The Developmental Disorders Genetics Research Program (DDGRP)

Rare Genetics Research Study: 17q12 CNVs

The Developmental Disorders Genetics Research Program (DDGRP) is conducting research to improve our understanding of how deletions and duplications in chromosome 17q12 relate to developmental disorders. Through this research, we hope to develop as much information as possible about rare genetic disorders for families and doctors.

We are inviting individuals and family members affected by 17q12 copy number variants (CNVs) to enroll in this study. Participation in the study consists of an interview, behavioral assessments, and provision of a blood sample to create stem cells.

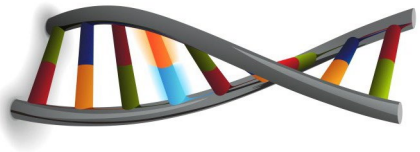
Participants are required to have prior clinical genetic testing and allow for contact with their doctors and medical record retrieval.

This study is led by:

Dr. Eric M. Morrow, MD, PhD

The Developmental Disorders Genetics Research Program

The Warren Alpert Medical School of Brown University and Emma Pendleton Bradley Hospital



To learn more about our program or to enroll, please contact us:

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