Cardiac CT provides uniquely accurate and comprehensive assessment of bioprosthetic aortic valve stenosis

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DESCRIPTION
An 81-year-old man was referred to the cardiology clinic with breathlessness and angina. His history included triple-vessel coronary artery bypass graft (CABG) plus St Jude Epic 21 mm bioprosthetic aortic valve replacement (AVR) surgery 7 years prior.

Transthoracic echocardiography (TTE) demonstrated severely elevated Doppler AVR velocities (VMax 4.7 m/s) and severe AVR stenosis (valve area 0.7 cm²). This was the suspected cause of symptoms, and transfemoral valve-in-valve AVR valve-in-valve transcatheter aortic valve implantation (VIV-TAVI) was being considered. However, TTE image quality was suboptimal due to echocardiographic windows and valve echogenicity, precluding accurate leaflet assessment (figure 1A–D). Transoesophageal echocardiography corroborated TTE findings (VMax 5.5 m/s) but failed to delineate the mechanism of AVR restriction (figure 1E–H). Degenerative leaflet calcification was evident on echocardiography, however echogenicity around the sewing ring prevented distinction between calcification and pannus, and ultrasound dropout precluded thrombus exclusion.

Invasive coronary angiography confirmed severe native coronary disease and patency of two bypass grafts, however the third graft could not be cannulated. Cardiovascular MRI confirmed inferolateral infarction, mild left ventricular systolic dysfunction and significant AVR stenosis. However, high antegrade AVR flow velocities and metallic valve stents resulted in signal loss and failure to delineate the mechanism of AVR restriction due to flow and susceptibility artefacts, respectively.

Given the limitations of echocardiography and angiography, a prospectively gated cardiac CT scan was performed (figure 2A,B). Its high spatial resolution and ability to characterise tissue based on CT absorption (attenuation) characteristics allowed accurate non-invasive AVR and coronary artery assessment. Absence of lung parenchyma and pulmonary embolic disease, and excellent patency of all bypass grafts were confirmed. Circumferential low attenuation (125 Hounsfield units; HU) around the underside of the sewing ring was consistent with pannus. There was accumulation of a large area of very low attenuation material (35 HU) over the non-coronary cusp and adjacent sewing ring segment. This was in keeping with significant thrombus burden causing symptoms due to leaflet restriction and orifice encroachment.

VIV-TAVI workup was postponed, and a repeat CT study after 3 months of warfarinisation confirmed thrombus resolution. Residual thin circumferential pannus and mild non-calcified degenerative leaflet thickening were confirmed (figure 2C,D). The patient was rendered asymptomatic with, corroboratively, significantly reduced TTE AVR velocities (VMax 3.8 m/s).

Figure 1  Top row (tranthoracic echocardiography): (A) Parasternal long-axis view with limited AVR visualisation (open-headed arrow) with increased echogenicity which can be due to pannus, calcification or metallic valve components; (B) turbulent antegrade colour-Doppler flow; (C) apical five-chamber view with limited AVR visualisation (open-headed arrow); (D) severely elevated antegrade AVR velocities on continuous-wave Doppler (VMax 6.4 m/s) suggestive of severe valve stenosis/obstruction. Bottom row (transoesophageal echocardiography): (E–G) Limited AVR visualisation (open-headed arrow) with dropout signal loss artefact (closed-headed arrow) precluding optimal valve leaflet visualisation; (H) severely elevated antegrade AVR velocities (VMax 5.5 m/s). AVR, aortic valve replacement.
Cardiac CT was the only modality adequately assessing the AVR, allowing appropriate therapy to be directed and preventing redo AVR surgery. Mild residual asymptomatic AVR stenosis due to mild leaflet degeneration and pannus was confirmed. The patient remains asymptomatic 6 months later on warfarin.

Prosthetic valve imaging using echocardiography can be limited by reverberation (reflective) and shielding (signal loss) artefacts from metal components which can also be challenging to differentiate from degenerative calcification. X-ray fluoroscopy can demonstrate leaflet mobility but is limited by its angle dependence and inability to visualise pannus and thrombus. Cardiac CT allows excellent non-invasive prosthetic valve assessment due to excellent spatial resolution and visualisation of leaflet motion. CT attenuation characteristics (HU) allow distinction between the key differentials of valve obstruction—thrombus and pannus—which can be notoriously challenging using echocardiography, X-ray fluoroscopy and cardiovascular MRI. CT also provides simultaneous excellent native coronary artery, CABG, aorta and pulmonary arterial imaging. Accurate CT aortic root assessment allows unrivalled assessment of endocarditic complications, including dehiscence, abscess, fistula and aneurysm formation. Cardiac CT is a valuable but currently underused tool in the multimodality imaging assessment of prosthetic cardiac valves, despite widespread availability.

**Learning points**

- Imaging of prosthetic cardiac valves using echocardiography can be significantly limited by reverberation and shielding artefacts, degrading leaflet assessment.
- Degenerative calcification, pannus and thrombus are key differentials for bioprosthetic valve obstruction and can be challenging to assess on echocardiography, X-ray fluoroscopy and cardiovascular MRI.
- Cardiac CT allows accurate non-invasive assessment of prosthetic valves with high spatial resolution. It allows excellent assessment of leaflet mobility, assessment for calcification, pannus and thrombus, visualisation of complications arising from endocarditis (abscesses, dehiscence) and also permits left ventricular and coronary artery assessment within the same study.

**Contributors**

All authors were heavily involved in production of the manuscript and images, and then finalising approval of the version published. All authors are agreeable to be accountable for the article and to ensure that all questions regarding the accuracy or integrity of the article are investigated and resolved. JNK: planning, conception and production of manuscript/images; clinical management of patient; undertook cardiac imaging (TTE, TOE, CMR, CT) and interpretation of data. BD: review and editing of manuscript; involved in cardiac imaging (CT) and interpretation of data. SV: review and editing of manuscript; clinical management of patient; involved in cardiac imaging (CMR, CT) and interpretation of data. BR: review and editing of manuscript; clinical management of patient; undertook cardiac imaging (TTE, TOE, CMR, CT) and interpretation of data. WM: review and editing of manuscript; involved in cardiac imaging (CMR, CT) and interpretation of data.

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