WHAT IS SICKLE CELL TRAIT?

Sickle Cell Trait (AS) is an inherited condition which affects the hemoglobin in your red blood cells.

It is important to know if you have sickle cell trait. Sickle cell trait is inherited from your parents, like hair or eye color. If one parent has sickle cell trait, there is a 50% (1 in 2) chance with each pregnancy of having a child with sickle cell trait. Sickle cell trait rarely causes any health problems. Some people may develop health problems under certain conditions, such as:

- Dehydration – from not drinking enough water
- Low oxygen – from over-exertion
- High altitudes – from low oxygen levels

HOW DO YOU KNOW IF YOU HAVE SICKLE CELL TRAIT OR DISEASE?

A simple blood test called a hemoglobin electrophoresis can determine your sickle cell status. Talk with your healthcare provider if you have more questions about sickle cell trait or want to be tested.

It is important to know if you have sickle cell trait before you decide to have children.

WHO CAN HAVE SICKLE CELL DISEASE AND SICKLE CELL TRAIT?

- It is estimated that SCD affects 90,000 to 100,000 people in the United States, mainly Blacks or African Americans.
- The disease occurs in about 1 of every 365 Black or African American births and in about 1 of every 36,000 Hispanic–American births.
- SCD affects millions of people throughout the world and is particularly common among those whose ancestors come from sub-Saharan Africa, regions in the Western Hemisphere (South America, the Caribbean, and Central America), Saudi Arabia, India, and Mediterranean countries such as Turkey, Greece, and Italy.
- About 1 of every 12 African Americans has sickle cell trait and about 1 of every 100 Hispanics has sickle cell trait.
- It is possible for a person of any race or nationality to have sickle cell trait.

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WHAT IS SICKLE CELL DISEASE?

Sickle cell disease (SCD) is a group of inherited red blood cell disorders.

• Hemoglobin is the protein in red blood cells that carries oxygen throughout the body.

• Each person inherits two hemoglobin genes, one from each parent. A normal gene will make normal hemoglobin (A). A specific change in the hemoglobin gene will make sickled hemoglobin (S).

• In SCD, the red blood cells become hard and sticky and look like a C-shaped farming tool called a sickle.

• Sickle cells can get stuck in small blood vessels and block the flow of blood and oxygen to organs within the body. These blockages can cause many problems.

• The most common types of sickle cell disease are hemoglobin SS, hemoglobin SC, and sickle beta thalassemia.

WHAT CAUSES SICKLE CELL DISEASE?

SCD is inherited in the same way that people get the color of their eyes, skin, and hair.

• A person with SCD is born with it.

• People cannot catch SCD from being around a person who has it.

• If a person has sickle cell disease, they inherit a sickle hemoglobin from each parent.

WHAT ARE THE POTENTIAL HEALTH PROBLEMS OF SICKLE CELL DISEASE?

Infections: The spleen helps the body fight infections. Sickle cells may get caught in the spleen, preventing it from working as well as it should. As a result, people with sickle cell disease are more likely to get infections. Children are put on antibiotics until their immune system matures.

Anemia: Sickle cells do not live as long as normal red blood cells. This causes anemia, or a low blood count. Anemia can cause weakness and fatigue.

Pain: Sickle cells that get caught in the small blood vessels of the body cause the interruption of oxygen and blood flow; these sickling crises can be very painful. This includes pain and swelling of the hands and feet in small children.

Acute Chest Syndrome: Blockage of the flow of blood to the lungs can cause acute chest syndrome (ACS). ACS is similar to pneumonia; symptoms include chest pain, coughing, difficulty breathing, and fever. It can be life threatening and should be treated in a hospital.

Stroke: Sickle cells can clog blood flow to the brain and cause a stroke. A stroke can result in lifelong disabilities and learning problems. Children under 16 are at the highest risk for stroke.

Jaundice: This is a yellowing of the eyes and skin. It is painless and occurs because of rapid breakdown and death of sickled red blood cells.

Leg Ulcers: Poor circulation due to blockage of blood vessels by sickle cells can lead to sores that are difficult to heal. These often occur around the ankles.

Complications During Pregnancy: A woman with sickle cell disease can have a healthy baby. However, risks are involved; both she and the baby should be closely monitored by a healthcare provider. Prenatal care is very important!

Other problems include:

• chest pain and trouble breathing
• organ damage
• gallstones
• blood in the urine
• eye disease
• painful erections in men

HOW IS SICKLE CELL DISEASE MANAGED?

Individuals with sickle cell disease should receive regular medical care from a hematologist and a general medical care provider.

Routine care is very important!

Treatments may include a medication called hydroxyurea which increases the body’s level of fetal hemoglobin which prevents sickling, as well as antibiotics for infections or prevention of infections, and medicines for painful events.

Everyone should eat a balanced diet, and receive regular check-ups and immunizations. Children with sickle cell disease require a few special immunizations. A yearly flu shot is also recommended.