

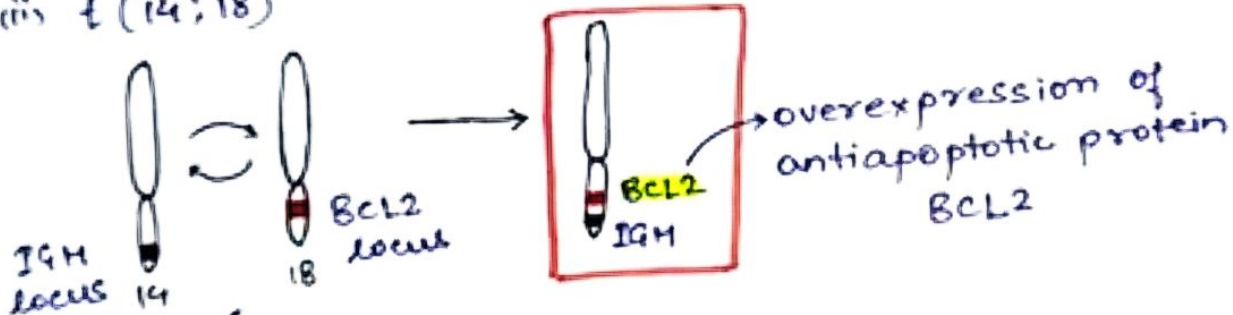
# DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL)

- most common<sup>A</sup> form of NHL
- median age of diagnosis = 60 yrs

## Pathogenesis:-

(i) Dysregulation of BCL<sup>A</sup>2  
↓  
Abnormal germinal center

(ii) t(14;18)



Similar translocation is seen in

Follicular lymphoma (30-50%)  $\xrightarrow{\text{transform}}$  DLBCL

# DIFFUSE LARGE B-CELL LYMPHOMA

→ WHO classification of Lymphoid Neoplasm

I) Precursor B-cell Neoplasm

↪ B-ALL (B-cell Acute Lymphoblastic Leukemia/Lymphoma)

II) Peripheral B-cell Neoplasm

↪ CLL/SLL (Chronic lymphocytic leukemia/small lymphocytic lymphoma)

→ B-cell prolymphocytic leukemia

→ Mantle cell lymphoma

→ Marginal zone lymphoma

→ Burkitt lymphoma

→ Diffuse large B-cell lymphoma

→ Hairy cell leukemia

↪ Follicular lymphoma

III) Precursor T-cell Neoplasm

↪ T-ALL (T-cell Acute Lymphoblastic leukemia/Lymphoma)

## Morphology:-

Diffuse Large B-cell lymphoma

Diffuse effacement of LN

Large lymphocyte  
(4x-5x of small lymphocyte)



## Immunophenotyping:-

- CD10+
- BCL6+
- Surface Ig+

## C/F:-

- Aggressive tumor

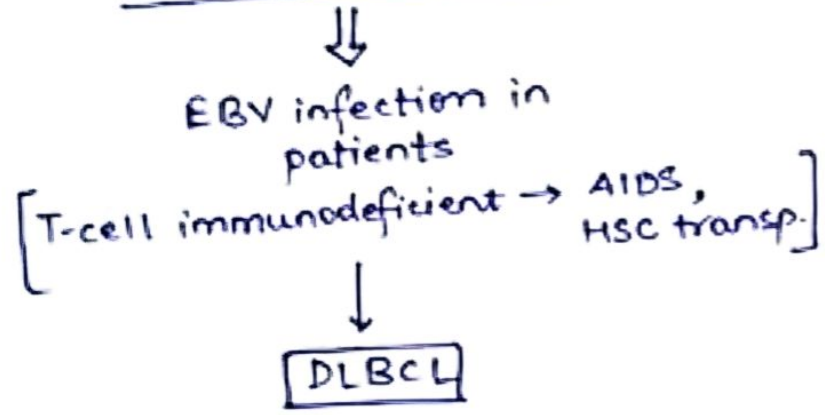
↓  
Rapidly enlarging @  
mass

Nodal site → Waldeyer's ring, tonsils, oropharyngeal lymphoid tissue, adenoids

↓  
Extranodal site → GIT, skin, bone, brain.

**Subtypes :-**

Immunodeficiency associated  
DLBCL



Primary effusion  
DLBCL

