Acute Ventilator Sedation Management Protocol

Inclusion Criteria:

- PICU patients; ED patients
- Acute respiratory failure requiring mechanical ventilation
- Exclusion Criteria:
- CVICU or NICU patients and/or chronic/ventilator dependent patients
- Neuromuscular respiratory failure

Assessment

- Monitor and document sedation assessment with the State Behavioral Sedation (SBS) tool a minimum of every 2 hours
- Discuss and identify the patient's trajectory of illness during multidisciplinary rounds every day
- Acute phase: Goal SBS -1 or -2: Patient critical and/or unstable, interventions escalating
- Titration phase: Goal SBS -1 or 0: Patient stable, interventions are not escalating
- Extubation phase: Goal SBS 0: Passed ERT and plan to attempt extubation
- Assess for extubation readiness daily using the Extubation Readiness Test (ERT)
- · Perform arousal and modified arousal assessments for patients in titration phase as indicated

Treatment

➢ If length of intubation/ventilation anticipated to be ≤ 2 days, begin: PRN doses of Benzodiazepines + narcotics:

- Midazolam 0.05 mg/kg IV Q 2 hours PRN agitation (max start dose 2 mg) OR
- Lorazepam 0.05 mg/kg NGT Q 4 hours PRN agitation (max dose 2mg)
- Fentanyl 1 mCg/kg IV Q 2 hours PRN pain (max start 50mCg) OR
- Morphine 0.1 mg/kg IV Q 2 hours PRN pain (max start 2 mg)

AND/OR Dexmedetomidine Infusion Start 0.2 – 0.5 mCg/kg/hr. Titrate to SBS goal.

OR Consider Propofol Infusion 25mCg/kg/min. Titrate to SBS goal.

- If length of intubation/ventilation anticipated to be >2 days, start continuous infusions with PRN doses equal to one hour of infusion dose: Dexmedetomidine +/- narcotics:
- Dexmedetomidine Infusion Start 0.2 0.5 mCg/kg/hr
- Fentanyl 0.5 mCg/kg/hr IV (max start 50mCg) OR
- Morphine 0.05 mg/kg/hr IV (max start 2 mg)
- > If additional sedation required, add Benzodiazepines:
- Midazolam 0.05 mg/kg/hr IV (max start dose 2 mg)

Titrate sedation according to the patient's trajectory/phase of illness to maintain goal SBS.

Continued Considerations

If extubation is planned within 24 hours and the patient is intolerant of an SBS of -1 or 0 then consider stopping versed and starting propofol 25 mCg/kg/min titrated to achieve desired SBS score, up to maximum of 100 mCg/kg/min. Discontinue propofol and extubate when SBS is 0.

Fentanyl should be used (over morphine) in patients with profound hypotension or unremitting reactive airway disease

In patients receiving neuromuscular blockade (these patients should all be managed in the acute phase) use: Assume Pain Present or Assume Agitation Present assessments

Monitor for delirium utilizing CAP-D screening tool, twice daily. If suspected, consider limiting the use of benzodiazepines (utilize dexmedetomidine as substitute) and/or consider the addition of an atypical antipsychotic



СНОС

Acute Phase (SBS -1, or -2) Goal is to maintain physiologic stability.

If patient's SBS is more positive than prescribed, exclude reversible causes and provide comfort measures. If ineffective, titrate dexmedetomidine. If ineffective, administer a fentanyl/morphine and/or versed PRN dose. If a total of 3 nonprocedural PRN doses are administered in ≤ 8 hours then increase fentanyl/morphine and/or versed infusion by 10-20%. Adjust PRN dose to equal one hour of infusion.

Titration Phase (SBS -1 or 0)

Goal is minimum yet effective dosing.

If SBS -3, complete an arousal assessment Turn off all sedation until SBS -1 or 0 then reduce sedation doses by 50%, restart and titrate

If SBS -2, complete a modified arousal assess. Reduce sedative doses by 50% then titrate

Every 8 hours adjust sedation. As in the acute phase, if patient's SBS is more positive than prescribed, exclude reversible causes and provide comfort measures. If ineffective, titrate dexmedetomidine. If ineffective, administer a fentanyl/morphine and/or versed PRN dose. If a total of 3 nonprocedural PRN doses are administered in ≤ 8 hours then increase fentanyl/morphine and/or versed infusion by 10-20%. Adjust PRN dose to equal one hour of infusion.

If the patient's SBS is more negative or as prescribed and if < 3 total nonprocedural PRN doses are given in an 8 hour period then decrease fentanyl/morphine and/or versed by 10-20%. Adjust PRN dose to equal one hour of infusion.

Extubation Phase (SBS 0)

Goal is to D/C sedation and extubate.

If patient on sedation for < 5 days then discontinue medications *OR*

If on sedation for ≥ 5 days identify target Withdrawal Assessment Tool (WAT-1) and begin methadone/ ativan wean

Methadone

0.1mg/kg IV q6h

5 mg IV q6h, Dosing Guidelines: > 50 kg; 0.1 mg/kg NGT q6h

5 mg NGT q6h, Dosing Guidelines: > 50 kg; Wean by 20% of initial dose Q other day, alternate w/ lorazepam wean day Morphine 0.1 mg/kg IV Q 4 hours, PRN WAT-1 >

identified target

Lorazepam starting dose = (total daily midazolam dose)/10 daily divided every 6 hours Wean by 20% of initial dose Q other day, alternate w/methadone wean day

Patient Education

- Review pain and sedation assessment and trajectory of illness with parents/care givers
- Discuss both nonpharmacologic and pharmacologic comfort measures to address pain and agitation



Approved Evidence Based Medicine Committee 1/15/14 Revised - 8/6/14; 12/23/14; 4/20/22 Reviewed - 11/15/17

Reassess the appropriateness of Care Guidelines as condition changes and 24 hrs after admission. This guideline is a tool to aid clinical decision making. It is not a standard of care. The physician should deviate from the guideline when clinical judgment so indicates.

Daily Extubation Readiness Test (ERT)





State Behavioral Scale (SBS) for Sedation Assessment

-3	Comatose and unresponsive
-2	Responsive to noxious stimulus
-1	Responsive to gentle touch or voice
0	Awake and able to calm
+1	Restless and difficult to calm
+2	Extremely agitated

Patient Identifier]						
	Date:							
	Time:							
Information from patient record	d, previous 12 hours							
Any loose /watery stools	No = 0 Yes = 1							
Any vomiting/wretching/gagging	No = 0 Yes = 1							
Temperature > 37.8°C	No = 0 Yes = 1							
2 minute pre-stimulus observa	tion							
State	$SBS^1 \le 0$ or asleep/awake/calm = 0 $SBS^1 \ge +1$ or awake/distressed = 1							
Tremor	None/mild = 0 Moderate/severe = 1							
Any sweating	No = 0 Yes = 1							
Uncoordinated/repetitive moveme	nt None/mild = 0 Moderate/severe = 1							
Yawning or sneezing	None or 1 = 0 >2 = 1							
1 minute stimulus observation								
Startle to touch	None/mild = 0 Moderate/severe = 1							
Muscle tone	Normal = 0 Increased = 1							
Post-stimulus recovery			94 G	 	 			
Time to gain calm state (SBS ¹ \leq 0)	< 2min = 0 2 - 5min = 1 > 5 min = 2							
Total Score (0-12)								

WITHDRAWAL ASSESSMENT TOOL VERSION 1 (WAT – 1) © 2007 L.S. Franck and M.A.Q. Curley. All Rights reserved. Reproduced only by permission of Authors.



References Acute Ventilator Sedation Management Care Guideline

- Anand, K. J., Willson, D. F., Berger, J., Harrison, R., Meert, K. L., Zimmerman, J., . . . Nicholson, C. (2010). Tolerance and withdrawal from prolonged opioid use in critically ill children. *Pediatrics*, 125(5), e1208-1225. https://doi:10.1542/peds.2009-0489 (Level II)
- Curley, M. A., Harris, S. K., Fraser, K. A., Johnson, R. A., & Arnold, J. H. (2006). State Behavioral Scale: A sedation assessment instrument for infants and young children supported on mechanical ventilation. *Pediatric Critical Care Medicine*, 7(2), 107-114. https://doi: 10.1097/01.PCC.0000200955.40962.38 (Level II)
- Curley, M. A., Wypij, D., Watson, R. S., Grant, M. J., Asaro, L. A., Cheifetz, I. M., ... Matthay, M. A. (2015). Protocolized sedation versus usual care in pediatric patients mechanically ventilated for acute respiratory failure: A randomized clinical trial. *The Journal of the American Medical Association*, 313(4), 379-389. https:// doi:10.1001/jama.2014.18399 (Level I)
- Deeter, K. H., King, M. A., Ridling, D., Irby, G. L., Lynn, A. M., & Zimmerman, J. J. (2011). Successful implementation of a pediatric sedation protocol for mechanically ventilated patients. *Critical Care Medicine*, 39(4), 683-688. https://doi:10.1097/CCM.0b013e318206cebf (Level III)
- Franck, L. S., Harris, S. K., Soetenga, D. J., Amling, J. K., & Curley, M. A. (2008). The Withdrawal Assessment Tool-1 (WAT-1): An assessment instrument for monitoring opioid and benzodiazepine withdrawal symptoms in pediatric patients. *Pediatric Critical Care Medicine*, 9(6), 573-580. https://doi:10.1097/ PCC.0b013e31818c8328 (Level II)
- Grant, M. J., Schneider, J. B., Asaro, L. A., Dodson, B. L., Hall, B. A., Simone, S. L., . . . Curley, M. A. (2016). Dexmedetomidine use in critically ill children with acute respiratory failure. *Pediatric Critical Care Medicine*, *17*(12), 1131-1141. https://doi: 10.1097/PCC.00000000000941 (Level I)
- Hayden, J. C., Doherty, D. R., Dawkins, I., Leacy, F. P., Healy, M., Breatnach, C. V., . . . Gallagher, P. J. (2019). The effectiveness of α2 agonists as sedatives in pediatric critical care: a propensity score-matched cohort study. *Critical Care Medicine*, *47*(7), e580-e586. https://doi:10.1097/CCM.00000000003789 (Level IV)
- Lebet, R., Hayakawa, J., Chamblee, T. B., Tala, J. A., Singh, N., Wypij, D., & Curley, M. A. (2017). Maintaining interrater agreement of core assessment instruments in a multisite randomized controlled clinical trial: the randomized evaluation of sedation titration for respiratory failure (RESTORE) trial. *Nursing Research*, *66*(4), 323-329. https://doi: 10.1097/NNR.0000000000224 (Level V)
- Sperotto, F., Mondardini, M. C., Dell'Oste, C., Vitale, F., Ferrario, S., Lapi, M., . . . Amigoni, A. (2020). Efficacy and safety of dexmedetomidine for prolonged sedation in the PICU: A prospective multicenter study (PROSDEX). *Pediatric Critical Care Medicine*, 21(7), 625-636. https://doi:10.1097/PCC.00000000002350 (Level II)
- Thomas, N. J., Shaffer, M. L., Willson, D. F., Shih, M. C., & Curley, M. A. (2010). Defining acute lung disease in children with the oxygenation saturation index. *Pediatric Critical Care Medicine*, 11(1), 12-17. https:// doi:10.1097/PCC.0b013e3181b0653d (Level II)
- Walker, T., & Kudchadkar, S. R. (2019). Pain and Sedation Management: 2018 Update for the Rogers' Textbook of Pediatric Intensive Care. *Pediatric Critical Care Medicine*, 20(1), 54-61. https://doi.org/10.1097/ PCC.000000000001765 (Level II)
- Watson, R. S., Asaro, L. A., Hertzog, J. H., Sorce, L. R., Kachmar, A. G., Dervan, L. A., . . . Curley, M. A. (2018). Long-term outcomes after protocolized sedation versus usual care in ventilated pediatric patients. *American Journal of Respiratory and Critical Care Medicine*, 197(11), 1457-1467. https://doi.org/10.1164/rccm.201708-1768OC (Level IV)

Reviewed by Evidence-Based Medicine Committee 11/15/17; Updated 4/20/22; 11/15/2017

