Randomized Clinical Evaluation of the AccuCinch Ventricular Restoration System in Patients who Present with Symptomatic Heart Failure with Reduced Ejection Fraction (HFrEF)

Status: Recruiting

Eligibility Criteria

Sex: Male or Female
Age Group: 18 years and over
This study is NOT accepting healthy volunteers

Inclusion Criteria:

- at least 18 years old - Ejection Fraction: between 20% and 40% measured by transthoracic echocardiography (TTE) - diagnosis and treatment for heart failure should be established at least 90 days before entering the study & should be on stable, optimal medical therapy for at least 30 days

Exclusion Criteria:

- myocardial infarction or any percutaneous cardiovascular intervention, cardiovascular surgery, or carotid surgery within 90 days prior to consent - any planned cardiac surgery or interventions within the next 180 days - women who are pregnant, planning to become pregnant, or are breast feeding - additional cardiac and medical diagnosis will exclude participation (study staff will review)

Conditions & Interventions

Interventions:
Device: AccuCinch Ventricular Restoration System, Drug: Guideline-Directed Medical Therapy
Conditions:
Heart & Vascular
Keywords:
Clinics and Surgery Center (CSC), Dilated Cardiomyopathy, Heart Failure, Heart Failure With Reduced Ejection Fraction (HFrEF)

More Information

Description: The objective of this study is to evaluate the safety and efficacy of the AccuCinch Ventricular Restoration System in patients with symptomatic heart failure with reduced ejection fraction (HFrEF).
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Phase: N/A
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