

**ແຜນການກວດພະຍາດເດັກນ້ອຍເກີດໃໝ່ປະຈຳຮຸ້ງເວີມອນ**  
**ລາຍການສະພາວະ**

ເດັກທຸກຄົນທີ່ເກີດໃນຮຸ້ງເວີມອນ ຕ້ອງໄດ້ຮັບການກວດຫາສະພາວະທີ່ມີລາຍການຢູ່ຂ້າງລຸ່ມນີ້. ລາຍການນີ້ແມ່ນຖືກຕ້ອງນັບຈາກວັນທີ 1 ເດືອນມີນາ ປີ 2008 ແຕ່ອາດມີການປ່ຽນແປງ ເນື່ອງຈາກ ມີການເພີ່ມ ຫຼື ເອົາລາຍການກວດອອກ ຈາກບັນຊີການກວດ. ຖ້າຫາກທ່ານມີຄໍາຖາມ, ກະລຸນາຕິດຕໍ່ຫາ ແຜນການກວດພະຍາດເດັກນ້ອຍເກີດໃໝ່ ປະຈຳຮຸ້ງເວີມອນ ທີ່ເບີໂທ (802) 951-5180.

- 3-Hydroxy-3-methylglutaryl-CoA lyase deficiency (HMG)
- 3-Methylcrotonyl-CoA carboxylase deficiency (3-MCC)
- Argininosuccinic acidemia (ASA)
- Beta-ketothiolase deficiency ( $\beta$ -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