

Clinical Guidance

Paediatric Critical Care: Total Anomalous Pulmonary Venous Drainage (TAPVD)

Summary

This is for staff to use to provide guidance regarding the diagnosis, management and perioperative care of TAPVD.

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Relevant external law, regulation, standards	
<p>This clinical guideline has been produced by the South Thames Retrieval Service (STRS) at Evelina London for nurses, doctors and ambulance staff to refer to in the emergency care of critically ill children.</p> <p>This guideline represents the views of STRS and was produced after careful consideration of available evidence in conjunction with clinical expertise and experience. The guidance does not override the individual responsibility of healthcare professionals to make decisions appropriate to the circumstances of the individual patient.</p>	

Glossary:

MAS = meconium aspiration syndrome
 PPHN = persistent pulmonary hypertension of the newborn
 PEEP = positive end expiratory pressure
 CXR = chest x-ray
 CVS = cardiovascular system
 RAI = right atrial isomerism

Change History		
Date	Change details, since approval	Approved by
03/23	Description expanded to reflect RBH guideline. Guidance agreed with RBH PICU team. Cardiac CT added.	RBH, PGC

Presentation:

Obstructed TAPVD may present in 1st hours of life

- Frequently undiagnosed lesion on antenatal scans
 - Baby may be compromised in utero, pass meconium during labour and be misdiagnosed as MAS/PPHN
 - Profound cyanosis, tachypnoea, tachycardia, poor volume pulses, hepatomegaly
 - Murmur not usually present
 - Early recognition/ consideration essential
- Unobstructed- symptoms less severe
- Present first weeks/ months of life
 - Failure to thrive, tachypnoea, recurrent chest infections, mild cyanosis, hepatomegaly.

Emergency management:

Obstruction is surgical emergency: urgent cardiac referral & transfer to surgical centre

- Intravenous/ umbilical/ intraosseous access
- Intubate & ventilate for hypoxia & cardiac support
 - Sats often do not improve
- May require inotropes to support blood pressure
- Cautious aliquots of anaesthetic induction agents
- Peak pressure to achieve good chest movement. PEEP 5-10cm H₂O.
- Dinoprostone (Prostaglandin E₂): if no improvement, do not escalate >10 nanogram/kg/min
- Standard sedation with Morphine
- Unobstructed: Rarely clinical emergency. May need ventilation, CVS support & diuretics.

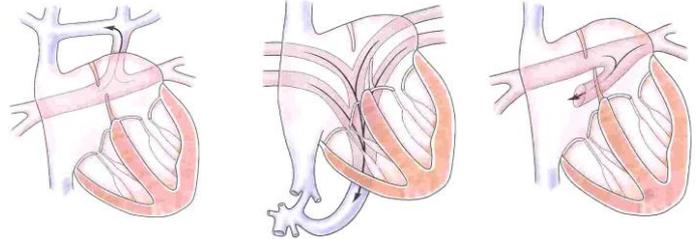
Special investigations

- **CXR:** heart size = small to normal if obstructed as left ventricle empty
SVC widening in supracardiac TAPVD; "snowman". Diffuse pulmonary oedema. Interstitial oedema identified by diffuse reticular pattern- "Snowstorm". Pleural effusion/s may be present.
- **ECG:** RV enlargement /R axis deviation. Tall, peaked P waves (not specific)
- ↑pO₂ in UVC gas diagnostic of infracardiac TAPVD
- Echo +/- Cardiac CT to clarify anatomy

Description:

Pulmonary veins (PV) do not return to L atrium: blood drains into R side of heart via systemic veins and then back to the L side via ASD/ VSD. If blood flow becomes obstructed the L side is underfilled and shock develops. Cyanosis is worsened by pulmonary oedema.

Classification



Supracardiac 49%
PV connect to Superior vena cava or innominate

Infracardiac 26%
PV connect to hepatic or inferior vena cava

Cardiac 16%
PV to coronary sinus to right atrium

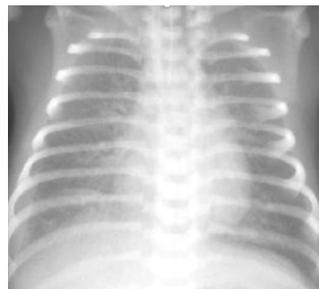
Mixed 9%: Combination of above

Obstruction seen in 75% infra-cardiac & in 50% supra-cardiac lesions
Obstruction develops or worsens over time

1/3 patients have no other cardiac anatomical defect

Frequently occurs with right atrial isomerism with asplenia, & heterotaxy

- 89% RAI associated with TAPVD



Obstructed infracardiac TAPVD
Small heart shadow, interstitial and alveolar oedema (plethoric, "snowstorm") with pleural effusion
R > L



Supracardiac TAPVD
Dominant right sided cardiac shadow. Enlarged mediastinum due to supracardiac draining vein & enlarged SVC. "Snowman". If obstructed, "snowman in snowstorm". Plethoric lung fields.

Peri-Operative management

Obstructed: (Low cardiac output state/shock)

- Needs urgent anatomical diagnosis-echo prior to emergency surgery. Urgent cross match.
- Proceed to definitive surgery. (ECMO considered on case by case discussion)
- **High risk surgical categories** <2.5kg, Evidence of in-utero pulmonary vein obstruction with restricted ASD (pleural effusions, ascites); Small left atrium, neonatal presentation with end organ dysfunction & profound cyanosis, hypoplastic pulmonary veins

Unobstructed:

- Cardiac failure management
- Accurate delineation of pulmonary veins & abdominal anatomy, cardiac CT increasingly performed to confirm anatomy.
- Delayed surgical intervention with surveillance for obstruction

Post op: Routine post cardiac surgical care in PICU

- Risk of pulmonary hypertensive crises ~50% so atrial communication should be left. Signs include ↓ETCO₂, desaturation, hypotension & ↑CVP. Causes RV failure. Prevent & treat with high FiO₂ (50-70%), minimal handling, sedation +/- muscle relaxation. Prime nitric oxide into ventilator circuit before return from theatre. Short term sildenafil may be required during NO wean. Potential for atrial arrhythmia due to surgical atriotomy
- High risk of organ dysfunction if obstructed- renal (dialysis), liver (glucose & coagulation) and delayed feeding (TPN)

Outcome/ long term prognosis

- Influenced by early recognition, timely intervention & severity of pulmonary hypertension post op.
- Long term morbidity due to progressive or recurrent pulmonary vein obstruction¹. Recurrence risk 15-20%
 - Infracardiac & mixed at greatest risk. Incidence at anastomotic site may be reduced by surgical 'sutureless' technique.
 - Vessel hypoplasia hard to diagnose pre op. Intrinsic lack of venous growth remains challenging with 3 year survival ~60%²