

## Programma di Screening dei neonati dello stato del New Hampshire Elenco delle malattie

Tutti i bambini nati nello Stato del New Hampshire vengono sottoposti a screening per verificare le malattie sotto elencate. Questo elenco è aggiornato in data 1 luglio 2007 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 New Hampshire (New Hampshire Newborn Screening Program) al numero (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