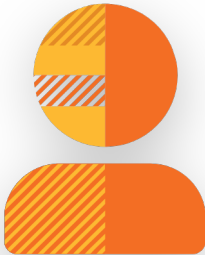


Carrier Status

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



<u>Canavan Disease</u>	>
<u>Variant not detected</u>	
<u>Congenital Disorder of Glycosylation Type 1a (PMM2-CDG)</u>	>
<u>Variant not detected</u>	
<u>D-Bifunctional Protein Deficiency</u>	>
<u>Variant not detected</u>	
<u>Dihydrolipoamide Dehydrogenase Deficiency</u>	>
<u>Variant not detected</u>	
<u>Familial Dysautonomia</u>	>
<u>Variant not detected</u>	
<u>Familial Hyperinsulinism (ABCC8-Related)</u>	>
<u>Variant not detected</u>	
<u>Familial Mediterranean Fever</u>	>
<u>Variant not detected</u>	
<u>Fanconi Anemia Group C</u>	>
<u>Variant not detected</u>	
<u>GRACILE Syndrome</u>	>
<u>Variant not detected</u>	
<u>Gaucher Disease Type 1</u>	>
<u>Variant not detected</u>	
<u>Glycogen Storage Disease Type Ia</u>	>
<u>Variant not detected</u>	

<u>Glycogen Storage Disease Type Ib</u>	>
<u>Variant not detected</u>	
<u>Hereditary Fructose Intolerance</u>	>
<u>Variant not detected</u>	
<u>Leigh Syndrome, French Canadian Type</u>	>
<u>Variant not detected</u>	
<u>Limb-Girdle Muscular Dystrophy Type 2D</u>	>
<u>Variant not detected</u>	
<u>Limb-Girdle Muscular Dystrophy Type 2E</u>	>
<u>Variant not detected</u>	
<u>Limb-Girdle Muscular Dystrophy Type 2I</u>	>
<u>Variant not detected</u>	
<u>MCAD Deficiency</u>	>
<u>Variant not detected</u>	
<u>Maple Syrup Urine Disease Type 1B</u>	>
<u>Variant not detected</u>	
<u>Mucopolidosis Type IV</u>	>
<u>Variant not detected</u>	
<u>Neuronal Ceroid Lipofuscinosis (CLN5-Related)</u>	>
<u>Variant not detected</u>	
<u>Neuronal Ceroid Lipofuscinosis (PPT1-Related)</u>	>
<u>Variant not detected</u>	

<u>Niemann-Pick Disease Type A</u>	>
<u>Variant not detected</u>	
<u>Nijmegen Breakage Syndrome</u>	>
<u>Variant not detected</u>	
<u>Nonsyndromic Hearing Loss and Deafness, DFNB1 (GJB2-Related)</u>	>
<u>Variant not detected</u>	
<u>Pendred Syndrome and DFNB4 Hearing Loss (SLC26A4-Related)</u>	>
<u>Variant not detected</u>	
<u>Phenylketonuria and Related Disorders</u>	>
<u>Variant not detected</u>	
<u>Pompe Disease</u>	>
<u>Variant not detected</u>	
<u>Primary Hyperoxaluria Type 2</u>	>
<u>Variant not detected</u>	
<u>Pyruvate Kinase Deficiency</u>	>
<u>Variant not detected</u>	
<u>Rhizomelic Chondrodysplasia Punctata Type 1</u>	>
<u>Variant not detected</u>	
<u>Salla Disease</u>	>
<u>Variant not detected</u>	
<u>Severe Junctional Epidermolysis Bullosa (LAMB3-Related)</u>	>
<u>Variant not detected</u>	

<u>Sickle Cell Anemia</u>	>
<u>Variant not detected</u>	
<hr/>	
<u>Sjögren-Larsson Syndrome</u>	>
<u>Variant not detected</u>	
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<u>Tay-Sachs Disease</u>	>
<u>Variant not detected</u>	
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<u>Tyrosinemia Type I</u>	>
<u>Variant not detected</u>	
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<u>Usher Syndrome Type 1F</u>	>
<u>Variant not detected</u>	
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<u>Usher Syndrome Type 3A</u>	>
<u>Variant not detected</u>	
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<u>Zellweger Spectrum Disorder (PEX1-Related)</u>	>
<u>Variant not detected</u>	