

Our research study "A Genomic Approach to Precision Medicine for Autism and Neurodevelopmental Conditions" is enrolling children and adults who have had diagnostic genetic testing that showed a deletion or a duplication in chromosome 17q12 to understand how genetics can impact their healthcare. The study will take approximately 6 hours and involves collecting health information and completing questionnaires and assessments online.

To participate, contact Dr. Daniel Moreno De Luca or the PRISMA research staff at +1 780 492 4467 or prisma@ualberta.ca





