Management of elective aortic valve replacement over the long term in the era of COVID-19

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As numerous patients await elective aortic valve intervention for aortic stenosis (AS) during this unprecedented outbreak of severe acute respiratory syndrome coronavirus 2 in 2019 (COVID-19), there is serious concern for the possibility of morbidity and mortality during prolonged wait-times. During this time period, the American College of Surgeons and Center for Disease Control released their recommendation to reschedule elective surgeries and to shift elective inpatient diagnostic and surgical procedures to the outpatient setting [1]. Accordingly, the Centers for Medicare & Medicaid Services provided a framework to further group elective surgeries into levels of urgency [2]. A tiered framework ensures that we are providing surgical services and procedures to those patients in whom the risk of delaying a procedure may lead to significant morbidity or mortality. As cardiovascular healthcare providers caring for patients with AS, we must now differentiate cases at a higher tier of urgency from the rest.

Current guidelines suggest treatment of severe AS when symptomatic (angina, heart failure and/or syncope), and there is now growing data to support intervention even before symptom onset [3]. Though most hospitals are equipped for urgent/emergency cases, the majority of aortic interventions are done on an elective basis, and therefore have been postponed due to the COVID-19 outbreak. However, there is a significant relationship between aortic valve replacement (AVR) wait-time and mortality as well as hospital readmission [4, 5]. Delaying AVR has been associated with poor operative outcomes and risk of mortality during the waiting period [6, 7]. Additionally, hospitalizations during wait-time and urgent/emergency AVR have been associated with worse short- and long-term outcomes [8]. It is, therefore, essential that our established multi-disciplinary heart team carefully reviews each patient individually, and determines who would likely benefit from an ‘early elective’ strategy. Timing of intervention when balanced with healthcare resources has not been a major focus among our current guidelines. Making with patients have been applied, particularly to patients with less cardiac reserve, it is imperative to negate the effects of AS before an event.

Symptom severity is generally the largest driver for an earlier AVR strategy. Patients with New York Heart Association (NYHA) IV symptoms and/or syncope clearly portend a worse prognosis than less symptomatic patients [9]. These patients should, therefore, be treated in a timely manner. The presence of angina is always concerning, although the natural history of patients with severe AS suggests that angina is not as ominous a sign as syncope. Yet, since the prevalence of concomitant coronary artery disease is as much as 50%, earlier strategies in patients with severe or unstable angina should be strongly considered [10].

There is a paucity of literature addressing clinical risks (i.e. non-invasive data, comorbidities and demographics) that are associated with higher clinical events during wait-time. Factoring in the severity of AS into our equation is important, as we know that patients who meet these criteria are at higher risk. Asymptomatic patients with indexed aortic valve area <0.4 cm² have a higher risk of events prior to intervention, and a peak jet velocity >5 m/s is an independent predictor of mortality [11, 12]. Another important echocardiographic finding is impaired ejec- tion fraction (EF). Patients managed conservatively with an impaired EF (EF < 60%) have been independently associated with poorer long-term outcomes, whilst an earlier AVR strategy has improved outcomes [13]. In these patients with less cardiac reserve, it is imperative to negate the effects of AS before an event.

Furthermore, the decision for type of intervention may also have significant impact. Transcatheter AVR (TAVR) indications have been expanded to include patients who are at low risk for surgical AVR complications. Strategies involving shared-decision-making with patients have been applied, particularly to patients <70 years old, remembering that the average age in the low-risk trials was 73 ± 6 years old. From these trials, TAVR did result in a shorter index hospitalization compared to surgical AVR (3 vs 7 days) [14]. Understanding the dynamic constraints on healthcare systems, minimalist TAVR can potentially help to further reduce post-care utilization of resources and allow early patient recovery at home [13]. Balloon aortic valvuloplasty as a bridge to TAVR has also shown an improved safety profile in the contemporary...
The era of TAVR [15]. Its role as a bridge for patients with anatomical challenges for TAVR, whether non-transfemoral or requiring adjunc- tive techniques (such as coronary protection with snorkel stenting or leaflet laceration, etc.) may be more important now than ever.

As the treatment for aortic intervention continues to improve both from surgical techniques to transcatheter technology, we need more focus on the timing and application of our interventions. For now, symptoms and echocardiographic criteria can drive how we deliver therapy (Fig. 1), but we also need to factor in other clinical information. Many questions remain; should baseline comorbidities such as chronic lung disease, renal impairment and Society of Thoracic Surgery PROM score push us to act more quickly, whereas other factors (i.e. immunocompromised patients) push us to delay? Do NYHA IIIb symptoms herald worse outcomes during waiting than NYHA IIIa? Should patients with NYHA III symptoms receive an ‘early elective’ strategy versus a patient with NYHA II symptoms? Do we need to factor in other non-invasive data such as degree of left ventricular hypertrophy, and biomarkers (including elevated natriuretic peptide) which may portend a higher rate of events during wait time? [16]

Should discussions with the multi-disciplinary team increase considerations for utilizing TAVR in the very low-risk AS patient and balloon aortic valvuloplasty as a bridge to TAVR in efforts to decrease length of stay, hospital resource utilization and patient/family exposure COVID-19? Further investigation is clearly needed, and we must work on developing a risk stratification system for suggested wait-times. Until then, we must balance the risk of delaying therapy against the availability of hospital resources and potential exposure of COVID-19 on a case-by-case basis.

**Conflict of interest:** Dr Chad A. Kliger is a consultant and receives speaking honoraria from Edwards Lifescience and Medtronic. All other authors declare no conflict of interest.

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