





## Adjudication for Paediatric Super-Urgent Listing

Additional questions were asked of the Working Group at the request the Director of CTAG(H) to address the point within POL 229/12: “For paediatric requests for urgent or super-urgent listing under categories 59 or 12 whereby a maximum acceptable donor size has been specified to be  $\geq 160\text{cm}$  in height or  $\geq 60\text{kg}$  in weight (*i.e. adult sizes*), the full CTAG Heart Adjudication Panel must be approached”, and some of the points that were raised when previous paediatric cases were adjudicated (e.g. data from right heart catheterisation).

- There was agreement that CTAG Heart Adjudication Panel approval is not required if the paediatric patient is on temporary MCS (e.g. VA ECMO) and there is agreement/ approval from the second paediatric centre (*i.e. both paediatric centres agree to SU listing*).
- There was agreement that CTAG Heart Adjudication Panel approval is required for paediatric patients with more than single organ failure/support on temporary MCS (e.g. VA ECMO but still intubated/ventilated and/or on dialysis), as it is currently applied to adult patients. However, several centres acknowledged some of the challenges in managing paediatric patients on MCS without sedation.
- There was agreement that right heart catheter is not required prior to SU listing in paediatric patients on temporary MCS. However, several centres indicated that right heart catheterisation should be considered in selected (older) paediatric patients, especially if transition to adult services prior to transplantation is likely.

## Summary and Recommendations

The donor heart allocation system was extensively discussed and debated, supported by a comprehensive review of relatively contemporaneous data. The FTWG did not achieve consensus in all the areas of discussion, which reflects the current limited evidence base of MCS in cardiogenic shock and more specifically MCS bridging to transplantation.

The SUHAS FTWG would like to make the following recommendations:

1. Adopt the proposed SUHAS criteria for patients with single ventricle physiology congenital heart disease.
2. Patients on isolated left ventricular support should not be excluded from the SUHAS.
3. The number of patients registered on the new SUHAS criteria for congenital heart disease and their outcomes should be reviewed at 12 or 24 months (depending on activity).
4. The number of patients on isolated LV support registered on the SUHAS and their outcomes should be reviewed at 12 or 24 months (depending on activity).
5. The Heart Allocation Scheme should be transitioned from a tier-based to a points-based system.
6. CTAG Heart Adjudication Panel approval should be removed for paediatric requests for urgent or super-urgent listing whereby a maximum acceptable donor size has been specified to be  $\geq 160$ cm in height or  $\geq 60$ kg in weight, if there is agreement/ approval from the second paediatric centre (i.e. both paediatric centres agree to SU listing).
7. CTAG Heart Adjudication Panel should be approached for paediatric requests for urgent or super-urgent listing whereby a maximum acceptable donor size has been specified to be  $\geq 160$ cm in height or  $\geq 60$ kg in weight with more than single organ failure/support on temporary MCS.

## Appendix

SUHAS criteria for congenital heart disease (single ventricle physiology)

Adult inpatient dependant on IV inotropes (on maximal tolerated dose of IV milrinone), worsening organ function and either ineligible or with high-risk anatomy for temporary MCS, AND one of the following:

- ICU admission and >1 inotrope with signs of hypoperfusion secondary to cardiogenic shock (lactate>2) and worsening renal (oliguria <1.0L/24 hours, eGFR<40), and/or liver function (bilirubin >2x upper limit of normal (40µmol/L) or transaminases >2x upper limit). There is normal renal size on imaging, eGFR>40 over the last 3 months. Liver size and hepatic function deemed acceptable for heart alone transplantation by pre-decompensation combined heart/liver MDT.
- Ineligible or too high-risk for EXCOR Revive/Fontan venous cannula VAD