

Identification of the Molecular and/or Pathophysiological Bases of Rare Diseases of Genetic Origin (or Rare Forms of Frequent Diseases Suspected of Being of Genetic Origin).

NCT03287193

Status RECRUITING
Sponsor Centre Hospitalier Universitaire Dijon
Enrollment 850 participants

Key Eligibility Criteria

Inclusion (7)

- Patients (children or adults) with a suspected rare disease (or rare form of a common disease) of genetic origin for which the molecular basis is not known, or for which the understanding of the pathophysiological mechanism is imperfectly known.
 - OR
 - Foetuses with developmental abnormalities for which the molecular basis is not known, or for which the understanding of the pathophysiological mechanism is imperfectly known.
 - OR
 - Apparently healthy relatives or controls AND
- ... and 2 more (see full listing online)

Exclusion (1)

- Patients without national health insurance cover

Locations (1 total)

Chu Dijon Bourgogne, Dijon, France