All Tested Health Conditions

- **Exercise Induced Metabolic Myopathy**
- **Respiratory Distress Syndrome**
- **Adverse Reaction to Certain Drugs, Variant 1**
- **Primary Open Angle Glaucoma, Variant 2**
- **Dental Hypomineralization**
- **Osteogenesis Imperfecta, Type III**
- **Gallbladder Mucoceles**
- **Bleeding Disorder due to P2Ry12 Defect**
- **Narcolepsy, Variant 1**
- **Progressive Retinal Atrophy, Variant 1**
- **Exercise-Induced Collapse**
- **Muscular Dystrophy, Variant 3**
- **Dilated Cardiomyopathy, Variant 2**
- **Benign Familial Juvenile Epilepsy**
- **Inherited Myopathy**
- **Myotubular Myopathy 1, Variant 1**
- **Unilateral Deafness and Vestibular Dysfunction**
- **Primary Open Angle Glaucoma, Variant 1**
- **Bardet-Biedl Syndrome 4**
- **Neonatal Encephalopathy with Seizures**
- **Beta Mannosidosis, Variant 1**
- **Osteogenesis Imperfecta, Variant 3**
- **Neuronal Ceroid Lipofuscinosis 6**
- **Renal Cystadenocarcinoma and Nodular Dermatofibrosis**
- **Malignant Hyperthermia**
- **Ichthyosis, Variant 1**
- **Early Cerebellar Ataxia**
- **Juvenile Cerebellar Ataxia**
- **Cone-Rod Dysplasia 3**

- **Nasal Parakeratosis, Variant 1**
- **Hypophosphatasia**
- **Hypomyelination of the Central Nervous System**
- **Progressive Retinal Atrophy GR2**
- **Degenerative Myelopathy, Variant 1**
- **Von Willebrand Disease II, Variant 1**
- **Hemophilia B, Variant 1**
- **Pituitary Dwarfism**
- **Ichthyosis, Variant 2**
- **Cystinuria Type 2A**
- **Polyneuropathy 2**
- **Generalized Myoclonic Epilepsy, with Photosensitivity**
- **Limb-Girdle Muscular Dystrophy Type 2F, Variant 1**
- **Duchenne Muscular Dystrophy, Variant 1**
- **Progressive Retinal Atrophy, Variant 2**
- **Limb-Girdle Muscular Dystrophy Type 2F, Variant 2**
- **Spinocerebellar Ataxia, Variant 1**
- **C3 Deficiency**
- **GM1 Gangliosidosis, Variant 1**
- **GM2 Gangliosidosis Type 1(Tay Sachs)**
- **Early Imerslund-Grasbeck Syndrome**
- **Late Imerslund-Grasbeck Syndrome**
- **Musladin-Lueke Syndrome**
- **Pyruvate Dehydrogenase Phosphatase Deficiency**
- **Familial Enamel Hypoplasia Amelogenesis Imperfecta, Variant 1**
- **Acral Mutilation Syndrome**
- **Hypothyroidism, Variant 1**
- **Skeletal Dysplasia 2(Disproportionate Dwarfism)**
- **Myotubular Myopathy 1, Variant 2**
- **Muscular Dystrophy, Variant 1**
- **Bilateral Deafness and Vestibular Dysfunction**

- **Primary Open Angle Glaucoma, Variant 3**
- **Encephalopathy**
- **Congenital Eye Malformation**
- **Polyneuropathy, Variant 1**
- **Multifocal Retinopathy 2(Cmr2)**
- **Multifocal Retinopathy 3(Cmr3), Variant 1**
- **Dandy Walker Syndrome**
- **Craniomandibular Osteopathy, Variant 1**
- **Menkes Disease**
- **Hypocatalasia**
- **Exfoliative Cutaneous Lupus Erythematosus**
- **Leukodystrophy**
- **Hemeralopia Achromatopsia(Cone Degeneration)**
- **Cystinuria**
- **Neuronal Ceroid Lipofuscinosis 5, Variant 1**
- **Neuronal Ceroid Lipofuscinosis 10**
- **Spinocerebellar Degeneration and Neuronal Vacuolation**
- **Neuronal Ceroid Lipofuscinosis A, Variant 1**
- **Oculoskeletal Dysplasia 1, Variant 1**
- **Ichthyosis, Variant 4**
- **Late Spinocerebellar Ataxia**
- **Spinocerebellar Ataxia with Myokymia and Seizure**
- **Spinocerebellar Ataxia, Variant 2**
- **Cone-Rod Dysplasia 1**
- **Progressive Retinal Atrophy With Progressive Rod-Cone Degeneration**
- **Von Willebrand Disease III, Variant 1**
- **Dystrophic Epidermolysis Bullosa, Variant 1**
- **Simplex Epidermolysis Bullosa**
- **Hemophilia A, Variant 1**
- **Hemophilia A, Variant 2**
- **Nasal Parakeratosis, Variant 2**

- **Neuronal Ceroid Lipofuscinosis 5, Variant 2**
- **Spondylocostal Dysostosis**
- **Limb-Girdle Muscular Dystrophy Type 2F, Variant 3**
- **Osteogenesis Imperfecta, Variant 1**
- **GM1 Gangliosidosis, Variant 2**
- **GM2 Gangliosidosis Type 0(Sandhoff Disease)**
- **Congenital Hypothyroidism with Goiter**
- **Congenital Dyshormonogenic Hypothyroidism with Goiter**
- **Hyperuricosuria**
- **Epidermolytic Hyperkeratosis**
- **Palmoplantar Hyperkeratosis, Variant 1**
- **Nemaline Myopathy**
- **Muscular Dystrophy, Variant 2**
- **Primary Open Angle Glaucoma, Variant 4**
- **Multifocal Retinopathy 1**
- **Multifocal Retinopathy 3(Cmr3), Variant 2**
- **Craniomandibular Osteopathy, Variant 2**
- **Leukoencephalomyelopathy**
- **Osteogenesis Imperfecta, Variant 2**
- **Achromatopsia**
- **Cystinuria Type 2B**
- **Neuronal Ceroid Lipofuscinosis 4A**
- **Neuronal Ceroid Lipofuscinosis 2**
- **Neuronal Ceroid Lipofuscinosis A, Variant 2**
- **Neuronal Ceroid Lipofuscinosis 8, Variant 1**
- **Chondrodysplasia**
- **Progressive Retinal Atrophy Type 1, Variant 1**
- **Progressive Retinal Atrophy, Variant 3**
- **Degenerative Myelopathy, Variant 2**
- **Von Willebrand Disease I**
- **Dystrophic Epidermolysis Bullosa, Variant 2**

- **Hemophilia A, Variant 3**
- **Hemophilia B, Variant 2**
- **Hemophilia B, Variant 3**
- **Progressive Retinal Atrophy Crd4/Cord1**
- **Oculoskeletal Dysplasia 1, Variant 2**
- **Polyneuropathy, Variant 2**
- **Cone-Rod Dystrophy 1**
- **Hypothyroidism, Variant 2**
- **Leukodystrophy(Krabbe Disease)**
- **Neuronal Ceroid Lipofuscinosis 1, Variant 2**
- **Neuronal Ceroid Lipofuscinosis 8, Variant 2**
- **Neuronal Ceroid Lipofuscinosis 8, Variant 3**
- **Cerebellar Ataxia**
- **Progressive Retinal Atrophy, Variant 4**
- **Von Willebrand Disease II, Variant2**
- **Progressive Retinal Atrophy Type 1, Variant 2**
- **Palmoplantar Hyperkeratosis, Variant 2**
- **Duchenne Muscular Dystrophy, Variant 2**
- **Cerebellar Abiotrophy**
- **Neonatal Cerebellar Cortical Degeneration**
- **GM1 Gangliosidosis, Variant 3**
- **Cone-Rod Dystrophy 2**
- **Cone-Rod Dysplasia 1A**
- **Mucopolysaccharidosis VI, Variant 2**
- **Severe Combined Immunodeficiency Disease, Variant 1**
- **Severe Combined Immunodeficiency Disease, Variant 2**
- **Pulmonary Surfactant Metabolism Dysfunction**
- **Phosphofructokinase Deficiency**
- **Pyruvate Kinase Deficiency, Variant 1**
- **Thrombopathia, Variant 1**
- **Macular Corneal Dystrophy**

- **Mucopolysaccharidosis IIIa**
- **Van Den Ende-Gupta Syndrome**
- **Primary Lens Luxation, Variant 1**
- **Recurrent Inflammatory Pulmonary Disease**
- **Nephritis(Alport Syndrome)**
- **Familial Nephropathy, Variant 1**
- **Congenital Macrothrombocytopenia**
- **Dry Eye Curly Coat Syndrome**
- **Myotonia Congenita, Variant 1**
- **Narcolepsy, Variant 2**
- **Diffuse Cystic Renal Dysplasia and Hepatic Fibrosis**
- **Polycystic Kidney Disease**
- **Methemoglobinemia**
- **Fecundity**
- **Vitamin D-Deficiency Rickets Type II**
- **May-Hegglin Anomaly**
- **Congenital Myasthenic Syndrome, Variant 1**
- **Congenital Myasthenic Syndrome, Variant 2**
- **Congenital Myasthenic Syndrome, Variant 3**
- **Intestinal Cobalamin Malabsorption**
- **Primary Lens Luxation, Variant 2**
- **Primary Ciliary Dyskinesia, Variant 2**
- **Primary Hyperoxaluria Type I(Oxalosis I)**
- **Thrombocytopenia**
- **Mucopolysaccharidosis VII, Variant 1**
- **Mucopolysaccharidosis VII, Variant 2**
- **Mucopolysaccharidosis I**
- **Factor VII Deficiency**
- **Spinal Dysraphism**
- **Pyruvate Kinase Deficiency, Variant 2**
- **Thrombasthenia, Variant 1**

- **L-2-Hydroxyglutaric Aciduria, Variant 1**
- **Cyclic Hematopoiesis**
- **Mucopolysaccharidosis VI, Variant 1**
- **Primary Ciliary Dyskinesia, Variant 1**
- **Familial Adenomatous Polyposis**
- **Familial Nephropathy, Variant 2**
- **Myotonia Congenita, Variant 2**
- **L-2-Hydroxyglutaric Aciduria, Variant 2**
- **Congenital Myasthenic Syndrome, Variant 4**
- **Pyruvate Kinase Deficiency, Variant 3**
- **Thrombopathia, Variant 2**
- **Familial Enamel Hypoplasia Amelogenesis Imperfecta, Variant 2**
- **Myotonia Congenita, Variant 3**
- **Severe Combined Immunodeficiency Disease, Variant 3**
- **Thrombopathia, Variant 3**
- **Thrombasthenia, Variant 2**
- **Multi-Drug Resistance**
- **Glycogen Storage Disease Type Ia**
- **Glycogen Storage Disease Type II(Pompe Disease)**
- **Periodic Fever Syndrome**
- **Persistent Mullerian Duct Syndrome**
- **Adverse Reaction to Certain Drugs, Variant 2**
- **Trapped Neutrophil Syndrome**
- **Stargardt Disease 1**
- **Lethal Acrodermatitis**
- **Laryngeal Paralysis**
- **Hereditary Cataracts, Variant 2**
- **Beta Mannosidosis, Variant 2**
- **Lysosomal Storage Disease**
- **Leukocyte Adhesion Deficiency Type I**
- **Hypotrichosis, Variant 1**

- **Scott Syndrome**
- **Juvenile Neuroaxonal Dystrophy**
- **Dilated Cardiomyopathy, Variant 1**
- **Ehlers-Danlos Syndrome Classic-Like Type 1, Variant 1**
- **Ehlers-Danlos Syndrome Classic-Like Type 1, Variant 2**
- **Ehlers-Danlos Syndrome Type VII**
- **Ectodermal Dysplasia**
- **Canine Paroxysmal Dyskinesia**
- **X-Linked Hypohidrotic Ectodermal Dysplasia**
- **Hypotrichosis, Variant 2**
- **Child-Like Syndrome**
- **Neuroaxonal Dystrophy, Variant 1**
- **Alexander Disease**
- **Ehlers-Danlos Syndrome Classic Type 1, Variant 1**
- **Ehlers-Danlos Syndrome Classic Type 1, Variant 2**
- **Neuroaxonal Dystrophy, Variant 2**
- **Dilated Cardiomyopathy, Variant 3**
- **Leukocyte Adhesion Deficiency Type III**
- **Cleft Palate With Syndactyly**
- **Shaking Puppy Syndrome**
- **Prekallikrein Deficiency**
- **Glycogen Storage Disease Type IIIa**