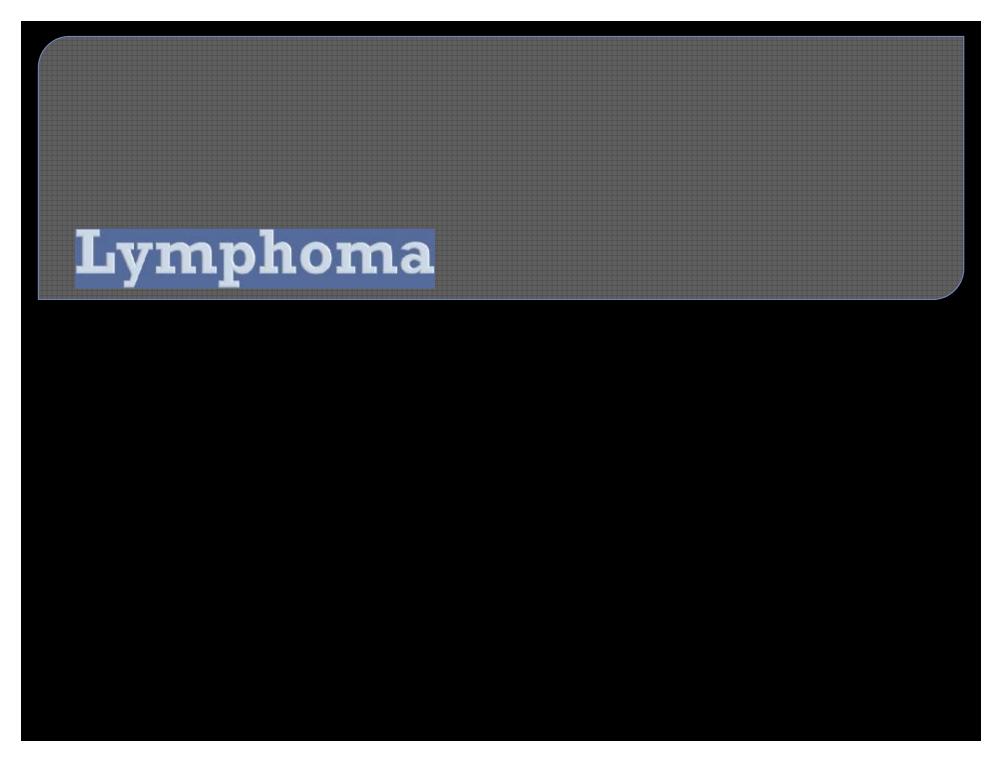
Case history

 30 years old male presented with history of fever, night sweating and weight loss.

 On examination: there are multiple bilateral inguinal and axillary lymphadenopathy Z Complete blood count:

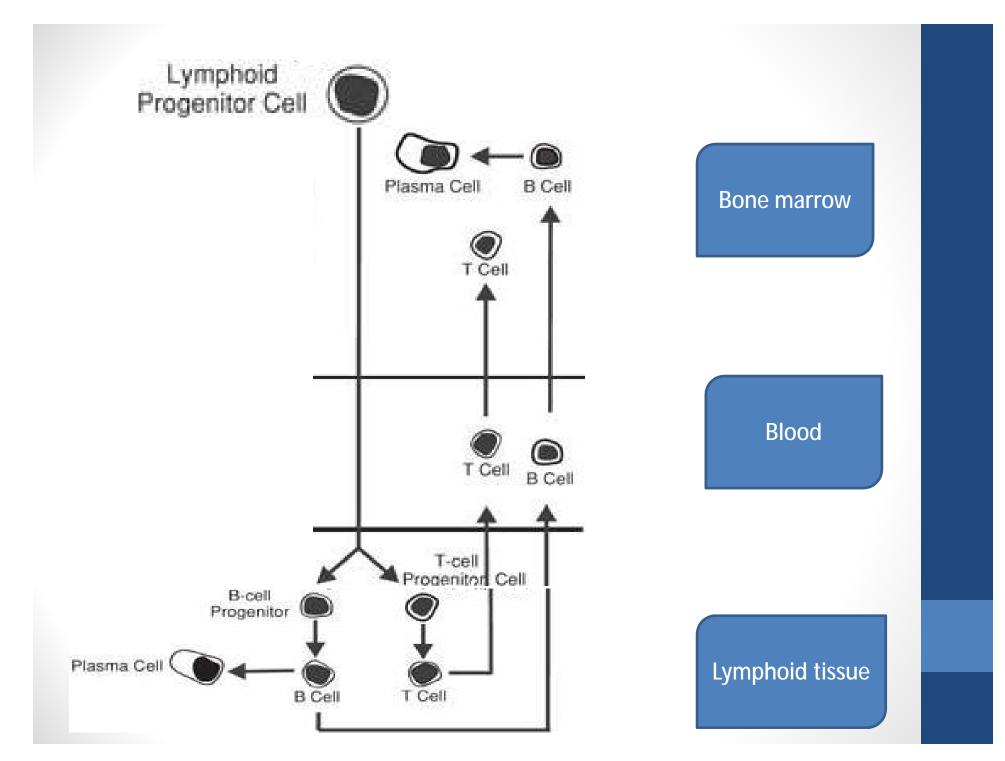
- \vee Hb= 9 gm/dl
- WBC count= 10000 cells/cmm
- Normal differential and no blast cells
- ≥ ESR= 90 mm/hr



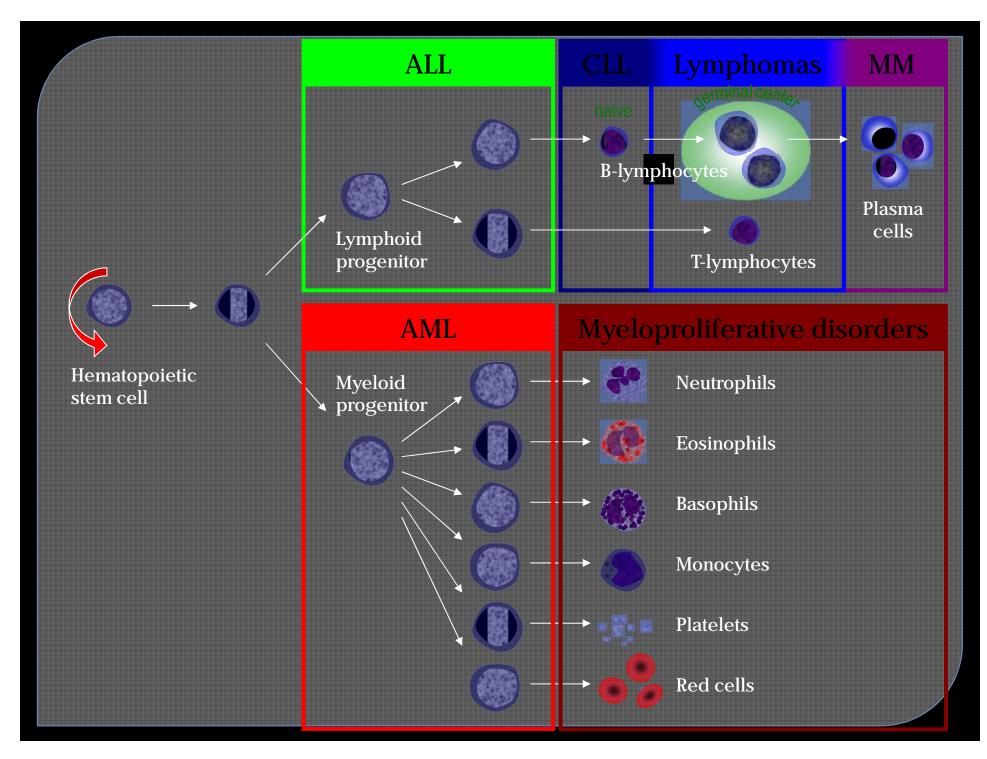
Introduction

These neoplasms arise from lymphoid tissues.

The majority are of B cell origin.



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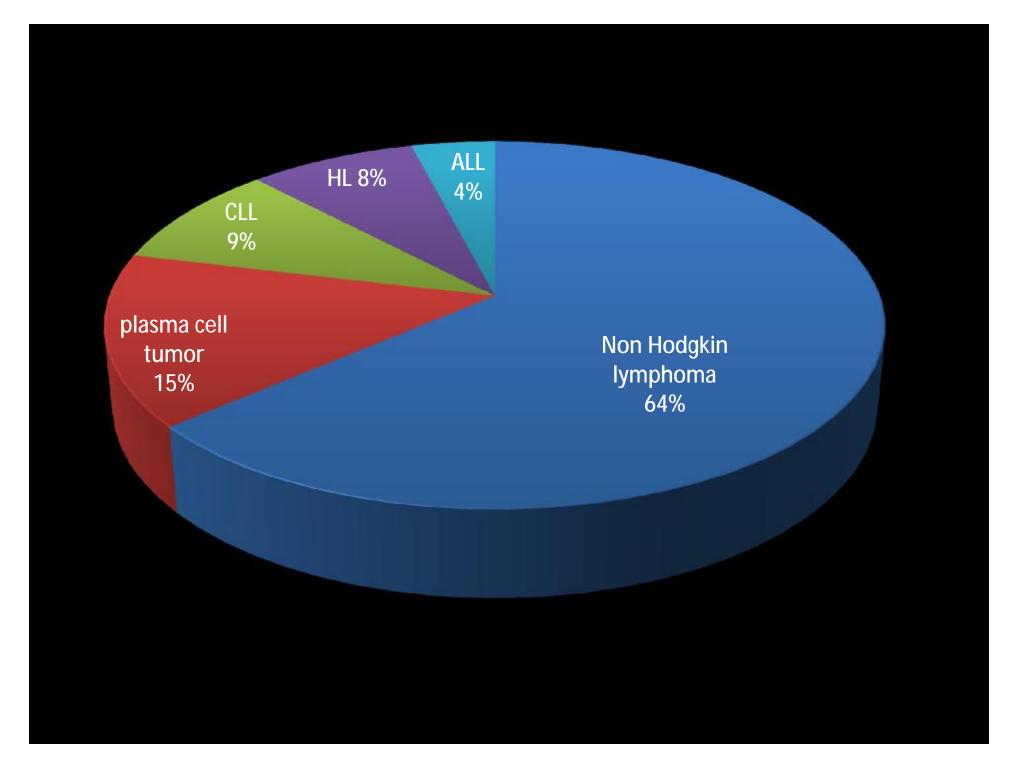
Histological Types

Hodgkin's lymphoma (HL)

Non-Hodgkin's lymphoma (NHL):

High-grade tumors divide rapidly.

Low-grade tumors divide slowly



Hodgkin's lymphoma

- HL is a neoplasm of lymphoid tissue in most cases derived from germinal center B cells.
- It has a characteristic neoplastic cell, the Reed-Sternberg cells.



Epidemiology

Male more common than female.

The disease has two peaks first peak at 20-35 years

and second at 50-70 years.

Clinical features

Asymptomatic

Constitutional symptoms called "B symptoms."

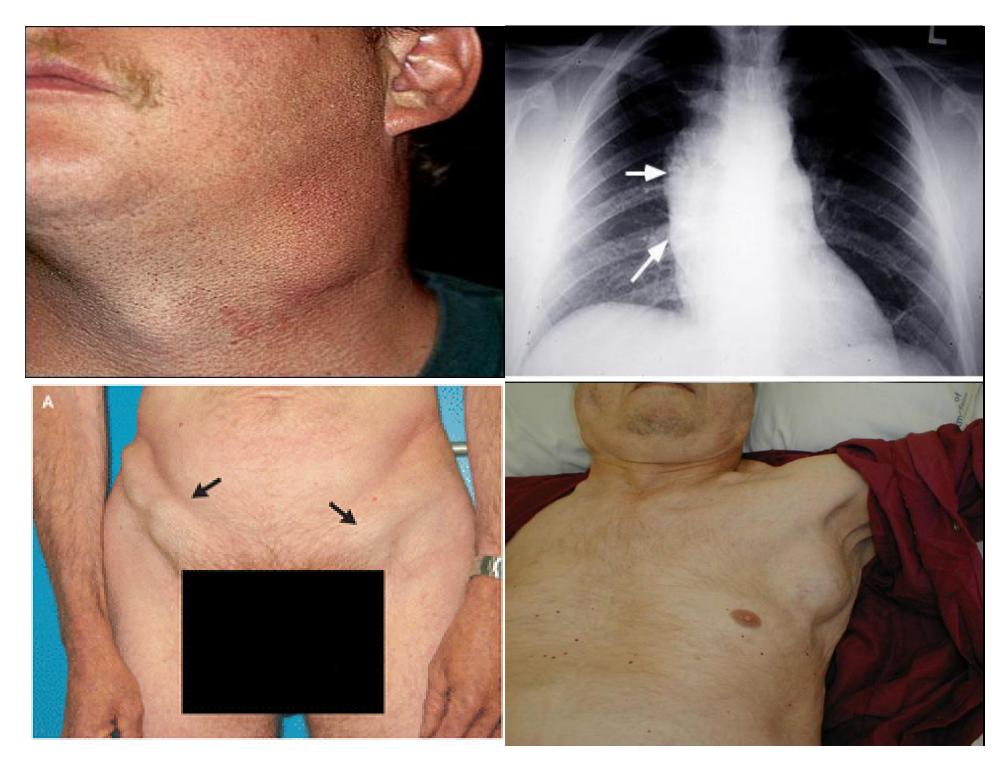
B- symptoms

Lymphadenopathy

u others



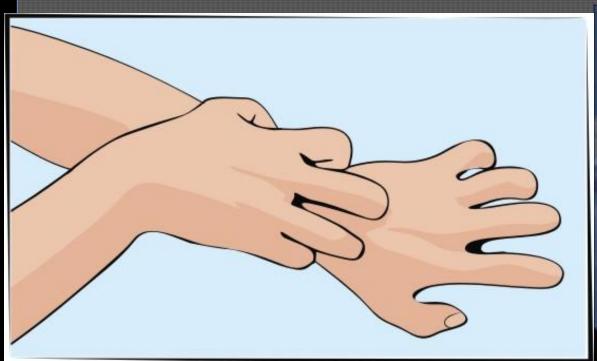
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- Intermittent fever (Pel-Ebstein fever)
- Patients may present with pruritus
- Extranodal disease, such as bone, or skin involvement, is rare

Splenomegaly and/or hepatomegaly may be present.





Investigations

- Confirm diagnosis: (lymph node biopsy)
- **z** Routine investigations:
 - CBC, LFT, RFT, & LDH
 - Echocardiography
- Staging:
 - CT scan of chest, abdomen and pelvis
 - PET- CT scan
 - Bone marrow biopsy

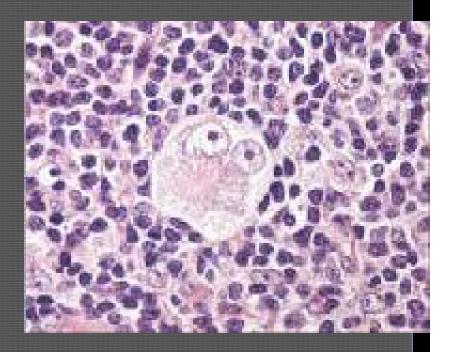
Diagnosis

Routine light microscopy (Reed

Sternberg cells) is almost

always augmented by

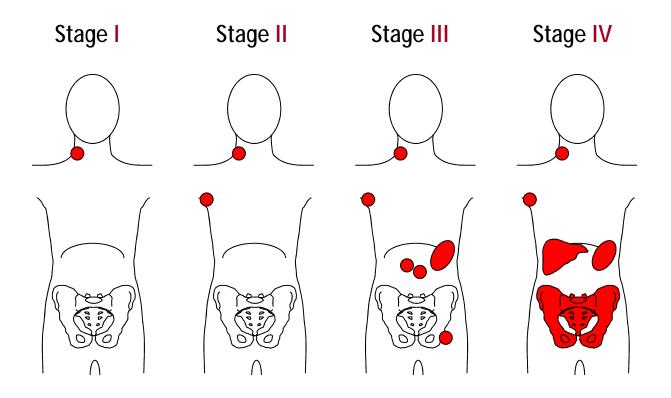
immunohistochemical staining.



Investigation

- CBC may be completely normal.
 - Normochromic, normocytic anemia,
 - Lymphopenia, eosinophilia or a neutrophilia may be present.
 - ESR may be raised.
- RFT & LFT.
- LDH: raised levels are an adverse prognostic factor.

Staging of lymphoma

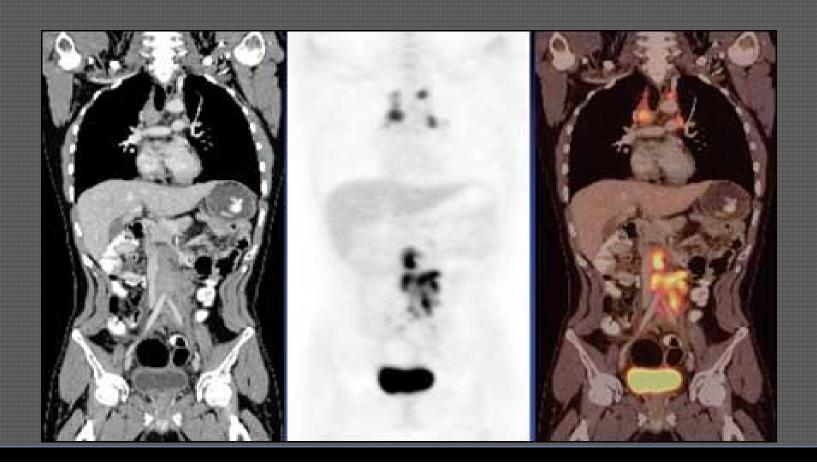


A: absence of B symptoms

B: fever, night sweats, weight loss

Ann Arbor staging classification for Hodgkin and non-Hodgkin lymphomas Stage I: Involvement of a single lymph node region (I) Involvement of two or more lymph node regions or lymphatic Stage II: structures on the same side of the diaphragm alone Stage III: Involvement of lymph node regions on both sides of the diaphragm Stage IV: Involvement of extranodal site(s) beyond that designated 'E' A: No symptoms B: Fever, drenching night sweats, weight loss > 10% in 6 months

- CT scan of chest and abdomen to permit staging.
- PET- CT scan.
- Bone marrow biopsy



Treatment

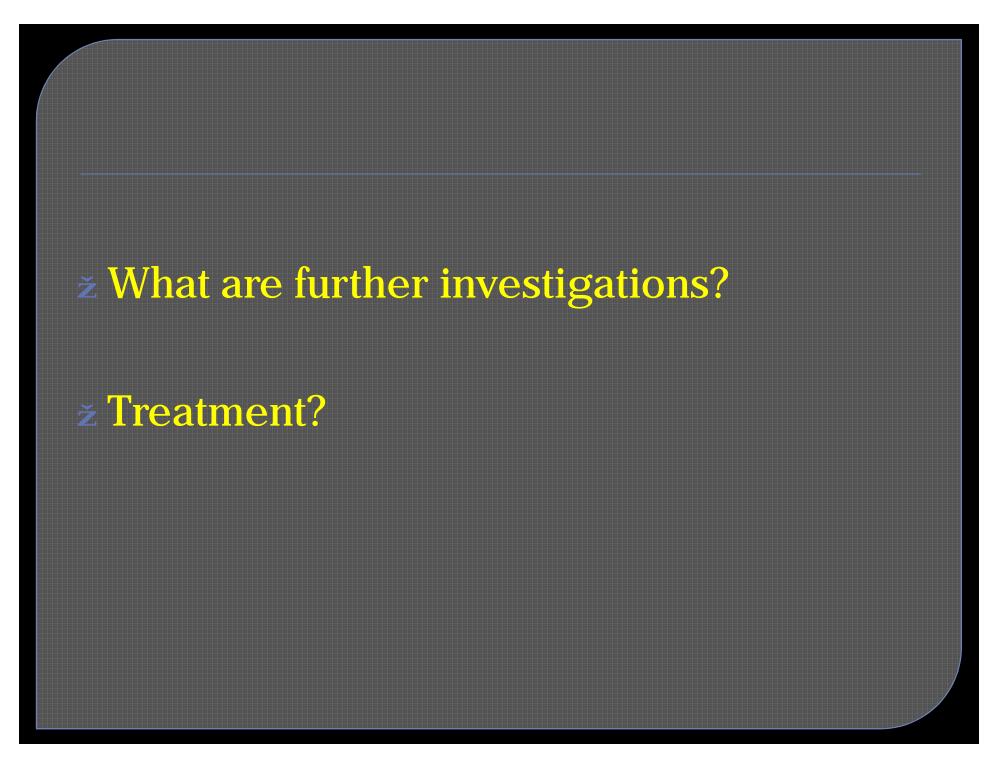
Characters	Treatment
Stage I or II	4 cycles of chemotherapy (ABVD) followed by radiotherapy (IF-RT).
Stage III or IV	6-8 cycles ABVD.
ABVD: (adriamycine/ bleomycin/ vinblastine/ dacarbazine)	

Case history

 30 years old male presented with history of fever, night sweating and weight loss.

 On examination: there are multiple bilateral inguinal and axillary lymphadenopathy





Non-Hodgkin lymphoma (NHL)

NHL a neoplasm of lymphoid tissue which consists of a diverse group of malignant solid tumors of the lymphoid tissues, it may be of B-cell (70%) or T-cell (30%) origin.

Clinically it is either high grade or low grade NHL.

Epidemiology

NHL account for about 4% of cases of new cancers

Slightly male more than female

The incidence rate increases dramatically with age, median age

of presentation 65-70 years.

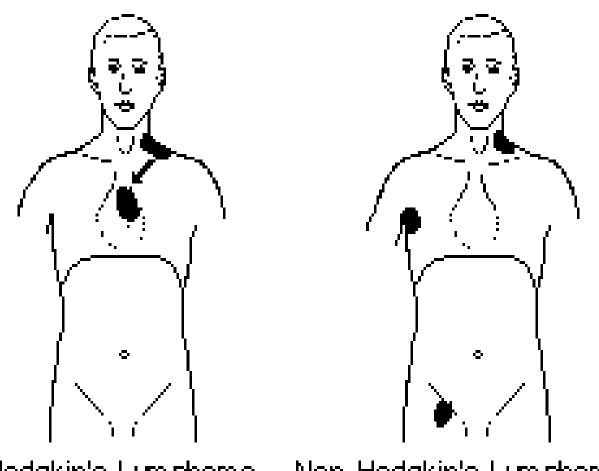
Clinical features

The most common presentation of NHL is <u>lymphadenopathy</u>.

Lymphadenopathy occurring in sites such as the cervical, axillary,

inguinal, mediastinum or retroperitoneum causes pressure

symptoms.



Hodgkin's Lymphoma Non-Hodgkin's Lymphoma

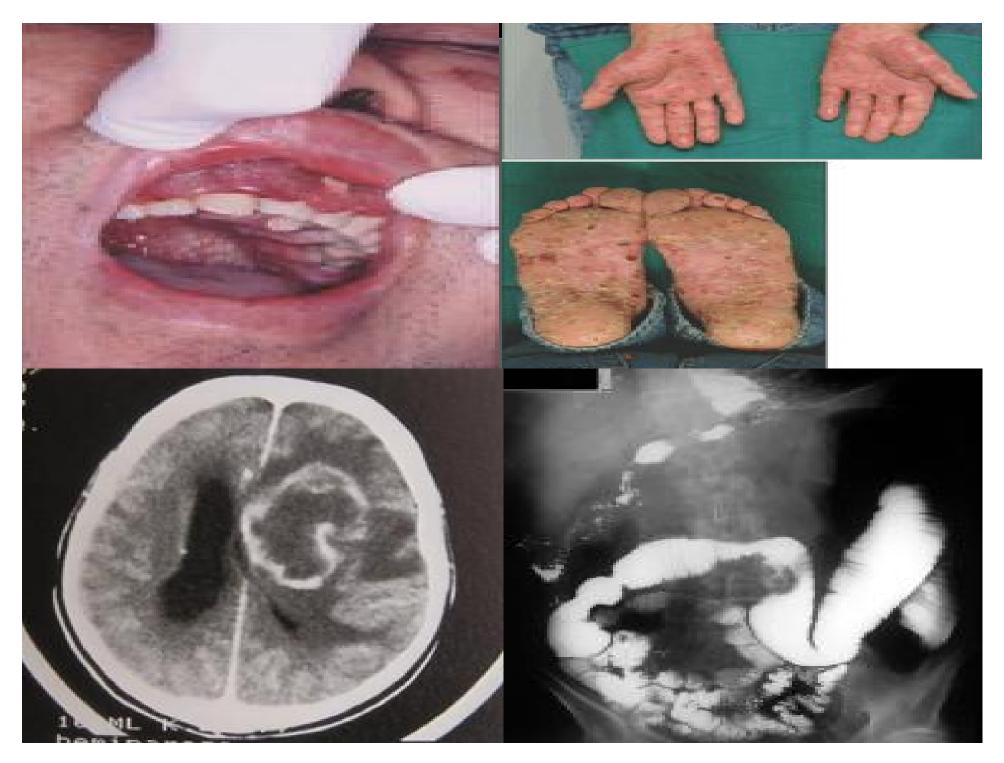
NHL are often associated with systemic

symptoms: fevers, night sweats, and

unexplained weight loss.

Other symptoms include fatigue, and pruritus.

 NHL can involve essentially any organ in the body: CNS: primary brain lymphoma, Shortness of breath with lymphomas in the lung, Bowel obstruction with small bowel lymphomas, Skin lesions with cutaneous lymphomas.



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High grade NHL present acutely with a rapidly growing mass,
and/or systemic B symptoms.

Indolent lymphomas are often insidious, presenting only with slow growing lymphadenopathy, hepatomegaly, splenomegaly, or cytopenias.

Investigation

Histological diagnosis from biopsy of a lymph node, bone

marrow or extranodal mass is essential, and confirmed by IHC

The investigation are as for HL but in addition the following

should be performed routine BM aspiration and trephine.

High-grade NHL

- All patients with high-grade NHL need treatment:
 - © Chemotherapy. The CHOP regimen (cyclophosphamide, doxorubicin, vincristine and prednisolone).
 - Monoclonal antibody therapy: rituximab.
 - Transplantation. Autologous SCT for relapsed cases.

Low grade NHL treatment

- Asymptomatic patients may not require therapy. Indications for treatment include:
 - Marked systemic symptoms,
 - Lymphadenopathy causing discomfort or disfigurement,
 - Bone marrow failure or
 - Compression syndromes.

Radiotherapy. This can be used for localized stage I disease. Chemotherapy: single or multidrug chemotherapy. Monoclonal antibody: The anti-CD20 antibody rituximab. Transplantation: autologous stem cell transplantation (SCT).