

A Prospective, Multi-Center, Open Label, Randomized Control Clinical Trial Evaluating the Safety and Efficacy of the Cordella™ Pulmonary Artery Sensor System in New York Heart Association (NYHA) Class II Heart Failure Patients (PROACTIVE- HF-2 Trial)

Status: Recruiting

Eligibility Criteria

Sex: Male or Female

Age Group: 18 years and over

This study is NOT accepting healthy volunteers

Inclusion Criteria:

- Heart Failure NYHA Class II or Class III

Exclusion Criteria:

- ACC/AHA Stage D refractory Heart Failure (HF) - history of multiple pulmonary embolism (PE) - resting systolic blood pressure less than 90 mmHg

Conditions & Interventions

Interventions:

Device: Cordella™ Pulmonary Artery Sensor System

Conditions:

Heart & Vascular

Keywords:

Cardiovascular, Clinics and Surgery Center (CSC), Heart Failure, HF

More Information

Description: The Cordella™ Pulmonary Artery Sensor System is a possible treatment for New York Heart Association (NYHA) Class II and III heart failure. The main purpose of this study is to investigate the safety and effectiveness of the study device in helping to reduce Heart Failure hospitalizations.

Study Contact: Michelle Pitt - henni032@umn.edu

Principal Investigator: Tamas Alexy

Phase: NA

IRB

Number: STUDY00021914

Thank you for choosing StudyFinder. Please visit <http://studyfinder.umn.edu> to find a Study which is right for you and contact sfinder@umn.edu if you have questions or need assistance.