

新罕布什爾州新生兒篩檢機構 篩檢疾病列表

在新罕布什爾州出生的每個嬰兒將接受下列疾病篩檢。這份列表提供的資訊準確至2007年7月1日，但可能會因為其他疾病被加入或刪出本篩檢組而有所變動。如果您有任何問題，請聯繫新罕布什爾州新生兒篩檢機構，電話：(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

Traditional Chinese
New Hampshire