

## New Hampshire Program za Ispitivanje Novorođjenac Lista Uzroka Stanja

Svaka novorođjena beba u New Hampshireu je testirana za uzroke koje su navedene dole. Ova lista je korigovana 1 Maja, 2007 ali je moguće da se promeni u zavisnosti dali su uzroci dodati ili skinuti sa ove baterije testova. Ako imate pitanja molimo vas pozovite Program za ispitivanje novorođjenac u New Hampshireu na broj (603) 271-4225.

3-hydroxy-3-methylglutaryl-CoA lysase deficiency  
3-methylcrotonyl-CoA carboxylase deficiency  
Argininemia  
Argininosuccinic aciduria  
Biotinidase deficiency  
Carnitine palmitoyltransferase II deficiency  
Carnitine uptake defect  
Citrullinemia I (ASA synthetase deficiency)  
Cobalamin A, B  
Congenital adrenal hyperplasia  
Congenital hypothyroidism  
Congenital toxoplasmosis  
Cystic fibrosis  
Galactosemia  
Glutaric aciduria type I  
Homocystinuria  
Hyperornithinemia, hyperammoninemia, homocitrullinemia syndrome  
Isovaleric acidemia  
Long chain 3-hydroxyacyl-CoA dehydrogenase (LCHAD) deficiency  
Maple syrup urine disease  
Medium-chain acyl-CoA dehydrogenase (MCAD) deficiency  
Methylmalonic acidemia  
Mitochondrial acetoacetyl-CoA thiolase deficiency  
Multiple acyl-CoA dehydrogenase deficiency  
Multiple carboxylase deficiency  
Phenylketonuria (PKU)  
Propionic acidemia  
Sickle cell disease/hemoglobin disorders  
Trifunctional protein deficiency  
Very long chain acyl-CoA dehydrogenase (VLCAD) deficiency