

PHENOXYMETHYLPENICILLIN

Trade Name	Phenoxyethylpenicillin Oral Solution (AFT Pharmaceuticals)	
Class	Beta-lactam antibiotic	
Mechanism of Action	Inhibits bacterial cell wall synthesis through interaction with penicillin-binding-proteins	
Indications	Treatment of infection of sensitive organisms (most commonly Group B Strep).	
Contraindications	Allergy to penicillins. Large doses may cause hypokalaemia and hypernatraemia	
Supplied As	Powder for reconstitution to make 125mg/5mL oral solution (orange flavour)	
Dilution	Water Added	Concentration
	See instructions on the bottle as different brands vary	125 mg / 5mL
Dosage	12.5 mg/kg/dose	
Interval	Every six hours	
Administration	Oral	
Compatible With	N/A	
Incompatible With	N/A	
Interactions	No known interactions that preclude concomitant use. Give at a different time of day to Gaviscon	
Monitoring	Urea, creatinine and electrolytes	
Stability	Discard after 10 days	
Storage	Powder – maintain at room temp; protect from light and moisture. Reconstituted solution –store in the fridge(2- 8 °C) Protect from light	
Adverse Reactions	Anaphylaxis and both immediate and delayed hypersensitivity reactions. High doses can cause CNS toxicity and seizures.	
Metabolism	Bioavailability = 60 – 70% Half life = 1-2 hrs in neonates (prolonged in renal impairment) Time to peak serum concentration = 30 – 60mins Renal excretion.	
Comments	0.35mmol of potassium per 125mg of phenoxyethylpenicillin Each 5mL of 125mg/5mL phenoxyethylpenicillin oral solution contains 2.8g of sugar	

References	<ol style="list-style-type: none">1. Paediatric Dosage Handbook Taketomo et al 16th Ed 2009.2. BNF for children 20093. www.medsafe.govt.nz
Updated By	A Lynn, B Robertshawe Dec 2011 A Lynn, B Robertshawe Dec 2012 (re-order profile) A Lynn, B Robertshawe Oct 2013 oral dilution instructions A Lynn, B Robertshawe Dec 2021 (routine review, no changes)