

# Genetic of Intellectual Deficiency and Autism Spectrum Disorders (RaDiCo-GenIDA)

NCT06871696

---

|            |   |
|------------|---|
| Status     | RECRUITING  |
| Sponsor    | Institut National de la Santé Et de la Recherche Médicale, France |
| Enrollment | 1,000 participants  |

## Key Eligibility Criteria

---

### Inclusion (6)

- Be a voluntary adult (aged 18 or older)
  - Be a family member (i.e., mother/father) of patients with intellectual disabilities and/or autism spectrum disorders of known genetic origin. This includes monogenic causes as well as recurrent copy number variations (CNVs) such as deletions or duplications. Note: we also allow adult patients to participate directly if they wish and have the capacity to do so.
  - Have knowledge of the genetic cause behind intellectual disabilities or autism spectrum disorders. An exception to this rule is possible for patients with a syndrome that includes intellectual disabilities or autism spectrum disorders, and for whom genetic investigation is considered, with the approval of the project's scientific council (which will define the syndromes eligible for this exception).
  - Have the intellectual and material capabilities to complete an internet questionnaire.
  - Have read the information sheet regarding the study and agreed to the general conditions of participation in the study.
- ... and 1 more (see full listing online)

### Exclusion (2)

- Patients affected by the presence of intellectual disability and/or an autism spectrum disorder of unknown genetic origin will not be able to participate in the study, except with the exception mentioned in the previous chapter.
- It is requested that only adults enter data. However, the collected data may pertain to a minor (in the case of a parent entering data about their minor child)

## Locations (1 total)

---

RaDiCo-GenIDA, Paris, Île-de-France Region, France