CLINICAL VIGNETTE

The bailout transseptal approach during valve-in-valve transcatheter aortic valve implantation with difficult crossing of the degenerated Mitroflow bioprosthetic valve

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Valve-in-valve (ViV) transcatheter aortic valve implantation (TAVI) is an accepted treatment in patients with structural valve deterioration (SVD) who are not good candidates for reoperation.^{1,2} A 70-year-old man with New York Heart Association class III heart failure was admitted with SVD of a Mitroflow 21-mm bovine pericardial prosthesis implanted 7 years ago concomitantly with coronary artery bypass grafting. Transthoracic echocardiography showed severe valve stenosis (effective orifice area, 0.7 cm²; mean gradient, 62 mm Hg) and left ventricular ejection fraction of 50%. Because of obesity (body mass index, 37.8 kg/m²) and high perioperative mortality risk (EuroSCORE, 6.31%), the Heart Team recommended transfemoral ViV TAVI after analyzing the multislice contrast computed tomography (FIGURE 1A and 1B).

The procedure was attempted by the right femoral approach under local anesthesia. Despite the use of several catheters (Amplatz left 1, Amplatz left 2, Judkins right [JR], pigtail, internal mammary artery), as well as various straight tip wires (0.035' soft diagnostic, hydrophilic coated, coronary), the valve crossing was impossible. The patient was switched to general anesthesia, intubated, and transesophageal echocardiography was used. The right femoral vein was canulated, and under the transesophageal echocardiography guidance, the atrial septal puncture was done. The transseptal sheath was introduced into the left atrium, the JR diagnostic catheter was pushed into the left ventricle

(FIGURE 1C), and the aortic valve was easily crossed anterogradely with a 0.035' straight tip diagnostic wire. After navigating the JR catheter into the ascending aorta, the soft guidewire was extended and snared using the snare introduced through the femoral artery. The JR catheter was pulled into the left ventricle along with the snare and guide catheter (FIGURE 1D-1E). It allowed retrograde placement of the stiff wire and implantation of a self-expanding Evolut R valve (FIGURE 1F). The procedure time was 3.5 hours. Control transthoracic echocardiography revealed mild paravalvular leak and reduced mean gradients to 13 mm Hg. The patient was discharged home without complications.

Although ViV TAVI is a widely accepted, less invasive alternative to reoperation in patients with SVD, it can pose some technical procedural challenges. Usually, crossing of the prosthesis is straightforward because in many cases, there is some degree of regurgitation and less calcifications than in the native valve which makes the leaflet opening more symmetrical. In our case, we found the crossing using variety of catheters and wires impossible. Antegrade aortic valve crossing via the transseptal access is a solution in such cases. This approach was used by Cribier et al³ during the first successful TAVI. Alternatively to our method, snaring of the retrograde wire can be done in the ascending aorta with a snare introduced by the antegrade catheter. Other complications associated with ViV TAVI are coronary artery obstruction, elevated

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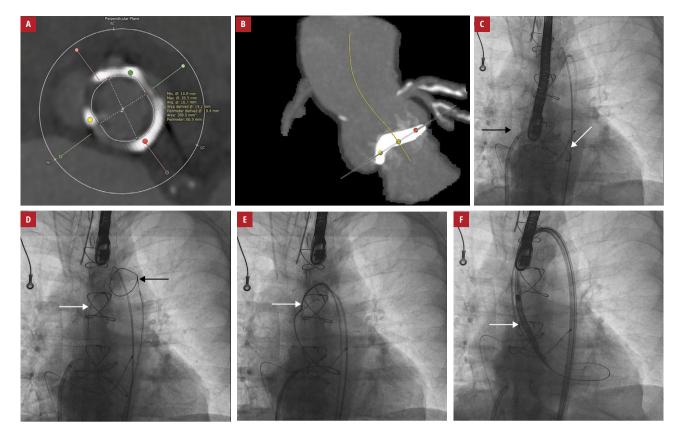


FIGURE 1 A – multislice contrast computed tomography of implanted bioprosthesis and the ascending aorta, the short axis view; **B** – multislice contrast computed tomography of implanted bioprosthesis and ascending aorta, the long axis view; **C** – fluoroscopy of the transeptal sheath (black arrow) in the left atrium and the diagnostic catheter (white arrow) in the left ventricle; **D** – fluoroscopy of the diagnostic catheter (white arrow) in the ascending aorta after crossing the bioprosthesis aortic valve and the snare (black arrow) introduced through the femoral artery; **E** – fluoroscopy of the snared antegrade wire (arrow) in the ascending aorta; **F** – fluoroscopy of the self-expandable valve (Evolut R; arrow) in the implantation position

postprocedural gradient, and malposition.^{2,4} According to Bapat et al,⁵ careful procedure planning includes type, size, true stent internal diameter, and fluoroscopic appearance of the degenerated bioprosthesis. Moreover, selection of the appropriate size and model of the new transcatheter heart valve is important for successful implantation and to reduce complications.

ARTICLE INFORMATION

CONFLICT OF INTEREST None declared.

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