



10 REASONS TO GIVE TO THE ANNUAL CAMPAIGN

MAY 1 - AUGUST 31

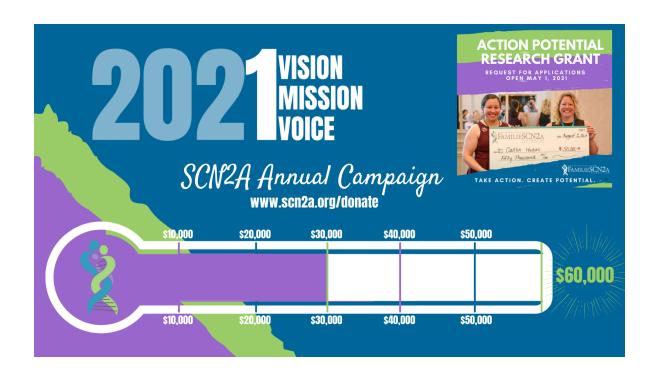
REASON #8 TO GIVE DONATIONS MATCHED!!



A GENEROUS DONOR WILL MATCH EVERY \$ RAISED BY AUG 31!

We haven't reached our \$60,000 goal, but it's not too late! A generous donor will match every dollar we raise by August 31 - JUST 4 WEEKS AWAY! **AND if we reach \$60,000, THE ENTIRE \$60K WILL BE MATCHED.** Please help us take advantage of this incredible opportunity to impact those with SCN2A related disorders. Share this with everyone you know!!

GIVE THE GIFT OF HOPE - DONATE TODAY



REASON #1 TO GIVE MEET BROOKS

REASON #2 TO GIVE GEARING UP FOR CLINICAL TRIALS PART 1



Brooks is almost 2 & is still waiting on a cure! Last year, his family gave back in a big way, hosting the top fundraiser during the Annual Campaign by sharing their SCN2A journey. <u>Visit Brook's Page</u> on our website and check out the toolkit for information on how to share your story.



Over the last 2 years, we have worked hard to fill the gaps in the process of getting to clinical trials. The FDA put out critical guidance on the very specific data they need to make decisions to approve new drugs and the biggest part, that we will answer with the CTRS, is how to define and demonstrate meaningful improvement. Learn more.

REASON #3 TO GIVE FINANCIAL SUPPORT FOR FAMILIES

REASON #4 TO GIVE GEARING UP FOR CLINICAL TRIALS PART 2



We help families access the proper equipment, therapy, or assistance they need to live their best lives. Read about how families have been impacted by this support.



At 17 months, Marty had his first seizure & life turned upside down for his family. Marty's Mom, Dad and big sister work hard every day to ensure Marty's VOICE is heard, most recently by participating in the SCN2A Clinical Trial Readiness Study. Visit Marty's Page.

REASON #5 TO GIVE SUPPORTING CONVERSATIONS BETWEEN PATIENT FAMILIES & RESEARCH/CLINICIANS

This year Families SCN2A is proud to host a virtual table talk series called RX Hope.

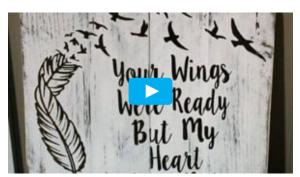
The purpose is to allow families unprecedented access to professional researchers & clinicians working diligently to find a cure for our children.



Virtual Table Talks

Families: <u>Click here</u> for the full schedule & to register for Virtual Table Talks.

REASON #6 TO GIVE IN MEMORY OF SCN2A ANGELS



REASON #7 TO GIVE MEET MATEO



Five-year-old Mateo is still waiting on a cure! Mateo's entire family advocates for SCN2A from the USA to Colombia! <u>Visit Mateo's Page</u> on our website and share your story, too.



The SCN2A Clinical Trial Readiness Study being led by Dr. Anne Berg at Ann & Robert H. Lurie Children's Hospital of Chicago

The purpose of this study is to provide information that will help researchers determine the performance of specific measures over time in children with SCN2A-DEE.

To learn more about the study and to begin your registration and consent process, click here.

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