

## ERYTHROMYCIN

<b>Trade Name</b>	E-Mycin (ethylsuccinate) Oral (Alphapharm) Erythrocin (lactobionate) IV (AFT Pharmaceuticals)										
<b>Class</b>	Macrolide antibiotic										
<b>Mechanism of Action</b>	Has bacteriostatic and bactericidal effects, partially by inhibiting protein synthesis. Motilin receptor agonist and induces stomach and small intestine motor activity.										
<b>Indications</b>	<p><b>Indication 1:</b> Treatment of infections caused by Chlamydia Ureaplasma, Mycoplasma.</p> <p><b>Indication 2:</b> Substitute for penicillin if significant allergy</p> <p><b>Indication 3:</b> Prophylactic treatment for Bordetella pertussis.</p> <p><b>Indication 4:</b> To treat gastric stasis or gut dysmotility</p>										
<b>Contraindications</b>	Hypersensitivity to erythromycin. Liver impairment In patients on cisapride, risk of life threatening arrhythmias.										
<b>Supplied As</b>	<p><b>Oral:</b> Erythromycin ethylsuccinate 200mg/5mL and 400mg/5mL</p> <p><b>IV:</b> Erythromycin lactobionate 1g vial</p>										
<b>Dilution</b>  <b>*Two dilution steps required*</b>	<p><b>Oral:</b> Prepare suspension as per manufacturers instructions</p> <p><b>IV:</b></p> <table border="1"> <thead> <tr> <th>Drug</th> <th>Water Added</th> <th>Volume</th> <th>Concentration</th> </tr> </thead> <tbody> <tr> <td>1g</td> <td>20mL</td> <td>20mL</td> <td><b>50mg/mL</b></td> </tr> </tbody> </table> <p><b>Then further dilute</b> by taking 1mL (50mg) and diluting with 9ml of sterile water to give a <b>final concentration of 5mg/mL</b></p>			Drug	Water Added	Volume	Concentration	1g	20mL	20mL	<b>50mg/mL</b>
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1g	20mL	20mL	<b>50mg/mL</b>								
<b>Dosage</b>	<p><b>Indication 1/2:</b> 12.5 mg/kg/dose for 14 days</p> <p><b>Indication 3:</b> 10mg/kg/dose for 14 days</p> <p><b>Indication 4:</b> 3 mg/kg/dose</p>										
<b>Interval</b>	6 hourly										
<b>Administration</b>	<p><b>Oral:</b> Shake well before giving, (administration with food increases absorption)</p> <p><b>IV:</b> Infusion over 60min</p>										

<b>Compatible With</b>	<p><b>Solution:</b> 0.9% sodium chloride and sterile water for injection.</p> <p><b>Terminal Y- site:</b> aciclovir, amiodarone, famotidine, heparin, lidocaine, lorazepam, magnesium sulfate, midazolam, morphine, nicardipine, penicillin G, pentobarbital, potassium chloride, ranitidine, sodium bicarbonate, and zidovudine.</p>
<b>Incompatible With</b>	<p><b>Solution:</b> dextrose 5% and 10% if exposed for longer than 2 hours, unless dextrose is buffered with sodium bicarbonate</p> <p><b>Terminal Y-site:</b> ampicillin, ceftazidime, chloramphenicol, flucloxacillin, fluconazole, frusemide, metoclopramide</p>
<b>Interactions</b>	<p>Erythromycin is a strong inhibitor of CYP3A4 and has the capacity to significantly increase serum concentrations of drugs which are metabolised by this enzyme eg amiodarone amlodipine, ciclosporin, cisapride, dexamethasone, hydrocortisone, midazolam, sildenafil, theophylline.</p> <p>Conversely carbamazepine and rifampicin may induce metabolism of erythromycin and reduce erythromycin serum concentrations.</p> <p>Erythromycin can cause QT interval prolongation and ventricular arrhythmias if administered too rapidly or used in combination with other medicines that also cause this eg cisapride, domperidone, fluconazole, sildenafil,</p> <p>Loop and thiazide diuretics may cause hypokalaemia leading to increased risk of QT interval prolongation.</p>
<b>Monitoring</b>	<p>Monitor heart rate and blood pressure during IV administration</p> <p>Routine monitoring of electrolytes to avoid hypokalaemia or hypomagnesaemia.</p> <p>Watch for abdominal discomfort</p>
<b>Stability</b>	<p><b>Oral:</b> Prepared suspension stable for 10 days.</p> <p><b>IV:</b> Discard opened vial immediately after use</p> <p>Discard unused reconstituted 5mg/mL solution immediately</p> <p>Use a new vial to draw up each dose</p>
<b>Storage</b>	<p><b>Oral:</b> Store in the fridge</p> <p><b>IV:</b> Unopened vials store at &lt;25 °C</p>

<b>Adverse Reactions</b>	<p><b>Oral:</b> nausea, vomiting, oral candida, cholestatic jaundice hypertrophic pyloric stenosis (RR 0.4%).</p> <p><b>IV:</b> bradycardia, hypotension, hypertrophic pyloric stenosis, (RR = 0.4%), intrahepatic cholestasis, loose stools, hearing loss (reversible), venous irritation.</p>
<b>Metabolism</b>	Half life of 2hrs. Protein binding 75 –90%. Drug penetrates CNS poorly, concentrates in liver, bile. Demethylated in liver. Secreted via the bowel.
<b>References</b>	<ol style="list-style-type: none"> <li>1. Neofax, 1999</li> <li>2. Medicines for Children, RCPCH, 1999</li> <li>3. Acta Paediatrica 87(10) 1079 – 84</li> <li>4. Lancet 1999; 354, 2101</li> <li>5. Trissel LA, Handbook of Injectable Drugs, ASHSP 2001</li> <li>6. BNF for Children 2011-2012*</li> <li>7. <a href="http://www.ANMFonline.com">www.ANMFonline.com</a></li> <li>8. nzfc in <a href="http://www.nzf.org.nz">www.nzf.org.nz</a></li> <li>9. Murchison L, De Coppi P, Eaton S. Post-natal erythromycin exposure and risk of infantile hypertrophic pyloric stenosis: a systematic review and meta-analysis. Pediatr Surg Int. 2016;32:1147-52</li> </ol>
<b>Updated By</b>	<p>Nicola Austin Mar 01, May 02  P Schmidt, B Robertshawe Dec 2004  A Lynn, B Robertshawe September 2009, August 2010  A Lynn, B Robertshawe Oct 2012 (re-order profile, 2 dilutions, discard vial)  A Lynn, B Robertshawe Feb 2017 change of liquid supplier &amp; expiry  A Lynn, M Wallenstein, B Robertshawe March 2021 (review/update)</p>