



# A Quadricuspid Aortic Valve Combined with Coronary Artery Disease

Živojin S Jonjev,<sup>1,2</sup> Novica Kalinić<sup>2</sup>

1. Institute for Cardiovascular Diseases of Vojvodina, Clinic of Cardiovascular Surgery, Sremska Kamenica, Serbia.
2. University of Banja Luka, Faculty of Medicine, University of Banja Luka, Banja Luka, the Republic of Srpska, Bosnia and Hercegovina.

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## Correspondence:

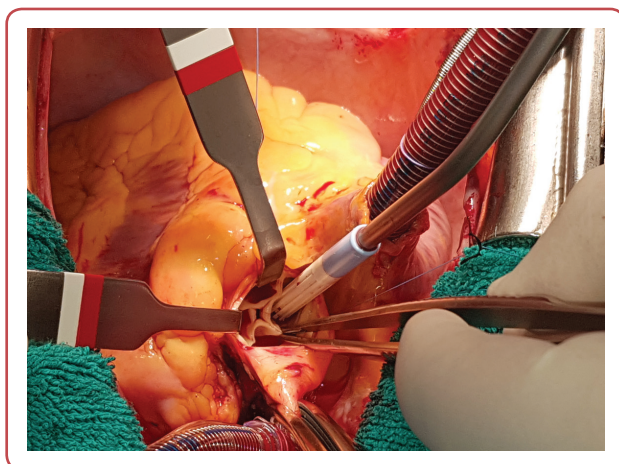
ŽIVOJIN S JONJEV  
M: + 381 64 218 23 67  
E: jonjevz@gmail.com

## Case history

Quadricuspid aortic valve (QAV) is a rare and purely understood congenital malformation with an estimated incidence of 0.013 % - 0.043 %. Here are presented images of an incompetent QAV found during coronary artery bypass grafting (CABG).

A 68 year old male was transferred from outside hospital with a sign of left heart failure. After admission, urgent coronarography was performed and significant stenosis ( $\geq 75$  %) was found on the proximal left anterior descending artery (LAD), ramus intermedius (RIM) and circumflex artery (RCx). Transoesophageal echocardiography revealed severe aortic regurgitation due to incompetent QAV, with hypokinetic anterior wall of the left ventricle.

After heart exposure, using cardiopulmonary bypass, cardiac arrest was induced as well as standard myocardial protection. A transversal aortotomy revealed enlarged annulus and four equal-sized aortic cusps (Type A)<sup>1</sup> with central gap and consequent severe malcoaptation of the QAV (Figure 1 and 2). The aortic valve was replaced with a #21 pericardial tissue prosthesis and myocardial revascularisation was successfully achieved with triple bypass using skeletonised left internal mammary artery to LAD and saphenous venous grafts to RIM and RCx. Patient had an uneventful postop course and was asymptomatic a year after surgery.



**Figure 1:** Intraoperative gross view of the quadricuspid aortic valve



**Figure 2:** Gross view of the specimen of the quadricuspid aortic valve

## Acknowledgements

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## Conflict of interest

None.

## References

1. Lin Y, Yin K, Wang Y, Yang D, Luo R, Dong L, et al. Clinical characteristics and surgical outcomes of dysfunctional quadricuspid aortic valve. *J Surg Res* 2018 Sep;229:223-9.