

AMIODARONE **This drug must be guardrailed**

Trade Name	IV: Amiodarone hydrochloride (Hameln) Oral: Amiodarone Liquid (prepared by pharmacy)			
Class	Antiarrhythmic Agent Class III			
Mechanism of Action	Amiodarone inhibits adrenergic stimulation of the myocardium, prolonging the action potential and refractory period. AV node conduction and sinus node function are also decreased.			
Indications	Management of life threatening resistant arrhythmias – ventricular and supraventricular			
Contraindications	Sinus node dysfunction/block; 2 nd or 3 rd degree A/V block; cardiogenic shock; severe hypotension, severe respiratory failure. Clearance of amiodarone is reduced in patients with impaired cardiac or hepatic function. Hypersensitivity to amiodarone. Precautions with hepatic, respiratory, thyroid disorders, electrolyte imbalance, hypotension, LV dysfunction, bradycardia, heart failure.			
Supplied As	IV: 50 mg/mL (150 mg/3 mL ampoule) Oral: 10 mg/mL			
Dilution	Drug	5% Dextrose Added	Final Volume	Concentration
	1mL (50mg)	24mL	25mL	2 mg/mL
	Dextrose 5% is the only compatible diluent See infusion sheet for charting and dosing rates Concentrations >2 mg/mL need central access Concentrations < 0.6 mg/mL are unstable and should not be used.			
Dosage *Must chart guardrail and use Alaris pump*	IV: Infuse 25 microgram/kg/min for 4 hours (gives a load of 6mg/kg) then 5-15 microgram/kg/min titrate to response (max 1.2g/24hr) Oral: 4mg/kg/dose 8 hourly for 1 week, then 4mg/kg/dose 12 hourly for 1 week, then, 4mg/kg/dose daily Consult cardiology for oral dose frequency when converting from iv to oral dosing			
Guardrails	Concentration: 2mg/mL Soft Min: 5 microgram/kg/min Hard Max: 30 microgram/kg/min Soft Max: 25 microgram/kg/min Default: 10 microgram/kg/min			
Interval	IV: Continuous infusion. Oral: Initially 12 hourly and then 24 hourly			
Administration	IV: Give via a central line to reduce risk of thrombophlebitis Oral: Give with food			

Compatible With	<p>Solution: 5% Dextrose (this is the only compatible fluid)</p> <p>Y-site :</p> <p>Adrenaline, alprostadil, amikacin, amphotericin (liposomal), atropine, benzylpenicillin, calcium chloride, calcium gluconate*, ciprofloxacin, cefuroxime*, dexmedetomidine, dopamine, dobutamine*, ephedrine, erythromycin, fluconazole, gentamicin, insulin*, lidocaine, metronidazole, midazolam, milrinone, morphine, naloxone, noradrenaline*, ondansetron, pancuronium, phenylephrine, potassium chloride*, smof lipid, tobramycin, TPN*, vancomycin, vasopressin, zidovudine.</p> <p>* Variable compatibility results with calcium gluconate, cefuroxime, dobutamine, insulin, noradrenaline, potassium chloride and TPN solutions, use a separate line if possible</p>
Incompatible With	<p>Do not mix with sodium chloride 0.9%, acyclovir, aminophylline, amoxicillin+clavulanate, azithromycin, calcium gluconate, cefazolin, cefotaxime, ceftazidime, dexamethasone, digoxin, flucloxacillin, furosemide, ganciclovir, heparin, hydrocortisone sodium succinate, imipenem+cilastin, meropenem, phenobarbital, phenytoin, piperacillin+tazobactam, potassium phosphates, ranitidine, sodium bicarbonate, sodium phosphates, sulfamethoxazole+trimethoprim</p>
Interactions	<p>Take care when administering with other medicines known to prolong QT interval eg: other antiarrhythmics, sotalol, erythromycin, sildenafil, domperidone, fluconazole, ciprofloxacin.</p> <p>Digoxin dose will need to be halved if given concurrently with amiodarone - amiodarone prevents elimination of digoxin.</p> <p>Flecainide will need a dose reduction of up to 50%.</p> <p>Clarithromycin, Erythromycin, Fluconazole and Phenytoin levels may increase and amiodarone levels may decrease when these agents are used concurrently.</p> <p>Amiodarone use may increase or decrease cyclosporin levels, with increased risk of nephrotoxicity. Increased monitoring is required.</p> <p>Diuretics, corticosteroids, amphotericin – risk of hypokalaemia, increasing the risk of torsades de pointes.</p> <p>Beta blockers and Ca channel blockers – increased risk of bradycardia, hypotension, AV block, myocardial depression</p> <p>Midazolam – amiodarone increases exposure.</p>
Monitoring	<p>Continuous ECG and BP, cardiorespiratory monitoring while on IV amiodarone</p> <p>Liver function tests including AST and ALT, thyroid function tests</p> <p>INR and electrolytes specifically Mg and K</p>
Stability	<p>IV: Amiodarone solutions are stable in 5% dextrose for up to 24 hours <u>only</u> if stored in glass or rigid PVC and polypropylene containers. We do not have these so IV amiodarone solutions should be discarded immediately after use and prolonged infusions should be avoided if possible (ie: >4-6 hours)</p> <p>Oral: 7 day expiry, shake well before use.</p>

Storage	IV: Below 25°C ; Do not refrigerate; Protect from light Oral: Store in the fridge (2 – 8 °C)												
Adverse Reactions	Bradycardia, hypotension, (possibly associated with rapid infusion rates), polymorphic ventricular tachycardia. Thrombophlebitis with prolonged IV use Skin discolouration (slate blue), photosensitivity, rash, Hypothyroidism, hyperthyroidism, hyperglycaemia, Hepatic toxicity, coagulation abnormalities Nausea, vomiting, constipation Optic neuritis, pulmonary fibrosis Contains benzyl alcohol –potential risk of Neonatal Gasping Syndrome												
Metabolism	Bioavailability ~50%, extensively metabolised in the liver by CYP3A4, active metabolite = N-desethylamiodarone, eliminated primarily by biliary excretion.												
Comments	Amiodarone contains 37.3% iodine - monitor thyroid function Protect skin from excess sunlight Each 1 ml of amiodarone injection contains 22.2mg of benzyl alcohol												
References	<ol style="list-style-type: none"> www.adhb.govt.nz/newborn/services/DrugProtocols NZHPA Notes on Injectable Drugs 5th Edition 2004 & www.noids.nz Lacy et al Drug Information Handbook 10th Edition 2003. Northern Network Formulary 11th Edition 2000 Neonatal Formulary 7th Edition Hammersmith Trust 2000 Trissell Hanbook of Injectable Drugs 10th Edition www.micromedexsolutions.com Medicines for Children RCPCH 1999. 												
Updated By	<table> <tr> <td>P Schmidt, B Robertshawe</td> <td>September 2006</td> </tr> <tr> <td>B Robertshawe, A Lynn</td> <td>July 2010</td> </tr> <tr> <td>B Robertshawe, A Lynn</td> <td>June 2012 (re-order profile)</td> </tr> <tr> <td>B Robertshawe, A Lynn</td> <td>Oct 2018 (dose alignment with Akld)</td> </tr> <tr> <td>A Lynn, M Wallenstein, B Robertshawe, A Evison</td> <td>May 2020 (review and update)</td> </tr> <tr> <td>A Lynn, B Robertshawe</td> <td>May 2023 (routine review)</td> </tr> </table>	P Schmidt, B Robertshawe	September 2006	B Robertshawe, A Lynn	July 2010	B Robertshawe, A Lynn	June 2012 (re-order profile)	B Robertshawe, A Lynn	Oct 2018 (dose alignment with Akld)	A Lynn, M Wallenstein, B Robertshawe, A Evison	May 2020 (review and update)	A Lynn, B Robertshawe	May 2023 (routine review)
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