SAPIEN 3 Transcatheter Aortic Valve Replacement Compared with Surgery Intermediate-Risk Patients: A Propensity Score Analysis

Vinod H. Thourani, MD on behalf of The PARTNER Trial Investigators

Edwards Lifesciences v. Boston Scientific Scimed IPR2017-00060, U.S. Patent 8,992,608 Exhibit 2052

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Disclosure Statement of Financial Interest

Vinod H. Thourani, MD

Within the past 12 months, I or my spouse/partner have h interest/arrangement or affiliation with the organization(s)

Affiliation/Financial Relationship

- Grant/Research Support
- Consulting Fees/Honoraria
- Major Stock Shareholder/Equity

Company

- Boston Scientific, Clare Lifesciences, Medtroni
- Abbott Vascular, Edwa St. Jude Medical
- None



Background

- The 3rd generation SAPIEN 3 transcatheter heademonstrated improved clinical outcomes in higher patients at 1 year.
- The 30-day outcomes in intermediate-risk patie with SAPIEN 3 in PARTNER 2 were...

All-cause Mortality: 1.1%

Disabling Stroke: 1.0%

 $-PVL \ge Moderate:$ 3.8%

 There is a paucity of longer-term data in interm patients with SAPIEN 3 and there have been n comparisons with surgery in intermediate-risk p



Purpose

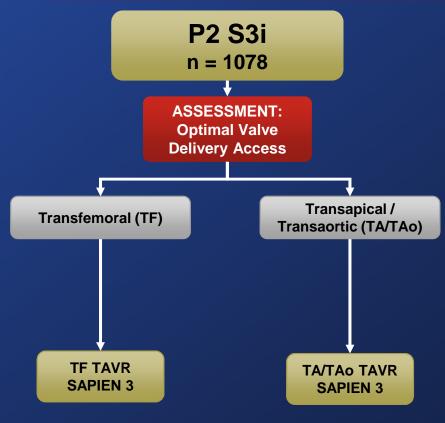
- To evaluate the 1-year clinical and echo out SAPIEN 3 TAVR in intermediate-risk patient
- To compare these intermediate-risk patient using SAPIEN 3 TAVR with surgery results intermediate-risk patients from the PARTNE using a rigorous pre-specified propensity so analysis.

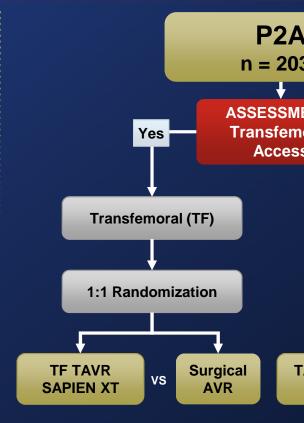


The PARTNER 2A and S3i Trials Study Design

Intermediate Risk Symptomatic Severe Aortic Stenos

Intermediate Risk ASSESSMENT by Heart Valve Team







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