

Natural History Study of Patients with HPDL Mutations

NCT05848271

Status RECRUITING
Sponsor University of California, San Diego
Enrollment 50 participants

Key Eligibility Criteria

Inclusion (6)

- Any individuals diagnosed with HPDL variants
- Clinical diagnosis can include:
 - HPDL-related hereditary spastic paraplegia (HSP)
 - HPDL-related neonatal mitochondrial encephalopathy
 - Spastic paraplegia -83 (SPG83)
- ... and 1 more (see full listing online)

Exclusion (2)

- Any known genetic abnormality (other than HPDL mutation)
- Any condition that, in the opinion of the Site Investigator, could put the participant at undue risk and/or would ultimately prevent the completion of study procedures

Locations (1 total)

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