Outcomes



#### **Definitive outcomes**

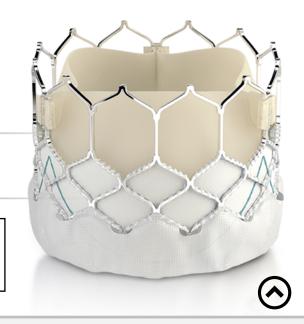


All-cause Mortality<sup>†</sup>

Disabling Stroke<sup>†</sup>

75<sup>%</sup> Lower than surgery

**Edwards Lifesciences v. Boston Scientific Scimed** IPR2017-00060, U.S. Patent 8,992,608 Exhibit 2033





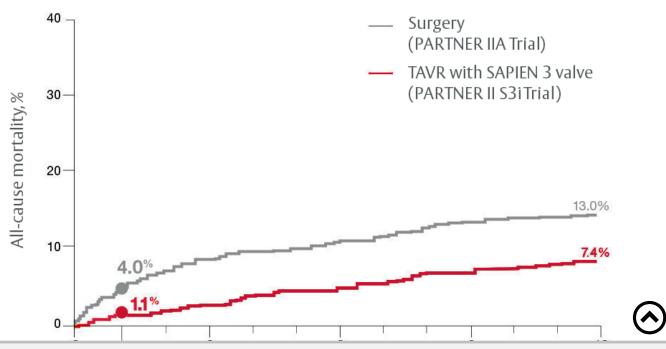
Outcomes

Design

Videos

Edwards SAPIEN valves are the only transcatheter heart valves approved for use in intermediaterisk patients in the United States

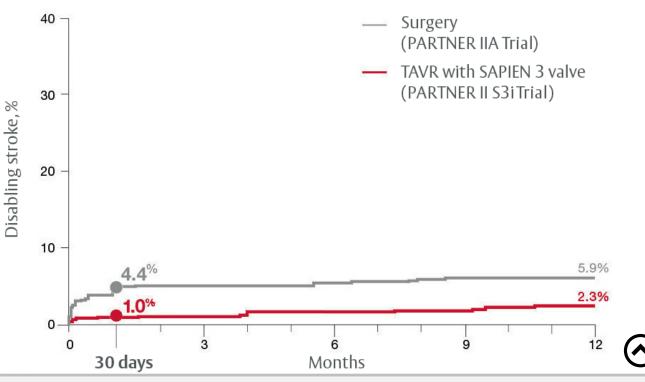
## The PARTNER II Trial intermediate-risk cohort clinical outcomes All-cause mortality<sup>‡</sup>





No. at risk:	Outcomes	Design	Videos		
Surgery	944	859	836	808	795
SAPIEN 3 TAVR	1,077	1,043	1,017	991	963

## The PARTNER II Trial intermediate-risk cohort clinical outcomes Disabling stroke<sup>‡</sup>





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SAPIEN 3 TAVR Outcomes 1,033 1,008 Videos 984 953

<sup>‡</sup>The PARTNER II Trial intermediate-risk cohort unadjusted clinical event rates, AT population.



(http://edwardsprod.blob.core.windows.net/media/Default/sapien3/the%20lancet%20article2.pdf)



SAPIEN 3 Valve Patient Brochure

(http://edwardsprod.blob.core.windows.net/media/Default/sapien3/sapien%203%20 patient brochure%20 ir%20 doc-0041194b%20.pdf)





(httn://edwardsnrod.hlob.core.windows.net/media/Default/saniena/final%20nn--us-



Outcomes

Videos



The PARTNER Trials represent the largest, most rigorous comparative body of evidence in the history of aortic valve replacement

## Transformational design

### Outer sealing skirt >

Low profile access >

Frame design >

Proven valve tissue ▶

Virtually eliminates moderate or greater paravalvular leak\*



\*PARTNER II Trial intermediate-risk SAPIEN 3 valve cohort, core lab assessed paravalvular leak at 30 days, n=1,106

# Complete range of valve sizes

20 0 23 0 26 0











# DOCKET

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