

# ALPROSTADIL (Prostaglandin E1) **Drug must be guardrailed**

|   |   |
|---|---|
| Trade Name  | Prostin VR (Pfizer)   |
| Class   | Prostaglandin.  |
| Mechanism of Action   | Direct vasodilation effect on all vascular smooth muscle – used for ductal effect   |
| Indications   | Temporary management of ductus arteriosus patency in duct-dependent congenital heart disease.<br>(Transposition of the great arteries, all right sided congenital heart defects associated with reduced pulmonary perfusion, left sided defects including hypoplastic left heart, coarctation of aorta and interrupted aortic arch.)  |
| Precautions   | Use with caution in:<br>Respiratory distress (can cause apnoeas)<br>Obstructed TAPVD (total anomalous venous drainage)<br>Infants with bleeding tendency (PGE1 inhibits platelets)<br>Seizure disorders   |
| Supplied As   | 500 microgram/mL 1mL ampoules   |
| Dilution  | <b>See Alprostadil Infusion sheet</b><br>Take 60 microgram/kg and make up to 50mL with 0.9% saline / 5% dextrose.<br><b>0.5mL/hr = 10 nanogram/kg/min</b><br>The final concentration of the infusion will vary depending on the weight of the baby but in most occasions the <b>concentration will be 3 – 6 microgram/mL</b><br>In rare situations, when a large dose is required and the volume infused needs to be restricted, the strength can be made up to a maximum of <b>20 microgram/mL</b> . This needs to be discussed with the Pharmacist as the solution is hyperosmolar. |
| Dosage<br><br><b>*Must chart guardrail and use Alaris pump*</b> | <b>Maintenance:</b> 10 – 100 nanogram/kg/min<br>(0.01-0.1 microgram/kg/min)<br>Start at <b>10 nanogram/kg/min</b> and increase by 10 nanogram/kg/min increments every 30 minutes.<br>Side effects are dose dependent.   |
| Guardrails<br><b>ALARIS PUMP</b>                                | Conc: Min – 0.36 microgram/mL Max – 20 microgram/mL<br>Soft Min: 5 nanogram/kg/min Hard Max: 100 nanogram/kg/min<br>Soft Max: 50 nanogram/kg/min Default: 10 nanogram/kg/min  |
| Guardrails<br><b>TRANSPORT PUMP</b>                             | Conc: Min – 0.36 microgram/mL Max – 20 microgram/mL<br>Soft Min: 0.005 microgram/kg/min Hard Max: 0.1 microgram/kg/min<br>Soft Max: 0.05 microgram/kg/min Default: 0.01 microgram/kg/min  |
| Interval  | Continuous infusion   |

|                          |   |
|--------------------------|---|
| <b>Administration</b>    | IV infusion via syringe pump.<br>Need separate IV access from maintenance fluids  |
| <b>Compatible With</b>   | <p><b>Solution:</b><br/>0.9% sodium chloride, 5% dextrose.</p> <p><b>Terminal Y-site:</b> adrenaline, amino acid solutions, atropine, benzylpenicillin, caffeine citrate, calcium gluconate, cefazolin, cefotaxime, chlorothiazide, digoxin, dobutamine, dopamine, fentanyl, flecainide, furosemide, gentamicin, glycopyrrolate, heparin, insulin, ketamine, methylprednisolone, midazolam, milrinone*, morphine, pancuronium, potassium chloride, ranitidine, sildenafil, tobramycin, vancomycin, vecuronium</p> <p>*milrinone only compatible at conc 0.5 mg/mL in 5% glucose</p> |
| <b>Incompatible With</b> | Inconclusive data available on compatibility with, 10% dextrose, SMOFlipid, noradrenaline, recommend to avoid infusing in same line as alprostadil  |
| <b>Monitoring</b>        | <p>Full cardiorespiratory monitoring required:<br/>Improved oxygenation and PaO<sub>2</sub> suggests duct has reopened and infusion rate may need to be decreased.<br/>BP, Temperature.<br/>IV access supervision.</p>  |
| <b>Stability</b>         | <p>Discard opened vial immediately after use<sup>0</sup><br/>Use a new vial for each dose.<br/>Continuous infusions need to be changed after 24 hours</p>   |
| <b>Storage</b>           | Below 8°C (but do not freeze). Discard any unused portion of the ampoule immediately after use.   |
| <b>Adverse Reactions</b> | <p>Apnoea, fever, bradycardia, flushing, seizures, hypotension, tachycardia, diarrhoea, bronchospasm, hypocalcemia.</p> <p>Less commonly: hyperkalemia, anemia, hypoglycemia, irritability/jitteriness, oedema, cardiac arrest, DIC, gastric outlet obstruction by stimulation of GIT smooth muscle (especially if PGE1 is infused for &gt;120 hours).</p> <p>Tissue extravasation may cause necrosis- central venous access recommended.</p> <p>Cortical peri-osteal reaction after prolonged (days) treatment (resolves 6-12 months after stopping treatment).</p>                |
| <b>Metabolism</b>        | Local: most tissues.  |
| <b>Comments</b>          | <p>Max drug effect usually seen in 30mins if cyanotic lesion; duration of effect short, so secure or reserve IV access vital, especially prior to transport.</p> <p>Most effective if given in first 4 days after birth.</p>  |

|                   |  |
|-------------------|--|
| <b>References</b> | <ol style="list-style-type: none"> <li>1. Young T.E. et al. Neofax 2000; 108-9.</li> <li>2. Neonatal Pharmacopoeia (1<sup>st</sup> edition 1998), RWH, Melb.</li> <li>3. Taketomo C. Paediatric Dosage Handbook 6<sup>th</sup> Edition.</li> <li>4. <a href="http://www.medsafe.govt.nz">www.medsafe.govt.nz</a></li> <li>5. <a href="http://www.adhb.govt.nz/newborn/DrugProtocols/">www.adhb.govt.nz/newborn/DrugProtocols/</a></li> <li>6. <a href="http://www.micromedexsolutions.com">www.micromedexsolutions.com</a></li> <li>7. <a href="http://www.anmfonline.org">www.anmfonline.org</a></li> </ol> |
| <b>Updated By</b> | <p>Jan Klimek Nov 2000.P Schmidt, B Robertshawe Aug 2005.<br/> A Lynn, B Robertshawe April 2009, July 2009, Sept. 2009<br/> A Lynn, B Robertshawe Nov 2012 (re-order profile, discard vial) Tx guardrail<br/> A Lynn, B Robertshawe Jan 2022 (update compatibility section, rename as alprostadil)<br/> A Lynn, B Robertshawe Jan 2025 (routine review, update compatibility)</p>  |