ANNUAL IMPACT REPORT 2025

COMMUNITY IMPACT

Cure Mito Foundation serves the global Leigh syndrome community by funding research, amplifying patient voices, and strengthening education and support.

LEIGH SYNDROME PATIENT REGISTRY

430+ registry participants

48 countries represented

3 peer-reviewed publications

2025 FUNDED PROJECTS

PROJECT	RESEARCH PARTNER	AMOUNT
Gene Replacement Therapy SURF1/ECHS1	University of Texas Southwestern Medical Center	$218,556
Mitochondrial Replacement Therapy	Washington University in St. Louis	$70,000 granted $140,000 committed
Gut Microbiome Study	Icahn School of Medicine	$34,000
Drug Repurposing	Unravel Bioscience	$33,250
Drug Repurposing	University of California	$17,651.38

FUNDING

$1,071,000 ASSETS AS OF 12/31/2025

$258,000 DONATIONS

$220,000 GRANTS

EMPOWER AND INSPIRE 4TH LEIGH SYNDROME SYMPOSIUM

- 460 registered participants
- 237 live attendees
- 24 countries
- 6 continents
- 36 U.S. states

SHARING OUR JOURNEY IN A NEW PUBLICATION

Our new publication is published in Journal of Research Involvement and Engagement, and highlighting our evolution from Cure SURF1 to Cure Mito Foundation and our growth as a patient-led research organization., and the progress we've made together. Read the paper: tinyurl.com/curemitopaper

CURE MITO FOUNDATION

info@curemito.org | curemito.org