

New Hampshire Newborn Screening Program

List of Conditions

Each baby born in New Hampshire is screened for the conditions listed below. This list is correct as of July 1, 2007 but may change as conditions are added to or removed from the testing panel. If you have any questions, please contact the New Hampshire Newborn Screening Program at (603) 271-4225.

- 3-hydroxy-3-methylglutaryl-CoA lysase deficiency
- 3-methylcrotonyl-CoA carboxylase deficiency
- Argininemia
- Argininosuccinic aciduria
- Biotinidase deficiency
- Carnitine palmitoyltransferase II deficiency
- Carnitine uptake defect
- Citrullinemia I (ASA synthetase deficiency)
- Cobalamin A, B
- Congenital adrenal hyperplasia
- Congenital hypothyroidism
- Congenital toxoplasmosis
- Cystic fibrosis
- Galactosemia
- Glutaric aciduria type I
- Homocystinuria
- Hyperornithinemia, hyperammoninemia, homocitrullinemia syndrome
- Isovaleric acidemia
- Long chain 3-hydroxyacyl-CoA dehydrogenase (LCHAD) deficiency
- Maple syrup urine disease
- Medium-chain acyl-CoA dehydrogenase (MCAD) deficiency
- Methylmalonic acidemia
- Mitochondrial acetoacetyl-CoA thiolase deficiency
- Multiple acyl-CoA dehydrogenase deficiency
- Multiple carboxylase deficiency
- Phenylketonuria (PKU)
- Propionic acidemia
- Sickle cell disease/hemoglobin disorders
- Trifunctional protein deficiency
- Very long chain acyl-CoA dehydrogenase (VLCAD) deficiency