Sickle Cell Anemia A Fictional Reconstruction Answer Key

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NATIONAL CENTER FOR CASE STUDY TEACHING IN SCIENCE ...
“Sickle Cell Anemia” by Debra Stamper Page 1 * Disclaimer: Th is case is a work of fi ction that refers to real events and people All of the discoveries mentioned in Section 1 were made by the individuals they are attributed to, as were the observations made by Dr Vernon Hahn described in Section 2
SCD 2-9-2017 final JJ & DH - NACNS
Clinicopathologic findings in sickle cell anemia The findings are a consequence of infarctions, anemia, hemolysis, and recurrent infection From Damjanov, 2000 Found on the free dictionary sight Psychosocial Life long illness with many interactions with health care providers
The Mystery of the Crooked Cell
A Closer Look at the Cause of Sickle Cell Anemia Your research has determined that sickle cell hemoglobin differs from normal hemoglobin in the net negative charge on the proteins This discovery is an important one; it identifies a characteristic that can be used to diagnose sickle cell anemia
Sickle Cell Trait and Sickle Cell Disease: A Case Study ...
Clinical case study: Sickle cell disease Shoumita Dasgupta, PhD Sickle Cell Trait and Sickle Cell Disease: A Facilitator’s Guide, Part I Abstract This case is a classical example of an African American individual who is a heterozygous for sickle cell disease and who does not manifest any symptoms

until

miniPR™ Sickle ell Genetics Lab: Diagnosing aby Marie

the sickle cell allele (referred to in shorthand as HbS) People who have either one or two copies of a normal beta-globin (HbA) allele will not be sick with sickle cell disease Even though heterozygous individuals, those with one HbS and one HbA allele, do not usually show signs of sickle cell anemia, they

A 19-year-old woman with sickle cell disease and pain

A 19-year-old woman with homozygous sickle cell disease presents to her family physician's office with acute generalized bone pain She fictional Correspondence to: Richard Ward, and acute painful crises in sickle cell anemia: effects on hospi-

Mystery of the Crooked Cell - Missouri University of ...

Sickle cell anemia is a genetic disease that affects the hemoglobin molecule in red blood cells laboratory where they apply the concepts acquired in the pre-lab to test a fictional patient for the presence of sickle cell hemoglobin using gel electrophoresis Mystery of the Crooked Cell: Background 2

Sickle Cell Today - University of South Alabama

flyers" with sickle cell disease that seemed to live in the emergency room most days of the month It was a standing joke amongst the doctors how two patients with sickle cell disease, Margaret and Bobby, were in the emergency room for their daily shots of demerol and how they were just addicts Every time I heard this it would make me so

Sample Background Answers to Questions in the Student Guide

There is very strong selection against sickle-cell anemia Malaria selects for the heterozygous condition, which does not produce sickle-cell anemia By selecting for the heterozygote, malaria increases the frequency of the allele for hemoglobin S, but the statement confuses the allele with the recessive phenotype of sickle-cell anemia 4

Title

Sickle cell syndrome is a genetic disease caused by a change in the which results in anemia With an abnormally high amount of red blood cells breaking down, there is an excess of diagnose a fictional patient to determine if the person has sickle cell syndrome, is a

Frequency measures of epidemiological studies

diabetes, hypertension, heart disease, malaria, sickle cell anemia, among others Polio is also one of the diseases suffered by children in some developing countries A lot of health-related problems have been bedeviling people all over the world How to measure the diseases, determining their causes and plan the appropriate means

CASE TEACHING NOTES for "Selecting the Perfect Baby: The ...

CASE TEACHING NOTES for "Selecting the Perfect Baby: Although the debate and doctors described are fictional, the case is based on actual events from the late (PKU), cystic fibrosis, and sickle cell anemia It is important for students to understand recessive-linked disorders in order to discuss why the procedures proposed in

SPEAKER PROFILES ** Alphabetized by first names**

SPEAKER PROFILES ** Alphabetized by first names** Adrienne Shapiro Co-founder, Axis Advocacy Alexis Wardlow is a Warrior living with Sickle Cell Anemia SS Although she has struggled with sickle cell, as well as two other chronic conditions (Crohn's Disease and Primary fictional novel about extraterrestrials who come to Earth to show

SCIENCE: BIOLOGY UNIT #3: EVOLUTION MECHANISMS (5 ...

population of fictional organisms to visualize the evolution of color in the population Students color a Worm Worksheet with the simulation or if using the manual version included at the site This simulation focuses on drift Sickle Cell Anemia attached, page 12-13 7

Disclosure of Confidential Public Health Information ...

Disclosure of Confidential Public Health Information, Records, or Data Policy #ES 01-16-002 I PURPOSE AND SCOPE OF POLICY The purpose of this policy is to outline the legal status of confidential public health information, records, or A seven year old black male in Taylor County has sickle cell anemia This

Medical School Introduces Teens To Biotechnology Careers

sickle cell anemia, or testing a fictional patient for HIV infection During "graduation," the junior scientists shared the results of their six weeks of exploration with family and friends at CityLab BIOTECHNOLOGY SCIENCE FAIR CityLab is also available to any high school or middle school student who wants to do a science fair project relat

Optimal mutation rate Dominance and its implications Why ...

Fictional data for sickle-cell hemoglobin (alleles A and S) in African-American adults Normal AA 400 Carrier AS 90 A ected SS 10 Suppose I told you: { How many people I sampled { How many of each allele I found { How many AS carriers I found Are there any possible surprises left in the data? (AA? SS?) This is why there is only 1 df