

## GENETICS OF SICKLE CELL ANEMIA%0A

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Sickle cell disease is a group of disorders that affects hemoglobin, the molecule in red blood cells that delivers oxygen to cells throughout the body. People with this disorder have atypical hemoglobin molecules called hemoglobin S, which can distort red blood cells into a sickle , or crescent, shape.

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### **Sickle Cell Disease Genetics News Medical**

Sickle-cell disease is caused by a gene mutation that leads to the production of Sickle haemoglobin, which affects the function of the red blood cells in the body. This mutation is inherited from

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### **Sickle Cell Anemia A Genetic Condition Explained YourDNA**

Sickle Cell Anemia or Sickle Cell Disease (SCD), is a genetic condition that affects the red blood cells and the ability to deliver oxygen to cells in the body. Over 100,000 people in the United States are affected with SCD 1 .

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### **SCD Genetics What is the Sickle Cell Trait Generation S**

An abnormal HbS gene causes red blood cells to become sickle shaped, which can limit the amount of oxygen the red blood cells can carry through the body. The hemoglobin gene inherited from each birth parent is very important because it can cause: Sickle cell disease Sickle cell trait, but not having sickle cell disease

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### **The Genetics of Sickle Cell Anemia A Literature Review**

Abstract Sickle cell disease (SCD) is the most common form of a genetic group of hemoglobinopathies in which normal human hemoglobin (HbA) is partially or completely replaced by mutant Much has been discovered about the genetics of sickle cell anemia and this is clearly seen by analyzing the articles, since the from 15 studies analyzed

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### **Sickle Cell Disease NHGRI genome gov**

Amniocentesis, a procedure in which a needle is used to take fluid from around the baby for testing, can show whether the fetus has sickle cell disease or carries the sickle cell gene. If the test shows that the child will have sickle cell disease, some parents may choose not to continue the pregnancy.

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### **Sickle cell anemia Diagnosis and treatment Mayo Clinic**

Sickle cell disease can be diagnosed in an unborn baby by sampling some of the fluid surrounding the baby in the mother's womb (amniotic fluid) to look for the sickle cell gene. If you or your partner has been diagnosed with sickle cell anemia or sickle cell trait, ask your doctor about whether you should consider this screening.

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### **Sickle cell anemia Symptoms and causes Mayo Clinic**

Sickle cell anemia is caused by a mutation in the gene that tells your body to make the red, iron-rich compound that gives blood its red color (hemoglobin). Hemoglobin allows red blood cells to carry oxygen from your lungs to all parts of your body.

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### **Sickle cell disease Wikipedia**

The allele responsible for sickle cell anaemia can be found on the short arm of chromosome 11, more specifically 11p15.5. A person who receives the defective gene from both father and mother develops the disease; a person who receives one defective and one healthy allele remains healthy, but can pass on the disease and is known as a carrier or heterozygote.

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### **Sickle cell disease in India A perspective**

Sickle cell disease is an inherited blood condition which is most common among people of African, Arabian and Indian origin. In disease of African origin, research has led to models of care which prevent serious complications, improve the quality of life, and increase survival 1. In India, the disease is largely undocumented.

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### **Genetics Sickle Cell Anaemia**

Sickle cell anaemia is an autosomal recessive disorder in which due to point mutation glutamic acid gets replaced by valine at 6th position in beta chain of haemoglobin. To help students search

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### **Genetics of sickle cell anemia**

Sickle cell disease is a disorder that affects the red blood cells, which use a protein called hemoglobin to transport oxygen from the lungs to the rest of the body. Normally, red blood cells are

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