SCN2A-RELATED AUTISM & EPILEPSY

FAMILIESCN2A®
FOUNDATION

FAMILY RESEARCH CHECKLIST

CHECK OFF ✓

- [ ] **Subscribe to the FamilieSCN2A email list**
- [ ] **Join the Global Family Network on Facebook**
- [ ] **Explore the new SCN2A.org website**
- [ ] **Create a unique CRID (Clinical Research ID)**
- [ ] **Learn about your genetic variant**
- [ ] **Gather and organize medical records**
- [ ] **Join the Global SCN2A DRAGONFLY Registry**
- [ ] **Register for the SCN2A Town Hall meetings**
- [ ] **Record baseline information**
- [ ] **Sign up for the Clinical Picture Maker**
- [ ] **Join Simons Searchlight Natural History Study**
- [ ] **Join the SCN2A Community on Citizen**
- [ ] **Watch educational webinars on YouTube, including "About SCN2A Series"**

Research is the key to unlocking future treatments and potential cures for SCN2A-related disorders.

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Learn About Your Genetic Variant

- ☐ Gain of Function (GoF)
- ☐ Loss of Function (LoF)
- ☐ Mixed Function
- ☐ Unsure

CRID – Clinical Research ID

Email: _____

Password: _____

The Dragonfly Study – SCN2A Patient Registry

- ☐ Enrollment
- ☐ 6 month follow up
- ☐ 18month follow up
- ☐ 12 month follow up
- ☐ 24 month follow up

Email: _____

Password: _____

Citizen

Username: _____

Password: _____

Simons Searchlight

Username: _____

Password: _____

Clinical Picture Maker (CPM)

Username: _____

Password: _____

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