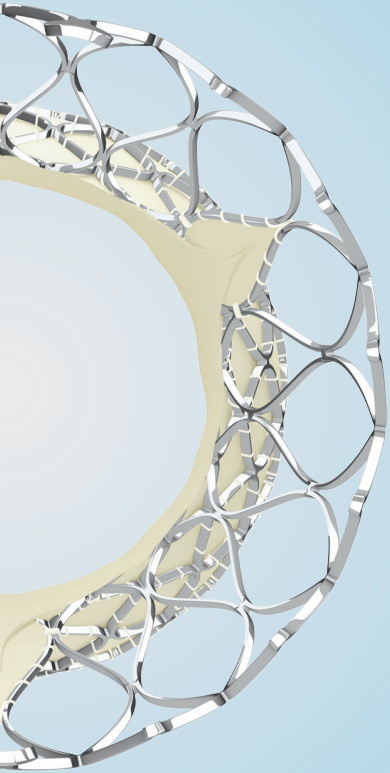


EVOLUT™ TAVI
HEALTH CANADA LICENCED FOR LOW RISK
AND ALSO
FIRST AND ONLY TAVI PLATFORM
HEALTH CANADA APPROVED FOR TREATMENT OF PATIENTS WITH SEVERE BICUSPID AORTIC STENOSIS*

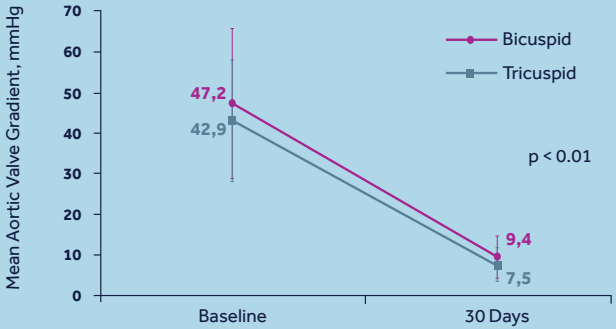
* Evolut™ TAVI Platform is indicated for treatment of patients with severe Bicuspid Aortic Stenosis who are at intermediate or greater risk for SAVR.

Medtronic
Further, Together

WHY EVOLUT™ IN BICUSPID PATIENTS?
DESIGNED FOR EXCELLENT PATIENT OUTCOMES

The Evolut™ platform offers patients with **BICUSPID** aortic valve disease excellent hemodynamics.[3]

MEAN AORTIC VALVE GRADIENT[2]

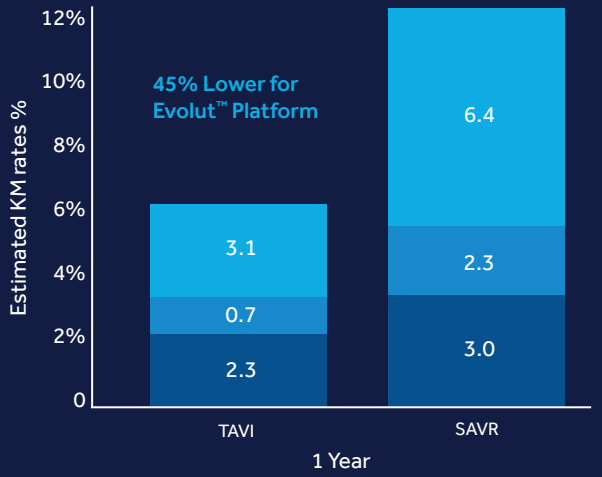


Number of Echos:		
Bicuspid	187	116
Tricuspid	6426	4267

The supra annular design of the Evolut™ TAVI platform allows for exceptional hemodynamics in challenging, bicuspid anatomies.[4]

EVOLUT™ TAVI COMPARED TO SAVR

TAVI showed lower mortality, fewer disabling strokes, and fewer heart failure rehospitalizations compared to SAVR.[1]



- HF Hospitalization
- Disabling Stroke
- Death

Difference = +4.5%
P = 0.002

TAVI 5.6% SAVR 10.2%
Composite Rates

SUPERIOR HEMODYNAMICS IN LOW RISK PATIENTS[2]

Evolut™ TAVI is currently **THE ONLY** TAVI device to demonstrate hemodynamics superiority in a low risk clinical trial vs. SAVR at one year.[2]

SUPERIOR
EOAs

Evolut™ TAVI
2.3 cm²
VS.
SAVR 2.0 cm²

SUPERIOR
Gradients

Evolut™ TAVI
8.6 mm Hg
VS.
SAVR
11.2 mm Hg

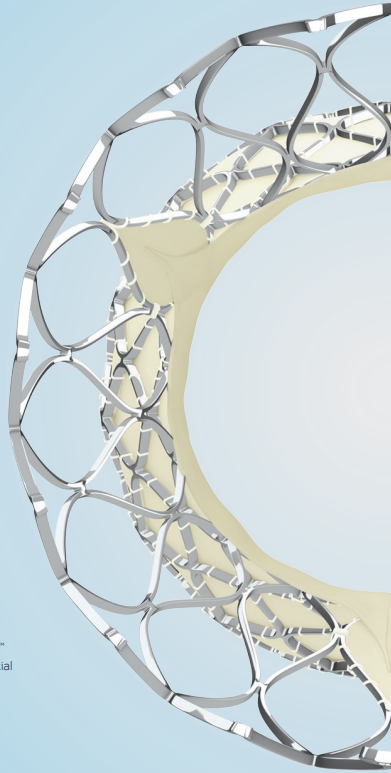
15%
LARGER EOAs

23%
LOWER GRADIENTS

HEMODYNAMICS **MATTER**

Patients benefit from larger EOAs, which decreases the risk of Patient Prosthesis Mismatch (PPM) and improves exercise capacity, helping them return to an active life.[5,6]

PPM CALCULATOR TOOL
PPMMATTERS.COM



Less Patient Prosthesis Mismatch, Better Patient Outcomes.[7**]

** Severe PPM is associated with increased mortality and HF rehospitalizations at 1 year.

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4. Philip F, Faza NN, Schoenhagen P, et al. Aortic annulus and root characteristics in severe aortic stenosis due to bicuspid aortic valve and tricuspid aortic valves: implications for transcatheter aortic.
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7. Herrmann HC, Daneshvar SA, Fonarow GC, et al. Prosthesis-Patient Mismatch in 62,125 Patients Following Transcatheter Aortic Valve Replacement: From the STS/ACC TVT Registry. J Am Coll Cardiol. Published online September 18, 2018.

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Please reference the CoreValve™ Evolut™ R and CoreValve™ Evolut™ PRO instruction for use for more information regarding indications, warnings, precautions, and potential adverse events. The commercial name of the Evolut™ R device is Medtronic CoreValve™ Evolut™ R System, and the commercial name of the Evolut™ PRO device is Medtronic Corevalve™ Evolut™ PRO System. See the CoreValve™ Evolut™ R and CoreValve™ Evolut™ PRO device manuals for detailed information regarding the instructions for use , indications, contraindications, warnings, precautions, and potential adverse events. For further information, contact your local Medtronic representative and/or consult the Medtronic website at medtronic.eu. For applicable products, consult instructions for use on manuals.medtronic.com. Manuals can be viewed using a current version of any major internet browser. For best results, use Adobe Acrobat® Reader with the browser.