

Sickle Cell Anemia: A Fictional Reconstruction*

by
Debra Stamper
Department of Biology
King's College



Part I – The Inquiry Begins

It was a brisk fall day in Boston—the type of day that Dr. William Castle preferred to start with a cup of coffee while he caught up on his correspondence, which often appeared to be an endless task. As a faculty member of Harvard Medical School, he had always received a fair amount of inquiries, but after he had published his data indicating that pernicious anemia was due to a vitamin B₁₂ deficiency, the amount of mail he received was sometimes overwhelming.

Sitting in his reclining chair he sorted through the large pile that had accumulated. He began to meticulously segregate it into smaller piles he would open in a prescribed order. Usually the delegation of a particular envelope was a relatively easy choice. There were a few that caused him to pause for a moment, such as the one he was currently holding. The return address indicated it was from an Irving Sherman at Johns Hopkins University. Since it was from someone he had never heard of, he was inclined to place it in the pile to be opened later. But on this day he decided to take another sip of coffee and see what Mr. Sherman had to say (see next page).



* *Disclaimer:* This case is a work of fiction 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. The time between discoveries has been dramatically condensed, however. Every effort has been made to present the scientific considerations concisely and accurately. Any errors should be attributed to the author and not the original investigators.

Sickle Cell Anemia A Fictional Reconstruction Answer Key

**Organisation for Sickle Cell Anaemia
Research**



Sickle Cell Anemia A Fictional Reconstruction Answer Key:

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 Anaemia* Organisation for Sickle Cell Anaemia Research, 1979 **Menace in My Blood** Ola Tamedu, 2011 **Sickle Cell Anaemia**, 1993

Sickle Cell Anemia A Fictional Reconstruction Answer Key Book Review: Unveiling the Magic of Language

In an electronic digital era where connections and knowledge reign supreme, the enchanting power of language has be more apparent than ever. Its capability to stir emotions, provoke thought, and instigate transformation is actually remarkable. This extraordinary book, aptly titled "**Sickle Cell Anemia A Fictional Reconstruction Answer Key**," compiled by a very acclaimed author, immerses readers in a captivating exploration of the significance of language and its profound affect our existence. Throughout this critique, we shall delve into the book is central themes, evaluate its unique writing style, and assess its overall influence on its readership.

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