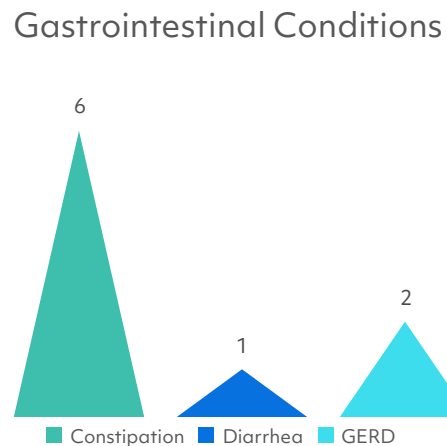
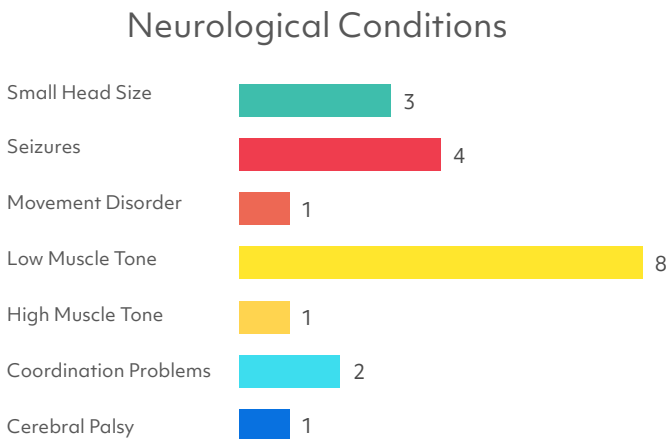
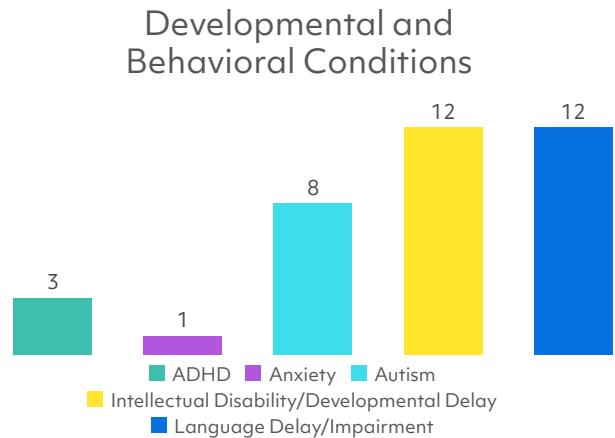
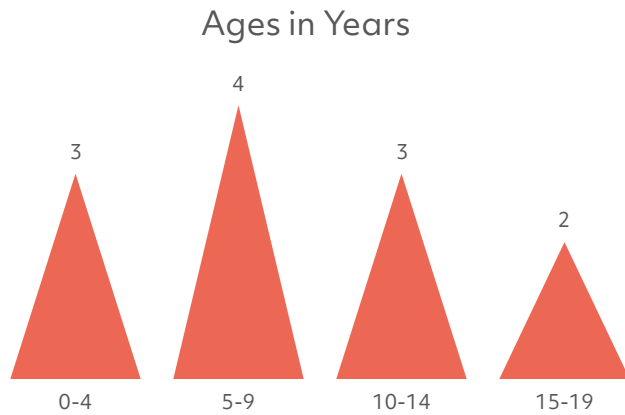


Simons Searchlight Registry Update EIF3F

Data in these four graphs are from the medical history information collected in Simons Searchlight from 12 participants with EIF3F-related syndrome (EIF3F-related Neurodevelopmental Disorder).

July 2025



NOTES: Graphs show counts of individuals in each category. Individual participants may appear in more than one category if they report multiple conditions. Graphs include individuals with pathogenic or likely pathogenic variants, and without additional variants.

How to participate?

The information in this report is made possible by the active participation of the EIF3F community! Progress for individuals in your community with EIF3F is shown below - log in to your simonssearchlight.org dashboard today to check for new surveys and tasks. Your data could hold the clues geneticists need to find answers.



STEP 1

Sign up
online

23



STEP 2

Provide your
genetic
lab report

17



STEP 3

Share your
important
medical history

13



STEP 4

Fill out
surveys

14



STEP 5

Provide a blood
sample if you are
interested

5



STEP 6

Update us
every year

Log in to see
next steps