Disorders screened for by the CT Newborn Screening Program

Amino Acid Disorders	**Screening Began**
Argininemia/Arginase Deficiency (ARG)	1/1/05
Citrullinemia, Argininosuccinic Acid Synthetase Deficiency. (ASD) Argininosuccinase Lyase Deficiency (ALD)	10/1/04
Hyperammonemia, hyperornithinemia, homocitrullinemia (HHH)	1/1/05
Homocystinuria / Hypermethionemia	1993
Ornithine Transcarbamylase Defiency (OTC)	1/1/05
Maple Syrup Urine Disease (MSUD)	1993
Nonketotic Hyperglycinemia (NKH)	1/1/05
Phenylketonuria (PKU)	1964
Tyrosinemia (TYR)	5/1/04

Fatty Acid Oxidation Disorders	**Screening Began**
Carnitine Palmitoyl Transferase Deficiency 1 (CPT I)	10/1/04
CPT II / Carnitine/Acylcarnitine Translocase (CACT)	10/1/04
SCADD / Ethylmalonic Acidemia (EDA)	1/1/05
Glutaric Acidemia II (GA II) / Multiple acyl-CoA Dehydrogenase Deficiency (MADD)	10/1/04
LCHADD / Trifunctional Protein Deficiency (TFP)	5/1/04
Medium Chain Acyl-CoA Dehydrogenase Deficiency (MCADD)	5/1/04
Very Long Chain Acyl-CoA Dehydrogenase Deficiency. (VLCADD)	10/1/04
2,4 Dienoyl CoA Reductase Deficiency (DCR)	11/01/04

Organic Acidurias	**Screening Began**
Beta-Ketothiolase Deficiency (BKT)	11/01/04
Glutaric Acidemia I (GA I)	11/01/04
3-Hydroxy-3-Methylglutaryl-CoA Lyase (HMG) / Multiple CoA Carboxylase (MCD) / 3-Methylcrotonyl-CoA Carboxylase (3MCC)	11/01/04
Isovaleric Acidemia (IVA)	11/01/04
Malonic Aciduria (MA)	1/1/05
Methylmalonic / Propionic Acidemia (MMA/PPA)	11/01/04

Other Diseases	**Screening Began**
Biotinidase Deficiency	1993
Congenital Adrenal Hyperplasia 17-OHP (CAH)	1997
Carnitine Transporter Deficiency	10/1/04
Hemoglobinopathies	1990
Congenital Hypothyroidism (CH) / TSH	1976
Galactosemia	1964