West of Scotland NEONATAL PARENTERAL Drug Monographs

Amiodarone

FORM Ampoules containing 150mg/3ml Concentrate solution for infusion

INDICATION Ventricular Arrhythmias

DOSE RANGE

AGE	DOSE	FREQUENCY	ROUTE
	5mg/kg over 4 hours	Loading dose	IV Infusion
1 to 6 months	Initially 5micrograms/kg/minute	Continuous	IV
	Increase according to response to a maximum dose of 15micrograms/kg/minute	Infusion	-

PRESCRIPTION OF CONTINUOUS INFUSION

SINGLE STRENGTH = 50mg/kg in 50ml Glucose 5%

This gives:-

• 5 microgram/kg/minute at 0.3ml/hour

DOUBLE STRENGTH (for babies weighing < 600grams)

= 100mg/kg in 50ml infusion fluid

This gives:-

• 10 microgram/kg/minute at 0.3ml/hour

(For stability purposes, a double strength infusion should be used for babies weighing less than 600grams)

RECONSTITUTION Not required

DILUTION For continuous Infusion

SINGLE STRENGTH
Using 50mg/ml injection

1 x wt (kg) is the number of ml of amiodarone 50mg/ml to be diluted up to 50ml total with infusion fluid (equivalent to **50mg/kg in 50ml**)

DOUBLE STRENGTHUsing 50mg/ml injection

2 x wt (kg) is the number of ml of amiodarone 50mg/ml to be diluted up to 50ml total with infusion fluid (equivalent to **100mg/kg in 50ml**)

METHOD OF ADMINISTRATION

Loading Dose

5mg/kg can be administered as a separate infusion over 4 hours ${\bf OR}$

given from the infusion as prepared above

SINGLE strength - Give 5ml (5mg/kg) over 4 hours DOUBLE strength - Give 2.5ml (5mg/kg) over 4 hours

For continuous Infusion

By continuous intravenous infusion, flow rate adjusted according to the baby's response (see prescription section for details).

Administration via a **central venous catheter** is recommended.

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COMPATIBILITY

Solution compatibility	glucose 5% ONLY
Solution incompatibility	sodium Chloride
IV Line compatibility	adrenaline, clarithromycin, dopamine, fluconazole, gentamicin, hydralazine, labetalol, linezolid, metronidazole midazolam, milrinone, , morphine, , vancomycin, vecuronium, Variable compatibility (dobutamine, insulin, magnesium sulphate, potassium chloride, sodium nitroprusside) Contact pharmacy for advice
IV Line incompatibility	Imipenem, furosemide, heparin, meropenam, phenobarbital, phenytoin, sodium bicarbonate

THIS LIST IS NOT EXHAUSTIVE PLEASE CONTACT PHARMACY FOR FURTHER INFORMATION ON COMPATIBILITY WITH ANY MEDICINES NOT INCLUDED

CAUTIONS, CONTRA-INDICATIONS AND SIDE EFFECTS

See Summary of Product Characteristics and most recent edition of BNF for Children (links below)

FURTHER INFORMATION

Monitoring requirements

- Liver function and thyroid-function tests required before treatment and then every 6 months.
- Serum potassium (risk of hypokalaemia)
- Blood pressure (circulatory collapse precipitated by rapid administration or over dosage).

Other Information

- This product may contain benzyl alcohol. Benzyl alcohol has been reported to be associated with a fatal "Gasping syndrome" in premature infants.
- Can cause photosensitivity (caution with phototherapy) and persistent slate-grey skin discolouration.

PH 3.5 - 4.5 (undiluted)

LICENSED STATUS Not licensed for use in children

LINKS BNF for Children / Electronic Medicines Compendium:

APPLICABLE POLICIES West of Scotland Neonatal Guidelines:

Consult local policy if applicable

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Prepared by:	WoS Neo Pharm Group	Checked by	Hazel Fisher		
Date prepared	May 2016	Date updated	September 2019		
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Administer reconstituted solutions immediately.

All vials, ampoules and infusion bags are for single use only unless otherwise stated.

Dose may vary depending on indication, age, renal function, hepatic function, and concomitant medications. This monograph should be used in conjunction with the package insert, BNF for Children, and Summary of Product Characteristics. For further advice contact your clinical pharmacist or pharmacy department.

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