CHAPTER 7

Major Histocompatibility Complex (MHC)

- ➢ What is MHC? − HLA
 - H-2
 -
 - Minor histocompatibility antigens
 Peter Gorer & George Sneell (1940)

Significance of the MHC

role in immune response
role in organ transplantation
role in predisposition to disease

- MHC molecules were initially discovered during studies aimed at understanding the molecules responsible for rejection of transplanted tissues.
- Hence the name "Major Histocompatibility Complex" (MHC).
- The term "Major Histocompatibility Complex" actually refers to a region of the genome that encodes a number of genes (hence Complex) that play an important (hence Major) role in tissue transplantation (hence Histocompatibility).
- The term "MHC molecule" or "MHC antigen" refers to a molecule encoded by a gene within this region.

louse II 2	H-2 complex Chromosome 17							
Complex	H–2							
MHC class	I II III III III			I				
Region	к	IA	IE	S		D L		
Gene	H-2K	IA	IE	C' proteins	TNF-α	H–2D H–2L		
products		αβ	αβ		TNF-β			
products Iuman HL Complex		αβ	αβ		HLA			
luman HL Complex		αβ	αβ		HLA		1	
luman HL		αβ	DR	Chro	mosome 6 HLA	В	I C	A

In humans:

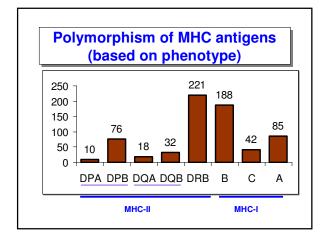
- Class I = A, B and C (also called HLA-A, HLA-B and HLA-C) - Ag (peptide) presentation to CD8+ cells
- Class II = DP, DQ and DR (also called HLA-DP, HLA-DQ and HLA-DR) - Ag (peptide) presentation to CD4+ cells
- Class III = Complement proteins, Tumor necrosis factor (TNFs)- α , β

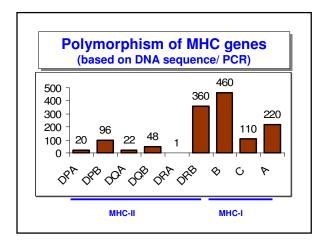
In the Mouse:

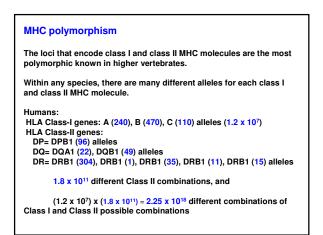
- Class I = K, D and L molecules (also called H-2D, H-2K and H-2L)
- Class II = A and E (also called I-A and I-E)
- Class III = Complement proteins, Tumor necrosis factor (TNFs)- α , β

MHC- Polimorphism

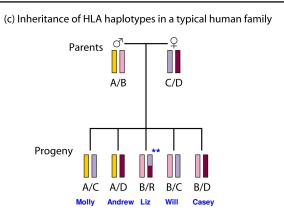
- MHC loci are highly polymorphic presence of many alternative forms of the gene or alleles in the population
- Inherited from mother and father
- New haplotypes are generated by <u>recombination</u>





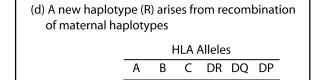






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7 w3 2 1

8

44

35

1

2

3

4

4

2

1

3

3

w2 3

4

7

w4

w1 7

w4

A 1

2

3

В

С

D 11

R 3 44

Haplotypes

Terminology:

- **Haplotype**: set of alleles present in each parental chromosome (two sets).
- **Inbred mouse strains**: same set of alleles (homozygous) at each locus (K, IA, IE, S, D).
- Inbred strains are **SYNGENIC** = identical at <u>all</u> genetic loci
- Inbred strains have been bred by brother-sister mating for > 20 generations
- Outbred mouse strains: different set of alleles at each locus ~ like humans.
- **Congenic strains** = genetically identical except at <u>a single</u> loci

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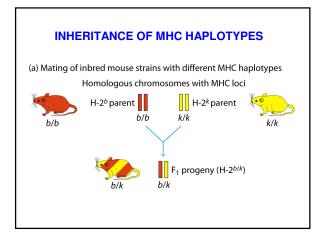
Mouse Strains

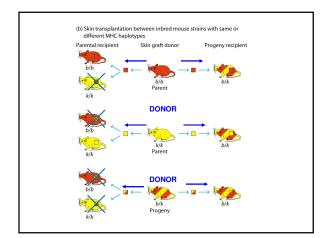
 Thus, the strain C57BL/6 was designated H-2^b haplotype and said to possess the 'b' allele at each MHC locus.

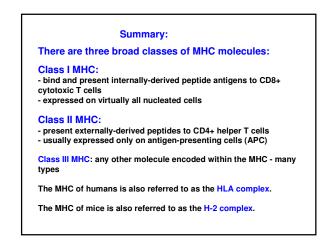
Thus, it is: $H-2b = K^b$, D^b , L^b , $I-A^b$, $I-E^b$

- Another strain, CBA/2 was found to possess different alleles than C57BL/10 and was arbitrarily designated as having the k haplotype (I.e. H-2^k).
- Thus, it is: $H-2k = K^k$, D^k , L^k , $I-A^k$, $I-E^k$

TABLE 7-1	I-2 Haplotypes of some mouse strai	ins		1943	H-2 ALLELE	5	
Prototype strain	Other strains with the same haplotype	Haplotype	ĸ	IA	IE	s	L
CBA	AKR, C3H, B10.BR, C57BR	k	k	k	k	k	,
DBA/2	BALB/c, NZB, SEA, YBR	d	d	d	d	d	6
C57BL/10 (B10)	C57BL/6, C57L, C3H.SW, LP, 129	ь	ь	Ь	ь	ь	Ł
A	A/He, A/Sn, A/Wy, B10.A	а	k	k	k	d	6
A.SW	B10.5, SJL	\$	5	5	5	\$	5
A.TL		<i>t1</i>	s	k	k	k	6
DBA/1	STOLI, B10.Q, BDP	9	q	9	9	9	9



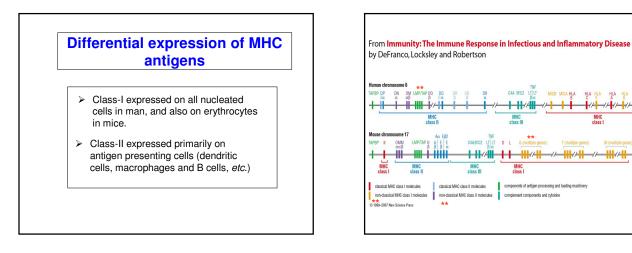


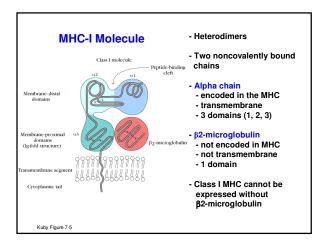


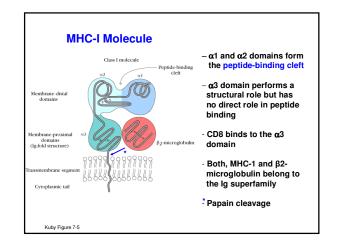
HLA

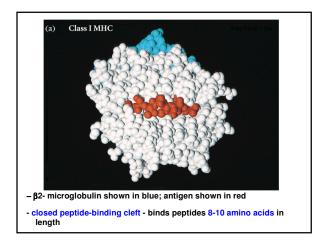
HH

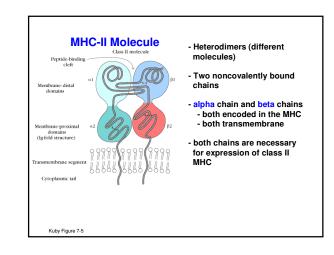
MHC class I

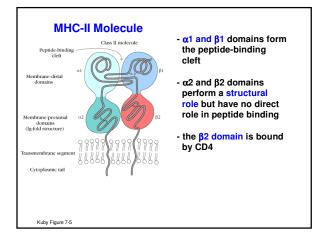


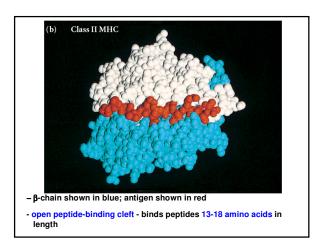


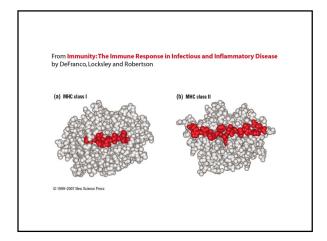


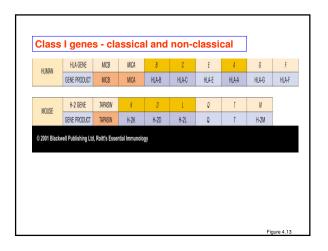




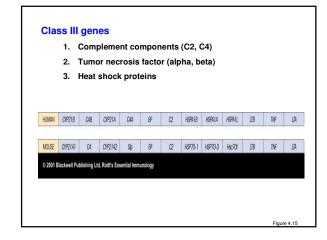








HUMAN GENE TADACIN DPB					2 DOB	DQB DQA	DRB DRA
	DP α DOα	DMα DMβ	PROTEASOME	PEPTIDE	DOβ	ΟQβ DQα	DRB DRa
PRODUCT	ILA-DP HLA-DO	HLA-DM	COMPLEX	TRANSPORT	R HLA-DO	HLA-DQ	HLA-DR
H-2 GENE Oa Ma	Mb2 Mb1	LMP2 TAP2	LMP7 TAP1	Ob A	b Aa	Eb	Ea
MOUSE GENE Oa DMa	DMβ2 DMβ1	PROTEASOME	PEPTIDE	βΑ	3 Aa	Εβ Ι	Εα
PRODUCT H-20	H-2DM	COMPLEX	TRANSPORTER	H-20	H-2A	H-2E	
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Peptide-MHC Interaction

- Peptide binding by MHC molecules is not as specific as antigen binding by antibodies or T cell receptors.
- Any particular MHC molecule will bind a large range of peptides but not all peptides.
- A given MHC molecule will bind peptides that have certain amino acids at key positions in the peptide (anchor residues).
- Each MHC molecule binds a unique set of peptides. Keep in mind that each allelic variant also binds a unique set of peptides!!

MHC-Peptide Interaction

MHC-I:

- Each unique molecule (A, B or C) binds a unique set of peptides
- Single nucleated cell express $10^{5}\ \text{of}\ \text{each}\ \text{class}\ I\ \text{molecule}$
- As few as 100 peptide-MHC complexes are enough to target a cell for killing by CD8+
- Requirements: 1) 8-10aa length, 2) key amino acids at positions 2 and 9

	Aromatic		Hydrophobic	
Elut fror H-2	$ \begin{array}{c} \text{H}_{3}N - V - G - (H_{3}N - S - (H_{3}N - S - (H_{3}N - S - (H_{3}N - S - (H_{3}N - H_{3}N - S - (H_{3}N - H_{3}N - H_{3}N - (H_{3}N - H_{3}N - H_$	3 4 5 6 P-Q-K-N-(P-R-K-A-(P-S-G-K-(P-E-R-1-($\begin{array}{cccccccccccccccccccccccccccccccccccc$	
Elut fror H-2	$ \begin{array}{c} \mathbf{n} \\ \mathbf{K}^{d} \\ \mathbf{H}_{3}\mathbf{N} \\ \mathbf{N} \\ N$	F-P-E-1-(Q-R-T-R-(1-G-S-1-(T)-H)-[]-coo- A)-[]-V)-coo- N)-(N)-[]-coo-	
	A = alanine E = glutamic acid F = phenylalanine G = glycine * H = histidine I = isoleucine *	K = lysine L = leucine \star N = asparagine P = proline Q = glutamine	$\begin{split} R &= arginine \\ S &= serine \\ T &= threeonine \\ V &= valine \\ Y &= tyrosine \end{split}$	

Peptide-binding grooves for class I and class II MHC are structurally similar

- Both have a peptide-binding groove
- Close-ended groove for class I MHC requires an 8-10 amino acid-length peptide to bind
- Open-ended groove for Class II MHC lets it bind a peptide 13-18 amino acids long, not all of which lie in the groove
- Anchor site rules apply to both classes in particular Class I MHC (P2 and P9)

Aspects of MHC

- 1. Recognition by T cells requires cell-cell contact.
- 2. Peptides from cytosol associate with class I MHC and is recognized by Tc cells.
- 3. Peptides from endocytic vesicles associate with class II MHC and is recognized by Th cells.

Aspects of MHC (continued)

- 3. Although there is a high degree of polymorphism for a species, an individual has maximum of six different class I MHC products and eight class II MHC products.
- 4. A peptide must associate with a given MHC of that individual, otherwise no immune response can occur. That is one level of control.

Aspects of MHC (continued)

- 4. Mature T cells must have a T cell receptor that recognizes the peptide associated with MHC. This is the second level of control.
- 5. Each MHC molecule has only <u>one</u> binding site. The different peptides a given MHC molecule can bind all bind to the same site, but only one at a time.

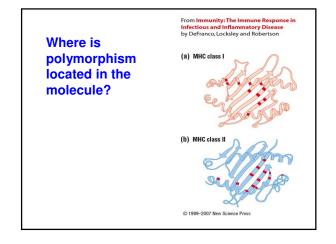
Aspects of MHC (continued)

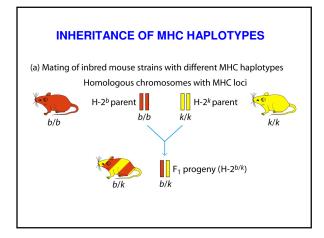
- MHC polymorphism is determined only in the germline. There are <u>no</u> recombinational mechanisms for generating diversity.
- 7. Because each MHC molecule can bind many different peptides, binding is termed **degenerate**.
- 8. Cytokines (especially interferon-γ) increase level of expression of MHC.

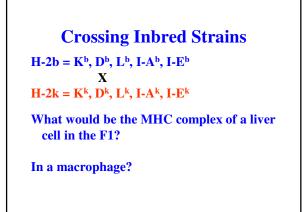
Aspects of MHC (continued)

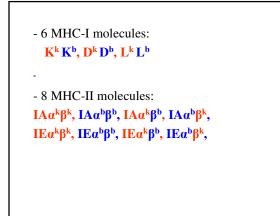
- 9. Alleles for MHC genes are **co-dominant**. Each MHC gene product is expressed on the cell surface of an individual nucleated cell.
- 10. Why the high degree of polymorphism?

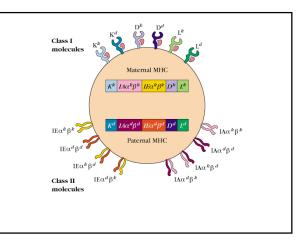
Survival of species!











Regulation of MHC Expression

- 1) Cytokines:
 - IFN-alpha, beta, gamma \uparrow Class-I expression.
 - IFN-gamma \uparrow Class-II expression in MO and DC
 - IL-4 \uparrow expression of MHC-II in resting B cells
 - IFN-gamma \downarrow expression of MHC-II in B cells
- 2) Corticosteroids and Prostaglandins •↓ expression of MHC-II
- 3) Viruses (↓ expression of MHC-I)
 - Human cytomegalovirus (CMV)
 - Hepatitis B virus (HBV)
 - Adenovirus 12 (Ad12)

MHC and immune responsiveness:

In many cases, the ability of an inbred mouse strain to respond to a given antigen will depend on which alleles the strain carries at its MHC loci.

The reason is that if an antigen cannot bind to an MHC molecule, it cannot be presented to T cells and therefore an immune response cannot be made to it.

To respond to an antigen, the **first criterion** that must be met is that the individual must have an MHC molecule that can bind and present the antigen.

The **second criterion** that must be met is that the individual must have T cells capable of responding to the antigen.



T cells are MHC-restricted i.e. they must recognize antigen presented on MHC.

CD4+ T cells are class II MHC-restricted i.e. they must recognize antigen presented on class II MHC.

CD8+ T cells are class I MHC-restricted i.e. they must recognize antigen presented on class I MHC.

A particular T cell clone may be I-E^k-restricted i.e. it recognizes its antigen ONLY when presented on I-E^k.

("restricted" = "recognizes antigen on...")

Associations between MHC and disease

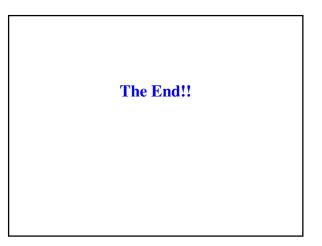
The risk of developing immunological diseases is often influenced by the presence or absence of specific MHC alleles.

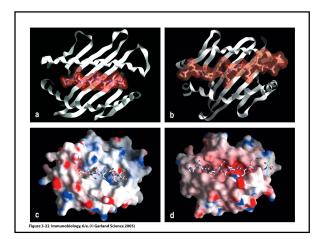
TABLE 7-4 SOME SIGNIFICANT ASSOCIATIONS OF HLA ALLELES WITH INCREASED RISK FOR VARIOUS DISEASES

	Associated HLA allele	Relative risk'
Ankylosing spondylitis	B27	90
Goodpasture's syndrome	DR2	16
Gluten-sensitive enteropathy	DR3	12
Hereditary hemochromatosis	A3	9.3
	B14	2.3
	A3/B14	90
Insulin-dependent diabetes mellitus	DR4/DR3	20
Multiple sclerosis	DR2	5
Myasthenia gravis	DR3	10
Narcolepsy	DR2	130
Reactive arthritis (Yersinia, Salmonella, Gonococcus)	B27	18
Reiter's syndrome	B27	37
Rheumatoid arthritis	DR4	10
Sjogren's syndrome	Dw3	6
Systemic lupus erythematosus	DR3	5

Associations between MHC and disease

Disease	Relative Risk	Allele
Ankylosing Spondylitis	90	B27
• Hereditary hemochromatosis	90	A3/B14
Narcolepsy	130	DR2





Self-MHC-restriction of T cells

Generally, T cells must recognize antigen on a self MHC allele and so are said to be self-MHC restricted.

This is because T cells are "tuned" to recognize antigen complexed with self-MHC during T cell maturation in the thymus.