

# Long-term Characterization of GORE® TAG® Conformable Thoracic Stent Graft With ACTIVE CONTROL System Performance

NCT06658730

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Status	RECRUITING
Sponsor	W.L.Gore & Associates
Enrollment	1,500 participants

## Key Eligibility Criteria

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### Inclusion (4)

- Patient or legally authorized representative (LAR) provides written authorization and/or consent per institution and geographical requirements.
- Patient has been or is intended to be treated with an eligible registry device.\*
- Patient is age e 18 years at time of informed consent signature.
- The intent to treat a patient with a Gore product must be made prior to soliciting for possible registry participation. If pre-procedure consent is not feasible due to emergent situation, consent prior to the time of discharge for the index procedure is acceptable.

### Exclusion (3)

- Patient who is, at the time of consent, unlikely to be available for standard of care (SOC) follow-up visits as defined by the site's guidelines and procedures.
- Patient is currently enrolled in or plans to enroll in any concurrent investigational drug and/or investigational device study\* within 12 months of Together Registry enrollment. Subjects cannot be enrolled in another Together Registry module protocol.
- The term "study" does not apply to other observational registries or quality improvement projects. Collection of Registry Device performance from interventional studies may be permissible provided device application is not investigational and there are no novel requirements that alter follow-up conduct (i.e., protocol-mandated interventions).

## Locations (10 total)

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Cedars-Sinai Medical Center, Los Angeles, California, United States  
Indiana University School of Medicine, Indianapolis, Indiana, United States  
Duke University, Durham, North Carolina, United States  
... and 7 more locations