Contact

Renée Clark Registry Coordinator

Toll-free phone 866-744-7879

Email
Renee_Clark@unc.edu



The Carolina Institute Research Registry

is housed in the Institute's Intellectual and Developmental Disabilities Research Center.

The CIDD is part of a national network of IDDRCs funded by the National Institute of Child Health and Human Development specifically to promote research on developmental disorders.

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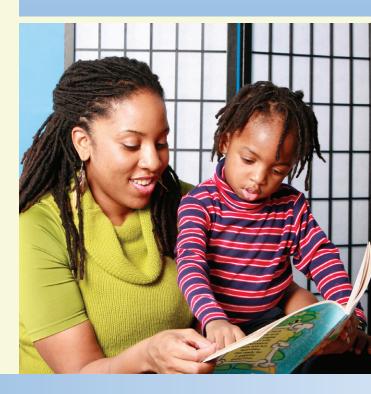




SERVICE | EDUCATION | RESEARCH

Research Registry

Links PEOPLE with opportunities for RESEARCH





The Carolina Institute Research Registry

Purpose

The Registry is a confidential database used to contact individuals or parents about research studies for which they may be eligible. Our goal is to mobilize studies and accelerate research on development and learning.

Participants

Children and adults who have a developmental disorder or condition such as developmental delay, ADHD, communication delay, autism spectrum disorder, intellectual disability, or a specific syndrome (e.g., Down, Prader-Willi, Turner, Williams, Angelman, Rett and others) may enroll.

Free, No Obligation, Private

The Registry is voluntary and you are not obligated to participate in studies. Your privacy is carefully safeguarded.

Contacts

The Registry will contact you about studies by mail, email, or phone.
Referrals are confidential and information about you is never released without your consent.

Studies

Research at The University of North Carolina at Chapel Hill covers a wide variety of topics such as genetics, language development, hearing, behavior and learning interventions, autism, clinical trials, risk factors, access to services, and more. Some studies involve surveys, assessments, or interviews. Some require travel; others do not.

Voluntary

You will be provided with the information you need to make an informed decision about participating in a research study.

How Do I Join?

You may enroll during your visit to the clinic at the Carolina Institute or request forms by calling the Registry Coordinator at 866-744-7879.

