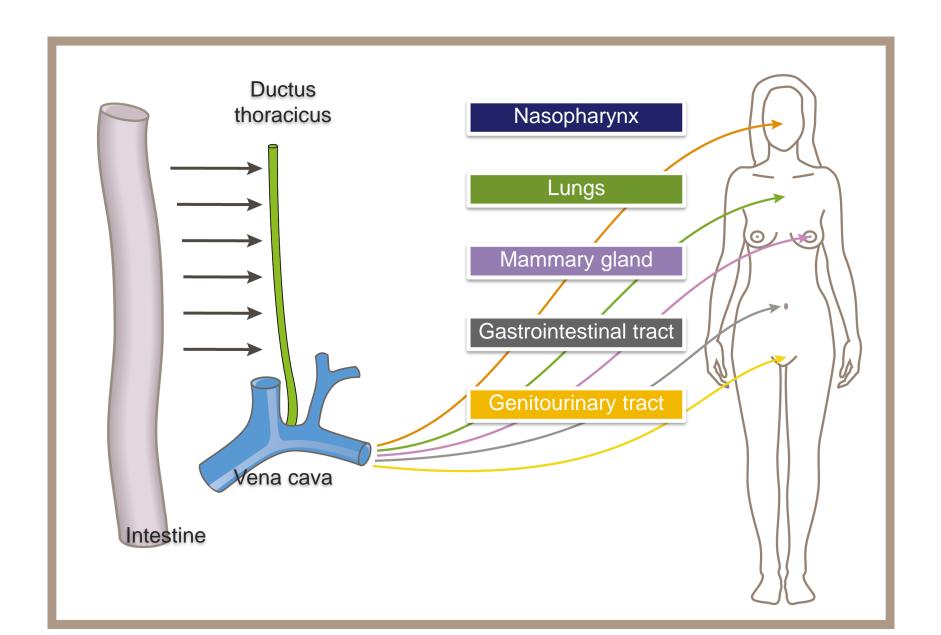
Mucosal immune system (MALT)

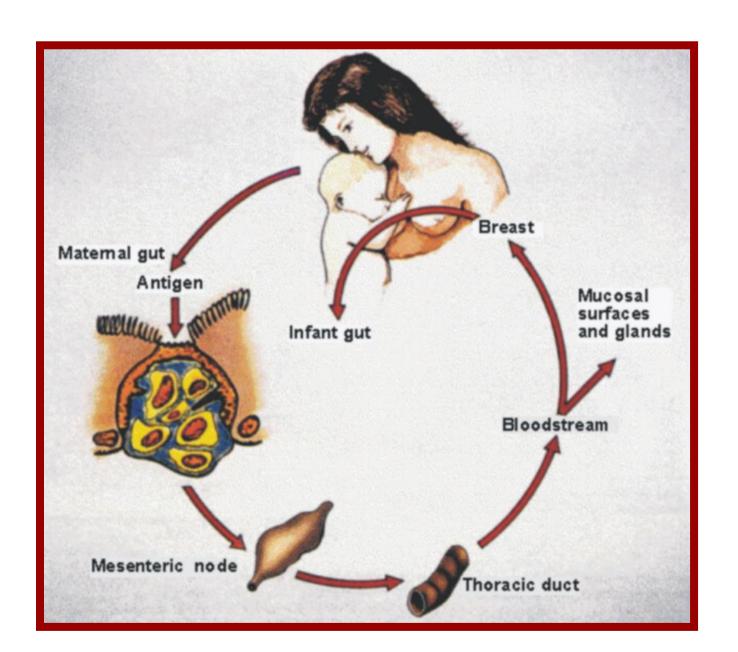
MALT (Mucous Associated Lymphoid Tissue)

- GALT (Gut Associated Lymphod tissue)
- BALT (Bronchi Associated Lymphoid Tissue)
- Immune tissues of the urinary tract, genital tract, conjunctiva, middle ear...
- Includes also brest gland!

Common immune system of mucous membranes



Role of brestfeeding in MALT



Imunologial aspects of brestfeeding

- IgA present mainly in colostrum, much less in maternal milk. IgA is not absorber in human GIT.
 The protective effect is limited to GIT.
- Presence of various immunoregulatory substances (cytokines, growth factors..).
- Antibacterial substances lysozyme, laktoferin.
- Markedly decreased koncentration of exgenous food allergens, however this is not an absolute elimination!

Homing of Lymphocytes

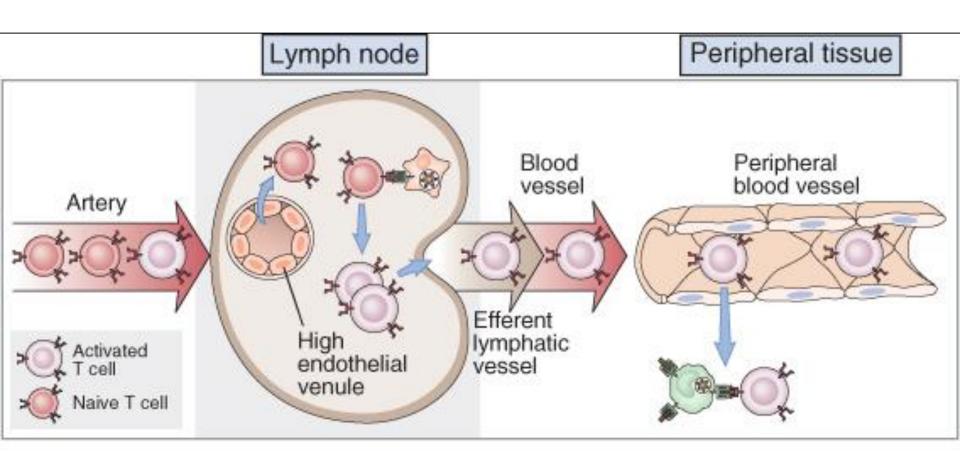
- The directed migration of subsets of circulating lymphocytes into particular tissue sites.
- Regulated by selective expression of adhesion molecules called homing receptors on lymphocytes.
- Tissue speciphic endothelial ligands are called addressins.

High Endotelial Venules

- Specialized venules. The site where lymphpocytes leave the blood stream and migrate into lymph nodes, spleen, organs of MALT.
- Adhesion molecules enable selective attachment of various types of lymphocytes.



Circulation of lymphocytes



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Mucosal immune system (MALT)

- Antigenic stimulation in one part of MALT leads to immune response also in other compartments of MALT.
- IgA is a predominant immunoglobulin secreted by the epitelial cells.
- Oral administration of antigens frequently leads to induction of immune tolerance.
- Specialized types of cells: Intraepitelial lymphocytes, M-cells.

Anatomy of MALT

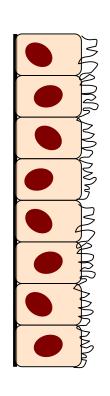
- Diffuse tissue containing lymphocytes and other cells of the immune system in submucosa.
- Specialized organs:
 - Waldeyer's ring
 - Payer's patches
 - Appnedix

Epitelial cells are intergal part of the immune system of mucous membranes

EXPRESION

- enzymes
- HLA antigens
- adhesion molecules
- receptors for:

mikrobes cytokines polymeric lg



PRODUCTION

- cytokines
 pro-inflammatory
 growth factors
 chemotactic
- antibiotic peptides
- various other mediators

INTERACTION WITH SPECIFIC IMMUNE SYSTEM

Antimicrobial nechanisms on mucous membranes

i actor	WECHAIIISIIIUS
Comensal bacteria	competition with pathogens production of antiinflammatory mediators
Tight epitelial junctions	protect from bacterial invasion into tissues
Cilia	bind and remove microbes
Mucin	bind microbes
Lysozyme	killing G+ bacteria
Laktoferin	iron binding (inhibition o microbial growth)

Mechanismus

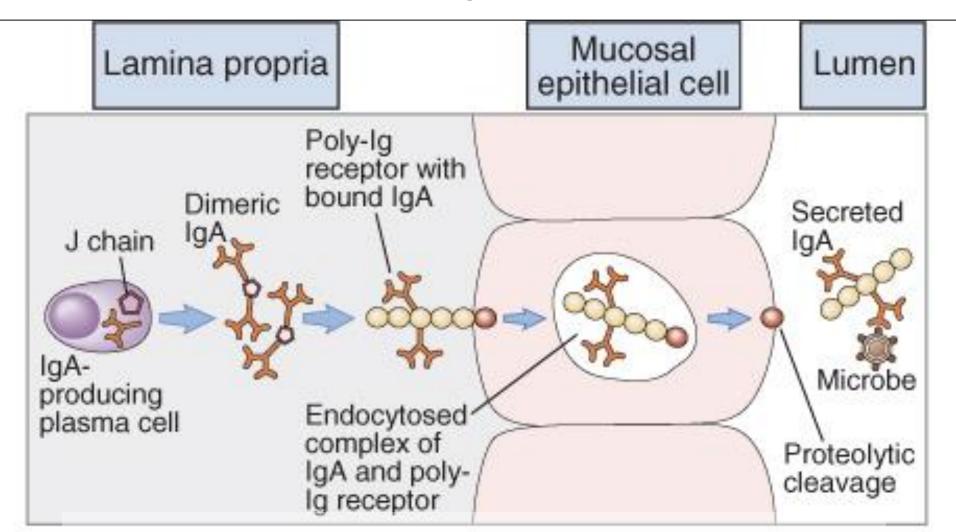
Antibiotic peptides killing microbes (mainly β defensins)

Factor

Secretory Ig Microbial adhesion blocade



Secretory IgA formation



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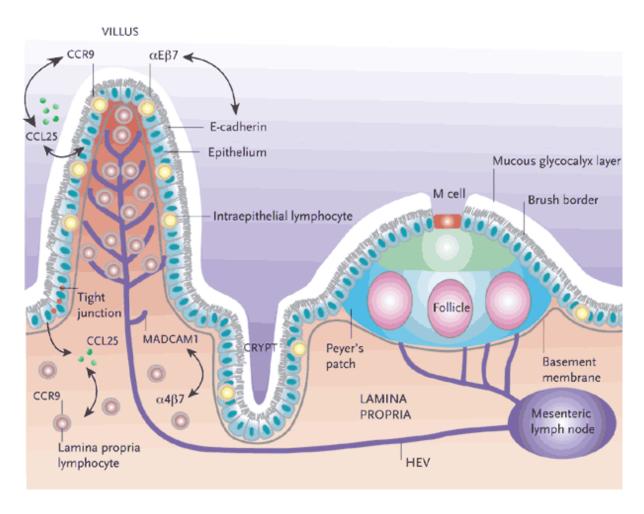
Intraepitelial T-lymphocytes

- TCR $\alpha\beta$ or $\gamma\delta$
- Extrathymic differentiation
- First line of specific immune response
- Predominantly CD8+
- Low antigenic specificity of TCR

M-cells

- Specialized enterocytes responsible for transport of antigens from the gut towards the immunocompetent cells inside the Payer's patches.
- Transport in mediated by transcytosis.

Lymphocyte circulation in GALT



Oral tolerance

- Stimulation of the GALT frequently leads to induction of immune tolerance to the stimulating antigen.
- This occurs mainly if the gut is in "normal, noninflammatory" conditions.
- Induction of Th3 cell is the main mechanism.
- The tolerance is important to avoid unnecessary reactions to non-pathogenic antigens.

Comensal (normal) microflore (of GIT)

- ~ 10¹⁴ microbial cells, ~ 1000 microbial species
- ~ 50% non cultivated
- Complex ecosystem
- Included in innate immunity of GIT
- Mutual interactions of microorganisms: competition, symbiosis..
- Interaction with macroorganism: symbiosis, commensalism, important in metabolic processes (production of vitamins etc.)
- Immune system modulation

Variability of comensal flora in various compartments of human body

