

# Biology 112 Unit 7

## Chapter Nine Patterns of Inheritance

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### I. Mendelian Genetics

- Gregor Mendel – monk who worked with pea plants
  - Strong training in the sciences and mathematics
  - Good experimental design and meticulous records
- Published paper in 1866
  - Said parents pass on discrete heritable factors

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### Opposed Two Theories of the Time

- 1) Pangenesis
  - Hippocrates theory
  - Acquired traits of the adult migrate to gametes
- 2) Blending Hypothesis
  - Early 19<sup>th</sup> century
  - Heritable traits from each parent blend in young

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## Why peas?

- You can easily trace 7 traits with 2 distinct forms each
- Can easily do both self-fertilization and cross-fertilization
  - Allowed him to keep track of parentage
- Kept going until he had true-breeding varieties
  - All identical to the parent plant (homozygous dominant/recessive)

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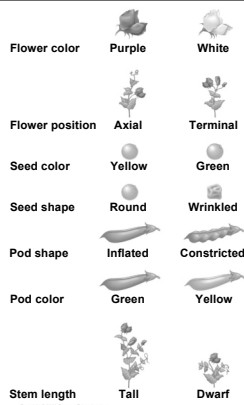
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## II. Monohybrid Crosses

- Track inheritance of a single trait
- Four Mendelian Hypotheses
  - 1) Different forms of a gene exist (allele)
  - 2) Each organism carries two copies (diploid)

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3) Gametes only carry one copy (haploid) due to Principle (or Law) of Segregation

Combining gametes (Offspring):

Homozygous – carry SAME form of the gene

Heterozygous – carry DIFFERENT forms of the gene

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4) Forms may differ in expression:

Dominant – Homozygous and Heterozygous

Recessive – Homozygous only

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## Solving Mendelian Inheritance

P generation X P generation

F<sub>1</sub> generation X F<sub>1</sub> generation

F<sub>2</sub> generation

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### Describe the organism two ways

- 1) Phenotype  
- What you look like
- 2) Genotype  
- your genetic makeup

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### Punnett Square

- Visually shows principle of Independent Assortment
- Steps in Solving
  - 1) Determine P generation genotypes (Diploid)
  - 2) Determine Gametes (Haploid)

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- 3) Cross gametes in square to yield offspring (Diploid)
- 4) Summarize

Example

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## Rules of Probability

- Rule of Multiplication
  - Product of each event  $\frac{1}{2} \times \frac{1}{2} = \frac{1}{4}$

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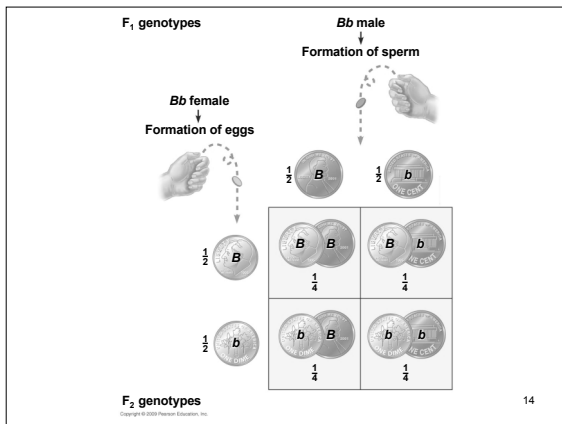
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## Testcross

- Used to determine unknown genotypes by crossing with homozygous recessive individual

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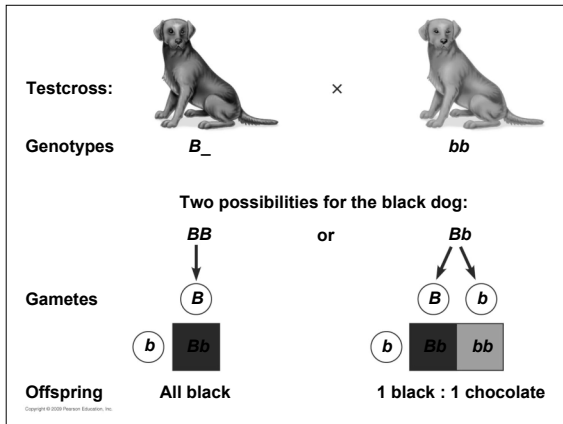
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### III. Dihybrid Cross

- Track the inheritance of two traits
- Steps to solving:
  - Determine P generation (diploid)
  - Determine gametes (haploid)
  - Cross gametes in square to yield offspring (diploid)
    - FOIL
  - Summarize

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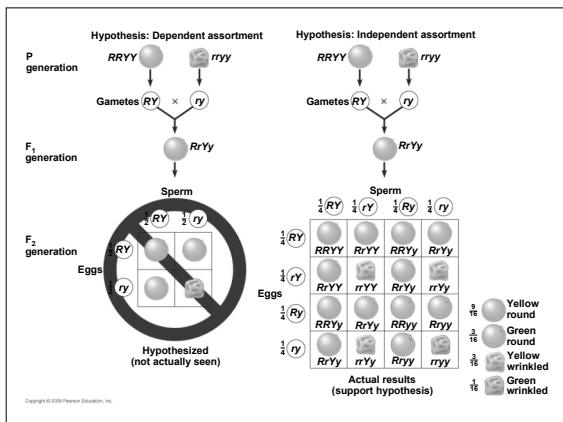
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## IV. Pedigrees

- Family genetic history illustrated
- Allows one to deduce pattern of inheritance
- Also to determine carriers – those that have the gene but are not affected by it

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	Female	Male
Normal	○	□
Affected	●	■

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First generation  
(grandparents)

□ *Ff*    ○ *Ff*    ■ *ff*    ○ *Ff*

Second generation  
(parents, aunts,  
and uncles)

○ *FF or Ff*    ■ *ff*    ■ *ff*    □ *Ff*    ○ *Ff*    ○ *ff*

Third generation  
(two sisters)

● *ff*    ○ *FF or Ff*

Female	Male	
●	■	Affected
○	□	Unaffected

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## Patterns of Inheritance

### Autosomal

#### 1. Simple Dominant / Recessive

#### Autosomal Recessive

Albinism, deafness, unattached earlobes,  
Cystic Fibrosis inflicts 1/1800 caucasian,  
1/25 are carriers

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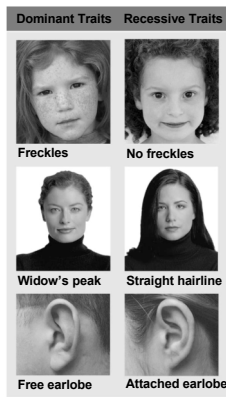
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### Autosomal Dominant

Extra digits, webbed digits, tongue rolling,  
dimples, achondroplasia (one form of  
dwarfism), Alzheimer's and Huntington's

Dominant doesn't necessarily mean most  
prevalent

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- 2) Incomplete dominance
- Intermediate phenotypes
  - Red x White = Pink
  - coat color (cats, cows), hypercholesterolemia, flower colors

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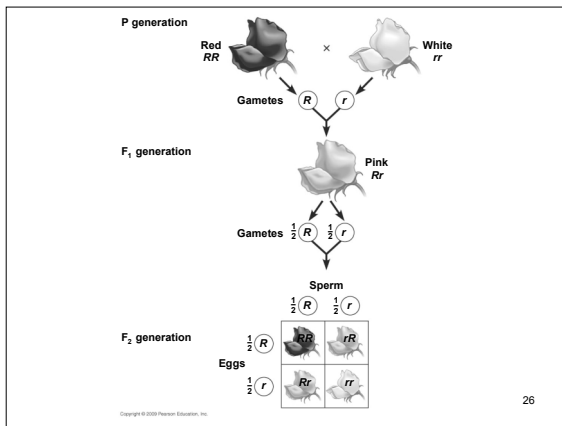
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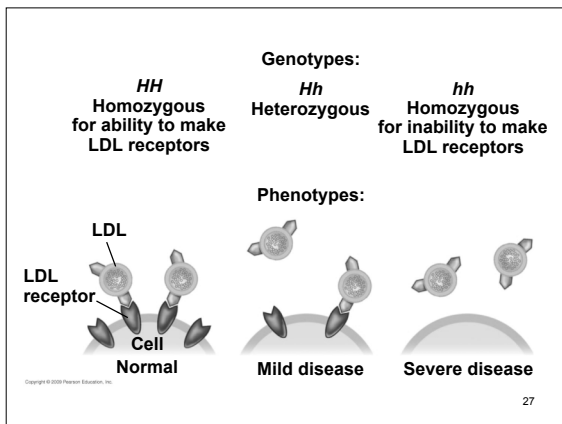
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- 3) Multiple Alleles:
- more than two alleles in the population
  - Example – ABO Blood Type of Humans (codominance)

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### Blood Type

Phenotype	Genotype
A	AA, AO
B	BB, BO
AB	AB
O	O

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Blood Group (Phenotype)	Genotypes	Red Blood Cells
O	$i i$	
A	$I^A I^A$ or $I^A i$	 Carbohydrate A
B	$I^B I^B$ or $I^B i$	 Carbohydrate B
AB	$I^A I^B$	

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4) Pleiotropy

- one trait affects more than one character
- PKU and Sickle cell

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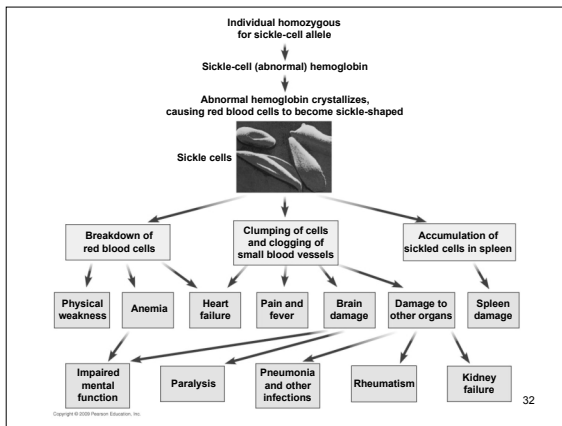
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5) Polygenic:

- additive effect of 2 or more genes on a single trait
- skin / hair/ eye colors and height in humans

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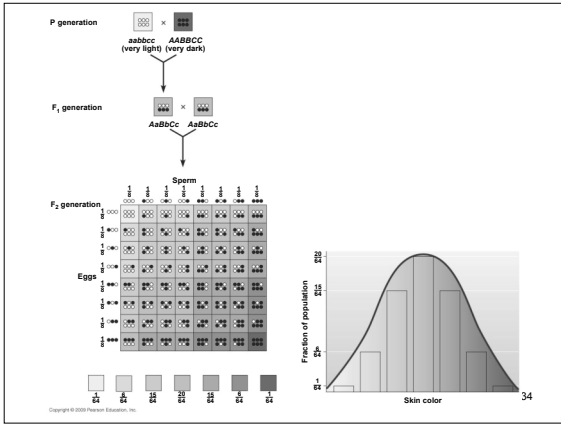
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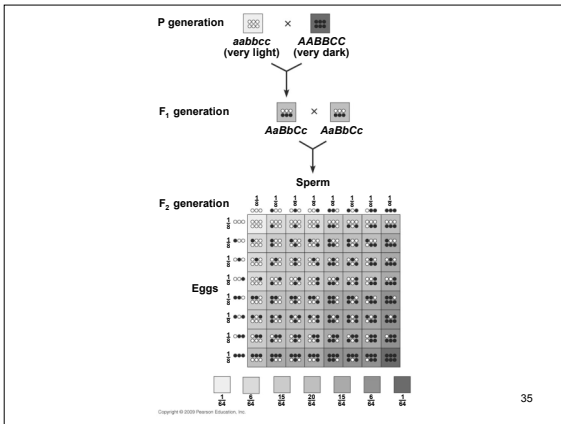
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## Sex Chromosomes

6) Sex-linked:

X-linked = colorblindness, hemophilia, heart-valve defect, MS, Fragile X syndrome, hypertrichosis

Y-linked = SRY gene (absence results in a female XY)

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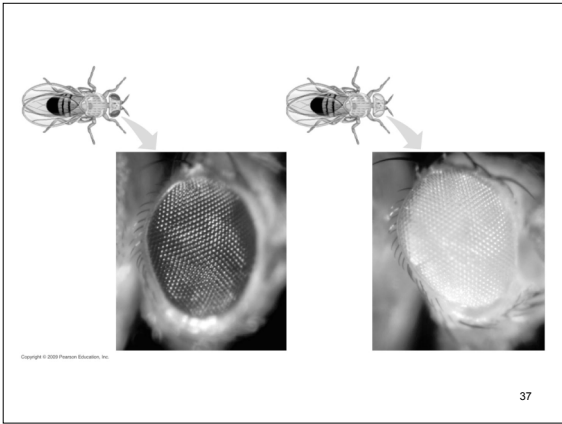
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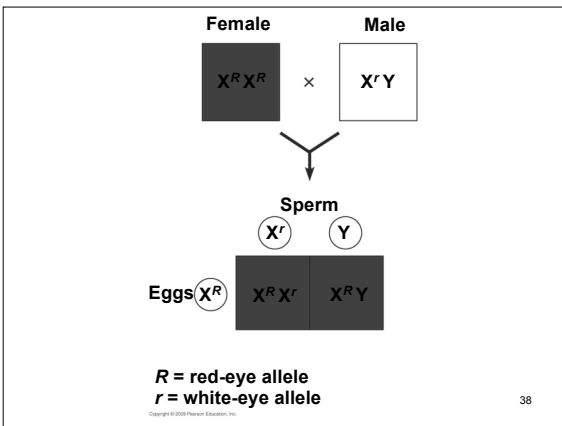
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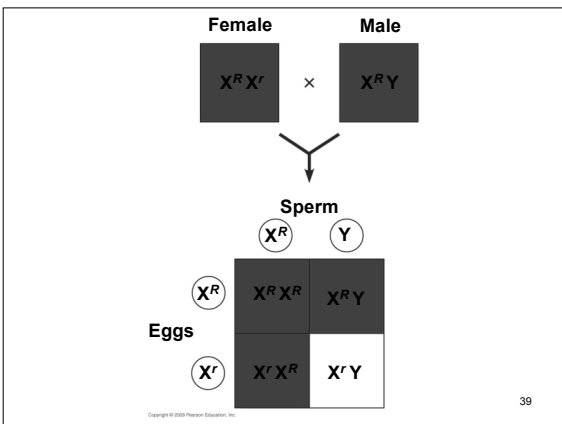
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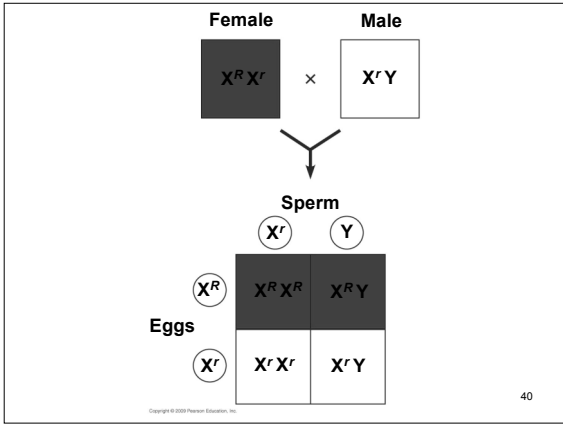
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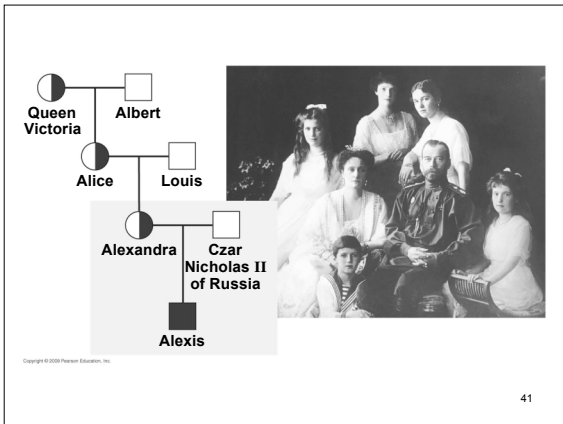
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