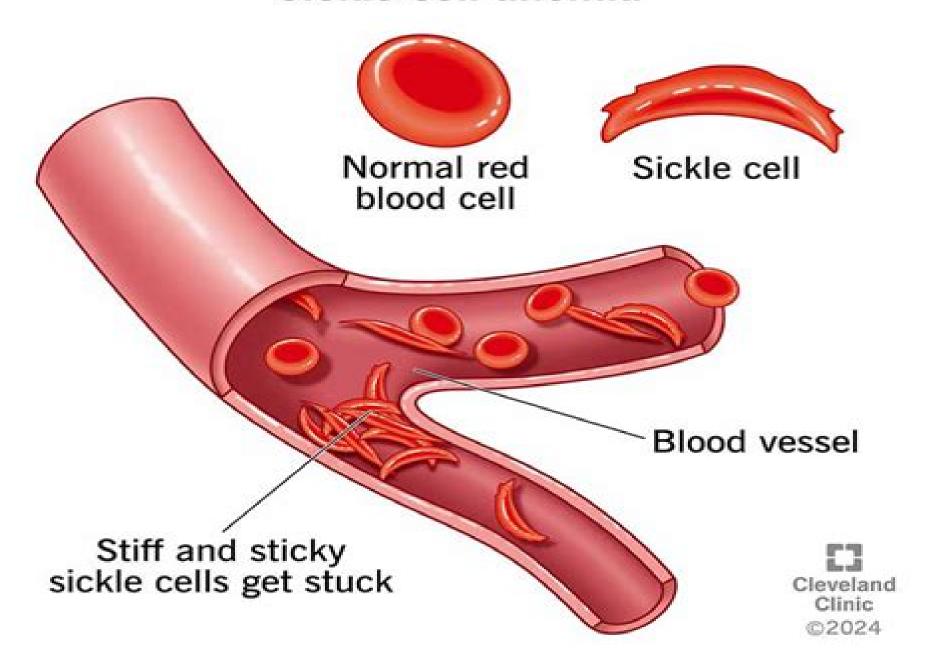
Sickle cell anemia



Sickle Cell Anemia

National Sickle Cell Disease Program, National Heart and Lung Institute

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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 therapy Sickle Cell Anemia Jane S. Lin-Fu,1975 Sickle Cell Anemia Fernando Ferreira Costa, Nicola Conran, 2016-03-29 Although sickle cell anemia was the first molecular disease to be identified its complex and fascinating pathophysiology is still not fully understood A single mutation in the beta globin gene incurs numerous molecular and cellular mechanisms that contribute to the plethora of symptoms associated with the disease Our knowledge regarding sickle cell disease mechanisms while still not complete has broadened considerably over the last decades Sickle Cell Anemia From Basic Science to Clinical Practice aims to provide an update on our current understanding of the disease s pathophysiology and use this information as a basis to discuss its manifestations in childhood and adulthood Current therapies and prospects for the development of new approaches for the management of the disease **Sickle Cell Anemia** Ruth Bjorklund, 2010-10-15 Provides comprehensive information on the causes are also covered treatment and history of sickle cell anemia Provided by publisher Sickle Cell Anemia ,1990 In the Blood Melbourne Tapper, 1999-02-04 Tapper shows how sickle cell anemia was used to promote the superiority of racial purity and to characterize the black body as contaminated Sickle Cell Anemia Jane S. Lin-Fu,1975 Sickle Cell Anemia: New Insights for the Healthcare Professional: 2011 Edition, 2012-01-09 Sickle Cell Anemia New Insights for the Healthcare Professional 2011 Edition is a ScholarlyBrief that delivers timely authoritative comprehensive and specialized information about Sickle Cell Anemia in a concise format The editors have built Sickle Cell Anemia New Insights for the Healthcare Professional 2011 Edition on the vast information databases of ScholarlyNews You can expect the information about Sickle Cell Anemia in this eBook to be deeper than what you can access anywhere else as well as consistently reliable authoritative informed and relevant The content of Sickle Cell Anemia New Insights for the Healthcare Professional 2011 Edition has been produced by the world's leading scientists engineers analysts research institutions and companies All of the content is from peer reviewed sources and all of it is written assembled and edited by the editors at ScholarlyEditions and available exclusively from us You now have a source you can cite with authority confidence and credibility More information is available at http www ScholarlyEditions com Fact Sheet, Sickle Cell Anemia National Sickle Cell Disease Program, National Heart and Lung Institute, 1975 Focus on Sickle Cell Research Ralph L. Plasmar, 2004 Sickle cell

anaemia is an inherited blood disorder characterised primarily by chronic anaemia and periodic episodes of pain and occurring in approximately 1 in every 400 African American infants born in the United States each year Individuals of Mediterranean Arabian Caribbean South and Central American and East Indian ancestry can also be affected The underlying problem involves haemoglobin a component of the red cells in the blood The haemoglobin molecules in each red blood cell carry oxygen from the lungs to the body organs and tissues and bring back carbon dioxide to the lungs In sickle cell anaemia the haemoglobin is defective After the haemoglobin molecules give up their oxygen some of them may cluster together and form long rod like structures These structures cause the red blood cells to become stiff and to assume a sickle shape Unlike normal red cells which are usually smooth and donut shaped the sickled red cells cannot squeeze through small blood vessels Instead they stack up and cause blockages that deprive the organs and tissue of oxygen carrying blood This process produces the periodic episodes of pain and ultimately can damage the tissues and vital organs and lead to other serious medical problems Unlike normal red blood cells which last about 120 days in the bloodstream sickled red cells die after only about 10 to 20 days Because they cannot be replaced fast enough the blood is chronically short of red blood cells a condition called anaemia Sickle cell anaemia is caused by an error in the gene that tells the body how to make haemoglobin The defective gene tells the body to make the abnormal haemoglobin that results in deformed red blood cells This book gathers the latest Sickle Cell Disease Charles F. Whitten, John F. Bertles, 1989 research in this important field **Explained** Dr. John Koramoa, 2014-04-11 Sickle cell disease is the most common genetic disease world wide For a person to suffer the disease he or she has to inherit the faulty gene from each parent It affects millions of people in the world It is a chronic illness of serious proportions Medical psychological and socioeconomic consequences place a great burden on sufferers and their Carers In the past sufferers of sickle cell anaemia survived for a few years only When only one sickle cell gene is inherited by a person with normal haemoglobin the condition is referred to as the carrier state or the trait Individuals with the trait have no medical problems under normal conditions of life This is how the gene has been able to survive and be passed on from generation to generation The trait can thus have serious implications for marriage and childbearing It is therefore imperative for people intending to have children to know about it in order to help prevent the disease Sickle cell disease which was once thought to be restricted to only a few areas especially among Black people is now known to be world wide It has been described as a world health problem This book provides information in an easily readable format for everybody to understand sickle cell and its related conditions There is the need for greater awareness and knowledge of the disease and the trait to avoid confusion misunderstanding and myths which hamper their proper management and prevention At present there is no readily available cure for sickle cell disease However with proper management of the many ill effects of the disease sufferers can be assisted to live comfortable lives and for many years Life style measures are paramount and are referred to in some detail in the book It is a great resource Sickle Cell Anemia George Beshore, 1994-01-01 Looks at

the history and symptoms of sickle cell anemia describes how it is treated and discusses current research *Understanding* Sickle Cell Disease Miriam Bloom, 2009-10-20 Although more is known about sickle cell disease than about any other inherited disease no cure for it exists In America alone about one in 375 who are of African ancestry is born with sickle cell disease A smaller number of Americans descended from families from the Mediterranean area the Middle East and India also are affected In addition about eight percent of black Americans who do not suffer from the disease itself carry the gene for it that can be transmitted to their children Sickle cell disease is of enormous biological social and historic importance It was first described in medical literature almost a century ago Improvements during the past two decades in our understanding of the disease and in medical care are permitting those afflicted to live longer more comfortable and more productive lives This book was written for all who are interested in this disease those who have it their families the carriers of the sickle cell gene teachers and those who wish to update their information about it This overview of sickle cell disease explains what it is and how it is inherited as well as the relationship between the sickle cell gene and its geographic origins the way the gene has been spread throughout history and the effect of sickle cell hemoglobin on red blood cells that carry it Understanding Sickle Cell Disease describes the variety of symptoms in both children and adults and details the emotional aspects of the disease Of particular interest is a chapter on the care especially the home care of those who are affected This book explains how it is possible today for couples carrying the genes to raise families free of the disease Although there is no known cure for sickle cell disease there is little doubt that one will ultimately be devised This volume surveys current research efforts and the Research, Treatment, and Prevention of Sickle Cell Anemia United States. Congress. House. promise they hold Committee on Interstate and Foreign Commerce. Subcommittee on Public Health and Environment, 1972 The Clinical Features of Sickle Cell Disease Graham R. Serjeant, 1974 This monograph summarizes the observations of a six year study of S hemoglobinopathies that was undertaken in Jamaica The principal aim of this research was the establishment of the true clinical picture of sickle cell anaemias as revealed by the study of a large group of adult patients A sickle cell clinic was formed in Kingston medical center and five peripheral clinics were established in rural hospitals The observations of the research team indicate that the symptoms and signs of classical sickle cell anaemia Hb S are similar to those observed in other parts of the world but that the disease is more benign the jaundice less marked the crises fewer and less severe and the expectancy of life longer when compared with the manifestations of sickle cell anaemia in Africa and the United States From Pathology of Sickle Cell Disease Joseph Song, 1971 Sickle-cell Anemia and Thalassemia Richard book jacket George Huntsman, Canadian Sickle Cell Society, 1987 Sickle Cell Anemia Alvin Silverstein, Virginia B. Silverstein, Laura Silverstein Nunn, 1997 A hereditary disease that is suffered by more than 60 000 Americans today sickle cell anemia affects red blood cells Includes the history of the disease case studies and treatments available today **New Developments in** Sickle Cell Disease Research Paul D. O'Malley, 2006

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