

Resurgence of the Ross procedure

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Innovation in transcatheter aortic valve replacement (TAVR), which has virtually transformed the care of patients with aortic stenosis (AS), has fueled an intense interest in the management of patients with aortic valve disease. Indications for TAVR have expanded to lowrisk symptomatic patients with AS, and clinical trials are currently in progress for asymptomatic patients with severe AS, testing the strategy of pre-emptive TAVR versus waiting for symptom onset or other indications for valve replacement.

However, expanding the eligibility for TAVR to lower risk, less symptomatic patients goes against the uncertainties regarding durability of transcatheter biologic valves, as such patients are generally much younger than those at higher surgical risk, for whom TAVR was initially targeted towards (1). The high rate of pacemaker requirement after TAVR is another limitation that must be acknowledged when TAVR is considered in younger lowrisk populations. Thus, the vast majority of patients with AS under the age of 60 requiring aortic valve replacement (AVR) remain within the realm of surgical intervention. In addition, there is no expectation in the near future of a transcatheter solution for patients with aortic regurgitation, who tend to be younger when AVR is warranted than patients with AS. Similarly, although there is emerging data on successful TAVR in patients of suitable age with AS and bicuspid aortic valves, many, if not most patients with bicuspid valves who require AVR fall under the age threshold below which surgery remains the best option.

In the midst of the excitement and din generated by TAVR, discussions concerning the most appropriate surgical valve substitute for those under the age of 60 years, has essentially been sidelined—yet this has been a debatable topic for decades. While it is clear that most cardiologists and cardiac surgeons shy away from recommending bioprosthetic heterografts and homografts for young adults in the 20–40 years age range due to accelerated structural valve deterioration, the long-term consequences of lifelong anticoagulant therapy with a vitamin K antagonist in patients with mechanical prostheses is equally daunting. These same issues carry over to patients in ages 40–60 years. There is no perfect valve substitute for young and middleaged patients with aortic valve disease. Against this continuing conundrum, the time is ripe for discussions on broadening the use of the Ross operation as a preferred option for selected young individuals who require AVR (2).

Following initial reports of success of the Ross operation, the procedure was adopted by many centers in the United States in the 1980s and 1990s. The initial enthusiasm was ultimately dampened by the procedure's technical demands, lengthy cardiopulmonary bypass times and, in some centers, poor long-term outcomes with deterioration of either the aortic autograft or the pulmonic replacement valve (heterograft or homograft) or both, and therefore, the need for complex repeat operations. Currently, there are vanishingly few expert centers remaining in the United States, resulting in limited training opportunities for junior surgeons, and the Ross procedure has thus virtually disappeared from the surgical armamentarium. It receives limited or absent endorsement in current societal guidelines, with a class IIb recommendation in the American College of Cardiology (ACC)/American Heart Association (AHA) guidelines for valvular heart disease only for young patients in whom anticoagulation is undesirable or contraindicated and is performed only at a comprehensive valve center by surgeons experienced in this procedure (3). Additionally, there is no mention in the European Society of Cardiology (ESC)/European Association for Cardio-Thoracic Surgery

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(EACTS) guidelines for valvular heart disease (4), and only passing mention with no specific recommendation in the updated ESC guidelines for adult congenital heart disease (5). It is likely that few practicing cardiologists in the United States can identify a Ross center of excellence for referral of patients, and the Ross procedure is thus beyond the bandwidth of most. The result is lack of access for suitable patients who would otherwise be excellent candidates for the only AVR operation that offers the patient a living aortic valve substitute. This represents a major lost opportunity (6).

However, several centers of excellence in Europe, Canada, and Australia have kept the flame alive and devoted the necessary time, resources and personnel to achieve exceptional expertise in perfecting the Ross operation, with long-term survival rates exceeding those of conventional AVR and with durability of both replacement valves, therefore yielding low rates of re-operation. A decade ago, a randomized clinical trial demonstrated superior tenyear survival with the Ross operation, as compared to a homograft AVR (7). While this is the only prospective randomized trial, more recent observational studies have reported higher long-term (15-20 years) survival rates in patients undergoing AVR with the Ross operation than with mechanical prosthetic valves (8-11) and, as anticipated, it presents lower risks of thromboembolic and bleeding complications related to chronic anticoagulation. Although the data is limited, the Ross operation also appears to provide superior survival and durability compared to AVR using a bioprosthetic heterograft (8,12). Additional prospective randomized trials are needed to validate these observational data, but it is apparent that the Ross operation is the only AVR procedure that results in long-term survival equivalent to that of an age and sex matched normal population (7,8,12,13).

These recent long-term outcome data has stimulated renewed interest in the Ross operation (2,14). The published data indicate what can be achieved when surgical teams operate in centers devoted to perfecting this procedure in appropriately selected patients. This experience is currently not generalizable. Whether these results can be broadened beyond this small network of specialized centers to other academic institutions and ultimately to larger community hospitals remains to be determined. There are a number of obstacles that need to be surmounted to achieve this potential. First and foremost is the need for educational and training opportunities for both cardiologists and cardiac surgeons, including surgical proctoring and hands-on experience. In the United States and other countries, regulatory issues and barriers related to insurance and reimbursement may limit the referral of patients from one hospital system to another in a more distant region. Further investigation is required to enhance longevity of the autograft and pulmonary homograft, such as strategies for optimal post-operative blood pressure control and optimal anti-inflammatory management. A limited supply of homografts may also slow progress in expanding access to the Ross operation; whether this can be overcome with biologically engineered human valves is yet to be determined (14). There are thus a number of technical, clinical, educational and regulatory challenges. There is great potential, but much more work needs to be done to realize a true Ross resurgence.

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Footnote

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