

缅因州新生儿筛检机构 筛检疾病列表

在缅因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 7 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，请联系缅因州新生儿筛检机构，电话：(207) 287-5357。

3-Hydroxy-3-methylglutaryl-CoA lyase deficiency
3-Methylcrotonyl-CoA carboxylase deficiency
Argininemia
Argininosuccinic acidemia
Beta-ketothiolase deficiency
Biotinidase deficiency
Carnitine palmitoyl transferase deficiency Type II
Citrullinemia
Congenital adrenal hyperplasia
Congenital hypothyroidism
Galactosemia
Glutaric acidemia type I
Glutaric acidemia type II
Homocystinuria
Hyperammonemia Hyperornithinemia Homocitrullinemia
(HHH Syndrome)
Isovaleric acidemia
Long-chain acyl-CoA dehydrogenase (LCAD) deficiency
Long-chain hydroxyacyl-CoA dehydrogenase (LCHAD) deficiency
Maple syrup urine disease
Medium-chain acyl-CoA dehydrogenase (MCAD) deficiency
Methylmalonic acidemia
Phenylketonuria (PKU)
Propionic acidemia
Short-chain acyl-CoA dehydrogenase (SCAD) deficiency
Sickle cell disease/hemoglobin disorders
Tyrosinemia type I
Tyrosinemia type II
Very long-chain acyl-CoA dehydrogenase (VLCAD) deficiency