

Pwogram Depistaj pou Timoun ki Fenk Fèt nan Maine Lis Kondisyon Anòmalye

Yo fè tès depistaj pou tout timoun ki fenk fèt nan Maine pou dekouvri si gen kondisyon anòmalye sa yo. Lis sa a valab jiska dat premye me 2007, men li ka chanje si gen kondisyon anòmalye yo mete ankò oswa yo retire nan pwogram pou tès yo. Si w gen nenpòt kesyon, kontakte Pwogram Depistaj pou Timoun ki Fenk Fèt nan Maine lan nan nimewo (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