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Commentary: Back to the future: Failed mitral valve bioprosthesis in the setting of mitral annular calcification

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Falasa and colleagues¹ present a case of mitral valve-in-valve (MViV) replacement in a patient with traumatic injuries who 8 years previously had an intra-atrial 25 mm Edwards pericardial mitral valve replacement (MVR) with a Dacron conduit due to extensive mitral annular calcification (MAC).¹ The authors should be applauded for innovative use of MViV with non-conventional anatomy and high surgical risk. Intra-atrial MVR is uncommon; however, a few technical points deserve mention.

Transeptal puncture must be altered from a conventional MViV (2 cm posterolateral to typical location), given the supra-annular prosthetic height and difficulty gaining height with the Edwards delivery system. Posterior bias on the septum remains critical for alignment. Second, navigating across stenotic valves typically requires a steerable guide as used here, facilitated by intraoperative 3-dimensional echocardiography. Finally, given the atrialization of the prosthesis, the risk of left ventricular outflow tract obstruction is low.

Another important consideration relates to patient-prosthesis mismatch. The patient originally received a 25-mm mitral prosthesis. In large analyses of the Valve-in-Valve International Data (VIVID) Registry, MViV with

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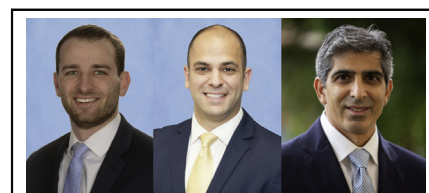
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Received for publication Aug 2, 2021; revisions received Aug 2, 2021; accepted for publication Aug 3, 2021; available ahead of print Aug 6, 2021.

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JTCVS Techniques 2021;10:269-70
2666-2507

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<https://doi.org/10.1016/j.jtc.2021.08.007>



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CENTRAL MESSAGE

This report highlights the safety of transcatheter MViV, even with challenging anatomy. It emphasizes that we need to consider lifetime management to optimize future interventions.

27 mm or smaller surgical valves increased risk of postprocedural stenosis and early failure. Subsequently, residual mitral stenosis carried a significantly greater risk of repeat MVR.²

The intra-atrial/nonannular mitral replacement also raises considerations. The goals of annular debridement of MAC are to (1) create a safe annulus for suture placement/valve seating and (2) create good left ventricular inflow by debriding sufficient mitral valve to avoid stenosis. Moreover, in patients with MAC or rheumatic mitral stenosis, we advocate aggressive debridement of the papillary muscles to minimize the risk of recurrent subvalvular stenosis.³ With the approach taken at the initial operation in this case, there remains risk of stenosis at the native annulus and subvalvular level, which would manifest as early prosthetic failure. No universal surgical approach to MAC has been accepted. Techniques range from complete resection and removal of all calcification,⁴ to using ultrasonic emulsification and aspiration of MAC,⁵ to placing a prosthesis in the left atrium as in this case, to bypassing the valve entirely with an external conduit.⁶

Given surgeons trepidation to aggressively debride MAC, other options have emerged, including: (1) percutaneous valve-in-MAC (ViMAC), (2) surgically deployed balloon expandable SAPIEN ViMAC,⁷ and (3) Tendyne-in-MAC.⁸ A 100-patient cohort who underwent ViMAC experienced significantly lower success rates compared with MViV (74% vs 91%, $P < .001$). ViMAC experienced a 10% rate of left ventricular outflow tract obstruction and 30-day mortality of 21.8%, both lower than previously reported rates for ViMAC, but significantly greater than the

MViV cohort.² Surgical transatrial deployment of balloon expandable valves can minimize left ventricular outflow tract obstruction risk by debridement of the anterior leaflet and septum.⁹ Finally, there is great enthusiasm for the Tendyne transcatheter valve in MAC due to report of 9 compassionate use cases, with only one mortality at 12-month follow-up despite a transapical approach.¹⁰

In summary, this report highlights the safety of transcatheter MViV, even with challenging anatomy. It also emphasizes that we, as surgeons, need to consider lifetime management to optimize future interventions, including adequate debridement, larger prostheses, and continued innovation.

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