Naltrexone Neuroimaging in Teens With Eating Disorders

NCT05509257

Status	RECRUITING
Phase	Early Phase 1
Sponsor	Children's Mercy Hospital Kansas City
Enrollment	60 participants

Key Eligibility Criteria

Inclusion (4)

- Adolescents and young adults aged 13-21 years
- Eating disorder diagnosis characterized by binge eating and/or purging (eg, Anorexia Nervosa-Binge/Purge, Bulimia Nervosa, Binge Eating Disorder, Other Specified Feeding/Eating Disorder) using Diagnostic and Statistical Manual of Mental Disorders, 5th edition (DSM-V) criteria.
- Stable medication regimen (no dose or drug changes in the past 4 weeks)
- Participant and parent/legal guardian (if under 18 years) are willing and able to provide informed permission/assent/consent for the study

Exclusion (6)

- Pregnant (via UCG)
- Prior hypersensitivity reaction to naltrexone (e.g., anaphylaxis)
- Non-removable metal in the body that is magnetic resonance imaging incompatible
- Current naltrexone use
- Self-reported opioid use in the past 7 days

... and 1 more (see full listing online)

Locations (1 total)

Children's Mercy Research Institute, Kansas City, Missouri, United States