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Commentary: Striking the right chord

Lawrence M. Wei, MD, and Vinay Badhwar, MD

In this issue of the *Journal*, Budra and colleagues¹ report the first mid-term results of transventricular mitral valve (MV) repair with the NeoChord DS1000 Artificial Chordae Delivery System (NeoChord, St Louis Park, Minn). This device aims to treat primary mitral regurgitation (MR) by implantation of expanded polytetrafluoroethylene (ePTFE) sutures to prolapsing or flail MV leaflet segments via a transapical approach on the beating heart. The highly experienced surgeon coauthors with access to the NeoChord device before current clinical trials were among the first to perform these operations. They are to be highly commended for their diligent and honest efforts to explore this novel technology.

The preferred therapy for severe primary degenerative MR is MV repair, receiving class 1 guideline recommendation.² Surgical MV repair technique has evolved to include a multitude of options, including implantation of artificial chordae, all with highly reproducible outcomes and durability, with repair rates >95% for focal disease in experienced centers.³⁻⁶ Multiple centers regularly use minimally invasive approaches, including robotic assistance, to achieve excellent short-term outcome, >95% repair, and 1-year durability with less than mild MR while eliminating the morbidity of sternotomy.^{4,5} Long-term freedom from recurrent moderate or severe MR following MV repair has been reported as high as 87.5% at 20 years.⁶ Newer technologies, such as the NeoChord device, laudably seek to achieve similar outcomes without the use of cardiopulmonary bypass. Implantation of chordae with the NeoChord device has achieved proof-of-concept as feasible, with



Vinay Badhwar, MD (left), and Lawrence M. Wei, MD (right)

CENTRAL MESSAGE

Mitral therapy with the NeoChord device has a high rate of failure at mid-term follow-up, even in experienced hands.

composite short-term survival and freedom of MR of 84%, but longer-term outcomes are not yet available.^{7,8}

Budra and colleagues provide longitudinal outcomes on 88 consecutive patients who underwent transventricular MV repair with the NeoChord device between 2011 and 2017 with a median follow-up of 42 months. Only 59 patients (67%) were free of recurrent MR >2+, and even the patients with the simplest pathology (isolated P2 prolapse) had freedom from MR >2+ of only 82%. These results did not improve with increasing experience of the surgical team. The authors describe several mechanisms of failure, including rupture of native or artificial chordae, neochordal elongation or dehiscence, and combinations of these mechanisms. Mitral pathology (rather than focal disease) and a dilated left ventricle were 2 factors that significantly increased the risk of MR progression. It might be hypothesized that following NeoChord implantation in the absence of annuloplasty, initial partial relief of MR resulted in remodeling of dilated ventricles and the chordae became elongated, resulting in recurrent MR.

This important experience demonstrates that MV therapy performed with the NeoChord device has an unacceptably high failure rate compared with conventional minimally invasive techniques. The authors are to be congratulated for their transparency and honesty in reporting these results despite being at the forefront of this technology and investing great time and effort in its development. While the outcome of a US randomized clinical trial is still pending, when faced with choices on the treatment of primary MR,

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it appears that even in experienced hands that performance of this technology fails to strike the right chord.

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