

Together, we are making a meaningful mark on the lives of patients with mitral and tricuspid regurgitation



Innovation for improved outcomes is what drives us. Now and in the future.

Mitral and tricuspid regurgitation are caused by compromised heart valves that allow blood to flow from the left ventricle to the left atrium and from the right ventricle to the right atrium, respectively.

These debilitating conditions affect more than 10% of people aged 65 and over in the US, creating a poor quality of life and high mortality rates. Yet fewer than 2% receive treatment intervention.*

*Based on 2010 US Census data and Stuge O, Liddicoat J. J Thorac Cardiovasc Surg. 2006;132:1258-1261.



Cardioband Tricuspid System

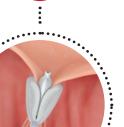
The first CE Mark transcatheter device for the treatment of tricuspid regurgitation reduces the tricuspid annulus to facilitate leaflet coaptation.



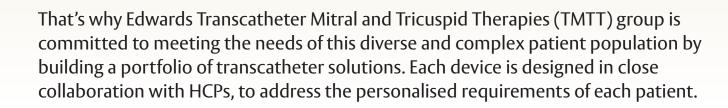


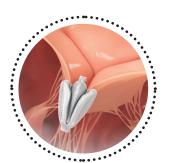
Cardioband Mitral System

Enables precise positioning to individual patient anatomy, and real-time adjustment for confirmation of results.



leaflet capture and enhanced coaptation to achieve effective mitral regurgitation reduction.





PASCAL System for Mitral and Tricuspid

Effectively bridge the mitral and tricuspid gap with independent, atraumatic leaflet grasping and central spacer to reduce stress.







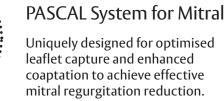
Future Innovations

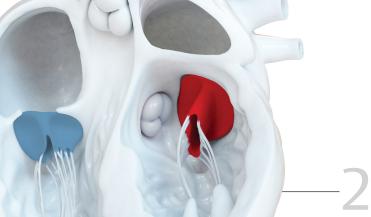
Looking to the future, Edwards TMTT is developing a portfolio of differentiated innovations from transcatheter repair to valve replacement that will continue to transform care for this underserved patient population. And we're just getting started.

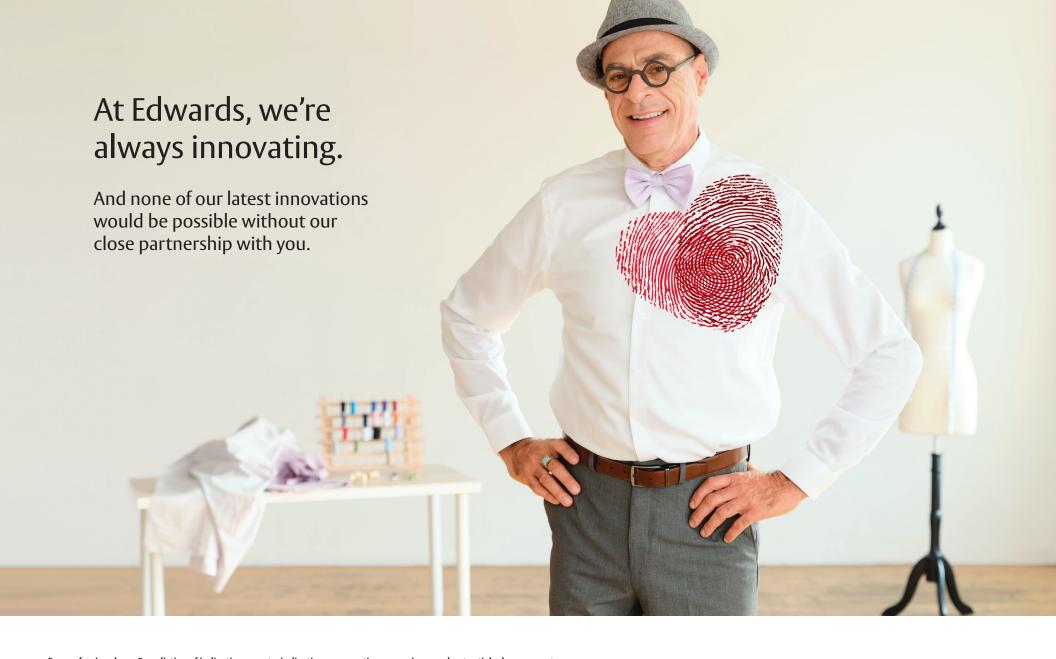


Expand leaflet repair treatment possibilities and the potential to achieve optimal MR/TR outcomes with the PASCAL Ace Implant System.









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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To learn more about our portfolio of mitral and tricuspid regurgitation solutions, visit www.makeamark.com

