Barnes-Jewish Hospital Adult Intubation Pharmacologic Considerations for ICU/ED Patient Care Areas

				Pretreatment (Optional)	
	Consid	er in patient	s with high ICP to blunt cated	cholamine mediated elevations in BP, HR and ICP in response to intubation	
Agent	Dose	Onset	Duration	Considerations	
Fentanyl	1-2 mcg/kg	<30 sec	0.5-1 hr	 Avoid: Decompensated shock or hemodynamic instability Caution: May blunt sympathetic response leading to cardiac arrest Obesity Dosing: Consider using AdjBW if BMI ≥30, otherwise use TBW 	
Induction					
Agent	Dose	Onset	Duration	Considerations	
Etomidate	0.2-0.3 mg/kg	10-15 sec	4-10 min	 <u>Caution</u>: may cause adrenal suppression ↑ EEG activity, ↓ seizure threshold, myoclonic movements, minimal cardiorespiratory depression Obesity Dosing: Consider using AdjBW if BMI ≥40, otherwise use TBW 	
Ketamine	1-2 mg/kg	10-30 sec	5-15 min	 Caution: Heart failure, hypertension, angina Consider use in shock or reactive airway disease May cause ↑ heart rate/blood pressure, emergence phenomenon, salivation, bronchorrhea Obesity Dosing: Consider using AdjBW if BMI ≥40, otherwise use TBW 	
Propofol	0.5-1.5 mg/kg [◊]	15-45 sec	3-10 min	 Caution: Profound hypotension (titration of induction dose helps prevent severe hemodynamic changes), pulmonary arterial hypertension Preferred induction agent for status epilepticus Consider lower dose in patients at increased risk for hypotension or premedicated with benzodiazepine and/or opioids Obesity Dosing: Consider using AdjBW if BMI ≥30, otherwise use TBW 	
Midazolam	0.2-0.3 mg/kg	60-90 sec	1-4 hr	 Consider lower dose in elderly patients Obesity Dosing: Consider using AdjBW if BMI ≥30, otherwise use TBW 	
				Paralysis (Optional)	
		ĺ	ndicated for use in rapid seq	uence intubation in patients without contraindications.	
	Consider use to	facilitate en	dotracheal intubation in pati	ients that can be rescued with bag mask ventilation AND without contraindications	
Agent	Dose	Onset	Time to return of first twitch on TOF	Considerations	
Rocuronium	0.6-1.2 mg/kg*	60-120 sec		 ↑ in duration with liver and renal dysfunction; May cause bronchospasm or anaphylaxis Obesity Dosing: Consider using IBW if BMI ≥30-39, AdjBW if BMI ≥50, otherwise use TBW Decrease dose by 50-90% in myasthenia gravis (use lower dose for more severe disease) Ensure proper deep sedation is initiated after administration given longer duration of action 	
	1-1.5 mg/kg	60 sec	3-5 min	 May cause arrhythmias, anaphylaxis, malignant hyperthermia, ↑ serum potassium 0.5-1 mEq/L[∆] Obesity Dosing: Consider using AdjBW if BMI ≥60, otherwise use TBW 	
Succinylcholine	neuromuscular j <u>AVOID</u> : Stroke, ii syndrome, meni	unction, or h mmobilizatio ngitis, necro	istory of malignant hyperthe n ≥7 days, any myopathy, sp tizing pancreatitis, catatonic	at predispose to hyperkalemia △, conditions that affect acetylcholine receptor expression at the	

Disclaimer: This document serves as an educational reference but should not supersede clinical judgment with respect to appropriate and necessary care for an individual patient.

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Reversal of Paralysis

Sugammadex

- <u>RESTRICTED USE:</u> Use in ED/ICU patient care areas is restricted to emergent reversal after failure to intubate/ventilate (i.e. impending hypoxic arrest) OR neurologic exam REQUIRED to prevent acute neurologic sequelae (i.e. impending herniation)
- CAUTION: Administration of sugammadex should NOT delay a surgical airway when appropriate

Level of NMB based on TOF Monitoring	Agent	Sugammadex Dose*			
TOF count ≥ 2	ROC or VEC	2 mg/kg			
TOF count = 0-1 [†]	ROC or VEC	4 mg/kg			
Reversal required for failed airway immediately after ROC administration	ROC only	16 mg/kg			
*Dosed on TBW. Round to nearest 200 mg					

- † PTC should be ≥ 1 prior to reversal if patient has 0 twitches on TOF
- Reversal typically occurs within 2-3 minutes. Inadequate dose for depth of blockade may result in delayed neuromuscular recovery (10-20 minutes)
- Adverse effects: Bradycardia, hypersensitivity reactions
- Drug interactions: Impairs effectiveness of hormonal contraceptives for up to 7 days therefore patients with childbearing potential must be counseled
- Recommend waiting at least 4 hours for re-paralysis with rocuronium or vecuronium after reversal with 2-4 mg/kg of sugammadex (in patients with normal renal function), otherwise, use cisatracurium or succinylcholine (if no contraindications)
- Contact central/satellite pharmacy for expedited delivery

Neostigmine (given with glycopyrrolate due to muscarinic adverse effects)

Level of NMB based on TOF monitoring	Neostigmine Dose*	
TOF count = 4 twitches (TOF ratio 0.4-0.9)	30 mcg/kg	
*Dosed on TBW. Max dose 5 mg.		

- Recommended for reversal of minimal depths of neuromuscular blockade only (TOF 4/4, no fade)
- Reversal typically occurs within 10 minutes if sufficient spontaneous recovery was achieved before administration. Deeper blocks will require more time to attain full neuromuscular recovery.
- Adverse effects: Bradycardia, increased oral and respiratory secretions, GI symptoms, hypersensitivity reactions
- Contraindications: Peritonitis, mechanical obstruction of intestinal or urinary tract

Glycopyrrolate (used to minimize neostigmine adverse drug reactions)

- Dose: 0.2 mg IV per 1 mg of neostigmine administered
- Give immediately prior to neostigmine dose

Succinylcholine Reversal

Rocuronium or

Vecuronium

Reversal

- No reversal agent
- Brief duration of paralysis (≤5 min) after one dose

ICP: intracranial pressure; BP: blood pressure; HR: heart rate; NMB: neuromuscular blockade; TOF: train of four; ROC: rocuronium; VEC: vecuronium; PTC: post-tetanic count; TBW: total body weight; AdjBW: adjusted body weight; IBW: ideal body weight

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