

Why Register?

CoRDS



By sharing your experience with Leigh syndrome, you can help researchers answer critical questions
curemito.org/registry

Who can sign up?

Anyone with Leigh syndrome



What Do I Need?

A computer with an internet connection & an email address



No cost to register & gain access to your records/insights

Will my information be private?

All records are de-identified for approved researchers



What information is collected?

Patient Reported- CoRDS
curemito.org/cords

Medical Records- AllStripes
curemito.org/allstripes



Power in Numbers

By coming together, we will advance understanding of LS and build a stronger community



Advance Research

De-identified information is used to understand the natural history of LS, notify patients of clinical trial opportunities, and improve treatment and care of patients with LS



To learn more & to register visit
curemito.org/registry