

Georgia's Newborn Screening Panel and Disorder Specific Information

<p>Organic Acid Disorders</p>	<p>Beta Ketothiolase (BKT)</p> <p>Cobalamin A and B Deficiency (Cbl A,B)</p> <p>Glutaric Acidemia type I (GA1)</p> <p>3-OH 3-CH Glutaric Aciduria (HMG)</p> <p>Isovaleric Acidemia (IVA)</p> <p>3 Methylcrotonyl-Co A Carboxylase Deficiency (3MCC)</p> <p>Multiple Carboxylase Deficiency (MCD)</p> <p>Methylmalonic Acidemia (MMA)</p> <p>Propionic Acidemia</p>
<p>Fatty Acid Oxidation Disorders</p>	<p>Carnitine Uptake Defect</p> <p>Long Chain 3 hydroxyl acyl-CoA dehydrogenase Deficiency (LCHADD)</p> <p>Medium Chain acyl-CoA dehydrogenase Deficiency</p> <p>Trifunctional Protein Deficiency (TFP)</p> <p>Very Long-chain acyl-CoA dehydrogenase Deficiency (VLCADD)</p>
<p>Amino Acid Disorders</p>	<p>Argininosuccinic Acidemia</p> <p>Citrullinemia</p> <p>Homocystinuria</p> <p>Maple Syrup Urine Disease (MSUD)</p> <p>Phenylketonuria (PKU)</p>

	Tyrosinemia
Hemoglobinopathy Disorders	Sickle Cell Anemia Sickle Beta Thalassemia Sickle C Disease Other Hemoglobin Variants
Other Disorders	Biotinidase Deficiency Congenital Adrenal Hyperplasia (CAH) Congenital Hypothyroidism (CH) Critical Congenital Heart Disease (CCHD) Cystic Fibrosis (CF) Galactosemia Hearing Impairment Severe Combined Immunodeficiency (SCID)