

Genetic Developmental and Epileptic Encephalopathy Natural History Study for Clinical Trial Readiness

NCT07413211

Status RECRUITING
Sponsor Weill Medical College of Cornell University
Enrollment 22,068 participants

Key Eligibility Criteria

Inclusion (29)

- Molecular diagnosis of a genetic disorder associated with DEE, as confirmed by the study investigators
- A neurological phenotype such as epilepsy or developmental delay as confirmed by the study investigators.
- English Speaking (Arms 1, 2, 4, 5). The registries may be completed by people who speak any language.
- ARM 1 (In person cohort)
- MEF2C
- ... and 24 more (see full listing online)

Exclusion (3)

- Presence of a significant non-DEE-related central nervous impairment/behavioral disturbance that would confound the scientific rigor or interpretation of results of the study
- History of prematurity (defined as gestational age <35 weeks), intraventricular hemorrhage, structural brain deficit, or congenital heart disease
- Presence of a clinical comorbidity deemed by the investigator to potentially confound the typical presentation of DEE

Locations (1 total)

Weill Cornell Medicine, New York, New York, United States