



A Card for Your Baby Book

This card is to be used in your baby book! Check the boxes next to each disorder so that you know which disorders your baby was screened for through the newborn screening process. The back side has important follow-up information that is easily available if you need it!

Newborn Screening


 Cut here!




Newborn Screening Report Card

For:

Disorders Screened For:

<ul style="list-style-type: none"> <input type="checkbox"/> Phenylketonuria (PKU) <input type="checkbox"/> Maple syrup urine disease (MSUD) <input type="checkbox"/> Homocystinuria (HCV) <input type="checkbox"/> Citrullinemia (CIT) <input type="checkbox"/> Argininosuccinic acidemia (ASA) <input type="checkbox"/> Tyrosinemia type I (TYR I) <input type="checkbox"/> Isovaleric acidemia (IVA) <input type="checkbox"/> Glutaric acidemia type I (GAI) <input type="checkbox"/> Hydroxymethylglutaric aciduria (HMG) <input type="checkbox"/> Multiple carboxylase deficiency (MCD) <input type="checkbox"/> Methylmalonic acidemia due to mutase deficiency (MUT) <input type="checkbox"/> Methylmalonic acidemia cblA and cblB forms (Cbl A, B) <input type="checkbox"/> 3-Methylcrotonyl-CoA carboxylase deficiency (3MCC) <input type="checkbox"/> Propionic acidemia (PROP) <input type="checkbox"/> Beta-Ketothiolase deficiency (BKT) <input type="checkbox"/> Medium-chain acyl-CoA dehydrogenase deficiency (MCAD) 	<ul style="list-style-type: none"> <input type="checkbox"/> Very long-chain acyl-CoA dehydrogenase deficiency (VLCAD) <input type="checkbox"/> Long-chain 3-OH acyl-CoA dehydrogenase deficiency (LCHAD) <input type="checkbox"/> Trifunctional protein deficiency (TFP) <input type="checkbox"/> Carnitine uptake defect (CUD) <input type="checkbox"/> Sickle cell anemia (Hb SS) <input type="checkbox"/> Hb S/Beta-Thalassemia (Hb S/Th) <input type="checkbox"/> Hb S/C disease (Hb S/C) <input type="checkbox"/> Congenital hypothyroidism (CH) <input type="checkbox"/> Biotinidase deficiency (BIOT) <input type="checkbox"/> Congenital adrenal hyperplasia (CAH) <input type="checkbox"/> Classical galactosemia (GALT) <input type="checkbox"/> Hearing loss (HEAR) <input type="checkbox"/> Cystic fibrosis (CF)
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 Fold here!

Reminder:

If your baby's initial test results require follow-up, take your doctor's advice and bring your baby for re-testing quickly. The results may indicate a disorder so it is important to follow through with prompt care.

For more information

- Talk to your pediatrician
- Call your State's Department of Health
- Visit the National Newborn Screening and Genetics Resource Center Website
www.genes-r-us.uthsca.edu
- Visit the New England Consortium of Metabolic's Program's Website
www.newenglandconsortium.org