

Donate Today

HOW ARE YOU RALLYING FOR SCN2A TODAY?

Send us a <u>message</u> or <u>post pictures</u>!! Every new person who learns about SCN2A Disorders helps to move us one step closer to our goal: A CURE! #cureSCN2A

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PAG APPLICATION NOW OPEN!

FAMILIESCN2A SUMMER SEMINAR JULY 31 -AUGUST 1 2020

Columbus, Ohio

2-Day Seminar will include:

- **✓ Family-oriented learning sessions**
- Research opportunities
- Family-friendly activities

Learn more at: www.scn2a.org



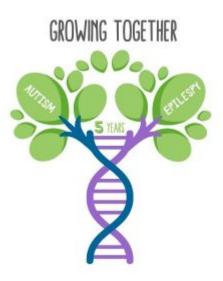




FamilieSCN2A Foundation moderates six private support networks for parents and caregivers of loved ones with SCN2A related disorders. These are a great resource for families to ask questions and find support. Click above to learn more!

2020 Annual Campaign

SCN2A Families: Hopefully you've read our recent email about streamlining 2020 fundraising efforts by launching our Annual Campaign TODAY! This campaign will solicit funds via email and snail mail letters and we need every one of you to participate by sharing this letter with your contacts. Think about people you may not have reached out to previously: neighbors, colleagues, businesses you frequent...



We are here to make this as easy as possible so, just send a list of mail/email contacts to info@scn2a.org and we'll send them on your behalf. Every dollar counts and if each family raised just \$100 we would easily reach our goal of raising \$50,000 by MAY 30! The first \$30,000 will go toward Patient Assistance Grants and everything after that will go toward research.

Researchers: Start thinking about your next SCN2A project!



RFA FOR 2020 GRANT CYCLE OPENS SOON!

IT'S NEVER TOO LATE TO ORDER AT THE SCN2A ONLINE STORE!

HELP US SPREAD AWARENESS!

The 2020 SCN2A Little Warriors Calendar is here! It's available in two sizes and can be shipped internationally. And, you can now order from our updated apparel shop year round! Click on the photos to reach the shop which can also be accessed using the STORE button in the top right corner of www.scn2a.org.



International SCN2A Awareness Day is TODAY, February 24.

Celebrate by ordering some new gear! We must raise awareness year round so whenever your items arrive, post pictures on your own social media of your family and friends wearing SCN2A shirts, bracelets or any purple/blue/green ANYtime!





Continue raising awareness even after today!

Rare Disease Day takes place on the last day of
February each year, this year being the rarest of
days: February 29! The main objective of Rare
Disease Day is to raise awareness among the
general public and decision-makers about rare
diseases and their impact on patients' lives. Click on
the logo to learn more.

So much news to share...quarterly newsletter coming soon!

Subscribe here

The FamilieSCN2A Foundation is a nonprofit 501(c)(3) organization. FEIN 47-3169795 Our mailing address is:

P.O. Box 82, East Longmeadow, MA 01028

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