



Research Opportunity: Calling parents of children who received a genetic diagnosis in the hospital (NICU, PICU, or CCU)

This study aims to explore parent/caregiver perceptions and emotions while receiving their child's genetic diagnosis in the hospital. Responses from this survey may be used to give guidance to medical providers on how to better give genetic diagnoses and support to families in the inpatient setting.

To participate, you must:

- Parents/caregivers 19 years or older who have a child that received a genetic diagnosis while in the NICU, PICU, or CCU.
- Your child's diagnosis cannot be Trisomy 21, Trisomy 18, Trisomy 13, or Turner syndrome as separate studies are likely more appropriate for parents of children with these conditions.

Please consider participating in this 10–15-minute survey using the HIPPA compliant Qualtrics survey link or QR code below.



https://unmcmmi.co1.qualtrics.com/jfe/form/SV_bgt7rdGDvSRBy1o

Questions? Contact Annika Paulsen at anpaulsen@unmc.edu and/or Kristen Fishler Malone kristen.fishler@unmc.edu

This study has been approved by the UNMC Office of Regulatory Affairs. (IRB 0385-25-EX)