

A Multicentric European Study to Promote Clinical Trial Readiness for STXBP1-related Disorders

NCT06625112

Status RECRUITING
Sponsor European STXBP1 Consortium
Enrollment 120 participants

Key Eligibility Criteria

Inclusion (1)

- participant has a (likely) pathogenic, disease-causing STXBP1 variant, according to the American College of Medical Genetics and Genomics (ACMG) criteria; or participant has a larger structural variant including the STXBP1 gene where STXBP1 is thought to be (one of) the culprit gene(s) causing the phenotype
- written informed consent from study participant and/or legal guardian.

Locations (1 total)

Universitair Ziekenhuis Antwerpen, Antwerp, Belgium