## **GENETIC DISORDERS--KEY**

Fill in the chart below to learn more about some of the possible genetic disorders a person may inherit.

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DISEASE/ BIRTH DEFECT	PEOPLE AFFECTED	WHEN APPEARS	METHOD OF INHERITANCE	EFFECT ON PERSON	TREATMENT
1. Cleft lip/pallet	ANYONE MORE COMMON IN ASIANS AND NATIVE AMERICANS	BIRTH	MULTI- FACTORAL	TWO SIDES OF LIP AND/OR PALLET NOT JOINED	SURGERY
2. Clubfoot	ANYONE	BIRTH	MULTI- FACTORAL	FOOT AND ANKLE TWISTED, MAKING IT IMPOSSIBLE TO WALK NORMALLY	SURGERY CORRECTIVE SHOES
3. Color blindness	MALES	BIRTH	X-LINKED	INABILITY TO DISTINGUISH CERTAIN COLORS	NONEJUST HELP ADAPT
4. Cystic fibrosis	ANYONE	BIRTH SOME IN ADULT- HOOD	RECESSIVE	LACK OF AN ENZYME. MUCOUS OBSTRUCTIONS IN BODY, ESPECIALLY LUNGS AND DIGESTION; GROWTH RETARDATION	RESPIRATORY HELP
5. Diabetes	ANYONE	BIRTH OR LATER	MULTI- FACTORAL	ABNORMAL METABOLISM OF SUGAR BECAUSE BODY DOESN'T PRODUCE ENOUGH INSULIN	INSULIN SHOTS/ORAL DIET
6. Down's Syndrome	MORE COMMON TO YOUNG OR OLD PAR- ENTS, OR IF YOU HAVE ALREADY HAD A DOWN'S BABY	BIRTH	CHROMO- SOMAL ERROR	SOME FORM OF MENTAL RETARDATION; OVAL SHAPED EYES; THICK, BIG TONGUE; SHORT NECK; BACK OF HEAD IS FLAT; EARS SMALL; NOSE FLAT AND WIDE; SHORTER, LOOSE JOINTS; HEART PROBLEMS	SURGERY FOR HEART THERAPY; SPECIAL ASSISTANCE HELPS FUNCTION BETTER
7. Hemophilia	MALES	BIRTH	X-LINKED (SOME SPONTAN- EOUS MUTATIONS	ABSENCE OF CLOTTING FACTOR IN BLOOD (BLEEDER'S DISEASE), CRIPPLING AND DEATH FROM INTERNAL BLEEDING	TRANS- FUSIONS; AVOID EMOTIONAL STRESS
8. Huntington's Disease (Chorea)	PARENT MUST HAVE RARE	MIDDLE AGES	DOMINANT	DETERIORATION OF BODY AND BRAIN IN MIDDLE AGE; DEATH	NONE
9. Hydro- cephalus	ANYONE SEVERE SPINA BIFIDA HIGHER CHANCE	BIRTH	MULTI- FACTORAL	OBSTRUCTION CAUSES WATER ON BRAIN; PRODUCE BRAIN DAMAGE AND DEATH	SURGERY TO PUT IN SHUNT

DIS	EASE/	PEOPLE	WHEN	METHOD OF	EFFECT ON PERSON	TREATMENT
	DEFECT	AFFECTED	APPEARS	INHERITANCE		11 1647 11 14 16 14 1
10. Ma	arfan's Indrome	PARENT MUST HAVE RARE PEOPLE WITH LONG, BONY LIMBS	BIRTH	DOMINANT	HEART MALFORMATION, HEARING LOSS, EYE WEAKNESS, IF SEVEREDEATH	HEART SURGERY, ANTIBIOTICS PHYSICAL THERAPY, REGULAR CHECKUPS
	uscular estrophy	ANYONE (X-LINKED)	SOME CHILD- HOOD OTHERS ADULT HOOD	MULTI- FACTORAL	WEAKENING OF MUSCLES. INABILITY TO WALK, MOVE, WASTING AWAY AND SOME- TIMES DEATH	NONE
	nenyl- etonuria PKU)	ANYONE	BIRTH	RECESSIVE	ABNORMAL DIGESTION OF PROTEIN; MENTAL RETARD- ATION, HYPERACTIVITY	PREVENT- ABLE DIET
13. Po	olydactyl	PARENT HASMORE IN BOYS	BIRTH	DOMINANT	EXTRA FINGERS AND TOES	CORRECTIVE SURGERY
	ickle Cell nemia	PRIMARILY AFFECTS BLACKS	BIRTH	RECESSIVE	ABNORMAL BLOOD CELLS; BOUTS OF PAIN, HEART AND KIDNEY FAILURE, LESS OXYGEN TO ALL PARTS OF THE BODY. DEATH IN CHILDHOOD. DAMAGE TO ADULT VITAL ORGANS	TRANS- FUSIONS
15. Sp Bi	pina ifida	ANYONE	BIRTH	MULTI- FACTORAL	VARYING DEGREESFROM SLIGHT CYST TO OPEN SPINE (SORES, INFERTILE, LEGS PARALYZED, POOR BLADDER AND BOWEL CONTROL)	SURGERY FOR SEVERE, PHYSICAL THERAPY
16. Ta	ay-Sachs	JEWS (ETHNIC, NOT RELIGION)	6 MONTHS DIES BY AGE 3	RECESSIVE	ENZYME DISEASE CAUSING INABILITY TO BREAK DOWN FATTY DEPOSITS IN BRAIN AND NERVE CELLS; CELLS CLOG AND SHUT DOWN NERVOUS SYSTEM; APPARENTLY HEALTHY AT BIRTH, DIES BY AGE 3	NO CURE
(C	halassemia Cooley's nemia)	GREEK AND ITALIAN	BIRTH	RECESSIVE	ABNORMAL BLOOD CELLS; PALENESS AND LISTLESSNESS, LOW RESISTANCE TO INFECT- ION; DO NOT PRODUCE ENOUGH RED BLOOD CELLS; LIVER AND SPLEEN ENLARGED	BLOOD TRANS- FUSION SURGERY LATER ON SPLEEN