

Division of Laboratory Services

630 Hart Lane Nashville, TN 37216 615-262-6300

https://www.tn.gov/newbornscreeninglab

Newborn Screening Disorder:

3 Methylglutaconyl CoA Hydratase Deficiency (3MGA)

| Alternate Name(s) | MGA type I 3 alpha methylglutaconic aciduria type I 3 methylglutaconyl CoA hydratase deficiency 3MG CoA hydratase deficiency 3-MGCA type I (3-MGCA-1) |
|---|---|
| Analyte(s) Tested | • C4-DC + C5-OH |
| Methodology | Tandem Mass Spectrometry |
| TDH Requisition Form | PH-1582 Form Requests: Contact state lab by email or fax. Include facility name, address, phone number and contact person on your request. Fax: (615) 262-6455 Email: DCLAB.supply@tn.gov |
| Acceptable Specimen | Dried blood spots on filter paper collected from infants less than 6 months of age |
| Collection Information | Optimal specimen: Collect at 24 hours + 1 minute of life Acceptable specimen: Collect 24-48 hours of life If transfused: Recollect 4 days post transfusion |
| Shipping Information | Health Departments and Birthing Hospitals Private Clinics and Midwives |
| Screening Results | Tennessee Newborn Screening's Secure Remote Viewer (SRV) Healthcare providers must be registered to view and print patient result reports. To Register: Complete and submit SRV Access Form (PH-3909). NBS List of Screened Disorders and Mailer Explanations |
| Laboratory Location Performing Testing | Nashville, TN |