

https://ataxia-global-initiative.net/
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Ataxia Global Initiative

Trial Site Registry

(AGI-TSR)

Dataset

Contact Details

Site name Address Zip-Code / City Country Homepage PI

- Title
- First name
- Last name
- E-Mail
- Phone

Patient Population

Genetic ataxia subtypes

In the fields underneath please fill in data and information regarding the cohort of ataxia patients you regularly see, e.g. at annual visits. Please be as accurate as possible.

| Disease name | Patients presents yes or no | Age groups 0-17 or 18+ | Ambulatory (number) | Consented to be contacted for clinical trials (both age groups) |
|------------------------|-----------------------------------|---------------------------|------------------------|---|
| autosomal-dominant ata | | | | |
| repeat SCAs | | | | |
| SCA1 | | | | |
| SCA2 | | | | |
| SCA3 | | | | |
| SCA6 | | | | |
| SCA7 | | | | |
| SCA 10 | | | | |



| SCA17 | | | | | | |
|---|------------------|--|--|--|--|--|
| SCA27b | | | | | | |
| DRPLA | | | | | | |
| channelopathies | | | | | | |
| KCNA1 (Episodic ataxia | | | | | | |
| type 1) | | | | | | |
| CACNA1A (Episodic ataxia type 2) | | | | | | |
| SCA19 | | | | | | |
| SCA29 | | | | | | |
| neurometabolic | | | | | | |
| Alexander disease | | | | | | |
| other SCAs | | | | | | |
| open entry field | | | | | | |
| Autosomal-recessive and | x-linked ataxias | | | | | |
| frequent ataxias (frequent, drugs in development and/or strong NHS consortia which will make trial-readiness in close future likely) | | | | | | |
| Friedreich's Ataxia (FA) | | | | | | |
| Ataxia Telangiectasia (AT) | | | | | | |
| Autosomal-recessive spastic ataxia type Charlevoix Saguenay (ARSACS) | | | | | | |
| RFC1 | | | | | | |
| mitochondrial (recurrent, drugs in development drugs in development and/or strong NHS consortia which will make trial-readiness in close future likely) | | | | | | |
| SPG7 | | | | | | |
| COQ8A | | | | | | |
| neurometabolic | | | | | | |
| Niemann Pick Type C (NPC) | | | | | | |
| Tay-Sachs Disease | | | | | | |
| Sandhoff Disease | | | | | | |
| Gaucher disease type 3 | | | | | | |
| Cerebrotendinous xanthomatosis (CTX) | | | | | | |
| Adrenoleukodystrophy (ALD) | | | | | | |
| Spastic ataxia associated with GBA2 | | | | | | |
| other | | | | | | |
| AOA1 | | | | | | |
| AOA2 | | | | | | |



Clinical Trials - Experience

Qualified for GCP (Good Clinical Practice) with documented certifications:

- number of clinicians
- number of study nurses

Number of clinical trials in which the site PI team participated in the last 5 years

| • | Phase I | (a) for neurological diseases | (b) for ataxias |
|---|-----------|-------------------------------|-----------------|
| • | Phase II | (a) for neurological diseases | (b) for ataxias |
| • | Phase III | (a) for neurological diseases | (b) for ataxias |
| • | Phase IV | (a) for neurological diseases | (b) for ataxias |

Clinical Trials - Equipment

- 1. Certified Laboratory
- 2. Centrifuge
- 3. Ice
- 4. Dry ice
- 5. Liquid nitrogen
- 6. Refrigerator lockable/not lockable with the following temperatures:
 - -80°C(-112°F)
 - -20°C (-4°F)
 - 2-8°C(36°F-46,4°F)
- 7. Lockable rooms or filling cabinets
- 8. Lockable safes to keep study drugs
- 9. Temperature recorder for the fridges and safes
- 10. Possibility for conducting trails in inpatient care setting
- 11. Dedicated clinical trials unit for children and/or adults
- 12. Emergency care unit
- 13. resuscitation team at the study site
- 14. MRI (at least 1,5 Tesla)
- 15. Pharmacy that is able to handle GMP grade drugs for trials
- 16. Possibility and previous experience with intrathecal administration of drugs
 - In clinical care setting
 - In trial setting