## **Newborn use only**

Alert	High risk medicine.				
	Antimicrobial Stewardship Team recommends this drug is listed as unrestricted.				
Indication	Treatment				
	1. Susceptible gram positive (including <i>Streptococcus</i> species, <i>Enterococcus faecalis</i> and <i>Listeria</i>				
	monocytogenes),				
	2. Susceptible gram-negative bacteria (some strains of <i>Escherichia coli</i> , non-beta-lactamase-				
	producing Haemophilus influenzae, Neisseria meningitidis, non-penicillinase-producing strains of Proteus and Salmonellae).				
	Empiric treatment of suspected early onset sepsis including meningitis, with an				
	aminoglycoside.				
	Prophylaxis				
	Urinary Tract Infection (UTI) prophylaxis				
	Asplenia/hyposplenism prophylaxis				
Action	Bactericidal – inhibits synthesis of the bacterial cell wall. Amoxicillin is hydrolysed by beta-lactamases				
	and therefore not effective against penicillinase-producing bacteria.				
Drug Type	Antibacterial – semi-synthetic, bactericidal aminopenicillin				
Trade Name	Multiple brands are available				
Presentation	IV: Amoxicillin 1 g vial.				
D	Oral: Syrup 125 mg/5 mL and 250 mg/5 mL; Paediatric drops 100 mg/mL.				
Dosage	Treatment - IV				
	Standard infections: 50 mg/kg/dose.  Meningitis: 100 mg/kg/dose.				
	Corrected gestational age/postmenstrual age*	Day of life	Interval		
	< 30 <sup>+0</sup> weeks	0–28 days	12 hourly		
	< 30 <sup>+0</sup> weeks	29+ days	8 hourly		
	30 <sup>+0</sup> –36 <sup>+6</sup> weeks	0–14 days	12 hourly		
	30 <sup>+0</sup> –36 <sup>+6</sup> weeks	15+ days	8 hourly		
	37 <sup>+0</sup> –44 <sup>+6</sup> weeks	0–7 days	12 hourly		
	37 <sup>+0</sup> –44 <sup>+6</sup> weeks	8+ days	8 hourly		
	*Also referred to as "current gestational age"		<del></del>		
	Treatment - ORAL				
	Treatment: 25–50 mg/kg/dose.				
	Corrected gestational age/postmenstrual age*	Day of life	Interval		
	37 <sup>+0</sup> –44 <sup>+6</sup> weeks	0–7 days	12 hourly		
	37 <sup>+0</sup> –44 <sup>+6</sup> weeks	8+ days	8 hourly		
	*Also referred to as "current gestational age"	· ·	<u>, , , , , , , , , , , , , , , , , , , </u>		
	Prophylaxis - ORAL				
	Urinary Tract Infection: 10–15 mg/kg/dose once a day Asplenia/hyposplenism: 20 mg/kg/dose once a day (14)				
Mayimum Daily	300 mg/kg/day	i uay (14)			
Maximum Daily	300 mg/ kg/ day				
Dose					
Route	IV IM (only if IV route not possible as intramuscular route is painful) ORAL				
Preparation	IV				
- I	Add 9.2 mL of water for injection to the 1 g vial to make 100 mg/mL solution.				
	FURTHER DILUTE				
	Draw up 5 mL (500 mg of amoxicillin) of the above solution and add 5 mL sodium chloride 0.9% to make				
	a final volume of 10mL with a final concentration of 50 mg/mL.  Use immediately as concentrated solution >30 mg/mL is not stable.(9)				
	IM				

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	Add 3.2 mL of water for injection or lidocaine (lignocaine) 1% to the 1g vial to make 250 mg/mL solution.  ORAL
	1. Syrup 125 mg/5 mL: Add 87 mL of water for irrigation to make a final volume of 100mL with a final
	concentration of 125mg/5mL of suspension.
	2 Syrup 250mg/5mL: Add 87 mL of water for irrigation to make a final volume of 100mL with a final concentration of 250mg/5mL of suspension.
	3. Paediatric drops 100 mg/mL: Add 18 mL water for injection to make a final volume of 21mL with a
_	final concentration of 100mg/mL.
Administration	IV: Infuse over 30 minutes into the proximal cannula site.
	Separate from aminoglycosides by clearing the lines with a flush as penicillins inactivate them.  IM injection: Only if IV route is not possible.
	PO: The liquid preparation should be shaken well. After mixing, administer immediately. The dose may
	be mixed with milk.
Monitoring	Monitoring is not required.
	Follow infectious disease/microbiology advice in case of poor therapeutic response.
Contraindications	Hypersensitivity to penicillins (unlikely to be an issue in neonates).
Precautions	Hypersensitivity to cephalosporins (unlikely to be an issue in neonates).
	In renal impairment, the excretion of amoxicillin will be delayed. In infants with severe renal
	impairment, it may be necessary to reduce the total daily dose.
<b>Drug Interactions</b>	IV: Aminoglycosides, including gentamicin, should not be mixed with amoxicillin when both drugs are
	given parenterally as inactivation of the aminoglycoside occurs. Ensure line is adequately flushed
	between antibiotics.
A d	PO: No significant drug-drug interaction found for neonates on oral amoxicillin.  Common: Diarrhoea, skin rash (erythematous maculopapular), phlebitis at the injection site,
Adverse	superinfection with resistant organisms during prolonged therapy.
Reactions	Uncommon/rare: Neurotoxicity, electrolyte disturbances e.g. hypernatraemia due to the sodium
	content (2.6 mmol per gram in Fisamox IV and 3.3 mmol per gram in Ibiamox IV), erythema
	multiforme, exfoliative skin lesions, <i>C. difficile</i> diarrhoea, pancytopenia, raised liver enzymes.
	Amoxicillin may result in a false positive for glucose in the urine due to excessive amounts of urinary
	amoxicillin.
Compatibility	Fluids: Glucose 5%, glucose 5% in sodium chloride 0.45% (less stable in carbohydrate solutions, it is
	preferable to avoid adding it to them) (13), sodium chloride 0.9%, water for injection (WFI)
	Vicitor No information (0)
Incompatibility	Y site: No information (9) Fluids: Blood products, dextran, fat emulsions, amino acid solutions
incompatibility	Traids. Blood products, dextrail, fat emaisions, amino dela solutions
	Y site: Amikacin, ciprofloxacin, gentamicin, imipenem-cilastatin, midazolam, potassium chloride,
	rocuronium, sodium bicarbonate, tobramycin (9)
Stability	IV: The reconstituted solution should be administered immediately; discard unused portion.
-	A transient pink or slight opalescence may appear during reconstitution. Do NOT administer if
	reconstituted solution is pink.
•	PO: The medication mixed with milk should be administered immediately.
Storage	IV: Store below 25°C. Protect from light. PO: Store unreconstituted powder for oral suspension at 20–25 °C. Reconstituted suspension is stable
	for 14 days at room temperature or if refrigerated. Refrigeration is preferred.
Special	Clearance is primarily by the renal route. Clearance increases with increasing gestational age and
Comments	postmenstrual age. Serum half-life is longer in premature infants and infants younger than 7 days.
Comments	Fisamox and Ibiamox 1g vial powder displacement volume ~ 0.8 mL.
Full day	Effectiveness:
Evidence	There are few studies of amoxicillin in the neonatal population to study effectiveness and the majority
	of the information is derived from studies of ampicillin. A study in two Estonian NICUs comparing
	ampicillin + gentamicin versus penicillin + gentamicin in the empiric therapy of neonates at risk of
	early-onset sepsis showed similar effectiveness in need to change antibiotics at 72 hours and/or 7-day

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all-cause mortality.(1) Subgroup analysis in ELBW neonates showed similar results, though NICU mortality was lower in the ampicillin group in < 26 weeks gestation neonates. (2)

In an RCT of amoxicillin prophylaxis for prevention of catheter-related infections in newborn infants with central venous catheters, bacterial contamination of the catheter tip at removal was significantly reduced in the amoxicillin group. No significant difference was found in the incidence of invasive infection.(3) In a randomised, open-label, equivalence trial in Africa, oral amoxicillin was found to be equivalent to injectable procaine penicillin plus gentamicin in the treatment of neonates and young infants with fast breathing.(4)

IV amoxicillin has similar properties to ampicillin and there is little to choose between the two when given by the IV route to treat susceptible organisms.(5) Amoxicillin achieves higher serum and CSF concentrations than ampicillin.(6) Oral amoxicillin has similar properties to ampicillin. Both the antibiotics are well absorbed when given by mouth, widely distributed in body tissues (including bronchial secretions) and rapidly excreted in the urine. Oral amoxicillin has better bioavailability but can be variable in young children.(5) Oral medication can nearly always be used to complete any sustained course of treatment.(12)

#### **Pharmacokinetics:**

Study of amoxicillin pharmacokinetics in preterm infants<sup>7</sup> has shown that a q12h schedule in the first week achieves serum concentrations well above the MIC for major micro-organisms in neonatal infections.(7) Another study in neonates older than 1 week showed that amoxicillin clearance was related to post-conceptional age and not to postnatal age with a rapid linear increase in clearance after 34 weeks post-conceptional age.(8)

In a study<sup>10</sup>, early switching to the oral route in asymptomatic full-term newborns with early onset GBS disease maintained serum amoxicillin concentrations within the therapeutic range.(10) The dose used in that study was 200–300 mg/kg/day in 4 divided doses. All the concentrations were in the therapeutic range with the lower dose. Another pharmacokinetic study in 6–13 days old neonates concluded that amoxicillin should be useful for oral treatment of neonatal infections caused by susceptible micro-organisms in infants who are not critically ill. The dose used was 50 mg/kg twice a day.(11)

#### Recommendation:

Amoxicillin can be used as a substitute for benzylpenicillin or ampicillin for suspected, early-onset, neonatal sepsis in combination with an aminoglycoside. When amoxicillin is used in combination with an aminoglycoside for the treatment of meningitis, it is recommended that the dose be doubled from 50 to 100 mg/kg/dose.(12) This is in keeping with similar recommendations for benzylpenicillin and ampicillin based on high minimum bactericidal concentration of group B streptococci and high inocula of the organisms in neonatal meningitis. (Level of evidence 5, Grade of recommendation D).

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