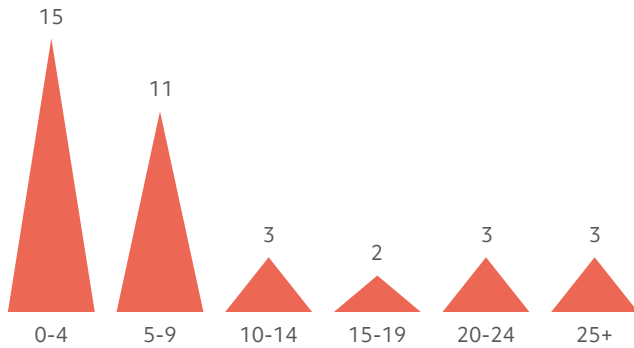


# Simons Searchlight Registry Update HNRNPH2

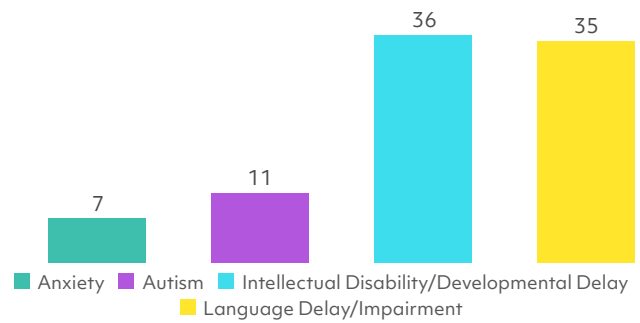
Data in these four graphs are from the medical history information collected in Simons Searchlight from 37 participants with HNRNPH2 (Bain Type of X-linked Syndromic Intellectual Disability).

July 2025

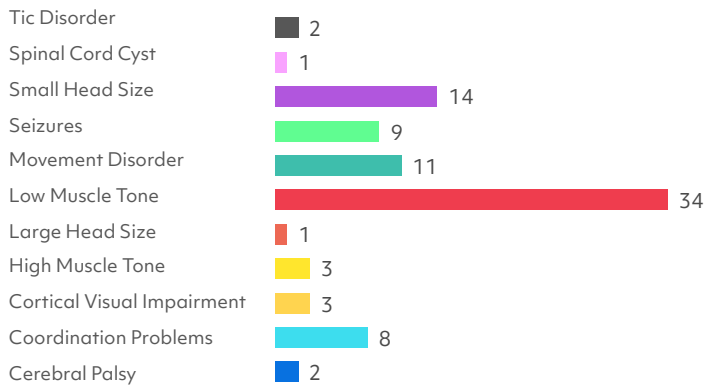
## Ages in Years



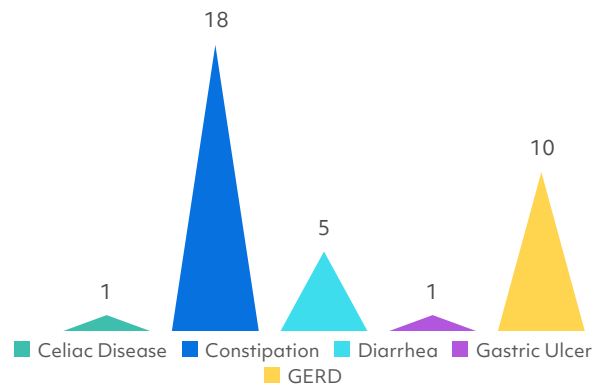
## Developmental and Behavioral Conditions



## Neurological Conditions



## Gastrointestinal Conditions



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 HNRNPH2 community! Progress for individuals in your community with HNRNPH2 is shown below - log in to your [simonssearchlight.org](https://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

74



### STEP 2

Provide your genetic lab report

58



### STEP 3

Share your important medical history

42



### STEP 4

Fill out surveys

50



### STEP 5

Provide a blood sample if you are interested

21



### STEP 6

Update us every year

Log in to see next steps