Carrier Status

Learn whether you have specific genetic variants that may not affect your health, but could affect your children's health.

Carrier Status Tutorial

Keep in mind that while our Carrier Status reports cover many variants, they don't include all possible variants associated with each condition. So it's still possible to be a carrier of a variant not included in our test. If you're considering having children, we encourage you to talk to a healthcare professional about your options for carrier screening.

Cystic Fibrosis
Variant detected

ARSACS
Variant not detected

Agenesis of the Corpus Callosum with Peripheral Neuropathy
Variant not detected

Autosomal Recessive Polycystic Kidney Disease
Variant not detected

Beta Thalassemia and Related Hemoglobinopathies
Variant not detected

Bloom Syndrome
Variant not detected

Canavan Disease
Variant not detected

Congenital Disorder of Glycosylation Type 1a (PMM2-CDG)
Variant not detected

D-Bifunctional Protein Deficiency
Variant not detected

Dihydrolipoamide Dehydrogenase Deficiency
Variant not detected

Familial Dysautonomia
Variant not detected

Familial Hyperinsulinism (ABCC8-Related)
Variant not detected

Familial Mediterranean Fever
Variant not detected

Fanconi Anemia Group C
Variant not detected

GRACILE Syndrome
Variant not detected

Gaucher Disease Type 1
Variant not detected

Glycogen Storage Disease Type Ia
Variant not detected

Glycogen Storage Disease Type Ib
Variant not detected

Hereditary Fructose Intolerance
Variant not detected

Leigh Syndrome, French Canadian Type
Variant not detected

Limb-Girdle Muscular Dystrophy Type 2D
Variant not detected

Limb-Girdle Muscular Dystrophy Type 2E
Variant not detected

Limb-Girdle Muscular Dystrophy Type 2I
Variant not detected

MCAD Deficiency
Variant not detected

Maple Syrup Urine Disease Type 1B
Variant not detected

Mucolipidosis Type IV
Variant not detected

Neuronal Ceroid Lipofuscinosis (CLN5-Related)
Variant not detected

Neuronal Ceroid Lipofuscinosis (PPT1-Related)
Variant not detected

Niemann-Pick Disease Type A
Variant not detected

Nijmegen Breakage Syndrome
Variant not detected

Nonsyndromic Hearing Loss and Deafness, DFNB1 (GJB2-Related)
Variant not detected

Pendred Syndrome and DFNB4 Hearing Loss (SLC26A4-Related)
Variant not detected

Phenylketonuria and Related Disorders
Variant not detected

Pompe Disease
Variant not detected

Primary Hyperoxaluria Type 2
Variant not detected

Pyruvate Kinase Deficiency
Variant not detected

Rhizomelic Chondrodysplasia Punctata Type 1
Variant not detected

Salla Disease
Variant not detected

Severe Junctional Epidermolysis Bullosa (LAMB3-Related)
Variant not detected

Sickle Cell Anemia
Variant not detected

Sjögren-Larsson Syndrome
Variant not detected

Tay-Sachs Disease
Variant not detected

Tyrosinemia Type I
Variant not detected

Usher Syndrome Type 1F
Variant not detected

Usher Syndrome Type 3A
Variant not detected

Zellweger Spectrum Disorder (PEX1-Related)
Variant not detected