

佛蒙特州新生兒篩檢機構 篩檢疾病列表

在佛蒙特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2008年3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，請聯繫佛蒙特州新生兒篩檢機構，電話：(802) 951-5180。

- 3-Hydroxy-3-methylglutaryl-CoA lyase deficiency (**HMG**)
- 3-Methylcrotonyl-CoA carboxylase deficiency (**3-MCC**)
- Argininosuccinic acidemia (**ASA**)
- Beta-ketothiolase deficiency (**β -KT**)
- Biotinidase deficiency (**BIOT**)
 - Includes multiple carboxylase deficiency (**MCD**)
- Carnitine uptake defect (**CUD**)
- Citrullinemia (**CIT or ASS**)
- Congenital adrenal hyperplasia (**CAH**)
- Congenital hypothyroidism (**CH**)
- Cystic fibrosis (**CF**)
- Galactosemia (**GALT**)
- Glutaric acidemia type I (**GA I**)
- Hearing loss (**HEAR**)
- Homocystinuria (**HCY or HCU**)
- Isovaleric acidemia (**IVA**)
- Long-chain hydroxyacyl-CoA dehydrogenase (**LCHAD**) deficiency
 - Includes trifunctional protein deficiency (**TFP**)
- Maple syrup urine disease (**MSUD**)
- Medium-chain acyl-CoA dehydrogenase (**MCAD**) deficiency
- Methylmalonic acidemia Cbl A, B (**MMA, Cbl A, B**)
 - Includes methylmalonic acidemia mutase deficiency (**MUT**)
- Phenylketonuria (**PKU**)
- Propionic acidemia (**PA**)
- Sickle cell disease/hemoglobin disorders
 - Includes sickle cell anemia (**SCA**), Hb S/Beta thalassemia (**Hb S/Th**), and Hb S/C disease (**Hb S/C**)
- Tyrosinemia type I (**TYR I**)
- Very long-chain acyl-CoA dehydrogenase (**VLCAD**) deficiency