5.3 Pedigrees

Tracking patterns of inheritance

Terms and Meanings

Today's Lesson:

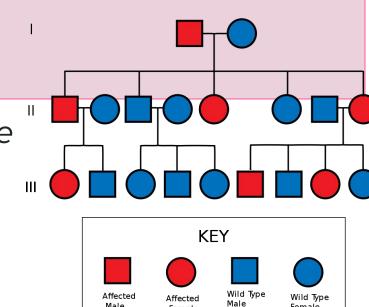
- → Pedigree: a diagram of an individual's ancestors used in human genetics to analyze the Mendelian inheritance of a certain trait; also used for selectively breeding of plants and animals.
- → Autosomal inheritance: inheritance of alleles located on autosomal (non-sex) chromosomes.

Next Lesson:

- → Sex-linked Inheritance: describes an allele that is found on one of the sex chromosomes, X or Y, and when passed on to offspring is expressed.
 - X-linked: phenotypic expression of an allele that is found on the X chromosome.
 - Y-linked: phenotypic expression of an allele that is found on the Y chromosome.

Pedigree

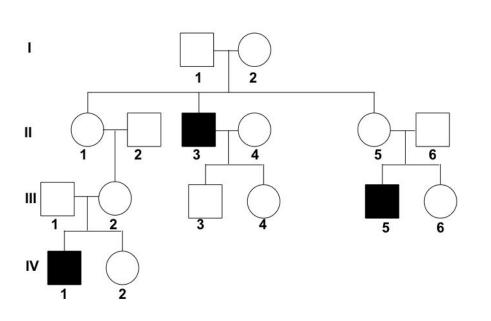
Tracking "bloodlines" can be done using traits other than blood type.



A pedigree: A type of flowchart that uses symbols to show the inheritance patterns of traits in a family over many generations.

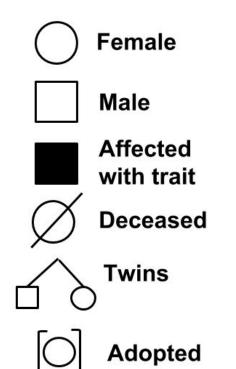
- Used to track breast cancer, Huntington's disease, cystic fibrosis, etc.
- Used in selective breeding of plants and animals.

The Rules

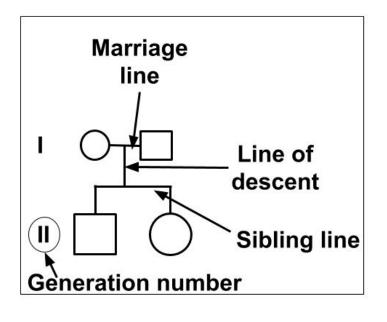


- Generations identified with Roman numerals
- Birth order is left to right (oldest to youngest - if possible)
- Arabic numbers for individuals in a generation
- Affected individual is shaded in; unaffected is un-shaded.

More Symbols

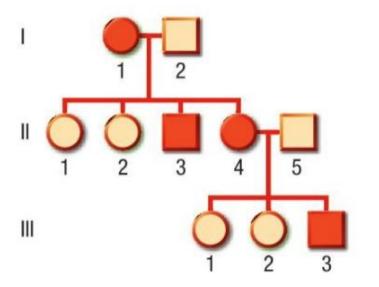


Miscarriage



Reading a Pedigree Chart

- This pedigree shows the presence of freckles in a family.
 - Freckles (F) allele is dominant
 - No freckles (f) is recessive



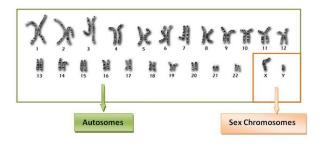
What is the relationship between (II-4) and (II-5)?

Who would be considered the grandmother in this pedigree?

Autosomal Inheritance

Autosomal Inheritance: Inheritance of traits whose

genes are found on the *autosomes *(chromosome sets 1 to 22 in humans)



- Autosomal dominant inheritance requires only one copy of the allele for trait for expression
- Autosomal recessive inheritance requires both copies of the allele for the trait to be present for expression (individuals are carriers if heterozygous)

You can determine if a trait is autosomal dominant or recessive just by looking at a pedigree chart!

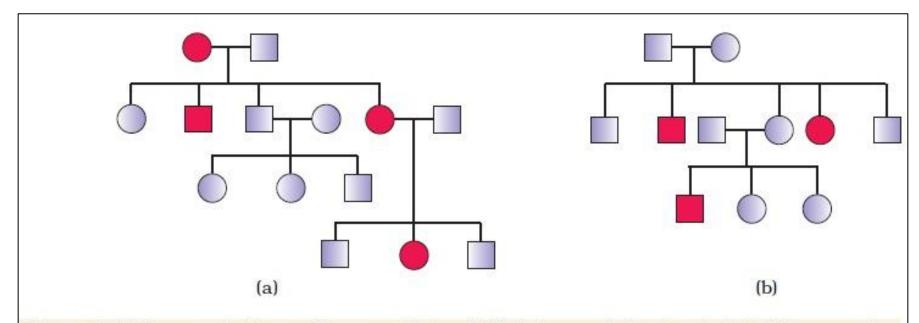
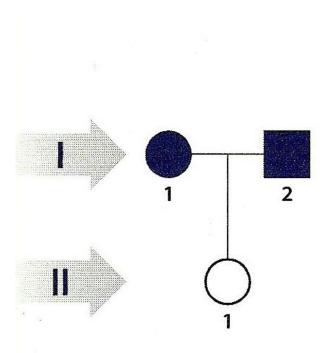
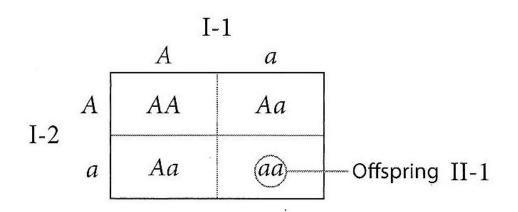


Figure 5.14 Representative pedigree analysis of (a) Autosomal dominant trait (for example: Myotonic dystrophy) (b) Autosomal recessive trait (for example: Sickle-cell anaemia)

Autosomal Dominant Inheritance





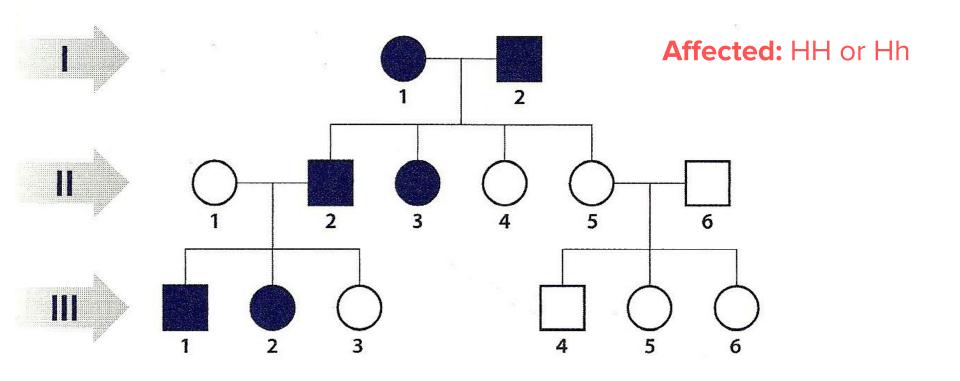
Key

AA = affected

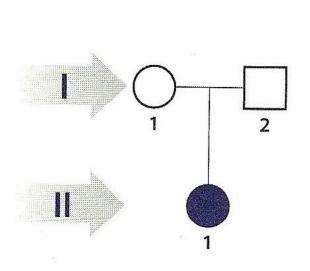
Aa = affected

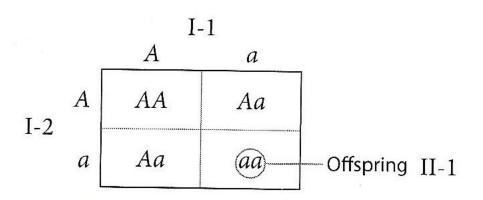
aa = unaffected

Autosomal Dominant Inheritance Huntington Gene



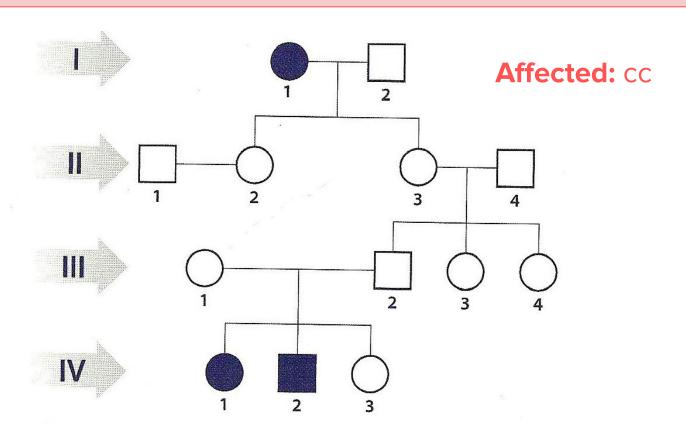
Autosomal Recessive Inheritance

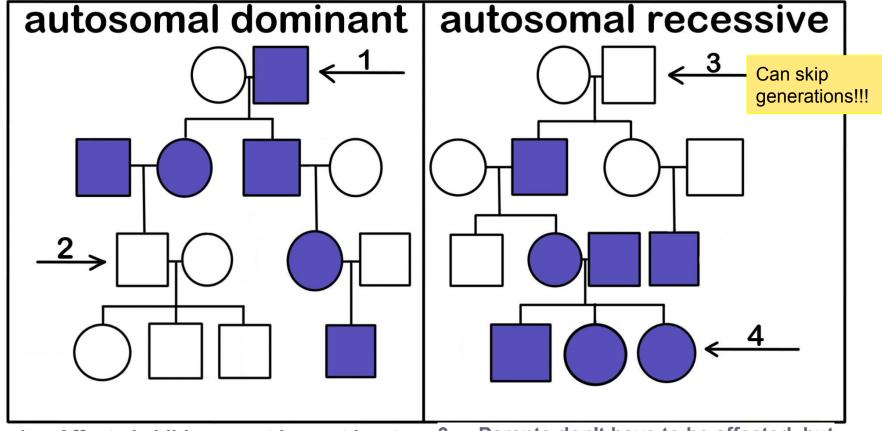




Aa = affected Aa = unaffected (carrier) AA = unaffected

Autosomal Recessive Inheritance Cystic Fibrosis

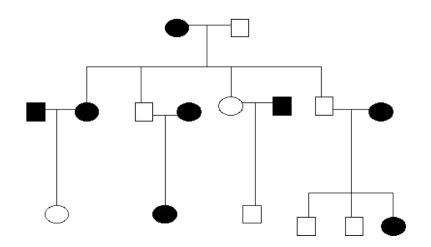


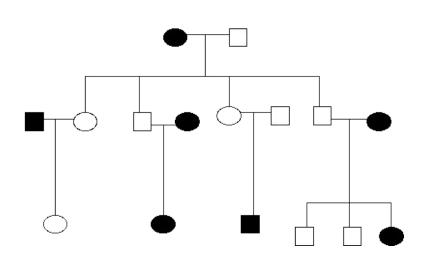


- 1. Affected children must have at least one affected parent.
- 2. Two affected parents can have an unaffected child.

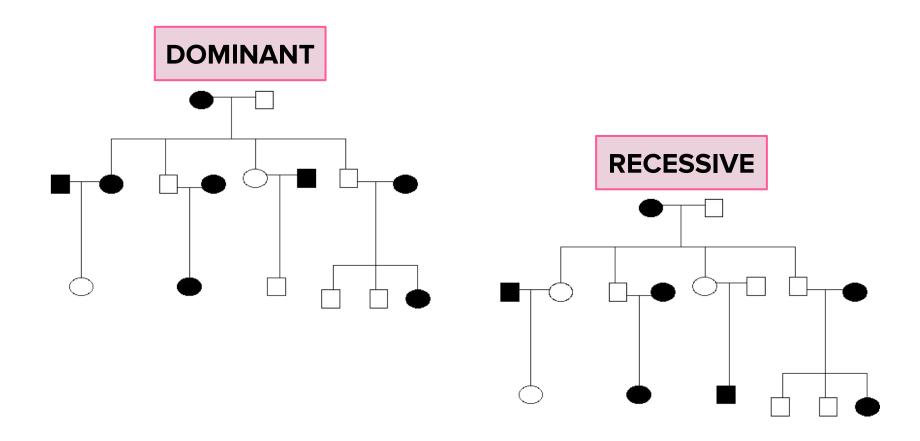
- 3. Parents don't have to be affected, but both must at least be carriers (heterozygous for the trait).
- 4. Two affected parents will only have affected children.

Which is Dominant? Which is Recessive?

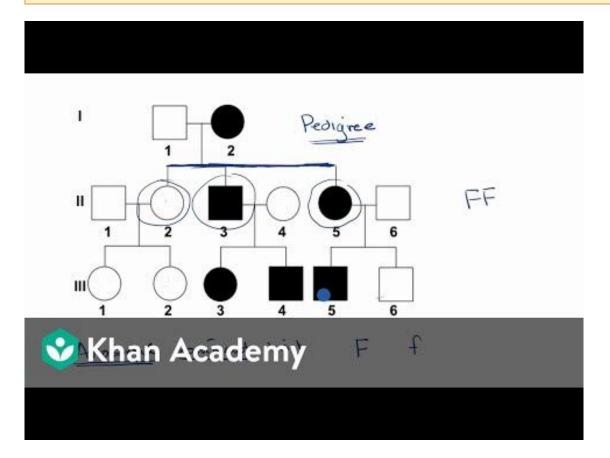




Which is dominant? Which is Recessive?



Practice Reading a Pedigree





Class/Homework

Worksheet Homework pg 201#1-5