

1 - Gaucher - L-myc gene (involved in small cell carcinoma) - Presenilin 2 (early alz.)	2	3 - Von Hippel Lindau (VHL) (3p26)	4 - Huntington (CAG) - Achondroplasia (FGFR3 gene)	5 - APC (FAP) (5q-) - Cri-Du-Chat (5p-) APC = "Colon" = 5 letters = chromosome 5!	6 Hemochromatosis HFE gene (C282Y, H63D) (example of Linkage Disequilibrium)
7 - Cystic Fibrosis (CFTR) - Williams syn. (7q-, elastin, "friendly elves", ↑ Ca ²⁺)	8 - C-myc gene (Burkitt's)	9 - Abl (bcr-abl) (9q34.1) - TSC1 (Hamartin) (9q34) - P16 tumor suppressor gene (Melanoma) - JAK2 (PV, ET, MMM) - Friedreich's Ataxia (GAA)	10	11 - ATM (11q22.3) - BCL-1 (cyclin D) - Wilms' tumor (deletion of tumor suppressor gene WT1) - Niemann Pick (sphingomyelinase) - β Globin Chain (β-Thal)	12
13 - Rb Gene (retino, osteo) - BRCA-2 - Patau Trisomy - Wilson's disease (ATP7b) Angelina Jolie = 13 letters = chromosome 13! (Lets pretend she's BRCA-2, not BRCA-1)	14 - Presenilin 1 (early alz)	15 - Tay-Sachs (Hexosaminidase A) - Marfan (FBN1) - Prader-Willi & Angelman (imprinting)	16 - TSC2 (tuberin) - PCKD1 (polycystin1) - α Globin Chain (α-Thal) Polycystic kidney = 16 letters = chromosome 16! Also 'alpha globin chain'	17 - P53 (Bax → bcl-2) - NF1 (Von Recklinghausen) - BRCA-1 Von Recklinghausen = 17 letters = chromosome 17!	18 - bcl-2 (apoptosis) - Edwards Trisomy - DCC (colon) - DPC4 (pancreatic)
19 - Familial Hyper Cholesterolemia (LDL receptor mutation) - ApoE4 / 2 (Late alz. / alz) (sporadic, not familial)	20	21 - Down's Trisomy - β-APP (very early alz.) - SOD1 (altered in ALS)	22 - NF2 (Merlin protein) - EWS (Ewing's sarc.) - DiGeorge syn. (11q-); Velocardiofacial syn.	23	

X	Quantitative sexual chromosome diseases	Y
<ul style="list-style-type: none"> - Menkes Syndrome (ATP7A copper Efflux Protein) - Fragile X Syndrome (CGG on gene FMR-1) - Bruton's (BTK gene) Agammaglobulinemia - Lesch-Nyhan (HPRT1) - Wiskott Aldrich (WASP Protein) - SCID (IL-2 receptor mutation) - Duchenne MD (Dystrophin) - Alport's (COL4A5) - Hemophilia A&B (F8&9) - Fabry's dz (telangiectasia & skin rashes, ↑ BUN) - Hunter's Syndrome (corneal clouding, pebbly skin) - G6PD deficiency - Ocular albinism - OTC deficiency (other urea cycle disease is AR) - Hypophosphatemic rickets (X-linked dominant) 	<ul style="list-style-type: none"> - Klinefelter Syndrome XXY (Barr Body) - Turner Syndrome XO (Shox Gene) 	

Translocations	
9:22	Philadelphia (Bcr-Abl over expression. P210KD Tyrosine Kinase. CML → better prognosis; ALL → worst prognosis)
8:14	Burkitt's Lymphoma (C-myc gene over expression)
11:14	Mantle Cell Lymphoma (Bcl-1 gene Aka Cyclin D)
14:18	Follicular NHL (Bcl-2)
15:17	AML M3 (Auer Rods, DIC)
11:22	Ewing's Sarcoma ("Patrick Ewing doubles your points")
12:21	Provides better prognosis in ALL

Taken and modified from www.Baronerocks.com