

# Genetic Disorder Sign Up Sheet

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Disorder	Students/Groups
Achondroplasia	
Adrenoleukodystrophy	
Albinism	
Alpha-1 Antitrypsin Deficiency	
Alzheimer's	
Angelman Syndrome	
Antiphospholipid Syndrome	
Autism	
Autoimmune Lymphoproliferative Syndrome (Alps)	
Autosomal Dominant Polycystic Kidney Disease	
Breast Cancer	
Burkitt Lymphoma	
Charcot-Marie-Tooth	
Colon Cancer	
Colorblindness	
Cooley's Anemia	
Cri Du Chat	
Cri-Du-Chat Syndrome	
Crohn's Disease	
Cystic Fibrosis	
Dercum Disease	
Diabetes Insipidus	
Down Syndrome	
Duane Syndrome	
Duchenne Muscular Dystrophy	
Edward's Syndrome/Patau Syndrome	
Epidermolysis	
Factor V Leiden Thrombophilia	

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Familial Hypercholesterolemia

Familial Mediterranean Fever

Fragile X Syndrome

Galactosemia

Gaucher Disease

Hemochromatosis

Hemophilia

Holoprosencephaly

Huntington's Disease

Klinefelter's syndrome

Left Lip/Palate

Marfan Syndrome

Myotonic Dystrophy

Neurofibromatosis

Noonan Syndrome

Osteogenesis Imperfecta

Parkinson's Disease

Phenylketonuria

Pituitary Dwarfism Syndrome

Poland Anomaly

Polycystic Kidney Disease

Polydactyl

Porphyria

Progeria

Prostate Cancer

Retinitis Pigmentosa

Sensorineural Deafness

Severe Combined Immunodeficiency (SCID)

Sickle Cell Disease

Skin Cancer

Spina Bifida/Anencephaly

Spinal Muscular Atrophy

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Tay-Sachs

Thalassemia

Trimethylaminuria

Turner's Syndrome

Velocardiofacial Syndrome

WAGR Syndrome

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