Mechanisms of Autoimmunity

Immunology Unit
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Reference Kuby Immunology 7th Edition 2013 Chapter 16 Pages 531-534

Objectives

- Autoimmunity results from activation of immune response against self antigens.
- To learn how immunological tolerance (central and peripheral) is induced against self antigens for maintaining normal health.
- To gain understanding of various factors contributing to the breakdown of immunological tolerance and development of autoimmunity.
- Gender predilection in autoimmunity is a well known phenomenon and is briefly described.

Autoimmunity

A condition that occurs when the immune system mistakenly attacks and destroys healthy body tissue

Autoimmunity

Immune system has evolved to discriminate between

Self and Non-self

Mediated by auto-reactive T cells and auto-reactive B cells (auto-antibodies)

Tolerance to self is acquired by:

- A) Deletion (clonal deletion) OR
- B) Functional inactivation (clonal anergy) of developing lymphocytes that possess antigenic receptors with high affinity for self-antigens.

Self-Tolerance

Central Tolerance
(Thymus & Bone marrow)

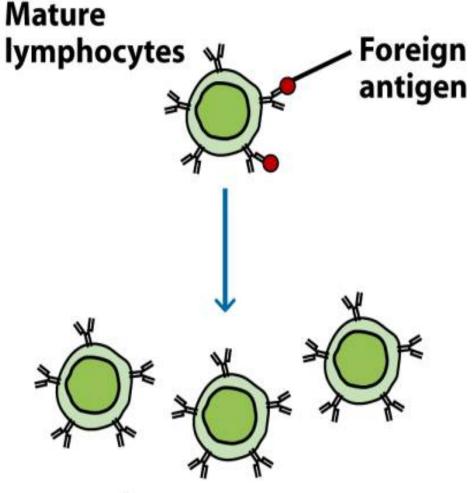
Peripheral tolerance (Peripheral tissues)

Central tolerance Lymphoid Newly emerged precursor (immature) clones of lymphocytes Self antigen present in generative lymphoid organ Maturation of clones Central tolerance: not specific for deletion of lymphocytes self antigens present specific for self antigens in generative organs present in generative organs

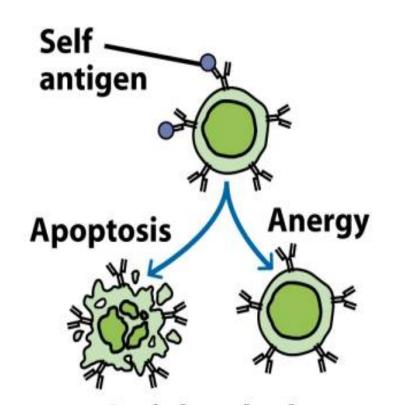
Figure 16-1a
Kuby IMMUNOLOGY, Sixth Edition
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Educated T-cell Autoreactive cell Central Tolerance

Peripheral tolerance

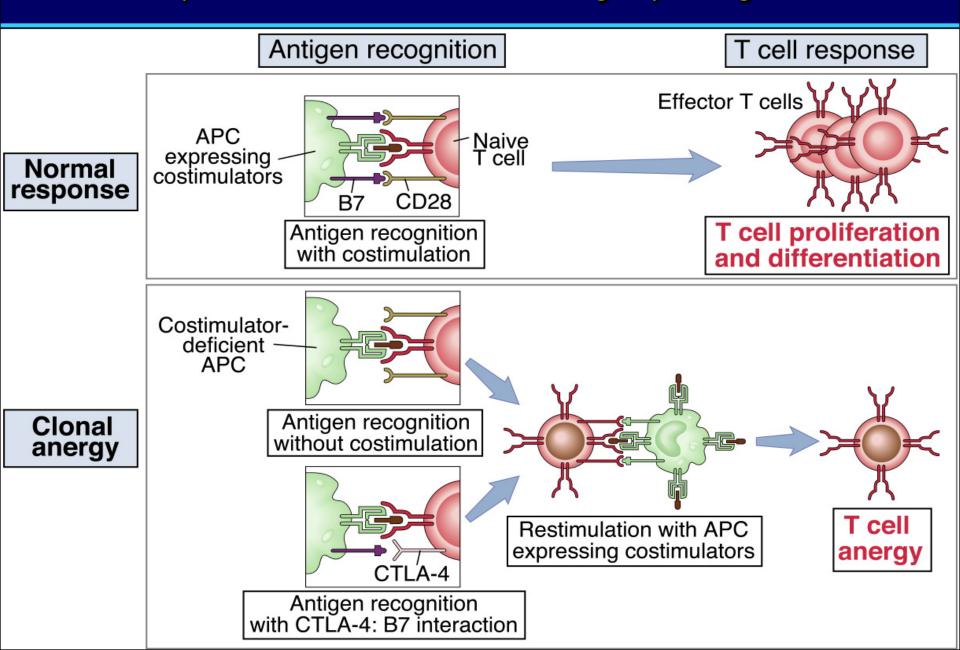


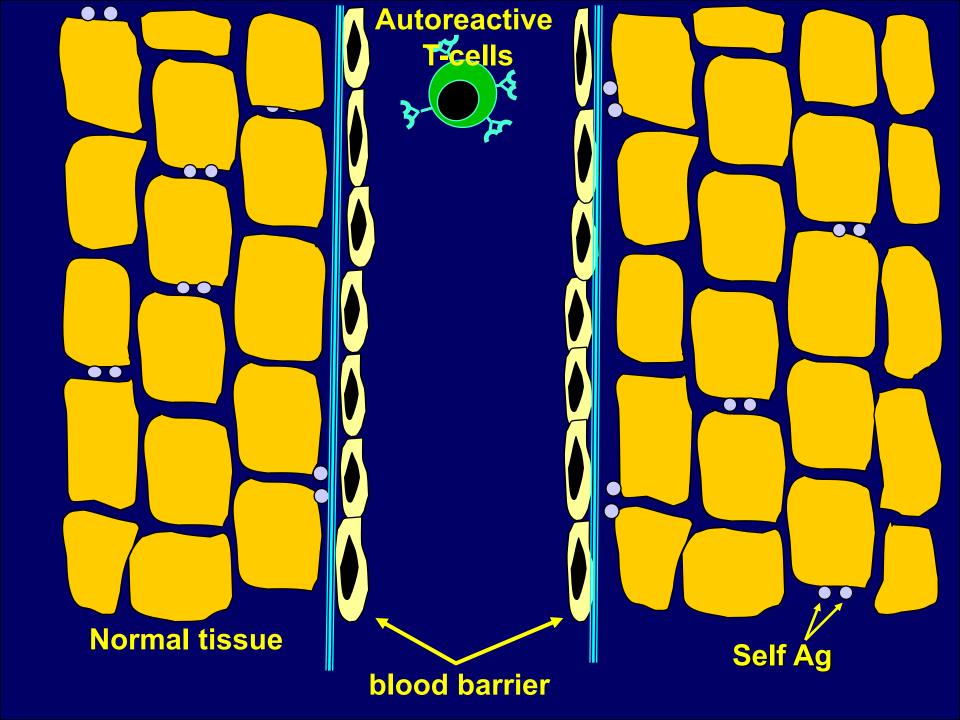
Immune response to foreign antigens



Peripheral tolerance: deletion or anergy of lymphocytes that recognize self antigens in peripheral tissues

Peripheral Tolerance of T Lymphocytes





Failure of Immune Tolerance (Development of Autoimmunity)

Induction of Autoimmunity "Proposed Mechanisms!"

- 1. Sequestered antigens
- 2. Molecular mimicry
- 3. Inappropriate class II MHC expression on none-antigen presenting cells
- 4. Polyclonal B cell activation

1. Sequestered antigens

- Some self-antigens are sequestered (hidden) in specialized tissues.
- These are not seen by the developing immune system – will not induce selftolerance.
- Exposure of T cells to these normally sequestered/tissue-specific selfantigens in the periphery results in their activation.

Examples of Sequestered Antigens

Myelin basic protein (MBP), associated with MS

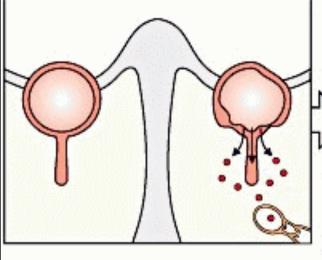
Sperm-associated antigens in some individuals following vasectomy

Lens and corneal proteins of the eye following infection or trauma

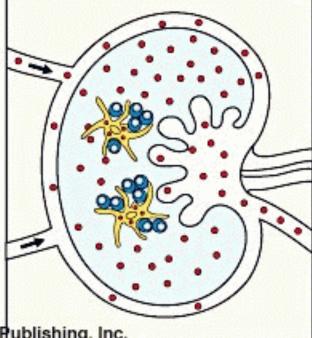
Heart muscle antigens following myocardial infarction

Sympathetic ophthalmia

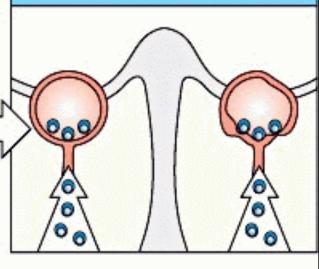
Trauma to one eye results in the release of sequestered intraocular protein antigens



Released intraocular antigen is carried to lymph nodes and activates T cells



Effector T cells return via bloodstream and encounter antigen in both eyes



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2. Molecular Mimicry (Cross-reacting Antigens)

 Viruses and bacteria possess antigenic determinants that are very similar, or even identical, to normal host cell components.

 This phenomenon, known as molecular mimicry, occurs in a wide variety of organisms.

 Molecular mimicry may be the initiating step in a variety of autoimmune diseases.

Examples of Molecular Mimicry

ORGANISMS AND HUMAN HOST PROTEINS

Protein*	Residue [†]	Sequence [‡]
Human cytomegalovirus IE2	79	PDPLGRPDED
HLA-DR molecule	60	VTELGRPDAE
Poliovirus VP2	70	STTKESRGTT
Acetylcholine receptor	176	TVIKESRGTK
Papilloma virus E2	76	SLHLESLKDS
Insulin receptor	66	VYGLESLKDL
Rabies virus glycoprotein	147	TKESLVIIS
Insulin receptor	764	NKESLVISE
Klebsiella pneumoniae nitrogenase	186	SRQTDREDE
HLA-B27 molecule	70	KAQTDREDL
Adenovirus 12 E1B	384	LRRGMFRPSQCN
α -Gliadin	206	LGQGSFRPSQQN
Human immunodeficiency virus p24	160	GVETTTPS
Human IgG constant region	466	GVETTTPS
Measles virus P3	13	LECIRALK
Corticotropin	18	LECIRACK
Measles virus P3	31	EISDNLGQE
Myelin basic protein	61	EISFKLGQE

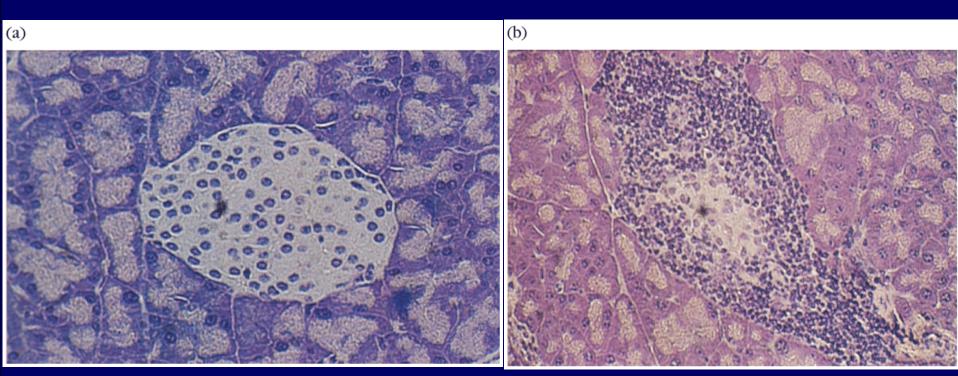
3. Inappropriate Expression of Class II MHC Molecules

- Class II MHC ordinarily expressed on antigen presenting cells, such as macrophages, dendritic cells and B cells.
- Abnormal expression of MHC determinants allows the recognition of these auto-antigens by selfreactive T cells.

Inappropriate Expression of Class II MHC Molecules

- This may occur due to the local production of IFN-γ, which is known to increase class II MHC expression on a variety of cells.
- The inducer of IFN-γ under these circumstances could be a viral infection.

Type I Diabetes: Pancreatic β cells express abnormally high levels of MHC I and MHC II (?)



Normal Pancreas

Pancreas with Insulitis

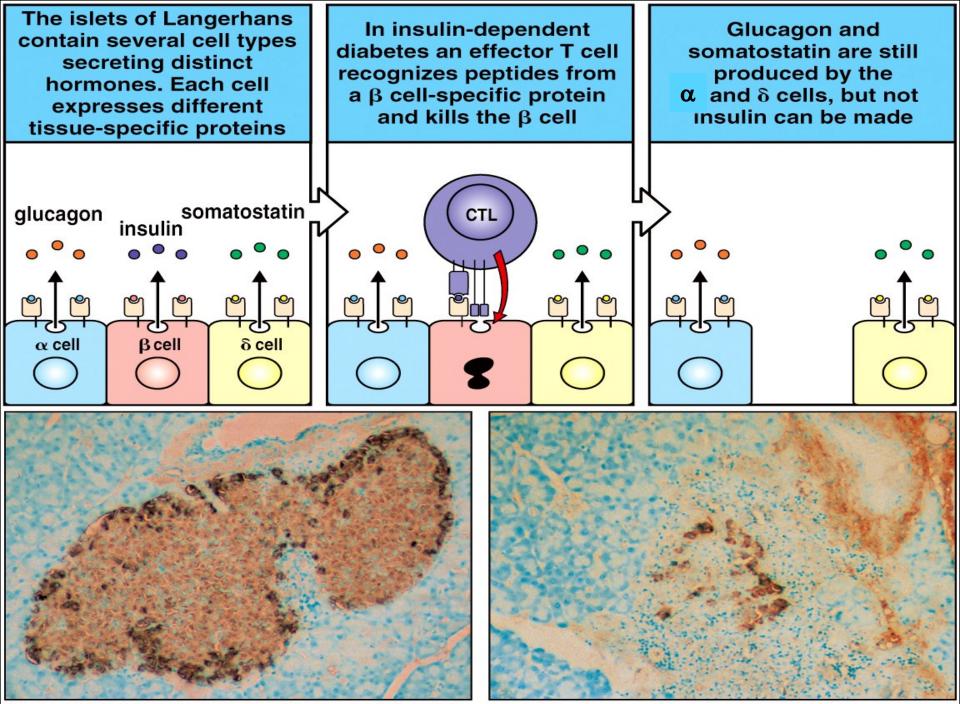


Figure 13-34 Immunobiology, 6/e. (© Garland Science 2005)

4. Polyclonal B Cell Activation

Viruses and bacteria can induce nonspecific polyclonal B cell activation, including:

- Certain gram negative bacteria
- Herpes simplex virus.
- Cytomegalovirus
- Epstein Barr Virus
- Human immunodeficiency virus (HIV)

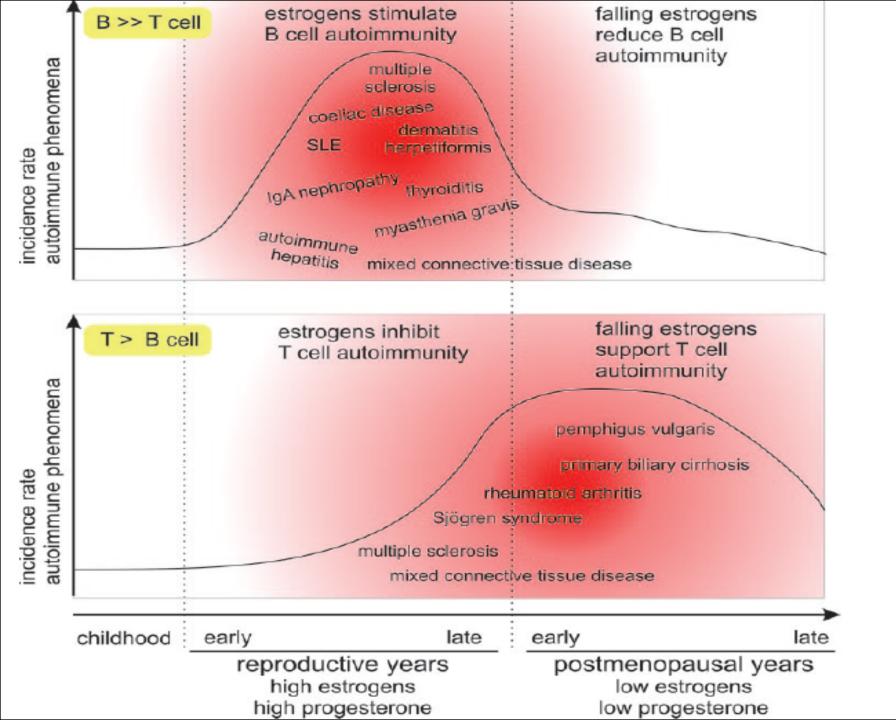
These viruses induce the proliferation of numerous clones of B cells to secrete IgM in the absence of a requirement for CD4 T cell help.

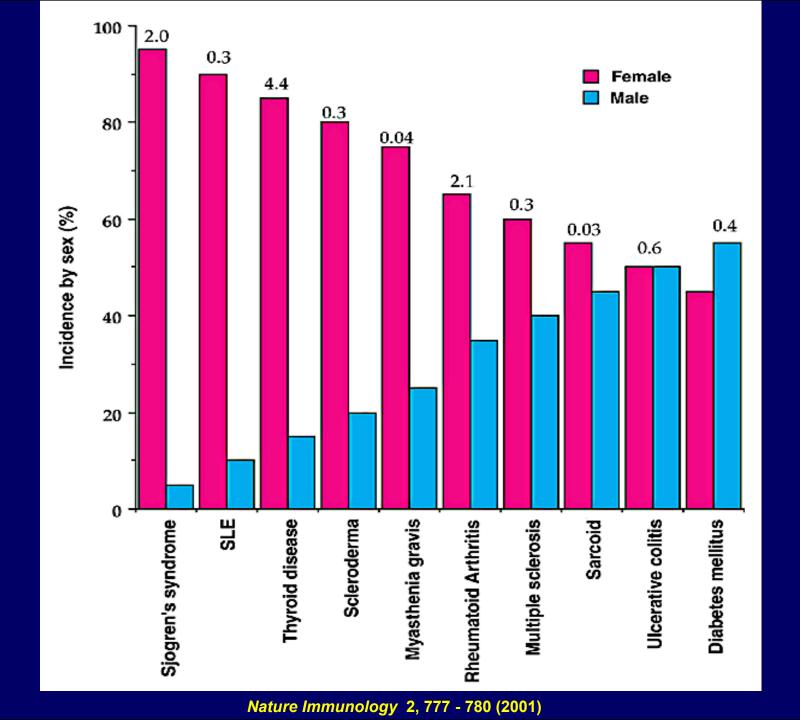
Polyclonal activation leads to the activation of self-reactive B cells and autoantibody production.

Patients with infectious mononucleosis (caused by EBV) and AIDS (HIV) have a variety of auto-antibodies.

Hormonal Factors

- About 90% of autoimmune diseases
 occur in women cause not known
- In animal models estrogen can induce B cells to enhance formation of anti-DNA antibodies
- SLE either appears or exacerbates during pregnancy





Take home message

- Normal healthy state is maintained by immunological tolerance against self antigens at central and peripheral levels
- Autoimmune diseases result from the breakdown of immunological tolerance to self antigens
- Certain autoimmune diseases exhibit strong association with female gender

Thank you