

CASE REPORT

ADVANCED

CLINICAL CASE

Radiofrequency Ablation for Prevention of Outflow Tract Obstruction in Percutaneous Mitral Valve-in-Valve Replacement



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ABSTRACT

Left ventricular outflow tract obstruction (LVOTO) can complicate percutaneous mitral valve replacement and may preclude patients considered high surgical risk from transcatheter therapies. We report a case of mitral valve-in-valve procedure in a patient at high risk for LVOTO. (**Level of Difficulty: Advanced.**) (J Am Coll Cardiol Case Rep 2022;4:1247-1251) © 2022 The Authors. Published by Elsevier on behalf of the American College of Cardiology Foundation. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

HISTORY OF PRESENTATION

An 82-year-old woman presented with worsening dyspnea on exertion for 1 month. She had been doubling her dose of furosemide without relief. Physical examination was grossly normal with no significant murmurs or peripheral edema, but right basilar rales. Blood pressure was 164/69 mm Hg with a regular pulse rate of 66 beats/min, respiratory rate of

18 breaths/min, and oxygen saturation of 98% on room air.

PAST MEDICAL HISTORY

She had a history of mitral valve (MV) endocarditis and underwent bioprosthetic MV replacement in 2015 out of state, and the operative report was not available.

DIFFERENTIAL DIAGNOSIS

Differential diagnosis included congestive heart failure exacerbation, chronic obstructive pulmonary disease exacerbation, coronary artery or valvular heart disease, including bioprosthetic MV regurgitation/stenosis.

INVESTIGATIONS

Recent pharmacologic nuclear stress test showed no ischemia. Labs revealed a high-sensitivity troponin T

LEARNING OBJECTIVES

- To be able to identify patients at excessive risk of surgical MV replacement and increased risk of LVOTO.
- To understand the role of novel therapies for neoLVOT enlargement in patients deemed not to be candidates for invasive surgical strategies or other less invasive therapies for LVOT enlargement such as ASA.

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**ABBREVIATIONS
AND ACRONYMS**

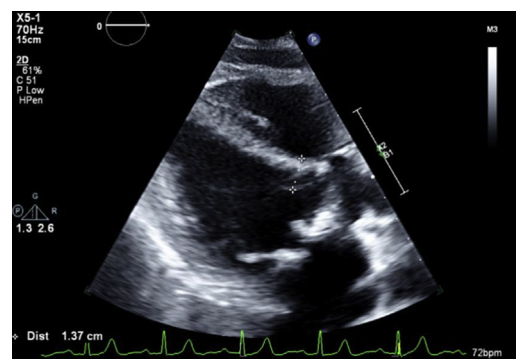
- ASA** = alcohol septal ablation
- CTA** = computed tomography angiogram
- ICE** = intracardiac echocardiography
- LVEF** = left ventricular ejection fraction
- LVOT** = left ventricular outflow tract
- LVOTO** = left ventricular outflow tract obstruction
- MAC** = mitral annular calcification
- MV** = mitral valve
- RFA** = radiofrequency ablation
- TMVR** = transcatheter mitral valve replacement
- TTE** = transthoracic echocardiogram
- VIV** = valve-in-valve

22 ng/L (≤ 14 ng/L) and Pro B-type natriuretic peptide 1,043 pg/mL. Transthoracic echocardiogram (TTE) showed normal left ventricular ejection fraction (LVEF) and severe bioprosthetic MV stenosis with a mean gradient of 12 mm Hg at a pulse rate of 74 beats/min and septal thickness 1.37 cm (Figure 1, Video 1). Transesophageal echocardiogram showed patchy calcification of the leaflets with partial fusion, trace regurgitation, and severe bioprosthetic MV stenosis. Gated cardiac computed tomography angiogram (CTA) revealed high-risk features for left ventricular outflow tract obstruction (LVOTO) with percutaneous mitral valve-in-valve (ViV) procedure with estimated neo-left ventricular outflow tract (LVOT) using a 26-mm valve of 149 mm² (Figure 2A), short prosthesis to septal distance, and relatively unfavorable aorto-mitral angle. Based on the fluoroscopy and preprocedural CT measurements, the valve appeared to be a 27-mm Magna Ease bioprosthetic valve.

MANAGEMENT

The heart team deemed her excessive risk for re-do surgical MV replacement, and septal reduction intervention was recommended for anticipated percutaneous mitral ViV procedure. Coronary angiography showed no obstructive coronary disease; however,

FIGURE 1 TTE Parasternal Long-Axis View Pre-Mitral ViV Procedure

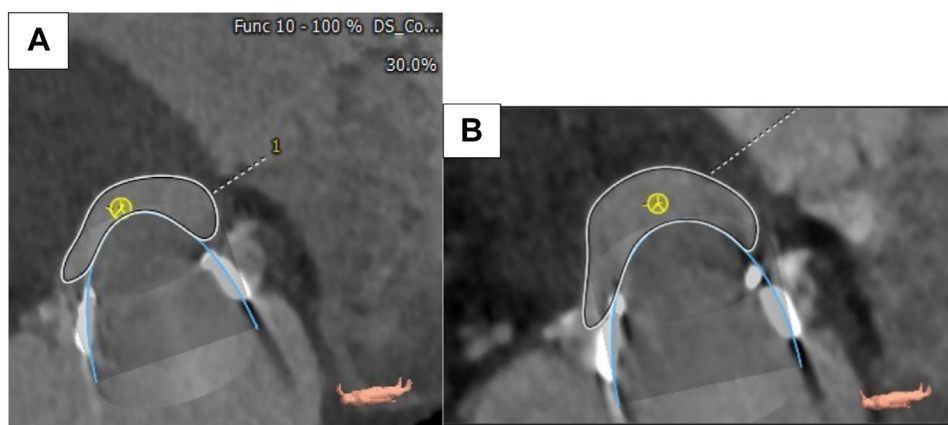


Septal thickness 1.37 cm. TTE = transthoracic echocardiogram; ViV = valve-in-valve.

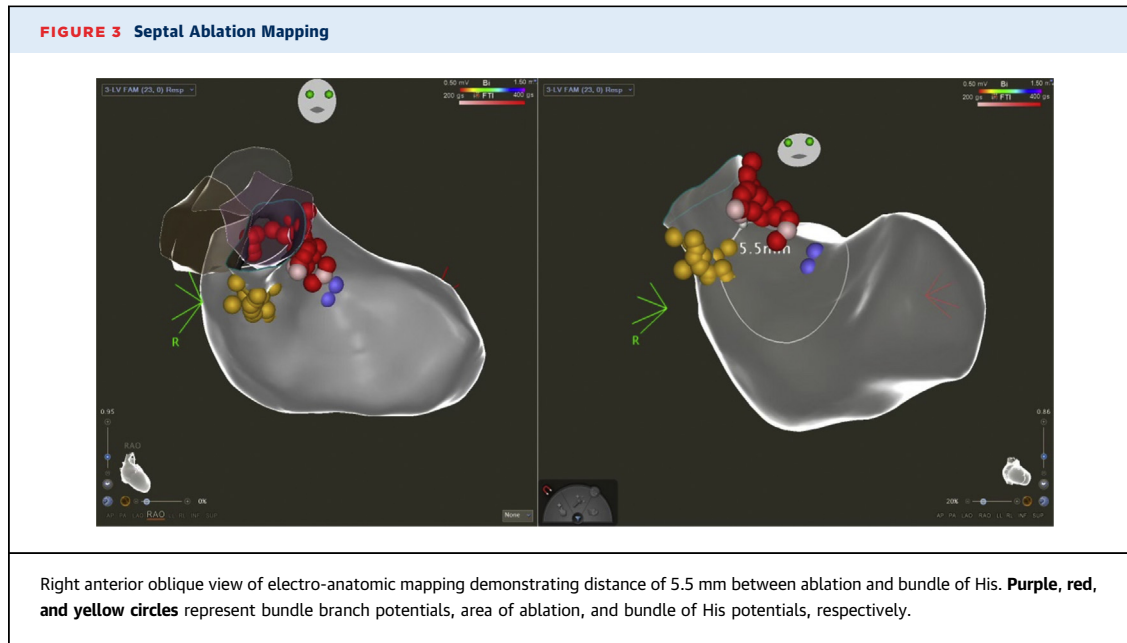
her anatomy was unsuitable for alcohol septal ablation (ASA) (Video 2). Therefore, she was referred for radiofrequency ablation (RFA) of septum to reduce the risk of LVOTO.

After obtaining informed consent, an electrophysiology study was performed with the patient under general anesthesia. Using intracardiac echocardiography (ICE), anatomic mapping of the aortic root, coronary cusps, LVOT, and the left ventricle was performed. Under the guidance of 3-dimensional

FIGURE 2 CTA Chest for Preprocedural Planning Mitral ViV Measurements and Post-RFA for Reassessment of neoLVOT



(A) Estimated neoLVOT at 30% systolic phase, using a 26-mm valve, of 149 mm², short prosthesis to septal distance, and unfavorable aorto-mitral angle. (B) Enlargement of neoLVOT at 40% systolic phase to 232 mm². CTA = computed tomography angiography; LVOT = left ventricular outflow tract; RFA = radiofrequency ablation; other abbreviation as in Figure 1.



electro-anatomic mapping system (CARTO 3, Biosense Webster), using the retrograde aortic approach under therapeutic anticoagulation, a 3.5-mm-tip open-irrigation catheter was used to perform a detailed anatomic and activation map of the His-Purkinje system, left anterior and posterior fascicle, septum, and LVOT. The MV apparatus-septal contact area was marked using ICE onto the anatomic map, and this region became the target for RFA. Avoiding the conduction system, ablation was initiated at 35 watts and 1-minute application time was used per lesion (Figure 3). A total of 47 minutes of energy was applied. Procedural endpoint included complete coverage of the valve-septal contact area, as well as basal septal akinesia on ICE imaging.

Postprocedure TTE showed a small pericardial effusion with normal LVEF 65% to 70%, and she was discharged with plan for elective mitral ViV procedure in 6 to 8 weeks. However, because of severely limiting symptoms, a repeat cardiac CTA was performed in 3.5 weeks confirming enlargement of neoLVOT to 232 mm² (Figure 2B). She subsequently underwent successful transcatheter mitral ViV procedure with 26-mm S3 Edwards Sapien valve via percutaneous transeptal approach (Videos 3 and 4).

DISCUSSION

LVOTO can complicate percutaneous MV replacement and may preclude patients considered high surgical risk from transcatheter therapies. Acute LVOTO after transcatheter mitral valve replacement (TMVR) is

associated with increased mortality. Preprocedure assessment and interventions to prevent this complication are vital. RFA is a novel technique that can be useful in patients unsuitable for other septal reduction strategies.

Repeat operation following MV replacement or repair can be required in a significant proportion of patients,¹ and with the emergence of patients with TMVR who may otherwise be considered high risk for surgical MV repair or replacement are now presented with a new treatment modality. Preprocedural evaluation of LVOT is critical because of the complex

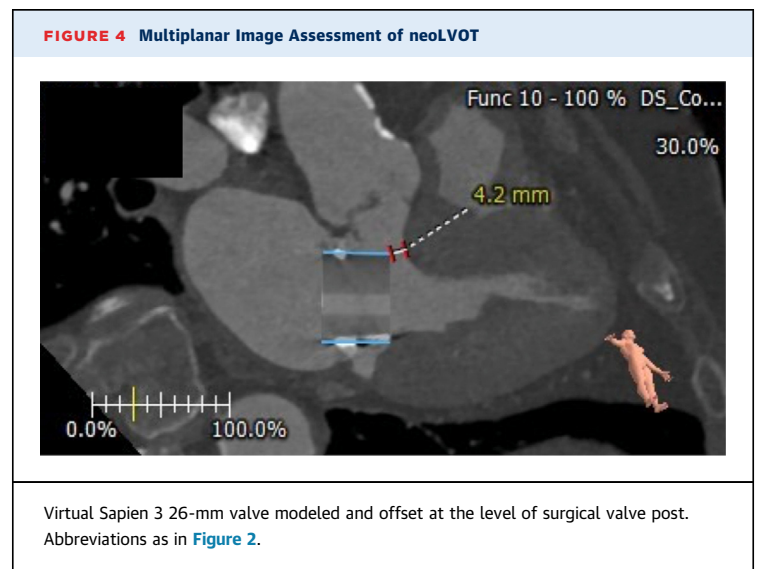
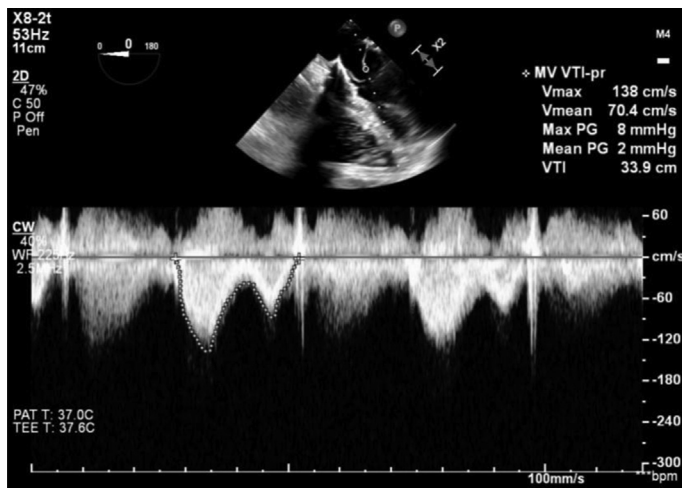


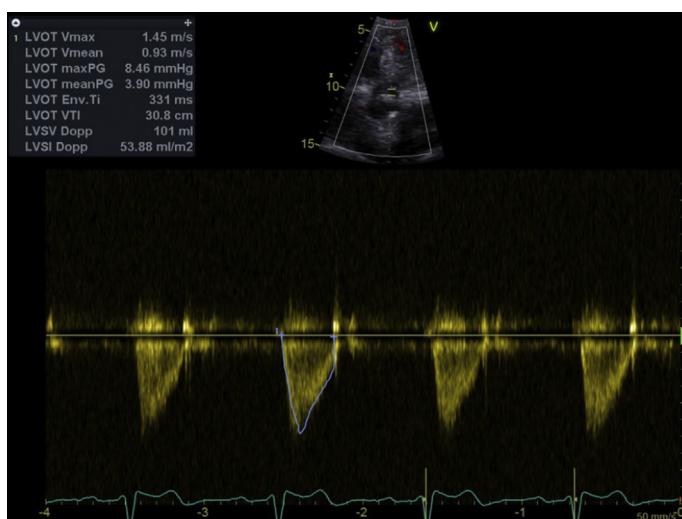
FIGURE 5 TEE Post Mitral ViV



Peak transmitral velocity 1.4 m/s and mean gradient across valve 2 mm Hg. TEE = transesophageal echocardiography; other abbreviation as in Figure 1.

anatomy of the MV, and risk of LVOTO with TMVR. LVOTO may have a low incidence, approximately 7% to 9%,² and with higher rates after valve in mitral annular calcification (MAC)³ or valve-in-ring procedures,⁴ but can be associated with significant peri-procedural morbidity and mortality.⁴ We measured neoLVOT from multiplanar images using a

FIGURE 6 TTE Post Mitral ViV



Abbreviations as in Figure 1.

proprietary software (3Mensio, Pie Medical) as recommended by the consensus document.⁵ Briefly, using multiplanar reconstruction, the segmentation of the basal ring of the mitral prosthesis was performed in mid-late systole. A virtual 26-mm valve was then modeled and offset at the level of surgical valve post and planimetry of the neoLVOT area was performed (Figures 2 and 4).

LVOTO following TMVR in a ViV procedure is caused by pushing the bioprosthetic leaflet into the LVOT covering the open cells of the balloon expandable valve. The size of the neoLVOT and the aorto-mitral angle are associated with increased risk of LVOTO after TMVR.⁶ ASA has been previously used in patients undergoing TMVR at high risk for LVOTO with success⁷; however, not all patients have suitable anatomy to undergo ASA with associated risk of need for permanent pacing and large myocardial infarction. In our patient, significant tortuosity of the left anterior descending artery and the absence of a single suitable proximal septal that supplied the area of interest, made her not a good candidate for ASA. She would have likely required injection of multiple septal branches and would have been at very high risk of pacemaker placement. RFA is a novel technique that has been previously used in patients unsuitable for LVOT enlargement with open procedures, such as surgical myomectomy or laceration of the anterior leaflet (LAMPOON) who require TMVR for native MV disease associated with MAC.⁸ To our knowledge, our case represents the first report of successful use of RFA for neoLVOT enlargement in anticipated TMVR for a mitral ViV procedure.

FOLLOW-UP

After the ViV procedure, postprocedural transesophageal echocardiogram and TTE showed normal LVEF and a mitral bioprosthesis with a mean gradient across the valve of 2 mm Hg with a pulse rate of 77 beats/min and peak LVOT velocity of 1.4 m/s (Figures 5 and 6). She had an uncomplicated post-procedural course and was discharged home the next day. At 4 months, she has had no recurrent hospitalizations and continues to report significant improvement in her ability to perform activities of daily living and overall quality of life.

CONCLUSIONS

Acute LVOTO after transcatheter mitral intervention is associated with increased mortality. Preprocedure assessment and interventions to prevent this complication are vital. RFA is a novel technique that

can be useful in patients unsuitable for other septal reduction strategies.

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The authors have reported that they have no relationships relevant to the contents of this paper to disclose.

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KEY WORDS ablation, mitral valve, valve replacement

APPENDIX For supplemental videos, please see the online version of this paper.