The secondary lymphoid



The structure of spleen

From where	Stroma	Parenchyma
Divide into	1- Capsule	1- White pulp
	2- Trabeculae	2- Red pulp
	3- Reticular CT	
Explanation of the	1-Capsule:	1) White pulp:
divisions	1- INICK 2- rich in collagonous, elastic fibers	1- lymphatic nodules (spienic maipignian corpuscies): aggregations of lymphocytes
	.smooth ms cells.	forming 1ry or 2ry nodules distributed
	2-Trabecula: are	throughout the parenchyma of the spleen
	1- short ones	
	2- extend from capsule.	2- Central arterioles (follicular arterioles):
	3- divide the spleen into incomplete	 Run at the periphery of the nodules
	compartment,	(eccentric). They are branches of splenic
	4- rich in elastic fibers & smooth ms. cells	artery
	3-Reticular CT:	• which give numerous branches before leaving the white pulp to enter the red pulp
	1- reticular cells and fibers	leaving the white pulp to enter the red pulp.
	2- form background	
		2) Red pulp: (79%) 1. Splania corda (Billroth corda):
		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 (vongus sinusos):
		wide spaces lined e fenestrated
		endothelium called stave cells which filter
		the blood & surrounded e Macrophages
		called Littoral cells
Organization of	• Periarteriolar lymphoid sheaths (PALS): mainly	
Cells in white pulp	T lymphocytes encircle the central arteriole and	
of spleen:	called (Thymus dependent zone of spieen)	
-	 Germinal center: lightly stained, contain 	
	activated B cells, plasma cells & macrophages	
	(located between PALS and marginal zone)	
	• Marginal zone at the periphery of W. pulp close	
	to red pulp has APCs & macrophages.	



Thymus

The lymphatic organs

The primary lymphoid

From where	Thymus	
Def	- is a 1ry lymphatic organ e an endocrine function	
	- Single bi-lobed structure, highly lobulated organ	
Location	behind the sternum in the mediastinum	
Development	1) Infant – ↑ in size	
_	2) Puberty – maximum size	
	3) Adult – ↓ in size	
Function	Differentiation and maturation of T cells	
Structure of thymus	1- Stroma	
	2- Parenchyma	

The structure of thymus

From where	Stroma	Parenchyma
Divide into	1- Capsule	1- Lymphocytes
	2- Trabeculae	2- Epithelial R cells
Explanation of the	1- Capsule: loose CT	1- Cortex:
divisions	2- Trabeculae (septa): <mark>Arise</mark> from capsule, penetrate its substance forming lobes, carry blood vessels. Each lobe is divided into incomplete lobules	 Der: Peripheral bark-stained zone, where i cell maturation occur Cortex contains thymocytes. The hematopoietic precursors which migrated from bone marrow → thymus. Thymocytes supported by a network of finely branched epithelial reticular cells
	3- Thymus has no reticular fibers . Reticulum is formed by the processes of 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
	2-Meduli	
	- Contair	is fully differentiated T lymphocytes, which leave medulla through venules.
	- T cells depende	travel to 2ry lymphatic organs (LN & spleen) where they settle in thymus nt zones
Hassall's corpu	- Contair scle consist μm in dia	s Hassall's corpuscles (diagnostic feature), which vary in size from 25 to 200 meter & are acidophilic in reaction.
mass of deg reticular cells s concentric l epithelial retio	enerated urrounded e ayers of cular cells	



T- lymphocytes

From where	T- lymphocytes		
Fun		Responsible for cell mediated immunity & also assist B lymphocytes in initiating the humoral response (T- helper)	
T- cells are se	everal	1- Naïve	
subtypes		2- Memory	
	3- Effector (T-helper, T- cytotoxic, T- suppressor (T reg cells) & T-killer cells)		
The	• The St	• The Stem cells from bone marrow travel to the thymus to reside in the outer part of cortex, once there they are called thymocytes	
progression	These	These thymocytes have neither CD4 nor CD8 surface markers (double –ve T cells)	
of T- cell	• Within	Within outer cortex the thymocytes will proliferate & undergo genetic arrangement & express 2 cell markers: TCP (T cell recorder)	
development	 ✓ ICK (I cell receptor) ✓ Cluster differentiation: CD4⁺ & CD8⁺ (double positive T cells) 		
	• Double	e positive T cells that don't recognize self –MHC epitope offered to them by cortical ER cells are forced into apoptosis	
	• (MHC:	is a large section on vertebrates DNA contains all genes that code for cell surface proteins)	
	• Still in cells that	cortex: double +ve cells that in contact e ER cells that carry MHC I will stop expressing CD4 ⁺ marker & become single +ve T at express only CD8 ⁺ maker	
	• Double marker	+ve T cells contact ER cells carry MHC-II stop expressing CD8 ⁺ marker & become single +ve T cells that express only CD4 ⁺	
	• By doi cells	ng that the T cells acquired the Thymic education which was done under the influence of hormones secreted by epithelia R	
	Only 1	- 3% of Double +ve T cells will survive the selection process and will allow to enter the medulla	
	• The pr	evious process is called positive selection and take place in the thymus cortex	
	• The fin	al step in maturation of T cells occurs in the medulla	
	• The me strongly	edullary ER cells will do another test & present self-epitopes of MHC-I & MHC-II to the single +ve T cells & those who bind / are forced to apoptosis	
	 It has t takes pl 	to be weak reaction to the MHC - epitopes complex to prevent autoimmune response. This called negative selection and ace in the Thymic medulla	
	• T cells	re-enter blood stream & travel to 2ry lymphatic organs (LN & spleen) where they settle in thymus dependent zones	
	• Epithe ✓ ✓	lial Reticular cells secrete thymic hormones that stimulate: T cell differentiation Expression of surface markers	
	• CD4+ o	cells called helper T cells: indirectly can kill cells indicated as foreign.	
	• CD8+ o	cells called cytotoxic T cells are able directly to kill virus infected & tumor cells	
	• MHC I • MHC II	molecule is expressed on all nucleated cells Except RBCs molecule is expressed on antigen presenting cells: macrophages , dendritic cells …etc	

Epithelial reticular cells (ERCs)

From where	Epithelial reticular cells (ERCs)
Char	1- Branched
	2- acidophilic cells e oval nuclei
	3- their long processes contain tonofilaments
	4- Also called thymic nurse cells
	5- They are connected together by desmosomes
	6- Do not produce reticular fibers.
	7- Found in both cortex & medulla (Cortical ERCs & medullary ERCs)
	8- Contain secretory granules which contain the thymic hormones
Fun	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- thymes barrier

From where	Blood- thymus barrier
Located	Barrier exists in the cortex only
Fun	 to separate the developing T-lymphocytes from antigens in blood
	 The barrier allow immature T lymphocytes to multiply & differentiate free from foreign Ags before they migrate to medulla & leave thymus to blood
formed by	 1-continuous capillary endothelium 2- pericytes 3-thick, continuous basal lamina around endothelium 4- perivacular space contains macrophages to deal e any antigen escape 5- complete layer of epithelial reticular cells around capillaries

Thymes gland of adult

From where	Thymus gland of adult
Formed by	 * Fibrous & adipose tissue. * Few lymphocytes, ↓ ER cells. * ↑ Hassall's corpuscles

Malt – mucous associated lymphoid tissue

From where	MALT- mucosa associated lymphoid tissue
Def	Collective name for the cells of the immune system in the mucosa of respiratory , alimentary , urogenital tracts
Fun	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.