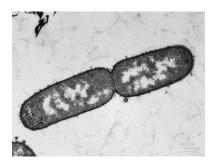
Meiosis and Sexual Reproduction

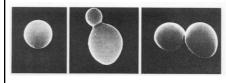
- I. Reproduction: new individual
 - sexual: by union (fertilization) of gametes 配子 offspring are genetically different from the parents
 - asexual: not involve the union of gametes
 offspring are genetically identical to the parent

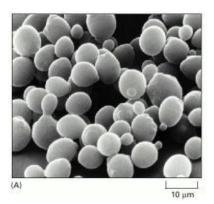
1

- I. Asexual reproduction
 - unicellular (single-cell) microorganisms
 - -> cell division -> reproduction e.g., bacteria, yeast



The yeast Saccharomyces cerevisiae.





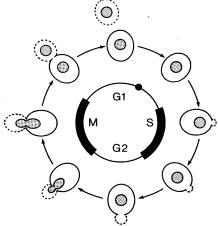
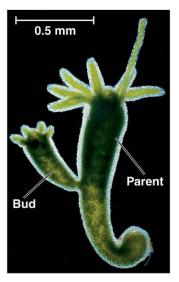


FIG. 1. S. cerevisiae mitotic cell cycle. The phases of the cell cycle are drawn in approximate proportion to their length. The mother cell is drawn with a solid line; the daughter bud and cell are drawn with a dotted line. The shaded material represents the cell nucleus. S. DNA synthesis; M, mitosis (nuclear division). The circle within G1 indicates the position at which yeast cells are arrested by mating factors. (Modified from reference 120 with permission.)

3

- -multicellular organisms
 - -> multiple cell division
 - -> differentiation 分化
 - -> individual
 - -> reproduction
 - e.g., Hydra水螅, plants



(a) Hydra

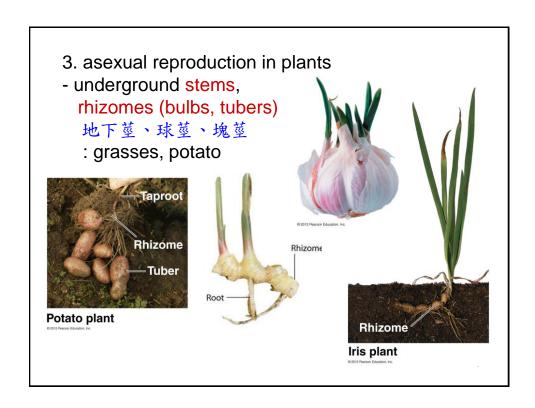
asexual reproduction in plants

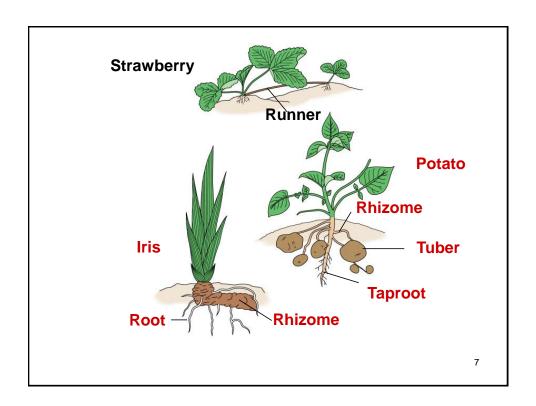
- stems above ground (runners or stolons)

走莖、匍匐莖

: strawberry





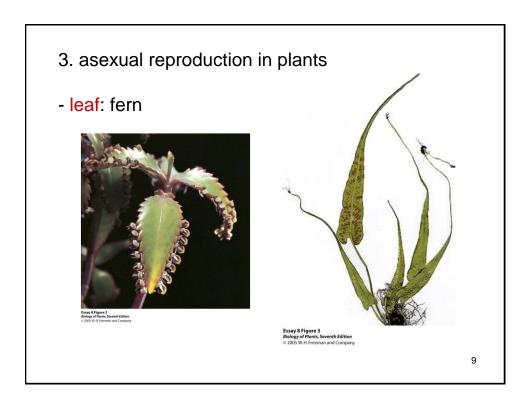


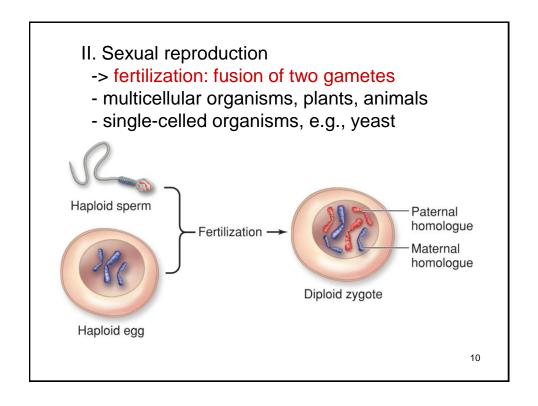
3. asexual reproduction in plants

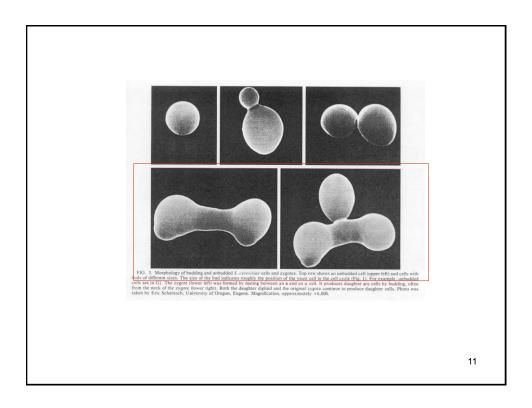
- roots: redwood, creosote bushes

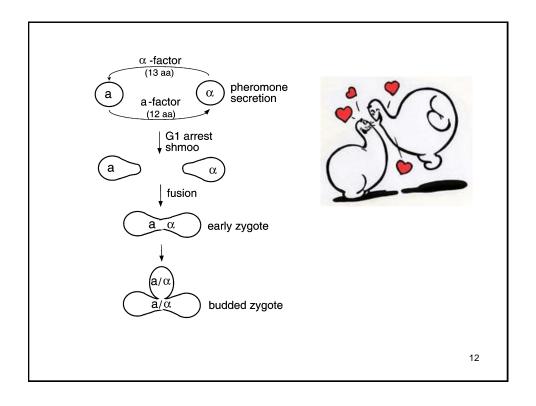


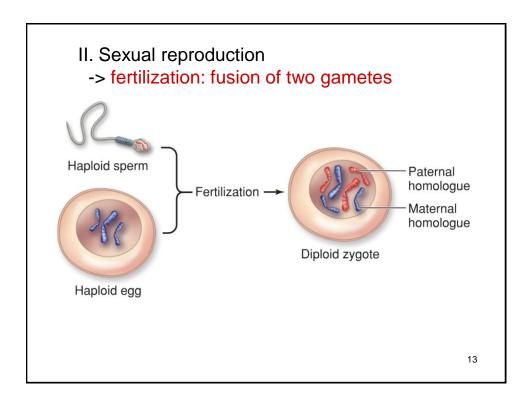












II. Sexual reproduction

- -> fertilization: fusion of two gametes
 - -> combination of genetic materials
- cause constant increase of chromosome number?

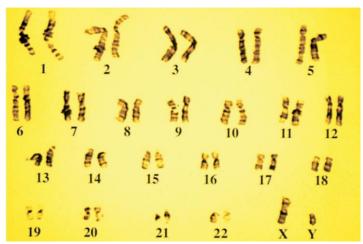
```
generation 1 1+1->2
generation 2 2+2->4
generation 3 4+4->8
generation 4 8+8->16
```

- every species has a fixed number of chromosomes in their somatic cells體細胞 and sex cells生殖細胞, respectively

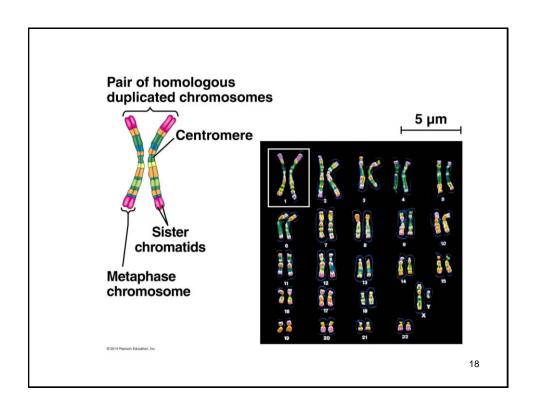
human: 46 / 23 maize: 20 / 10

15

- chromosomes vary in size, location of the centromere
- -> array of an individual's chromosomes (metaphase)
 - => karyotype 核型



- 1). haploid and diploid
- a. diploid
- somatic cells contain 2 copies (1 pair) of each particular chromosome, similar in appearance (size, centromere position, staining pattern)
- => homologous chromosomes (homologs) 同源染色體
- -> carry the same set of genetic information (genes)



- 1). haploid and diploid
 - somatic cells -> diploid cells 二倍體細胞
 - the number of total chromosomes
 - => diploid number (2n),

fixed for each species and constant form generation to generation

human somatic cell:

46 chromosomes (2n=46, n=23)

44 autosomes, 2 sex chromosomes: XX or XY

體染色體 性染色體

23 pairs of homologous chromosomes

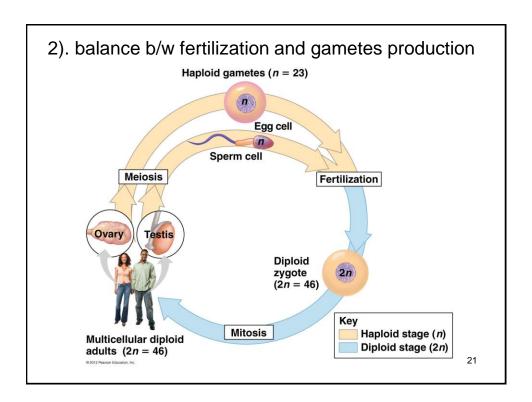
19

b. haploid

- sex cells (gametes) contain only 1 copy of each pair of homologous chromosomes
- the number of chromosomes is half of the somatic cells

```
=> haploid number (n)
sex cells -> haploid cells 單倍體細胞
human sperm or egg:
```

23 chromosomes haploid number n=23



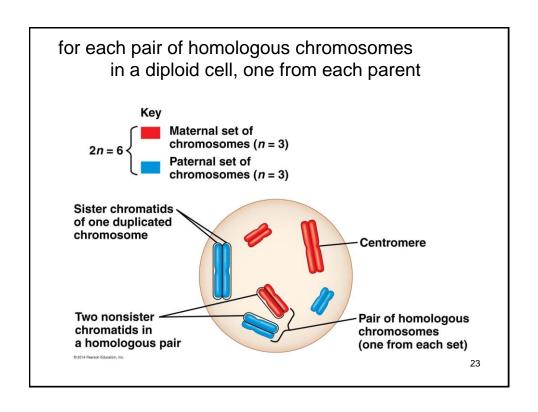
2). balance b/w fertilization and meiosis

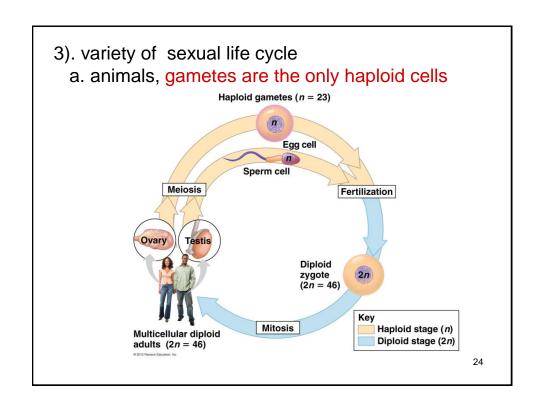
meiosis: the process to produce haploid gametes

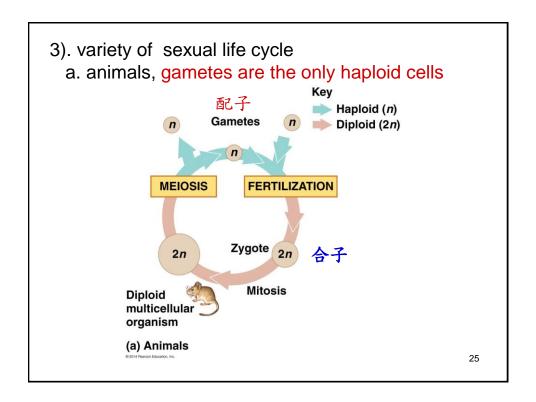
減數分裂 from a diploid cell

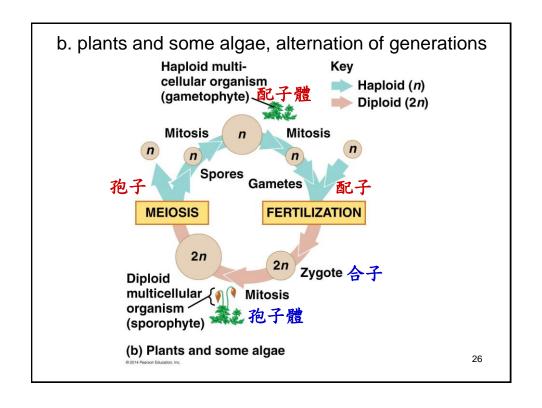
diploid cell --> gametes 2n 1n

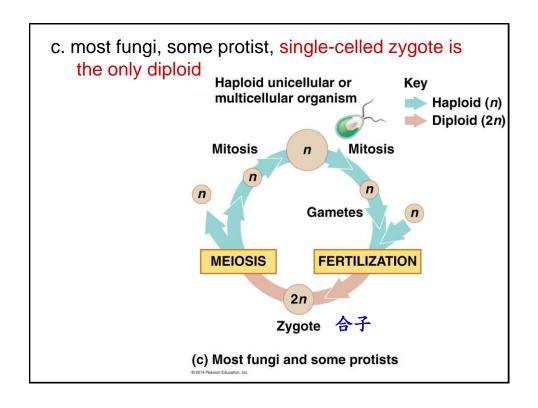
fertilization: the fusion of two haploid gamete to form a diploid zygote

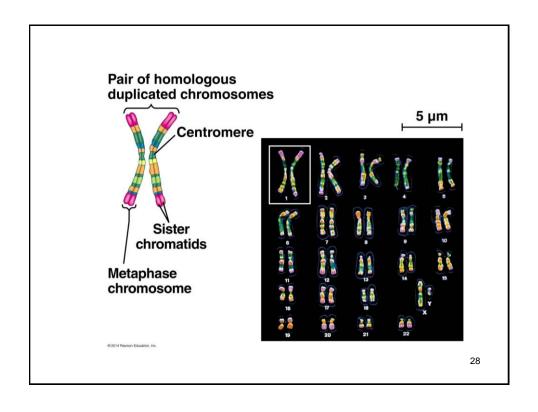












II. Meiosis

1). key points

A. a diploid cell (2n)

-> 1x chromosome duplication

-> 2x nuclear divisions (chromosome segregations)

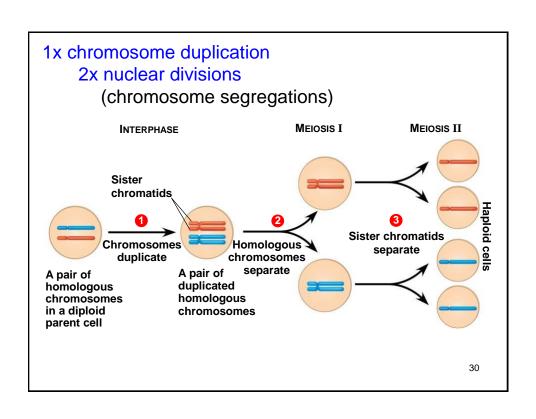
meiosis I: separation of homologous

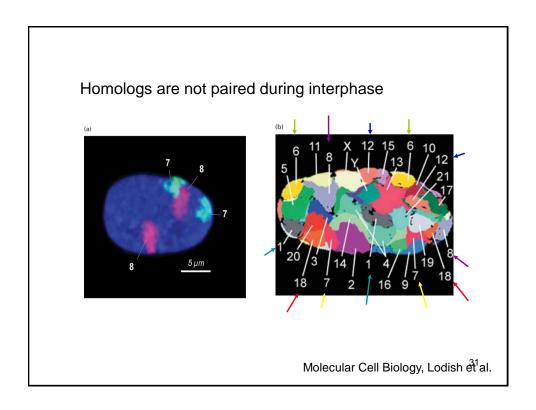
chromosomes

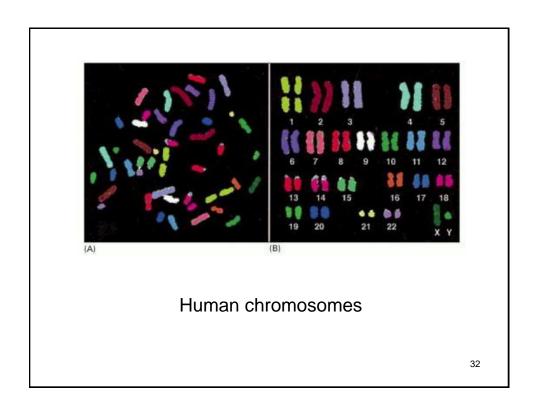
meiosis II: separation of

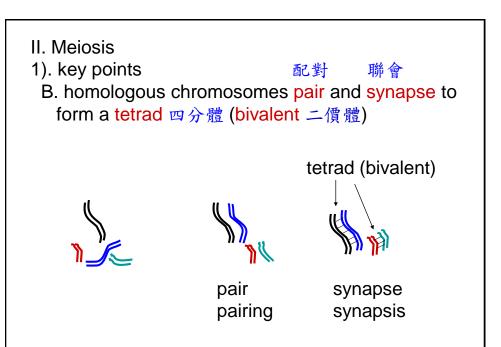
sister chromatids (= mitosis)

-> 4 daughter cells (1n)









key events:

- homologs pairing and synapsis

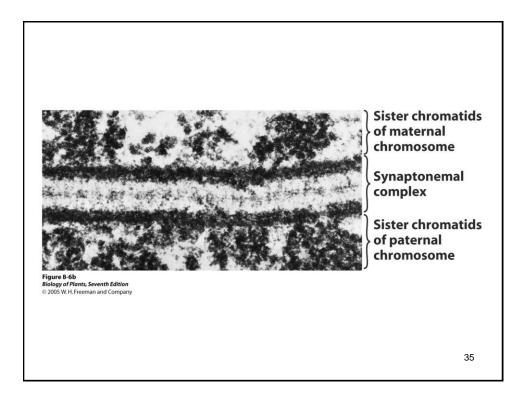
(via synaptonemal complex)

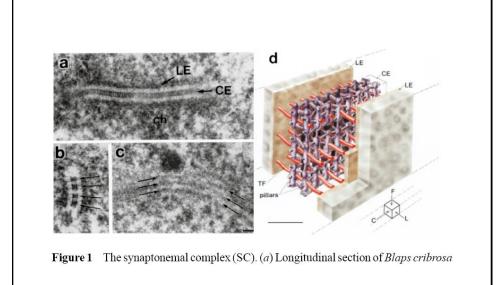
聯會複合體

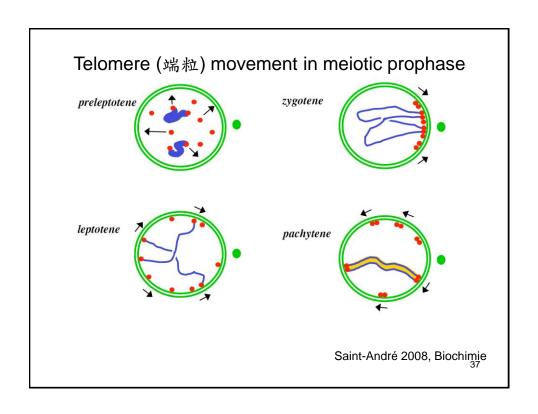
-> form a bivalent (tetrad), as one unit

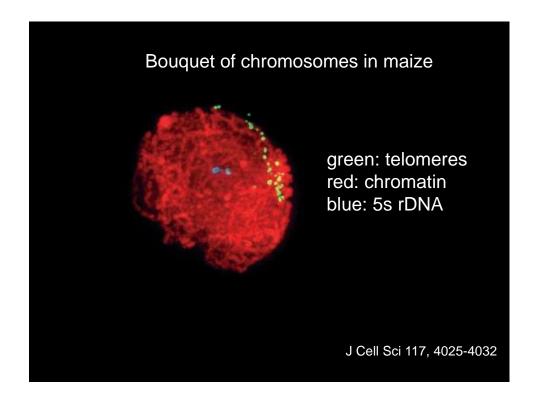
pairing ≠ synapsis
close together connected by a structure

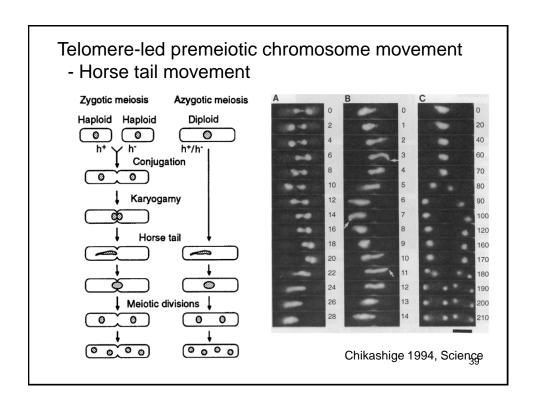
34

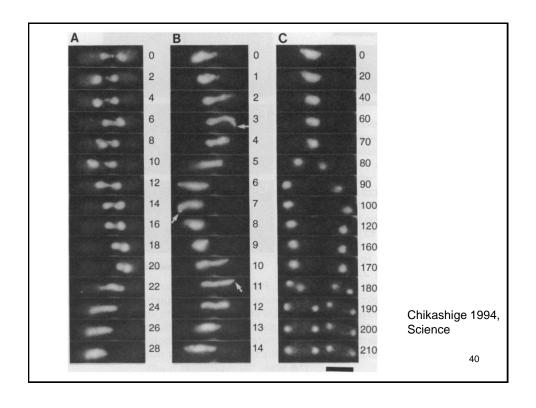




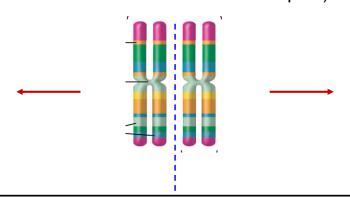


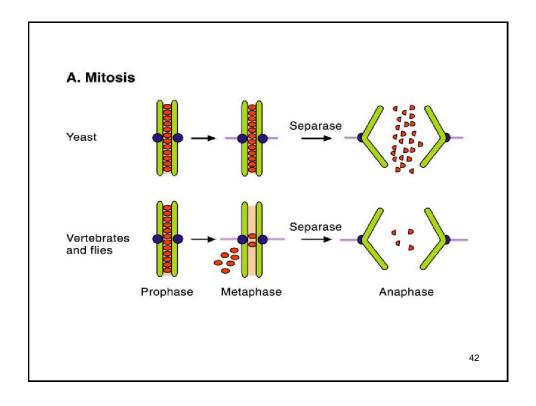


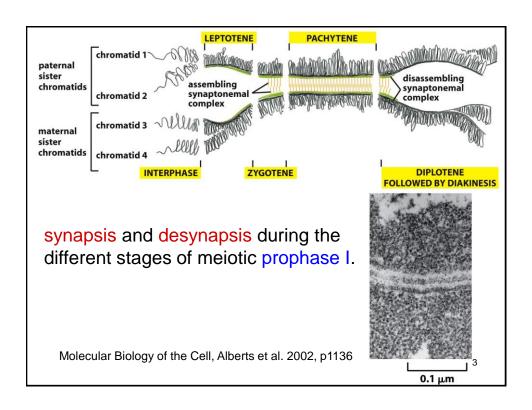


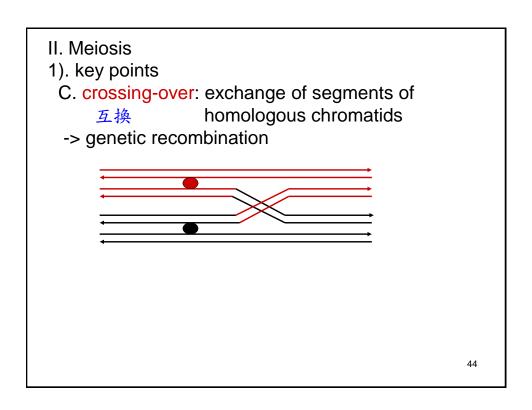


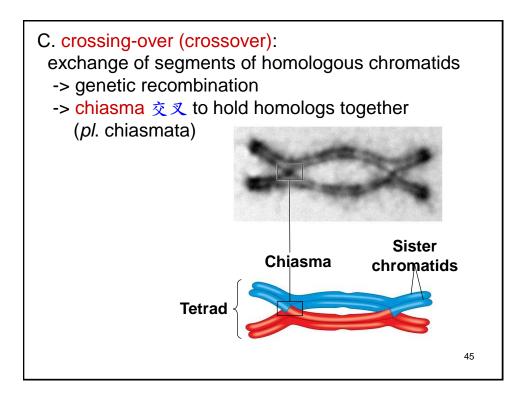
- II. Meiosis
- 1). key points
- B. homologous chromosomes pair and synapse to form a tetrad -> as one unit at prophase I and metaphase I
- -> homologous chromosomes separate at meiosis I, (sister chromatids move to the same pole)

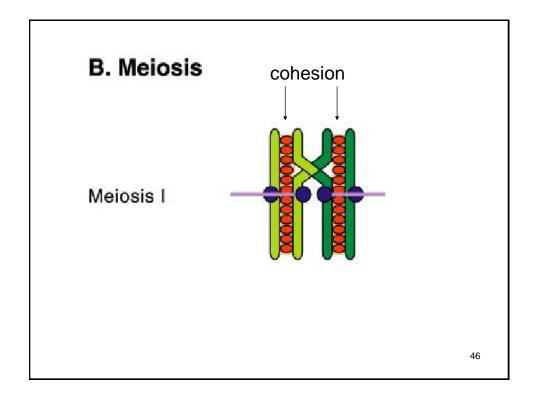


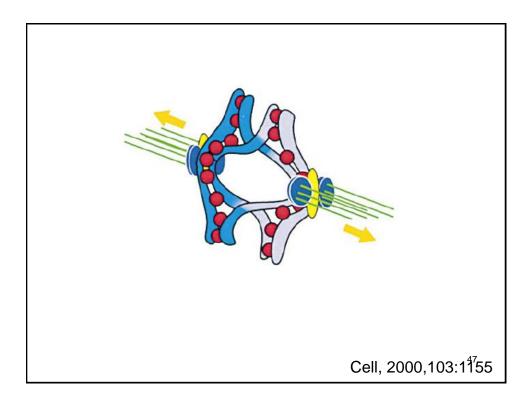












- II. Meiosis
- 1). key points
- A. a diploid cell (2n)
 - -> 1x chromosome duplication
 - -> 2x nuclear divisions (chromosome segregations)
- B. homologous chromosomes pair and synapse to form a tetrad -> as one unit at prophase I and metaphase I
- C. crossing-over: exchange of segments of homologous chromatids
 - -> genetic recombination
 - chiasma to hold homologous chromosomes together

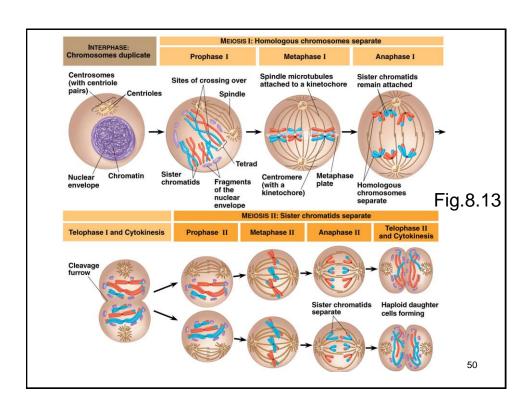
2). stages:

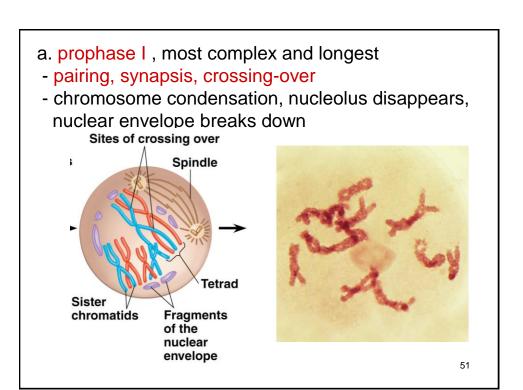
DNA replication

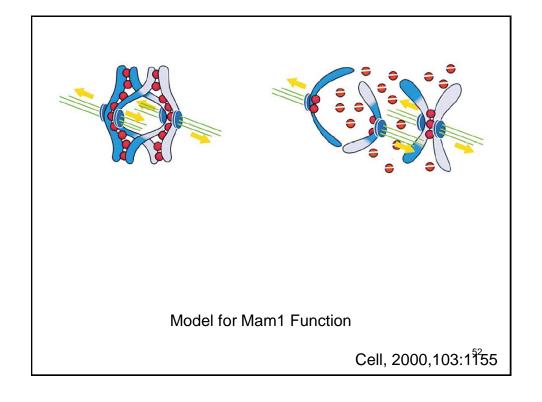
meiosis I

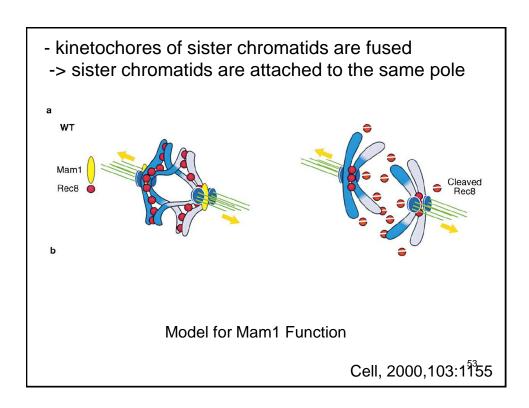
- a. prophase I, most complex and longest
- b. metaphase I
- c. anaphase I
- d. telophase I (+ cytokinesis in some species)

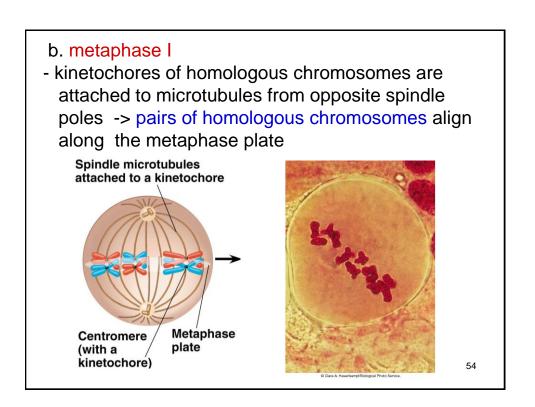
meiosis II, similar to mitosis

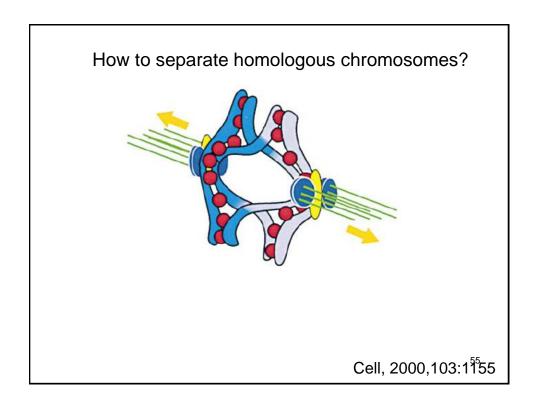


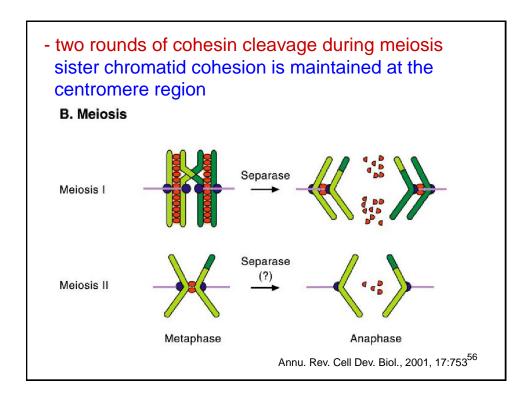


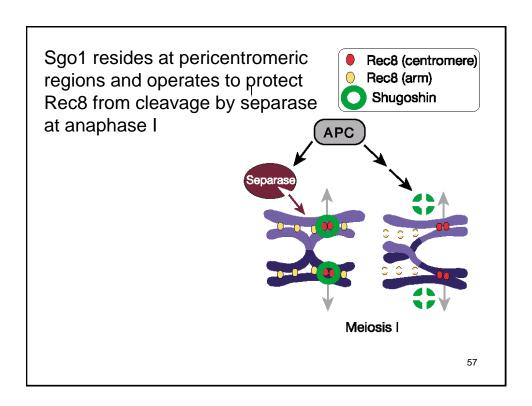


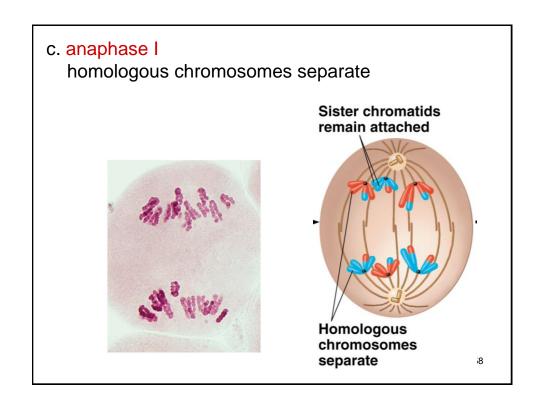


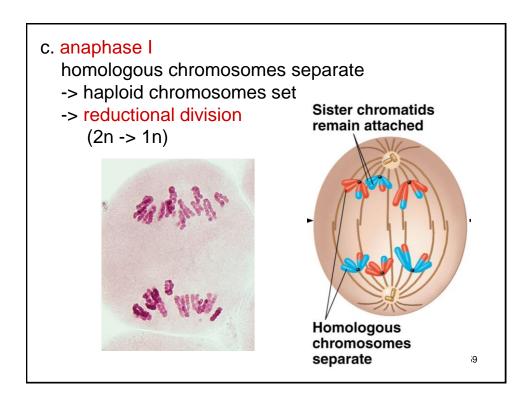


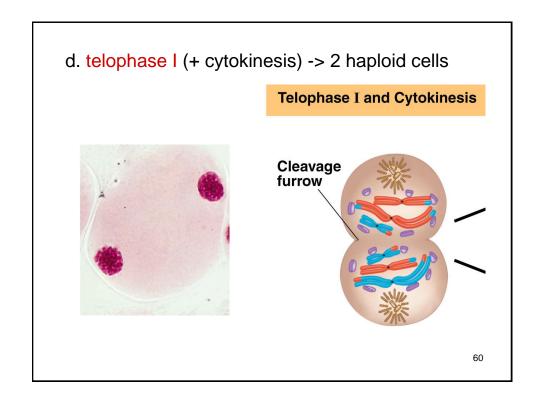




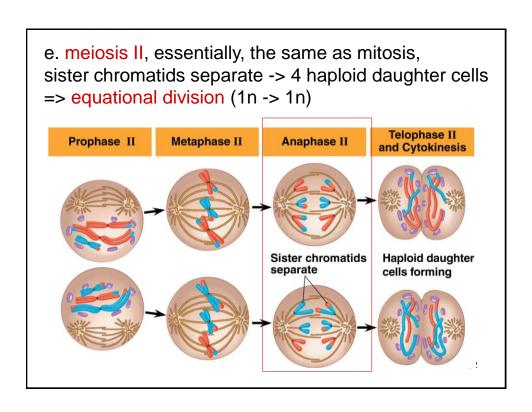


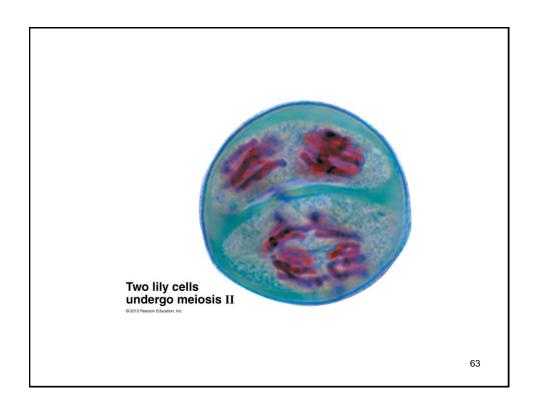


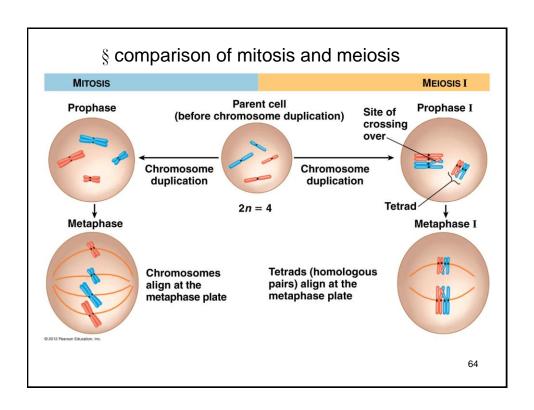


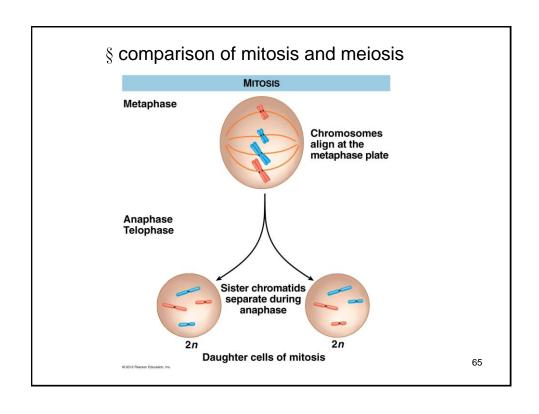


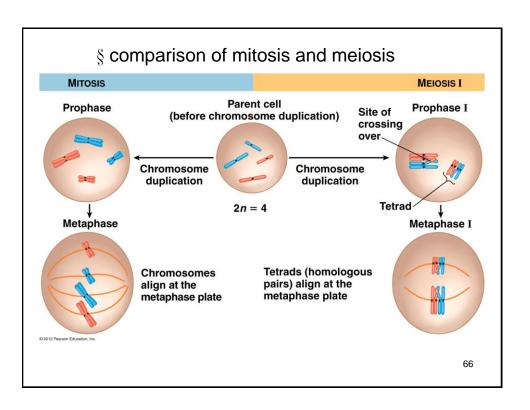
- d. telophase I (+ cytokinesis) -> 2 haploid cells
- § continue meiosis II, or enter into interphase
- § no DNA replication before meiosis II

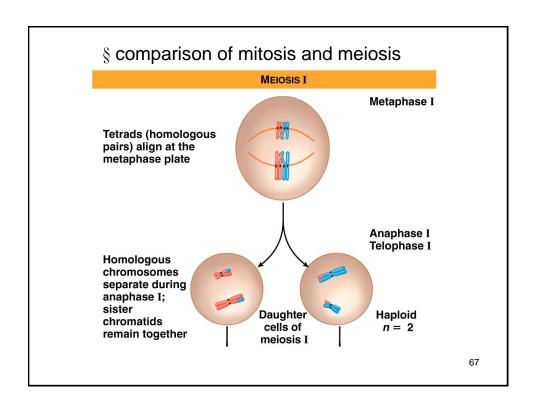


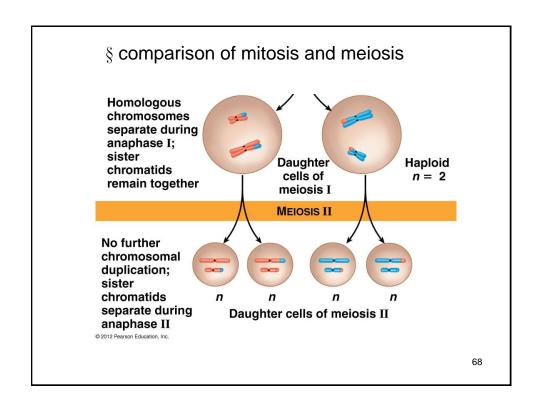








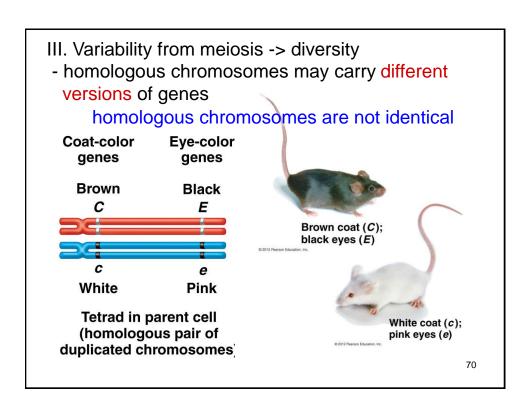


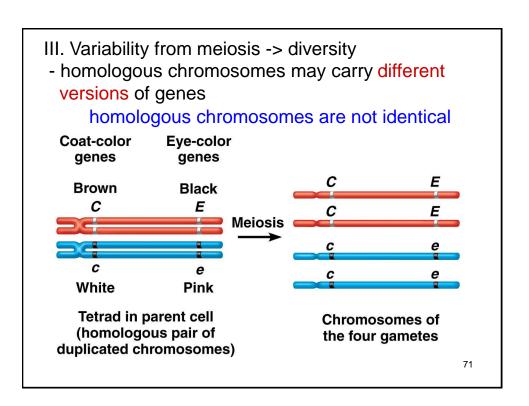


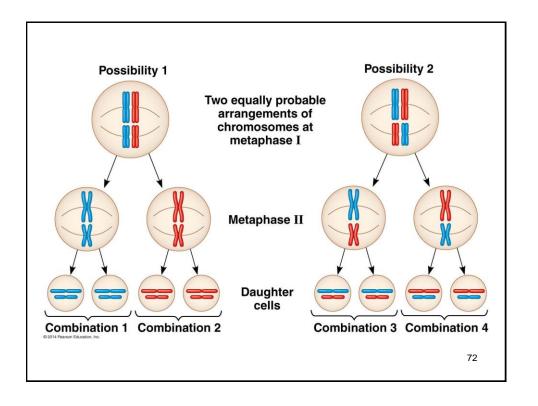
§ comparison of mitosis and meiosis

mitosis produces daughter cells genetically identical to their parent cell and to each other

meiosis produces cells genetically different from their parent cell and to each other

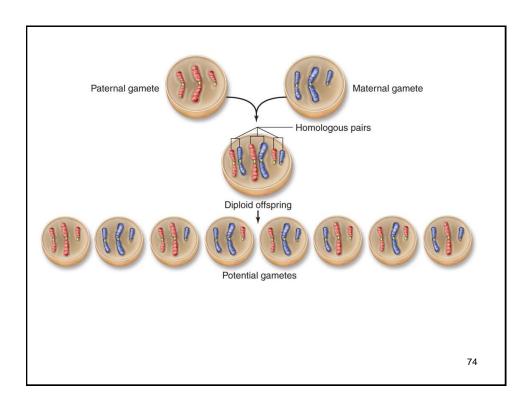






homologous chromosomes are not identical

- III. Variability from meiosis --> diversity
- independent assortment
 when ≥ 2 pairs of homologous chromosomes are
 considered
- random orientation of tetrads at metaphase I
- random combination of chromosomes after meiosis I



1. independent assortment

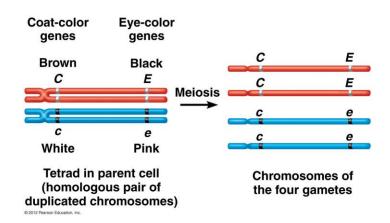
when ≥ 2 pairs of homologous chromosomes are considered

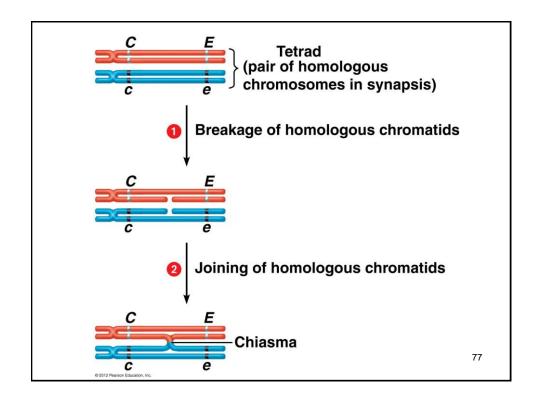
- random orientation of tetrads at metaphase I
- random combination of chromosomes after meiosis I
- possible combination of chromosomes in gametes 2^n , n = haploid number

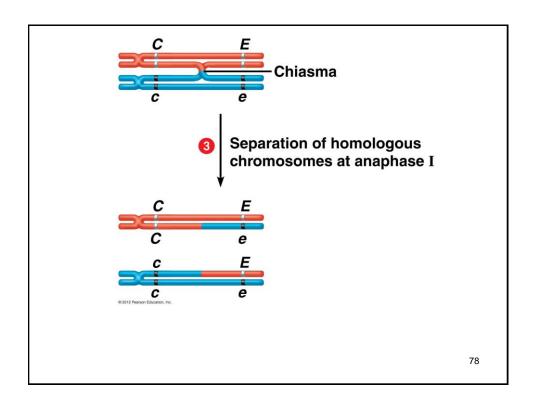
human: $2^{23} = 8,388,608$

75

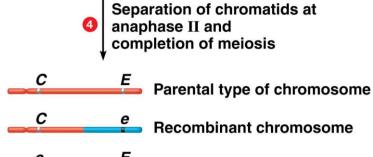
2). crossing-over: genetic recombination, adds more variety in meiosis







2). crossing-over: genetic recombination, adds more variety in meiosis



c E Recombinant chromosome

c e Parental type of chromosome

Gametes of four genetic types

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- 3). random fertilization
 - -> more variability for sexual reproduction
 - fertilization
 - -> random combination of sperms and eggs

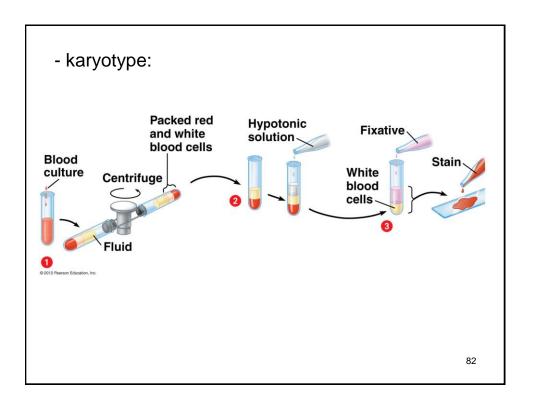
of possible zygote
= (possible # of sperm) x (possible # of egg)

human: $2^{23} \times 2^{23} = 2^{46} \approx 7 \times 10^{13}$

You are unique!

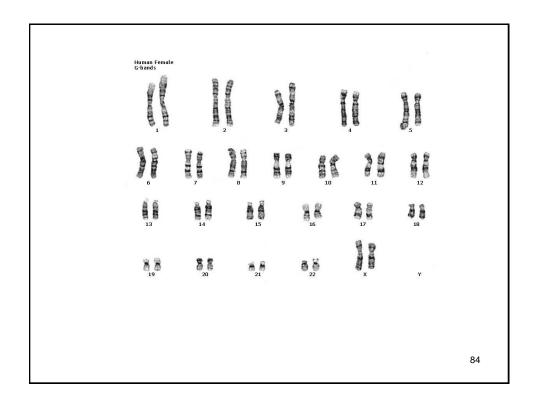
IV. Alterations of chromosome number and structure

- karyotype:
 - a display of microphotographs of metaphase chromosomes of a cell, arranged by size and centromere position, often used lymphocyte (a type of white blood cell), pre-grow and arrest at metaphase
- -> an overview of a person's genome
- -> to screen for abnormal numbers of chromosomes and for defective chromosomes



- karyotype:





1). abnormal numbers of chromosomes

-> aneuploid (非整倍性)

2n-1 -> monosomy (e.g., XO)
2n+1 -> trisomy, most spontaneous abortions, few live-born (e.g., trisomy 8, 13, 18, 21, Y, or X),

85

- a. trisomy 21 (Down syndrome)
- most common serious birth defect 1 in 850 (USA)



Trisomy 21



A person with Down syndrome

- a. trisomy 21 (Down syndrome)
- syndrome:

characteristic facial features, short stature, heart defects,

susceptibility to diseases (respiratory infection,

leukemia,....)

mental retardation short life span



87

- a. trisomy 21 (Down syndrome)
- syndrome:

characteristic facial features, short stature, heart defects, susceptibility to diseases (respiratory infection, leukemia,....), mental retardation, short life span

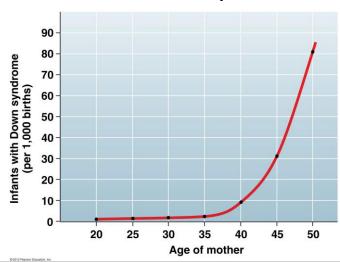


widely separated first and second toes and increased skin creases

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- a. trisomy 21 (Down syndrome)
- older age of the mother, higher the risk of Down syndrome children

under 30: < 0.05% 40 yr.: ~1%



- in female, meiosis begins before birth, but arrested at prophase I; only one egg matures every month
- -> maybe, longer the time lag, greater the chance that eggs to be damaged (meiosis or checkpoint)

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human somatic cell:

46 chromosomes (2n=46)

44 autosomes, 2 sex chromosomes: XX or XY

體染色體 性染色體

23 pairs of homologous chromosomes

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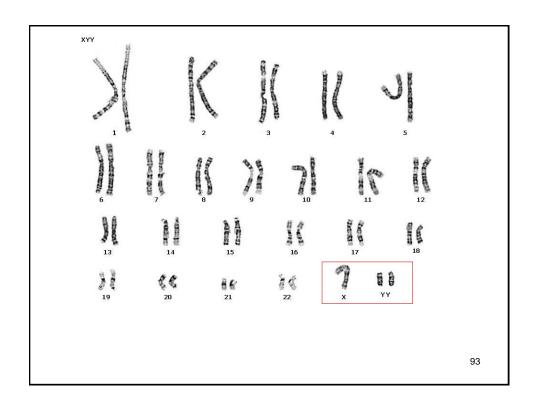
- b. aneuploid of sex chromosome
- less harmful than autosome aneuploid
- relative few genes on Y chromosome, regulation on X -> only one X is active

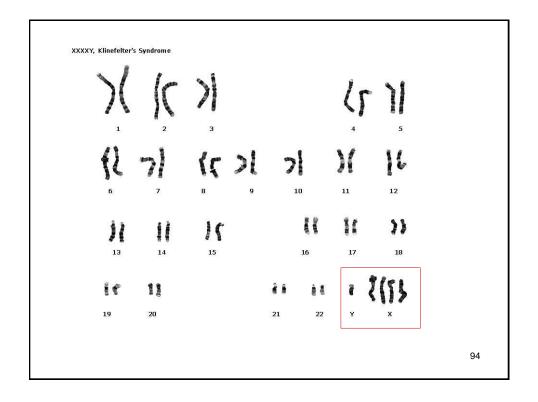
XO Turner syndrome (female) sterile

XXX metafemale syndrome limited fertility

XXY Klinefelter syndrome (male) sterile

XYY normal male normal





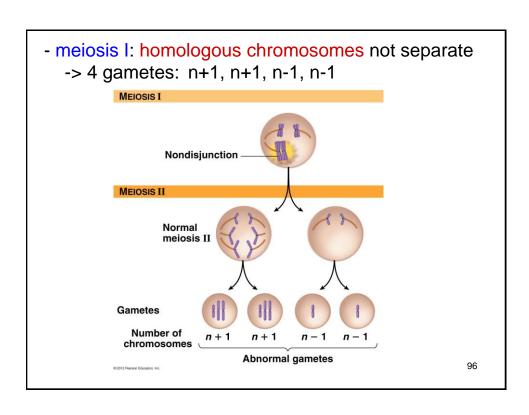
- c. causes of aneuploid:
 - chromosome nondisjunction during meiosis
 - -> members of a chromosome pair fail to separate
 - meiosis I:

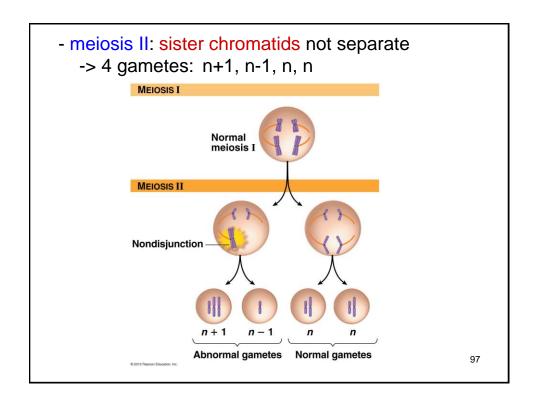
homologous chromosomes not separate

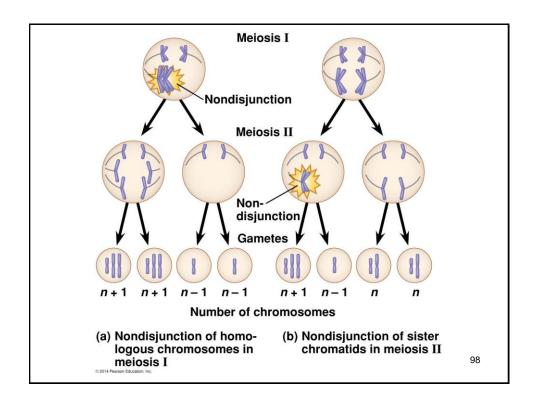
- -> 4 gametes: n+1, n+1, n-1, n-1
- meiosis II:

sister chromatids not separate

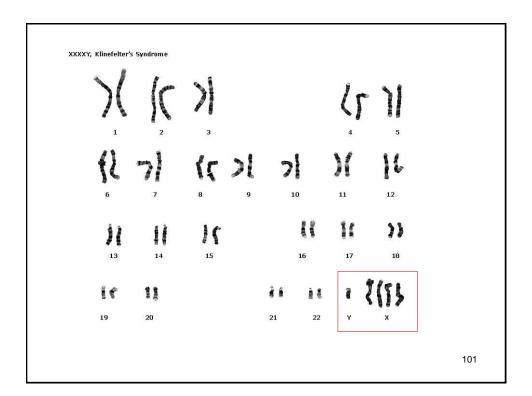
-> 4 gametes: n+1, n-1, n, n







Homework-2 due date 4/12		
type	sperm or egg	nondisjunction at meiosis I, II, or I or II
XXY	sperm egg	l I or II
XYY	sperm	?
XXX	?	?
ХО	?	?
		100



1). abnormal numbers of chromosomes

-> aneuploid (非整倍性)

2n-1 -> monosomy (e.g., XO)

2n+1 -> trisomy, most spontaneous abortions, few live-born (e.g., trisomy 8, 13, 18, 21, Y, or X),

compare to polyploidy (e.g., 3n, 4n)

d. polyploidy:

more than two complete chromosome sets

- triploidy (3n), tetraploidy (4n)
- common in plants, e.g., wheat (6n), strawberry (8n)
- polyploidy fish and amphibian
- gametogenesis by mitosis or zygote failed to divide after DNA replication

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2). Alterations of chromosome structure

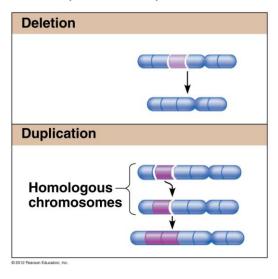
on single or b/w homologous chromosomes

- deletion 缺失
- duplication 重複
- inversion 倒位

between nonhomologous chromosome

- translocation 易位

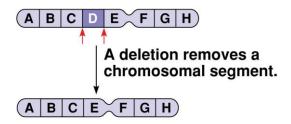
- 2). Alterations of chromosome structure
- deletion: loss of a segment of a chromosome
- duplication: repetition of part of a chromosome



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- 2). Alterations of chromosome structure
- deletion: loss of a segment of a chromosome

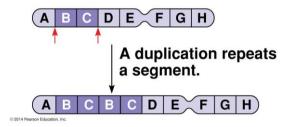
(a) Deletion



2). Alterations of chromosome structure

- duplication: repetition of part of a chromosome

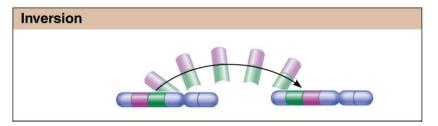
(b) Duplication



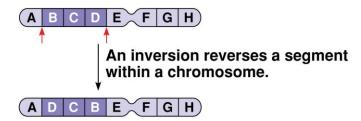
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2). Alterations of chromosome structure

- deletion: loss of a segment of a chromosome
- duplication; repetition of part of a chromosome
- inversion: change in the orientation of a segment of a chromosome, less harmful



- 2). Alterations of chromosome structure
- inversion: change in the orientation of a segment of a chromosome, less harmful
 - (c) Inversion



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- 2). Alterations of chromosome structure
- translocation: one fragment of a chromosome breaks off and becomes attached to a nonhomologous chromosome, reciprocal or nonreciprocal

