STUDY REGISTRY: CENTER FOR INNOVATIVE TRIALS IN CHILDREN AND ADULTS (TRIDENT) COVID-19

FREQUENTLY ASKED QUESTIONS (FAQ)

Information: Trident Study link https://www.mednax.com/about/education/creqs/ and then click on Neonatology.

Do we need hospital/IRB permission to handout the brochure?

- Your hospital may want to review the brochure prior to your clinical staff handing out the brochure. Each hospital site is unique, so if you are unsure, then it is best to reach out to the IRB and/or hospital compliance department.

Is this study IRB approved?

- Yes, this study has IRB approval through The Duke Health Institutional Review Board and is designed to encompass hospital sites following the specific components of the study.

What is the target population for brochure handouts?

- Any mother or infant who tests positive for COVID-19 during their hospitalization for delivery and/or after delivery or at discharge. This includes well baby nursery and NICU.

How does someone register?

- Simply refer the study subject to the study website or the QR Code on the brochure. This directs the participant to the screening questions.

Is their risk to any privacy information?

- This study complies with all HIPAA as mandated in 45 CRF Parts 160 and 164.