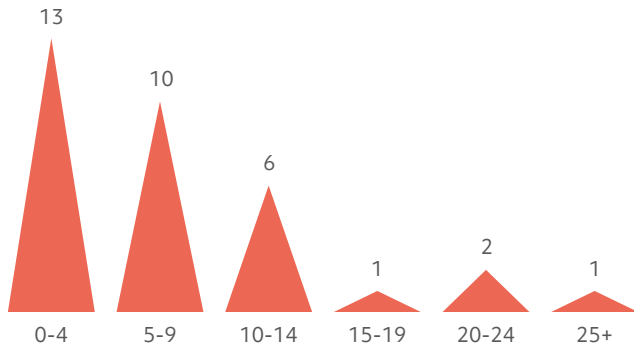


Simons Searchlight Registry Update ARID1B

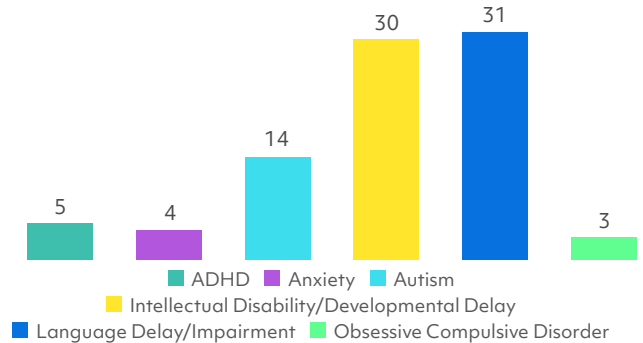
Data in these four graphs are from the medical history information collected in Simons Searchlight from 33 participants with ARID1B-related syndrome (Coffin-Siris Syndrome 1).

July 2026

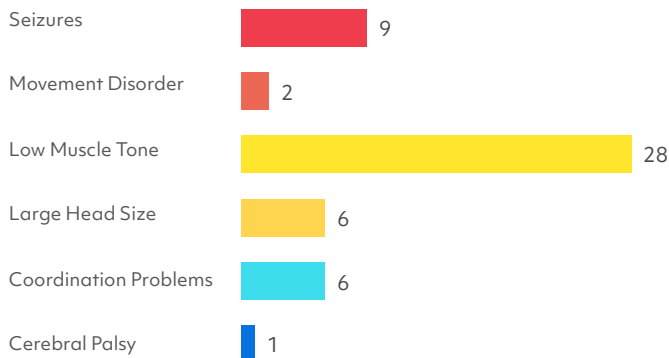
Ages in Years



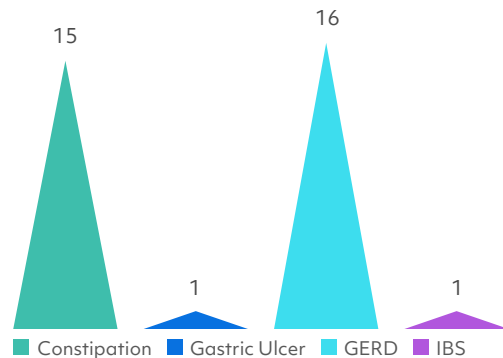
Developmental and Behavioral Conditions



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 only 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 ARID1B community! Progress for individuals in your community with ARID1B 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

78



STEP 2

Provide your
genetic
lab report

59



STEP 3

Share your
important
medical history

44



STEP 4

Fill out
surveys

47



STEP 5

Provide a blood
sample if you are
interested

7



STEP 6

Update us
every year

Log in to see
next steps