AFFIX: Post-market Study of MaxTack™ Motorized Fixation Device in Subjects Undergoing Ventral Hernia Repair

NCT06710795

Status	RECRUITING
Phase	Not Applicable
Sponsor	Medtronic - MITG
Enrollment	110 participants

Key Eligibility Criteria

Inclusion (7)

- Subject has provided informed consent (IC)
- Subject is 18 years of age or older at the time of consent
- Subject is able and willing to comply with the study requirements and follow-up schedule
- Subject is undergoing an elective, single-stage, primary or incisional ventral hernia repair
- Subject is undergoing minimally invasive ventral hernia repair procedure using the MaxTack™ Motorized Fixation Device

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

Exclusion (22)

- Subject is undergoing an emergency surgery (e.g., lifesaving procedures performed where subject is in imminent danger of death, strangulated hernia, etc.)
- Subject has history of 3 or more hernia repair procedures
- Subject has existing mesh in the space where the physician needs to apply the new mesh to be fixated with the MaxTack™ Motorized Fixation Device
- Subject is scheduled (or anticipated to be scheduled) for additional surgery, and subsequent surgery would jeopardize previous application of study treatment
- Subject has history of allergic reactions to Poly (Glycolide-co-L-lActide) (PGLA)

... and 17 more (see full listing online)

Locations (3 total)

Cleveland Clinic - Weston Hospital, Weston, Florida, United States
Cleveland Clinic - Ohio, Cleveland, Ohio, United States
The Ohio State University, Columbus, Ohio, United States