

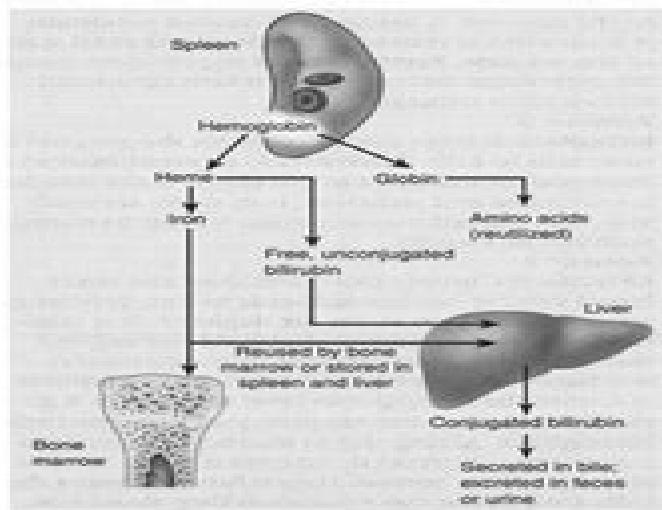
## CHAPTER 13

### SECTION I: ASSESSING YOUR UNDERSTANDING

#### Activity A

- biconcave, cell membrane
- iron
- nucleus
- 4
- glycolytic
- methemoglobin
- red blood cell count (RBC)
- hematocrit
- mean corpuscular hemoglobin concentration (MCHC)
- Anemia
- hypoxia
- Hemolytic
- sickle cell, thalassemias
- spherocytosis
- $\beta$ -Thalassemias,  $\alpha$ -thalassemias
- glucose-6-phosphatase (G6PD)
- Iron deficiency
- chronic blood loss
- Megaloblastic
- Pernicious
- Aplastic anemia
- Polycythemia
- oxygen
- conjugate
- anemia

#### Activity B



#### Activity C

1. e	2. a	3. f	4. c	5. i
6. j	7. b	8. d	9. h	10. g

#### Activity D

- The hemoglobin molecule is composed of two pairs of structurally different  $\alpha$  and  $\beta$  polypeptide chains. Each of the four-polypeptide chains consists of a globin (protein) portion and heme unit, which surrounds an atom of iron that binds oxygen. Thus, each molecule of hemoglobin can carry four molecules of oxygen. The binding that occurs is cooperative, or allosteric. When one oxygen molecule binds, it makes it easier for the next to bind. The process also works in reverse.
- A group of large phagocytic cells found in the spleen, liver, bone marrow, and lymph nodes facilitates the destruction of RBCs. These phagocytic cells recognize old and defective red cells and then ingest and destroy them in a series of enzymatic reactions. During these reactions, the amino acids from the globulin chains and iron from the heme units are salvaged and reused. The bulk of the heme unit is converted to bilirubin, which is insoluble in plasma and attaches to plasma proteins for transport. Bilirubin is removed from the blood by the liver and conjugated with glucuronic acid to render it water soluble so that it can be excreted in the bile.
- (1) Manifestations of impaired oxygen transport and the resulting compensatory mechanisms, (2) reduction in red cell indices and hemoglobin levels, and (3) signs and symptoms associated with the pathologic process that is causing the anemia.
- Premature destruction of the cells due to the rigid nondeformable membrane occurs in the spleen, causing hemolysis and anemia due to a decrease in red cell numbers. Secondly, vessel occlusion, a complex process involving an interaction among the sickle cells, endothelial cells, leukocytes, platelets, and other plasma proteins, will interrupt blood flow. The adherence of sickle cells to the vessel endothelium causes endothelial activation with liberation of inflammatory mediators and substances that increase platelet activation and promote blood coagulation.
- Exposure to high doses of radiation, chemicals, and toxins that suppress cellular activity directly or through immune mechanisms is the standard cancer treatment. Chemotherapy and irradiation commonly result in bone marrow depression, which causes anemia, thrombocytopenia, and neutropenia. Identified toxic agents including benzene, the antibiotic chloramphenicol, and the alkylating agents and antimetabolites used in the treatment of cancer will decrease bone marrow of stem cells, thus affecting the production of RBCs.
- Viscosity rises exponentially with the hematocrit and interferes with cardiac output and blood flow. Hypertension is common, and there may

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**James Howard Otto**

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## **Table of Contents Chapter 13 Study Guide Answers Biology**

1. Understanding the eBook Chapter 13 Study Guide Answers Biology
  - The Rise of Digital Reading Chapter 13 Study Guide Answers Biology
  - Advantages of eBooks Over Traditional Books
2. Identifying Chapter 13 Study Guide Answers Biology
  - Exploring Different Genres
  - Considering Fiction vs. Non-Fiction
  - Determining Your Reading Goals
3. Choosing the Right eBook Platform
  - Popular eBook Platforms
  - Features to Look for in an Chapter 13 Study Guide Answers Biology
  - User-Friendly Interface
4. Exploring eBook Recommendations from Chapter 13 Study Guide Answers Biology
  - Personalized Recommendations
  - Chapter 13 Study Guide Answers Biology User Reviews and Ratings
  - Chapter 13 Study Guide Answers Biology and Bestseller Lists
5. Accessing Chapter 13 Study Guide Answers Biology Free and Paid eBooks

- Chapter 13 Study Guide Answers Biology Public Domain eBooks
- Chapter 13 Study Guide Answers Biology eBook Subscription Services
- Chapter 13 Study Guide Answers Biology Budget-Friendly Options

6. Navigating Chapter 13 Study Guide Answers Biology eBook Formats

- ePUB, PDF, MOBI, and More
- Chapter 13 Study Guide Answers Biology Compatibility with Devices
- Chapter 13 Study Guide Answers Biology Enhanced eBook Features

7. Enhancing Your Reading Experience

- Adjustable Fonts and Text Sizes of Chapter 13 Study Guide Answers Biology
- Highlighting and Note-Taking Chapter 13 Study Guide Answers Biology
- Interactive Elements Chapter 13 Study Guide Answers Biology

8. Staying Engaged with Chapter 13 Study Guide Answers Biology

- Joining Online Reading Communities
- Participating in Virtual Book Clubs
- Following Authors and Publishers Chapter 13 Study Guide Answers Biology

9. Balancing eBooks and Physical Books Chapter 13 Study Guide Answers Biology

- Benefits of a Digital Library
- Creating a Diverse Reading Collection Chapter 13 Study Guide Answers Biology

10. Overcoming Reading Challenges

- Dealing with Digital Eye Strain
- Minimizing Distractions
- Managing Screen Time

11. Cultivating a Reading Routine Chapter 13 Study Guide Answers Biology

- Setting Reading Goals Chapter 13 Study Guide Answers Biology
- Carving Out Dedicated Reading Time

12. Sourcing Reliable Information of Chapter 13 Study Guide Answers Biology

- Fact-Checking eBook Content of Chapter 13 Study Guide Answers Biology
- Distinguishing Credible Sources

13. Promoting Lifelong Learning

- Utilizing eBooks for Skill Development

- Exploring Educational eBooks
- 14. Embracing eBook Trends
  - Integration of Multimedia Elements
  - Interactive and Gamified eBooks

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