Santé	REGIONAL ADULT PARENTERAL DRUG MONOGRAPH		
Southern Sud	GENERIC NAME sugammadex		
Effective Date: Mar 14 2018	CLASSIFICATION	OTHER NAMES	PAGE
Revised Date: Nov 2024	Antidote (for neuromuscular blocker)	Bridion	1 of 1
ADMINISTRATION POLICY:			
IV Bolus - Administration restricted to anesthetists in OR			
<b>RECONSTITUTION/DILUTION/ADMINISTRATION:</b> <b>Available as:</b> 100 mg/mL - 2 mL, 5 mL single dose vial PROTECT FROM LIGHT (When not			
Available as: 100 mg/mL - 2 mL, 5 mL single dose vial PROTECT FROM LIGHT (When not protected from light, the vial should be used within 5 days.)			
<b>IV Bolus:</b> Undiluted into an infusing IV, based on monitoring for twitch responses and the			
	extent of spontaneous reco	overy	
Maximum rate:	10 seconds		
Maximum concentration:	100 mg/mL		
DOSAGE:			
Routine reversal of rocuronium induced blockage: 2 to 4 mg/kg IV once Immediate reversal of rocuronium induced blockade: 16 mg/kg IV once			
NOTE: Dosing is based on actual body weight			
STABILITY/COMPATIBILITY:			
<b>Compatibility:</b> Compatible with D5W, normal saline, combination dextrose-saline solutions and ringer's			
lactate.			
PRECAUTIONS, POTENTIAL ADVERSE REACTIONS:			
• An anaphylactic hypersensitivity reaction has been identified as a rare but significant adverse reaction			
associated with this agent. Nausea and vomiting are common after reversal, rates are similar to those seen with neostigmine reversal. Have EPINEPHRrine readily available for each dose			
Contraindications: Previous hypersensitivity reaction			
ADDITIONAL NOTES AND NURSING CONSIDERATIONS:			
• <b>Bradycardia:</b> Marked bradycardia and bradycardia with cardiac arrest have been reported, usually within minutes after administration. Monitor closely for hemodynamic changes during and after reversal of			
neuromuscular blockade.			
• <b>Respiratory monitoring:</b> Ventilatory support is mandatory for patients until adequate spontaneous respiration			
<ul> <li>is restored and the ability to maintain a patent airway is assured.</li> <li>Hemostatic and coagulation parameters in patients with risk for impaired hemostasis</li> </ul>			
<ul> <li>Hemostatic and coagulation parameters in patients with risk for impaired hemostasis</li> <li>For re-administration of rocuronium a reversal time of 24 hours is suggested.</li> </ul>			
<ul> <li>May diminish effect of hormonal contraceptives (oral or non-oral). Patients on hormonal contraceptives should</li> </ul>			
use a nonhormonal contraceptive method for 7 days after sugammadex treatment.			
Some patients experience a delayed or minimal response to recommended doses of sugammadex.			