NEWLY DIAGNOSED?

We are here for you and want you to know that you are not alone. Welcome to the FamilieSCN2A Foundation!

Who Are We?

We are an organization created by parents of children diagnosed with rare forms of Epilepsy and Autism as a result of a change in the SCN2A gene.

Our vision is to find effective treatments and a cure for SCN2A related disorders. Our mission is to improve the lives of those affected by SCN2A related disorders through research, public awareness, family support and patient advocacy.

We are a registered 501(c)(3) organization run almost entirely by parent volunteers.

We offer several programs to our global community. Visit our website to learn more: www.scn2a.org

PROGRAMS AVAILABLE

Global SCN2A Support Network is a private network of Facebook pages established by the FamilieSCN2A Foundation for parents, caregivers, patients and families of children with SCN2A related disorders.

SCN2A Family Event Grants serve to intentionally bring SCN2A families around the world together to cultivate a stronger community and deepen relationships.

SCN2A Patient Assistance Grants are in place to offer small grants to patients with SCN2A Disorders for necessary medical equipment, therapy devices, and educational aids associated with these conditions that are not covered through private insurance or other assistance programs.

Birthday Club is a way for us to show our love and celebrate our kids on their special day!

Newsletter and Emails help you to stay current on all that the Foundation is up to, SCN2A related research, and important SCN2A news by signing up to receive our newsletter and emails.

Warrior Wednesdays we welcome you to post pictures of your Warrior and share a story about how they / you are “fighting on”.

WWW.SCN2A.ORG