

# MYT1L Syndrome: a Rare Paediatric Genetic Syndrome Responsible for a Neurodevelopmental Disorder

NCT07008612

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**Status** RECRUITING  
**Sponsor** University Hospital, Rouen  
**Enrollment** 50 participants

## Key Eligibility Criteria

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### Inclusion (9)

- MYT1L Group Patients
- Language: French
- Consent of parents or legal guardian
- Social security coverage required
- Prosody Group Patients
- ... and 4 more (see full listing online)

### Exclusion (8)

- MYT1L Group patients
- Unaided visual or hearing impairment making assessments impossible
- Non-French speaking patients
- Patients with a dual molecular genetic diagnosis also causing a neurodevelopmental disorder
- Acquired neurological disorder
- ... and 3 more (see full listing online)

## Locations (1 total)

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University Hospital of Rouen, Rouen, France