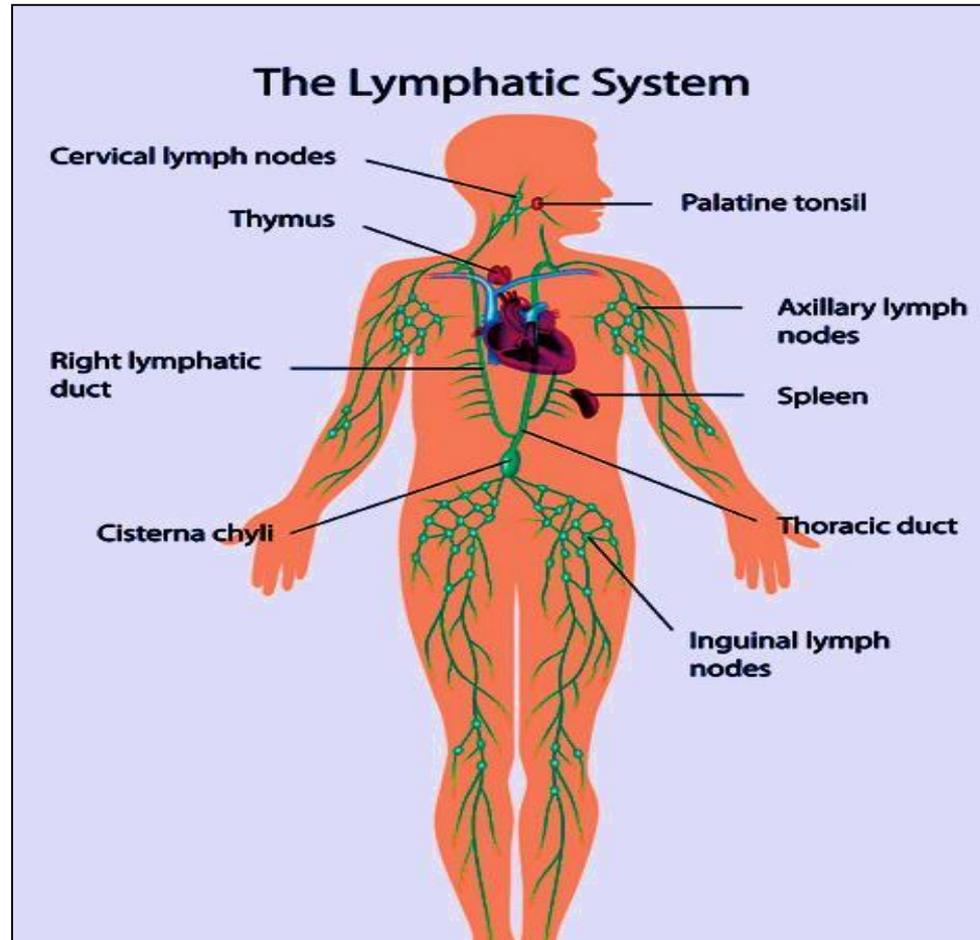
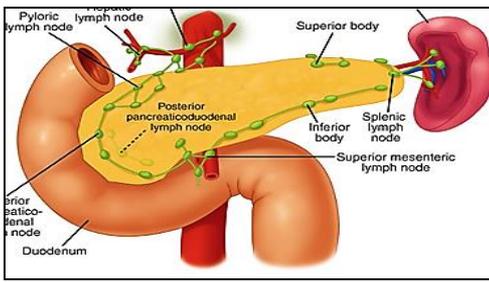


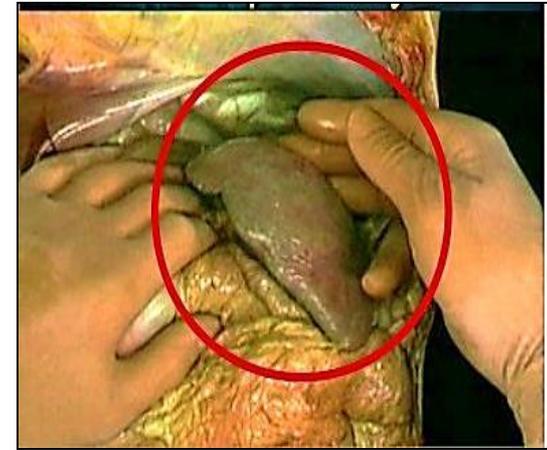
The lymphatic system (Part II)

Medical students / First Year





Spleen



- Largest single hemo-lymphatic organ
- Important blood filter. Is the site of destruction of aged RBCs & recycling of iron
- Immunological function through B & T cells (humoral & cell mediate immunity)
- A site of hematopoiesis in the fetus, and stores RBCs & platelets (blood reservoir in animals).

Spleen

A- Stroma

Capsule

Trabeculae
=septa

Reticular CT

B- Parenchyma

1-
White pulp

Lymphatic
nodules

2-
Red pulp

splenic
cords

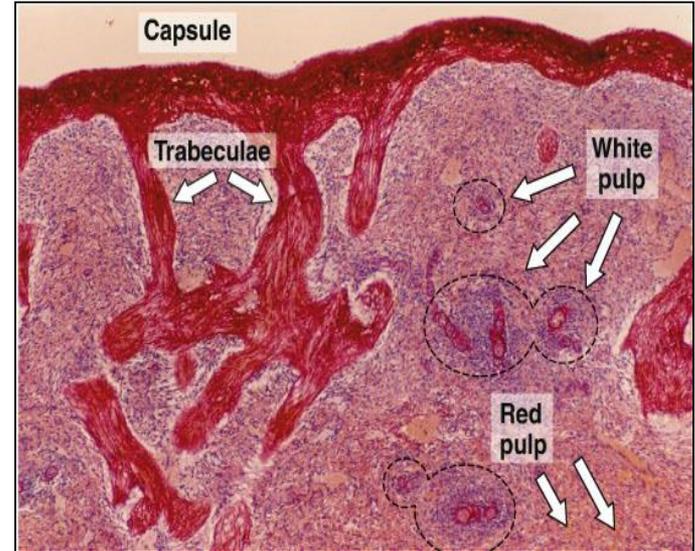
Blood
sinusoids



Structure of spleen

A-Stroma

1-Capsule: thick, rich in collagenous, elastic fibers & **smooth ms cells.**

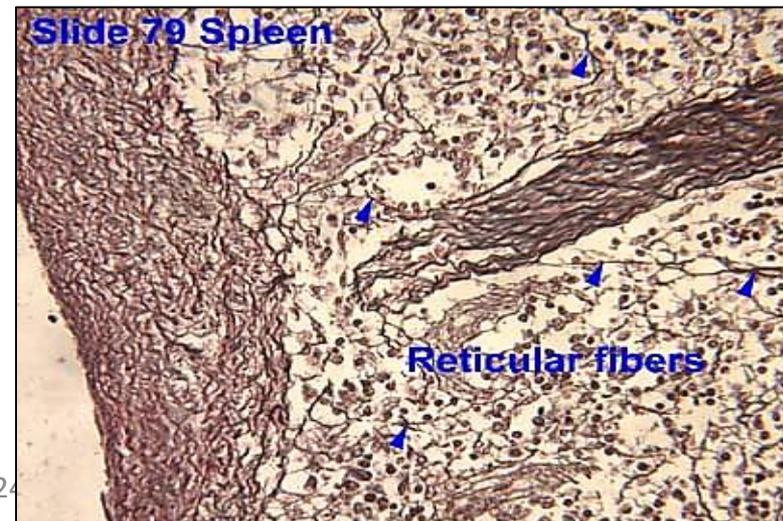


2-Trabecula: are short ones, extend from capsule.

divide the spleen into incomplete compartment, rich in elastic fibers & smooth ms. cells

3-Reticular CT:

reticular cells and fibers, form network



B- parenchyma

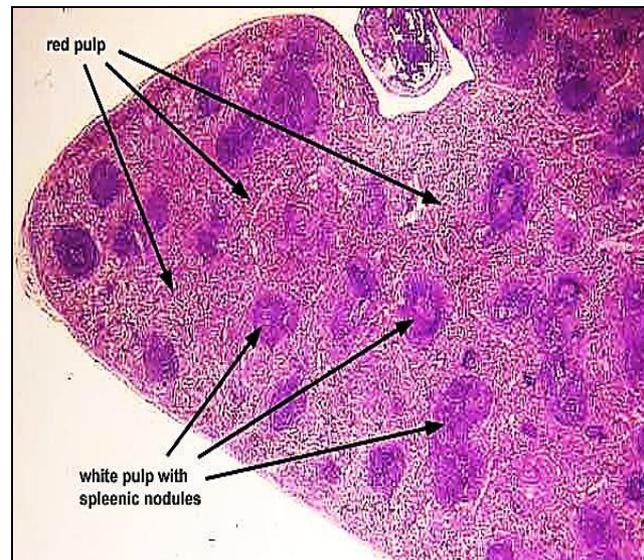
White pulp

Red pulp

Lymphoid
nodules

PALS

Peri-arteriolar
lymphatic sheath



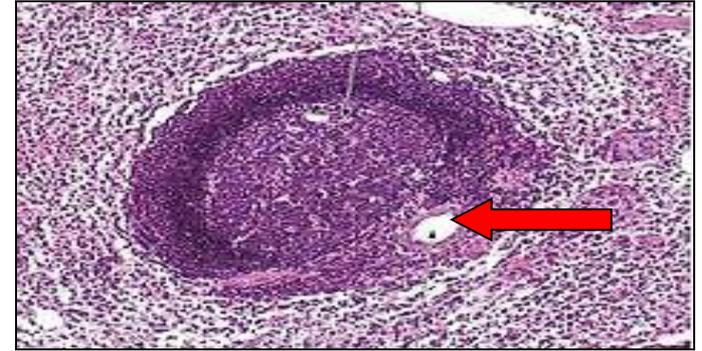
Blood
sinusoids

Splenic
cords

I- white pulp

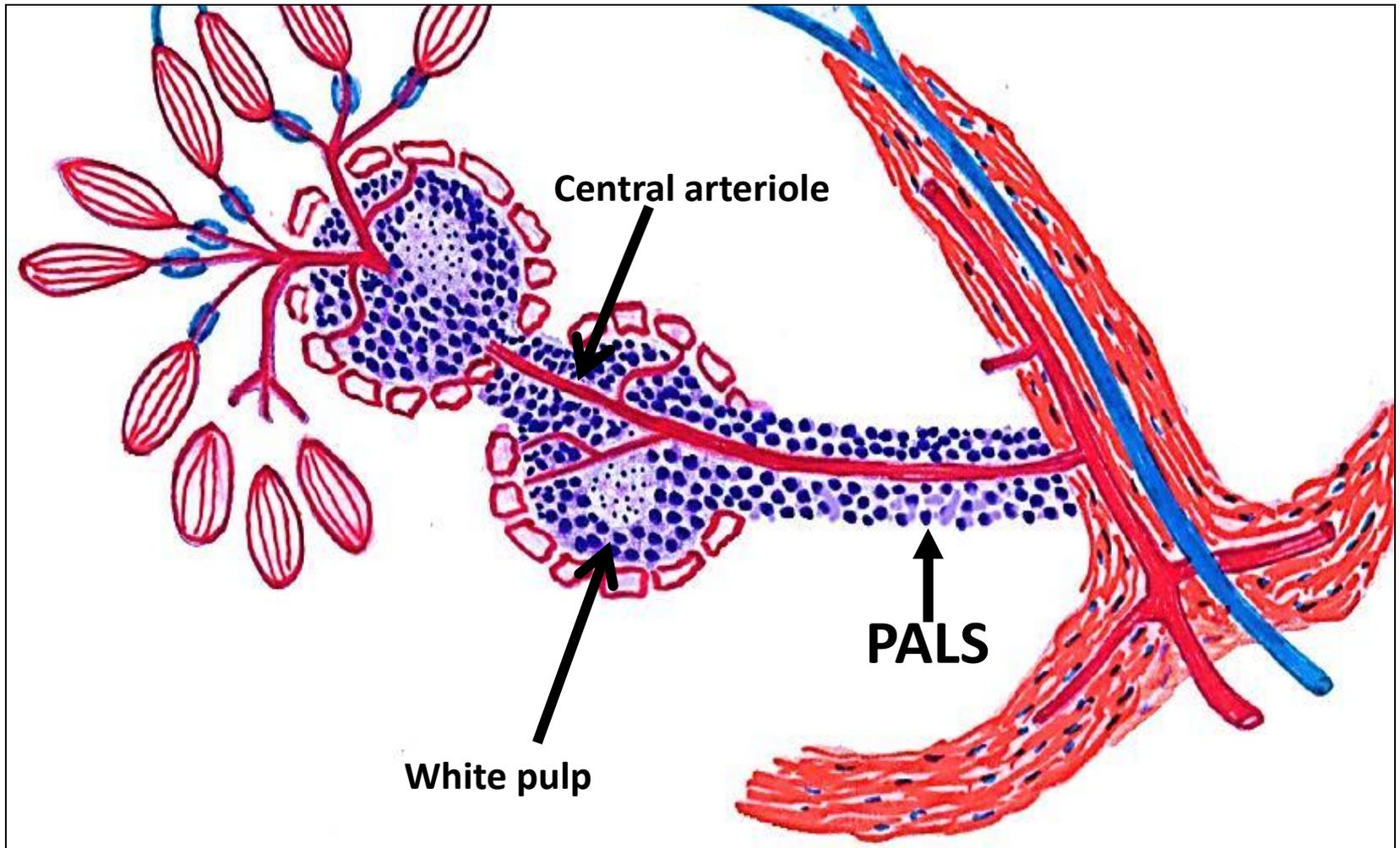
1- lymphatic nodules (Splenic Malpighian corpuscles):

aggregations of lymphocytes forming 1ry or 2ry nodules distributed throughout the parenchyma of the spleen



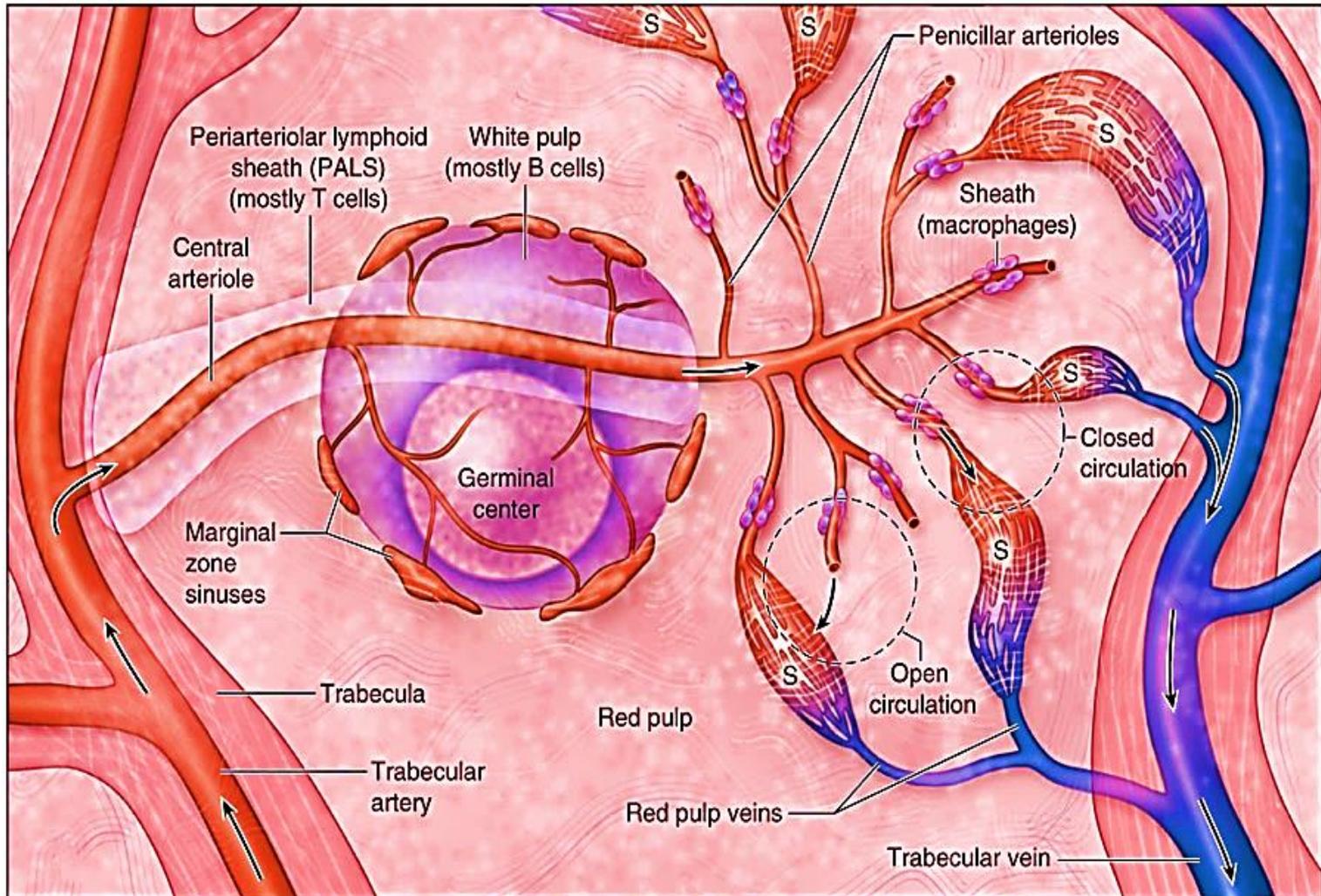
2- Periarteriolar lymphoid sheaths (PALS):

- mainly T lymphocytes encircle the central arteriole and called (Thymus dependent zone of spleen)
- Central arteriole runs at the **periphery** of the nodules (**eccentric**). They are branches of splenic artery which give numerous branches before leaving the white pulp to enter the red pulp.



The sketch shows the lay out of the blood supply of the spleen

- **Germinal center** : lightly stained, contain activated B cells, plasma cells & macrophages
- **Marginal zone** at the periphery of white pulp close to the red pulp has APCs & macrophages.

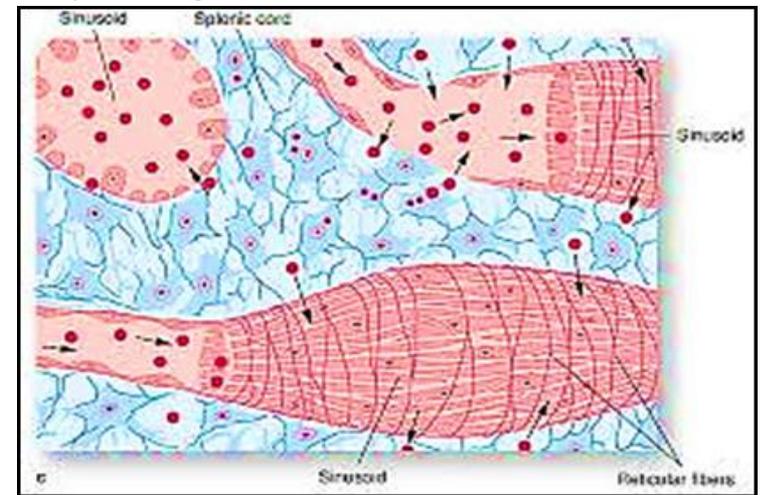
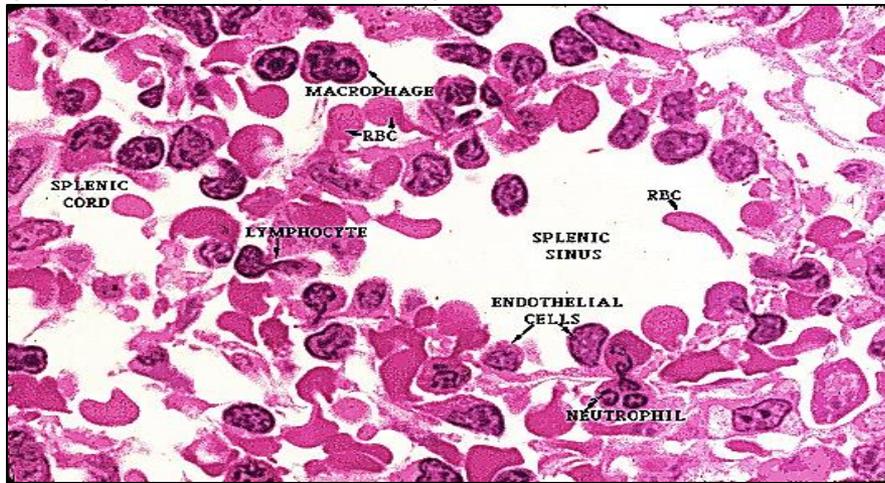


Organization of Cells in white pulp of spleen

II- Red pulp (79%)

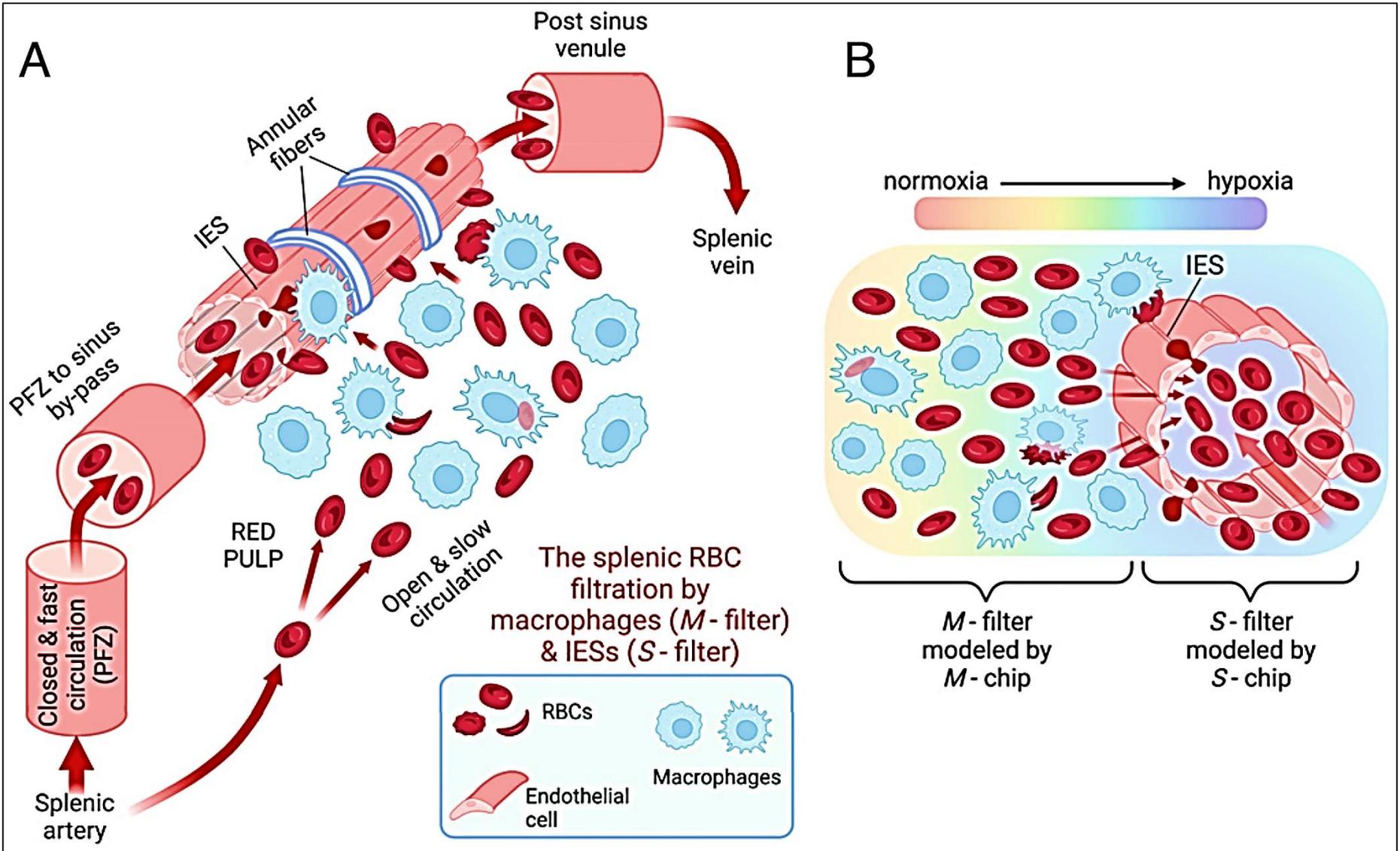
1-Splenic cords (Billroth cords):

- Network of reticular fibers between blood sinusoids to support the free cells found e.g. blood cells, T & B lymphocytes, plasma cells, macrophages



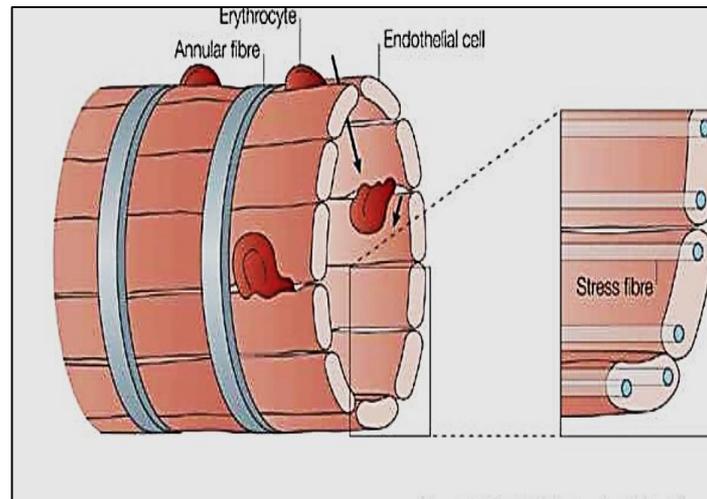
2-Blood sinusoids (venous sinuses):

- wide spaces lined with fenestrated endothelium called stave cells which filter the blood & surrounded with *Macrophages called Littoral cells*



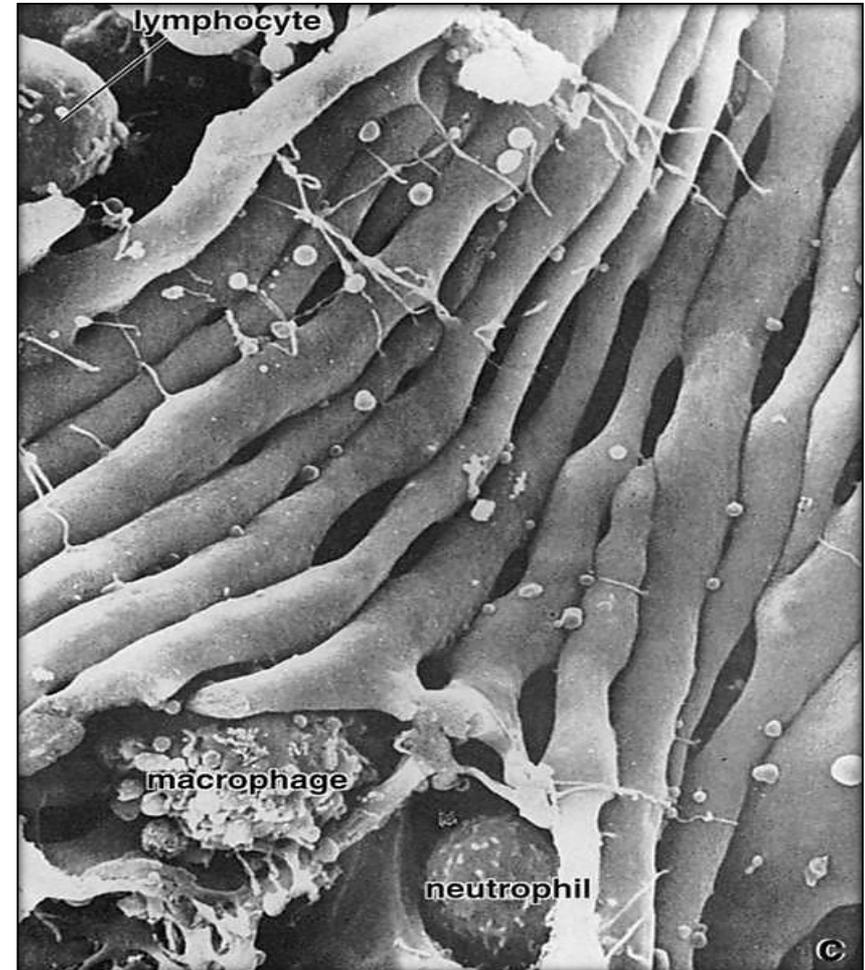
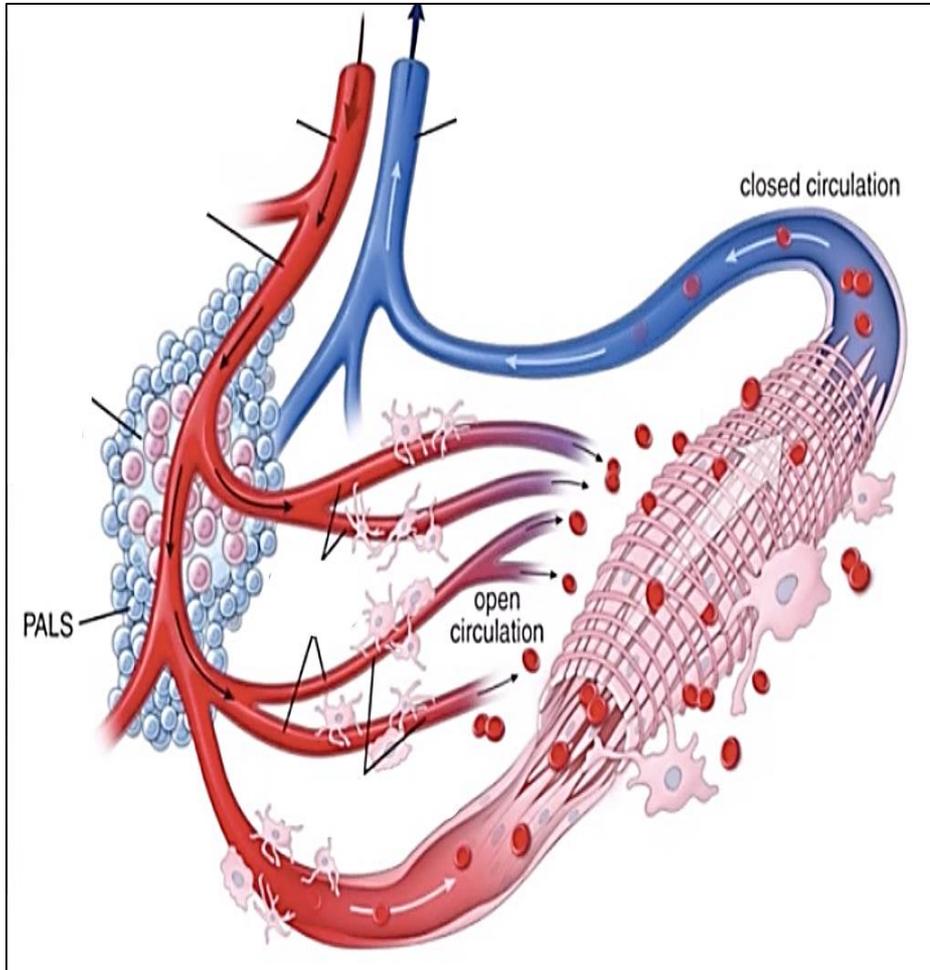
Destruction of red blood cells in the spleen

- **Stave cells**, unusual elongated endothelial cells(rod-like) oriented parallel to the sinusoidal blood flow
- These cells have discontinuous basement membrane which wrap the cells cross wise



- The gaps between the endothelial cells mechanically filter the blood cells.. Old or abnormal RBCs attempting to squeeze through the endothelial gaps become badly damaged and subsequently removed by macrophages

After about 120 days the erythrocytes undergo membrane changes & swell , signals for their engulfment by macrophages in the cords of the reticular between the venous sinuses



The lining of splenic sinusoids and the EM of Stave cells

Thymus

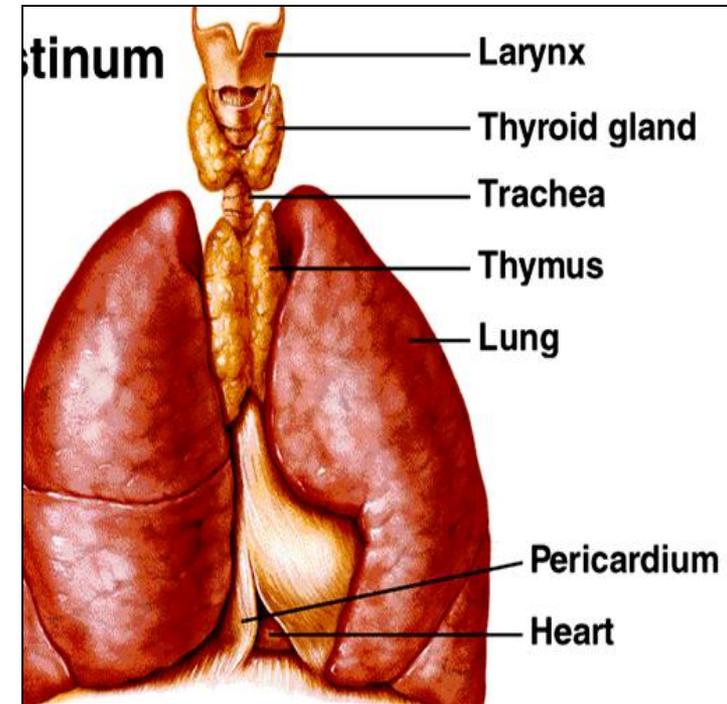
- is a **1ry** lymphatic organ with an endocrine function
- Location: behind the sternum in the mediastinum
- Single bi-lobed structure, highly lobulated organ

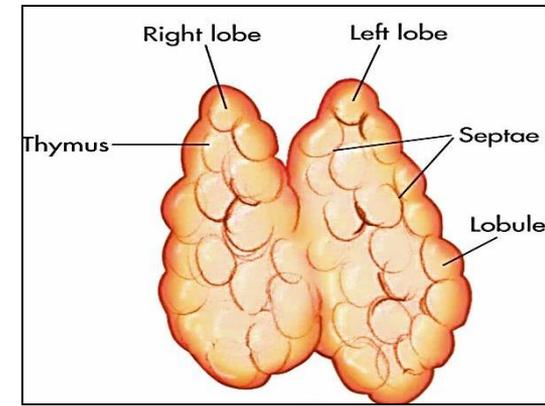
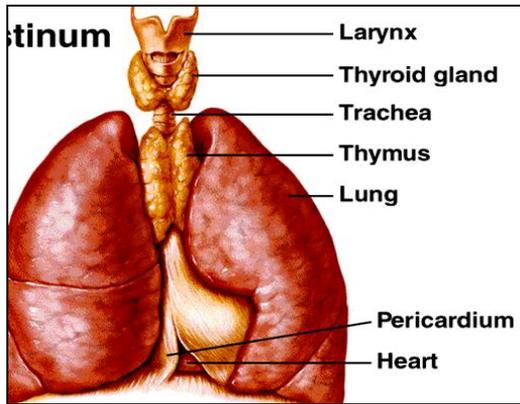
- Development:
 - Infant – ↑ in size
 - Puberty – maximum size
 - Adult – ↓ in size

- **Function**

Differentiation and maturation of T cells

Antigen-independent maturation





Thymus

A-Stroma

- 1-Capsule
- 2-Trabeculae

B-Parenchyma

- 1- Lymphocytes
- 2- Epithelial R cells

1.

Cortex

2.

Medulla

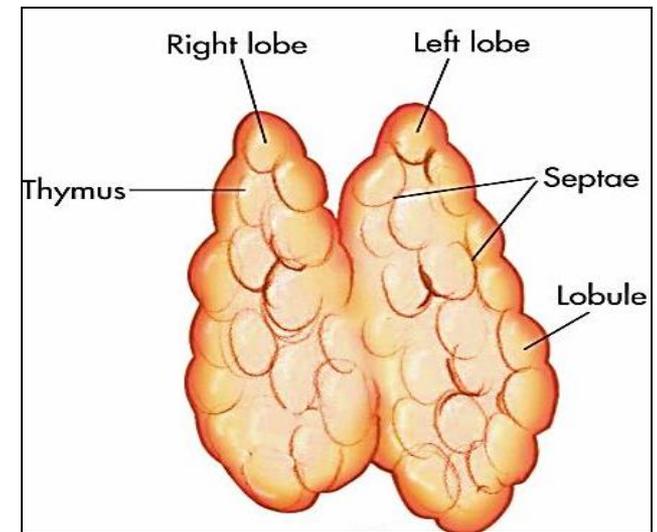
A- Stroma:

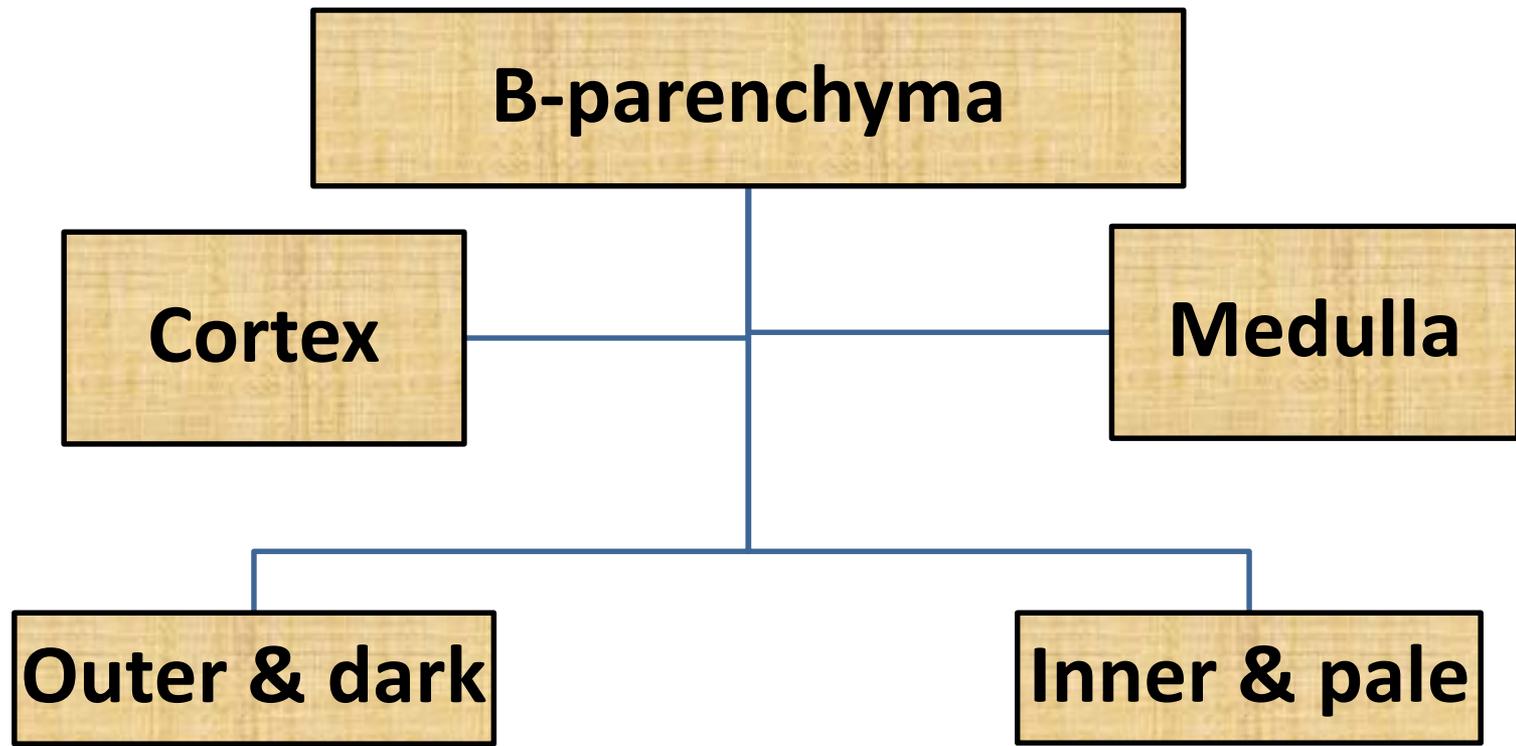
1- Capsule: loose CT

2- Trabeculae (septa):

Arise from capsule, penetrate its substance forming lobes, carry blood vessels. Each lobe is divided into incomplete lobules

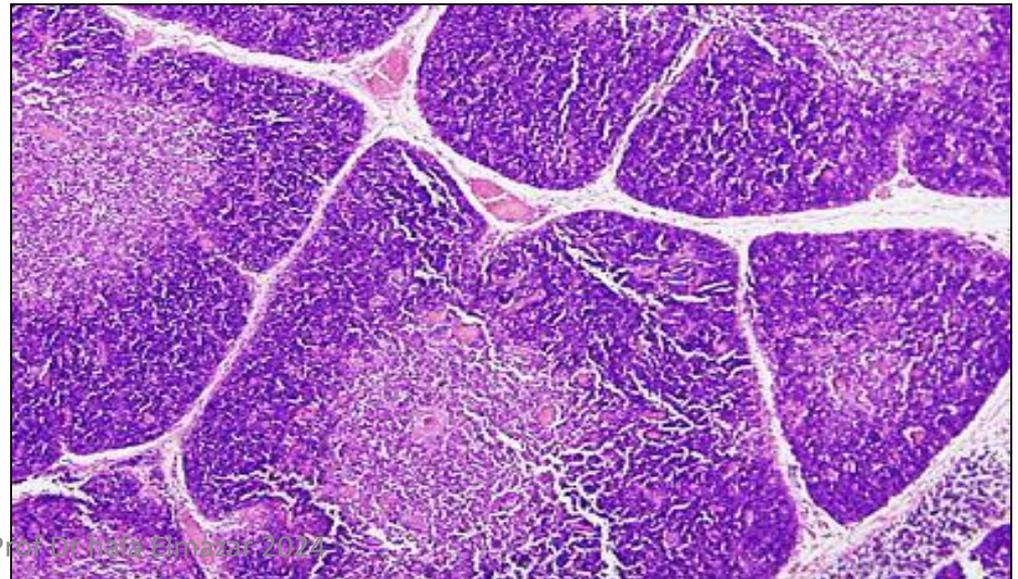
3- Thymus has no reticular fibers . Reticulum is formed by the processes of epithelial reticular cells





Both contain:

- 1- T. Lymphocytes.**
- 2- Epithelial reticular cells.**
- 3- Few macrophages.**
- 4- Blood capillaries**



1- Cortex:

- Peripheral dark-stained zone, where T cell maturation occur

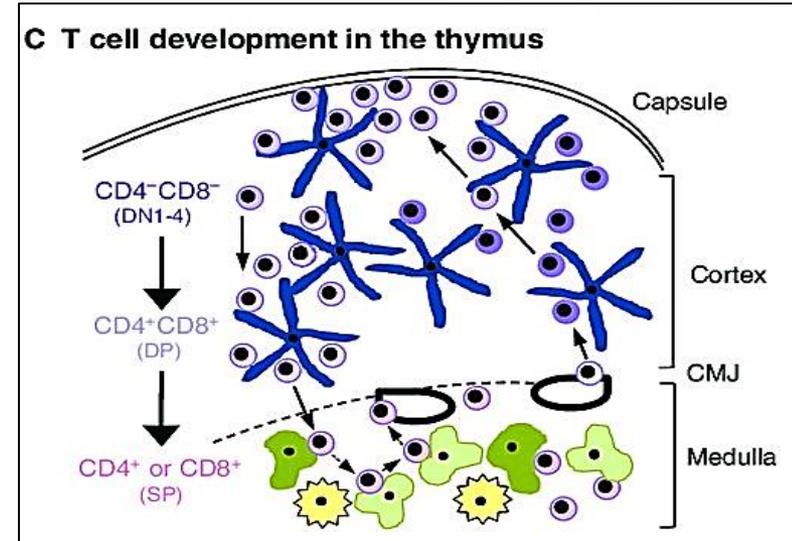
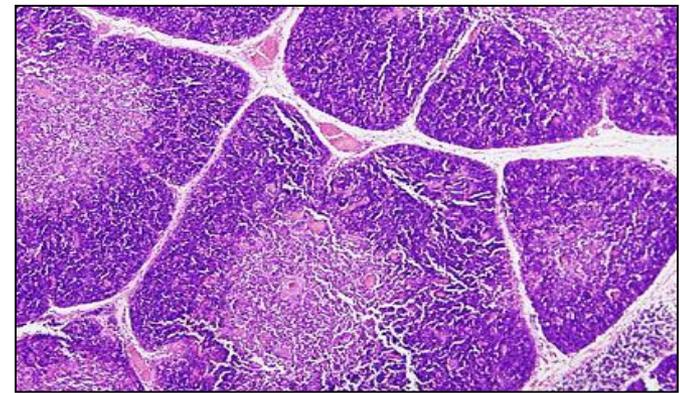
- Cortex contains thymocytes.

The hematopoietic precursors which migrated from bone marrow → thymus. Thymocytes is supported by a network of finely branched

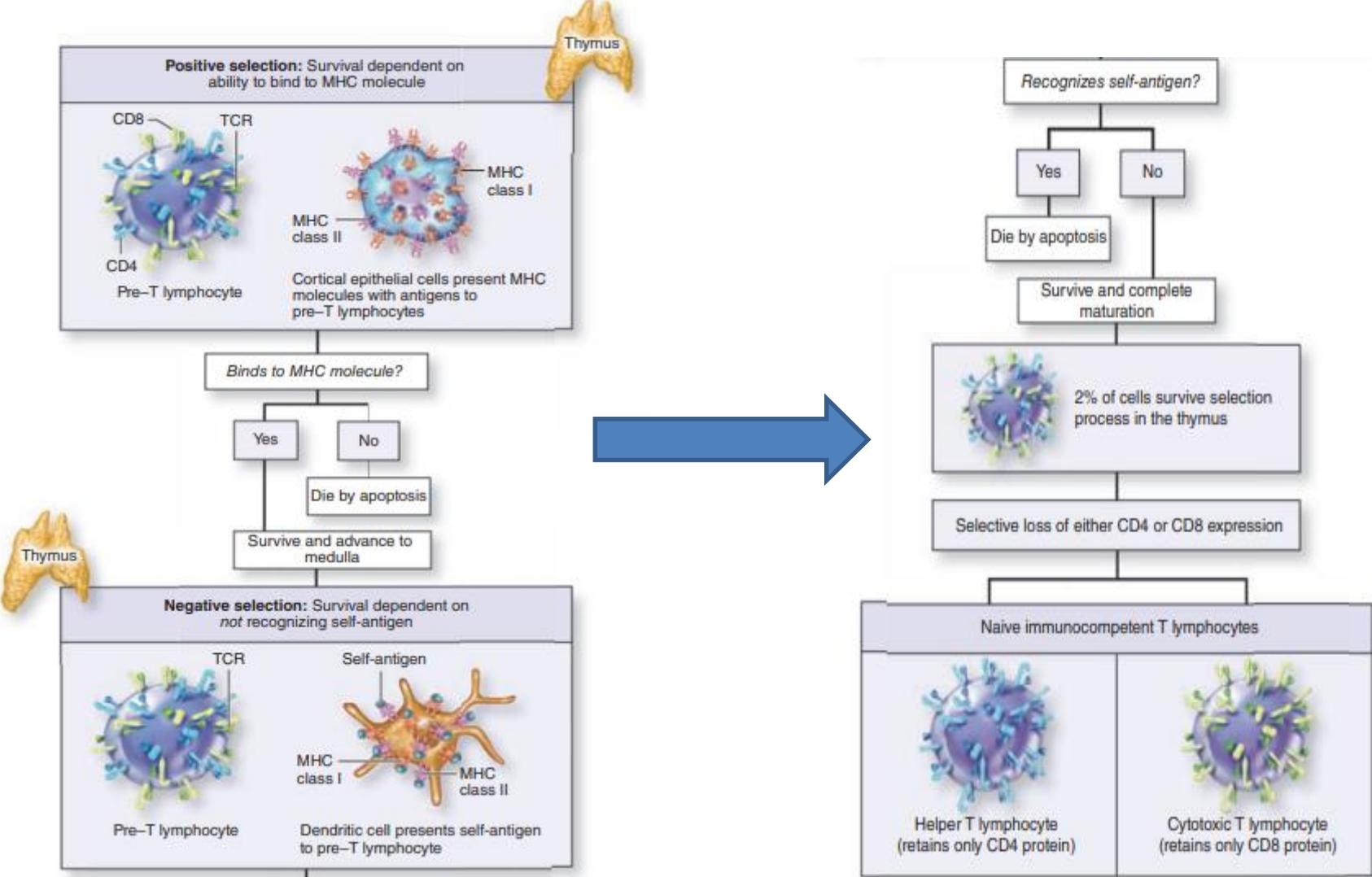
epithelial reticular cells

- Thymocytes are completely surrounded epithelial reticular cells

- The cortex is the site of **earliest events in thymocyte development**, where T cell receptor mature & positive selection take place
- Mature T lymphocytes** leave the **cortex** → **the medulla**.



Positive selection occurs in the cortex and allows survival only of T cells with functional TCRs that recognize MHC class I and class II molecules. **Negative selection** occurs in the medulla and allows survival only of T cells that do not tightly bind self-antigens presented on dendritic cells there

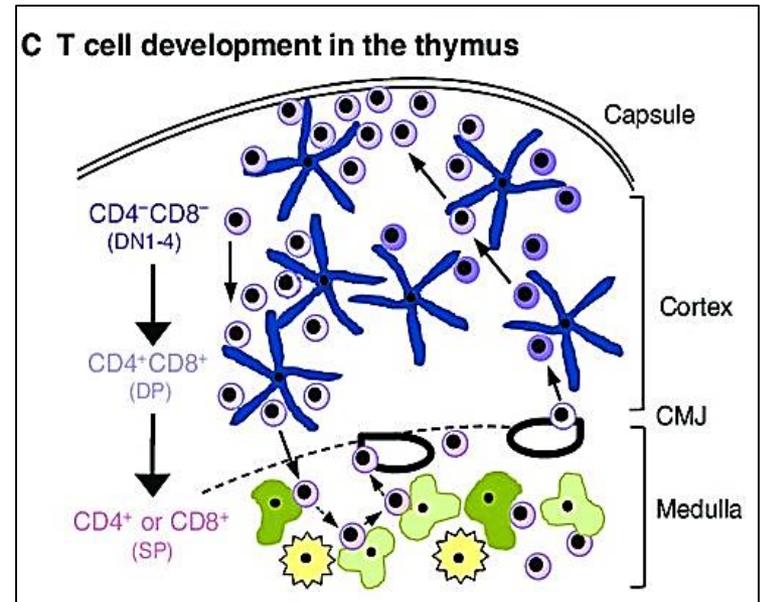


T- lymphocytes:

- Responsible for cell mediated immunity & also assist B lymphocytes in initiating the humoral response (**T- helper**)
- T- cells are several subtypes:
 - **Naïve (how they leave the thymus) (HEV) (simple cuboidal epith.)**
 - **Effector** (T- helper, T- cytotoxic , T- suppressor (T reg cells) & T- killer cells)
 - **Memory**
- T cells re-enter blood stream & travel to 2ry lymphatic organs (LN & spleen) where they settle in **thymus dependent zones**

Epithelial reticular cells (ERCs) :

- Branched, acidophilic cells e oval nuclei, their long processes contain tonofilaments (Keratin filaments)
- Also called thymic **nurse cells**
- They are connected together by desmosomes
- Do not produce reticular fibers.
- Found in both cortex & medulla (Cortical ERCs & medullary ERCs)
- Contain secretory granules which contain the thymic hormones



Functions of ERCs:

1- nursing cells for T cells during their differentiation

2- Secrete the thymic hormones

- Thymulin
- Thymopoietin
- Thymosins
- Thymic humoral factor

3- Share in the blood-thymus barrier

4- Antigen presenting cells for developing T lymphocytes

5- in medulla form Hassall's corpuscles

Blood- thymus barrier

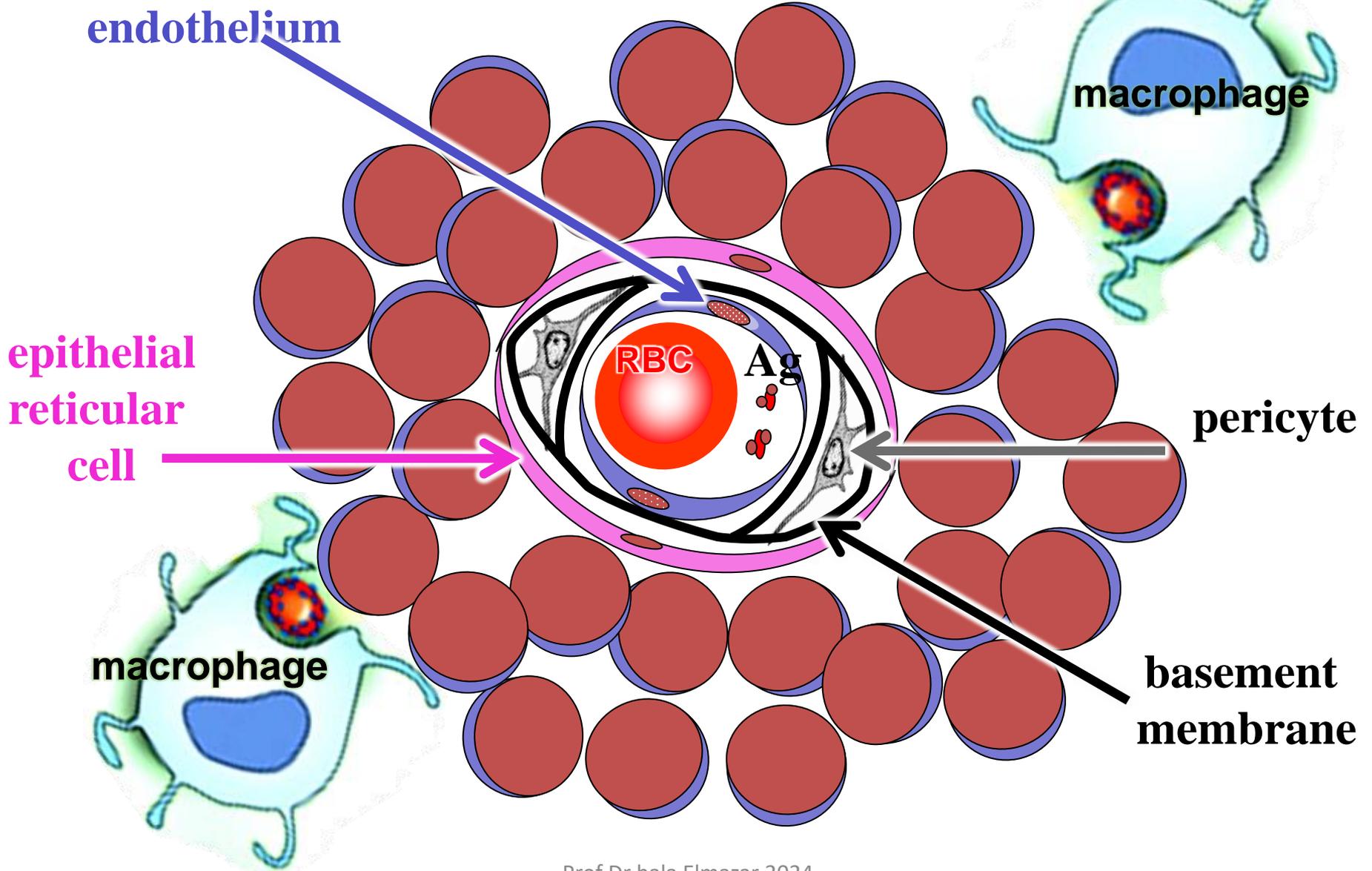
Barrier exists in the cortex only to separate the developing T-lymphocytes from antigens in blood

The barrier is formed by:

- 1- Continuous capillary endothelium
- 2- Pericytes
- 3- Continuous basal lamina around endothelium
- 4- Perivascular space contains macrophages to deal e any antigen escape
- 5- Complete layer of epithelial reticular cells around capillaries

The barrier allow immature T lymphocytes to multiply & differentiate free from foreign Ags before they migrate to medulla & leave thymus to blood

Blood thymic barrier



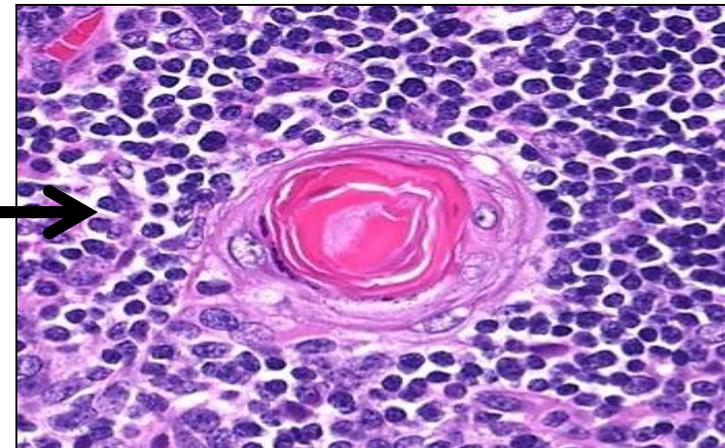
2-Medulla:

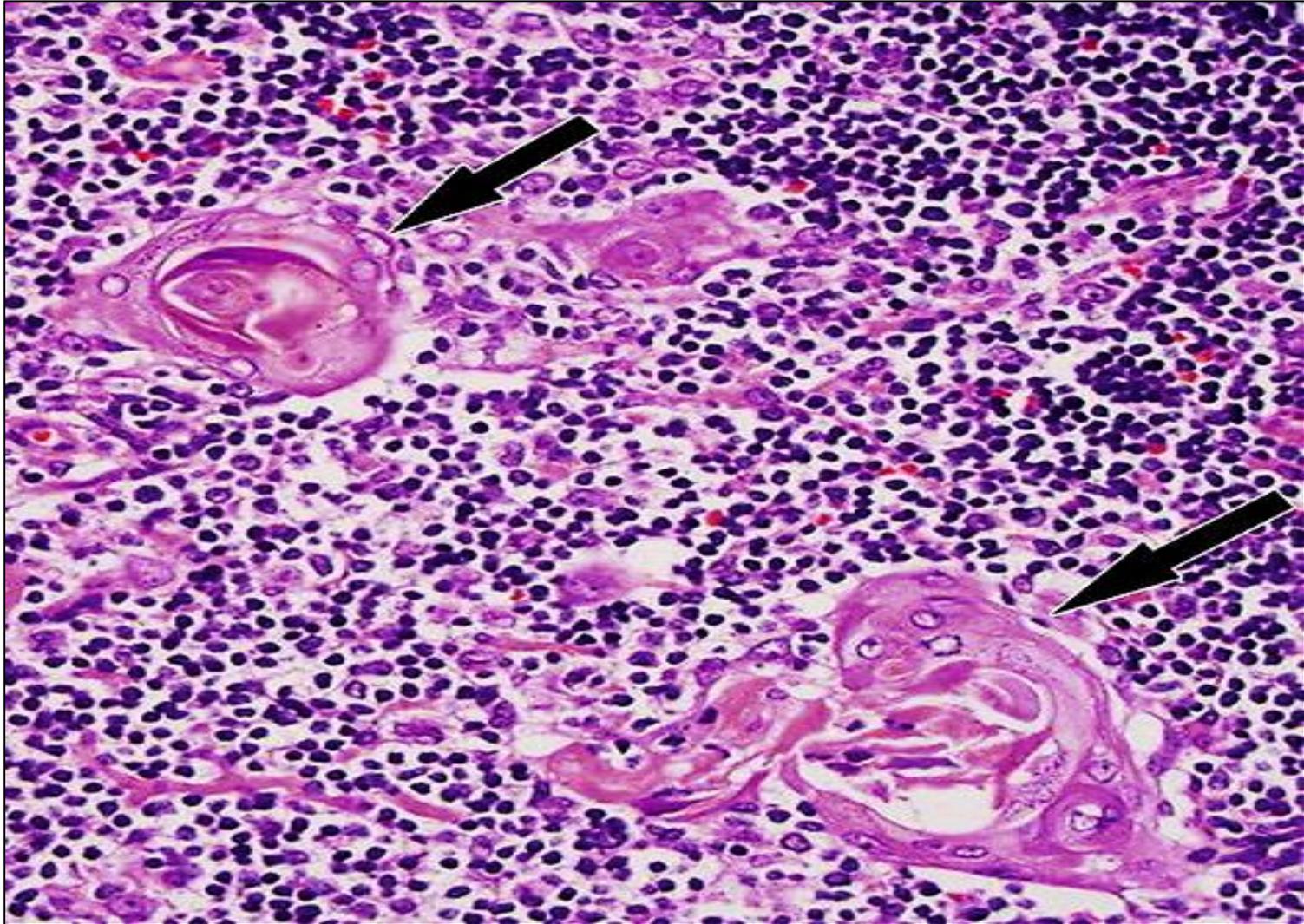
Contains fully differentiated T lymphocytes, which leave medulla through post capillary venules.

T cells will travel to 2ry lymphatic organs (LN & spleen) where they settle in thymus dependent zones

Contains **Hassall's corpuscles** are acidophilic structureless mass surrounded by concentric layers of epithelial reticular cells responsible for the release of cytokines that regulate dendritic activity.

Hassall's corpuscle





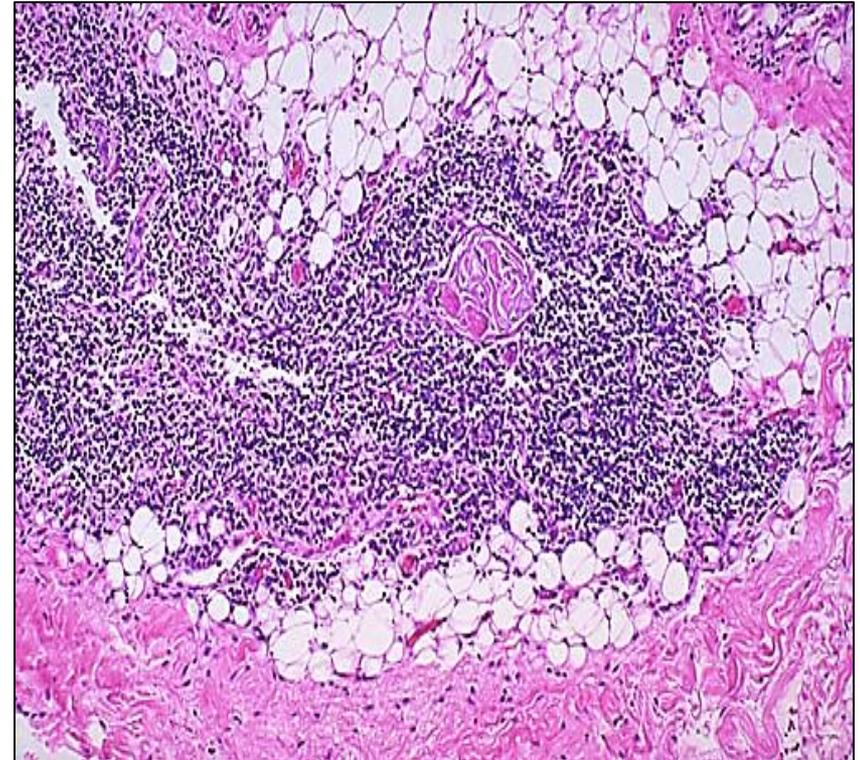
Thymus gland showing Hassall's corpuscles

Hassall's corpuscles provide developing thymocytes with paracrine and juxtacrine signals to ensure their proper functional maturation

Thymus gland of adult

Formed by:

- * Fibrous & adipose tissue.
- * Few lymphocytes, ↓ ER cells.
- * ↑ Hassall's corpuscles

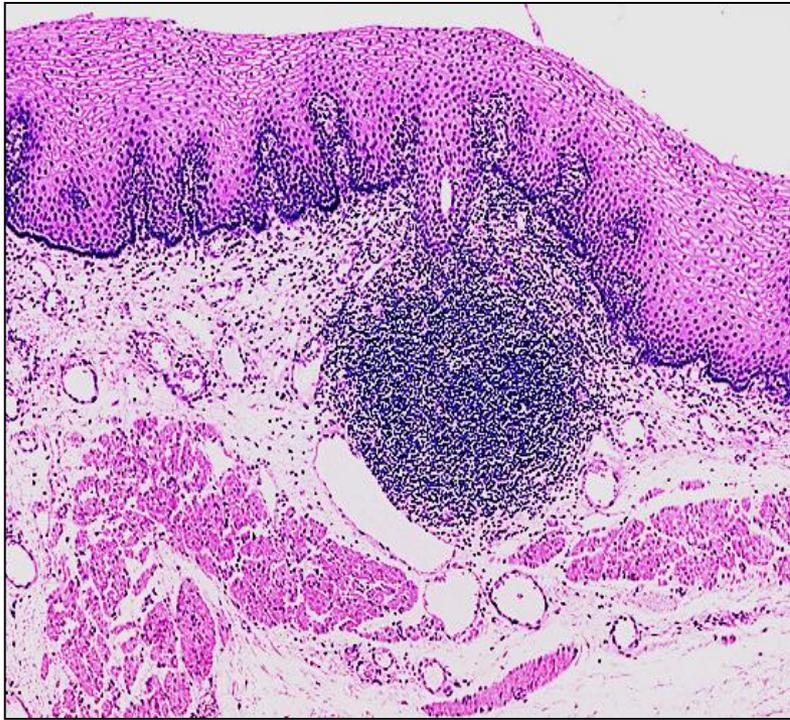


MALT- mucosa associated lymphoid tissue

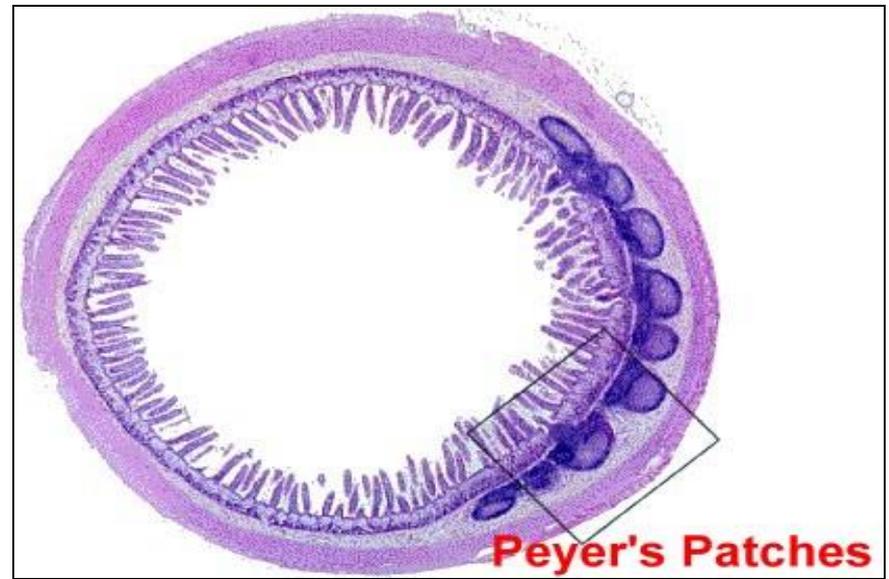
- Collective name for the cells of the immune system in the mucosa of respiratory , alimentary , urogenital tracts
- Function : is to augment the mechanical & chemical barriers of surface mucosal epithelium
- Distribution :
 - ✓ Tonsil
 - ✓ Bronchus : BALT
 - ✓ Gut: GALT

MALT Examples are:

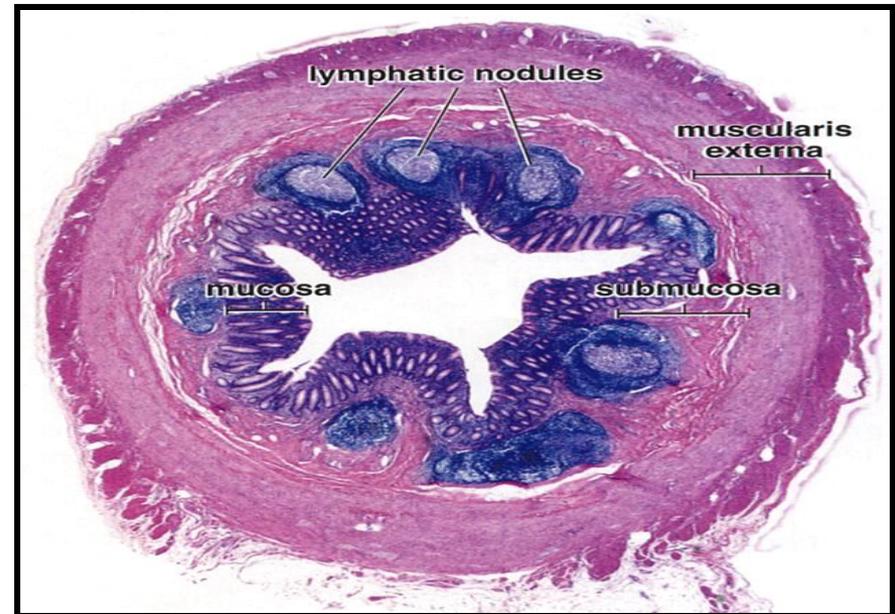
1. Payer's patches of ileum .
2. MALT of appendix.



MALT in wall of esophagus



MALT in ileum



MALT in appendix

Thank you

