NEONATAL West of Scotland Drug Monographs PARENTERAL Routes

Dinoprostone (Prostin) FIXED CONCENTRATION

Take special care with units, calculating doses and administration volumes Frequently involved in medication errors, care with similar named drugs.

FORM Pre-made syringe containing 50micrograms in 50ml

OR

Ampoules containing 1mg/ml dinoprostone (if no pre-made syringe is

available or preparing a high concentration syringe)

INDICATION Maintaining patency of the Ductus Arteriosus

SEEK CONSULTANT ADVICE BEFORE PRESCRIBING. WHERE POSSIBLE, DO NOT START OR STOP THIS TREATMENT WITHOUT

FIRST CONSULTING A CARDIOLOGIST.

DOSE RANGE

AGE	DOSE	FREQUENCY	ROUTE
	Initially 5nanograms/kg/MINUTE		
0-6 months	Gradually increase as necessary in 5nanogram/kg/minute increments to 20nanograms/kg/minute.	Continuous Infusion	IV
	Doses of up to 100 nanogram/kg/minute have been used but are associated with an increased incidence of side effects.		

PRESCRIPTION OF CONTINUOUS INFUSION

SPECIAL CARE with PRESCRIBING.

Flow rates are expressed as ml/kg/hour NOT ml/hr

LOW CONCENTRATION (if doses less than

20nanograms/kg/min are required)

50 micrograms in 50ml infusion fluid

This gives:-

- 5nanogram/kg/MINUTE at 0.3ml/**KG**/hr
- 10nanogram/kg/MINUTE at 0.6ml/KG/hr
- 20nanograms/kg/MINUTE at 1.2ml/KG/hr

<u>HIGH CONCENTRATION</u> (if doses greater than or equal to 20nanograms/kg/min are required)

200micrograms in 50ml infusion fluid

This gives:-

- 20nanograms/kg/MINUTE at 0.3ml/KG/hr
- 50nanograms/kg/MINUTE at 0.75ml/KG/hr

Meomaral

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IF NO PRE-MADE SYRINGE IS AVAILABLE PREPARE A SYRINGE OF either 50micrograms in 50ml (low concentration) or 200micrograms in 50ml (high concentration) USING A 1mg/ml AMPOULE AS

FOLLOWS;

RECONSTITUTION

Already in solution

DILUTION

Initial Dilution for both concentrations:

Dinoprostone Injection (1mg/ml)	0.5ml	
Water for Injection.	up to 20ml total	

This gives a 25microgram in 1ml solution.

Final Dilution

LOW CONCENTRATION

- 1. Draw up 2ml of DILUTED DINOPROSTONE 25micrograms in 1 ml solution.
- 2. Make up to 50ml with Glucose 5% or Sodium Chloride 0.9%.
- 3. Mix well
- 4. This gives a solution containing **50 micrograms in 50ml** (1microgram per 1ml)

HIGH CONCENTRATION

- Draw up 8ml of DILUTED DINOPROSTONE 25micrograms in 1 ml solution.
- 2. Make up to 50ml with Glucose 5% or Sodium Chloride 0.9%.
- 3. Mix well
- 4. This gives a solution containing **200 micrograms in 50ml** (4micrograms per 1ml)

METHOD OF ADMINISTRATION

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

WARNING: DO NOT FLUSH THROUGH A LINE CONTAINING DINOPROSTONE AS IT IS A POTENT DRUG AND MAY CAUSE SERIOUS PROBLEMS.

COMPATIBILITY

Solution compatibility	Sodium chloride 0.9%, Glucose 5%.	
Solution incompatibility	No information	
IV Line compatibility	No other drugs or fluids in same line at same time	
IV Line incompatibility	All other drugs	

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

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FURTHER INFORMATION

During the infusion of dinoprostone, monitoring of heart rate, blood pressure, respiratory rate and core body temperature are required. Respiratory depression and apnoea can also occur.

In the event of complications such as apnoea, profound bradycardia or severe hypotension consult with senior medical staff.

IV Dinoprostone can cause respiratory depression and apnoea. Facilities

for intubation and ventilation must be available and should be considered for babies being transferred to another site with a

Dinoprostone infusion.

STORAGE Dinoprostone is stored in a refrigerator. Once diluted the injection

solution is stable for 24 hours at room temperature.

PH No information

LICENSED STATUS Not licensed for use in children.

APPLICABLE POLICIES West of Scotland Neonatal Guidelines

Consult local policy if applicable

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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 terms of reference document prepared by the West of Scotland Pharmacist Network. Information is correct at the time of publication and as per local practice agreement. For further advice please contact your clinical pharmacist or pharmacy department