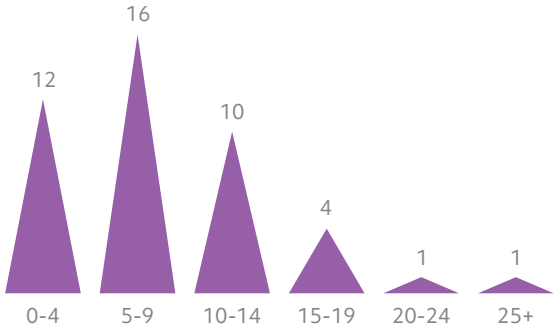
Simons Searchlight Registry Update CSNK2A1

July, 2022

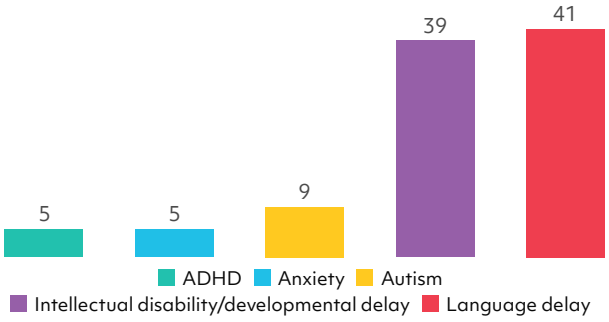
SIMONS SEARCHLIGHT

Data in these four graphs are from the medical history phone interviews collected in Simons Searchlight on 44 participants with CSNK2A1 (Okur-Chung neurodevelopmental syndrome).

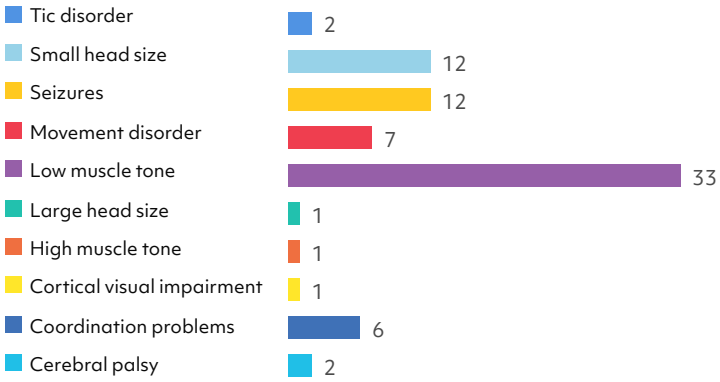
Ages in Years



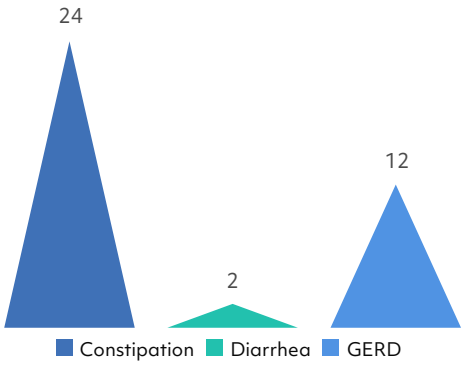
Developmental and Behavioral Conditions



Neurological Problems



Gastrointestinal Problems



Notes:

- Graphs show counts of individuals in each category. Individual participants may appear in more than one category if they report multiple conditions.

How to Participate

The information in this report is made possible by the active participation of the CSNK2A1 community! Progress for individuals in your community with CSNK2A1 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	STEP 2	STEP 3	STEP 4	STEP 5	STEP 6
Sign up online	Provide your genetic lab report	Share your important medical history	Fill out surveys	Provide a blood sample if you are interested	Update us every year
83	67	51	60	16	Log in to see next steps