

Confirmed Disease Identified

The Tennessee Newborn Screening (NBS) test results can be affected by baby's age, medical or treatment status at the time of specimen collection; the quality and quantity of the specimen and/or other variables and may not detect all affected babies. The possibility of false negative or false positive results must always be considered when screening children for metabolic/genetic disorders.

if you have diagnosed a child with one of the Metabolic/Genetic Disorders on the NBS Panel, but were not notified by the Program, please complete the form below.

Thank you!

Child's First Name:

Child's Last Name:

Child's Date of Birth:

Was a referral received for this infant?

☐ Yes

☐ No

Hospital of Birth:

- ☐ Athens Regional Medical Center
- ☐ Baby & Co - Knoxville
- ☐ Baby & Co - Nashville
- ☐ Ballad Health - Bristol Regional Medical Center
- ☐ Baptist Dekalb Hospital
- ☐ Baptist Health System of East TN
- ☐ Baptist Memorial Hospital - Union City
- ☐ Baptist Memorial Hospital - Memphis
- ☐ Baptist Memorial Hospital - Tipton
- ☐ Big South Fork Medical Center
- ☐ Blanchfield Army Community Hospital
- ☐ Blount Memorial Hospital
- ☐ Cookeville Regional Medical
- ☐ Cumberland Medical Center
- ☐ East TN Children's Hospital
- ☐ Erlanger East Women's Services
- ☐ Erlanger Medical Center
- ☐ The Farm Midwifery Center
- ☐ Franklin Woods Community Hospital
- ☐ Ft. Sanders Regional Medical Center
- ☐ Greeneville Community Hospital - East
- ☐ Hardin Medical Center
- ☐ Hendersonville Medical Center
- ☐ Henry County Medical Center
- ☐ Indian Path Community Hospital
- ☐ Infinity Birthing Center
- ☐ Jackson-Madison County General Hospital
- ☐ Jellico Community Hospital
- ☐ Johnson City Medical Center
- ☐ Lakeway Regional Hospital
- ☐ LeBonheur Children's Hospital
- ☐ Leconte Medical Center
- ☐ Lincoln Medical Center
- ☐ Lisa Ross Birth and Women's Center
- ☐ Livingston Regional Hospital
- ☐ Maury Regional Hospital
- ☐ McKenzie Regional Hospital
- ☐ McNairy Regional Hospital
- ☐ Methodist Healthcare - Germantown
- ☐ Methodist Hosp - South
- ☐ Methodist Medical Center - Oak Ridge
- ☐ Metro Nashville General Hospital
- ☐ Morristown-Hamblen Healthcare
- ☐ Natchez Trace Maternity Center
- ☐ Northcrest Medical Center
- ☐ Park Ridge East Hospital
- ☐ Parkwest Medical Center
- ☐ Private Residence
- ☐ Regional Medical Center at Memphis
- ☐ Riverview Regional Medical Center
- ☐ Roane Medical Center
- ☐ Southern TN Regional Health - Lawrenceburg
- ☐ Southern TN Regional Health - Pulaski
- ☐ Southern TN Regional Health - Winchester
- ☐ St. Francis Hospital
- ☐ St. Francis Hospital - Bartlett
- ☐ St. Thomas - Midtown
- ☐ St. Thomas - River Park Hospital
- ☐ St. Thomas - Rutherford Hospital
- ☐ Stonecrest Medical Center
- ☐ Sumner Regional Medical Center
- ☐ Sweetwater Hospital Association
- ☐ Sycamore Shoals Hospital
- ☐ Takoma Regional Hospital
- ☐ Tennova Healthcare - Newport Medical Center
- ☐ Tennova Healthcare - Clarksville
- ☐ Tennova Healthcare - Cleveland
- ☐ Tennova Healthcare - Jamestown
- ☐ Tennova North Knoxville Medical Center

- ☐ Tennova Physicians Regional Medical Center
- ☐ Tennova Turkey Creek Medical Center
- ☐ Tristar Centennial Medical Center
- ☐ Tristar Horizon Medical Center
- ☐ Tristar Southern Hills Medical Center
- ☐ Tristar Summit Medical Center
- ☐ UT Medical Center
- ☐ Vanderbilt Tullahoma Harton Hospital
- ☐ Vanderbilt University Hospital
- ☐ Vanderbilt Wilson County Hospital
- ☐ Wellmont Holston Valley Medical Center
- ☐ West TN Healthcare - Dyersburg Hospital
- ☐ West TN Healthcare - North Hospital
- ☐ West TN Healthcare - Volunteer Hospital
- ☐ Williamson Medical Center
- ☐ Women's Wellness & Maternity
- ☐ Other

If other birth location, please elaborate:

Mother's First Name:

Mother's Last Name:

City of Mother's Residence:

State or Territory of Mother's Residence:

- ☐ Tennessee
- ☐ Alabama
- ☐ Alaska
- ☐ Arizona
- ☐ Arkansas
- ☐ California
- ☐ Colorado
- ☐ Connecticut
- ☐ Delaware
- ☐ District of Columbia
- ☐ Florida
- ☐ Georgia
- ☐ Hawaii
- ☐ Idaho
- ☐ Illinois
- ☐ Indiana
- ☐ Iowa
- ☐ Kansas
- ☐ Kentucky
- ☐ Louisiana
- ☐ Maine
- ☐ Maryland
- ☐ Massachusetts
- ☐ Michigan
- ☐ Minnesota
- ☐ Mississippi
- ☐ Missouri
- ☐ Montana
- ☐ Nebraska
- ☐ Nevada
- ☐ New Hampshire
- ☐ New Jersey
- ☐ New Mexico
- ☐ New York
- ☐ North Carolina
- ☐ North Dakota
- ☐ Ohio
- ☐ Oklahoma
- ☐ Oregon
- ☐ Pennsylvania
- ☐ Rhode Island
- ☐ South Carolina
- ☐ South Dakota
- ☐ Texas
- ☐ Utah
- ☐ Vermont
- ☐ Virginia
- ☐ Washington
- ☐ West Virginia
- ☐ Wisconsin
- ☐ Wyoming
- ☐ American Samoa
- ☐ Guam
- ☐ Northern Mariana Islands
- ☐ Puerto Rico
- ☐ U.S. Virgin Islands

Disease Group that Diagnosis belongs to:

- ☐ Amino Acid Disorder
- ☐ Fatty Acid Disorder
- ☐ Organic Acid Disorder
- ☐ Galactosemia
- ☐ Hemoglobinopathy
- ☐ Lysosomal Disorder
- ☐ Endocrine Disorder
- ☐ Immune Deficiency Disorder
- ☐ Other

Amino Acid Disorder Diagnosis (Check all that apply):

- ☐ Argininemia
- ☐ Argininosuccinic Aciduria (Argininosuccinate lyase deficiency)
- ☐ β -Ketothiolase Deficiency (aka SKAT)
- ☐ Citrullinemia Type I (Classic - Argininosuccinate Synthetase Deficiency)
- ☐ Citrullinemia Type II (Citrin Deficiency)
- ☐ Homocystinuria
- ☐ Hypermethioninemia
- ☐ Hyperornithinemia -Hyperammonemia-Homocitrullinuria
- ☐ Hyperornithinemia with Gyrate Atrophy
- ☐ Hyperphenylalaninemia due to Phenylalanine Hydroxylase Deficiency
- ☐ Hyperphenylalaninemia due to GTP Cyclohydrolase I Deficiency
- ☐ Hyperphenylalaninemia due to Pterin-4-Alpha-Carbinolamine Dehydratase Deficiency
- ☐ Hyperphenylalaninemia due to 6-Pyruvoyltetrahydropterin Synthase Deficiency
- ☐ Hyperphenylalaninemia - Defects of biopterin co factor biosynthesis
- ☐ Hyperphenylalaninemia - Defects of biopterin co factor regeneration
- ☐ Hyperphenylalaninemia-Dihydropteridine reductase
- ☐ Maple Syrup Urine Disease
- ☐ Nonketotic Hyperglycinemia
- ☐ Ornithine Transcarbamylase Deficiency
- ☐ Phenylketonuria
- ☐ Tyrosinemia Type I
- ☐ Tyrosinemia Type II
- ☐ Tyrosinemia Type III

Fatty Acid Disorder Diagnosis (Check all that apply):

- ☐ 2,4 Dienoyl CoA Reductase Deficiency
- ☐ Carbamoyl Phosphate Synthetase I Deficiency I
- ☐ Carnitine Palmitoyltransferase Type I Deficiency
- ☐ Carnitine Palmitoyltransferase Type II Deficiency
- ☐ Carnitine Acylcarnitine Translocase Deficiency
- ☐ Systemic Primary Carnitine Deficiency or Carnitine Uptake Defect/Carnitine Transport Defect
- ☐ Glutaric Acidemia Type II (aka Multiple AcylCoA Dehydrogenase Deficiency)
- ☐ Long-Chain 3-Hydroxyacyl-CoA Dehydrogenase Deficiency
- ☐ Medium-Chain Acyl-CoA Dehydrogenase Deficiency
- ☐ Medium/Short Chain 3-hydroxyacyl-CoA Dehydrogenase Deficiency
- ☐ Short Chain Acyl-CoA Dehydrogenase Deficiency
- ☐ Trifunctional Protein Deficiency
- ☐ Very Long Chain Acyl-CoA Dehydrogenase Deficiency

Organic Acid Disorder Diagnosis (Check all that apply):

- ☐ 2-Methyl-3-Hydroxy Butyric Aciduria
- ☐ 2-Methylbutyryl Glycinuria
- ☐ 3-Hydroxy-3-Methylglutaryl Aciduria
- ☐ 3-Methylcrotonyl-CoA Carboxylase Deficiency
- ☐ 3-Methylglutaconic Aciduria
- ☐ Glutaric Acidemia Type I
- ☐ Isobutyrylglycinuria
- ☐ Isovaleric Acidemia
- ☐ Malonic Aciduria
- ☐ Methylmalonic Acidemia due to Methylmalonyl-CoA Mutase Deficiency
- ☐ Methylmalonic Acidemia - Cobalamin A (CblA) or Cobalamin B (CblB) Deficiency
- ☐ Methylmalonic Acidemia with Homocystinuria (Cbl C or D Type)
- ☐ Multiple CoA Carboxylase Deficiency
- ☐ Propionic Acidemia
- ☐ Ethylmalonic Encephalopathy
- ☐ Methylmalonic Acidemia (Methylmalonyl-CoA Mutase Deficiency)
- ☐ Methylmalonic Acidemia (Cobalamin Disorders)
- ☐ Succinyl-CoA Synthetase Deficiency (SUCLA2)
- ☐ Multiple Carboxylase Deficiency (Holocarboxylase Synthase Deficiency)

Galactosemia Disorder Diagnosis (Check all that apply):

- ☐ GALT Deficiency (Classical Galactosemia)
- ☐ Galactokinase Deficiency
- ☐ Epimerase Deficiency

Hemoglobinopathy Disorder Diagnosis (Check all that apply):

- ☐ Sickle Cell Anemia
- ☐ Sickle Beta Thalassemia
- ☐ Sickle Cell Disease
- ☐ Various Other Hemoglobinopathies

Lysosomal Disorder Diagnosis (Check all that apply):

- ☐ Pompe
- ☐ Fabry
- ☐ Gaucher
- ☐ Krabbe
- ☐ Mucopolysaccharidosis Type I

Endocrine Disorder Diagnosis (Check all that apply):

- ☐ Congenital Adrenal Hyperplasia
- ☐ Congenital Hypothyroidism

Immune Deficiency Disorder Diagnosis (Check all that apply):

- ☐ Severe Combined Immunodeficiency
- ☐ Other T-Cell related lymphocyte deficiencies

Other Disorder Diagnosis (Check all that apply):

- ☐ X-Linked Adrenoleukodystrophy
- ☐ Biotinidase Deficiency
- ☐ Cystic Fibrosis
- ☐ Spinal Muscular Atrophy

Diagnosis Date:

Confirmatory Laboratory Studies:

Upload of Confirmatory Lab Studies:

Clinical Condition of Child that Led to Evaluation:

- ☐ Well
- ☐ Unwell
- ☐ Hospitalized
- ☐ Deceased

Current Status of Child (Check All that Apply):

- ☐ Well
- ☐ Unwell
- ☐ Hospitalized
- ☐ Deceased

Treatment:

Treatment Date Start:

Treating Specialist:

Provider Name:

Provider Phone:

Submitted by (Name):

If you would like a confirmation email of this form submission emailed to you, please include an email address here:
