Amikacin

Newborn use only

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Aicit	Amikacin and gentamicin are both AMINOGLYCOSIDE antibiotics and MUST NOT be prescribed at the same time.					
	The Antimicrobial Stewardship Team has listed this drug under the following category:					
	Restricted.					
Indication	Treatment of suspected or proven gram-negative infection resistant to other aminoglycosic					
	Used in combination with a beta-lactam antibiotic for sepsis in the newborn.					
Action	Bactericidal agent that acts by inhibiting protein synthesis in susceptible bacteria.					
Drug Type	Aminoglycoside					
Trade Name	DBL Amikacin, Amikacin SXP, Amikacin Wockhardt.					
Presentation	500 mg/2 mL Excipients: Sodium citrate, sodium metabisulfite.					
Dosage/Interval	Postmenstrual age/corrected gestational age	Postnatal age	Dose	Interval		
	≤29 weeks	0–7 days	14 mg/kg	48-hourly		
		8–28 days	12 mg/kg	36-hourly		
		≥29 days	12 mg/kg	24-hourly		
	30–34 weeks	0–7 days	12 mg/kg	36-hourly		
		≥8 days	12 mg/kg	24-hourly		
	≥35 weeks	All	12 mg/kg	24-hourly		
	Infants with perinatal asphyxia	and on therapeuti	c hypothermia:	Increase dose interval by 12		
	hours [1-3].					
	Infants treated with cyclo-oxygenase inhibitors (indomethacin or ibuprofen): Increase dose					
	interval by 12 hours [1-3]					
Maximum daily dose	Intervariant inferior					
Route	Intravenous infusion					
Preparation/Dilution	Intramuscular injection tion Two-step dilution:					
r reparation, bilation	Step 1: Add 1 mL (250 mg) of amikacin to 9 mL of sodium chloride 0.9% to make a 25 mg/mL					
	solution. Step 2: FURTHER DILUTE 1 mL (25 mg) of this solution to 9 mL of sodium chloride 0.9%					
	2.5 mg/mL solution.					
Administration	IV infusion over 60 minutes using the proximal IV port.					
	IM: May be given if IV route not available.					
Monitoring	Routine therapeutic drug monitoring for ≤48 hours duration of therapy is not necessary unless					
_	renal function is impaired.					
	For infants on continuing treatment, perform early trough and peak levels (prior to and 1 hour					
	after the second amikacin dose). Target peak levels 24–35 mg/L and troughs <5 mg/L [2].					
	ssess renal function.					
Contraindications	Hypersensitivity to amikacin or other aminoglycosides.					
	Myasthenia Gravis ¹³					
Precautions	Treatment with amikacin for more than 14 days has not been established as being safe.					
	CAUTION in patients with pre-existing renal impairment, auditory or vestibular impairment,					
	hypocalcaemia, depressed neuromuscular transmission. Gastrointestinal: Amikacin has been associated with <i>Clostridium difficile</i> diarrhoea; discontinue					
	use if suspected.					
	Immunological: Allergic-type reactions, including anaphylaxis and life-threatening or less severe					
	asthmatic reactions, may occur in patients with sulfite sensitivity as preparation contains sodium					
	metabisulfite.					
	Neurological: Use caution in pati	ents with parkinso	nism; muscle we	akness may be aggravated.		
Drug Interactions	Diuretics may cause ototoxicity of					
J	concentrations.	- 0	- 1			

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	Neurotoxic and/or nephrotoxic agents: Avoid concurrent or sequential use of other neurotoxic and/or nephrotoxic antibiotics, including other aminoglycosides, polymyxin B, colistin, cisplatin, vancomycin, amphotericin B, clindamycin and cephalosporins. Anaesthetics/neuromuscular blocking agents or medications with neuromuscular blocking activity: succinylcholine, tubocurarine, decamethonium, halogenated hydrocarbon inhalation anaesthetics, opioid analgesics and massive transfusions with citrate anticoagulated blood may increase neuromuscular blockade. Treatment with anticholinesterase agents or calcium salts may help to reverse the blockade. Penicillins: Aminoglycosides are inactivated by solutions containing penicillins. Ensure line is adequately flushed between antibiotics.
Adverse Reactions	Serious reactions include neuromuscular blockade with subsequent respiratory paralysis,
Compatibility	ototoxicity and nephrotoxicity (see evidence review). Fluids: Glucose 5%, glucose 10%, glucose 20%, sodium chloride 0.9%, amino acid solutions.
Incompatibility	Aciclovir, amiodarone, atenolol, atracurium, atropine, aztreonam, buprenorphine, calcium chloride/gluconate, caspofungin, cefazolin, cefotaxime, cefoxitin, ceftazidime, ceftriaxone, chloramphenicol, cimetidine, clindamycin, dexamethasone, dexmedetomidine, digoxin, dobutamine, adrenaline (epinephrine), epoetin alfa, erythromycin, esmolol, fentanyl, filgrastim, fluconazole, foscarnet, furosemide (frusemide), gentamicin, isoprenaline, ketamine, labetalol, lidocaine (lignocaine), linezolid, magnesium sulfate, methadone, methylprednisolone, midazolam, milrinone, morphine, glyceryl trinitrate, noradrenaline (norepinephrine), octreotide, ondansetron, pancuronium, pethidine, phenobarbital (phenobarbitone), piperacillin, piperacillin-tazobactam, potassium chloride, procainamide, propranolol, protamine, pyridoxine, ranitidine, remifentanil, rocuronium, sodium acetate, sodium bicarbonate, succinylcholine, vancomycin, vasopressin, vecuronium, warfarin, zidovudine
Stability	Penicillins and cephalosporins, amphotericin, azathioprine, azithromycin, diazepam, diazoxide, folic acid, ganciclovir, heparin, hydralazine, ibuprofen, indomethacin, insulin, pentamidine, pentobarbital (pentobarbitone), phenytoin, potassium chloride, propofol, sulfamethoxazoletrimethoprim, teicoplanin Administer immediately, discard unused portion.
Stability	The diluted solution is stable for 24-hours at room temperature.
Storage	Store below 25°C.
Special Comments	
Evidence summary	Refer to full version.
References	Refer to full version.

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