

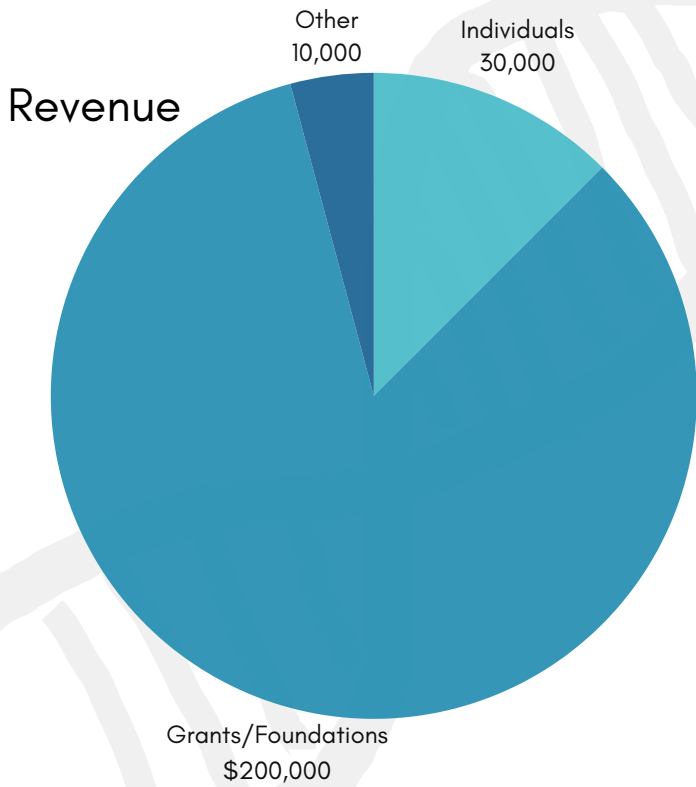


The TBCK Foundation is dedicated to serving families impacted by TBCK Syndrome through accelerating patient-led research, advocacy, education, and a dynamic system of support.

15 new families joined our Community Registry

ANNUAL REPORT

 Emily Durham, PhD joins as Research Engagement Director



4 TBCK Community Newsletters

\$15,795 raised for memorial fundraisers 

Hosted Bilingual Virtual TBCK Conference 

2 New Board Members

Launched collaborative Patient Registry with: **RAREX**

Funded over **\$110,000** of TBCK Research since 2020

 Hosted TBCK Community Monthly Chats

5th Annual Race for Rare Kids

Advocated for **100+** patients in **20+** countries 

1 New Scientific Advisory Board Member

2/1: Hosted TBCK Syndrome Awareness Day 

Attended 2 Rare Disease Conferences

 New logo!