

Coordination of Rare Diseases at Sanford (CoRDS) is a disease agnostic platform with data for 2,029 rare diseases, 91 partner groups, 15,149 participants, 50 states, and 87 countries represented - as of June, 2022

Participants respond to 2 surveys - General and Leigh syndrome specific survey



Results and findings are always shared with the community. First results: curemito.org/results

## LEIGH SYNDROME GLOBAL PATIENT REGISTRY

Help us bring patients, families, and researchers together to gain a better understanding of Leigh syndrome



The participant owns his/her personal data and can withdraw the data from the registry at any time



For more information and to register visit:

curemito.org/ leighsyndromeregistry





ClinicalTrials.gov
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