Ohio Newborn Screening Program:
CONFIRMED CASE REPORTING FORM

Practitioner Name:	**Practitioner Address:**	**Date:**
Baby's Last Name:	**First Name:**	**Date of Birth:**
Mother's Name:	**Birth Hospital:**	**NBS kit number:**

Check the appropriate final diagnosis and return this form with documentation of the diagnosis, confirming laboratory results, and treatment information to the Ohio Newborn Screening Program, 8955 East Main Street, Bldg 22, Reynoldsburg OH 43068 or Fax to 614-644-4648.

Amino Acid Disorders

- ☐ Argininemia (ARG)
- ☐ Argininosuccinic Aciduria (ASA)
- ☐ Citrullinemia Type I (CIT)
- ☐ Citrullinemia Type II (CIT II)
- ☐ Homocystinuria (HCY)
- ☐ Hypermethioninemia (MET)
- ☐ Hyperphenylalaninemia - Benign (H-PHE)
- ☐ Hyperphenylalaninemia - Biopterin Cofactor Biosynthesis Defect (BIOPT BS)
- ☐ Hyperphenylalaninemia - Biopterin Cofactor Regeneration Defect (BIOPT REG)
- ☐ Maple Syrup Urine Disease (MSUD)
- ☐ Phenylketonuria (PKU)
- ☐ Tyrosinemia Type I (TYR I)
- ☐ Tyrosinemia Type II (TYR II)
- ☐ Tyrosinemia Type III (TYR III)

Organic Acid Disorders

- ☐ 2-Methylbutryl-CoA Dehydrogenase Deficiency (2MBG)
- ☐ 3-hydroxy-3-methylglutaryl-CoA lyase deficiency (HMG)
- ☐ 3-Methylcrotonyl-CoA Carboxylase Deficiency (3-MCC)
- ☐ β-Ketothiolase Deficiency (BKT)
- ☐ Glutaric Acidemia Type I (GA1)
- ☐ Isobutyryl-CoA Dehydrogenase Deficiency (IBG)
- ☐ Isovaleric Acidemia (IVA)
- ☐ Methylmalonic Acidemia (cobalamin disorders) (Cbl A, B)
- ☐ Methylmalonic Acidemia (methylmalonyl-CoA mutase) (MUT)
- ☐ Methylmalonic Acidemia with Homocystinuria (Cbl C, D)
- ☐ Multiple Carboxylase deficiency (MCD)
- ☐ Propionic Acidemia (PROP)

Not Otherwise Listed

- ☐ Other (specify):

Fatty Acid Disorders

- ☐ Carnitine Acylcarnitine Translocase Deficiency (CACT)
- ☐ Carnitine Palmitoyl Transferase Deficiency Type II (CPT II)
- ☐ Carnitine Uptake Defect (CUD)
- ☐ Glutaric Acidemia Type II (GA2)
- ☐ Long-Chain Hydroxyacyl-CoA Dehydrogenase Deficiency (LCHAD)
- ☐ Medium-Chain Acyl-CoA Dehydrogenase Deficiency (MCAD)
- ☐ Short Chain Acyl-CoA Dehydrogenase Deficiency (SCAD)
- ☐ Trifunctional Protein Deficiency (TFP)
- ☐ Very Long-Chain Acyl-CoA Dehydrogenase Deficiency (VLCAD)

Endocrine Disorders

- ☐ Congenital Adrenal Hyperplasia (CAH)
- ☐ Congenital Hypothyroidism (CH)

Lysosomal Storage Disorders

- ☐ Glycogen Storage Disease Type II (Pompe) (GSD II)
- ☐ Krabbe Disease
- ☐ Mucopolysaccharidosis type I (MPS-1)

Hemoglobin Disorders

- ☐ Hemoglobinopathy - Sickle Cell Anemia (Hb SS)
- ☐ Hemoglobinopathy - Sickle β-Thalassemia (Hb S/β Th)
- ☐ Hemoglobinopathy - Sickle C Disease (Hb S/C)
- ☐ Hemoglobinopathy - Other (specify):

Other Conditions

- ☐ Biotinidase Deficiency (BIOT)
- ☐ Cystic Fibrosis (CF)
- ☐ Galactosemia - Classic (GALT)
- ☐ Severe combined immunodeficiency (SCID)
- ☐ T-Cell Related Lymphocyte Deficiencies (specify)