

Genetic Disorder Sign Up Sheet

Disorder	Students/Groups
Achondroplasia	
Alpha-1 Antitrypsin Deficiency	
Antiphospholipid Syndrome	
Autism	
Autoimmune Lymphoproliferative Syndrome (ALPS)	
Autosomal Dominant Polycystic Kidney Disease	
Breast Cancer	
Charcot-Marie-Tooth	
Colon Cancer	
Cri Du Chat	
Crohn's Disease	
Cystic Fibrosis	
Dercum Disease	
Down Syndrome	
Duane Syndrome	
Duchenne Muscular Dystrophy	
Factor V Leiden Thrombophilia	
Familial Hypercholesterolemia	
Familial Mediterranean Fever	
Fragile X Syndrome	
Gaucher Disease	
Hemochromatosis	
Hemophilia	
Holoprosencephaly	
Huntington's Disease	
Klinefelter Syndrome	
Marfan Syndrome	
Myotonic Dystrophy	
Neurofibromatosis	

Noonan Syndrome

Osteogenesis Imperfecta

Parkinson's Disease

Phenylketonuria

Poland Anomaly

Porphyria

Progeria

Prostate Cancer

Retinitis Pigmentosa

Severe Combined Immunodeficiency (SCID)

Sickle Cell Disease

Skin Cancer

Spinal Muscular Atrophy

Tay-Sachs

Thalassemia

Trimethylaminuria

Turner Syndrome

Velocardiofacial Syndrome

Alzheimer's

Albinism

Burkitt Lymphoma

Diabetes Insipidus

Fragile X Syndrome

Epidermolysis

Duchenne Muscular Dystrophy

Thalassemia

Edward's Syndrome/Patau Syndrome

Turner's Syndrome

Polycystic Kidney Disease

Klinefelter's Syndrome

Cooley's Anemia

Gaucher's Disease

Hemophilia

Neurofibromatosis

Marfan Syndrome

Cri-Du-Chat Syndrome

Parkinson's Disease

Cystic Fibrosis

Colorblindness

Down Syndrome

Pituitary Dwarfism Syndrome

Angelman Syndrome

Adrenoleukodystrophy

Galactosemia

Sensorineural Deafness

Spina Bifida/Anencephaly

Osteogenesis Imperfecta

Polydactyl

Left Lip/Palate

WAGR Syndrome

<http://www.genome.gov/10001204>