1964 PKU

1978 Congenital Hypothyroidism

1987 Hemoglobinopathies

1992 Galactosemia

1994 Congenital Adrenal Hyperplasia

1997 Voice Response System (VRS)

04/2004 Biotinidase Deficiency

10/2004 **Amino Acid Disorders:**
Citrullinemia (CIT)
Homocystinuria (HCY)
Maple Syrup Urine Disease (MSUD)
Tyrosinemia (TYR)
Argininosuccinate aciduria (ASA)

Organic Acid Disorders:
Propionic Acidemia (PROP)
Methylmalonic Acidemia (Vitamin B12 Disorders) (CBL, A,B)
Methylmalonic Acidemia (methylmalonyl-CoA mutase) (MUT)

Fatty Acid Disorders:
Medium chain acyl-CoA dehydrogenase deficiency (MCAD)
Carnitine Uptake Defect (CUD)

10/2006 **Organic Acid Disorders:**
Glutaric Acidemia (GA-1)
Isovaleric Acidemia (IVA)
Multiple carboxylase (MCD
3-Hydroxy 3-methylglutaric Aciduria (HMG)

04/2007 **Fatty Acid Disorders:**
Very long chain acyl-CoA dehydrogenase deficiency (VLCAD)
Long chain 3-hydroxyacyl-CoA dehydrogenase deficiency (LCHAD)
Trifunctional Protein Deficiency (TFP)

Organic Acid Disorders:
3-Methylcrotonyl-CoA carboxylase (3-MCC)
Beta ketothiolase (BKT)
Carnitine palymitoyltranferase II (CPT II)

01/2008 Universal Newborn Hearing Screening*

04/2008 Cystic Fibrosis (CF) (IRT/DNA)

2009 Cord Blood collection and testing discontinued

06/2013 Critical Congenital Heart Disease (CCHD)

10/2018 Severe Combined Immunodeficiency

***started voluntarily in 2001/ mandated 2008**