

Management of Dyssynergic Defecation

NCT07225803

Status	RECRUITING
Phase	Not Applicable
Sponsor	Mayo Clinic
Enrollment	200 participants

Key Eligibility Criteria

Inclusion (2)

- Men and women aged 18-80 years who are able and willing to provide written informed consent and to comply with the requirements for the entire study
- Satisfy Rome III criteria for functional constipation and/or irritable bowel syndrome as assessed by questionnaires and objective evidence of dyssynergic defecation on anorectal testing

Exclusion (11)

- Clinical evidence of significant cardiovascular, respiratory, renal, hepatic, gastrointestinal, hematological, neurological, psychiatric or other diseases that may interfere with the objectives of the study and/or pose safety concerns.
 - History of total, hemi- or subtotal colectomy at any time or other abdominal operations within 60 days prior to entry into the study. Segmental resection of the colon (e.g. sigmoid resection) for diverticulitis will be permitted.
 - Current anal fissure
 - Clinically significant structural abnormalities (eg, rectocele \gt 4cm, large enterocele, full thickness rectal prolapse) that, in the opinion of the investigator, are likely contributing to symptoms
 - Current or past history of colon or rectal cancer, scleroderma, inflammatory bowel disease, congenital anorectal abnormalities, solitary rectal ulcer syndrome, rectal resection or pelvic irradiation
- ... and 6 more (see full listing online)

Locations (1 total)

Mayo Clinic, Rochester, Minnesota, United States