

ONLINE EVENT

MARCH 27, 2024  
930am-12pm



# A DISCUSSION ABOUT FAMILY EXPERIENCES AFTER NEWBORN SCREENING

## FOR WHO?

Parents or caregivers of children who had a medical condition diagnosed from newborn screening (bloodspot, hearing, or cardiac tests) in DE, DC, MD, NJ, NY, PA, PR, USVI, VA, or WV

## WHAT?

A 2.5 hour online conversation with a small group of other families

## WHY?

To learn from families about what aspects of their family's life were impacted by newborn screening, how families are doing years after the diagnosis, and how they can be better supported

Spanish  
interpretation  
and closed  
captioning

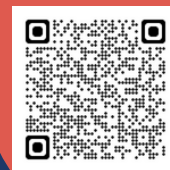
If you are interested, please fill out the  
form and we will contact you if you are  
selected to participate  
(click below or use QR code)

<https://forms.gle/2zFyPPmQztA5Wzgb6>



Families who are selected  
for the session will receive a  
stipend for their time

¿Interesado?  
Haga clic  
aquí para  
español



For More Information about Newborn Screening:

<https://www.babysfirsttest.org/newborn-screening/about-newborn-screening>



[NYMAC@ferre.org](mailto:NYMAC@ferre.org)



[NYMACgenetics.com](https://NYMACgenetics.com)

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