

ICU Pain/Agitation/Delirium Reference

Goal of Sedation/Analgesia

- 1. Pain level <3 using Visual Analog Scale or Cognitively Impaired Scale
- 2. No delirium according to the Modified Confusion Assessment Method for the ICU (CAM ICU)
- 3. Richmond Agitation Sedation Scale (RASS) of 0 to -2.

Assess patient's sedation and pain level every four hours.

Begin awakening trials for Midazalam and Lorazepam continuous infusions at 4:30am daily. Hold infusions until patient reaches RASS score 0 to +1. Do not institute daily awakening trials if patient is receiving a neuromuscular blocking (NMBA) or paralytic agent.

Begin awakening trials for propofol continuous infusions at 6:00am daily. Hold infusions until patient reaches RASS score of 0 to +1. Do not institute daily awakening trials if patient is receiving a NMBA or paralytic agent.

STEP 1 Assess Patient's Pain using Visual Analog or Cognitively Impaired Scale

I. Rule out environmental or reversible causes
Reposition
Hot and cold therapy
Temperature control

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- ☐ MORPHINE (Drug of choice in patient hemodynamically stable and adequate renal function)
 - 1. Initial dosing: 2mg IVP every 10 min until pain level is less than 4/10
 - 2. Intermittent dosing: 2 mg IVP every 2 hours PRN to maintain pain level less than 4/10. May increase by 1mg every 2 hours to maximum of 4mg every 2 hours prn.
 - 3. If intermittent dosing required more often than every 2 hours, begin continuous infusion of 2mg/hour and titrate to pain level less than 4/10
- ☐ FENTANYL (May be used in patients who are hemodynamically unstable and/or have severe renal dysfunction)
 - 1. Initial dosing: 50mcg IVP every 10 min until pain level is less 4/10.
 - 2. Intermittent dosing: 50 mcg every 2 hours PRN to maintain pain level less than 4/10.
 - 3. If intermittent dosing required more often than every 2 hours, begin continuous infusion of 50 mcg/hour and titrate to pain level less than 4/10

- ☐ HYDROMORPHONE (May be used in patients who are hemodynamically unstable and/or have severe renal dysfunction)
 - 1. Initial dosing: 0.25mg IV every 5 minutes until pain level is less than 3/10
 - 2. Intermittent dosing: 1mg IV every 4 hours PRN to maintain pain level less than 3/10
 - 3. If intermittent dosing required more often than every 4 hours, begin continuous infusion of 1mg/hour and titrate in 0.5 mg/hour increments to maintain pain level less than 3/10

If under medicated, rebolus using intermittent dosing as prescribed above and then increase continuous infusion dose by 50%

If overmedicated, hold continuous infusion until at analgesia goal then restart continuous infusion at a dose decreased by 50%

III. Assess pain/sedation level every 4 hours.

STEP 2 Assess for delirium using the Modified Confusion Assessment Method for the ICU (CAM ICU)

Administer Haloperidol 5mg IVP every 2 hours PRN if delirium is noted on CAM ICU assessment (notify physician if > 100mg administered over 48 hours)

STEP 3 Assess patient's anxiety using the Richmond Agitation Sedation Scale (RASS)

- I. Adequate sedation is mandatory during use of NMBA or paralytic therapy. EEG monitoring is recommended for assessment in patients with NMBA or paralytic infusions
- II. Rule out environmental or reversible causes

Hypoxemia Pain

Hypercarbia Alcohol withdrawal Hyperglycemia Drug withdrawal

Hypoglycemia Electrolyte abnormalities Hypotension Avoid excessive stimulation

III. Use sedatives to achieve goal of 0 to -2 on RASS. (EEG monitoring may be used to supplement RASS. Must be ordered separately)

If frequent neurological assessment is required or sedation is planned to be less than 72 hours:

□ PROPOFOL:

- 1. Initial dosing: 25mg IVP one time bolus
- 2. Start continuous infusion at 5mcg/kg/min.
- 3. Titrate by 5mcg/kg/min every 5 min to RASS goal or max of 80 mcg/kg/min
- 4. Monitor triglycerides 48 hours after infusion started if propofol still infusing. Notify physician for triglyceride level greater than 400mg/dl
- ☐ MIDAZOLAM (should not be used in patients with severe renal dysfunction and/or for more than 72 hours)
 - 1. Initial dosing: 2mg IVP PRN every 20 min to achieve sedation goal
 - 2. Intermittent dosing: 2mg every 1 hour PRN to maintain sedation goal
 - 3. If intermittent dosing is required more often than every hour, begin continuous infusion of 1mg/hour and titrate by 1mg/hr every 10 minutes to RASS goal. Maximum dose is 0.2mg/kg/hr

If less frequent neurological assessment is required or sedation is expected to last longer than 72 hours:

- ☐ LORAZEPAM (Drug of choice for long term sedation)
 - 1. Initial dosing: 2mg IVP PRN every 20 min to achieve sedation goal
 - 2. Intermittent dosing: 2mg every 2 hours PRN to maintain sedation goal
 - 3. If intermittent dosing is required more often than every 2 hours, begin continuous infusion of 1mg/hour titrate by 1mg/hr every hour to sedation goal.

If under medicated, rebolus using intermittent dosing as prescribed above and increase continuous infusion dose by 50%.

If overmedicated, hold continuous infusion until at sedation goal then restart continuous infusion at a dose decreased by 50%.

If patient has been on Propofol for longer than three days and does not have compromised ICP, it is recommended that sedation be changed to Lorazepam.

III. Assess pain/sedation level every 4 hours

STEP 4 Perform awakening trial daily

- I. All sedation should be stopped at time daily awakening trial is performed
- II. Allow patient to awaken to RASS 0 to +1 before restarting

III.	Alert MD for assessment			
IV.	If necessary to restart sedation/analgesia, restart at 50% prior dose			
V.	Do not attempt awakening trial in patients with NMBA or paralytic infusions			
Date:	Physician Signature:			