

My baby

ACOG-Recommended Conditions

Beta-thalassemia
Canavan disease
Cystic fibrosis
Familial dysautonomia
Sickle cell disease
Tay-Sachs disease

Additional Carrier Status Conditions

21-Hydroxylase-deficient congenital adrenal hyperplasia
3-Methylcrotonyl-CoA carboxylase deficiency
Achromatopsia
Acrodermatitis enteropathica
Alkaptonuria
Alpha-1 antitrypsin deficiency
Alpha-mannosidosis
Amyotrophic lateral sclerosis
Andermann syndrome
Argininosuccinate lyase deficiency
ARSACS
Aspartylglucosaminuria
Ataxia with vitamin E deficiency
Ataxia-telangiectasia
Autoimmune polyglandular syndrome, type I
Bardet-Biedl syndrome, BBS1-related
Bartter syndrome type 4A
Beta-ketothiolase deficiency
Biotinidase deficiency
Carnitine deficiency, primary systemic
Carnitine palmitoyltransferase I deficiency
Cartilage-hair hypoplasia
Cerebrotendinous xanthomatosis
Choroideremia
Citrullinemia, type I
Cohen syndrome
Combined pituitary hormone deficiency, PROP1-related
Congenital disorder of glycosylation type Ia
Costeff optic atrophy syndrome
Crigler-Najjar syndrome
Cystinosis
Diabetes, permanent neonatal
Dubin-Johnson syndrome
Ehlers-Danlos syndrome, dermatosparaxis

Ashkenazi Jewish Conditions

Bloom syndrome
Canavan disease
Cystic fibrosis
Dihydropyrimidine dehydrogenase deficiency
Factor XI deficiency
Familial dysautonomia
Fanconi anemia

Ehlers-Danlos syndrome, hypermobility
Ehlers-Danlos syndrome, kyphoscoliotic
Factor V Leiden thrombophilia
Familial Mediterranean fever
Galactokinase deficiency
Galactosemia
Glutaric acidemia, type 1
Glycogen storage disease, type Ib
Glycogen storage disease, type III
Glycogen storage disease, type V
GM1-gangliosidosis
Hearing loss, DFNB1 and DFNB9 nonsyndromic
Hearing loss, DFNB59 nonsyndromic
Hemochromatosis
Hemoglobin C
Hemoglobin D
Hemoglobin E
Hemoglobin O
Hereditary Fructose Intolerance
Herlitz junctional epidermolysis bullosa, LAMA3-related
Herlitz junctional epidermolysis bullosa, LAMB3-related
Herlitz junctional epidermolysis bullosa, LAMC2-related
HMG-CoA lyase deficiency
Homocystinuria, cb1E type
Homocystinuria, classic
Hurler syndrome
Hypophosphatasia, autosomal recessive
Inclusion Body Myopathy 2
Juvenile retinoschisis, X-linked
Krabbe disease
Lipoamide dehydrogenase deficiency
Lipoprotein lipase deficiency, familial
Medium-chain acyl-CoA dehydrogenase deficiency
Megalencephalic leukoencephalopathy with subcortical cysts
Metachromatic leukodystrophy

Gaucher disease
Glycogen storage disease, type 1A
Maple syrup urine disease
Mucopolidosis IV
Niemann-Pick disease
Tay-Sachs disease
Tyrosinemia
Usher syndrome, type 1F

Methylmalonic acidemia
Mucopolidosis II
Mucopolidosis III
Multiple carboxylase deficiency
Nephrotic syndrome, steroid-resistant
Neuronal ceroid lipofuscinosis, CLN3-related
Neuronal ceroid lipofuscinosis, CLN5-related
Neuronal ceroid lipofuscinosis, CLN8-related
Neuronal ceroid lipofuscinosis, PPT1-related
Neuronal ceroid lipofuscinosis, TPP1-related
Nijmegen breakage syndrome
Pendred syndrome
Phenylketonuria
Polycystic kidney disease
Pompe disease
Prekallikrein deficiency
Primary hyperoxaluria, type 1
Primary hyperoxaluria, type 2
Primary hyperoxaluria, type 3
Propionic acidemia
Prothrombin deficiency
Rhizomelic chondrodysplasia punctata type 1
Rh-null syndrome
Rickets, pseudovitamin D-deficiency
Salla disease
Sandhoff disease
Short-chain acyl-CoA dehydrogenase deficiency
Sick sinus syndrome
Smith-Lemli-Opitz Syndrome
Spherocytosis, hereditary
Tay-Sachs pseudodeficiency
Thrombocytopenia, congenital amegakaryocytic
Tyrosine Hydroxylase Deficiency
Very long-chain acyl-CoA dehydrogenase deficiency
Von Willebrand disease type 2 Normandy
Von Willebrand disease type 3
Wilson disease
Zellweger syndrome spectrum, PEX1-related