LECTURE PRESENTATIONS

For CAMPBELL BIOLOGY, NINTH EDITION

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Chapter 13



Overview: Variations on a Theme

- Living organisms are distinguished by their ability to reproduce their own kind
- Genetics is the scientific study of heredity and variation
- Heredity is the transmission of traits from one generation to the next
- Variation is demonstrated by the differences in appearance that offspring show from parents and siblings

Figure 13.1



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Offspring acquire genes from parents by inheriting chromosomes

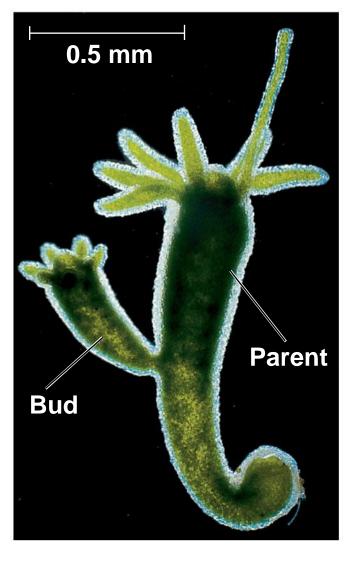
- In a literal sense, children do not inherit particular physical traits from their parents
- It is genes that are actually inherited

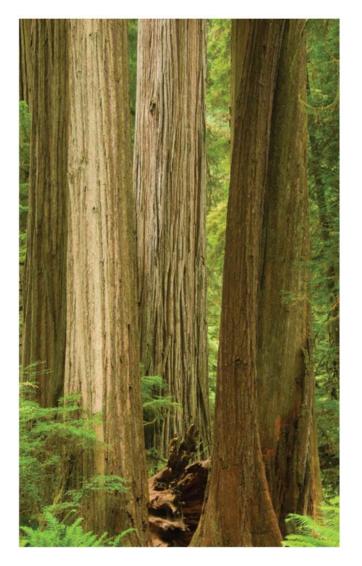
Inheritance of Genes

- Genes are the units of heredity, and are made up of segments of DNA
- Genes are passed to the next generation via reproductive cells called gametes (sperm and eggs)
- Each gene has a specific location called a locus on a certain chromosome
- Most DNA is packaged into chromosomes

Comparison of Asexual and Sexual Reproduction

- In asexual reproduction, a single individual passes genes to its offspring without the fusion of gametes
- In sexual reproduction, two parents give rise to offspring that have unique combinations of genes inherited from the two parents





(a) Hydra
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(b) Redwoods

Asexual Reproduction

Fertilization and meiosis alternate in sexual life cycles

 A life cycle is the generation-to-generation sequence of stages in the reproductive history of an organism

Sets of Chromosomes in Human Cells

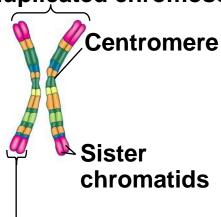
- Human somatic cells (any cell other than a gamete) have 23 pairs of chromosomes
- A karyotype is an ordered display of the pairs of chromosomes from a cell
- The two chromosomes in each pair are called homologous chromosomes, or homologs
- Chromosomes in a homologous pair are the same length and shape and carry genes controlling the same inherited characters

APPLICATION



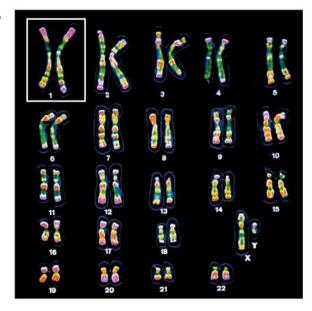
TECHNIQUE

Pair of homologous duplicated chromosomes



Metaphase chromosome

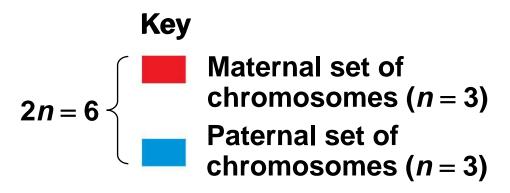
5 μ**m**

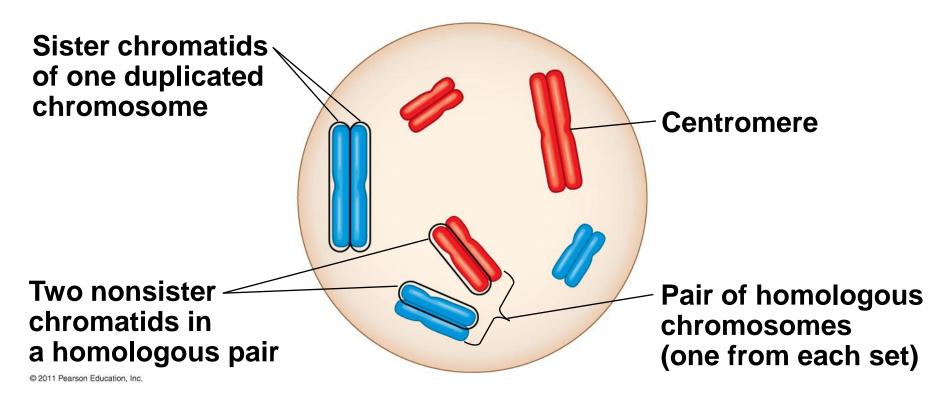


- The sex chromosomes, which determine the sex of the individual, are called X and Y
- Human females have a homologous pair of X chromosomes (XX)
- Human males have one X and one Y chromosome
- The remaining 22 pairs of chromosomes are called autosomes

- Each pair of homologous chromosomes includes one chromosome from each parent
- The 46 chromosomes in a human somatic cell are two sets of 23: one from the mother and one from the father
- A diploid cell (2n) has two sets of chromosomes
- For humans, the diploid number is 46 (2n = 46)

- In a cell in which DNA synthesis has occurred, each chromosome is replicated
- Each replicated chromosome consists of two identical sister chromatids



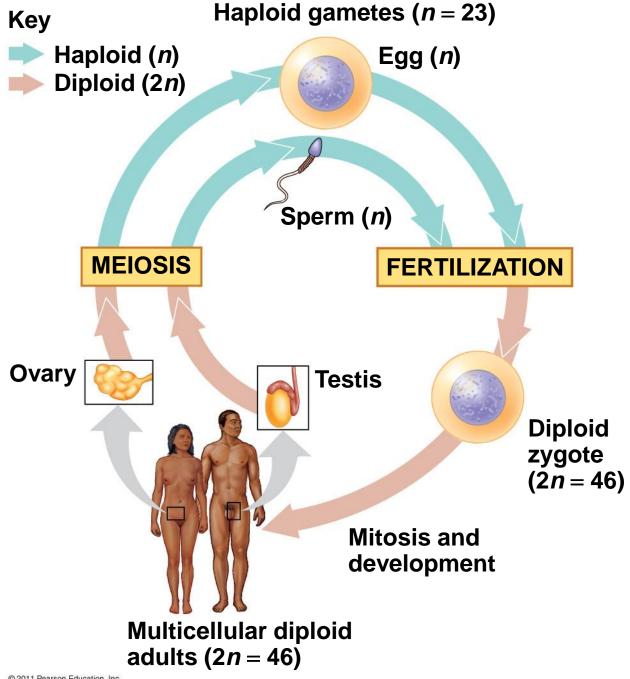


- A gamete (sperm or egg) contains a single set of chromosomes, and is haploid (n)
- For humans, the haploid number is 23 (n = 23)
- Each set of 23 consists of 22 autosomes and a single sex chromosome
- In an unfertilized egg (ovum), the sex chromosome is X
- In a sperm cell, the sex chromosome may be either X or Y

Behavior of Chromosome Sets in the Human Life Cycle

- Fertilization is the union of gametes (the sperm and the egg)
- The fertilized egg is called a zygote and has one set of chromosomes from each parent
- The zygote produces somatic cells by mitosis and develops into an adult

- At sexual maturity, the ovaries and testes produce haploid gametes
- Gametes are the only types of human cells produced by meiosis, rather than mitosis
- Meiosis results in one set of chromosomes in each gamete
- Fertilization and meiosis alternate in sexual life cycles to maintain chromosome number



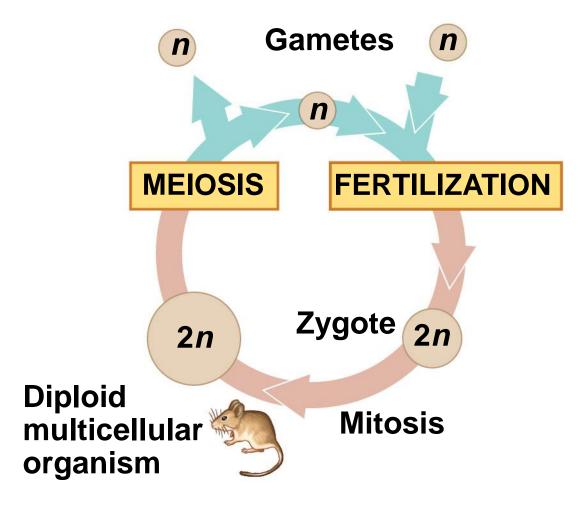
The Variety of Sexual Life Cycles

- The alternation of meiosis and fertilization is common to all organisms that reproduce sexually
- The three main types of sexual life cycles differ in the timing of meiosis and fertilization

- Gametes are the only haploid cells in animals
- They are produced by meiosis and undergo no further cell division before fertilization
- Gametes fuse to form a diploid zygote that divides by mitosis to develop into a multicellular organism

Key

- Haploid (n)
- Diploid (2*n*)



(a) Animals

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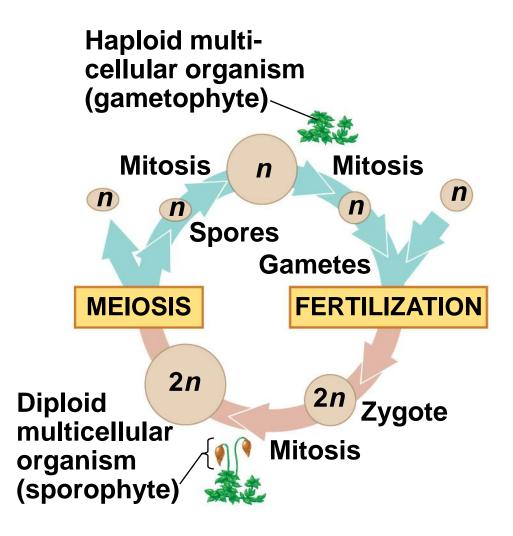
- Plants and some algae exhibit an alternation of generations
- This life cycle includes both a diploid and haploid multicellular stage
- The diploid organism, called the sporophyte, makes haploid spores by meiosis

- Each spore grows by mitosis into a haploid organism called a gametophyte
- A gametophyte makes haploid gametes by mitosis
- Fertilization of gametes results in a diploid sporophyte

Key

Haploid (n)

Diploid (2*n*)



(b) Plants and some algae

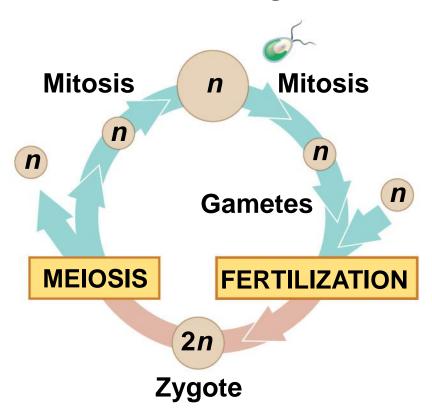
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- In most fungi and some protists, the only diploid stage is the single-celled zygote; there is no multicellular diploid stage
- The zygote produces haploid cells by meiosis
- Each haploid cell grows by mitosis into a haploid multicellular organism
- The haploid adult produces gametes by mitosis

Key

- Haploid (n)
- Diploid (2*n*)

Haploid unicellular or multicellular organism



(c) Most fungi and some protists

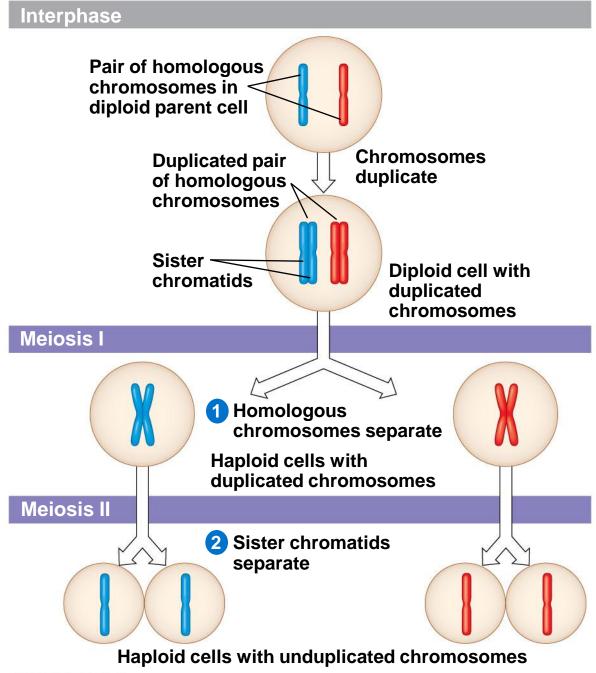
- Depending on the type of life cycle, either haploid or diploid cells can divide by mitosis
- However, only diploid cells can undergo meiosis
- In all three life cycles, the halving and doubling of chromosomes contributes to genetic variation in offspring

Meiosis reduces the number of chromosome sets from diploid to haploid

- Like mitosis, meiosis is preceded by the replication of chromosomes
- Meiosis takes place in two sets of cell divisions, called meiosis I and meiosis II
- The two cell divisions result in four daughter cells, rather than the two daughter cells in mitosis
- Each daughter cell has only half as many chromosomes as the parent cell

The Stages of Meiosis

- After chromosomes duplicate, two divisions follow
 - Meiosis I (reductional division): homologs pair up and separate, resulting in two haploid daughter cells with replicated chromosomes
 - Meiosis II (equational division) sister chromatids separate
- The result is four haploid daughter cells with unreplicated chromosomes



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- Meiosis I is preceded by interphase, when the chromosomes are duplicated to form sister chromatids
- The sister chromatids are genetically identical and joined at the centromere
- The single centrosome replicates, forming two centrosomes

- Division in meiosis I occurs in four phases
 - Prophase I
 - Metaphase I
 - Anaphase I
 - Telophase I and cytokinesis

Prophase I

- Prophase I typically occupies more than 90% of the time required for meiosis
- Chromosomes begin to condense
- In synapsis, homologous chromosomes loosely pair up, aligned gene by gene

- In crossing over, nonsister chromatids exchange DNA segments
- Each pair of chromosomes forms a tetrad, a group of four chromatids
- Each tetrad usually has one or more chiasmata, X-shaped regions where crossing over occurred

Metaphase I

- In metaphase I, tetrads line up at the metaphase plate, with one chromosome facing each pole
- Microtubules from one pole are attached to the kinetochore of one chromosome of each tetrad
- Microtubules from the other pole are attached to the kinetochore of the other chromosome

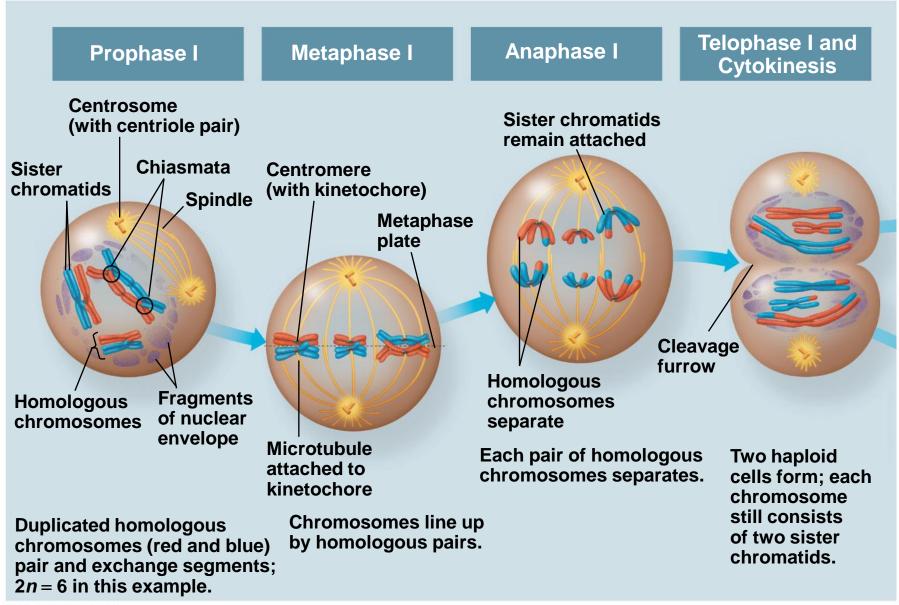
Anaphase I

- In anaphase I, pairs of homologous chromosomes separate
- One chromosome moves toward each pole, guided by the spindle apparatus
- Sister chromatids remain attached at the centromere and move as one unit toward the pole

Telophase I and Cytokinesis

- In the beginning of telophase I, each half of the cell has a haploid set of chromosomes; each chromosome still consists of two sister chromatids
- Cytokinesis usually occurs simultaneously, forming two haploid daughter cells

- In animal cells, a cleavage furrow forms; in plant cells, a cell plate forms
- No chromosome replication occurs between the end of meiosis I and the beginning of meiosis II because the chromosomes are already replicated



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- Division in meiosis II also occurs in four phases
 - Prophase II
 - Metaphase II
 - Anaphase II
 - Telophase II and cytokinesis
- Meiosis II is very similar to mitosis

Prophase II

- In prophase II, a spindle apparatus forms
- In late prophase II, chromosomes (each still composed of two chromatids) move toward the metaphase plate

Metaphase II

- In metaphase II, the sister chromatids are arranged at the metaphase plate
- Because of crossing over in meiosis I, the two sister chromatids of each chromosome are no longer genetically identical
- The kinetochores of sister chromatids attach to microtubules extending from opposite poles

Anaphase II

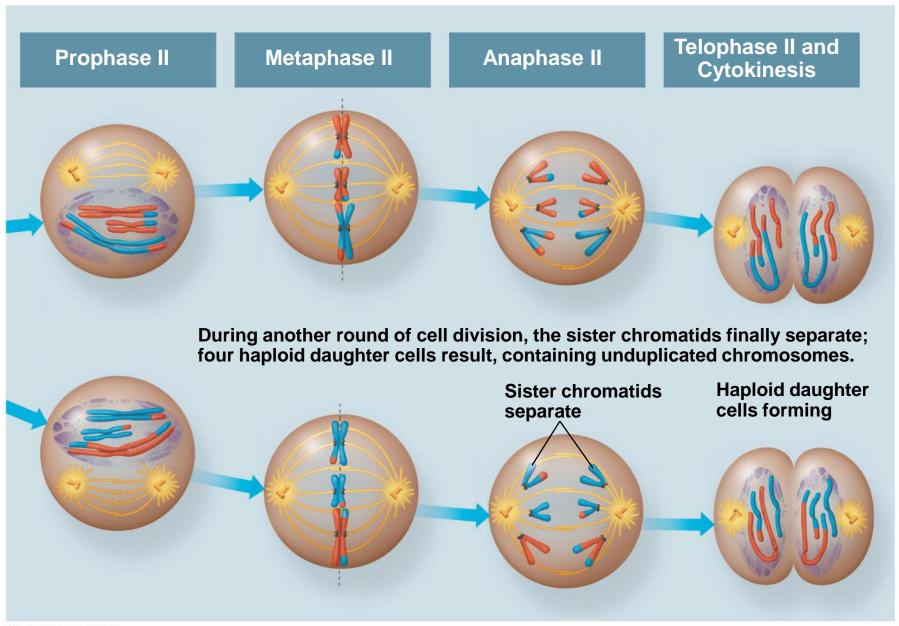
- In anaphase II, the sister chromatids separate
- The sister chromatids of each chromosome now move as two newly individual chromosomes toward opposite poles

Telophase II and Cytokinesis

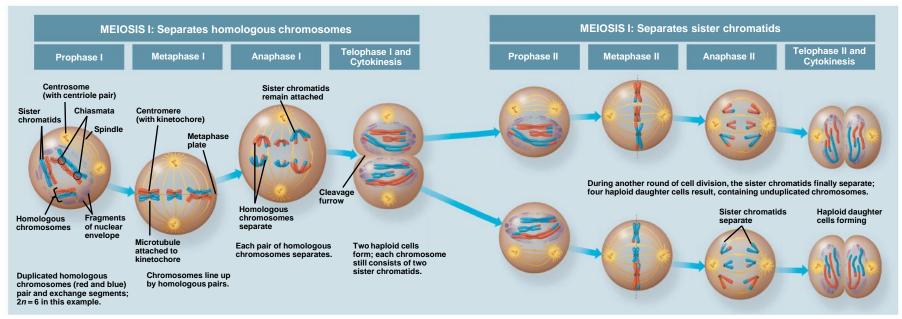
- In telophase II, the chromosomes arrive at opposite poles
- Nuclei form, and the chromosomes begin decondensing

- Cytokinesis separates the cytoplasm
- At the end of meiosis, there are four daughter cells, each with a haploid set of unreplicated chromosomes
- Each daughter cell is genetically distinct from the others and from the parent cell

Figure 13.8b



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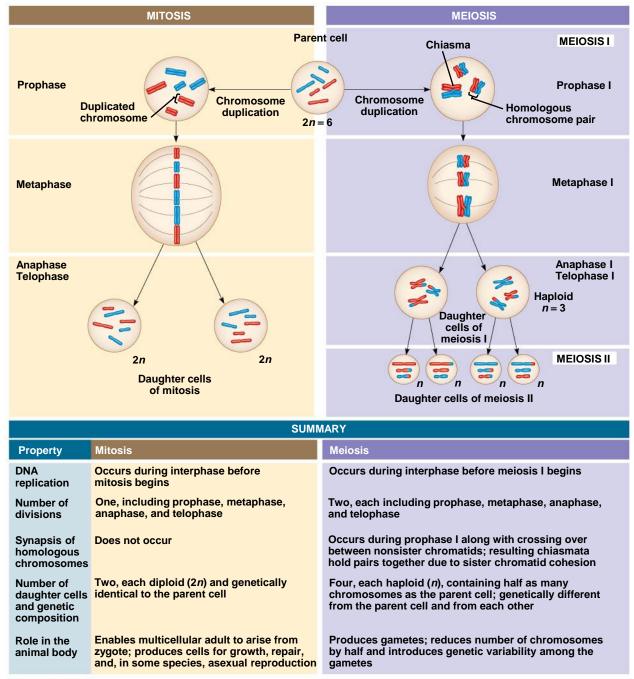


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A Comparison of Mitosis and Meiosis

- Mitosis conserves the number of chromosome sets, producing cells that are genetically identical to the parent cell
- Meiosis reduces the number of chromosomes sets from two (diploid) to one (haploid), producing cells that differ genetically from each other and from the parent cell

Figure 13.9



- Three events are unique to meiosis, and all three occur in meiosis I
 - Synapsis and crossing over in prophase I:
 Homologous chromosomes physically connect and exchange genetic information
 - At the metaphase plate, there are paired homologous chromosomes (tetrads), instead of individual replicated chromosomes
 - At anaphase I, it is homologous chromosomes, instead of sister chromatids, that separate

Genetic variation produced in sexual life cycles contributes to evolution

- Mutations (changes in an organism's DNA) are the original source of genetic diversity
- Mutations create different versions of genes called alleles
- Reshuffling of alleles during sexual reproduction produces genetic variation

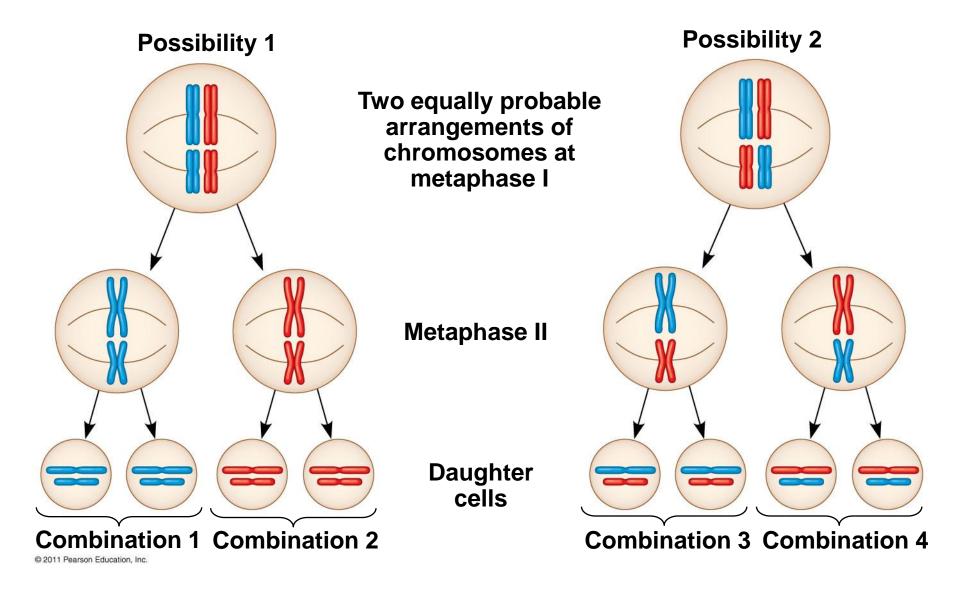
Origins of Genetic Variation Among Offspring

- The behavior of chromosomes during meiosis and fertilization is responsible for most of the variation that arises in each generation
- Three mechanisms contribute to genetic variation
 - Independent assortment of chromosomes
 - Crossing over
 - Random fertilization

Independent Assortment of Chromosomes

- Homologous pairs of chromosomes orient randomly at metaphase I of meiosis
- In independent assortment, each pair of chromosomes sorts maternal and paternal homologues into daughter cells independently of the other pairs

- The number of combinations possible when chromosomes assort independently into gametes is 2ⁿ, where n is the haploid number
- For humans (n = 23), there are more than 8 million (2^{23}) possible combinations of chromosomes

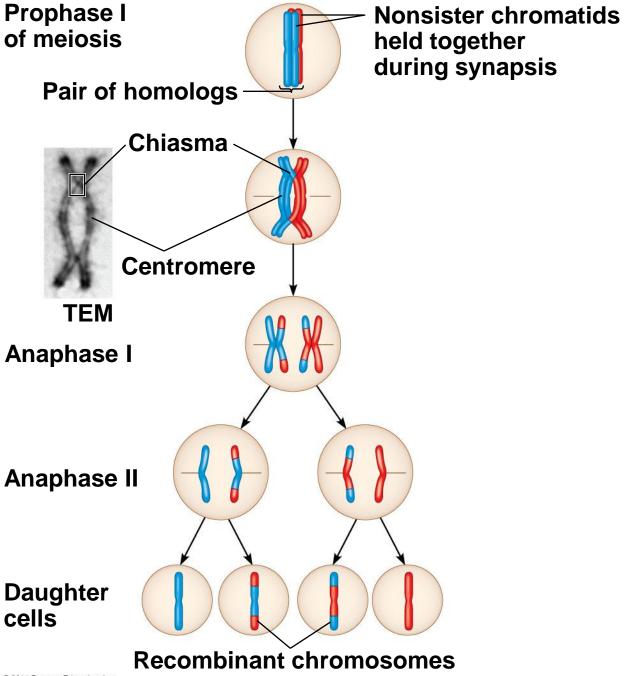


Crossing Over

- Crossing over produces recombinant chromosomes, which combine DNA inherited from each parent
- Crossing over begins very early in prophase I, as homologous chromosomes pair up gene by gene

- In crossing over, homologous portions of two nonsister chromatids trade places
- Crossing over contributes to genetic variation by combining DNA from two parents into a single chromosome

Figure 13.11-5



Random Fertilization

- Random fertilization adds to genetic variation because any sperm can fuse with any ovum (unfertilized egg)
- The fusion of two gametes (each with 8.4 million possible chromosome combinations from independent assortment) produces a zygote with any of about 70 trillion diploid combinations
- Crossing over adds even more variation
- Each zygote has a unique genetic identity

The Evolutionary Significance of Genetic Variation Within Populations

- Natural selection results in the accumulation of genetic variations favored by the environment
- Sexual reproduction contributes to the genetic variation in a population, which originates from mutations