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Multimodality imaging in heart valve disease

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openheart Multimodality imaging in heart valve disease

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ABSTRACT

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Professor John B Chambers; john.chambers@gstt.nhs.uk In patients with heart valve disease, echocardiography is the mainstay for diagnosis, assessment and serial surveillance. However, other modalities, notably cardiac MRI and CT, are used if echocardiographic imaging is suboptimal but can also give complementary information to improve assessment of the valve lesion and cardiac compensation to aid the timing of surgery and determine risk. This statement discusses the way these imaging techniques are currently integrated to improve care beyond what is possible with echocardiography alone.

INTRODUCTION

Heart valve disease is common and a major indication for imaging in all cardiac centres. Imaging needs to assess: (1) valve morphology to determine the aetiology and suitability for invasive intervention; (2)haemodynamic severity; (3) remodelling of the left ventricle (LV) and right ventricle (RV); (4) involvement of the aorta and (5) the prediction of adverse cardiovascular events. Echocardiography will continue as the first-line technique for diagnosis and is likely to remain the mainstay for assessment and serial surveillance in most cases. However, other modalities, notably cardiac MRI (CMR) and CT, are used if echocardiographic imaging is suboptimal and can give complementary information, particularly to aid risk assessment.

The purpose of this statement is to summarise those multimodality approaches to valve disease currently available and how these might develop in the future. It is not intended as a systematic review. The aim is to encourage thinking beyond echocardiography towards an integrated approach to imaging in valve disease that uses each technique to its best advantage.

Aortic valve disease

Transthoracic echocardiography (TTE) is used for assessing the aetiology and severity of the valve disease and the remodelling response of the LV. If echocardiographic image quality is poor (table 1), CT or CMR can image the valve (figure 1) while CMR can measure velocities across the valve.¹ Anatomical area on CT or CMR can be useful and, although not the same as effective orifice area, it is possible that a combination of anatomical area and calcium score will give a reliable measure of the grade of stenosis² (figure 1).

In aortic stenosis, the echocardiographic minimum data set is the peak transaortic velocity, the mean gradient and the effective orifice area,³ but these measures sometimes give discordant results.² CT can help resolve this. First, the LV outflow tract may be oval rather than circular⁴ and CT can assess the size and shape of the LV outflow tract to allow correction of the continuity equation. This can also be carried out using threedimensional (3D) echocardiography provided that image quality is adequate.⁵ Second, CT can give a calcium score for the valve which can differentiate severe from more moderate stenosis^{6–8} with one suggested cut-off of 2065 Agatston units (AU) in males and 1275 AU in females.

The risk of events in aortic stenosis is dependent on a number of factors including the results of exercise testing. TTE suggests a high risk of events if the Vmax has increased by >0.3 m/s in a year with heavy calcification as assessed subjectively,⁹ but quantification of calcification by CT is likely to be more accurate.⁶ ⁷ Fused positron emission tomography (PET)/CT imaging with 18F-fluoride could assess both the burden of valve calcification (CT calcium score) and its activity (18F-fluoride PET) and is of potential use in refining the prediction of aortic stenosis progression.¹⁰ ¹¹

Estimation of the grade of aortic regurgitation (AR) using multiple echocardiographic modalities remains standard.¹² However, some situations, for example, a bicuspid



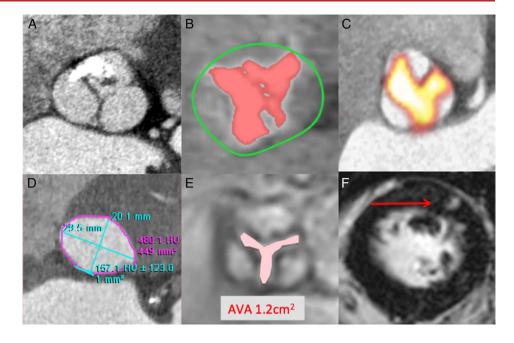
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Table 1 Role of CT and CMR beyond those provided by echocardiography		
Aortic stenosis		
>CT	Imaging of valve if echo window suboptimal	
	Imaging aorta if not seen clearly on echocardiography	
	LV outflow area to improve accuracy of continuity equation if grade uncertain	
	Aortic valve calcium (as a sign of severe aortic stenosis, eg, in low gradient low flow and as a marker of a	
	high risk of events on follow-up)	
	CT/PET to assess calcification and activity	
	Preintervention—coronary anatomy, detection of porcelain aorta, peripheral vessel anatomy, height of	
	coronary artery ostia above the valve	
CMR	Imaging of valve if echo window suboptimal	
Civin	Imaging or value if echo window suboplimation in a	
	LV mass if required for research, eg, to measure rate of regression after surgery	
	Future possibility of LV fibrosis as a sign suggesting early surgery	
Aortio rogurgit		
Aortic regurgita		
U1	Imaging of valve if echo window suboptimal	
	Imaging aorta if not seen clearly on echocardiography	
CMR	Imaging of valve if echo window suboptimal Imaging aorta if not seen clearly on echocardiography	
Mitral value die	Future possibility of surgery guided by LV volumes and early myocardial fibrosis	
Mitral valve dis	sease	
-	Imaging of volvo if only window subortimal	
CT/CMR MR	Imaging of valve if echo window suboptimal	
CT	Refining imaging of subvalve apparatus before percutaneous mitral valve procedures	
01	Predicting LV outflow tract obstruction after percutaneous mitral valve procedures	
	Quantifying mitral annulus calcification before mitral valve repair	
CMR	Imaging of valve if echo window suboptimal	
CIMIN	Future possibility of surgery guided by LV volumes and evidence of early fibrosis	
Right-sided va		
TR		
СТ	No current indications	
CMR	RV volumes to guide surgery	
CIVIN	Tricuspid annulus diameter to guide repair at the same time as left-sided surgery	
PR	Thouspid annulus diameter to guide repair at the same time as left-sided surgery	
CT	No current indications	
CMR		
CIVIN	Better than echo for imaging the valve and detecting obstruction above or below the valve, and branch artery stenoses	
	Quantification of pulmonary regurgitation	
Deplessment	Quantification of serial RV volumes to guide surgery	
Replacement I CT		
U1	Imaging of leaflets or occluder to differentiate patient-prosthesis mismatch from pathological obstruction	
	Detection of pannus	
CMR	Quantification of regurgitation	
Endocarditis	Detection of variation, or an heavily calcified values	
СТ	Detection of vegetation, eg, on heavily calcified valves	
	May detect aortic root abscess missed by echo	
	Coronary anatomy to avoid invasive angiography before surgery	
	CT/PET to detect endocarditis, eg, on replacement valves or electrical devices	
CMR	May detect root abscesses missed on echo	
CMR, cardiac magnetic resonance imaging; MS, mitral stenosis; LV, left ventricle; PET, positron emission tomography; PR, pulmonary regurgitation; TR, tricuspid regurgitation; RV, right ventricle.		
regurgitation, rrn, treuspid regurgitation, rrv, right ventricle.		

valve with an eccentric jet, may be difficult to assess and CMR quantification may provide useful complementary information. In addition, there is evidence that CMR identifies patients at high risk of events better than does TTE¹³ (figure 2). The quantification of AR appears to differ between echocardiography and CMR, with a significant overlap between grades, and a suggestion of lower quantities by CMR than TTE (figure 2). This

may be due to differing techniques for each method, and/or the potential for mild underestimation of AR by CMR.

CT (figure 3) or CMR (figure 4) is already used in clinical practice if the ascending aorta is not well imaged on TTE or to check the aortic diameter when approaching a surgical threshold. It may be better to use CT when considering referral for aortic valve surgery Figure 1 Multimodality imaging of aortic stenosis. (A) Contrast CT imaging of the aortic valve can provide detail regarding valve morphology and the distribution of calcification. (B) CT calcium scoring allows reproducible quantification of the calcific burden, which acts as a marker of disease severity. (C) Fused positron emission tomography and CT imaging with 18F-fluoride provides an indication of ongoing calcification activity in the valve. CT has an important clinical role in the workup of patients prior to transcatheter aortic valve implantation, providing accurate dimensions of the annulus for valve sizing (D) while cardiac MRI can be used to planimeter the aortic valve area (E) and to detect replacement myocardial fibrosis, red arrow (F).



since this can also detect calcium in the ascending aorta and identify porcelain aorta as an indication for a transcatheter procedure instead of conventional surgery. In those at no more than moderate coronary risk (eg, younger patients with a bicuspid aortic valve), CT can produce an assessment of the aorta and coronary arteries in one study. CT is also useful in the workup towards intervention. For transcatheter aortic valve implantation, it provides information additional to echo including the degree of calcification of the leaflets, the distance to the coronary arteries and the calibre, tortuosity and calcific burden of the peripheral vessels.¹⁴

Current thresholds for surgery in AR are based on LV diameters on TTE, but these may be unreliable

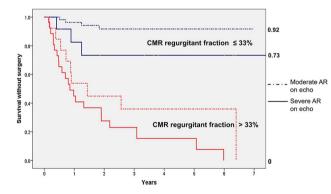


Figure 2 Comparison of CMR and echocardiography in aortic regurgitation. In 109 asymptomatic patients with moderate or severe aortic regurgitation on echocardiography, prognosis was better related to the regurgitant fraction on CMR with a cut-point of 33%. The graph shows CMR regurgitant fraction \leq 33% in blue and >33% in red. The regurgitation was either moderate or severe by echocardiography in both of these two CMR groups.

since the LV becomes more spherical in severe AR and a linear dimension may not be representative of the whole LV. There is still little information on volumetric thresholds for surgery, but it is possible that LV volumes by CMR will be better than those by TTE. It is most likely that myocardial fibrosis detected by CMR^{15–17} will provide a powerful predictor of events in aortic stenosis and regurgitation, but further work is required before this can be recommended clinically. CMR is far more accurate than TTE for quantifying LV mass and is therefore useful in research studies to document the regression of LV hypertrophy after surgery,¹⁸ although it has not yet been shown to be a prognostic marker.

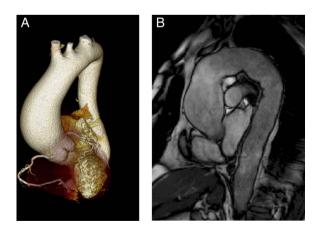


Figure 3 Imaging the aorta using CT and cardiac MRI (CMR). On the left is a reconstructed three-dimensional-rendered CT scan of the heart with a dilated ascending aorta and on the right is a steady-state free precession (SSFP) image on CMR of a moderately dilated ascending aorta.

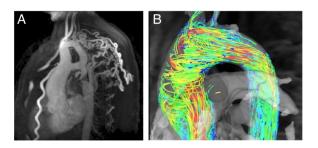


Figure 4 The aorta using cardiac MRI (CMR). (A) A contrast MR angiogram showing critical coarctation and very dilated thoracic collateral vessels. (B) A four-dimensional CMR flow image showing very helical flow in the ascending aorta in a patient with a bicuspid aortic valve.

Mitral valve disease

TTE is the mainstay for assessing the aetiology, morphology and grade of stenosis and regurgitation as well as the adaptation of the right and left heart. A 3D echo, particularly using transoesophageal echocardiography (TOE), improves identification of the minimum orifice for planimetry in mitral stenosis¹⁹ (figure 5) and may also improve the assessment of valve anatomy and function of the valve in mitral regurgitation caused by mitral prolapse¹⁹ (figure 6).

CMR is already used to assess mitral valve morphology if the TTE window is suboptimal and quantification can

Figure 5 A three-dimensional (3D) echocardiogram showing planimetry of a stenotic mitral valve. The 3D image allows alignment of the plane to ensure that planimetry is performed at the minimum orifice (courtesy Dr Stam Kapetanakis).

be performed by CMR if the TTE is uncertain or produces discrepant results. In mitral regurgitation, thresholds for surgery are based on a linear diameter on TTE, but volumes are expected to be more reliable.²⁰ This is a potential role for CMR, although this is not a clinical routine. It is possible that risk stratification by CMR assessment of the grade of MR and LV volumes may aid the frequency of follow-up clinically and by echocardiography and the need for functional tests.

CT can provide an assessment of mitral annular calcification if this appears severe on TTE to help determine the feasibility and safety of mitral repair or replacement. For percutaneous mitral valve interventions, CT may also be useful in further defining mitral valve anatomy and the subvalve structures including false tendons and hypertrabeculation. CT may be useful for predicting LV outflow tract obstruction after percutaneous mitral valve implantation.

Tricuspid and pulmonary disease

TTE is used for initial diagnosis and will retain a firstline role into the future. However, CMR is used routinely for the assessment of the RV in adult congenital disease,²¹ and we believe it should be used more frequently in severe tricuspid or pulmonary regurgitation (table 1) where decisions for surgery rest on an accurate assessment of RV volume, or a serial change in RV size or function. It is also better for refining the description

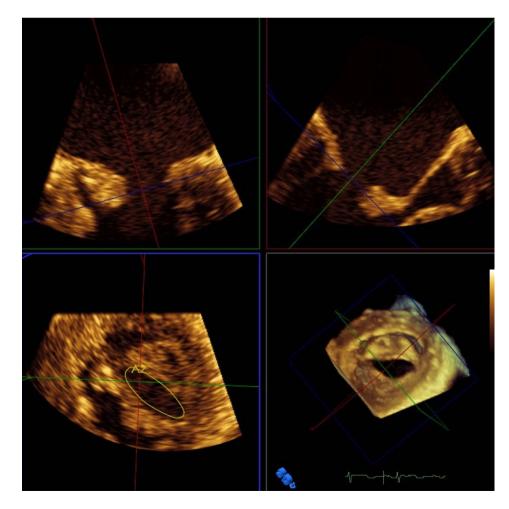
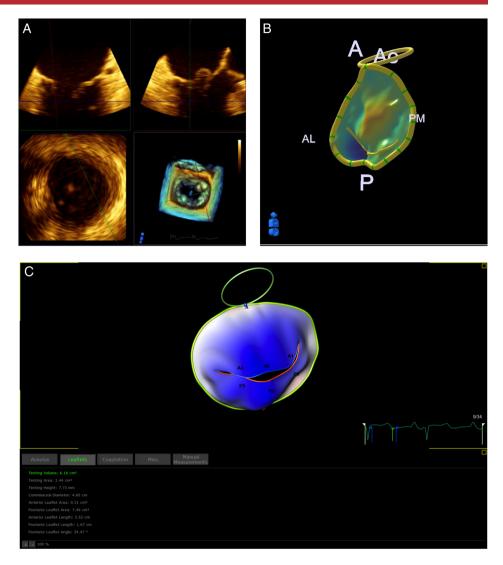


Figure 6 Three-dimensional (3D) in mitral prolapse. (A) A 3D image of the valve showing prolapse of the middle scallop, P2 using the Carpentier classification. (B) A colour-contoured map with prolapsing areas in red and restricted areas in blue. The main lesion is prolapse of the middle portion of the anterior leaflet. (C) A colour-contoured map in a patient with functional mitral regurgitation showing restriction of both mitral leaflets (courtesy Dr Stam Kapetanakis). A, anterior; AL, anterolateral; Ao, aorta; P, posterior; PM, posteromedial.



of the pulmonary valve, quantifying pulmonary regurgitation looking for obstruction above and below the valve and defining branch pulmonary stenoses²¹ (figure 7). CMR is particularly useful in larger adult patients in whom assessment of the pulmonary valve with echocardiography may be difficult.

Tricuspid valve repair may be recommended at the same time as left-sided surgery depending on the grade of tricuspid regurgitation and the annulus diameter. The latter is better assessed using 3D echocardiography than 2D and also using CMR. CT is not usually helpful for right-sided valve disease apart from excluding lung pathology contributing to elevated pulmonary artery pressures.

Replacement heart valves

TTE can usually make the initial diagnosis of valve obstruction or abnormal regurgitation. However, TOE is often needed to quantify paraprosthetic mitral regurgitation to determine the cause of mitral obstruction.

CT is likely to have a key role in patients with unexpectedly high gradients, especially when immediate postimplantation echocardiograms are not available (table 1). In the early recovery period after surgery in biological replacement valves, high gradients may be caused by small thrombi at the base of the cusps and these can be detected by CT^{22} ²³ but not by TTE or TOE. Thereafter, the differentiation of patient-prosthesis mismatch from primary failure in the aortic position may be difficult on TTE or even TOE since the cusps or occluder may not be imaged fully. CT can then be used for imaging the leaflet motion of bileaflet mechanical valves, although fluoroscopy may be still better for a single tilting disc and caged-ball valves. Pannus is also difficult to detect by TTE and TOE, but there is growing evidence that CT is useful²⁴ (figure 8).

In mechanical replacement valves CMR can assess forward flow patterns and localise and quantify regurgitant flow. This is especially so for paravalvular leaks (both in conventional and transcatheter replacement valves), which can be challenging to assess with echocardiography. CMR is also useful for ventricular volumes and function if TTE images are suboptimal.

Endocarditis

The complementary use of TTE and TOE has a high sensitivity and specificity for vegetations and local



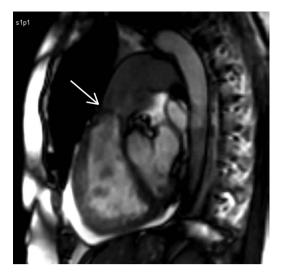


Figure 7 Severe congenital pulmonic stenosis. This is a cardiac MRI image (steady-state free precession, SSFP) in a sagittal view through the right ventricular outflow tract, demonstrating mobile leaflets but fused tips of the pulmonic valve (arrow).

complications.²⁵ However, CT may be of value in detecting vegetations on heavily calcified valves²⁶ ²⁷ and may detect abscesses missed by TOE. CMR is not ideal for detecting small vegetation due to their small size and chaotic, highly mobile motion which presents problems for the images acquired over several cardiac cycles. Both CT and CMR are useful if aortic root pathology is not well seen on TTE and if TOE is not feasible, or if there is complex pathology (eg, false aneurysms or complex abscesses). Coronary CT angiography can be considered in place of invasive coronary angiography to avoid the risk of dislodging vegetation.

In the future, combined PET-CT is likely to be useful for diagnosing endocarditis in difficult cases, for example, infection of implantable electrical devices²⁸ and replacement heart valves,²⁹ although issues with a myocardial uptake of this tracer will need to be resolved. This indication is now included in the most recent European Society of Cardiology guidelines on infective endocarditis.³⁰

Conclusion including future work

Multimodality imaging implies using each technique where it has most to offer rather than simply repeating

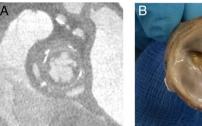




Figure 8 Pannus related to a stented biological valve. (A) CT and (B) surgical finding in the same patient.

full studies by each technique. Examples are surgery based on a combination of severe aortic stenosis on TTE and high CT calcium score, or trials of early surgery in aortic and mitral regurgitation based on CMR assessment of regurgitant fraction and LV volumes. This approach has obvious advantages but has not been tested clinically and is largely unresearched. This statement is intended to draw attention to the exciting possibilities and to act as a call for the research community to plan collaborative projects.

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Competing interests GJM-H receives lecture fees from GE for cardiac CT education.

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REFERENCES

- Myerson SG. Heart valve disease: investigation by cardiovascular magnetic resonance. J Cardiovasc Magn Reson 2012;14:7.
- Dweck MR, Chin C, Newby DE. Small valve area with low-gradient aortic stenosis: beware the hard hearted. J Am Coll Cardiol 2013;62:2339–40.
- Baumgartner H, Hung J, Bermejo J, et al. Echocardiographic assessment of valve stenosis: EAE/ASE recommendations for clinical practice. Eur J Echocardiogr 2009;10:1–25.
- De Vecchi C, Caudron J, Dubourg B, et al. Effect of the ellipsoid shape of the left ventricular outflow tract on the echocardiographic assessment of aortic valve area in aortic stenosis. J Cardiovasc Comput Tomogr 2014;8:52–7.
- Gutiérrez-Chico JL, Zamorano JL, Prieto-Moriche E, *et al.* Real-time three-dimensional echocardiography in aortic stenosis: a novel, simple, and reliable method to improve accuracy in area calculation. *Eur Heart J* 2008;29:1296–306.
- Messika-Zeitoun D, Bielak LF, Peyser PA, et al. Aortic valve calcification: determinants and progression in the population. Arterioscler Thromb Vasc Biol 2007;27:642–8.
- Clavel MA, Pibarot P, Messika-Zeitoun D, *et al.* Impact of aortic valve calcification, as measured by MDCT, on survival in patients with aortic stenosis: results of an international registry study. *J Am Coll Cardiol* 2014;64:1202–13.
- Kamperidis V, van Rosendael PJ, Katsanos S, *et al.* Low gradient severe aortic stenosis with preserved ejection fraction: reclassification of severity by fusion of Doppler and computed tomographic data. *Eur Heart J* 2015;36:2087–96.
- Rosenhek R, Binder T, Porenta G, et al. Predictors of outcome in severe asymptomatic aortic stenosis. N Engl J Med 2000;343:611–17.
- 10. Dweck MR, Jones C, Joshi NV, *et al.* Assessment of valvular calcification and inflammation by positron emission tomography in patients with aortic stenosis. *Circulation* 2012;125:76–86.
- Dweck MR, Jenkins WS, Vesey AT, et al. 18F-NaF uptake is a marker of active calcification and disease progression in patients with aortic stenosis. Circ Cardiovas Imaging 2014;7:371–8.
- Lancellotti P, Tribouilloy C, Hagendorff A, et al. European Association of Echocardiography recommendations for the assessment of valvular regurgitation. Part 1: aortic and pulmonary regurgitation (native valve disease). Eur J Echocardiogr 2010;11:223–44.
- Myerson SG, d'Arcy J, Mohiaddin R, *et al.* Aortic regurgitation quantification using cardiovascular magnetic resonance: association with clinical outcome. *Circulation* 2012;126:1452–6.
- Achenbach S, Delgado V, Hausleiter J, et al. SCCT Expert consensus document on computed tomography imaging before transcatheter aortic valve implantation (TAVI)/transcatheter aortic valve replacement (TAVR). J Cardiovasc Comput Tomogr 2012;6:366–80.

- Weidemann F, Herrmann S, Störk S, *et al.* Impact of myocardial fibrosis in patients with symptomatic severe aortic stenosis. *Circulation* 2009;120:577–84.
- Dweck MR, Joshi S, Murigu T, *et al.* Mid-wall fibrosis is an independent predictor of mortality in patients with aortic stenosis. *J Am Coll Cardiol* 2011;58:1271–9.
- Chin CW, Shah AS, Vassiliou V, et al. Left ventricular hypertrophy with strain and aortic stenosis. *Circulation* 2014;130:1607–16.
- de Marvao A, Dawes TJW, Shi W, *et al.* Population-based studies of myocardial hypertrophy: high resolution cardiovascular magnetic resonance atlases improve statistical power. *J Cardiovasc Magn Reson* 2014;16:16. http://jcmr-online.com/content/16/1/16
- Zamorano J, Cordeiro P, Sugeng L, *et al.* Real-time three-dimensional echocardiography for rheumatic mitral valve stenosis evaluation: an accurate and novel approach. *J Am Coll Cardiol* 2004;43:2091–6.
- Marsan NA, Westenberg JJ, Ypenburg C, et al. Quantification of functional mitral regurgitation by real-time 3D echocardiography: comparison with 3D velocity-encoded cardiac magnetic resonance. JACC Cardiovasc Imag 2009;2:1245–52.
- Baumgartner H, Bonhoeffer P, De Groot NM, et al. ESC guidelines for the management of grown-up congenital heart disease (new version 2010). Eur Heart J 2010;31:2915–57.
- Habets J, Symersky P, van Herwerden LA, et al. Prosthetic heart valve assessment with multidetector-row CT: imaging characteristics of 91 valves in 83 patients. Eur Radiol 2011;21:1390–6.
- Pache G, Blanke P, Zeh W, et al. Cusp thrombosis after transcatheter aortic valve replacement detected by computed tomography and echocardiography. Eur Heart J 2013;34:3546.

- Teshima H, Hayashida N, Yano H, et al. Obstruction of St Jude Medical valves in the aortic position: histology and immunohistochemistry of pannus. J Thorac Cardiovasc Surg 2003;126:401–7.
- Habib G, Badano L, Tribouilloy C, *et al.* Recommendations for the practice of echocardiography in infective endocarditis. *Eur J Echocardiogr* 2010;11:202–19.
- Feuchtner GM, Stolzmann P, Dichtl W, et al. Multislice computed tomography in infective endocarditis: comparison with transesophageal echocardiography and intraoperative findings. J Am Coll Cardiol 2009;53:436–44.
- Fagman E, Perrotta S, Bech-Hanssen O, *et al.* ECG-gated computed tomography: a new role for patients with suspected aortic prosthetic valve endocarditis. *Eur Radiol* 2012;22:2407–14.
- Sarrazin JF, Philippon F, Tessier M, et al. Usefulness of fluorine-18 positron emission tomography/computed tomography for identification of cardiovascular implantable electronic device infections. J Am Coll Cardiol 2012;59:1616–25.
- Saby L, Laas O, Habib G, et al. Positron emission tomography/ computed tomography for diagnosis of prosthetic valve endocarditis: increased valvular 18F-fluorodeoxyglucose uptake as a novel major criterion. J Am Coll Cardiol 2013;61:2374–82.
- Habib G, Lancellotti P, Antunes MJ, et al., Authors/Task Force Members. 2015 ESC Guidelines for the management of infective endocarditis: The Task Force for the Management of Infective Endocarditis of the European Society of Cardiology (ESC) Endorsed by: European Association for Cardio-Thoracic Surgery (EACTS), the European Association of Nuclear Medicine (EANM). *Eur Heart J* 2015;36:3075–128.

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