

**CEFUROXIME**

<b>Trade Name</b>	Cefuroxime Actavis® or Cefuroxime AFT®																		
<b>Class</b>	Antibiotic - second generation cephalosporin.																		
<b>Mechanism of Action</b>	<p>Inhibition of bacterial cell wall synthesis through binding to essential proteins.</p> <p>Resistant to most <math>\beta</math>-lactamases and has activity against a wide range of gram-positive and gram-negative organisms including <i>H.influenzae</i>.</p>																		
<b>Indications</b>	<p>Broad-spectrum antibiotic indicated for empiric treatment of infection or where sensitivity is established.</p> <p>Biliary tract infection, meningitis, septicaemia, pneumonia, peritonitis, UTIs.</p> <p>Infection prophylaxis or treatment after surgery.</p>																		
<b>Contraindications</b>	<p>Known hypersensitivity to cefuroxime or other cephalosporins.</p> <p>Caution in patients with a known penicillin allergy.</p>																		
<b>Supplied As</b>	750mg powder vials																		
<b>Dilution</b>	<p>Cefuroxime Actavis</p> <table border="1"> <thead> <tr> <th>Vial</th> <th>Water Added</th> <th>Final Volume</th> <th>Concentration</th> </tr> </thead> <tbody> <tr> <td>750mg</td> <td>7mL*</td> <td>7.5mL</td> <td>100mg/mL</td> </tr> </tbody> </table> <p>*Displacement value = 0.5mL</p> <p>Cefuroxime AFT</p> <table border="1"> <thead> <tr> <th>Vial</th> <th>Water Added</th> <th>Final Volume</th> <th>Concentration</th> </tr> </thead> <tbody> <tr> <td>750mg</td> <td>6.7mL*</td> <td>7.5mL</td> <td>100mg/mL</td> </tr> </tbody> </table> <p>*Displacement value = 0.8mL</p> <p>Reconstituted solution is pale yellow</p>			Vial	Water Added	Final Volume	Concentration	750mg	7mL*	7.5mL	100mg/mL	Vial	Water Added	Final Volume	Concentration	750mg	6.7mL*	7.5mL	100mg/mL
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<b>Dosage</b>	<p><b>Prophylaxis:</b> 25mg/kg/dose</p> <p><b>Severe infection:</b> 50mg/kg/dose</p>																		
<b>Interval</b>		<b>Postnatal Age (days)</b>	<b>Interval (hourly)</b>																
		0-7	12																
		8-14	8																
		>14	6																
<b>Administration</b>	<p><b>IV:</b> Slow push over 3-5 minutes</p> <p><b>IM: Not recommended in infants &lt;1 month of age</b></p>																		

<b>Compatible With</b>	<p><b>Solution:</b> sodium chloride 0.9%, dextrose 5%, dextrose 10%, dextrose saline, lactated ringers, potassium chloride</p> <p><b>Y-site:</b> aciclovir, adrenaline, amikacin*, aminophylline, amphotericin liposomal, atropine, benzyl penicillin, calcium gluconate, cefazolin, cefotaxime, ceftazidime, clindamycin, dexmedetomidine, digoxin, dopamine, ephedrine, epoetin alfa, erythromycin, famotidine, flucloxacillin, furosemide, fentanyl, gentamicin*, glycopyrrolate, heparin, hydrocortisone, indometacin, insulin, lidocaine, meropenem, methylprednisolone, metronidazole, milrinone, morphine, noradrenaline, octreotide, pancuronium, paracetamol, piperacillin, phenylephrine, potassium chloride, propofol, pyridoxine, ranitidine, sodium acetate, thiamine, ticarcillin, tobramycin*, vasopressin, TPN</p> <p>*Some sources suggest theoretical interaction between cephalosporins and aminoglycosides if possible separate administration by one hour.</p>
<b>Incompatible With</b>	Amoxicillin, calcium chloride, dexamethasone, diazoxide, dobutamine, doxapram, fluconazole, ganciclovir, magnesium sulphate, midazolam, phenobarbital, phenytoin, sodium bicarbonate, sulphamethoxazole-trimethoprim, vancomycin (variable). Note: Compatibility of cefuroxime with SMOF lipid has not been tested and hence it is not possible to recommend giving cefotaxime in the same line as lipid solution
<b>Monitoring</b>	N/A
<b>Stability</b>	Discard remaining solution in vial after reconstitution Use a new vial for each dose Vials are not designed for multidosing
<b>Storage</b>	Room temperature for powder vials. Protect from light
<b>Adverse Reactions</b>	Thrombophlebitis may occur following IV injection. Hypersensitivity: rashes, pruritis and fever. Gastrointestinal: diarrhoea, nausea and vomiting. Neutropenia, susceptibility to fungal super infection Rarely: transient increase in LFTs.
<b>Metabolism</b>	33-50% protein bound. Cefuroxime does not undergo metabolism and 85-90% is excreted unchanged in urine (in first 24 hours). Half-life 1-6 hours depending on postnatal age
<b>Comments</b>	Modify dose in patients with severe renal impairment. Each 750mg vial of cefuroxime contains 1.8mmol of sodium.
<b>References</b>	<ol style="list-style-type: none"> <li>1. Shann F., Drug Doses, 11<sup>th</sup> edition. 2001.</li> <li>2. Medicines for Children, 1999.</li> <li>3. Paediatric Dosage Handbook, 8<sup>th</sup> edition. 2001-2002.</li> <li>4. BNF March 2003, 45<sup>th</sup> edition.</li> <li>5. Notes on Injectable Drugs, 4<sup>th</sup> edition.</li> <li>6. Trissel LA, Handbook on Injectable Drugs, 12<sup>th</sup> Ed, 2003</li> <li>7. <a href="http://www.medsafe.govt.nz">www.medsafe.govt.nz</a></li> <li>8. <a href="https://www.starship.org.nz/guidelines/cefuroxime-sodium/www.nzf.org.nz">https://www.starship.org.nz/guidelines/cefuroxime-sodium/www.nzf.org.nz</a></li> <li>9. <a href="http://www.micromedexsolutions.com">www.micromedexsolutions.com</a> (online access to Trissell )</li> </ol>

<b>Updated By</b>	A Lynn, B Robertshawe, F Robertson	May 2009 (new pumps)
	A Lynn, B Robertshawe	June 2012 (re-order profile), Nov 2012 (discard vial)
	A Lynn, M Wallenstein, B Robertshawe	May 2021 brands
	A Lynn, B Robertshawe	Oct 23 – routine review