# Sickle Cell Anemia questions with correct answers

What is sickle cell anemia? - Answer An autosomal recessive disorder resulting in a mutated form of hemoglobin (HbS)

What causes sickle cell TRAIT? - Answer ONE allele on the beta globin gene carries the sickle mutation, one normal allele for HbA (heterozygous)

What causes sickle cell DISEASE? - Answer TWO alleles on the beta globin gene carries mutation for sickle mutation (homozygous)

What does the sickle cell mutation cause resistance to? - Answer Plasmodium falciparum (malaria)

How common is sickle trait and disease in African Americans? - Answer Trait: 1 in 12

Disease: 1 in 500

Newborn screening for sickle cell disorders is optional. True or false? - Answer False (screening is mandatory but parents can opt out)

To diagnose sickle trait, what is seen on Hb electrophoresis? - Answer HbA > HbS (50-60% A, 20-45% S)

At what age will HbS amounts increase? - Answer 6 months (don't utilize the beta globin gene - where the mutation is - until 6 months old)

# **What Causes Sickle Cell Anemia Yahoo Answers**

**JE Gale** 

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Chaplaincy: What in "H" Was I Thinking? Susan R. Street-Beavers D.Min., 2015-11-25 For many the route of coming into and through CPE and the process of self discovery is mysterious and puzzling fraught with fear and uncertainty With amazing clarity and insightful wisdom Dr Susan Street Beavers demystifies the CPE process You are guided step by step with humor poetry prayer scripture and thought provoking questions through the story of Dr Susie's journey to wholeness Whether you are a nurse social worker therapist pastor elder leader in your church community or already a chaplain or just thinking and praying about it this book offers skills to enhance self awareness growth in interpersonal relationships and the strengthening of your ministry An absolute must read for seminary students church leaders and all who might consider participating in Clinical Pastoral Education CPE and desire their experience to be a journey of wholeness Dr Frank A Thomas In this book Dr Susie gives the reader a view of her personal journey through the clinical pastoral education CPE process From this book the reader will learn that many people become chaplains as a result of going through the CPE process Whether one becomes a chaplain or serves in another ministry the self reflection and growth experienced from the CPE process help make us better people Dr Susan Street Beavers shares that the growth that she experienced caused her to write What in h was I thinking The h means heaven The benefits gained by going through the CPE process can help to aid the CPE journeyer in her his walk with God as we strive to share divinity with those that we are called to serve Dr James A Nooks The Digital Divide United States. Congress. House. Committee on Small Business. Subcommittee on Empowerment, 2001 Newswatch ,2004

Men Like Us Daniel Wolfe, Gay Men's Health Crisis, Inc, 2000 For nearly two decades GMHC has provided vital support education and health information to gay men Now with Men Like Us their guidance and the insights of hundreds of gay men across America will help readers everywhere Practical down to earth and accessible this authoritative health resource covers such topics as body basics exercise and diet relationships and intimacy sexual pleasure medical care prevention and healing mental health and therapy spirituality and community and much much more Filled with expert advice from leading doctors lawyers therapists and fitness instructors to ordinary gay men whose stones provide important voices of experience Men Like Us opens a window onto the ways gay men in all their diversity care for themselves and each other **Encyclopedia of** Associations ,2006-12 **The News** ,2003 Library Journal Melvil Dewey, Richard Rogers Bowker, L. Pylodet, Charles Ammi Cutter, Bertine Emma Weston, Karl Brown, Helen E. Wessells, 2006 Includes beginning Sept 15 1954 and on the 15th of each month Sept May a special section School library journal ISSN 0000 0035 called Junior libraries 1954 May 1961 Also Obanta Newsday ,2006 Answers to Common Ouestions about Sickle Cell Disease ,1973 issued separately

Answers to Frequently Asked Questions about Sickle Cell Disease National Heart and Lung Institute,1973 Sickle

Cell Anemia and Your Child Roland B. Scott, Althea D. Kessler,1960 Questions & Answers Missouri. Department of Health, Missouri. Sickle Cell Anemia Program,1987\* Answers to Common Questions about Sickle Cell Disease, 1973

**Sickle Cell Anemia** Jane S. Lin-Fu,1975 Sickle Cell Anemia Judy Monroe Peterson, 2008-08-15 Sickle cell anemia is a genetic disease of the blood It is caused by a defect in one gene of a person Genes are the elements in cells that carry the information that determines traits such as hair or eye color In sickle cell anemia a defect in the gene controls how hemoglobin is made This defect can be passed from parents to their children Students explore the history of sickle cell anemia the pioneering doctors who studied its cause and early treatments They also investigate hemoglobin S who gets sickle cell and how the gene mutation is inherited. They learn about the different types of sickle cell disease and treatments including blood transfusions and bone marrow transplants and some of the promising new research in medicines and gene Sickle cell anemia Rumi Michael Leigh, 2022-08-21 Sickle Cell Anemia Things You Should Know Questions and therapy Answers explains this inherited blood disorder in a simple question and answer format covering causes symptoms complications and treatments Inside you will learn what sickle cell anemia is how it affects red blood cells and hemoglobin and why these cells become rigid sticky and sickle shaped The book explains how sickle cell anemia shortens the life of red blood cells and disrupts normal blood flow leading to serious health problems It covers symptoms such as extreme pain fatigue infections slow growth delayed puberty swelling in the hands and feet and vision problems Complications are also discussed including stroke pulmonary hypertension organ damage blood clots gallstones priapism ulcers blindness and pregnancy risks Readers will also find explanations of diagnostic methods from blood tests to prenatal testing and treatment options such as medications blood transfusions and bone marrow transplant This book is suitable for anyone who wants to understand sickle cell anemia its effects on the body and current approaches to treatment in a direct and easy to read way

Questions and Answers about Sickle Cell Disease Illinois. Department of Public Health,1972 Answers to Questions Frequently Asked about Sickle Cell Disease ,1973 Answers to Frequently Asked Questions about Sickle Cell Disease ,1973 Sickle Cell Anemia: From Basic Science to Clinical Practice Elena Ledger,2021-11-16 A sickle cell disease is a group of blood disorders that a person inherits from parents They usually occur when the person inherits two abnormal copies of the hemoglobin gene Sickle cell anemia is the most common type of sickle cell disease It causes an abnormality in hemoglobin the oxygen carrying protein found in red blood cells The problems due to sickle cell anemia begin to appear around 5 to 6 months of age It leads to problems such as sickle cell crisis swelling in hands and feet stroke and bacterial infections The care of people suffering from sickle cell anemia includes infection prevention with vaccination and antibiotics folic acid supplementation and pain medication A bone marrow transplant is also used in certain cases This book consists of contributions made by international experts It contains some path breaking studies in sickle cell anemia It will serve as a valuable source of reference for graduate and post graduate students

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