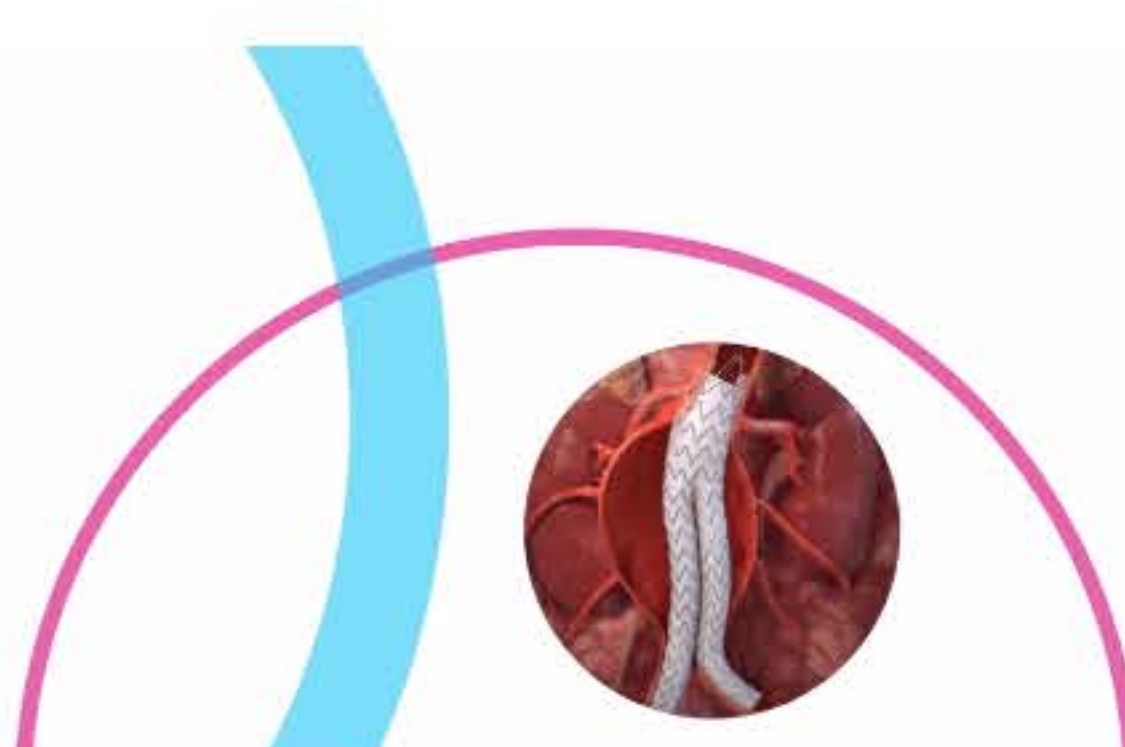


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EnCHEVAR Elevating CHEVAR therapy

Check out latest clinical data



Indications, Safety, and Warnings



- Discover EnCHEVAR
- Explore latest clinical release
- Watch experts testimonials
- View stent graft sizing guidelines

EnCHEVAR with Endurant™ II/IIIs stent graft system + Radiant™ balloon-expandable covered stent

Proven device combination for CHEVAR.

EnCHEVAR with Endurant™ II/IIIs system + Radiant™ covered stent demonstrates durable clinical outcomes (mean f/u 24.6 ± 17.4 months)¹:

- 1.6% new late onset type Ia endoleak benefits

Play video



Proven, safe, and effective way to lengthen aortic sealing zones

Learn how EnCHEVAR with Endurant™ II/IIIs system + Radiant™ stent provides durable outcomes in challenging hostile sealing zones.

EnCHEVAR deployment

Step-by-step deployment of the Endurant™ II/IIIs and Radiant™ stent grafts for CHEVAR.¹

Clinical outcomes

ENCHANT Trial³ 1-year data release

EnCHEVAR (Endurant II/IIIs stent graft system + Radiant¹ balloon-expandable covered stent) analysis (n = 62)³

ENCHANT is a multicentre, prospective, single arm, post-market study to assess the clinical outcomes, safety, and performance of the Endurant II/IIIs system used with balloon-expandable covered stents for treatment of juxtarenal aortic aneurysms with a short infrarenal neck in a real-world setting.

Key outcomes of the one year Kaplan-Meier analyses.

- 29 centres in EU and Russia
- Freedom from all cause mortality: 87.0 ± 4.3%
- Freedom from aneurysm rupture: 100%
- Freedom from conversion: 100%
- Freedom from all secondary procedures: 88.0 ± 4.3%
- Freedom from secondary procedures to treat type Ia endoleak: 100%



97.6% chimney graft patency at one year.

One year sac status by Core Lab (n = 41)⁴

- 2.4% sac increase
- 68.3% sac stable
- 29.3% sac decrease

ENCHANT Trial³ (pdf)

Download ENCHANT brochure



PROTAGORAS study

The PROTAGORAS study¹ evaluated the performance of the Endurant™ II/IIIs stent graft for patients with pararenal pathologic processes that were treated by the chimney/snorkel endovascular technique.

Standardised use of the Endurant™ II/IIIs stent graft system and Radiant™ covered stent for CHEVAR demonstrated:

N = 128 patients¹

- 24.6 months mean radiologic follow-up (range, 0-61 months)
- 100% technical success
- New late onset of type Ia endoleaks: 1.6%
- Primary patency of chimney grafts: 95.7%
- AAA sac diameter regression/stability: 90.6%
- Freedom from chimney graft related reinterventions: 93.1%

Radiant™ covered stent shows significantly better performance in CHEVAR compared to other grafts, with 94.6% patency at midterm compared to 84.1% patency with other covered stents.²

View brochure for full details.

Download brochure



Experts testimonials

Our experts tell it best. Watch their testimonials



Our experts tell it best. Clinician experiences on EnCHEVAR

Prof. Reijnen and Prof. Donas share their clinical experience about EnCHEVAR technique with Endurant™ II/IIIs system and Radiant™ covered stent.

Stent graft sizing

Expand All

- Endurant™ II/IIIs stent graft sizing guidelines
- Radiant™ stent graft sizing guidelines
- Radiant™ stent graft size configurations

Use this tool to see full ordering information and plan case specifics such as sizing and placement.

Sizing sheet

Preparing for the case

Recommended ancillary equipment[§]:

Soft wire for easy access to renal arteries

- 0.035" soft Terumo GLIDEWIRE™ hydrophilic-coated guidewire

Renal wire balanced between stiffness and flexibility; atraumatic short tip

- Cook Rosen™ wire guide

7 F sheath, 90 cm length with atraumatic tip for renal entry

- Medtronic TourGuide™ steerable sheath 7 F, 90 cm
- Cook Flexor™ Shuttle™ guiding sheath, Terumo Destination™ guiding sheath

Shorter 16 F sheath or larger may be used for access during a two-chimney case.

Catheter – 5 F, minimum length of 110 cm to 125 cm

Must be able to pass through and extend beyond 90 cm sheath; variety of angles to support renal cannulation; vertebral multipurpose.

Balloons

- Medtronic Reliant™ stent graft balloon catheter to achieve good conformability of the Medtronic Endurant II/IIIs stent graft system with the aortic wall and the renal stents
- Medtronic Admiral Xtreme™ PTA balloon catheter OTW 0.035" assist in access to renal (4 mm x 40 mm)

Inflation devices

- 20 cc syringe
- 20 cc inflation device equipped with manometer

Related pages

- Endurant™ II/IIIs Stent Graft System
- Radiant™ Balloon-Expandable Covered Stent
- Clinical outcomes

Aortic catalog

Choose from a complete portfolio of market-leading aortic vascular therapies to treat aneurysm disease.

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[§] Third-party brands are trademarks of their respective owners.

¹ Patients in this dataset were treated with Advanta V12, which was re-labelled as Radiant and received CE mark approval in September 2022 for use in CHEVAR.

² Denominator is the number of subjects with Core Lab-reported maximum aneurysm diameter at both one month post-implant and one year follow-up.

³ Other covered stents include: GORE Viabahn®, Bentley BeGraft™, BD LifeStream™.

⁴ All indications related to the Endurant™ II/IIIs and Radiant™ stent grafts are not approved globally. Please check your local regulatory approval status. Refer to the complete Instructions for Use manual of the Endurant™ II/IIIs and Radiant™ stent grafts for the indications approved in your geography.

References

¹ Donas KP, Torsello GB, Piccoli G, et al. The PROTAGORAS study to evaluate the performance of the Endurant stent graft for patients with pararenal pathologic processes treated by the chimney/snorkel endovascular technique. J Vasc Surg. January 2016;43(1):1-7.

² Psooulas GA, Torsello G, Austermann M, et al. Outcomes of elective use of the chimney endovascular technique in pararenal aortic pathologic processes. J Vasc Surg. February 2021;73(2):433-442.

³ Prof. Giovanni Torsello. "ENCHANT Trial interim analysis: Endurant CHEVAR technique for juxtarenal aneurysms." Presented at LINC, June 8, 2023; Leipzig, DE.

Brief statement: See the device manual for detailed information regarding the instructions for use, the implant procedure, 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 www.medtronic.com/manuals. Manuals can be viewed using a current version of any major internet browser. For best results, use Adobe Acrobat® Reader with the browser.

Important Reminder: This information is intended only for users in markets where Medtronic products and therapies are approved or available for use as indicated within the respective product manuals. Content on specific Medtronic products and therapies is not intended for users in markets that do not have authorization for use.