





PRODUCT BROCHURE

Custom-Made Relay® Branch

Built to Accommodate



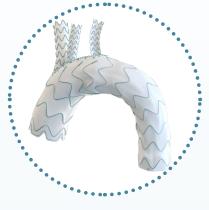




Addressing Challenges in the Aortic Arch

The Custom-made Relay®Branch provides an effective solution for the treatment of aortic arch disease¹.

Depending on the pathology and the targeted supra aortic vessel(s), this program can provide:



Triple Branch

The complete endovascular solution for the repair of the aortic arch







Double Branch

Two tunnels to be deployed under the BCT and LCCA







Single Branch

One tunnel to be deployed under the LSA





The Single Branch Customisation may include a proximal scallop to proximally extend the landing zone





Van der Weijde et al. (2019) - 'Total Endovascular Repair of the Aortic Arch: Initial Experience in the Netherlands', *The Annals of Thoracic Surgery*.

Case images courtesy of Prof. Piotr Szopinski, Institute of Hematology and Transfusion Medicine. Warsaw, https://www.vumedi.com/video/future-of-tevar-the-evolution-from-single-to-triple-branch-technology/?list=30f43900-2c2b-4624-830e-46dac385494e



Custom Relay®Branch Key Benefits

The Custom-made Relay®Branch builds upon the core features of the standard Relay®Pro platform, incorporating the same advanced technology and design principles, but with added customisation to address a specific patient need. The benefits of this platform include:



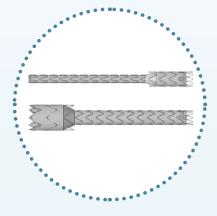
Graft

- Potential customisation options
- Wide window customisation
- Lock stent technology (where necessary)



Delivery System

- Dual sheath technology
- Pre-curved nitinol inner catheter
- Support wires
- Asymmetrical proximal clasping



Branches

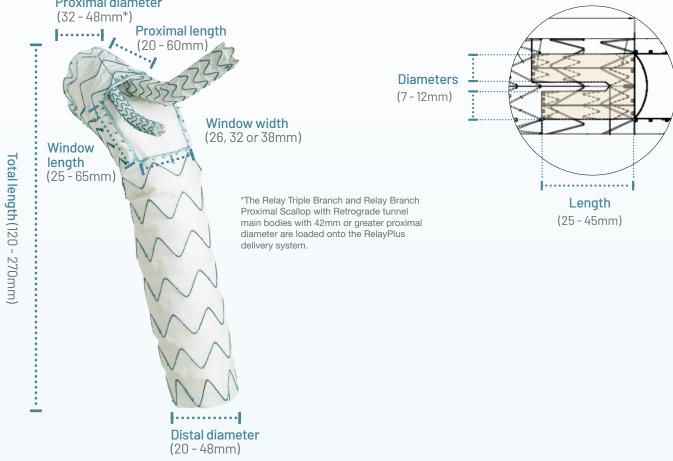
- TREO-based limb design
- Proximal clasping and short nose cone delivery system
- Controlled deployment
- 14 Fr O.D. delivery system

Case images courtesy of Prof. Piotr Szopinski, Institute of Hematology and Transfusion Medicine. Warsaw https://www.vumedi.com/video/relayr-double-and-triple-branch-case-overview/
Case images courtesy of Dr. Martin Funovics, Cardiovascular and Interventional Radiology Medical University of Vienna, https://www.vumedi.com/video/future-of-tevar-the-evolution-fromsingle-to-triple-branch-technology/?list=30f43900-2c2b-4624-830e-46dac385494e
Case images courtesy of Dr. Florian Elger, UniversitätSmedizin Göttingen



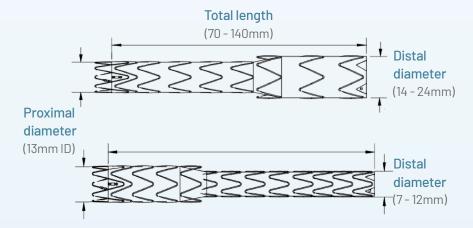
Potential Customisation Options Tailored Design For Every Need

Main Body Internal Tunnels Proximal diameter



Note: Measurements may vary depending on the specific configuration and the antegrade or retrograde position of the tunnels.

Branches





Wide Window Customisation

Designed for Easy Cannulation

▶ The cannulation window is designed to align to the outer curvature of the aorta to help **simplify branch** cannulation

Mean Operative Time Including Cervical

Bypassing 6

40min Fluoroscopy Time 7



"The large window[...] makes cannulation easy and fast while maintaining cerebral perfusion "7

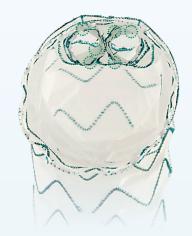
Lock Stent Technology*

Designed to prevent disconnection

▶ Dull barbs facing towards lumen of the tunnel designed to prevent potential potential disconnection of the branches 7

0% Type III Endoleak 7 At 18-month mean followup

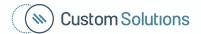
0% Branch disconnection or migration 7 At 18-month mean followup



"The presence of rounded barbs inside the tunnels (lock stent system) may also explain the absence of migration or disconnection phenomenon and subsequently of type III endoleak ** 7

*Lock stents are available upon request on every configuration and are limited to 12mm tunnel diameter. They are required when a TREO-based branch limb design is requested as a bridging stent.

^{6.} Kudo et al. (2020) - Early and midterm results of thoracic endovascular aortic repair using a branched endograft for aortic arch pathologies: A retrospective single-center study, JTCVS Techniques
Ferrer et al. (2019) - ITalian Registry of doUble inner branch stent graft for arch PatHology (the TRIUmPH Registry) Journal of Vascular Surgery

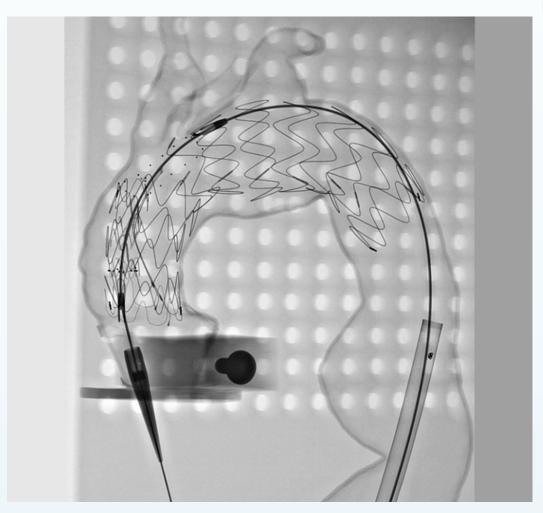


Dual Sheath Technology and Pre-Curved Nitinol Inner Catheter

Navigating the arch with care, designed for self-alignment

- ▶ Dual sheath delivery system, to facilitate the atraumatic advancement into zone 0
- ▶ Pre-curved nitinol inner catheter, to align the cannulation window to the outer curvature of the aorta

"[...]a "self-righting" inner sheath that aligns the large branch cannulation window to the outer curve [...]." 8



Benchtop Model

8.3% Stroke rate at 30-day follow-up ⁹

^{*} Custom-made double branch devices are based on an evolutionary device Terumo Aortic has developed that was part of a limited, regulated IDE clinical study in the USA focused on advanced disease states that has enrolled 30 subjects. As with any endovascular surgical repair involving the aortic arch, implanting this type of multi-branch device may lead to a neurological event for the patient. Whilst indicating overall effectiveness so far, we are aware that a significant proportion of the subjects who have been implanted have experienced a neurological event. While these events have differed in severity (disabling and non-disabling) and timing (<24 hours to 4.5 years) post implantation and many have resolved, the overall rate is ~60%. Note that the subjects enrolled in the study had to meet specific inclusion/exclusion criteria, specifically those patients that are at very high risk or prohibitive risk for open surgical repair, which may not be applicable to an individual patient and custom-made device. After extensive and continued review of the events and potential contributing factors. mitigations have been implemented and Terumo Aortic, in conjunction with its physician advisory committee, continue to seek additional methods and technologies to further reduce the event rate. There is no further enrollment in this study.

^{3.} McClure et al. (2021) - Zone 0 Aortic Arch Reconstruction Using the RelayBranch Thoracic Stent Graft CJC Open

^{9.} Iglesias Iglesias et al. (2023) - An early single-center experience with the Relay double inner-branch arch endograft, Journal of Thoracic Disease



Support Wires and Asymmetrical Proximal Clasping Precise and Controlled Deployment

- Two support wires guide the inferior portion toward the inner aortic wall, keeping it aligned with the landing zone, minimising the risk of retroflex
- Two clasped stent apices, both located on the outer curve, allow for precise and controlled deployment

"The presence of pared "driving-wires" (support wires) allows for a precise proximal landing in Zone 0 and for progressive apposition of the proximal stent-graft segment against the aortic wall. 99 10



Early Type la Endoleak 11

100%

Bridge Stents

- TREO-based Design
- Proximal Clasping and Short Nose Cone delivery system
- 14 Fr O.D. with 49cm long detachable sheath delivery system
- "Mechanical advantage" featured in the delivery system for controlled deployment

"The branches are customized versions of the iliac limbs used with the Treo abdominal endovascular stent graft system "1



Van der Weijde et al. (2019) - 'Total Endovascular Repair of the Aortic Arch: Initial Experience in the Netherlands', The Annals of Thoracic Surgery. Iglesias Iglesias et al. (2023) - An early single-center experience with the Relay double inner-branch arch endograft, Journal of Thoracic Disease Riambau et al. (2015) - Application of the Bolton Relay Device for Thoracic Endografting In or Near the Aortic Arch, AORTA Czerny et al. (2021) - Results of endovascular aortic arch repair using the Relay Branch System, European Journal of Cardio-Thoracic Surgery

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View Custom-Made device IFU at eifu.terumoaortic.com for more information on use, indications, contraindications and warnings/precautions.

Custom made devices are specifically made in accordance with a written prescription of any person authorised by national law by virtue of that person's professional qualifications; which gives (1) specific design characteristics provided under that person's responsibility and (2) is intended for the sole use of a particular patient exclusively to meet their individual conditions and needs. Custom made devices are not available in the US and availability is subject to local regulatory approval.

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