Santé	REGIONAL ADULT PARENTERAL DRUG MONOGRAPH GENERIC NAME erythromycin lactobionate			
Southern Sud				
Effective Date: Dec 2012	CLASSIFICATION Antibiotic	other names Erythrocin	PAGE	
Revised Date: Nov13 2013 Review Date: Mar 14 2018	Antibiotic	Li yun ochi	1 of 1	
ADMINISTRATION POLICY:				
IV Intermittent - May be administered by a nurse				
IV Bolus - Not recommended				
IM Injection - Not recommended				
RECONSTITUTION/DILUTION/ADMINISTRATION:				
Available as:1 gram vial: Add 20 mL sterile water for injection.				
Do not use diluents containing preservatives.				
Final Volume: 20 mL Final Concentration: 50 mg/mL				
IV Intermittent: Dilute ordered dose in 250 mL normal saline, and administer over 60 minutes.				
EXCEPTION: May administer centrally in not less than 100 mL of compatible diluent for a fluid restricted patient.				
Maximum rate:Over 20 minutes (slow infusions preferred to minimize local irritation)				
Maximum concentration: Peripheral: 5 mg/mL				
DOSAGE:	Central: 10 mg/ml	ـــــــــــــــــــــــــــــــــــــ		
Usual:	500 to 1000 mg IV every 6 hours.			
Maximum single dose:	1000 mg			
Maximum daily dose:				
STABILITY/COMPATIBILITY:				
Stability of Reconstituted Solution: 24 hours at room temperature				
Stability of Final Admixture	e: 8 hours at room te	8 hours at room temperature in normal saline.		
Compatibility:	Compatible with	normal saline, D5W, D5W-normal saline	alutions	
Company.		Lactated Ringer		
	Lactated Kinger			
PRECAUTIONS, POTENTIAL ADVERSE REACTIONS:				
• Thrombophlebitis (pain, and inflammation at the injection site) particularly with higher concentrations.				
• Decreasing the rate of infusion or using more dilute solutions may minimize local irritation.				
Hypersensitivity reactions: skin rash, urticaria				
Nausea, vomiting, abdominal discomfort, diarrhea, increased liver enzymes.				
• Reversible hearing loss (rare; usually occurs with doses greater than or equal to 4 grams/day)				
ADDITIONAL NOTES AND NURSING CONSIDERATIONS:				
• IM use or IV Bolus not recommended since extravasation of drug into tissues may cause tissue injury.				
• No dosage adjustment necessary in patients with mild-moderate hepatic or renal impairment. Use with caution				
in patients with significant hepatic or renal disease				