

Barnaamijka Baadhitaanka Ilmaha Dhawaan Dhashay ee Vermont Liiska Xaaladaha

Ilmo kasta oo ku dhasha Vermont waxa laga baadhaa xaaladaha lagu taxay hoos. Liiskani waxa uu sax ahaa laga bilaabo 1-da Bisha Saddexaad 2008 laakiin waxa ku dhici kara isbeddel hadba marka lagu daro xaalado kale ama qaar laga saaro liiska baadhitaanka. Haddii aad qabtid wax su'aalo ah, fadlan la soo xidhiidh Barnaamijka Baadhitaanka Ilmaha Dhawaan Dhashay ee Vermont (Vermont Newborn Screening Program) oo leh lambarka (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