REASON #7 TO GIVE
MEET MATEO
Five-year-old Mateo is still waiting on a cure! Mateo's entire family advocates for SCN2A. His Mom serves on the FamilieSCN2A Foundation Board and loves sending out welcome packets to new families. His Dad often helps with our A/V needs and his big brother spreads awareness every chance he gets. Mateo's grandpa hosted a charity golf tournament for SCN2A in Colombia! Visit Mateo's Page on our website and check out the toolkit for information on how to share your story.

GIVE THE GIFT OF HOPE - DONATE TODAY
REASON #1 TO GIVE
MEET BROOKS

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. Visit Brook's Page on our website and check out the toolkit for information on how to share your story.

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

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.

REASON #6
IN MEMORY OF OUR SCN2A ANGELS

Families: Click here for the full schedule & to register for Virtual Table Talks.
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.