

## Ventricular assist devices in pediatric cardiomyopathy and congenital heart disease patients: an analysis from the German National Register for Congenital Heart Defects

Lammers A. (1), Sprenger K.(1), Diller G.P. (2), Helm P.C. (3), Asfour B.(4), Abdul-Khaliq H.(5), Uebing A.(1), Ewert P.(6), Bauer U.M.M.(3), Tutarel O.(6)

Department of Paediatric Cardiology, University Hospital Münster, Münster/Germany (1); Division of Adult Congenital and Valvular Heart Disease, Department of Cardiovascular Medicine, University Hospital Münster, Münster/Germany (2); National Register for Congenital Heart Defects and Competence Network for Congenital Heart Defects, Berlin/Germany, DZHK (German Centre for Cardiovascular Research), Berlin/ Germany (3); Asklepios Klinik Sankt Augustin GmbH, German Pediatric Heart Center, Sankt Augustin/Germany (4); Department of Pediatric Cardiology, Saarland University Medical Center, Homburg/Germany (5); Department of Paediatric Cardiology and Congenital Heart Disease, German Heart Centre Munich, Technical University of Munich, Munich/Germany (6)

### Introduction

Heart failure is a major contributor to morbidity and mortality in pediatric cardiomyopathy and congenital heart disease (CHD) patients. Advanced therapies for heart failure like ventricular assist devices (VAD) are increasingly used in patients with acquired heart disease. Limited data exists, however, on the use and outcome of this technology in pediatric cardiomyopathy and CHD patients.

### Methods

Patients who were supported with a VAD were identified from the German National Register for Congenital Heart Defects. We report patient demographics, underlying cardiac defect, status of surgical repair/palliation, associated conditions, complications, and 30-day and overall mortality in this population. Univariate regression analysis was performed to identify predictors of an adverse outcome.

### Results

Overall, 91 patients (median age 4.29 years [IQR, 0.44-15.96], 45.1% female) received a VAD. 45 pat. (49.5%) had a congenital heart defect, while 46 (50.5%) were reported as cardiomyopathy. The type of VAD implanted was Berlin Heart in 31 pat. (34.1%), Heartware in 16 (17.6%), Levitronix in 9 (9.9%), unknown in 28 (30.8%). In 67 pat. (73.6%) the systemic ventricle was supported, in 4 (4.4%) the subpulmonary ventricle, and in 20 (22.0%) both. Reasons for implantation were heart failure in 46 (50.5%), post-surgery in 24 (26.4%), transfer from ECMO in 16 (17.6%), and miscellaneous in 5 (5.5%). At least one complication occurred in 54 pat. (59.3%) such as bleeding (n=37), thrombosis (n=15), embolic (n=19), 20 of which involved cerebral complications. Cause for explantation was clinical improvement in 22 (24.2%), death in 22 (24.2%), heart transplantation in 39 (42.9%), and change to ECMO in 3 (3.3%). Overall, 30-day mortality rate was 19.8%. Longer term all-cause mortality rate was 39.6%. On univariate regression analysis, no significant predictor could be identified for 30-day mortality. For all-cause mortality, age at implantation (OR:1.12; 95%CI 1.06 to 1.18, p<0.0001) and the presence of a congenital heart defect (OR: 4.92; 95%CI 1.97 to 12.32, p<0.001) were the only univariate predictors.

### Conclusions

The use of VAD for heart failure in pediatric cardiomyopathy and CHD patients is increasing. Even though, mortality remains high in this cohort, reflecting the complexity of this particular patient population presenting with heart failure.