Confidence for Endovascular Treatment of Type B Dissection

The Conformable GORE® TAG® Device is specifically engineered to treat compromised thoracic aortas:

- No barbs or bare springs
- Low spring-back force
- Fully covered distal end provides a transition between the stent frame and the septum, decreasing the risk of septum perforation
- Provides the greatest options for oversizing of any commercially available device

Retrograde Type A Dissection (RTAD)

Gore devices compared to devices with proximal bare springs

Publications[1]	Device-related RTAD Incidence[3]	% RTAD w/Proximal Bare Springs	% RTAD w/Gore Devices
J Vasc Surg. 2012;55(5):1255-1262.	1.27%	50%	0%
Eur J Cardiothorac Surg. 2012;42(3):566-570.	N/A	100%	0%
J Vasc Surg. 2010;52(6):1450-1457.	2.46%	100%	N/A
Circulation. 2009;120(11)Suppl:S276-S281.	0.80%[2]	93%	N/A

Stent-Induced New Entry Tear (SINE)

Gore devices compared to devices with proximal bare springs

Publications[1]	Distal SINE Incidence	SINE w/Proximal Bare Springs	SINE w/Gore Devices
Catheter Cardiovasc Interv. 2015;85(2):E43-E53.	6.3%	100%	0%
J Vasc Surg. 2010;52(6):1450-1457.	1.2%	71.4%	N/A

1. Complete reference on file.
2. Derived from 48 patients with complete data sets on patient and procedural characteristics, presentation, management, and outcomes from retrograde ascending aortic complications.
3. Retrograde Type A dissection determined to be caused by the endovascular device.

To view further dissection information, visit goremedical.com/aortic/tevar

CONFORMABLE

GORE TAG® THORACIC ENDOPROSTHESIS

Products listed may not be available in all markets.

PERFORMANCE through experience