

Office of Newborn Screening  
Confirmed Case Data Report--2012

<b>DISORDER</b>	<b>Total-Primary Disorders</b>	<b>Secondary Findings/Traits</b>
Congenital Hypothyroidism	59	8 Transient
Congenital Adrenal Hyperplasia	8	
Sickle Cell Anemia	8	
Sickle Beta Thalassemia	1	879 combined hemoglobin traits
Sickle C Disease	2	
Biotinidase Deficiency	7	6 partial
Galactosemia	0	
Phenylketonuria	6	
Maple Syrup Urine Disease	0	
Homocystinuria	0	
Citrullinemia	1	
Argininosuccinic Acidemia	1	
Tyrosinemia Type 1	0	
Carnitine Uptake Defect	1	
Medium-chain Acyl-CoA Dehydrogenase Deficiency	5	
Very Long-chain Acyl-CoA Dehydrogenase Deficiency	2	
Long-chain 3-OH Acyl-CoA Dehydrogenase Deficiency	0	
Trifunctional Protein Deficiency	0	
Isovaleric Acidemia	0	
Glutaric Acidemia Type 1	0	
3-Hydroxy-3-Methylglutaric Aciduria	0	
3-Methylcrotonyl CoA Carboxylase Deficiency	2	
Multiple Carboxylase Deficiency	0	
Methylmalonic Acidemia (mutase deficiency)	0	
Methylmalonic Acidemia (cobalamin defects)	2	
Propionic Acidemia	1	
Beta-Ketothiolase Deficiency	0	
Cystic Fibrosis	25	106 carrier, 1 non-classical
<b>Total Primary Disorders</b>	<b>131</b>	
<b>Total Secondary Findings</b>		<b>1,000</b>

  

Occurrent Births 2012	87,274	
Primary Bloodspot Disorder Total	131	
Secondary Disorders Detected	1,000	
Total Confirmed:		
Primary & Secondary Disorders	1,131	
Total Confirmed Hearing Loss	133	