

Programma di Screening dei neonati dello Stato del Vermont Elenco delle malattie

Tutti i bambini nati nello Stato del Vermont vengono sottoposti a screening per verificare le malattie sotto elencate. Questo elenco è aggiornato in data 1 marzo 2008 ma può cambiare se delle malattie vengono aggiunte o rimosse dal pannello dei test. Se dovessero esserci domande, mettersi in contatto con il Programma di Screening dei neonati dello Stato del Vermont (Vermont Newborn Screening Program) al numero (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