

The emerging roles of cardiovascular magnetic resonance imaging in transcatheter aortic valve implantation (TAVI)



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Transcatheter aortic valve implantation (TAVI) is now well established as an alternative to aortic valve replacement (AVR) in patients who are considered high-risk for surgery¹, or who are inoperable². Subsequent data have underscored the longevity of TAVI³, and the procedure has become so successful that trials are now ongoing (e.g., SURTAVI) to examine its potential use in an intermediate-risk population. Cardiovascular imaging techniques, particularly echocardiography and CT, play a vital role in TAVI⁴. However, the literature examining the role of cardiovascular magnetic resonance (CMR) in TAVI is currently limited to only a handful of papers.

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In this edition of EuroIntervention, Ribeiro et al⁵ describe their findings using CMR to look for myocardial injury in patients who had recently undergone TAVI. In a group of 37 patients, only one of whom did not show a post-procedural rise in troponin, the authors found that the only new areas of gadolinium enhancement seen were at the ventricular apex in patients who had undergone a transapical (TA) TAVI; this is similar to the apical scar often seen following insertion of a left ventricular assist device⁶ (**Figure 1**). A TA approach involves the insertion of a large calibre (≥ 18 Fr) sheath into the ventricular apex, and some data have shown an increased mortality risk with this approach⁷. Although this may largely be driven by higher rates of bleeding and stroke⁷, the results of the current paper would suggest that some of the risk of the transapical approach may be due to the long-term myocardial injury caused; TA patients were noted to have lower left ventricular ejection fraction at follow-up.

The lack of gadolinium enhancement seen in any other myocardial areas is an encouraging finding given that even minor necrosis of the order of 1.4% of ventricular myocardium can be detected with CMR. However, this finding must be contrasted with the results of MRI studies of the brain, which have shown that the vast majority of TAVI patients suffer some form of cerebral microembolic damage⁸. Therefore, given the large volume of embolic debris released during TAVI, some passage of this material down the coronary arteries – leading to post-procedural myocardial necrosis beyond the resolution of CMR – remains a possibility⁹. Indeed, previous studies that have performed CMR following PCI have shown that even relatively large embolic loads (coronary plaques up to 50 mm³ in volume) may be treated without causing any subsequent myocardial enhancement¹⁰.

In addition, many patients with severe aortic stenosis have significant left ventricular hypertrophy which predisposes to subendocardial myocardial ischaemia¹¹, particularly during placement of the TAVI prosthesis using a period of rapid ventricular pacing: a recent publication by Selle et al¹² has demonstrated that even short periods of rapid ventricular pacing can lead to microcirculatory arrest. This propensity to subendocardial ischaemia may explain why Yong et al found that a longer procedural duration and the absence of beta-blockade were associated with myocardial injury¹³. Therefore, although a lack of myocardial gadolinium enhancement on post-TAVI CMR may be reassuring, this does not mean that no permanent myocardial injury has taken place. Newer CMR imaging

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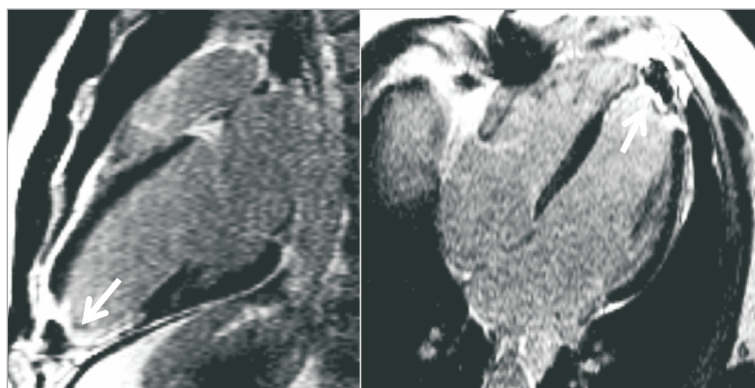


Figure 1. Example of transmural apical scarring (arrows) following left ventricular assist device insertion depicted on the late gadolinium enhancement images acquired in the vertical and horizontal long-axis views of the left ventricle (from Moon et al 2003), similar to that seen in the study by Ribeiro⁵. Reproduced in accordance with BioMed Central's Open Access Charter.

techniques, such as T1 mapping, may reveal global diffuse myocardial interstitial fibrosis that is not currently visualised using late gadolinium enhancement¹⁴.

A couple of limitations of the study must be highlighted, not least the small numbers of patients studied. A larger study, encompassing patients with and without significant left ventricular hypertrophy and perhaps confined to transfemoral cases only, would help to clarify the true incidence of CMR-detected myocardial injury following TAVI. Such a study must include not only T1 mapping but also T2-weighted myocardial oedema imaging sequences; the latter were not used in the current study but have widely been considered to display “area at risk” in CMR studies following myocardial infarction¹⁵.

What is the role of CMR in TAVI?

Looking forward, the use of CMR in patients undergoing, or being considered for, TAVI is likely to expand. Prior to TAVI, MRI can provide several important pieces of structural information: in addition to precise quantification of ventricular volumes and function, aortic valve planimetry can be performed to determine valve area accurately, and late gadolinium enhancement can show areas of myocardial scarring. Furthermore, work from our institution has shown that MRI can provide accurate aortic annulus measurements which, in terms of predicting the presence and severity of aortic regurgitation after TAVI, are comparable to those obtained using CT¹⁶. Also, in cases where CT angiography is not available or is deemed inappropriate, magnetic resonance angiography of the peripheral vasculature can be performed to assess access routes.

Following TAVI, CMR can be used to provide an accurate estimate of ventricular function, and a recent publication has also shown the accuracy of CMR to detect and quantify paravalvular leak¹⁷. Ultimately, however, unless any post-procedural CMR-specific measurements can be shown to lead to treatment changes that affect prognosis, it will remain highly debatable whether it is worth the time and expense to perform CMR on a routine basis after TAVI; echocardiography may be sufficient for the majority of cases.

In the longer term, however, a role may arise for percutaneous valve interventions such as TAVI to be completely MRI guided; real-time transarterial implantation of the Medtronic CoreValve (Medtronic, Minneapolis, MN, USA), using a modified, MRI-compatible delivery device, has already been described in a swine model¹⁸.

In summary, although larger studies are needed, the paper by Ribeiro et al reassures us that contemporary TAVI practice, outside of TA procedures, generally causes little significant macroscopic myocardial injury. As the volume of TAVI procedures being performed year-on-year continues to grow, and its application is extended to younger patients, it is likely that larger CMR series will be reported, further emphasising its importance as an imaging tool in the diagnosis and management of structural heart disease.

Conflict of interest statement

The authors have no conflicts of interest to declare.

References

1. Smith CR, Leon MB, Mack MJ, Miller DC, Moses JW, Svensson LG, Tuzcu EM, Webb JG, Fontana GP, Makkar RR, Williams M, Dewey T, Kapadia S, Babaliaros V, Thourani VH, Corso P, Pichard AD, Bavaria JE, Herrmann HC, Akin JJ, Anderson WN, Wang D, Pocock SJ; PARTNER Trial Investigators. Transcatheter versus surgical aortic-valve replacement in high-risk patients. *N Engl J Med.* 2011;364:2187-98.
2. Leon MB, Smith CR, Mack M, Miller DC, Moses JW, Svensson LG, Tuzcu EM, Webb JG, Fontana GP, Makkar RR, Brown DL, Block PC, Guyton RA, Pichard AD, Bavaria JE, Herrmann HC, Douglas PS, Petersen JL, Akin JJ, Anderson WN, Wang D, Pocock S; PARTNER Trial Investigators. Transcatheter aortic-valve implantation for aortic stenosis in patients who cannot undergo surgery. *N Engl J Med.* 2010;363:1597-607.
3. Mack MJ, Leon MB, Smith CR, Miller DC, Moses JW, Tuzcu EM, Webb JG, Douglas PS, Anderson WN, Blackstone EH, Kodali SK, Makkar RR, Fontana GP, Kapadia S, Bavaria J, Hahn RT, Thourani VH, Babaliaros V, Pichard A, Herrmann HC,

Brown DL, Williams M, Davidson MJ, Svensson LG; PARTNER 1 trial investigators, Akin J. 5-year outcomes of transcatheter aortic valve replacement or surgical aortic valve replacement for high surgical risk patients with aortic stenosis (PARTNER 1): a randomised controlled trial. *Lancet*. 2015 Mar 15. [Epub ahead of print].

4. Delgado V, Ng AC, Shanks M, van der Kley F, Schuijf JD, van de Veire NR, Kroft L, de Roos A, Schalij MJ, Bax JJ. Transcatheter aortic valve implantation: role of multimodality cardiac imaging. *Expert Rev Cardiovasc Ther*. 2010;8:113-23.

5. Ribeiro HB, Larose E, de la Paz Ricapito M, Le Ven F, Nombela-Franco L, Urena M, Allende R, Amat-Santos I, Dahou A, Capoulade R, Clavel MA, Mohammadi S, Paradis JM, De Larochellière R, Doyle D, Dumont E, Pibarot P, Rodés-Cabau J. Myocardial injury following transcatheter aortic valve implantation: insights from delayed-enhancement cardiovascular magnetic resonance. *EuroIntervention*. 2015;11:205-13.

6. Moon JC, Sievers B, Pennell DJ, Yacoub MH, Mohiaddin RH. Myocardial scarring caused by left ventricular assist device (LVAD) insertion demonstrated by cardiovascular magnetic resonance. *J Cardiovasc Magn Reson*. 2003;5:361-3.

7. Conrotto F, D'Ascenzo F, Francesca G, Colaci C, Sacciatella P, Biondi-Zoccai G, Moretti C, D'Amico M, Gaita F, Marra S. Impact of access on TAVI procedural and midterm follow-up: a meta-analysis of 13 studies and 10,468 patients. *J Interv Cardiol*. 2014;27:500-8.

8. Astarci P, Price J, Glineur D, D'Hoore W, Kefer J, Elkhoury G, Grandin C, Vanoverschelde JL. Cerebral embolization during percutaneous valve implantation does not occur during balloon inflation valvuloplasty: prospective diffusion-weighted brain MRI study. *J Heart Valve Dis*. 2013;22:79-84.

9. Rodés-Cabau J, Gutiérrez M, Bagur R, De Larochellière R, Doyle D, Côté M, Villeneuve J, Bertrand OF, Larose E, Manazzoni J, Pibarot P, Dumont E. Incidence, predictive factors, and prognostic value of myocardial injury following uncomplicated transcatheter aortic valve implantation. *J Am Coll Cardiol*. 2011;57:1988-99.

10. Porto I, Selvanayagam JB, Van Gaal WJ, Prati F, Cheng A, Channon K, Neubauer S, Banning AP. Plaque volume and occurrence and location of periprocedural myocardial necrosis after percutaneous coronary intervention: insights from delayed-enhancement magnetic resonance imaging, thrombolysis in myocardial infarction myocardial perfusion grade analysis, and intravascular ultrasound. *Circulation*. 2006;114:662-9.

11. Attarian DE, Jones RN, Currie WD, Hill RC, Sink JD, Olsen CO, Chitwood WR Jr, Wechsler AS. Characteristics of chronic left ventricular hypertrophy induced by subcoronary valvular aortic stenosis. II. Response to ischemia. *J Thorac Cardiovasc Surg*. 1981;81:389-95.

12. Selle A, Figulla HR, Ferrari M, Rademacher W, Goebel B, Hamadanchi A, Franz M, Schlueter A, Lehmann T, Lauten A. Impact of rapid ventricular pacing during TAVI on microvascular tissue perfusion. *Clin Res Cardiol*. 2014;103:902-11.

13. Yong ZY, Wiegerinck EM, Boerlage-van Dijk K, Koch KT, Vis MM, Bouma BJ, Henriques JP, Cocchieri R, Piek JJ, de Mol BA, Baan J Jr. Predictors and prognostic value of myocardial injury during transcatheter aortic valve implantation. *Circ Cardiovasc Interv*. 2012;5:415-23.

14. Lee SP, Lee W, Lee JM, Park EA, Kim HK, Kim YJ, Sohn DW. Assessment of diffuse myocardial fibrosis by using MR imaging in asymptomatic patients with aortic stenosis. *Radiology*. 2015;274:359-69.

15. Dall'Armellina E, Karia N, Lindsay AC, Karamitsos TD, Ferreira V, Robson MD, Kellman P, Francis JM, Forfar C, Prendergast BD, Banning AP, Channon KM, Kharbada RK, Neubauer S, Choudhury RP. Dynamic changes of edema and late gadolinium enhancement after acute myocardial infarction and their relationship to functional recovery and salvage index. *Circ Cardiovasc Imaging*. 2011;4:228-36.

16. Jabbour A, Ismail TF, Moat N, Gulati A, Roussin I, Alpendurada F, Park B, Okoroafo F, Asgar A, Barker S, Davies S, Prasad SK, Rubens M, Mohiaddin RH. Multimodality imaging in transcatheter aortic valve implantation and post-procedural aortic regurgitation: comparison among cardiovascular magnetic resonance, cardiac computed tomography, and echocardiography. *J Am Coll Cardiol*. 2011;58:2165-73.

17. Crouch G, Tully PJ, Bennetts J, Sinhal A, Bradbrook C, Penhall AL, De Pasquale CG, Baker RA, Selvanayagam JB. Quantitative assessment of paravalvular regurgitation following transcatheter aortic valve replacement. *J Cardiovasc Magn Reson*. 2015;17:32.

18. Kahlert P, Parohl N, Albert J, Schäfer L, Reinhardt R, Kaiser GM, McDougall I, Decker B, Plicht B, Erbel R, Eggebrecht H, Ladd ME, Quick HH. Real-time magnetic resonance imaging-guided transarterial aortic valve implantation: in vivo evaluation in swine. *J Am Coll Cardiol*. 2012;59:192-3.