

Programa de Triagem de Recém-Nascidos de Vermont Lista das Condições Médicas

Cada bebê nascido em Vermont é testado para as condições médicas enumeradas abaixo. Esta lista estava vigente em 01 de março de 2008, mas poderá mudar caso condições médicas sejam adicionadas ou retiradas da triagem. Caso tiver perguntas, entre em contato com o Programa de Triagem de Recém-Nascidos de Vermont através do telefone (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

Portuguese