEP have not been compared • Normal ranges of EIT indices unknown and may vary between patients	• Technological improvement and/or increase in number of electrodes on belt • Further studies on potential benefit of combining apical EIT analysis with conventional one • Best EIT indices for each purpose need to be identified in patients with ARDS • “Normal ranges” should be accurately evaluated

Table 3. Main Advantages and Limitations of EIT and Future Improvements to Address Limitations

Main Advantages	Main Limitations	Direct Impact on Clinical Practice	Potential Future Improvements
EIT can be coupled with ventilator for further physiological analysis	Lung perfusion assessment insufficiently developed	- Most EIT devices still do not perform lung perfusion analysis - Evaluation currently requires apnea and hypertonic saline bolus	- Integration of lung perfusion module in all EIT devices. - Promising method for lung perfusion without saline bolus should be further investigated
Noninvasive continuous bedside monitoring	Unknown clinical benefit of daily EIT-guided mechanical ventilation strategy	Clinical impacts of ventilation strategy guided by EIT remain unknown	Few clinical trials are ongoing (NCT04247477, NCT03793842, NCT03112512) to assess potential benefit of EIT-guided mechanical ventilation on outcomes in patients with ARDS

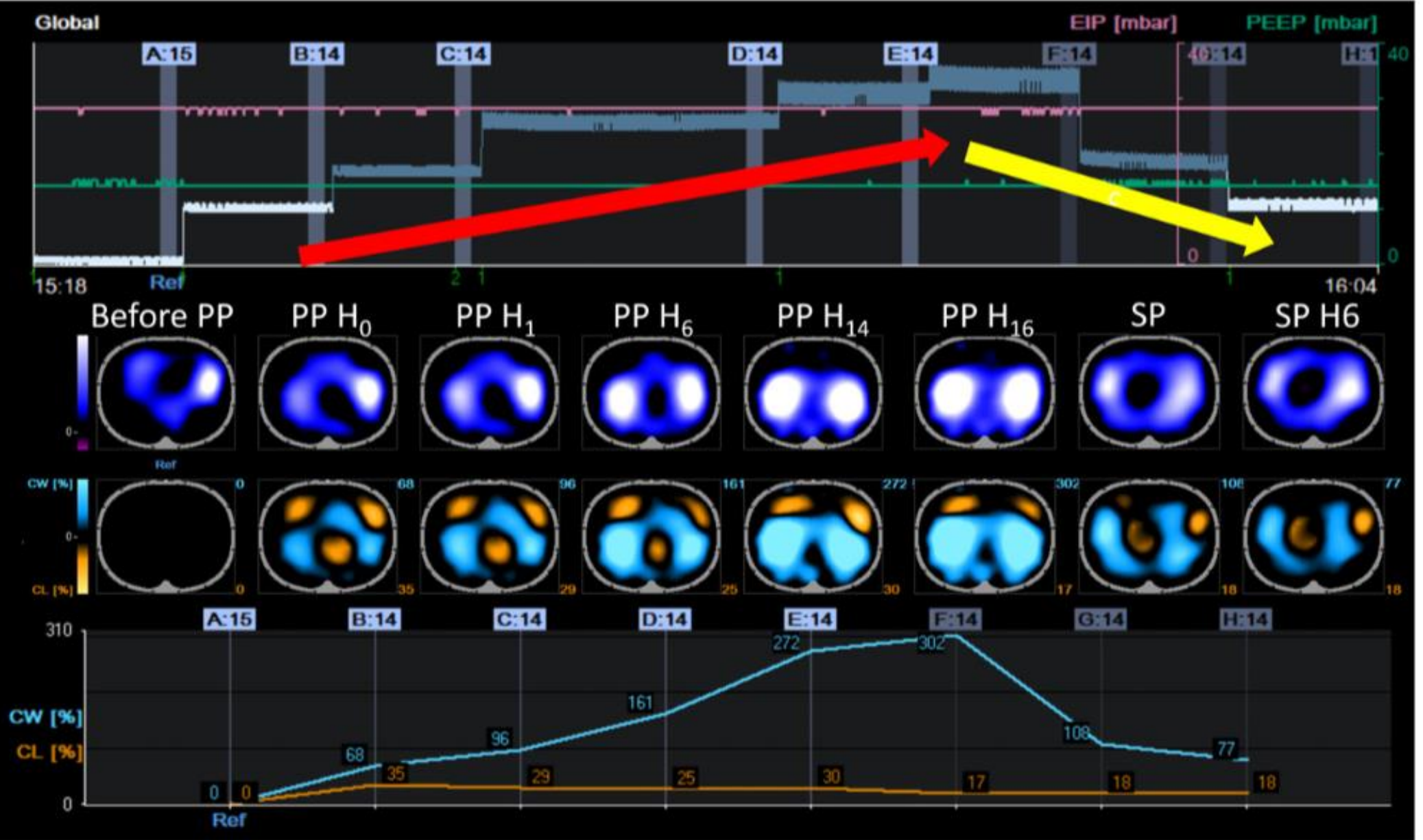




Tidal impedance and EELI variations

Local tidal impedance variation

Local compliance variation



Step 1. Setup and Calibration

- Place **16–32 electrodes** around the thorax (4th–5th intercostal space).
- Ensure good **skin contact**; avoid bandages or subcutaneous emphysema.
- Keep patient position stable to reduce motion artifacts.
- Calibrate baseline end-expiratory lung impedance (EELI).

Step 2. Baseline Monitoring

- Observe **tidal image (TIV)** and **end-expiratory impedance (EELI)** in real time.
- Identify regions of:
 - Hypoventilation or collapse.
 - Overdistension or pneumothorax.
 - Tube malposition or regional derecruitment after suctioning/BAL.

Step 3. PEEP Titration using EIT

- **Goal:** Find the compromise between alveolar recruitment and overdistension.
 1. Perform a **decremental PEEP trial** (keep other settings constant).
 2. For each PEEP level, calculate:
 1. Regional compliance changes.
 2. Percentage of overdistension vs. collapse.
 3. Plot both curves → Choose the **intersection point** as *optimal PEEP* ($\approx +2$ cmH₂O transpulmonary pressure).
 4. Confirm with improved dorsal ventilation and reduced heterogeneity.

Step 4. Adjust Tidal Volume (VT)

- Compare **regional compliance** before and after halving driving pressure.
 - ↑ Compliance → Overdistension → Decrease VT.
 - ↓ Compliance → Collapse → Consider higher PEEP.
- Target: Better oxygenation + lower regional ventilation delay.

Step 5. Positioning Strategies

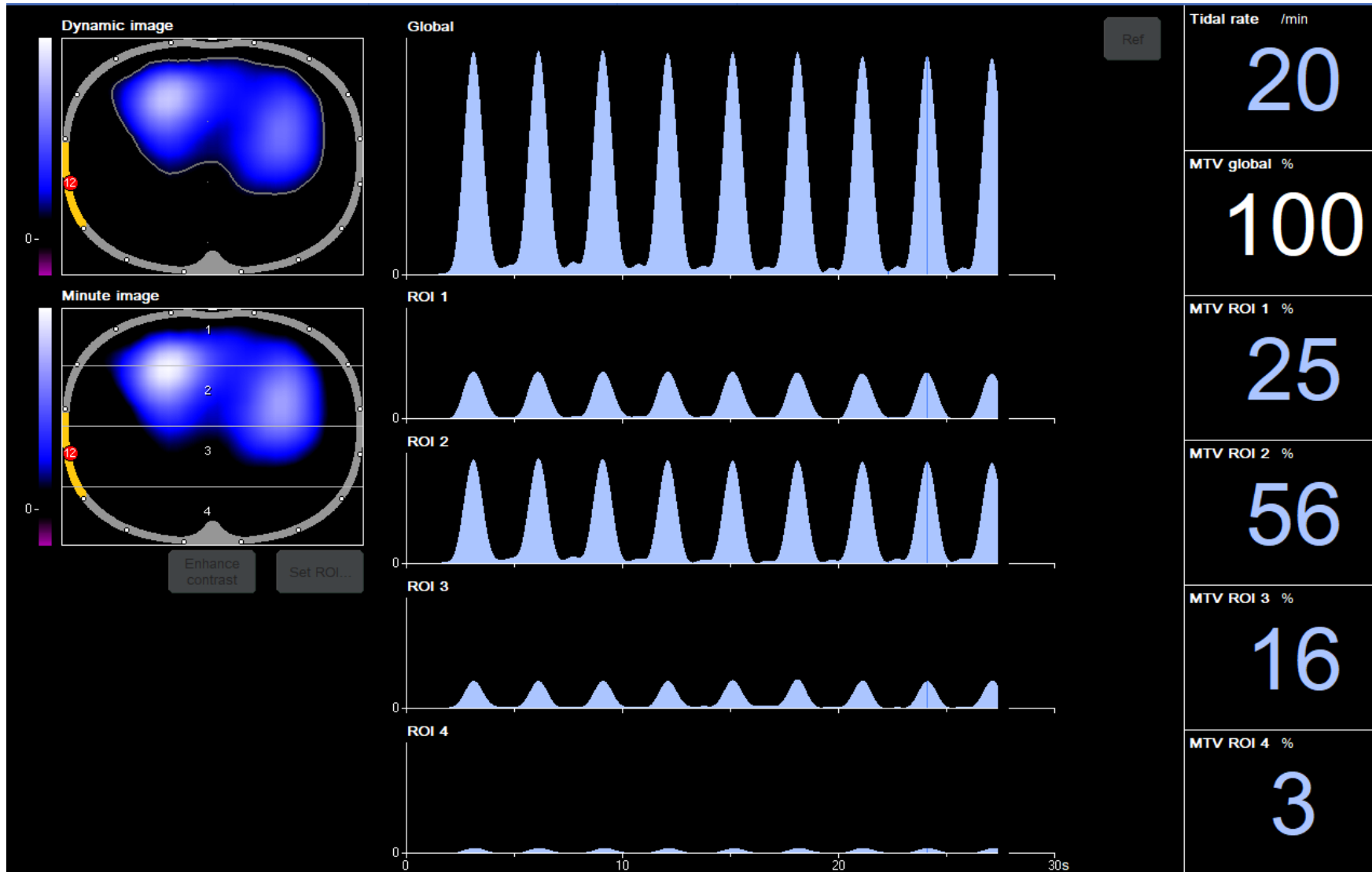
- **Prone position:** Improves dorsal ventilation and compliance.
- **Lateral position:** May promote dorsal recruitment without high pressure.
- Monitor EELI & TIV to detect derecruitment during repositioning.

Step 6. Weaning & Spontaneous Breathing Trial

- Use EIT to track changes in:
 - **Global Inhomogeneity Index (GI)** – rising GI predicts SBT failure.
 - **EELI** – drop indicates derecruitment.
- Early warning of regional instability and failed extubation risk.

GI = 各像素 TIV 與全肺中位數 TIV 的絕對差總和/全肺所有像素 TIV 的總和。
結果反映「**通氣分布的離散程度**」。

Day 1



- Dorsal part atelectasis (+) under PEEP 14.
- RM was approached with 2 steps incremental PEEP until PIP 40 cmH₂O.
- Decremental PEEP trial then started with 2 cmH₂O stepwise until PEEP 12.

PEEP

26

24

22

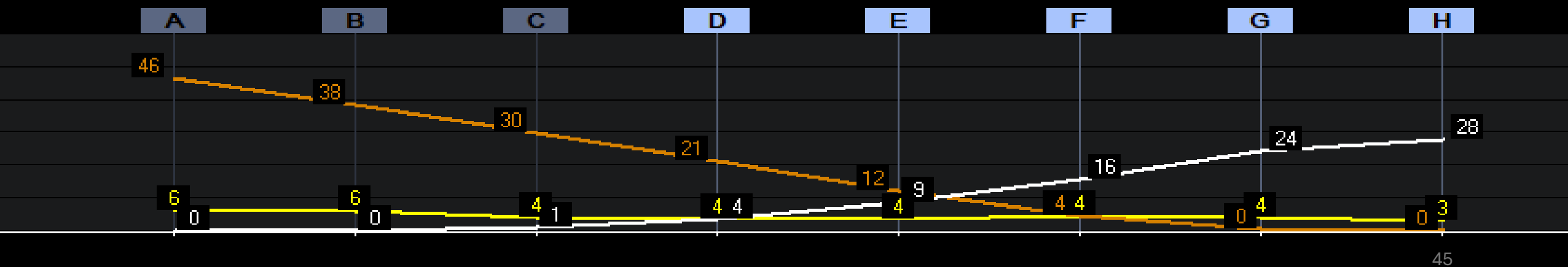
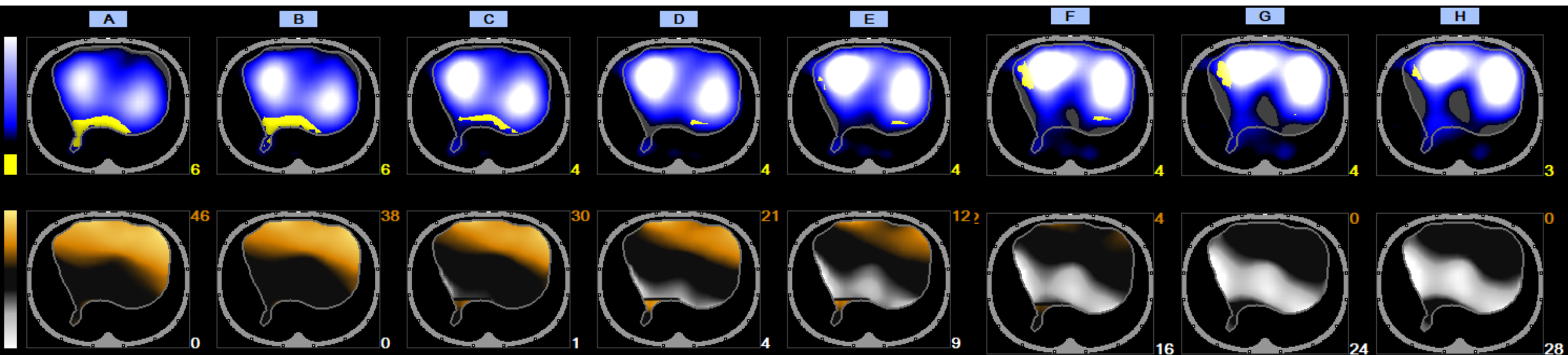
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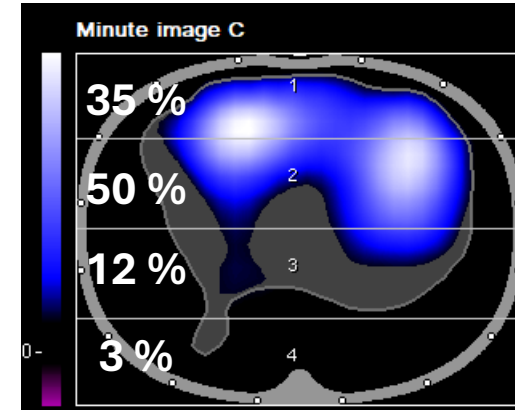
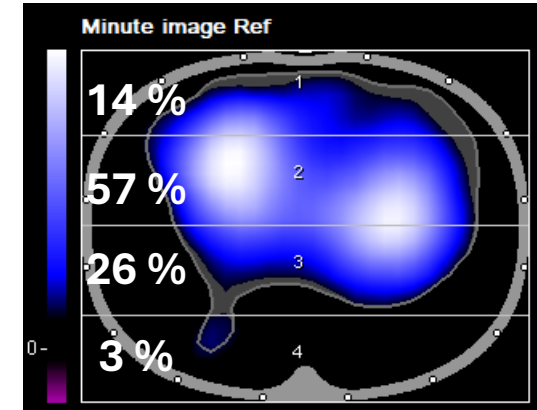
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12

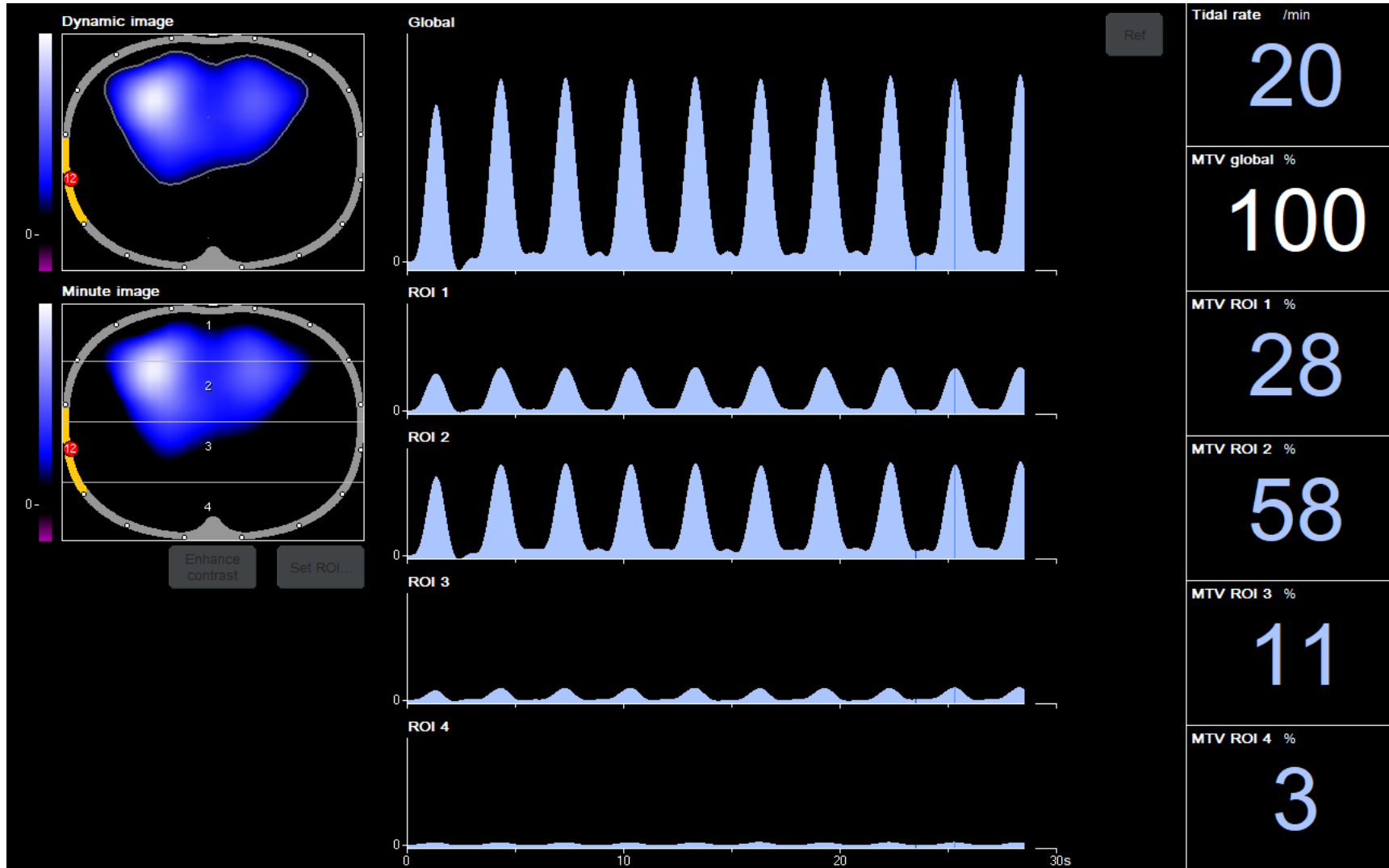


Day 1 – analysis result



- Ventilation of ROI 4 (most dorsal part) remains low even at highest PEEP.
- ROI 3 is recruitable (12 → 26 %) but overdistension is noted over ROI 1 (by decreasing ventilation from 35 to 14%).
- Optimal PEEP (balance overdistension & collapse) is closed to 18 but with ECMO in place, PEEP was set at 14 based on minimal overdistension strategy.
- Interestingly, shearing force could be seen over high PEEP indicates further recruitment possibility by using higher PEEP.

Day 2



- Similar image as compared to yesterday under PEEP 14.
- RM was approached with 2 steps incremental PEEP until PIP 40 cmH₂O.
- Decremental PEEP trial then started with 2 cmH₂O stepwise until PEEP 10.

PEEP

24

22

20

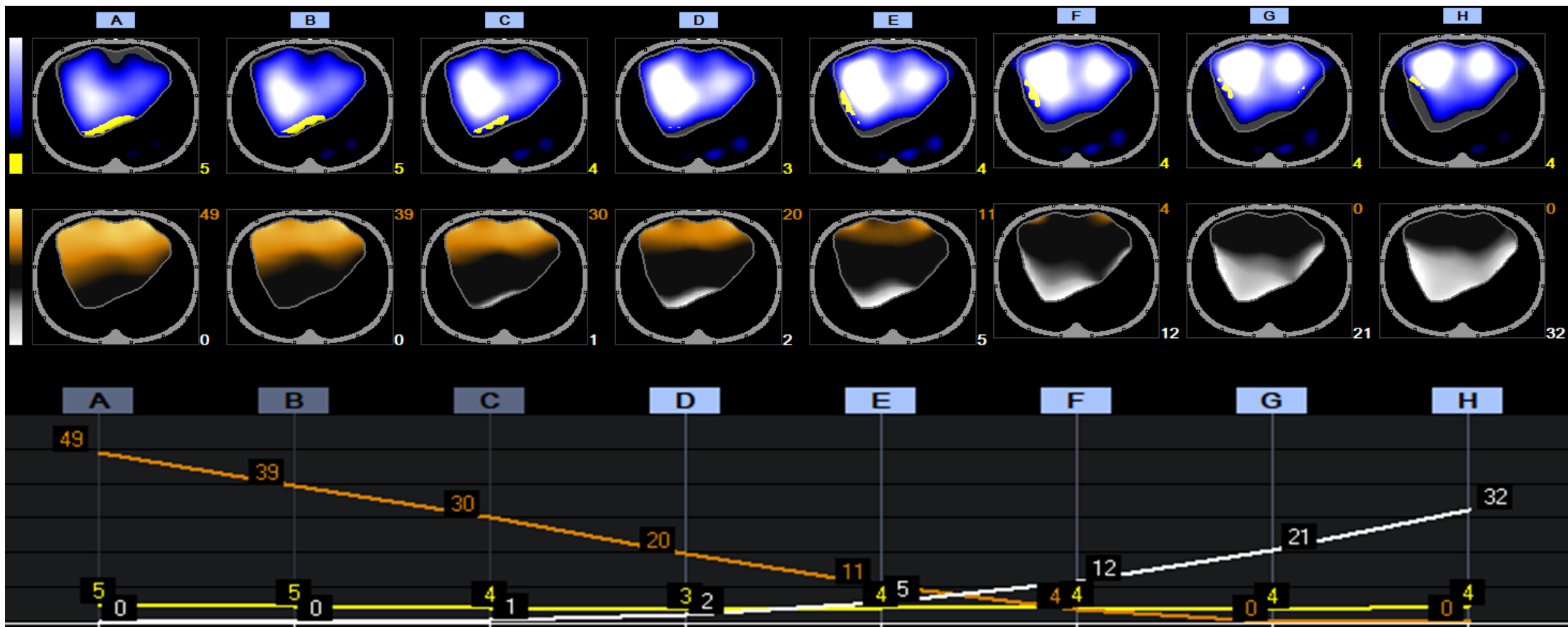
18

16

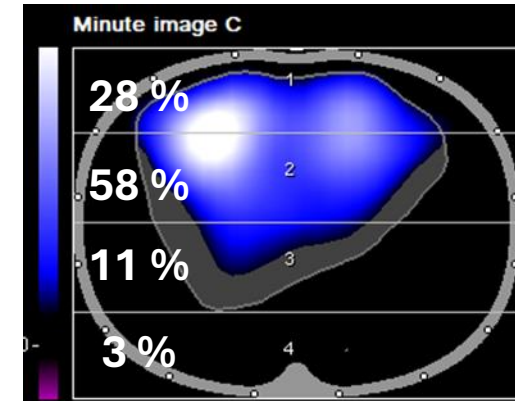
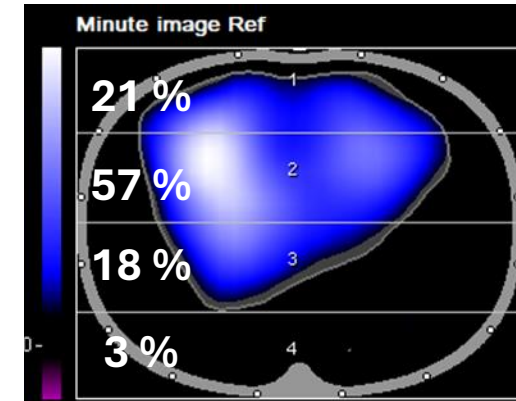
14

12

10



Day 2 – analysis result

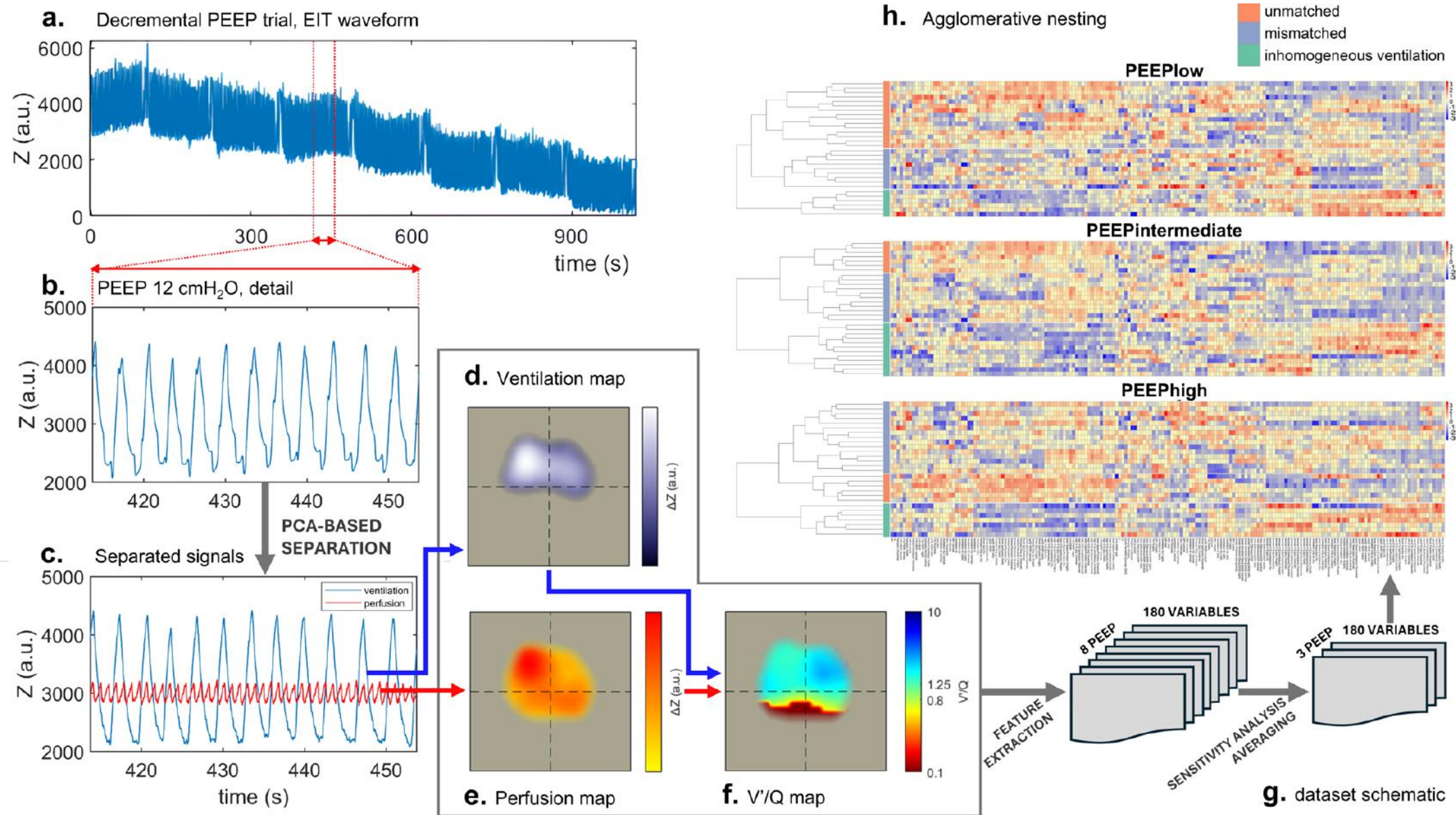


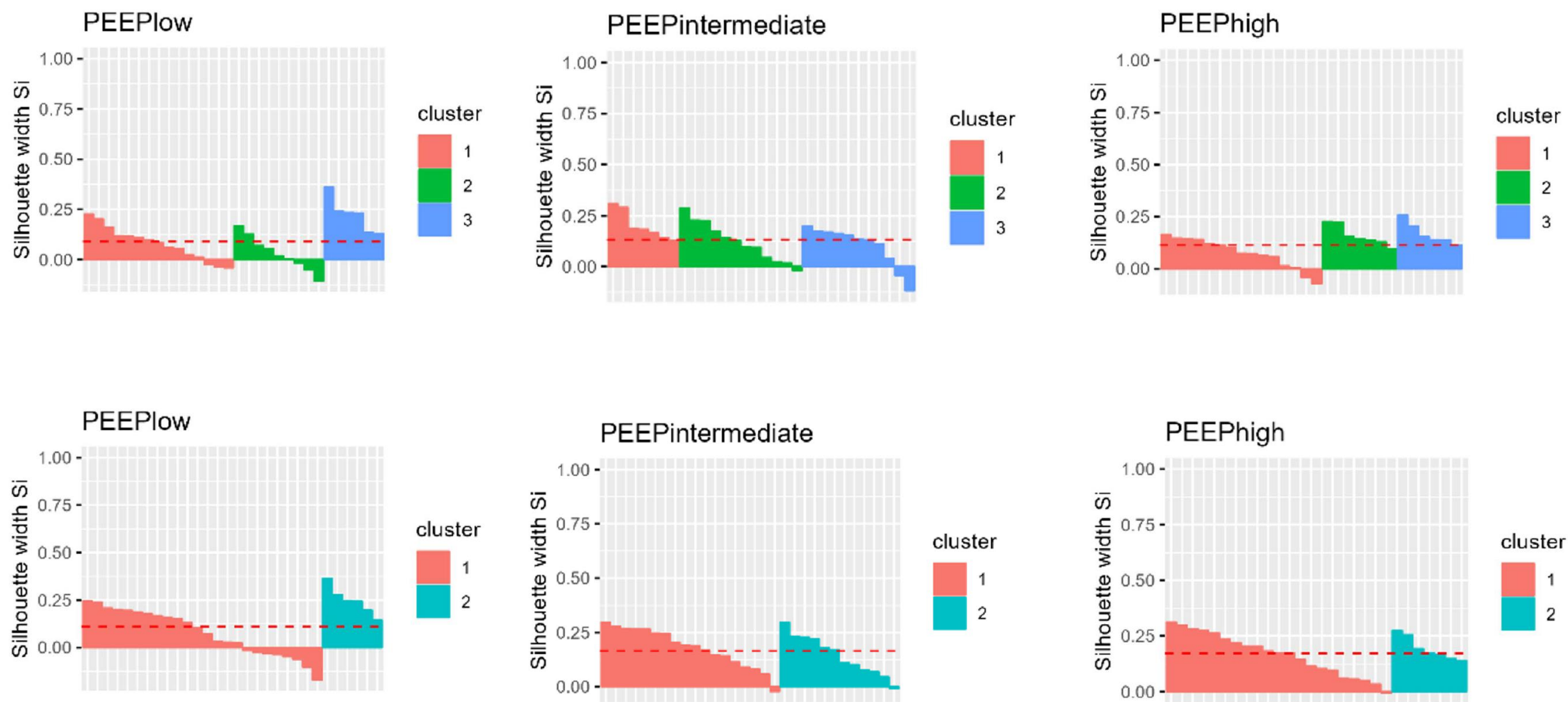
- Same as yesterday, ventilation of ROI 4 (most dorsal part) is not responsive to RM.
- ROI 3 is recruitable (18 → 11 %) but overdistension is noted over ROI 1 (by decreasing ventilation from 28 to 21%). Both minor changes as compared to yesterday.
- Optimal PEEP (balance overdistension & collapse) is reduced to 15 and both 14 or 12 are suitable for minimal overdistension strategy.
- Still, shearing force could be seen over high PEEP same as yesterday.

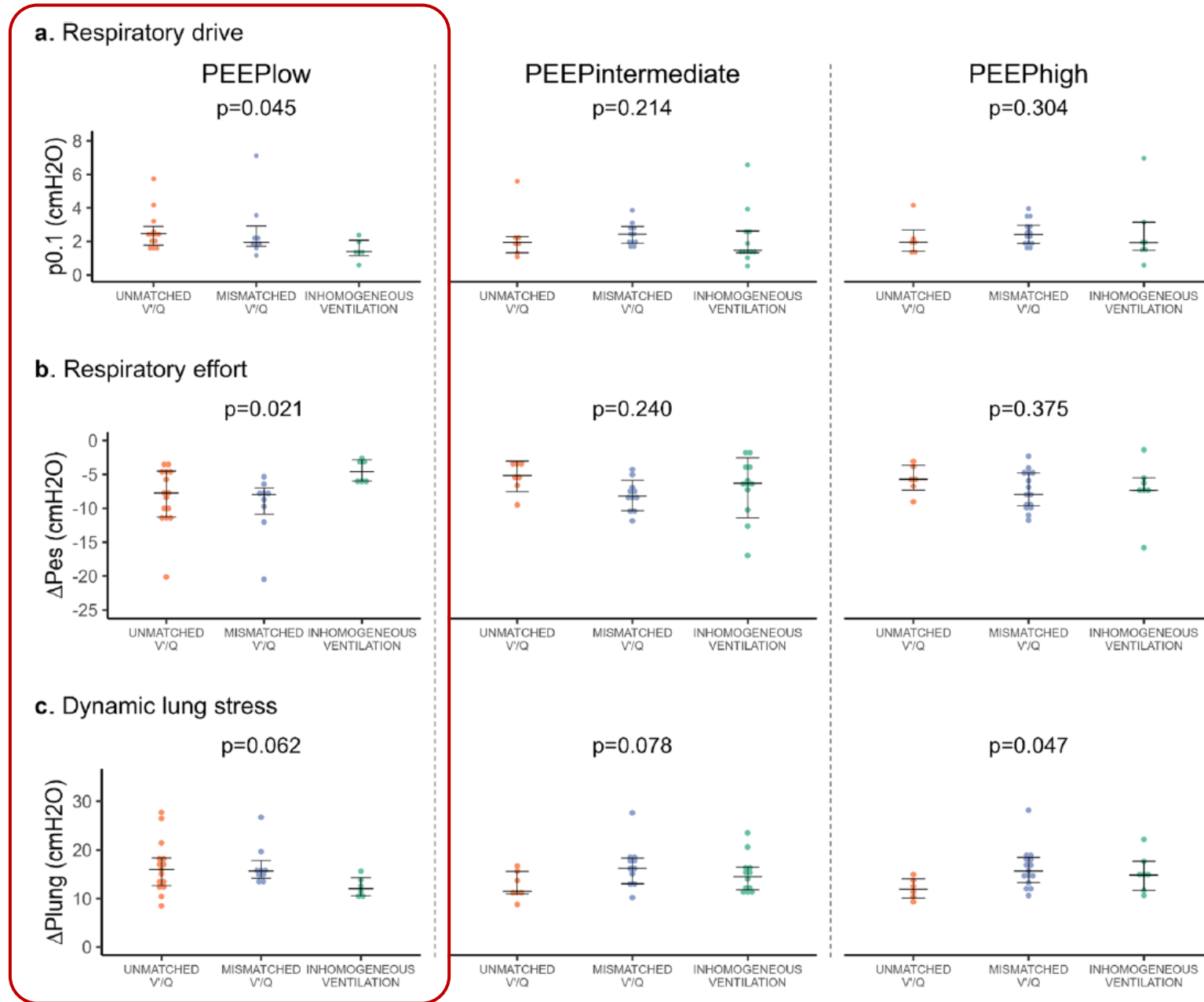
EIT-omics

不是只看 1–2 個 EIT 指標，
而是同時把通氣、灌流
(以脈動訊號作 proxy)、
以及 V' /Q 匹配等 180 個
變數一起分析，再用階層
式分群 (AGNES，
Ward's linkage) 把病人
在不同 PEEP 下的狀態分
出「生理亞型」。目標是
找出呼吸驅動/努力 (P0.1、
ΔPes) 較高、可能較容易
P-SILI 的族群，並觀察
PEEP 調整能否「消弭差異」

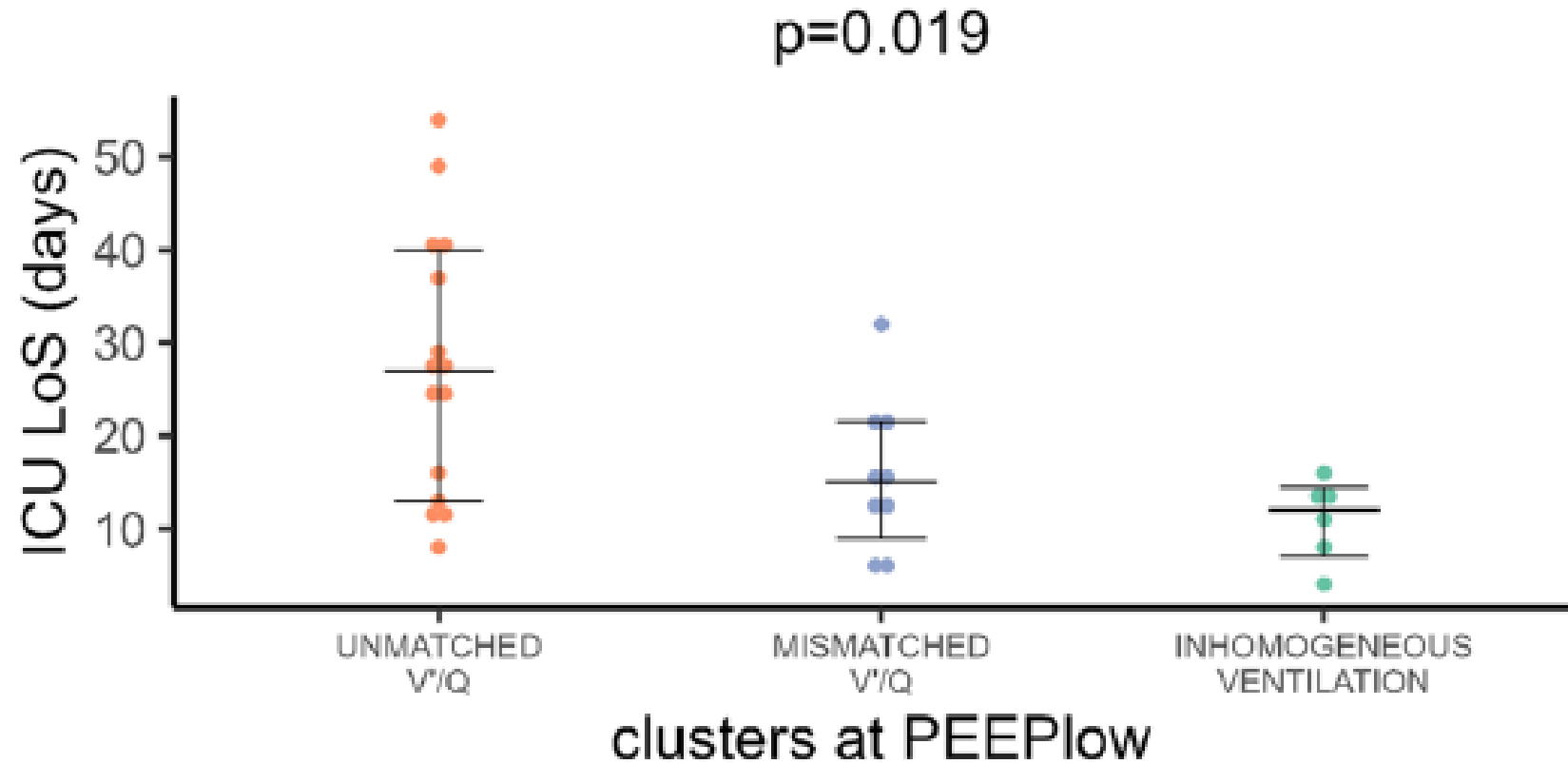
	PEEP _{low}				
	ALL (n = 30)	UNMATCHED (n = 15)	MISMATCHED (n = 9)	INHOMOG. VENTILATION (n = 6)	p-value
age (years)	64 ± 14	59 ± 15	68 ± 12	71 ± 12	0.178
gender (male / female)	23 / 7 (77% / 13%)	12 / 3 (80% / 20%)	7 / 2 (78% / 22%)	4 / 2 (67% / 33%)	0.805
BMI (kg/m ²)	28 ± 5	28 ± 6	27 ± 3	27 ± 5	0.944
P/F (mmHg)	205 ± 55	198 ± 58	202 ± 43	228 ± 66	0.515
PaCO ₂ (mmHg)	45 ± 6	47 ± 5	43 ± 7	43 ± 6	0.239
pH	7.44 ± 0.04	7.43 ± 0.04	7.44 ± 0.06	7.43 ± 0.03	0.914
Ventilatory Ratio	1.69 ± 0.46	1.70 ± 0.49	1.58 ± 0.29	1.81 ± 0.62	0.642
Vt/PBW (ml/kg)	7.88 ± 1.8	7.75 ± 1.46	8.57 ± 2.19	7.19 ± 1.94	0.333
Pulmonary ARDS	22 (73%)	12 (80%)	7 (77.78%)	3 (50%)	0.350
Infectious ARDS	22 (73%)	11 (73.33%)	8 (88.89%)	3 (50%)	0.249
	PEEP _{intermediate}				
	ALL (n = 30)	UNMATCHED (n = 7)	MISMATCHED (n = 11)	INHOMOG. VENTILATION (n = 12)	p-value
age (years)	64 ± 14	63 ± 18	66 ± 8	63 ± 17	0.887
gender (male / female)	23 / 7 (77% / 13%)	5 / 2 (71% / 29%)	10 / 1 (91% / 9%)	8 / 4 (67% / 33%)	0.363
BMI (kg/m ²)	26.64 (24.61–30.86)	26 (25–29)	27 (26–31)	27 (23–31)	0.638
P/F (mmHg)	205 ± 55	227 ± 50	178 ± 47	217 ± 58	0.110
PaCO ₂ (mmHg)	45 ± 6	44 ± 5	46 ± 8	45 ± 5	0.751
pH	7.44 ± 0.04	7.43 ± 0.04	7.46 ± 0.04	7.41 ± 0.04	0.022
Ventilatory Ratio	1.61 (1.38–1.87)	1.48 (1.32–1.65)	2.00 (1.73–2.26)	1.46 (1.27–1.65)	0.017
Vt/PBW (ml/kg)	7.57 (6.66–8.54)	7.72 (6.8–8.07)	7.23 (6.74–8.3)	8.04 (6.52–9.9)	0.786
Pulmonary ARDS	22 (73%)	5 (71.43%)	10 (90.91%)	7 (58.33%)	0.209
Infectious ARDS	22 (73%)	5 (71.43%)	7 (63.64%)	10 (83.33%)	0.561
	PEEP _{high}				
	ALL (n = 28)	UNMATCHED (n = 6)	MISMATCHED (n = 15)	INHOMOG. VENTILATION (n = 7)	p-value
age (years)	64 ± 14	62 ± 19	64 ± 14	61 ± 11	0.910
gender (male / female)	23 / 7 (77% / 13%)	5 / 1 (83% / 17%)	13 / 2 (87% / 13%)	5 / 2 (71% / 29%)	0.683
BMI (kg/m ²)	26.64 (25.04–30.86)	26 (25–27)	26 (25–30)	29 (27–33)	0.411
P/F (mmHg)	205 ± 55	233 ± 51	199 ± 57	204 ± 58	0.473
PaCO ₂ (mmHg)	45 ± 6	44 ± 6	46 ± 7	43 ± 4	0.534
pH	7.44 ± 0.05	7.45 ± 0.04	7.45 ± 0.04	7.39 ± 0.05	0.015
Ventilatory Ratio	1.61 (1.38–1.93)	1.53 (1.40–1.69)	1.76 (1.47–2.17)	1.43 (1.30–1.65)	0.434
Vt/PBW (ml/kg)	7.57 (6.66–8.56)	7.78 (7.32–8.19)	6.87 (6.55–8.3)	8.51 (7.57–9.94)	0.251
Pulmonary ARDS	21 (75%)	3 (50%)	14 (93.33%)	4 (57.14%)	0.053
Infectious ARDS	21 (75%)	5 (83.33%)	11 (73.33%)	5 (71.43%)	0.864







e. Clinical outcomes



Each EIT variable was tested for differences between clusters by bonferroni-corrected ANOVA/Kruskal-Wallis tests, significant features at PEEP_{low} are here represented with their p-values and intra-class correlation coefficients (ICC)

f. Significant features

variable	unmatched	mismatched	inhomogeneous ventilation	Bonferroni-corrected p-value	ICC
Dead Space Fraction (%)	85 ± 7	61 ± 12	57 ± 8	<0.001	80
High V'/Q [ventilation] (%)	72 ± 16	55 ± 14	31 ± 7	0.003	71
Normal V'/Q [perfusion] (%)	16 ± 7	34 ± 7	31 ± 7	0.001	75
Wasted Perfusion (%dorsal)	70 ± 15	38 ± 12	34 ± 19	0.005	70

Take home message

- EIT is an emerging **functional lung imaging tool** with great potential.
- EIT Provides **noninvasive, continuous bedside monitoring** of lung ventilation and perfusion dynamics.
- The main challenges lie in **perfusion measurement and clinical validation**.
- Future progress depends on **integrating ventilation–perfusion analysis, improving bedside usability, and generating solid clinical evidence**.

Thanks for your attention