TRITON-PN: A Study to Evaluate the Efficacy and Safety of Nucresiran in Patients With Hereditary Transthyretin Amyloidosis With Polyneuropathy

NCT07223203

Status	RECRUITING
Phase	Phase 3
Sponsor	Alnylam Pharmaceuticals
Enrollment	125 participants

Key Eligibility Criteria

Inclusion (4)

- Has documented diagnosis of hATTR-PN
- Has a diagnosis of hATTR amyloidosis with polyneuropathy with a documented TTR gene variant
- Has a neuropathy impairment score (NIS) of 5 to 130 (inclusive)
- Has a Karnofsky Performance Status (KPS) of e60%

Exclusion (7)

- Has had a liver transplant or is likely, in the opinion of the Investigator, to undergo liver transplantation during the Treatment Period of the study
- Has known other (non-hATTR) forms of amyloidosis or clinical evidence of leptomeningeal amyloidosis
- Has a New York Heart Association (NYHA) heart failure classification \>2
- Has alanine aminotransferase (ALT) or aspartate aminotransferase (AST) \>2.5 upper limit of normal (ULN)
- Has total bilirubin \>1.5 ULN

... and 2 more (see full listing online)

Locations (1 total)

Clinical Trial Site, Boston, Massachusetts, United States