

ແຜນການກວດພະຍາດເດັກນ້ອຍເກີດໃໝ່ປະຈຳຮຸ້ງຊາວ ລາຍການສະພາວະ

ເດັກທຸກຄົນທີ່ເກີດໃນຮຸ້ງຊາວ ຕ້ອງໄດ້ຮັບການກວດຫາສະພາວະທີ່ມີລາຍການຢູ່ຂ້າງລຸ່ມນີ້. ລາຍການນີ້ແມ່ນຖືກຕ້ອງນັບຈາກວັນທີ 1 ເດືອນພຶດສະພາ ປີ 2007 ແຕ່ອາດມີການປ່ຽນແປງ ເນື່ອງຈາກ ມີການເພີ່ມ ຫຼື ເອົາລາຍການກວດອອກ ຈາກບັນຊີການກວດ. ຖ້າຫາກທ່ານມີຄໍາຖາມ, ກະລຸນາຕິດຕໍ່ຫາ ແຜນການກວດພະຍາດເດັກນ້ອຍເກີດໃໝ່ປະຈຳຮຸ້ງຊາວ ທີ່ເບີໂທ (207) 287-5357.

- 3-Hydroxy-3-methylglutaryl-CoA lyase deficiency
- 3-Methylcrotonyl-CoA carboxylase deficiency
- Argininemia
- Argininosuccinic acidemia
- Beta-ketothiolase deficiency
- Biotinidase deficiency
- Carnitine palmitoyl transferase deficiency Type II
- Citrullinemia
- Congenital adrenal hyperplasia
- Congenital hypothyroidism
- Galactosemia
- Glutaric acidemia type I
- Glutaric acidemia type II
- Homocystinuria
- Hyperammonemia Hyperornithinemia Homocitrullinemia
(HHH Syndrome)
- Isovaleric acidemia
- Long-chain acyl-CoA dehydrogenase (LCAD) deficiency
- Long-chain hydroxyacyl-CoA dehydrogenase (LCHAD) deficiency
- Maple syrup urine disease
- Medium-chain acyl-CoA dehydrogenase (MCAD) deficiency
- Methylmalonic acidemia
- Phenylketonuria (PKU)
- Propionic acidemia
- Short-chain acyl-CoA dehydrogenase (SCAD) deficiency
- Sickle cell disease/hemoglobin disorders
- Tyrosinemia type I
- Tyrosinemia type II
- Very long-chain acyl-CoA dehydrogenase (VLCAD) deficiency

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