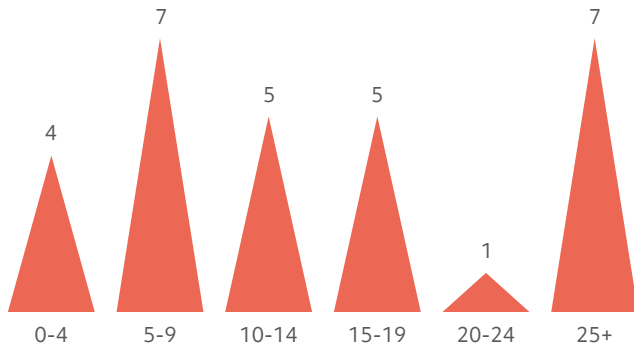


Simons Searchlight Registry Update 2p16.3 deletion & NRXN1

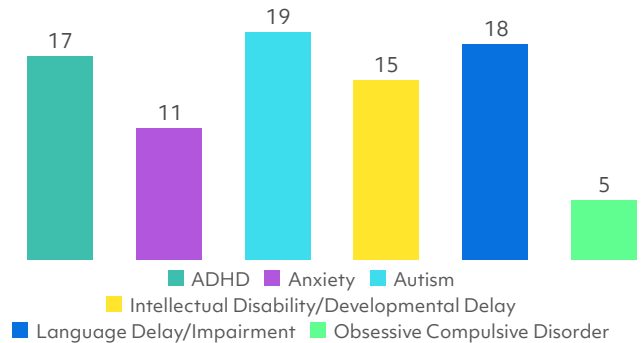
Data in these four graphs are from the medical history information collected in Simons Searchlight from 29 participants with a 2p16.3 deletion or NRXN1-related syndrome.

October 2025

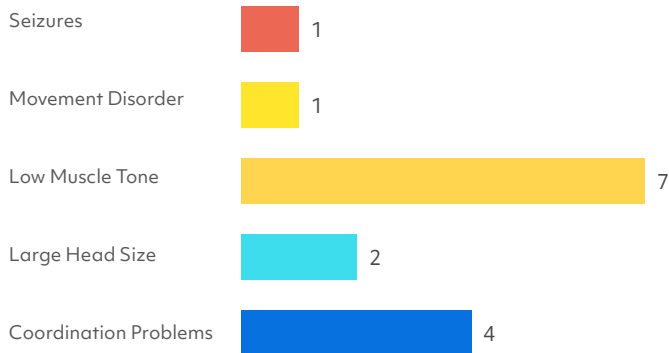
Ages in Years



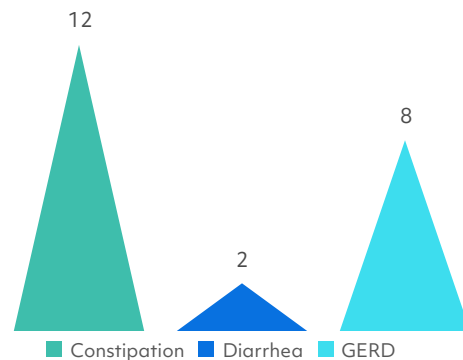
Developmental and Behavioral Conditions



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 2p16.3 deletion/NRXN1 community! Progress for individuals in your community with 2p16.3 deletion/NRXN1 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

98



STEP 2

Provide your
genetic
lab report

58



STEP 3

Share your
important
medical history

42



STEP 4

Fill out
surveys

46



STEP 5

Provide a blood
sample if you are
interested

5



STEP 6

Update us
every year

Log in to see
next steps