

# Southampton Oxford Neonatal Transport Guideline

## DUCT DEPENDENT CONGENITAL HEART DISEASE

### Additional equipment:

- Prostin: PGE2 Dinoprostone
- **DO NOT use other prostaglandins – only Dinoprostone**
- Nitric oxide
- i-STAT machine (if available)

- **TIME CRITICAL/ Immediate Dispatch\*\*if baby is unstable and not responding to appropriate management**
- **DO NOT spend a long time trying to stabilise patient. You may have to scoop and run. Discuss regularly with MDT team.**

Duct dependent – systemic circulation	Duct dependent pulmonary circulation	Differential diagnoses
<ul style="list-style-type: none"> <li>• Co-arcuation of the aorta</li> <li>• Critical aortic stenosis</li> <li>• Hypoplastic left heart syndrome</li> </ul>	<ul style="list-style-type: none"> <li>• Pulmonary atresia</li> <li>• Critical pulmonary stenosis</li> <li>• Tricuspid atresia</li> </ul>	<ul style="list-style-type: none"> <li>• Pulmonary hypertension (C18)</li> <li>• Sepsis (C20)</li> <li>• Metabolic disorders (C19)</li> <li>• Primary lung pathology</li> <li>• Obstructed TAPVD – CXR plethoric</li> </ul>
<b>Duct dependent systemic and pulmonary circulations</b>	<ul style="list-style-type: none"> <li>• Transposition of the great vessels with restrictive circulation</li> </ul>	
Aim for: palpable pulses, resolving acidosis	Aim for: sats 75-85%, lactate < 2 mmol/L	Aim for: improved oxygen, BP, acidosis

### Initial assessment

(NB: may be no murmur)

- Antenatal and family history: antenatal scans, if antenatal Δ ?plan for care after birth
- History of labour and delivery: risk factors for sepsis, condition at birth, resuscitation.
- Time course of presentation and predominant symptoms / signs (cyanosis, shock, acidosis, ↑WOB)
- Management to date and rationale for therapeutic decisions
- **ABC:** Current clinical parameters and progression: RR, WOB, HR, MBP, perfusion, saturations, ↑liver size
- **ABCDEF:** acidosis, lactate, BS, electrolytes, CRP, temperature, what have parents been told?
- Assess need for further urgent intervention (see below)
- Discuss findings and provisional plans with supervising neonatal consultant and paediatric cardiologist

### Respiratory support

(In stable babies with prostin of 10 nanogram/kg/min or below, the risk of apnoea is low and they do not routinely need intubation for transfer)

- If apnoea, or if symptoms of respiratory or cardiac failure: intubate and ventilate.
- Add oxygen, if required, to achieve saturations of 75-85% (to avoid pulmonary over-circulation)
- If ventilated, give analgesia with morphine, +/- muscle relaxation
- Monitor pre and post ductal saturations. Aim: PaO<sub>2</sub> 5 kPa, PaCO<sub>2</sub> 5 kPa

### Cardiovascular

- Establish secure access. Two peripheral cannulae minimum. **Do not delay transfer for central access.**
- Treat hypotension with 10 ml/kg fluid boluses and reassess.

### Commence Prostin (make up carefully to avoid errors)

- If antenatal diagnosis and baby well and non-acidotic – start at 10 nanogram/kg/min
  - If presentation of absent femoral pulses but well and non-acidotic - start at 20 nanogram/kg/min
  - If acidotic, unwell infant (late presentation) with suspected CHD – 50 nanogram/kg/min
- Double the dose every 20 minutes if no improvement** (maximum of 100 nanograms/kg/min- discuss with SONeT consultant or cardiologist if considering >50nanograms/kg/min)

**Prostin can be given peripherally or centrally**

### Other considerations

- Blood cultures, antibiotics. (Remember that sepsis can co-exist with antenatally diagnosed CHD)
- Check blood sugars regularly. Correct hypocalcaemia and hypomagnesaemia.
- Consider causes of acidosis and optimise.
- If PPHN suspected: start inhaled nitric oxide.