

I-InTERACT Preterm Parenting

Status: RECRUITING

Eligibility Criteria

Sex: ALL

Age: 3 years to 8 years old

Healthy Volunteers: This study is also accepting healthy volunteers

Inclusion Criteria:

- Born at < 32 weeks gestational age.
- Total T score of > 55 on the Child Behavior Checklist Total or Externalizing Behavior Scales OR Total T score of > 55 on the Eyberg Child Behavior Inventory total problem- or total intensity-scale.
- English is the primary spoken language in the home.

Exclusion Criteria:

- Is not 18 years or older.
- Participant will be excluded from the study if the child does not reside with the caregiver at least half-time; the caregiving situation is not stable (i.e., there must be no scheduled custody hearings).
- English is not the primary language spoken in the home.
- Caregivers with a psychiatric hospitalization in the past year.

Conditions & Interventions

Interventions:

BEHAVIORAL: I-InTERACT Parenting Intervention (I2P) and coaching sessions, BEHAVIORAL: I-InTERACT Parenting Microlearning Intervention (I2P Micro) and coaching sessions, OTHER: Internet Resources

Conditions:

Child Behavior Problem, Preterm, Parent-Child Relations, Parenting

Keywords:

telehealth, online learning, microlearning

More Information

Description:

Many children born very preterm experience behavior problems, and existing resources for parenting these children are lacking. A pilot trial established the effectiveness of a preterm parenting intervention, I-Interact Preterm (I2P). This study proposes a three-arm randomized controlled trial (RCT) comparing the established seven-session I2P program, a microlearning delivery mode (I2P-Micro), and an internet resource comparison group (IRC). Outcomes will be assessed at pretreatment, post-treatment (12 weeks later), and at an extended follow-up six months post-randomization. These outcomes include parenting behaviors, child behavior problems, and parent distress. It is anticipated that both I2P and I2P-Micro will result in significant improvements relative to the IRC condition, with greater utilization expected in the I2P-Micro group.

Contact(s): Zyah Flagg - zyah.flagg@cchmc.org

Principal Investigator:

Principal Investigator ID:

Phase: NA

IRB Number:

System ID: NCT06767293

Thank you for choosing StudyFinder. Please visit <http://studyfinder.cchmc.org> to find a Study which is right for you and contact clinicalstudies@cchmc.org if you have questions or need assistance.