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Commentary: The pericardial autologous solution

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Infective aortic valve endocarditis with the involvement of mitro-aortic continuity and the anterior mitral valve leaflet is a severe and devastating disease associated with high mortality and morbidity. It occurs in 10.6% of cases of left-sided infective endocarditis, and early mortality ranges from 10% to 32%.¹⁻³ Surgical operation is complex and requires radical debridement and reconstruction. Several surgical techniques have been proposed.⁴ Replacement of aortic and mitral valve with reconstruction of the mitro-aortic continuity is the most common technique of choice, but the presence of 2 prosthetic valves may increase the risk of recurrent infection. Homografts and stentless xenografts represent excellent solutions, offering a low rate of recurrent infection and a low transvalvular gradient.^{5,6} However, they are technically more demanding, and future reoperation is a great challenge due to calcified structural degeneration. A monoblock aorto-mitral homograft has been also suggested as a surgical option for extensive bivalvular infective endocarditis.⁷

In this issue of *JTCVS Technique*, Benedetto and colleagues⁸ propose a “biological” solution for the treatment of aortic and mitral endocarditis. Using autologous pericardium, the aortic valve is replaced according to the Ozaki procedure, and the mitral valve is repaired with a patch via a transaortic approach. This strategy is interesting and combines the benefits of a stentless valve without the use of prosthetic material. In addition, the transaortic approach avoids left atrial opening with consequently less surgical



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CENTRAL MESSAGE

Aortic valve neocuspidualization and transaortic mitral valve repair with autologous pericardium is a new technique for treating aortomitral endocarditis.

trauma and bleeding. The Ozaki procedure was developed mainly to treat degenerative aortic valve disease in patients with a small aortic annulus and patients who had rejected a mechanical valve.⁹ The indications were then expanded for aortic regurgitation and infective endocarditis. Midterm results have been excellent, with a 4.2% cumulative incidence of reoperation at 8 years reported. Nevertheless, 13 of 15 patients requiring reoperation (87%) had infective endocarditis, with a rate of 0.3% per patient-year. The annual incidence rate for prosthetic valve endocarditis ranges between 0.12% and 0.35% per patient-year, raising some concerns regarding the benefit of aortic valve neocuspidualization in infective endocarditis.^{10,11} In addition, at 8-year follow-up, patients at risk for cumulative incidence of reoperation were only 54 out of 850, so a higher incidence of endocarditis might be expected.⁹

Finally, although autologous pericardium has good long-term durability in term of calcification, stiffness, and tears, most patients undergoing the Ozaki procedure will require a new operation at some point in their life. Reoperation is technically more demanding than primary operation, because of adhesions and the risk of iatrogenic injury to cardiac structures, especially in patients with less pericardium. Furthermore, the lack of prosthetic support might preclude the transcatheter aortic valve replacement (TAVR) approach. Several studies have shown that valve-in-valve TAVR is a safe and effective procedure with similar mortality as surgical AVR, at least in the short term.¹² To date, no data have been reported on TAVR in a AVNeo (Ozaki) procedure. Regardless, my congratulations to Benedetto and colleagues for

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adding a new technique to the list of surgical treatments for aortomitral valve endocarditis: the pericardial autologous solution.

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