

Access Cannulation Trial II

NCT05490225

Status	RECRUITING
Phase	Not Applicable
Sponsor	Voyager Biomedical
Enrollment	100 participants

Key Eligibility Criteria

Inclusion (36)

- The subject's AVF is deemed uncannulatable because:
 - The subject's anticipated cannulation zone(s) is/are ≥ 6 mm in depth from the surface of the skin to the anterior wall of the access vein as confirmed by ultrasound within 8 weeks prior to device implantation (each zone, Arterial/Pull and Venous/Push, shall be assessed independently of one another for device placement):
 - Arterial/Pull Zone: _____ mm deep
 - Venous/Push Zone: _____ mm deep
 - OR
- ... and 31 more (see full listing online)

Exclusion (2)

- The subject's access vein is ≥ 15 mm in depth at either cannulation zones as measured by ultrasound within 8 weeks prior to device implantation.
- Arterial/Pull Zone: _____ mm in depth

Locations (8 total)

Trinity Research Group, Dothan, Alabama, United States
Apex Research, Riverside, California, United States
Brigham and Women's Hospital, Boston, Massachusetts, United States
... and 5 more locations