

#CureSCN2A

International
CHROMOSOME
LOCATION
2q24.3
SCN2A
Awareness
February 24th
DAY!

www.scn2a.org

Start spreading the news...

International SCN2A
Awareness Day

Why February 24th?

The SCN2A gene is located on the long (q)
arm of Chromosome 2, at position 24.3



2/24

www.scn2a.org

We're celebrating our day...

'Celebrate' may not seem like the right word to use when talking about disease, but there is much to celebrate in the SCN2A Community and spreading awareness is ONE of the most important things we can do in building towards a cure. Please join us in celebrating our:

- Vibrant community of caring families who gladly share information and support one another so that no one feels alone
- Strong leaders who have propelled research and secured a \$600K [CZI grant](#) to expand our capacity to find cures
- Brilliant scientific partners dedicated to SCN2A research
- Possible Clinical Trials later this year!



**INTERNATIONAL SCN2A DAY IS RIGHT
AROUND THE CORNER. ORDER YOUR
APPAREL TODAY!**

Visit our store:
www.scn2a.org/shop.html

You've GOT to BE a part of it!

- Wear your SCN2A gear and post pix to social media. Tag #CureSCN2A #SCN2A #SCN2AAwarenessDay
- Order 2022 SCN2A Warrior Calendars [here](#). Give one to a friend!
- Ask your friends to follow us on social media
- Create a [Facebook fundraiser](#), it's easy and effective
- For a sample letter to send to friends and family, [email us](#)
- [Share your story](#)
- For more ideas, check out this [video](#) of the SCN2A community spreading awareness last year



USE this community-building grant to meet up with other SCN2A families in your area. Plan now, go whenever. **Get together or just announce it on 2/24!** [Email us for info.](#)

HOW CAN YOU GET INVOLVED?



SPREAD THE WORD

Share FamilieSCN2A materials, updates and events on social media and email within your network. Talk to people about our mission. Spread the word about SCN2A.



#SCN2A #CureSCN2A



@FamilieSCN2AFoundation



familiescn2afoundation



@FamilieSCN2A

Finding effective treatments and a cure is very expensive!
Simply put: To achieve our mission and vision, we need monetary funds.
Direct donation is the single action which has the biggest impact. Every donation matters, every dollar counts.

DONATE NOW



Donate at: www.scn2a.org

START FUNDRAISING

Without funding research cannot happen. We ask for your help as families, caregivers and friends of those affected by SCN2A to help the cause of finding effective treatments and a cure. Many people are happy to give; help provide them with the opportunity to do so by starting your own fundraising event.



Get started today! Email us at: info@scn2a.org

The FamilieSCN2A Foundation is almost completely operated by volunteers. In order to continue serving the needs of our growing community, we need your help. Join us and make a difference today!

GET INVOLVED



To join our team, email us at: info@scn2a.org

KICKING OFF THE 2022 ANNUAL CAMPAIGN

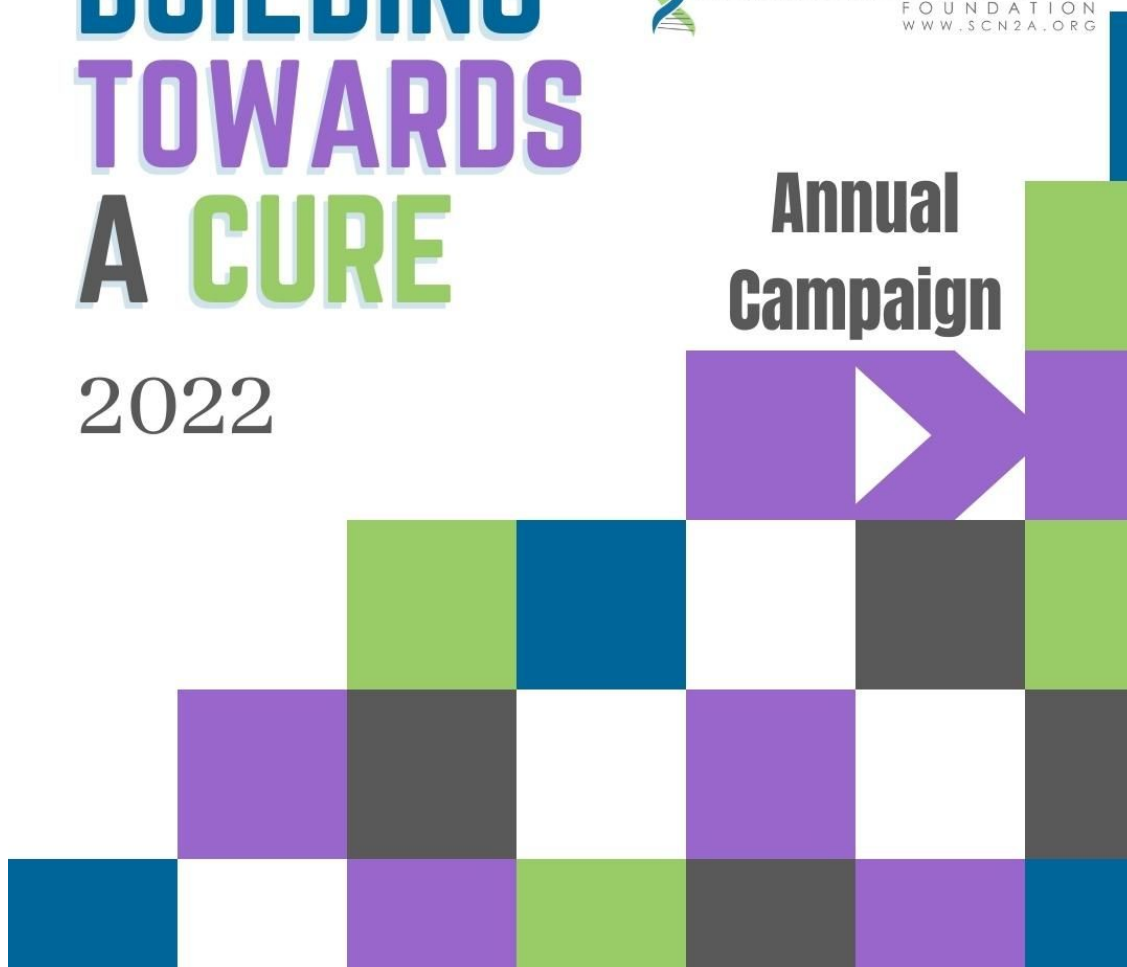
February kicks off our Annual Campaign! Thanks to you, every year we've met our goal, steadily building towards a cure. Now in our 7th year, the bar is raised to a goal of **\$70,000** in the next 3 months (Feb to April). Your donations will be put to work right away as we open the next round of Action Potential Research Grants on International SCN2A Awareness Day, 2/24. [Why wait? Give Today.](#)

BUILDING TOWARDS A CURE

2022



Annual
Campaign





Donate



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