

Edwards SAPIEN 3 Ultra valve

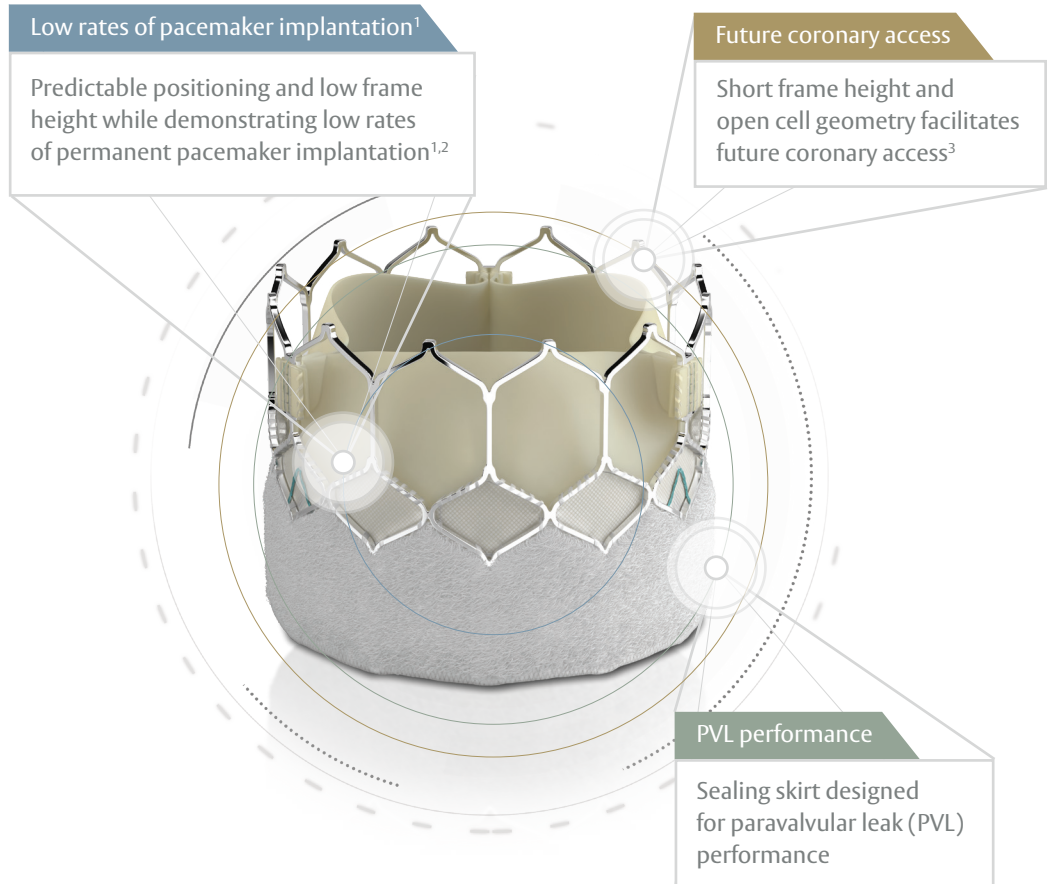
# Now approved for low-risk severe aortic stenosis

## Low rates of pacemaker implantation<sup>1</sup>

Predictable positioning and low frame height while demonstrating low rates of permanent pacemaker implantation<sup>1,2</sup>

## Future coronary access

Short frame height and open cell geometry facilitates future coronary access<sup>3</sup>



Low-risk patients deserve the lowest-risk procedure

Give your patients SAPIEN 3 Ultra TAVI



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# Low-risk patients deserve the lowest-risk procedure

SAPIEN 3 Ultra demonstrates low rates of death and stroke at 30 days.<sup>1</sup>

Your valve choice today can impact your patients' future



## Coronary access

- Post-TAVI patients are at risk of developing coronary artery disease (CAD) following implantation<sup>4</sup>



## Paravalvular leak

- Paravalvular leak (PVL) has been associated with long-term patient cardiovascular mortality after TAVI<sup>5</sup>



## Conduction disturbances

- Long-term complications from pacemaker implantation negatively impact patient quality of life and mortality<sup>6</sup>

### References:

1. STS/ACC TVT Registry, Data on File at Edwards Lifesciences.
2. Webb J, Gerosa G, Lefèvre T, Leipsic J, Spence M, Thomas M, et al. Multicenter evaluation of a next-generation balloon-expandable transcatheter aortic valve. *J Am Coll Cardiol.* 2014;64(21):2235–43.
3. Tarantini G, Fovino LN, Leprince P, et al. Predictors, feasibility and outcomes of coronary intervention up to 3 years after TAVI with a balloon-expandable valve: results from a large European multicenter registry. Presented at: ESC Congress 2019; September 2019; Paris, France.
4. Yudi MB, et al. Coronary Angiography and percutaneous coronary intervention after transcatheter aortic valve replacement. *JACC Vol 71, No 12, 2018.*
5. Kapadia-, S. et al. (2015). 5-year outcomes of transcatheter aortic valve replacement compared with standard treatment for patients with inoperable aortic stenosis (PARTNER 1): a randomised controlled trial. *The Lancet.* 385 (9986), 2485–91.
6. Fujita B, et al. Impact of new pacemaker implantation following surgical and transcatheter aortic valve replacement on 1-year outcome. *Eur J Cardiothorac Surg* 2019.

Available in 20, 23, and 26 mm sizes.

For professional use. For a listing of indications, contraindications, precautions, warnings, and potential adverse events, please refer to the Instructions for Use (consult [eifu.edwards.com](http://eifu.edwards.com) where applicable).

Edwards devices placed on the European market meeting the essential requirements referred to in Article 3 of the Medical Device Directive 93/42/EEC bear the CE marking of conformity.

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