Join our study on Neurexin 1 Deletion (2p16.3 deletion)

Study title:
Relative Diversity associated with Neurexin Trajectories (RaDiaNT)

Principal Investigator:

Dr. Louise Gallagher

Interested? To ask questions contact:

The Beacon Group:

recruitment.beacon@sickkids.ca

Version:2_29Nov2024

REB: 1000080839

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Are you someone with NRXN1 deletion or a family member (parent/sibling) of someone with NRXN1 deletion? Consider participating in our study:

What is the study about?

We are doing this study to better understand NRXN1 deletion to help us understand the variability in thinking skills, language, social skills and mental health outcomes. Family members (parents/siblings) are invited to participate to help us examine if other genetic factors outside of the NRXN1 gene might help to explain some of the variability in diagnosis and behaviours between individuals.

Who can participate?

We are looking for Individuals with NRXN1 deletion and their family members (parents/siblings)

What's involved?

We ask participants to:

- Provide a blood draw or saliva sample
- Complete some skills assessments
- Complete questionnaires and interviews

Are there benefits to participating?

- Participants will be given a gift certificate in recognition of their contribution.
- You can also receive volunteer hours for school.
- Your involvement will help doctors better understand NRXN1 deletion and why some people develop neurodevelopmental and mental health conditions and others do not.