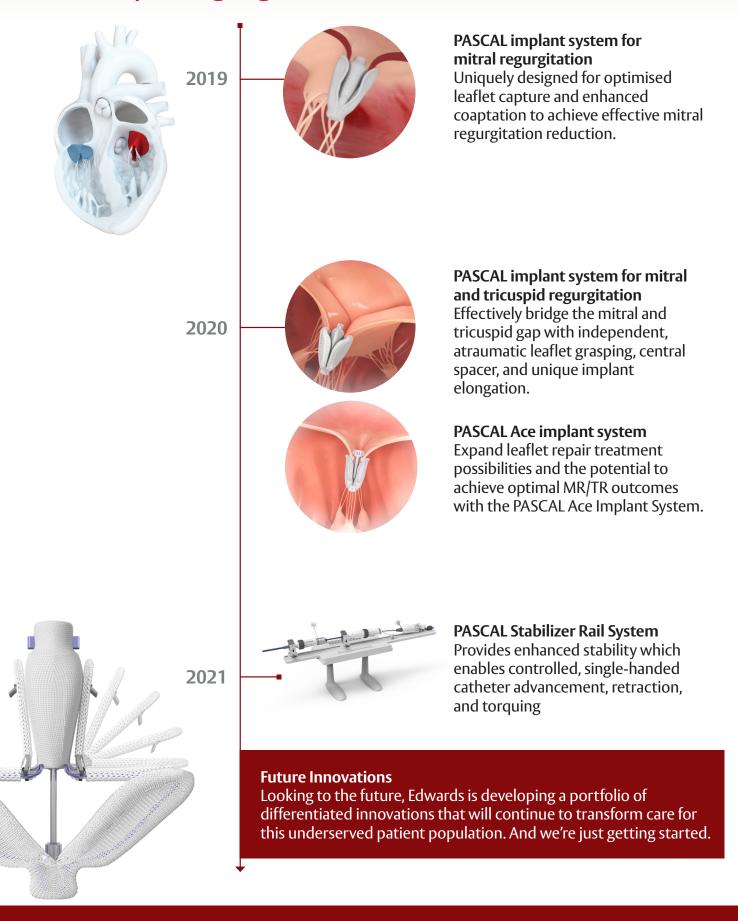
Edwards PASCAL Transcatheter Valve Repair System

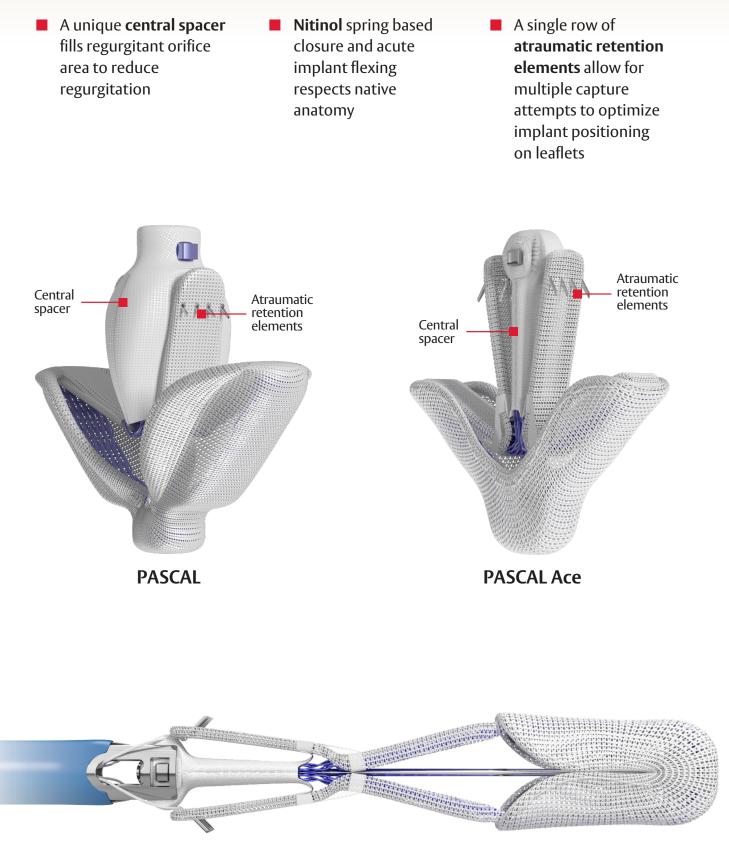
In your hands we can create amazing.



The PASCAL repair system is just the beginning of our dedication to treating a large and complex patient population with mitral and tricuspid regurgitation



PASCAL repair system designed for predictable capture, positioning, and release in patients with **mitral and tricuspid** regurgitation



PASCAL and PASCAL Ace feature implant elongation

PASCAL Ace implant system designed for even challenging tricuspid anatomies to treat more patients with **tricuspid** regurgitation

Dense chordae



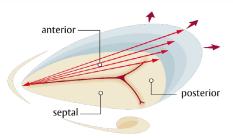
High variability of chordae: quantity, density, and location^{1,2}

Thin and variable leaflets



Thin, translucent, and more delicate leaflets with deep clefts and folds^{1,2,3}

Large annulus

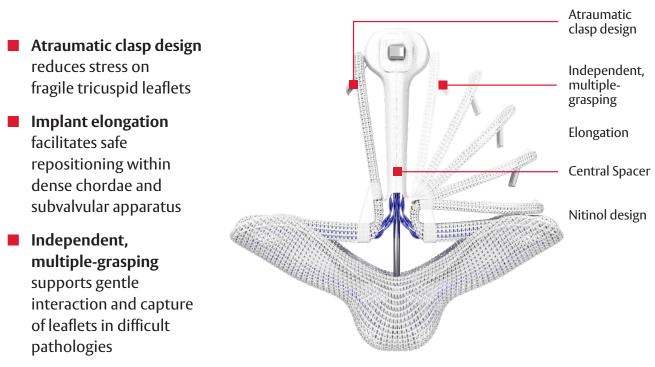


90% of TR is Functional TR (FTR), presenting with annular dilatation⁴

References:

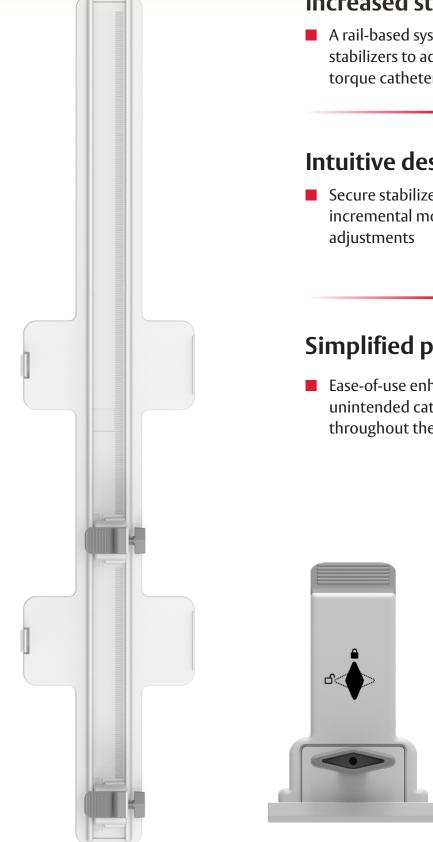
1. Pozzoli A, et al. Frontiers in Cardiovascular Medicine. 2018; 5. 1-9; 2. Dahou A, et al. JACC, 2019; 12: 458-68; 3. Athavale S, et al. Anat Cell Biol, 2017;50:1–6; 4. Tomos Mas P, et al. Heart. 2015; 101:1840-1848

PASCAL Ace implant system A narrower profile designed for precision and ease-of-navigation in complex anatomies



PASCAL Ace

The PASCAL Stabilizer Rail System designed to add stability to enable controlled catheter movements



Increased stability

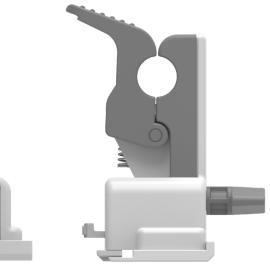
A rail-based system with multiple stabilizers to advance, retract, and torque catheters

Intuitive design

Secure stabilizers allow for stable, incremental movements and fine

Simplified process

Ease-of-use enhancements help reduce unintended catheter movements throughout the procedure





Visit Edwards.com/PASCAL to learn more.

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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