

TSICU/BICU Routine Ventilator Management Protocol

INITIATE VENTILATOR SET UP:

- On arrival place patient on APV/SIMV and initiate Pulse Oximetry with the following settings:
- Initial Tidal Volume is 7milliliters per kilogram of Ideal Body Weight (IBW).
- Initial Ventilator Rate 14
- FiO₂ 100% initially, Weaning FiO₂ to 40% in 10-15 minute increments, while monitoring the Pulse Oximeter. Auto-wean FiO₂ to maintain SPO₂ greater than or equal to 88%.
- PEEP of 5cmH₂O *May increase PEEP if the patient cannot be weaned to FiO2 goal and has a systolic blood pressure that is greater than 80mmHg. Follow the FiO2/ PEEP scale*
- Set initial PS to 10cmH₂O
- I:E 1:1.0-1:3.0

ARTERIAL OXYGENATION GOAL: Pa02 55 – 80 mm Hg or Sp02 >88%

Fi02	0.3	0.4	0.4	0.5	0.5	0.6	0.7	0.7	0.7	0.8	0.9	0.9	0.9	1.0
PEEP	5	5	8	8	10	10	10	12	14	14	14	16	18	18-24
High PEEP/Low FiO2 Scale														
Fi02	0.3	0.3	0.3	0.3	0.3	0.4	0.4	0.5	0.5	0.5-0.8	0.8	0.9	1.0	1.0
PEEP	5	8	10	12	14	14	16	16	18	20	22	22	22	24

Low PEEP/HIGH FiO2 Scale

ABG will be drawn one hour after mechanical ventilation has been initiated

A.Respiratory Acidosis Management:

<u>lf pH 7.15 – 7.30:</u>

Increase set RR until pH greater than 7.30 or $PaCO_2$ less than 25 (**Maximum** Set RR = 30)

<u>If pH less than 7.15:</u> Increase set RR (maximum 30).

If set RR has been increased to 30 and pH 7.15; tidal volume may be increased in1 ml/ kg increments (not to exceed 8ml/kg IBW) until pH greater than 7.15

B.Respiratory Alkalosis Management: (pH greater than 7.45): Decrease set RR (minimum set rate of 6) Decrease tidal volume by 1ml/kg IBW (no lower than 5 ml/kg)

ABG will be drawn 30 minutes after any Minute Ventilation change. *If the ABG has critical values, notify the ordering physician*

1. MINIMUM QShift SBTs will be performed between the hours of 0900-1000 and 2000-2100. For possible liberation from the vent (<u>All</u> of the following conditions must be met)

- a. Fi 0_2 less than or equal to 40%
- b. PEEP less than or equal to 5 cmH₂0)
- c. Intact airway reflexes (Cough or gag)
- d. Hemodynamic stability
- e. Adequate level of consciousness
- 2. Trial of spontaneous breathing: Reference SAT/SBT chart (see below) to make sure all parameters are met.

PERFORM CUFF LEAK TEST PRIOR TO PLACING PATIENT ON A SBT. If the patient is negative for a cuff leak, notify the ordering physician and document accordingly.

CPAP 5 with PS of 5-10 cmH₂0 for **20-30** minutes

*If the patient is on ASV, lower the %MV to 40%. If the patient is not triggering spontaneous breaths, lower the %MV to 25% (Do not lower beyond 25% MV. Closely monitor patient). IF the patient is spontaneously breathing and the PS that is being delivered via the lowered %MV is less than or equal to 10 cmH2O, the patient is considered to be performing an SBT.

3. Trial will be stopped if <u>ANY</u> of the following conditions are observed:

- a. RR > 35 breath/minute for greater than **5** minutes
- b. Sa0₂ < 90%
- c. Sustained increase or decrease in HR by greater than or equal to 20% of baseline.
- d. Systolic BP > 180 mmHg or < 90 mmHg
- e. If the patient is performing a SBT on ASV, discontinue if the patient is not meeting the target MV goal with the lowered %MV. This is indicative of apnea or inability to generate large enough VTs to maintain adequate MV.
- If any of the above conditions are observed the trial is considered unsuccessful, place patient on previous ventilator settings. *Notify ordering physician and document accordingly.
- If any of the above conditions are not observed, the trial is considered successful; obtain weaning parameters. *Notify ordering physician and document accordingly.

Weaning Parameters:

- NIF better than -20cmH₂O
- Spontaneous tidal volume > 5mL/kg IBW
- Spontaneous respiratory rate < 25 breaths per minute
- Spontaneous minute volume < 10 L per minute
- Vital capacity >10mL/kg IBW

- RSBI less than or equal to 90 (Respiratory Rate / Tidal Volume)
- 4. Physician will be notified after the spontaneous breathing trial and a decision for extubation will be discussed.
- **NOTE:** If the trial is stopped, the patient will be reassessed for weaning tolerance at the next scheduled SBT time. A SBT may be performed at any time in ADDITION to the scheduled trials if the physician deems the patient ready.