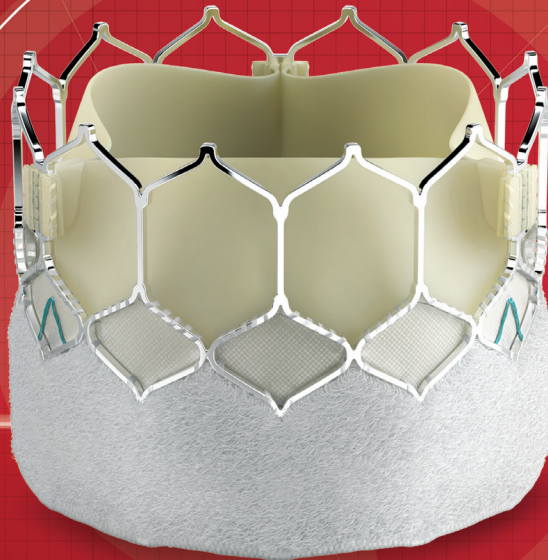
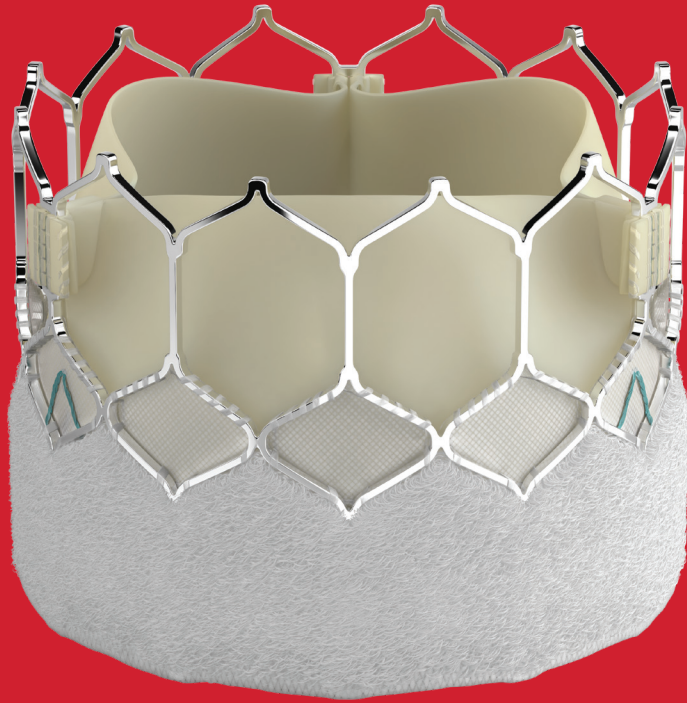


Introducing the **SAPIEN 3 Ultra valve**

Edwards Engineering for Tomorrow's TAVI

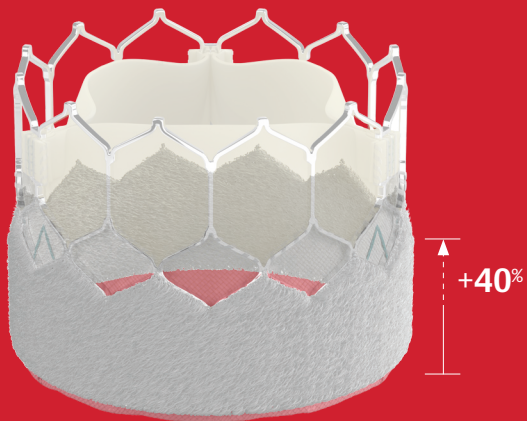




An evolution of the valve you know and trust

**Edwards Lifesciences' commitment to innovation
continues with the SAPIEN 3 Ultra valve.**

Built upon the proven SAPIEN platform, the SAPIEN 3 Ultra transcatheter heart valve was designed with patient needs in mind.

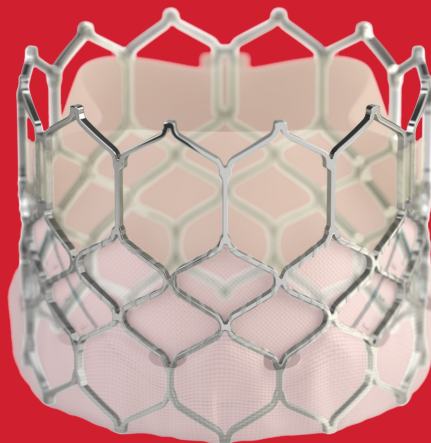


Sealing skirt technology

SAPIEN 3 Ultra valve features an approximately 40% taller*, textured PET outer skirt

Enhanced frame design

Frame geometry designed for high radial strength; short frame height facilitates future coronary access²



- **Valve tissue**

Uses same bovine pericardium tissue and process as the SAPIEN 3 valve

*Compared with the Edwards SAPIEN 3 valve. Available in 20, 23 and 26mm sizes.



For predictable, proven results

Edwards Commander Delivery System

- Stable, precise deployment delivering 99% first-time accuracy¹
- Low delivery profile
- Dual articulation for coaxiality
- Optimal positioning control

Edwards eSheath Introducer Set

- 14F compatibility
- Dynamic Expansion Mechanism (DEM) for low-profile access
- Hydrophilic coating designed for ease of entry



The SAPIEN 3 Ultra valve is available with the Edwards Commander delivery system and Edwards eSheath Introducer helping you achieve consistent performance and controlled deployment.



Tomorrow's TAVI is here today.

- Future TAVI patients need systems designed for performance. Choose SAPIEN 3 Ultra TAVI for your patients and their severe aortic stenosis.

References:

1. The PARTNER II Trial intermediate-risk cohort 30-day unadjusted clinical event rates for TAVR with the SAPIEN 3 valve, AT population (n=1077).
2. 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.

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 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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