Right ventricular assist device implantation for right ventricle

®Boško Skorić*

University Hospital Centre Zagreb, Zagreb, Croatia KEYWORDS: right ventricular failure, right ventricular assist device, ProtekDuo cannula.

CITATION: Cardiol Croat. 2021;16(9-10):309. | https://doi.org/10.15836/ccar2021.309

*ADDRESS FOR CORRESPONDENCE: Boško Skorić, Klinički bolnički centar Zagreb, Kišpatićeva 12, HR-10000

Zagreb, Croatia. / Phone: +385-1-236-7501 / Email: bskoric3@yahoo.com

ORCID: Boško Skorić, https://orcid.org/0000-0001-5979-2346

Right ventricular failure (RVF) refractory do medical therapy carries a high mortality risk, and circulatory support with right ventricular assist device (RVAD) may be a lifesaving option. 1-3 While surgical RVAD represents a more invasive therapeutical approach that carries about 50% mortality, the use of percutaneous RVAD support may improve outcomes. Acute RVF may result from 1) decreased contractility due to RV myocardial infarction or myocarditis; 2) volume overload, i.e. increase in preload due to excessive transfusions or following LVAD implantation; 3) pressure overload, i.e. increase in afterload during hypoxia, acidosis, or positive pressure ventilation. RV displays distinctive features in comparison to LV, including high complaint and low resistant pulmonary circulation. Accordingly, RV is more sensitive to an increase in afterload. Treatment of acute RVF includes specific causative treatment, optimization of preload, management of arrhythmia, reduction of afterload, and augmentation of contractility. If RVF is refractory, it is crucial to consider mechanical circulatory support on time. Percutaneous RV support includes VA-ECMO, Impella RP axial pump, and ProtekDuo cannula with an extracorporeal centrifugal pump. The selection of the device depends on whether RVF is isolated or part of biventricular failure, and whether it is a consequence of pulmonary artery (PA) disease. Percutaneous or surgically implanted RVAD is an option in isolated RV failure. Although ECMO can be also used in this setting, it is the device of choice if RVF is a consequence of PA disease. In the case of biventricular failure, a patient can be supported with paracorporeal surgically implanted or percutaneous BiVAD or ECMO. Support in RVF may be a bridge-to-recovery, bridge-to-candidacy, bridge-to-heart transplantation, or bridge-to-permanent RVAD. RECOVER RIGHT study showed 73% survival in patients supported with Impella RP. The Protek Duo cannula with internal jugular cannulation allows early ambulation during support. It is crucial to understand the pathophysiology of RVF, i.e. recognize ventricular interdependence with impaired LV filling to avoid futile therapeutical strategies like excessive volume infusion, and to start with mechanical support before irreversible multiorgan failure develops.

RECEIVED: July 28, 2021 ACCEPTED: August 5, 2021



- Kremer J, Farag M, Brcic A, Zubarevich A, Schamroth J, Kreusser MM, et al. Temporary right ventricular circulatory support following right ventricular infarction: results of a groin-free approach. ESC Heart Fail. 2020 Oct;7(5):2853-2861. https://doi.org/10.1002/ehf2.12888
- Anderson MB, Goldstein J, Milano C, Morris LD, Kormos RL, Bhama J, et al. Benefits of a novel percutaneous ventricular assist device for right heart failure: The prospective RECOVER RIGHT study of the Impella RP device. J Heart Lung Transplant. 2015 Dec;34(12):1549-60. https://doi.org/10.1016/j.healun.2015.08.018
- Konstam MA, Kiernan MS, Bernstein D, Bozkurt B, Jacob M, Kapur NK, et al; American Heart Association Council on Clinical Cardiology; Council
 on Cardiovascular Disease in the Young; and Council on Cardiovascular Surgery and Anesthesia. Evaluation and Management of Right-Sided
 Heart Failure: A Scientific Statement From the American Heart Association. Circulation. 2018 May 15;137(20):e578-e622.
 https://doi.org/10.1161/CIR.0000000000000560