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Commentary: Transventricular mitral valve repair: Are we ready to move in reverse to repair P2 prolapse?

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Budra and colleagues¹ report their single center, midterm experience with the DS1000 NeoChord (NeoChord Inc, St Louis Park, Minn) transventricular mitral valve (MV) repair system.¹ This is a beating-heart, transapical device that captures and attaches polytetrafluoroethylene chords to the prolapsing segment of the MV and exteriorizes the chords to the left ventricular (LV) epicardium while providing tension to the repaired prolapse leaflet. They reported outcomes of 89 patients who underwent the procedure between 2011 and 2017, with a wide range of prolapse pathology, including 22 isolated P2 segment prolapses (type A), 47 multisegment posterior leaflet prolapses (type B), 15 bi-leaflet and/or commissural prolapses (type C), and 4 isolated anterior leaflet prolapses (type D). Immediate procedural reduction of mitral regurgitation (MR) to $\leq 2+$ was noted in 98%, and trace to no MR was noted in 69.3%. Operative mortality was 1% due to cardiac tamponade in 1 patient on postoperative day 2 leaving 88 patients for follow-up. Overall freedom from $\geq 2+$ MR at 1, 6, 12, 24, and 36 months was 86%, 77%, 72%, 59%, and 49%, respectively. Acceptable MR reduction was achieved in 82% of patients with type A prolapse, whereas only 20% of patients with type C prolapse were free of $\geq 2+$ MR. The authors found that patients with an increased risk of

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CENTRAL MESSAGE

More data are needed to determine the optimal anatomical characteristics for successful transventricular MV repair. We are encouraged by this new technology and look forward to this burgeoning field.

recurrent MR included type C pathology and patients with pre-existing LV enlargement.

This is an important study and we congratulate the investigators for this seminal work and detailed follow-up. Their results correlate outcomes reported by Colli and colleagues^{2,3} from a multi-center registry demonstrating 74% and 72% freedom from $\geq 2+$ MR, respectively. However, the current study lacks some important technical data such as chordal placement on the LV access site (eg, true-apex vs anterior vs posterior ventricular approach). The access location might not only allow the operator to better reach the prolapsing segment, but also may provide an ability to adjust the degree of and location of displacement of the prolapsing segment, which could influence durability. An example of this is a posterior lateral approach with over-tensioning of the prolapsing segment that may allow the chords to remain under optimal tension even after ventricular reverse remodeling has occurred. This was highlighted in this study where risk of recurrent MR was increased if there was significant LV enlargement before the procedure. Because the chords are attached to the outer ventricular wall farther away from the MV annulus relative to native chordae, reverse remodeling of the ventricle could lead to laxity of the neochords and recurrence of prolapse. An anterior approach, on the other hand, could allow better coaptation of the repaired segment with the anterior leaflet because it draws the posterior leaflet toward the middle of the valve, mimicking an annuloplasty effect. Additional data that

could also determine durability are the number of chords used per prolapsing segment. More chords may better distribute tension and prevent tearing, as well as provide a more uniform correction of prolapse.

The 2 transapical systems currently being investigated in the United States are the DS1000 Neochord (used in the current study) and the Harpoon Medical Device (Edwards Life-Sciences, Irvine, Calif).^{4,5} The ongoing Randomized Trial of the Neochord DS1000 System Versus Open Surgical Repair (ReChord) trial, a randomized trial comparing transapical MV repair with the DS1000 Neochord device with conventional surgery may help determine differences in durability between the 2 groups. The RESTORE trial using the Harpoon Medical Device is a prospective multicenter single-arm trial with a safety end point (comparing with conventional MV surgery as matched via the Society of Thoracic Surgeons database) and an effectiveness end point (comparing echocardiograms for recurrence of MR from patients within a National Institutes of Health trial on mitral and tricuspid valve repair).

There remains skepticism among many cardiac surgeons on the utility of transapical mitral repair; mostly due to the inability to place a band or ring for support of the MV annulus. Furthermore, some may view these data as not at par with some excellent prior surgical series.⁶ However, it

should be noted that there remains a paucity of adjudicated echocardiogram follow-up in the modern surgical era for degenerative surgical repair and a true incidence of freedom from MR has not been well evaluated.

We congratulate Budra and colleagues¹ on their work. Although more data are needed to determine the optimal anatomical characteristics for successful durable transapical MV repair, we are encouraged by this new technology and look forward to this chapter in the field of mitral valve surgery.

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