

# UNC MEDICAL CENTER GUIDELINE

# Dosing Guideline for Intravenous Patient Controlled Analgesia in Patients with Limited Opioid Exposure (Age 6 Years or Above)

This should be used as a guide for the initiation and management of intravenous patient controlled analgesia (PCA-IV) therapy for patients who are *opioid naïve*. Patients who are considered *opioid-tolerant* are those who have been taking, for a week or longer, at least 60 mg of oral morphine daily or 30 mg of oral oxycodone daily or an equianalgesic dose of another opioid. Patients who do not meet the definition of *opioid-tolerant*, are considered *opioid naïve*. PCA doses for patients with chronic pain or cancer pain should be based on prior narcotic requirements and response to therapy; consider consulting the Chronic Pain Service for assistance. PCA use by caregivers is in accordance with the <u>Caregiver and Nurse</u> <u>Controlled Analgesia (CNCA) for Pediatric Patients Policy</u>. **HYDROmorphone is the preferred agent for PCA-IV at UNC Medical Center**. Morphone or fentaNYL are alternatives when certain conditions are present (see dosing charts below).

## Initial Dosing Adjustments

Dosing adjustments are recommended if patients meet one or more of the following criteria:
Age > 60 years
Obesity (BMI > 30 kg/m<sup>2</sup>)
Pulmonary Impairment
Hepatic Impairment
Renal Impairment
Obstructive Sleep Apnea

Recommended dose adjustment guidelines include the following:

- Reduce loading dose by 50%, and reduce intermittent dose by 25%.
- Avoid use of continuous infusions.
- If a continuous infusion is prescribed, monitor closely for acute changes in functional status and vital signs.



## Patient Assessment During PCA-IV Therapy

- Patients should be assessed for pain relief within 60 minutes of starting therapy or dosage changes.
  - o See Pain Management (nursing) and Patient Controlled Analgesia Policies
- If dose needs to be adjusted use the following guideline:
  - Too much drug, decrease dose by 25%.
  - Too little drug, increase intermittent dose by 50%.

# Morphine

• Morphine should be dosed based on lean body weight (see appendix 1)

Lean Body Weight	Morphine Loading Dose (mg)		Morphine Intermittent	Morphine Usual Lockout	Morphine Continuous Infusion
	Initial Dose	Max Initial Dose	Dose (mg)	Interval* (min)	Rate (mg/hour) Only if needed
Usual weight-based	0.05 mg/kg	0.1 mg/kg	0.02 mg/kg		0.02 – 0.04 mg/kg/h
14 – 17 kg	0.7 mg – 0.9 mg	1.4 mg – 1.7 mg	0.3 mg	10 minutes	0.3 mg – 0.6 mg
17.1 – 22 kg	0.8 mg – 1.1 mg	1.7 mg – 2.2 mg	0.4 mg	10 minutes	0.4 mg – 0.8 mg
22.1 – 27 kg	1.1 mg – 1.3 mg	2.2 mg – 2.7 mg	0.5 mg	10 minutes	0.5 mg – 1 mg
27.1 – 32 kg	1.3 mg – 1.6 mg	2.7 mg – 3.2 mg	0.6 mg	10 minutes	0.6 mg – 1.2 mg
32.1 – 37 kg	1.6 mg – 1.8 mg	3.2 mg – 3.7 mg	0.7 mg	10 minutes	0.7 mg – 1.4 mg
37.1 – 42 kg	1.8 mg – 2.1 mg	3.7 mg – 4 mg	0.8 mg	10 minutes	0.8 mg – 1.6 mg
42.1 – 47 kg	2.1 mg – 2.3 mg	4 mg	0.9 mg	10 minutes	0.9 mg – 1.8 mg
> 47 kg	2 mg	4 mg	1 mg	10 minutes	1 mg – 2 mg

#### TABLE 1: INITIAL MORPHINE PCA-IV DOSING GUIDANCE

\*may use lockout as low as 6 minutes for extreme cases of pain; 4h Lockout Limits should be considered for patients with impaired pulmonary, renal or liver function or who are receiving continuous infusion and PCA bolus dosing



## HYDROmorphone

• HYDROmorphone should be dosed based on lean body weight (see appendix 1)

#### TABLE 2: INITIAL HYDROmorphone PCA-IV DOSING GUIDANCE

Lean Body Weight	HYDROmorphone	HYDROmorphone	HYDROmorphone Usual	HYDROmorphone Continuous Infusion
	Loading Dose (mg)	Intermittent Dose (mg)	Lockout Interval* (min)	Rate (mg/hour) Only if needed
Usual weight-based	Max = 0.02 mg/kg	0.003 – 0.004 mg/kg		0.003 – 0.005 mg/kg/h
14 – 17 kg	0.3 mg (max)	0.06 mg	10 minutes	0.04 mg – 0.08 mg
17.1 – 22 kg	0.4 mg (max)	0.08 mg	10 minutes	0.05 mg – 0.11 mg
22.1 – 27 kg	0.5 mg (max)	0.1 mg	10 minutes	0.06 mg – 0.13 mg
27.1 – 32 kg	0.6 mg (max)	0.12 mg	10 minutes	0.08 mg – 0.16 mg
32.1 – 37 kg	0.7 mg (max)	0.14 mg	10 minutes	0.1 mg – 0.18 mg
37.1 – 42 kg	0.8 mg (max)	0.16 mg	10 minutes	0.11 mg – 0.2 mg
42.1 – 47 kg	0.9 mg (max)	0.18 mg	10 minutes	0.12 mg – 0.23 mg
> 47 kg	0.5 mg usual	0.2 mg	10 minutes	0.2 mg – 0.4 mg
	(1 mg max)	_		

\*may use lockout as low as 6 minutes for extreme cases of pain; 4h Lockout Limits should be considered for patients with impaired pulmonary, renal or liver function or who are receiving continuous infusion and PCA bolus dosing

#### fentaNYL

• fentaNYL should be dosed based on lean body weight (see appendix 1)

#### TABLE 3: INITIAL fentaNYL PCA-IV DOSING GUIDANCE

Lean Body Weight	fentaNYL Loading	fentaNYL Intermittent Dose	fentaNYL Usual Lockout	fentaNYL Continuous Infusion Rate
	Dose (mcg)	(mcg)	Interval* (min)	(mcg/hour) Only if needed
Usual weight-based	Max = 1 mcg/kg	0.2 mcg/kg		0.2 – 0.4 mcg/kg/h
14 – 17 kg	17 mcg (max)	3 mcg	8 minutes	3 mcg – 6 mcg
17.1 – 22 kg	22 mcg (max)	4 mcg	8 minutes	4 mcg – 8 mcg
22.1 – 27 kg	27 mcg (max)	5 mcg	8 minutes	5 mcg – 10 mcg
27.1 – 32 kg	32 mcg (max)	6 mcg	8 minutes	6 mcg – 12 mcg
32.1 – 37 kg	37 mcg (max)	7 mcg	8 minutes	7 mcg – 14 mcg
37.1 – 42 kg	42 mcg (max)	8 mcg	8 minutes	8 mcg – 16 mcg
42.1 – 47 kg	47 mcg (max)	9 mcg	8 minutes	9 mcg – 18 mcg
> 47 kg	50 mcg usual	10 mcg	8 minutes	10 mcg – 20 mcg
_	(100 mcg max)			

\*may use lockout as low as 6 minutes for extreme cases of pain; 4h Lockout Limits should be considered for patients with impaired pulmonary, renal or liver function or who are receiving continuous infusion and PCA bolus dosing



### **APPENDIX 1: LEAN BODY WEIGHT CALCUALTIONS**

**Males**: (0.73 x Height in cm) – 59.42

**Females**: (0.65 x Height in cm) – 50.74

Note: To convert inches to cm: inches x 2.54 = centimeters

## REFERENCES

- 1. Cravero JP, et al. The Society for Pediatric Anesthesia recommendations for the use of opioids in children during the perioperative period. Paediatr Anaesth. 2019 Jun;29(6):547-571.
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- 5. Practice guidelines for acute pain management in the perioperative setting: an updated report by the American Society of Anesthesiologists Task Force on Acute Pain Management. Anesthesiology. 2012 Feb;116(2):248-73.