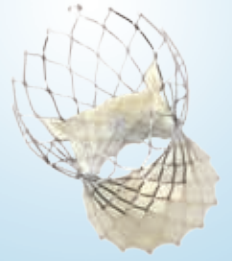


# HIGH POST-TAVI PCI SUCCESS RATE

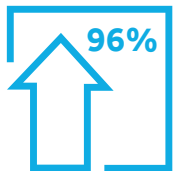
## REVIVAL REGISTRY DATA



### TREATING UNPLANNED PCI AFTER TAVI HAS A HIGH SUCCESS RATE

Unplanned PCI after TAVI is a rare but serious situation. A 15,325-patient multicenter international registry has demonstrated that successful treatment and outcomes are likely, regardless of the TAVI device used.<sup>1</sup>

### FINDINGS FROM REVIVAL REGISTRY<sup>1</sup>



Very high success rate (96.6%) for PCI post-TAVI.



No statistical difference in success rate for balloon-expandable versus self-expanding.



No statistical difference in the number of diagnostic or guiding catheters used, total fluoroscopy time, or vascular access route during PCI among patients treated with the two valve types.



The incidence of unplanned PCI post-TAVI was rare (0.9%, with a median follow-up of 191 days).



“The highest incidence of unplanned PCI is in the first week post-TAVI, and then there is a progressive decline.”

## REFERENCE

<sup>1</sup>Stefanini GG, Cerrato E, Pivato CA, et al. Unplanned Percutaneous Coronary Revascularization After TAVR. *JACC Cardiovasc Interv.* January 25, 2021;14(2):198-207.

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](http://medtronic.eu).

For applicable products, consult instructions for use on [manuals.medtronic.com](http://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.

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.

# Medtronic

## Europe

Medtronic International  
Trading Sàrl.  
Route du Molliau 31  
Case postale  
CH-1131 Tolochenaz  
Tel. +41 (0)21 802 70 00  
Fax +41 (0)21 802 79 00

[medtronic.eu](http://medtronic.eu)

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