# ADULT RESPIRATORY DISTRESS SYNDROME

A form of non-cardiogenic pulmonary oedema that is a result of the non-specific response of the lung to a variety of insults.

# **DEFINITION.**

- Respiratory failure requiring mechanical ventilation
- PaO2/FiO2 ratio ≤ 200
- New, bilateral [diffuse, patchy or homogenous] pulmonary infiltrates on CXR.
  - No clinical evidence of heart failure/fluid overload/chronic lung disease.
- One or more recognised risk factors.

BOX 74-1	CONDITIONS ASSOCIATED WITH ACUTE RESPIRATORY DISTRESS SYNDROME									
Sepsis										
Shock	en en else interletten									
Ioxic gas	or smoke innalation									
Gastric	contents									
Near-d	Near-drowning									
Hydrod	carbons/solvents									
Pneumor	iia									
Drug read	ction									
Salicyla	ates									
Opiate	S									
Iricycli	c antidepressants									
Amiod	Johne									
Cancer	chemotherapeutic agents (e.g., bleomycin)									
Hvdroo	chlorothiazide									
Trauma										
Burns										
Transfusio	on reaction									
Radiation	injury									
Pancreati	tis									
Ihrombo	embolism									
Air embo	lism									
Amniotic	fluid embolism									
Eclampsia	a									
Neuroger	nic (e.g., subarachnoid hemorrhage, head trauma)									
Dissemin	ated intravascular coagulation									
High-altit	ude exposure									
Oxygen t	oxicity									
Cardiopu	Imonary bypass									

**TREATMENT.** Largely supportive.

## LUNG-PROTECTIVE VENTILATOR STRATEGY.

- Tidal volume:
  - 6mL/kg of ideal body weight
  - Initial RR set to target minute ventilation.
- Oxygenation goal:
  - PaO2 55-80mmHg OR SpO2 88-95%.
- Plateau pressure goal:
  - ≤ 30 cmH2O
- pH goal:
  - 7.30-7.45



NIH NHLBI ARDS Clinical Network Mechanical Ventilation Protocol Summary

#### **INCLUSION CRITERIA: Acute onset of**

- $PaO_2/FiO_2 \le 300$  (corrected for altitude)
- Bilateral (patchy, diffuse, or homogeneous) infiltrates consistent with 2. pulmonary edema
- 3. No clinical evidence of left atrial hypertension

#### PART I: VENTILATOR SETUP AND ADJUSTMENT

- Calculate predicted body weight (PBW) 1. Males = 50 + 2.3 [height (inches) - 60]
- Females = 45.5 + 2.3 [height (inches) -60]
- Select any ventilator mode 2
- Set ventilator settings to achieve initial  $V_T = 8 \text{ ml/kg PBW}$ 3.
- Reduce  $V_{T}$  by 1 ml/kg at intervals  $\leq$  2 hours until  $V_{T}$  = 6ml/kg PBW. 5. Set initial rate to approximate baseline minute ventilation (not > 35
- bpm).
- Adjust  $V_T$  and RR to achieve pH and plateau pressure goals below. 6.

## pH GOAL: 7.30-7.45

#### Acidosis Management: (pH < 7.30) If pH 7.15-7.30: Increase RR until pH > 7.30 or PaCO<sub>2</sub> < 25 (Maximum set RR = 35).

If pH < 7.15: Increase RR to 35.

If pH remains < 7.15,  $V_T$  may be increased in 1 ml/kg steps until pH > 7.15 (Pplat target of 30 may be exceeded). May give NaHCO

Alkalosis Management: (pH > 7.45) Decrease vent rate if possible.

I: E RATIO GOAL: Recommend that duration of inspiration be < duration of expiration.

#### PART II: WEANING

- A. Conduct a SPONTANEOUS BREATHING TRIAL daily when: 1.  $FiO_2 \le 0.40$  and  $PEEP \le 8$ .
  - 2. PEEP and  $FiO_2 \leq$  values of previous day.
  - Patient has acceptable spontaneous breathing efforts. (May 3. decrease vent rate by 50% for 5 minutes to detect effort.)
  - 4. Systolic BP  $\geq$  90 mmHg without vasopressor support.
  - 5. No neuromuscular blocking agents or blockade.

#### OXYGENATION GOAL: PaO<sub>2</sub> 55-80 mmHg or SpO<sub>2</sub> 88-95% Use a minimum PEEP of 5 cm H<sub>2</sub>O. Consider use of incremental FiO<sub>2</sub>/PEEP combinations such as shown below (not required) to achieve goal.

#### Lower PEEP/higher FiO2

FiO <sub>2</sub>	0.3 0.4		0.4	0.5	0.5	0.6	0.7	0.7			
PEEP	5	58		8 10		10	10	12			
	0 7	~ ~	~ ~	~ ~	~ ~	1 0					

FIU <sub>2</sub>	0.7	0.0	0.9	0.9	0.9	1.0
PEEP	14 14		14	16	18	18-24

#### **Higher PEEP/lower FiO2**

FiO <sub>2</sub>	0.3	0.3	0.3		0.3		0.3		).4	0.4	0.5
PEEP	5	8	10		12		14		14		16
FiO <sub>2</sub>	0.5	0.5-0.8		0	0.8 (		0.9		0	1.0	
PEEP	18	20	0		2	22		22		24	

#### PLATEAU PRESSURE GOAL: ≤ 30 cm H<sub>2</sub>O

Check Pplat (0.5 second inspiratory pause), at least q 4h and after each change in PEEP or V<sub>T</sub>

If Pplat > 30 cm H<sub>2</sub>O: decrease  $V_T$  by 1ml/kg steps (minimum = 4 ml/ka).

If Pplat < 25 cm H<sub>2</sub>O and V<sub>T</sub>< 6 ml/kg, increase V<sub>T</sub> by 1 ml/kg until Pplat > 25 cm  $H_2O$  or  $V_T = 6$  ml/kg.

If Pplat < 30 and breath stacking or dys-synchrony occurs: may increase V<sub>T</sub> in 1ml/kg increments to 7 or 8 ml/kg if Pplat remains  $\leq$  30 cm H<sub>2</sub>O.

#### B. SPONTANEOUS BREATHING TRIAL (SBT):

If all above criteria are met and subject has been in the study for at least 12 hours, initiate a trial of UP TO 120 minutes of spontaneous breathing with FiO2 < 0.5 and PEEP < 5:

1. Place on T-piece, trach collar, or CPAP  $\leq$  5 cm H<sub>2</sub>O with PS  $\leq$  5

- 2. Assess for tolerance as below for up to two hours.
  - $SpO_2 \ge 90$ : and/or  $PaO_2 \ge 60$  mmHg a. b.
    - Spontaneous  $V_T \ge 4 \text{ ml/kg PBW}$  $\dot{RR} \le 35/min$
  - c. . pH ≥ 7.3 d.
  - e.
    - No respiratory distress (distress= 2 or more) HR > 120% of baseline
      - Marked accessory muscle use
    - Abdominal paradox
    - Diaphoresis
    - Marked dyspnea
- 3. If tolerated for at least 30 minutes, consider extubation.
- 4. If not tolerated resume pre-weaning settings.

## Definition of UNASSISTED BREATHING (Different from the spontaneous breathing criteria as PS is not allowed)

- Extubated with face mask, nasal prong oxygen, or 1. room air, OR
- 2 T-tube breathing, OR
- 3. Tracheostomy mask breathing, OR
- CPAP less than or equal to 5 cm H<sub>2</sub>0 without 4. pressure support or IMV assistance.