

Paralytic Dosing & Reversal Management

Utilizing quantitative TOF monitoring

Intubation & Intraoperative Management

Rocuronium Dosing Guidelines – Elective Intubation (Non-RSI)

Intubation Dose: Rocuronium Bromide 0.6 mg/kg (IBW)

- Dose adjusted for sex: Dose reduction females (15%)
- Dose adjusted for age: For each year > 55 (1%/year)

Incremental Dosing

- 20% of ED₉₅ Dose (ED₉₅ Dose Rocuronium = 0.3 mg/kg)

Redosing Indications via Quantitative Monitoring

Redose @ TOFC of 3

- If profound block is necessary, redose @ PTC at count of 1

Avoid total twitch loss (PTC of 0)

- Avoid redosing in last 30 minutes of the procedure

Extubation Timing

- TOFR ≥ 90% via Quantitative Monitor

Reversal Management

Quantitative TOF Value	Reversal Agent and Dose [†]
TOFR ≥ 90%	Reversal not required
TOFR > 40%	Neostigmine 40 mcg/kg (Max 5mg)*
TOFR < 40% TOFC 3, 2, or 1	Sugammadex 2 mg/kg**
TOFC 0 PTC ≥ 1 PTC 0 (non-emergent)	Sugammadex 4 mg/kg**
PTC 0 (emergent) Can't intubate/ventilate	Sugammadex 16 mg/kg**

[†]After Rocuronium or Vecuronium

This guide is based on published literature. Clinicians should use their own judgment.

*Neostigmine dosed on Ideal Body Weight; Neostigmine requires time to work; time reversal with expected extubation time. If ≥ 90% TOFR is not achieved in **15 minutes**, dose with 2 mg/kg Sugammadex and confirm 90%

**Sugammadex dosed on Actual Body Weight.

1. S. R. Thilen et al., *Anesthesiology*. 138, 13–41 (2023).
2. Merck & Co., Inc., Bridion® dosing considerations.