PIXI
PARENTS AND INFANTS INTER(X)ACTION INTERVENTION

PIXI: Parent and Infant Inter(X)action Intervention

Our team is inviting families to engage in a research study to help us better understand the needs of families and infants with neurogenetic conditions. Participants will receive 20 weekly early intervention support. Please contact us if you have a child under the age of 18 months with a neurogenetic condition. We are especially interested in:

- Prader-Willi syndrome
- Down syndrome
- Angelman syndrome
- 22q deletion syndrome (DiGeorge)
- Fragile X syndrome
- Dravet syndrome

PIXI sessions and resources are provided at no cost to families

Participation Includes

- Weekly virtual visits (1 hour, via Zoom)
- Parent coaching on ways to support your child in play and daily routines
- Answering questions about your child's development and your experiences

Our Phone
(919) 248-8533

Email: Sam Scott at Sjscott@rti.org

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The University of North Carolina at Chapel Hill

For questions or concerns about your rights as a research subject, please contact the Institutional Review Board at 919-966-3113 or by email to IRB_subjects@unc.edu.