

Subject: Scope of Practice/Procedure – ALS
Ketamine Hydrochloride (Ketalar)

Associated Policies:

- I. Class:
 - A. Anesthetic (high dose); analgesic (low dose)
- II. Indications:
 - A. Patient in pain
- III. Therapeutic Effects:
 - A. Blocks impulses of pain perception.
 - B. Suppresses spinal cord activity.
 - C. Affects CNS transmitter systems.
 - D. Provides anesthesia with profound analgesia and minimal respiratory depression and minimal skeletal muscle relaxation.
- IV. Contraindications
 - A. Known hypersensitivity to Ketamine
 - B. Patients with known schizophrenia
 - C. Neonates
- V. Relative:
 - A. Pregnancy.
- VI. Adverse Effects:
 - A. Dysrhythmia, tachycardia, bradycardia
 - B. Nausea/ vomiting
 - C. Emergence reactions including confusion, hallucinations and/or delirium.
 - 1. Less common in Pediatric patients and patients > 65 years of age
 - 2. Administering concomitant benzodiazepines may blunt the emergence phenomena.
- VII. Administration and Dosage – ADULTS and PEDIATRICS
 - A. Analgesia
 - IV: 0.3mg/kg diluted in at least 100cc NS and infused over 10 minutes
 - IM: 0.3 mg/kg
 - IN: 0.5mg/kg
 - Maximum dose of 30 mg.
- VIII. Special Information:
 - A. Patients should be advised of medication administration and subsequent side effects.
 - B. Continuous respiratory and cardiac monitoring is required.
 - C. Monitor closely for signs of emergence phenomena and administer benzodiazepines for severe anxiety per Policy # 6552.

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- D. Monitor closely for adverse side effects.
- E. Do not administer more than 1cc volume via IN.