



Letter to the Editor

Reply to “a pediatric perspective on the TAPSE/PASP ratio in pulmonary arterial hypertension”

Khodr Tello^{a,*}, Robert Naeije^b, Hossein A. Ghofrani^{a,c,d}, Henning Gall^{a,e}, Manuel J. Richter^a^a Department of Internal Medicine, Justus-Liebig-University Giessen, Universities of Giessen and Marburg Lung Center (UGMLC), Member of the German Center for Lung Research (DZL), Giessen, Germany^b Erasme University Hospital, Brussels, Belgium^c Department of Pneumology, Kerckhoff Heart, Rheuma and Thoracic Center, Bad Nauheim, Germany^d Department of Medicine, Imperial College London, London, UK^e Department of Epidemiology, Erasmus MC, University Medical Center Rotterdam, Rotterdam, the Netherlands

ARTICLE INFO

Article history:

Received 3 December 2018

Accepted 12 December 2018

Keywords:

Pulmonary arterial hypertension

Right ventricular function

Pediatric

TAPSE

PASP

Right ventricular-pulmonary arterial coupling

To the editor,

Many thanks to Koestenberger and co-workers for their correspondence emphasizing the pediatric perspective on the tricuspid annular plane systolic excursion/pulmonary arterial systolic pressure (TAPSE/PASP) ratio, an echocardiographic surrogate for right ventricular (RV)-pulmonary arterial (PA) coupling in patients with pulmonary arterial hypertension (PAH) [1]. We agree that adult TAPSE/PASP values should not be adopted as reference values for the pediatric age group [2], as acquisition might be challenging and development-dependent. To overcome some of these limitations, the TAPSE/pulmonary artery acceleration time ratio was introduced as a possible substitute for TAPSE/PASP in children [3]. In addition, RV fractional area change to invasive mean pulmonary artery pressure [4] and the ratio of RV area change to RV end-systolic area [5] were described in adult PAH as promising non-invasive surrogates of RV-PA coupling. However, caution should be exercised when using echocardiographic parameters as surrogates, because they have not yet been validated against invasive pressure-volume loop-derived measurements of RV-PA coupling. The

inverse relationship of TAPSE/PASP with pulmonary vascular resistance and its strong association with PA capacitance (stroke volume/pulse pressure) in patients with PAH [1] could suggest that TAPSE/PASP predominantly reflects afterload. TAPSE is an indirect measure of RV longitudinal shortening and consequently RV volume change, and PASP is associated with pulse pressure, which might explain the strong association of TAPSE/PASP with PA capacitance as well as the prognostic capacity of TAPSE/PASP.

In summary, knowledge gaps remain, and we further encourage Koestenberger and co-workers to explore echocardiographic parameters as surrogates for RV-PA coupling in children.

Funding

This work was funded by the Excellence Cluster Cardio-Pulmonary System (ECCPS) and the Collaborative Research Center (SFB) 1213 - Pulmonary Hypertension and Cor Pulmonale. For the letter, editorial assistance was provided by Dr. Claire Mulligan (Beacon Medical Communications Ltd., Brighton, UK), funded by the University of Giessen.

Conflict of interest

The authors report no relationships that could be construed as a conflict of interest.

References

- [1] K. Tello, J. Axmann, H.A. Ghofrani, et al., Relevance of the TAPSE/PASP ratio in pulmonary arterial hypertension, *Int. J. Cardiol.* 266 (2018) 229–235.
- [2] M. Koestenberger, A. Avian, M. Cantinotti, G. Hansmann, N. European Pediatric Pulmonary Vascular Disease, Tricuspid annular plane systolic excursion (TAPSE) in pediatric pulmonary hypertension: Integrating right ventricular ejection efficiency (RVEe) into advanced multi-parametric imaging, *Int. J. Cardiol.* 274 (2019) 296–298.
- [3] P.T. Levy, A. El Khuffash, K.V. Woo, G.K. Singh, Right ventricular-pulmonary vascular interactions: an emerging role for pulmonary artery acceleration time by echocardiography in adults and children, *J. Am. Soc. Echocardiogr.* 31 (2018) 962–964.

* Corresponding author at: Department of Internal Medicine, Justus-Liebig-University Giessen, Klinikstrasse 32, 35392 Giessen, Germany.

E-mail address: khodr.tello@innere.med.uni-giessen.de (K. Tello).

- [4] K.W. Prins, S.L. Archer, M. Pritzker, et al., Interleukin-6 is independently associated with right ventricular function in pulmonary arterial hypertension, *J. Heart Lung Transplant.* 37 (2018) 376–384.
- [5] S. French, M. Amsallem, N. Ouazani, et al., Non-invasive right ventricular load adaptability indices in patients with scleroderma-associated pulmonary arterial hypertension, *Pulm. Circ.* 8 (2018) (2045894018788268).