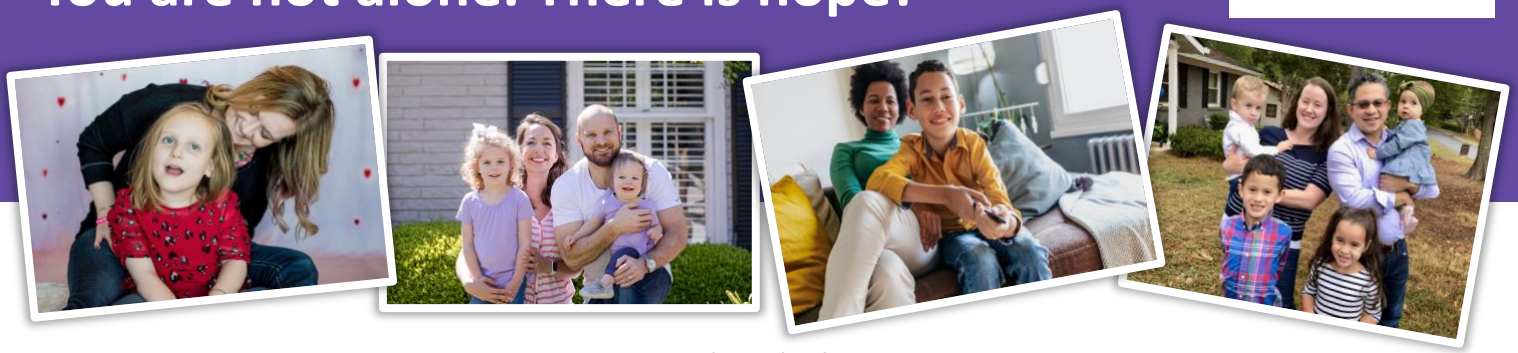
Welcome

You are not alone. There is hope!



Getting a rare disease diagnosis can be a scary time in a family's life. Know that you are not alone. Those with CACNA1A variants have different sets of symptoms and no two patients are exactly the same. Early intervention, intensive therapies and appropriate seizure control are critical factors in patient outcomes. We are a small but mighty community, and we are in this fight together.

Sign Up for Our Newsletter

Sign up with your email address to receive news about the latest CACNA1A research and Foundation announcements.

Take our Community Survey

Help us get to know you better. You will be given the option to upload your genetic testing report.

Follow Us on Social Media

Like, comment and share our posts on Instagram, Twitter & Facebook to raise CACNA1A awareness.

Start Your Own Fundraiser

When you're ready, check out the fundraising opportunities we have to help us accelerate research for potential therapies and ultimately a cure.

Visit the Newly Diagnosed Page

Our website includes information about understanding your genetic testing report, the various CACNA1A symptoms, figuring out your next steps and finding parent support.

Download Professional Brochure

Because CACNA1A variants are rare, many physicians are unfamiliar with CACNA1A symptoms and emergencies that may occur. Use this brochure to inform your providers.

Email info@cacna1a.org to schedule a call with a CACNA1A Foundation Board Member to learn more.