

新罕布什尔州新生儿筛检机构 筛检疾病列表

在新罕布什尔州出生的每个婴儿将接受下列疾病筛检。这份列表提供的信息准确至 2007 年 9 月 1 日，但可能会因为其它疾病被加入或删出本筛检组而有所变动。如果您有任何疑问，请联系新罕布什尔州新生儿筛检机构，电话：(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

Simplified Chinese
New Hampshire