ALPROSTADIL (prostaglandin E₁)

	Indication	 Temporarily maintain patency of ductus arteriosus¹ Use with ductal dependent congenital heart defects¹ Can be considered for reducing right ventricular afterload in persistent pulmonary hypertension^{2,3} 					
	Presentation	Ampoule: 500 microgram in 1 mL (500,000 nanogram in 1 mL)					
	Dosage (initial)	 0.01–0.02 microgram/kg/minute (10–20 nanogram/kg/minute) Titrate according to response in consultation with paediatric cardiologist 					
SU	Dosage (maintenance)	 Usual range 0.01–0.02 microgram/kg/minute (10–20 nanogram/kg/minute) Can be given at range of 0.005–0.1 microgram/kg/minute (5–100 nanogram/kg/minute)⁴ 					
INTRAVENOUS	Preparation	 Minimise contact time of undiluted alprostadil with the plastic syringe Refer to stability Draw up 9 mL of 0.9% sodium chloride Draw up 500 microgram (1 mL) alprostadil and immediately add to the 9 mL 0.9% sodium chloride (10 mL total volume) <i>Concentration now equal to 50 microgram/mL</i> From the 50 microgram/mL solution, draw up 18 microgram/kg and make up to 30 mL total volume with 0.9% sodium chloride <i>Concentration now equal to 18 microgram/kg in 30 mL</i> 					
	Administration	 Infuse using a medication infusion pump at prescribed rate 18 microgram/kg in 30 mL infusion at 1 mL/hour is equivalent to 0.01 microgram/kg/minute (10 nanogram/kg/minute) 					
 Special considerations Consult with paediatric cardiologist first Apnoea is common and intubation may be required Caution If bleeding disorders¹ If respiratory distress syndrome¹ 		 Apnoea is common and intubation may be required Caution If bleeding disorders¹ 					
	Monitoring	 Continuous cardio-respiratory monitoring (apnoea frequent side-effect)¹ BP¹, temperature, pulse oximetry Improvement in blood oxygenation, systemic BP and blood pH demonstrates efficacy¹ Extravasation risk: can cause tissue sloughing and necrosis⁵ 					
	Compatibility • Fluids • 5% glucose ⁵ , 0.9% sodium chloride ⁵ • Y-site • Do not give other drugs via same line Incompatibility • No information ⁵						
	Interactions	 Concurrent use with heparin may result in increased risk of bleeding⁶ 					
	 Undiluted solution Store in refrigerator at 2–8 °C1 If undiluted alprostadil comes into contact with plastic, the solution may turn hazy and must then be discarded1 Diluted solution Stable for 24 hours at 25 °C, then discard1 						



	 Blood pathology: disseminated intravascular coagulation (DIC)¹ Circulatory: hypotension¹, bradycardia¹, tachycardia¹, cardiac arrest¹, oedema¹ 	
	 Digestive: diarrhoea¹, gastric outlet obstruction secondary to antral hyperplasia¹ (prolonged treatment) 	
Side effects	 Musculo-skeletal: widened fontanels, pretibial and soft tissue swelling (associated with prolonged duration)¹ 	
	• Nervous: cutaneous flushing (related to infusion rate) ¹ , fever ¹ , urticaria ⁷ , seizures ⁷	
	 Respiratory: apnoea appearing during first hour of infusion (more common in babies weighing less than 2 kg)¹ 	
	 Causes vasodilation of all arterioles (i.e. ductus arteriosus as well as ductal tissue surrounding the duct)⁷ 	
Actions	 Inhibits platelet aggregation⁷ 	
	• Short half-life of 5–10 minutes necessitates infusion rather than bolus administration ¹	
	 Maximum effect observed within 96 hours after birth¹ 	
Abbreviations	BP: blood pressure, DIC: disseminated intravascular coagulation	
Keywords	Prostaglandin E1, Prostin VR, alprostadil, PDA, patent ductus arteriosus, duct dependent congenital heart defect, PGE1	

The Queensland Clinical Guideline *Neonatal Medicines* is integral to and should be read in conjunction with this monograph. Refer to the disclaimer. Destroy all printed copies of this monograph after use.

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