

**Unit 7: Hematology**  
**Chapter 33 & 34**  
**ONLINE CONTENT (2H)**

**Complete the worksheet and submit in the Unit 7: Hematology dropbox by March 17, 2025 at 0800. Please be sure to bring a copy to class on March 17, 2025.**

| Table 1                        | <b>Iron Deficiency Anemia</b>   | <b>Thalassemia</b>  | <b>Cobalamin (Vitamin B<sub>12</sub>) Deficiency</b>                 | <b>Folic Acid Deficiency</b>  |
|--------------------------------|---|---|--|---|
| <b>Etiology</b>                | May develop from inadequate diet intake, absorption, blood loss or hemolysis. Normal iron intake is usually enough to meet the needs of men and older women. It may be adequate for people with higher needs (ex: pregnant women, and menstruating) | Group of diseases involving inadequate production of Hgb, which decreases RBC production. | Caused by an absence of intrinsic factor.                            | Folic acid is needed for DNA synthesis leading to RBC formation and maturation. |
| <b>Clinical Manifestations</b> | May be asymptomatic but could have pallor changes, glossitis, cheilitis, headache, and burning sensation of the tongue.   | Cardiac complications, lung disease, hypertension, and endocrine problems                 | Tissue hypoxia, GI upset, weakness, impaired cognition, ataxia, etc. | GI problems, dysphagia, diarrhea, neuro problems                                |
| <b>Diagnostic Studies</b>      | Hemoglobin, Hematocrit, RBC, Serum Iron, Occult blood   | RBC, Hgb, Iron, Bilirubin, B12  | Hgb, Hct, RBC, Iron, MCV, bilirubin, B12                             | Serum folate levels, serum cobalamin  |
| <b>Drug Therapy</b>            | Iron, Vitamin C   | Blood transfusion, Exjade, Deferoxamine   | Cobalamin, B12   | Replacement therapy with folic acid   |
|                                |   | Thalassemia minor   | Assess neuro   | Teach patient to eat  |

|                           |   |   |  |                          |
|---------------------------|---|---|--|--------------------------|
| <b>Nursing Management</b> | Treat underlying problem that is causing iron loss such as malnutrition, alcohol use, or poor iron absorption | does not require treatment because the body adapts to the reduction of normal Hgb | problems, reduce risk for injury, and protect the patient from burns, falls, and trauma. | foods high in folic acid |
|---------------------------|---|---|--|--------------------------|

| Table 2                        | <b>Anemia of Chronic Disease</b>   | <b>Aplastic Anemia</b>   | <b>Acute Anemia due to Blood Loss</b>  | <b>Chronic Anemia due to Blood Loss</b>                                    |
|--------------------------------|--|--|--|--|
| <b>Etiology</b>                | Chronic disease associated with underproduction of RBC and mild shortening of RBC survival | Due to autoimmune activity by autoreactive T lymphocytes.  | Acute blood loss due to trauma, Sx, and problems that disrupt the vascular system                                    | The effects of chronic blood loss are commonly due to depleted iron stores |
| <b>Clinical Manifestations</b> | Bleeding, cancer, infectious diseases, HF  | Suppression of bone marrow elements, fatigue, dyspnea  | SOB, hypotension, pain, shock, numbness  | Poor renal function, SOB, weakness, fatigue, Jaundice                      |
| <b>Diagnostic Studies</b>      | RBC, WBC, Iron, Folate, Cobalamin  | Hgb, WBC, platelet values, RBC, bone marrow biopsy   | RBC, Hgb, HCT  | RBC, Hgb, HCT, BUN, Creatinine   |
| <b>Drug Therapy</b>            | Blood transfusions, EPO therapy  | HSCT, blood transfusion, cyclosporine  | Blood transfusion  | Iron, blood transfusions,  |
| <b>Nursing Management</b>      | Identifying and correcting the cause of RBC deficiency                                     | Removing the causative agent when possible and providing support until the pancytopenia resolves | Replace blood volume to prevent shock, promote coagulation to prevent further bleeding, find the source of the bleed | Identify the source and stopping of bleeding, pt may need iron supplements |

| Table 3                        | Acquired Hemolytic Anemia   | Hemochromatosis  | Polycythemia   |
|--------------------------------|---|--|--|
| <b>Etiology</b>                | Results from hemolysis of RBC's from extrinsic factors. These factors include physical destruction, antibody reactions, and infectious agents | Iron overload disorder characterized by increased intestinal iron absorption       | Production and presence of increased RBC. The increase can be so great that blood circulation is impaired because of the increased blood viscosity |
| <b>Clinical Manifestations</b> | Fatigue, Dyspnea, Infection, Pain   | Fatigue, arthralgia, impotence, abdominal pain, and weight loss                    | Hypertension, hypervolemia, vertigo, tinnitus, dizziness, visual changes   |
| <b>Diagnostic Studies</b>      | RBC, WBC, Hgb, HCT  | Serum iron, TIBC, Serum Ferritin, Liver biopsy                                     | Hgb, HCT, RBC, bone marrow examination   |
| <b>Drug Therapy</b>            | Glucocorticoids, Rituximab  | Iron-chelating drugs, deferoxamine, Deferiprone                                    | Myelosuppressive agents such as Hydroxyurea, or busulfan   |
| <b>Nursing Management</b>      | General supportive care until the causative agent can be eliminated or at least made less injurious to the RBC's                              | Manage problems from organ involvement with the usual treatment for these problems | Assess pt's nutrition status, I/O, GI symptoms, and pain. Begin activity and drug therapy to decrease thrombus formation                           |

***In order to receive full credit (2H class time) for this assignment, it must be completed in its entirety by the due date/time assigned. Any assignment not completed in its entirety by the due date and time will result in missed class time and must be completed by the end of the semester to pass the course.***