

WWW.GRIN2B.COM

2018

FEBRUARY

APPROVED BY STATE OF ILLINOIS AS A VERIFIED NONPROFIT



MARCH

ORGANIZED FIRST GRIN2B
AWARENESS MONTH

RAISED OVER \$9000
IN FIRST MONTH FUNDRAISING

LAUNCHED PODCAST

2018

MAY

FORMED MEDICAL ADVISORY BOARD WITH FIRST 3 MEMBERS

SEPTEMBER

FIRST GRIN2B FOUNDATION FAMILY CONFERENCE IN ATLANTA,

attended by 23 families & 14 members of the Medical & Research community

BLOOD SAMPLES COLLECTED FROM 10 FAMILIES

2018

OCTOBER

ATTENDED GLOBAL GENES RARE PATIENT ADVOCACY SUMMIT

NOVEMBER

FORMED FUNDRAISING COMMITTEE

NOVEMBER-DECEMBER

RAISED OVER \$40,000 FOR RESEARCH GRANTS



ADDITIONAL 2018 ACCOMPLISHMENTS



SUPPORT

Private parent support group grew from 192 to 364 members, representing 39 countries.

Created interactive map on website.

Shared 3 blog posts on grin2b.com.

Two stories written by GRIN2B board members featured on The Mighty.

2018

AWARENESS





Created and sold GRIN2B Awareness apparel and accessories.

700+ followers on public Facebook page.

Joined Twitter and Instagram.

Shared 1st Newsletter - Nov 2018.

Created YouTube channel 1st Awareness video shared Nov 2018.

2018

RESEARCH

Recruited 2 additional members to our Medical Advisory Board.

Encouraged families to enroll in Simons VIP Connect registry.

GRIN2B added to GARD Genetic and Rare Diseases Information Center.

Began preliminary discussions with Advisors on Research grant application process.

ON THE HORIZON FOR 2019



Establish & fund first research grant cycle.

Collaborate with medical community on funding & organizing registry for GRIN-related disorders.

Plan various fundraisers with fundraising committee.

2nd GRIN2B Awareness month in March.

Investigate more Rare Disease Conferences to attend.

Create more newsletters, Awareness videos, podcasts and blog posts.

Design a brochure & other relevant informational materials (with translated versions).

New social media campaign featuring children impacted by a GRIN2B diagnosis.

2019