

GORE® TAG®
Conformable Thoracic
Stent Graft



U.S. IDE Clinical Trials

Five-year data across etiologies

GET STARTED



Together, improving life



Longterm results you can trust

U.S. IDE Clinical Trials: Five year data across ALL etiologies

ZERO

Migrations*
Fractures
Compressions

Type III endoleaks
Erosions

100%

Procedural
survival

↓↓ Low
rates of

Endoleaks
requiring
reintervention

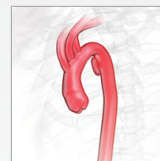
Device-related
reinterventions



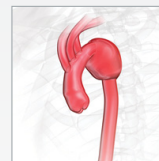
View trials



Acute Complicated
Type B Dissection »



Traumatic Aortic
Transection »

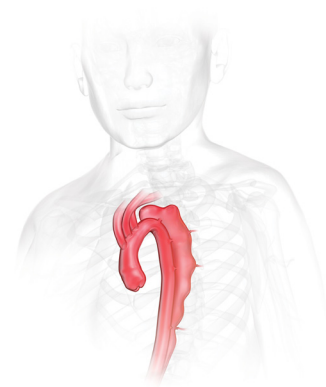


Thoracic Aortic
Aneurysm »

* Migrations are reported as those requiring reintervention.

Acute Complicated Type B Dissection

Long-term outcomes in a prospective multicenter trial



U.S. IDE Clinical Trial



50

patients
enrolled

100%

procedural
survival

5-year data

91%

of patients exhibited
an increase in true
lumen diameter*

87%

complete
thrombosis
observed

96%

of patients did not
experience late deaths
attributable to aortic
pathology after 30 days†

ZERO

Migrations‡

Type III endoleaks

Erosions

Compressions

Fractures

Conversions to open
surgical repair

* At five years.

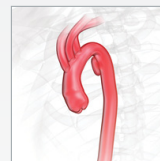
† 8% 30-day mortality rate.

‡ Migrations are reported as those
requiring reintervention.

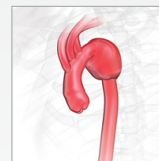
View trials



Acute Complicated
Type B Dissection »



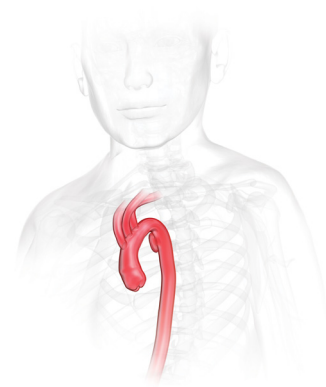
Traumatic Aortic
Transection »



Thoracic Aortic
Aneurysm »

Traumatic Aortic Transection

Long-term outcomes in a prospective multicenter trial



U.S. IDE Clinical Trial



101

patients
enrolled

100%

procedural
survival

5-year data

100%

Freedom from
aortic-related
mortality

99%

did not experience
a device/procedure-
related stroke*

95%

Freedom from
all-cause mortality
at 30 days

89%

Freedom from
all-cause mortality
at 5 years

ZERO

Reinterventions

Endoleaks[†]

Ruptures[‡]

Fractures

Lumen obstructions

Thrombus related
events

Conversions to open
surgical repair

Compressions

Migrations[§]

Erosions



As published in:
The Annals of Thoracic Surgery (2021).

* n = 1 post procedural, ischemic, stroke.

† No serious endoleaks reported. Two minor
endoleaks were reported which resolved
without reintervention.

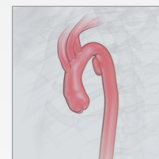
‡ Post procedure.

§ Requiring reinterventions.

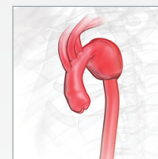
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Acute Complicated
Type B Dissection »



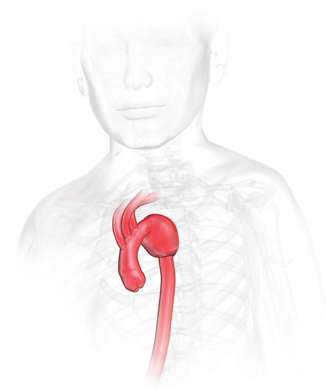
Traumatic Aortic
Transection »



Thoracic Aortic
Aneurysm »

Thoracic Aortic Aneurysm

Long-term outcomes in a prospective multicenter trial



U.S. IDE Clinical Trial



66
patients
enrolled

100%
procedural
survival

5-year data

88%

Freedom from
device-related
reinterventions

82%

Stable aneurysm
sac diameters

93%

Freedom from
aneurysm-related
mortality

ZERO

Fractures
Compressions
Migrations*
Type III endoleaks

Conversions to open
surgical repair
Erosions



**Low
rates of**

Reintervention†
Proximal leaks‡
Rupture§



As published in:
Journal of Vascular Surgery (2021).

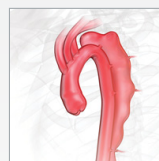
* With clinical sequelae resulting in reinterventions through five-years.

† Device related reinterventions: 7.6% (n=5)
four of the five were treatments for endoleaks
and one for a distal thoracic pseudoaneurysm.

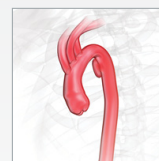
‡ Type Ia was rare n = 1.

§ 1.5% n = 1.

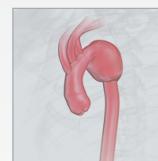
View trials



Acute Complicated
Type B Dissection »



Traumatic Aortic
Transection »



Thoracic Aortic
Aneurysm »

Contact a Gore representative to learn more.
tevar.tv

References

OP869 Farber MA, Krishnasastry KV, Desai N, Starnes BW, Matsumura JS, Tohill BC; TAG 08-02 Clinical Trial Investigators. Five-year outcomes with Conformable GORE® TAG® endoprosthesis used in traumatic aortic transections. *Annals of Thoracic Surgery*. In press.
<https://www.sciencedirect.com/science/article/pii/S0003497521010134>

OP871 Jordan WD, Desai N, Letter AJ, Matsumura JS. Long-term outcomes of the CTAG thoracic endograft in a prospective multicenter trial. *Journal of Vascular Surgery* 2021;74(5):1491-1498.
<https://www.sciencedirect.com/science/article/pii/S0741521421007485>

W. L. Gore & Associates. Evaluation of the GORE Conformable TAG® Thoracic Endoprosthesis for Treatment of Acute Complicated Type B Aortic Dissection. Bethesda, MD: National Library of Medicine; 2009. NLM Identifier: NCT00908388. Published May 25, 2009. Updated October 27, 2017. Accessed March 1, 2022. Available from: <https://clinicaltrials.gov/ct2/show/NCT00908388>

 Consult Instructions
for Use
eifu.goremedical.com

INDICATIONS FOR USE IN THE U.S.: The GORE® TAG® Conformable Thoracic Stent Graft is intended for endovascular repair of all lesions of the descending thoracic aorta, including: Isolated lesions in patients who have appropriate anatomy, including: Adequate iliac/femoral access; Aortic inner diameter in the range of 16-42 mm; ≥ 20 mm non-aneurysmal aorta proximal and distal to the lesion. Type B dissections in patients who have appropriate anatomy, including: Adequate iliac/femoral access; ≥ 20 mm landing zone proximal to the primary entry tear; proximal extent of the landing zone must not be dissected; Diameter at proximal extent of proximal landing zone in the range of 16-42 mm. **CONTRAINDICATIONS:** The GORE® TAG® Conformable Thoracic Stent Graft is contraindicated in: Patients with known sensitivities or allergies to the device materials (Table 1); Patients who have a condition that threatens to infect the graft. Refer to *Instructions for Use* at eifu.goremedical.com for a complete description of all applicable indications, warnings, precautions and contraindications for the markets where this product is available. **Rx Only**

INDICATIONS FOR USE UNDER CE MARK: The GORE® TAG® Conformable Thoracic Stent Graft is indicated for endovascular repair of the descending thoracic aorta. **CONTRAINDICATIONS:** Patients with known sensitivities or allergies to the device materials; patients with a systemic infection who may be at increased risk of endovascular graft infection. Refer to *Instructions for Use* at eifu.goremedical.com for a complete description of all applicable indications, warnings, precautions and contraindications for the markets where this product is available. **Rx Only**

Products listed may not be available in all markets.

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