

Learning Objectives After this lesson, you will be able to identify disorders that are caused by chromosomal changes, explain the difference between autosomal dominant and autosomal recessive inheritance, and describe the inheritance pattern of sickle cell anemia.

Disorder	Cause	Inheritance
Down Syndrome (trisomy 21)	Chromosomal: Down's is the most common genetic cause of intellectual disability. The risk increases with maternal age.	Autosomal recessive
Phenylketonuria (PKU)	Enzyme deficiency: PKU is caused by a deficiency of the enzyme phenylalanine hydroxylase, which is responsible for breaking down the amino acid phenylalanine.	Autosomal recessive
Sickle Cell Anemia	Red blood cell shape: Sickle cell anemia is a genetic disorder that affects the shape of red blood cells, causing them to become rigid and sticky.	Autosomal recessive
Tay-Sachs Disease	Enzyme deficiency: Tay-Sachs is a rare genetic disorder that causes a deficiency of the enzyme hexosaminidase A, which is responsible for breaking down certain lipids in the brain.	Autosomal recessive

Lesson Summary

Chromosomal Disorders

Chromosomal disorders occur as the result of errors in chromosomal structure or number. The most common chromosomal disorders are Down syndrome, which is caused by an extra copy of chromosome 21, and sickle cell anemia, which is caused by a change in the structure of the hemoglobin gene. Chromosomal disorders can be inherited or acquired.

Autosomal recessive disorders require two copies of the defective gene, one from each parent.

Human Genetic Disorders Answer Key Chapter 13

J Dewey



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Unveiling the Power of Verbal Art: An Mental Sojourn through **Human Genetic Disorders Answer Key Chapter 13**

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