

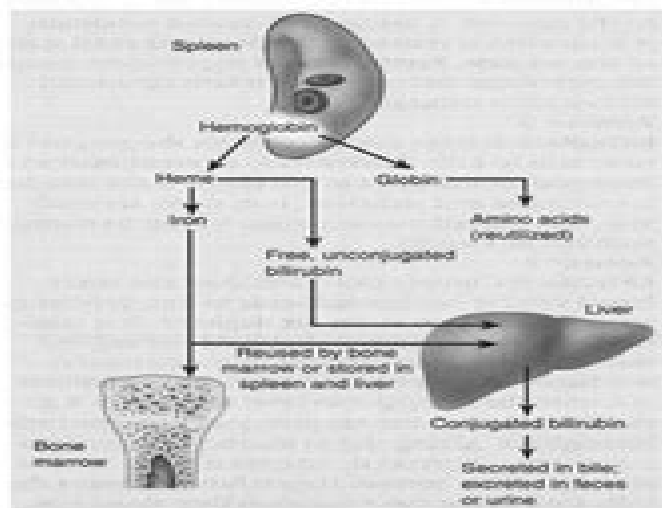
CHAPTER 13

SECTION I: ASSESSING YOUR UNDERSTANDING

Activity A

1. biconcave, cell membrane
2. iron
3. nucleus
4. 4
5. glycolytic
6. methemoglobin
7. red blood cell count (RBC)
8. hematocrit
9. mean corpuscular hemoglobin concentration (MCHC)
10. Anemia
11. hypoxia
12. Hemolytic
13. sickle cell, thalassemias
14. spherocytosis
15. β -Thalassemias, α -thalassemias
16. glucose-6-phosphatase (G6PD)
17. Iron deficiency
18. chronic blood loss
19. Megaloblastic
20. Pernicious
21. Aplastic anemia
22. Polycythemia
23. oxygen
24. conjugate
25. 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

1. The hemoglobin molecule is composed of two pairs of structurally different α and β 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.
2. 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 glucuronide to render it water soluble so that it can be excreted in the bile.
3. (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.
4. 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.
5. 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.
6. 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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