

# Low-Dose Ketamine Infusion for Analgesia



## Inclusion Criteria:

- Suspected or potential opioid induced hyperalgesia
- Acute pain in patients on chronic high dose opioids
- Neuropathic pain resistant to standard treatments
- Patients with cancer and chronic opioid requirements
- As a component of palliative or end of life care for analgesia

## Exclusion Criteria:

- Allergy to ketamine
- Liver failure
- Myocardial ischemia
- < 3 years of age

## Caution in patients with:

- Uncontrolled hypertension
- Severe hypovolemia
- Psychotic disorders
- Increased ICP in patients not receiving mechanical ventilation
- Cerebral vascular disease and aneurysms
- Renal or hepatic impairment (dose reduction maybe required)
  - Cardiac failure
  - Pulmonary hypertension
- Poorly controlled seizure disorders
  - Thyrotoxicosis

## Pre-assessment

- Obtain baseline vital signs including SPO2 and sedation score prior to start of infusion.
- Continuous pulse oximetry and cardiorespiratory monitoring.

## Dosing

### Analgesia

#### Continuous Infusion

- *Recommended Starting Dose:* 0.05 – 0.1 mg/kg/hr
- *Dosing Range:* 0.05 – 0.3 mg/kg/hr – unless permission given by Pain Management or Palliative Care Teams.
- *Maximum Suggested Dose:* 40 mg/hr - unless permission given by Pain Management or Palliative Care Teams.

### End of Life Care

#### Continuous Infusion

- *Recommended Starting Dose:* 0.05 – 0.1 mg/kg/hr
- Dosing is based on titration to clinical effect/comfort and to avoid occurrence of any undesirable dose limiting side effects.
- Maximal doses will be determined per patient's clinic effect, by the palliative care team.

## Recommendations/Considerations

Treatment or prevention of excessive oral secretions:

- *Recommend:* glycopyrrolate
- *Starting dose:* 0.004 mg/kg/dose IV every 6 hours as needed for hypersalivation.
- IV dose may be increased to a maximum of 0.01 mg/kg/dose (1.5 mg/dose maximum) or 20–40 mcg/kg/dose PO every 6-8 hours.
- Frequency of assessment may be increased based on the patients' response to the ketamine infusion.

## Assessment

- Assess and document respiratory rate, SPO2 and sedation score every 30 minutes x 2, then 1 hour x 1, then every 4 hours (per floor policy and times) after starting ketamine infusion.
  - i.e. if started at 8am, vitals at 0830, 0900, 1000, then 1200, 1600, 2000, 0000.
- For CHOC ketamine titrations in rate: Q30min x 2, then Q4 per unit policy.
- Monitor for possible side effects including:
  - Fatigue
  - Drowsiness
  - Dizziness
  - Vivid dreams
  - Misperceptions or confusion
  - Hallucinations

❖ **Ketamine infusions and titrations are only started during the hours while Pain Service is physically available.**

## Patient Education

- Low-Dose Ketamine Infusion: Patient/Family Education

## Interventions

If patient experiences dysphoria or hallucinations:

- Reduce dose of ketamine and prescribe benzodiazepine (e.g. lorazepam 0.025 mg/kg every 12 hours)
- Consider scheduled low-dose benzodiazepines (lorazepam 0.03 – 0.05 mg/kg/dose IV every 6 hours) to prevent or mitigate psychomimetic side effects, if they occur.

Refer to Patient Care Policy F968  
Low-Dose Ketamine Infusion for Analgesia

## Discharge Criteria

- Completion of weaning protocol for patients on infusion greater than 5 days.
- Successful transition to oral regimen and maintained for 24 hours.

## Pasero Opioid-Induced Sedation Scale (POSS)

**S = Sleep, easy to arouse**

**1 = Awake and alert**

**2 = Slightly drowsy, easily aroused**

**3 = Frequently drowsy, arousable, drifts off to sleep during conversation**

**4 = Somnolent, minimal or no response to verbal and physical stimulation**

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See: Pasero, C. (2009). Assessment of sedation during opioid administration for pain management. *Journal of PeriAnesthesia Nursing*, 24(3), 186-190.

## References

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