

Hematology and Hematologic Malignancies

Cancer of the formed elements of the blood

What is a hematological malignancy?

- A hematologic malignancy is a malignancy (or cancer) of any of the formed elements in the blood.
- The malignancies may be classified into
 - **Lymphomas**
 - Hodgkins versus non-Hodgkins
 - **leukemias**
 - Chronic versus acute

Etiology of hematological malignancies

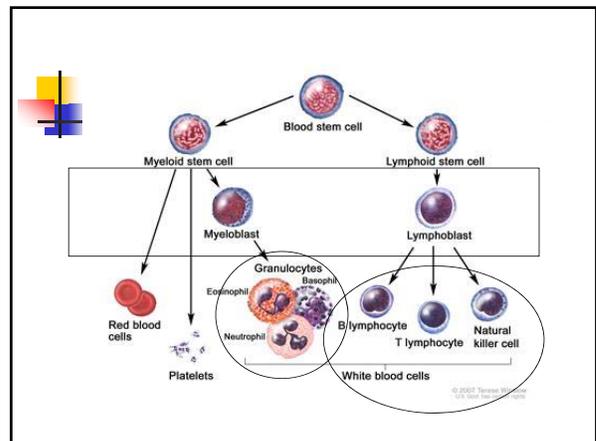
1. **Host Factors**
 - a. Hereditary
 - b. Chromosomal abnormalities
 - c. Immunodeficiency
2. **Environmental Factors**
 - a. Ionizing radiation
 - b. Chemicals
 - c. Drugs
 - d. Viruses

Leukemias

- Divided as
- Acute or chronic
- Myeloid or lymphoid

Lymphocytic and myelocytic

- The lymphocytic leukemias are caused by cancerous production of lymphoid cells
- myelogenous leukemia, begins by cancerous production of young myelogenous cells-precursors of WBC other than lymphocytes



Acute vs Chronic

- **Chronic** in which the onset is gradual ,the disease is **less** aggressive, and the cells involved are usually more mature cells.
- **Acute** in which the onset is usually rapid, the disease is **very** aggressive, and the cells involved are usually poorly differentiated with many **BLASTS** Clinically, acute leukemia is defined as a disease in which the patient die within 6 months without treatment.

Leukemias

- ALL – acute lymphocytic leukemia
- CLL – chronic lymphocytic leukemia
- AML-acute myeloid leukemia
- CML-chronic myeloid leukemia

Acute vs Chronic Leukemia

	Acute	Chronic
Age	All ages	Usually adults
Clinical Onset	sudden	insidious
Course (untreated)	6 mo	2-6 yrs
Leukemic cells	Blasts	More mature cells
WBC count	Variable	High

Acute vs Chronic

- Acute Leukemia= Blasts in marrow and often blood.
- Chronic Leukemia= mature appearing cells in marrow and blood.

Acute leukemia

- Rapid progression of symptoms
- Uncontrolled proliferation of blast cells in bone marrow results in bone marrow failure
- Blast cells infiltrate organs causing problems

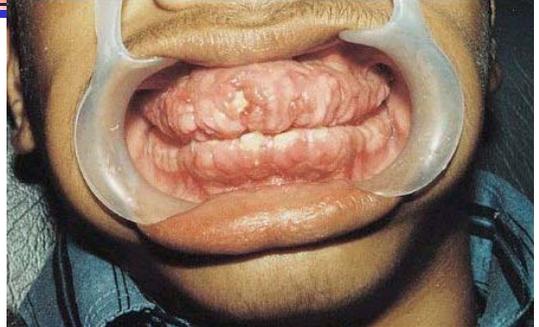
Bone Marrow Failure

- **Leukopenia** : Infections, sepsis
- **Anemia**: Fatigue, Pallor
- **Thrombocytopenia**: Bleeding

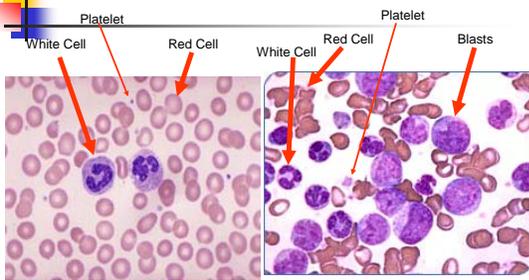
Furthermore, Infiltration of tissues/organs causes:

- Enlargement of liver, spleen, lymph nodes
- Gum hypertrophy
- Bone pain
- Other organs: CNS, skin, testis, any organ

Gum Hypertrophy



Pictures Of Blood



Normal human blood

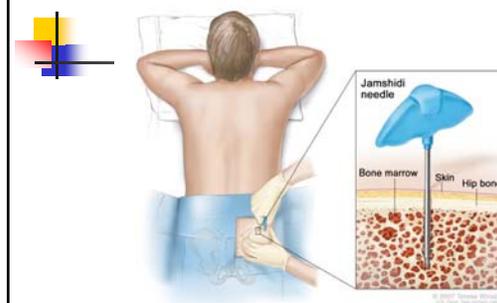
Blood with leukemia

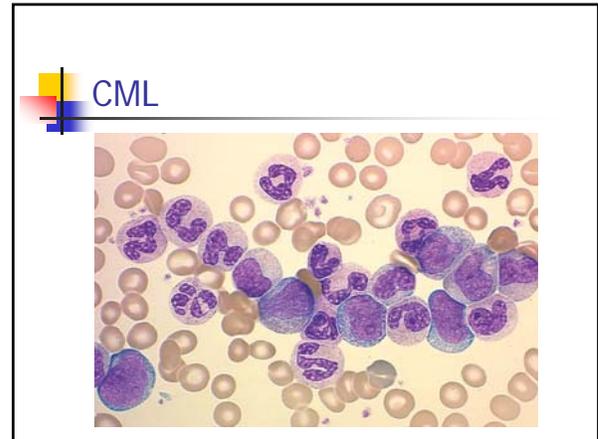
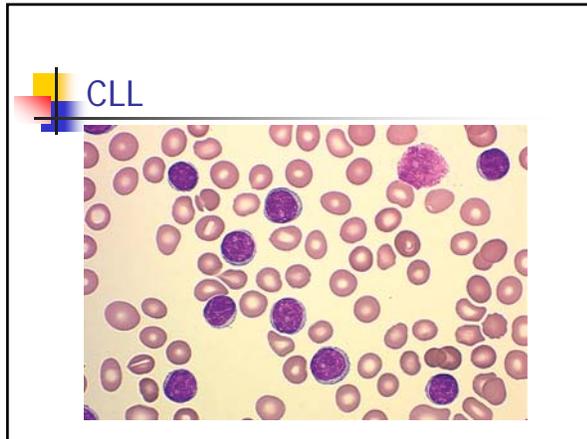
Sources from Arghineh.umdnj.edu

Sources from beyond2000.com

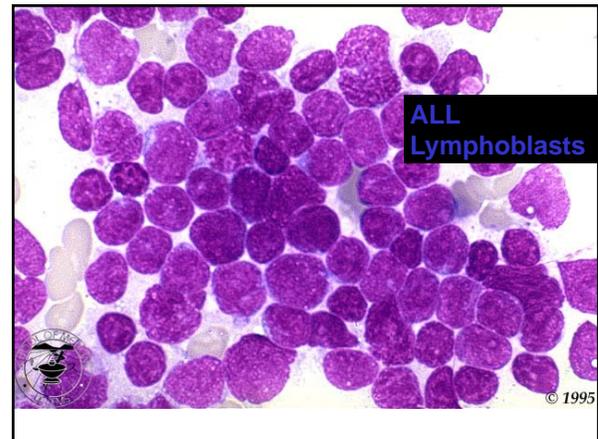
Symptoms

- Anemia
- Infections
- Hepatosplenomegaly
- Fever
- Night sweats
- Enlarged lymph nodes





- Treatment of chronic leukemias**
- Chronic lymphocytic leukemia: rituximab, steroids, fludarabine, bendamustine, ibrutinib.
 - Chronic myeloid leukemia: Tyrosine kinase inhibitors eg gleevec.
 - Polycythemia vera: phlebotomy and hydroxyurea.



- Treatment of acute leukemias**
- Induction chemotherapy: clear bone marrow of all leukemia.
 - Consolidation chemotherapy to keep the disease from coming back.
 - If the risk of relapse is very high, then one would consider an allo transplant as consolidation.

- Treatment of Acute Myeloid Leukemia**
- Typical induction chemotherapy is Idarubicin (or Daunorubicin) daily x 3 days and cytarabine by civi x 7 days (7+3).
 - Check cardiac function before anthracycline chemotherapy.
 - Cyatarabine can be associated with cerebellar and ocular toxicity but not typically at this dose.

Consolidation chemo for AML

- Consolidation often consists of high dose of cytarabine q 12 hrs x6.
- Pts needs cerebellar checks prior to each chemo dose and steroid eye drops because of risk of conjunctivitis.

Treatment of acute lymphoblastic lymphoma

- Induction chemotherapy is more complex than AML.
- Use multiple drug combinations.
- All patients with ALL need CNS prophylaxis eg IT methotrexate and/or IT cytarabine.



Presentation of lymphomas

- Painless lymphadenopathy
- Enlarged liver and spleen
- Recurrent fever and infections
- Night sweats
- weight loss

Leukemia vs Lymphoma

- Leukemia= increased WBC in blood and marrow
- Lymphoma usually starts in secondary lymphoid tissues, especially lymph nodes, so usually the patient is presented to the physician with lymphadenopathy

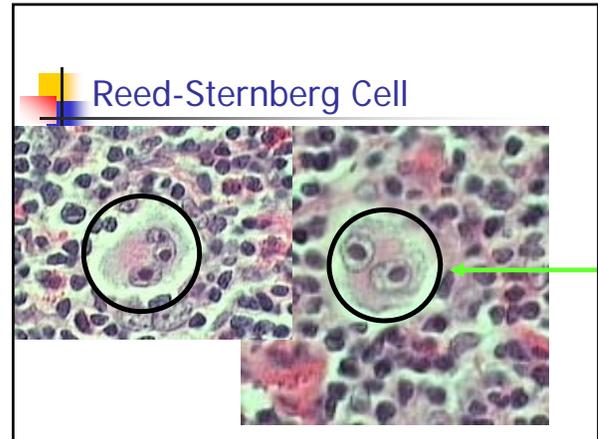
Lymphomas

- Could be classified as:
 - Hodgkin's Lymphoma
 - Non-Hodgkin's Lymphoma

Hodgkin's Lymphoma

- It is characterized by:
 - B cell in origin, and the presence of:

REED-STERNBERG CELLS

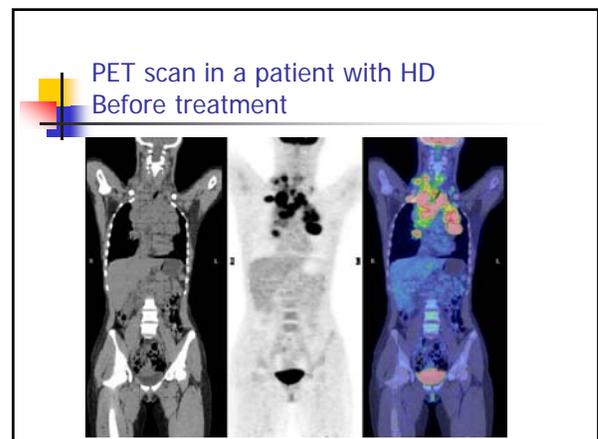


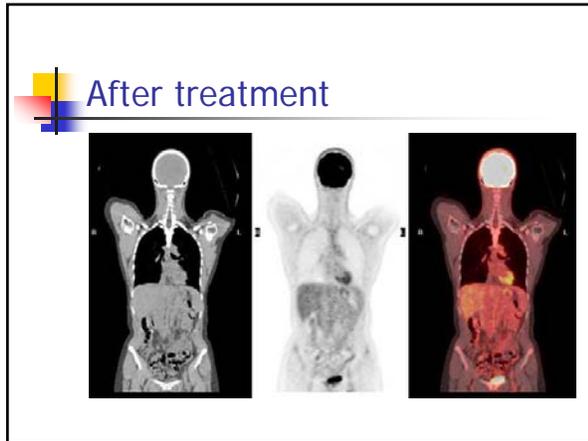
Non-Hodgkin's Lymphoma

The term is used to characterize all other lymphomas in which the description of Hodgkin's lymphoma is not applied, i.e. whenever there is **NO** REED-Sternberg cells, it could be due to B or T or NK cells involvement.

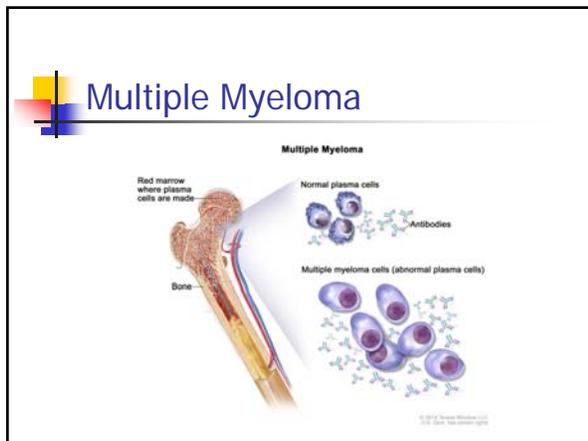
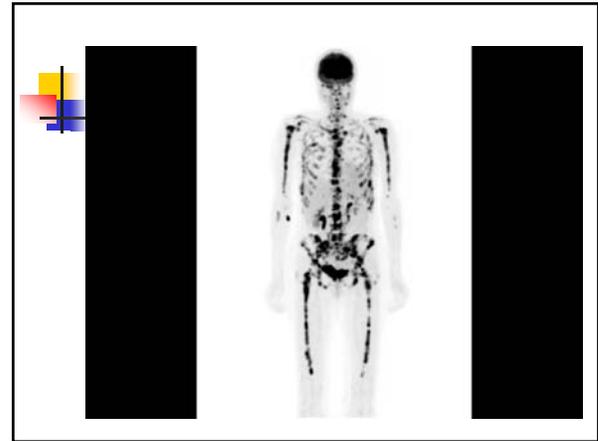
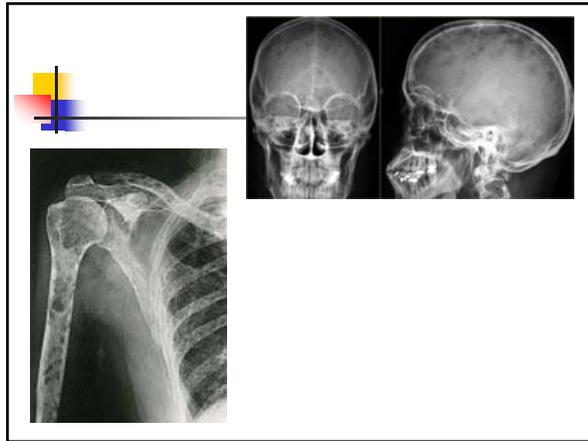
Comparison of Hodgkin Lymphoma and Non-Hodgkin Lymphoma

Feature	Hodgkin Lymphoma	Non-Hodgkin Lymphoma
Nodal involvement	Localized to a specific group of nodes	Usually disseminated among > 1 node group
Spread	Tends to spread in an orderly, contiguous fashion	Spreads noncontiguously
Extranodal involvement	Infrequent	Frequent
Stage at diagnosis	Usually early	Usually advanced
Histologic classification in children	Usually one with a favorable prognosis	Usually high grade





- ### Treatment of lymphoma
- Hodgkins disease: ABVD or BEACOPP.
 - Diffuse large B cell lymphoma: R-CHOP.
 - Relapsed lymphoma: R-ICE or R-DiHAP.

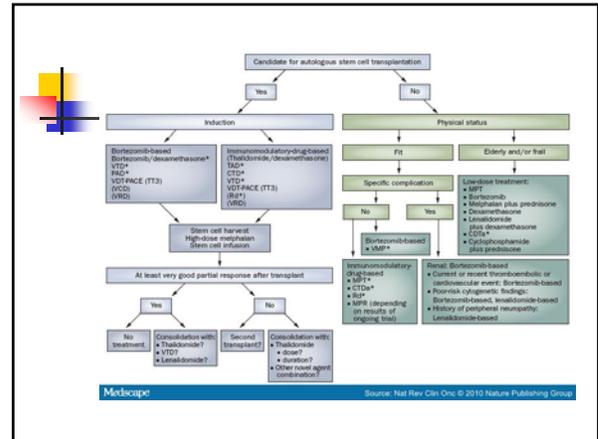


What is Active Myeloma?

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Presentation of multiple myeloma

- Acute renal failure.
- Proteinuria.
- Anemia.
- Bone pain. Pathological fractures.
- Hypercalcemia.
- Monoclonal protein in the blood or the urine.



Febrile Neutropenia

- Medical Emergency
- Neutropenia defined as ANC <0.5.
- Temp >100.5 deg F (oral).
- Panculture: blood, urine, sputum, etc.
- CXR.
- Start on abs asap. Gram negative coverage is imperative (eg cefepime). Gram positive coverage if patients have a central line (eg vancomycin).