

ERYthromycin

Indication	<ul style="list-style-type: none"> • Treatment of infections due to susceptible gram-positive and gram-negative organisms¹ <ul style="list-style-type: none"> ○ Chlamydia trachomatis, mycoplasma, ureaplasma² ○ Azithromycin or clarithromycin preferred for pertussis treatment or prevention 														
ORAL	Presentation	<ul style="list-style-type: none"> • Powder for oral solution: 200 mg in 5 mL 													
	Dosage ^{3,4}	<table border="1" data-bbox="391 443 1300 607"> <thead> <tr> <th>Dose</th> <th>Day of life</th> <th>Frequency</th> </tr> </thead> <tbody> <tr> <td rowspan="3">10 mg/kg</td> <td>0–7 days</td> <td>every 12 hours</td> </tr> <tr> <td>8–28 days</td> <td>every 8 hours</td> </tr> <tr> <td>29 or more</td> <td>every 6 hours</td> </tr> </tbody> </table>	Dose	Day of life		Frequency	10 mg/kg	0–7 days	every 12 hours	8–28 days	every 8 hours	29 or more	every 6 hours		
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Preparation	<ul style="list-style-type: none"> • Add water for injection to dry powder according to label instructions • Shake vigorously 														
Administration	<ul style="list-style-type: none"> • Draw up prescribed dose into oral/enteral syringe • Oral/OGT/NGT 														
INTRAVENOUS	Presentation	<ul style="list-style-type: none"> • Vial: 1 g 													
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Preparation (Step 1: all babies)	<ul style="list-style-type: none"> • Add 3 mL of water for injection to 1 g vial • Withdraw entire contents and make up to 20 mL total volume with water for injection⁵ <ul style="list-style-type: none"> ○ Concentration now equal to 50 mg/mL • Draw up DOUBLE the prescribed dose from the 50 mg/mL solution • Then prepare according to either criteria A or criteria B 														
Preparation (Criteria A)	<ul style="list-style-type: none"> • Baby: 2.5–5 kg without CVL <ul style="list-style-type: none"> ○ Further dilute to a maximum concentration⁶ of 5 mg/mL with 0.9% sodium chloride ○ Prime the infusion line and reduce total syringe volume by half ○ Prescribed dose in remaining half of syringe volume 														
Preparation (Criteria B)	<ul style="list-style-type: none"> • Baby 2.5–5 kg with CVL OR baby less than 2.5 kg (with or without CVL) <ul style="list-style-type: none"> ○ Further dilute to a maximum concentration of 10 mg/mL with 0.9% sodium chloride ○ Prime the infusion line and reduce total syringe volume by half ○ Prescribed dose in remaining half of syringe volume 														
Administration	<ul style="list-style-type: none"> • IV infusion via syringe driver pump over 60 minutes <ul style="list-style-type: none"> ○ Via PVL⁶: maximum concentration of 5 mg/mL ○ Via CVL⁶: maximum concentration of 10 mg/mL • On completion, disconnect syringe and infusion line • Flush access port at same rate as infusion 														
Special considerations	<ul style="list-style-type: none"> • Higher dosage may be required for chlamydia infection (12.5 mg/kg every 6 hours)² <ul style="list-style-type: none"> ○ Seek expert advice (e.g. consultant/pharmacist) before prescribing • Caution <ul style="list-style-type: none"> ○ If co-administered with any drug metabolised by cytochrome P450 enzyme (CYP3A4)¹ ○ Cardiac toxicity reported in preterm neonates⁷ ○ Serious ventricular arrhythmias reported in adults with IV doses⁵ of 1 g ○ Avoid in acute porphyria⁸ • UAC route: Consult with neonatologist/paediatrician and refer to Queensland Clinical Guideline: <i>Neonatal medicines</i>⁹ 														

Monitoring	<ul style="list-style-type: none"> Extravasation risk: may cause severe tissue damage⁶ High doses IV may cause pain along the vein⁶
Compatibility	<ul style="list-style-type: none"> Fluids <ul style="list-style-type: none"> 0.9% sodium chloride⁶ Y-Site <ul style="list-style-type: none"> Aciclovir⁶, amikacin⁶, amiodarone⁶, atracurium⁶, azathioprine⁶, calcium gluconate⁶, cefotaxime⁶, ceftriaxone⁶, dexmedetomidine⁶, esmolol⁶, fluconazole⁶, foscarnet⁶, gentamicin⁶, hydrocortisone sodium succinate⁶, hydromorphone⁶, midazolam⁶, morphine sulfate⁶, suxamethonium⁶, tobramycin⁶, verapamil⁶, zidovudine⁶
Incompatibility	<ul style="list-style-type: none"> PN and fat emulsion: co-infusion with ERYthromycin not recommended (evidence limited). If unavoidable, seek pharmacist advice first, filter infusion and flush before and after Fluids <ul style="list-style-type: none"> Glucose solutions⁴ Y site <ul style="list-style-type: none"> Cefalotin⁶, cefazolin⁶, dexamethasone⁶, flucloxacillin⁶, furosemide (frusemide)⁶, heparin sodium⁶, linezolid⁶, phenobarbital (phenobarbitone)⁶, phenytoin⁶, rocuronium⁶
Interactions	<ul style="list-style-type: none"> Increases concentration of digoxin, midazolam and carbamazepine⁸
Stability	<ul style="list-style-type: none"> Vial <ul style="list-style-type: none"> Store below 25 °C⁵ Oral solution <ul style="list-style-type: none"> Store reconstituted solution in the fridge at 2–8 °C and use within 10 days
Side effects	<ul style="list-style-type: none"> Circulatory: prolonged QT interval and ventricular arrhythmias associated with IV use, particularly related to rapid administration and high doses⁷ Digestive: vomiting¹, diarrhoea¹, infantile hypertrophic pyloric stenosis (associated with longer term use of ERYthromycin in neonates, especially those aged less than 2 weeks¹) Nervous: ototoxicity (after high doses in patients with impaired renal or hepatic function²) Hypersensitivity: rashes, pruritus, urticaria or angioneurotic oedema. Anaphylaxis has been reported¹
Actions	<ul style="list-style-type: none"> Macrolide antibiotic that inhibits protein synthesis and thereby bacterial growth¹
Abbreviations	IV: intravenous, CVL: central venous line, OGT: orogastric tube, PVL: peripheral venous line NGT: nasogastric tube
Keywords	Erythromycin, ERYthromycin, antibiotic, macrolide

The Queensland Clinical Guideline *Neonatal Medicines* is integral to and should be read in conjunction with this monograph. Refer to the disclaimer. Destroy all printed copies of this monograph after use.

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