Join the NF Registry

When you join the NF Registry you have access to the latest discoveries about the many ways living with NF can affect individuals and families, which helps you and your family find the best possible care.

As an NF Registry participant, you complete a yearly health survey. This data helps researchers study how NF affects everyone differently and how NF changes over time. You can then choose whether to receive personalized emails about:

- Clinical trials and research studies relevant to you or your child
- Updates to NF care recommendations
- Research announcements and news
- Surveys designed to get patient input on key NF challenges
- Educational materials specific to you
- Resources to help you on your path with NF

Even if you choose not to be contacted, your participation helps researchers learn from the real experts – NF patients and families.

Join at [nfregistry.org](http://nfregistry.org) today

“Knowing my son is part of the NF Registry helps me to trust this journey, even when I do not understand it.”

HEATHER FRAZIER AND HER SON RYAN who lives with NF1

“Sometimes there are good days and sometimes there are bad days, but it’s a matter of taking it one step at a time. Take the NF Registry on your journey with you.”

SEQUOYAH DANIEL-ROBINSON who lives with NF2

Join at [nfregistry.org](http://nfregistry.org) today
How do we know the NF Registry is working? Has it had results?

Yes, absolutely! In addition to connecting NF patients with specific research studies and clinical trials, NF Registry studies have provided critical information about where and what type of NF care is available to patients, especially for adults with NF. This is helping doctors and researchers better plan for future care and treatment options for patients with all forms of NF. Increased knowledge about specific manifestations of NF is also helping researchers improve treatment considerations, and patients are reporting back improved lifestyles. For example, an adult NF2 patient maintained his hearing because of the chance to use a specific drug, and the parent of an NF1 patient was given a more comprehensive range of treatment possibilities for her young child.

How private is private?

Your NF history is very personal and important to you. The NF Registry protects you by using the highest standards for data security (HIPAA, GDPR) and ethical use (Western Institutional Review Board). When filling out the patient questionnaire, you are free to answer (or not answer) the questions at your personal comfort level. The NF Registry will then only contact you if you give permission. Your information is not shared with the Children’s Tumor Foundation or any other organization, and it will never be used for fundraising or marketing.

Who should I contact?

For questions about the NF Registry, please email nfregistry@ctf.org.

Researchers: To apply to the Data Usage Committee, go to nfregistry.org and click “For Researchers.”

The NF Registry is a patient-driven resource for accelerating research and finding treatments for all forms of NF, including neurofibromatosis type 1 (NF1), and all types of schwannomatosis (SWN), including NF2-related schwannomatosis (NF2), formerly known as neurofibromatosis type 2. Sponsored by the Children’s Tumor Foundation, the NF Registry is a safe and effective tool to empower NF patients and their caregivers by inviting them to take an active role in advancing NF research.

If you or your child is living with NF, unite with thousands of others by joining the NF Registry today. Go to nfregistry.org and click on “Create a New Account” to get started.