

Cells can distribute genetic info to their daughter offsprings.

- Every cell comes from a cell
- Cell division distributes DNA to daughter cells
- Mitosis: division -> identical 2n somatic cells
- Meiosis: germ cell -> 4 gametes half the chrom
- DNA has to be copied before divided
- Chromosome organization of (2m) DNA
- Chromosome chromatin: DNA + proteins only
- Somatic (2x23 humans), gametes (1x23)

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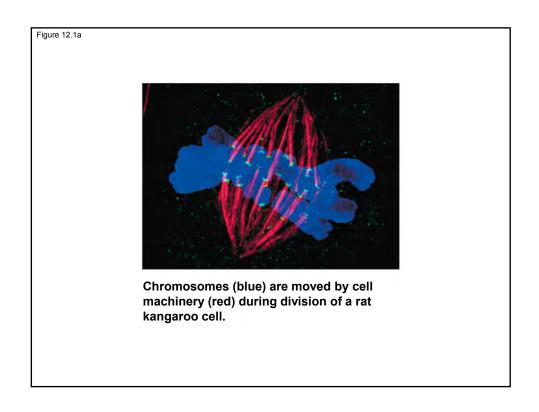
Cells can distribute genetic info to their daughter offsprings.

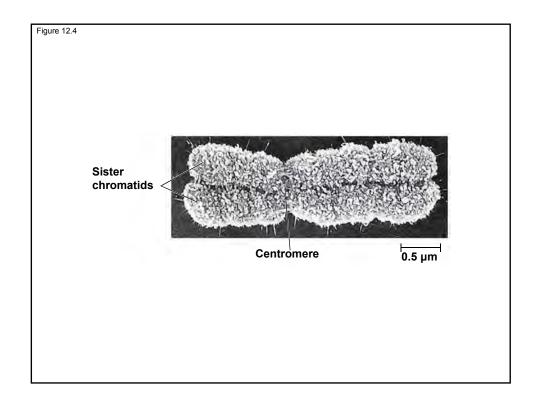
- During division chromosomes condense
- Sister chromatids joined by cohesins
- Centromere: region of DNA where chromatids are attached to each other
- Cytokinesis: division of cytoplasm of cell
- Mitosis: daughter cells get full 2x 23 set
- Meiosis: daughter cells get one set of 23
- One centromere per duplicated chromosome

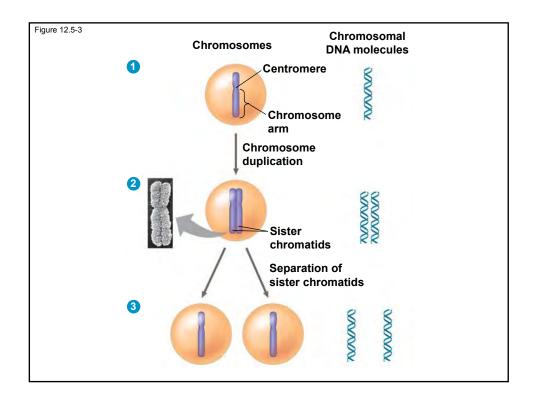
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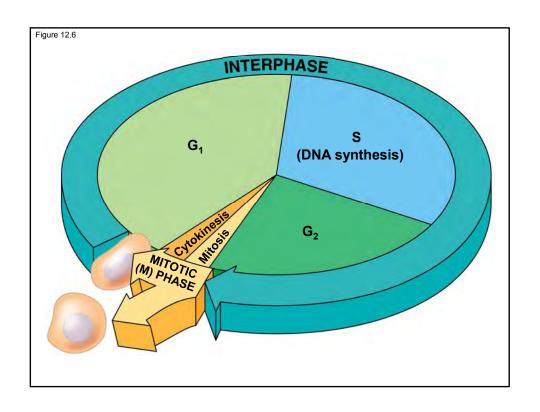
Mitosis alternates with longer interphase.

- Interphase G1, S, G2
- S (synthesis) duplication of chromosomes
- Mitosis + cytokinesis M 1 hr, S 10 to 12 hrs
- G1 variable length (doing job), G2 5 hr prep
- G2 nuceolus envelope, uncondensed chrom
- Normal function and growth is in G1 (what we know cells to be at is G1)

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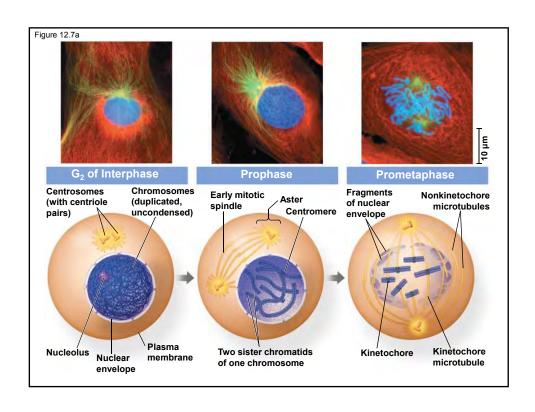
Substages of Mitosis.

- Prophase: condensed chromatin, nucleolus disappear, chromatids joined with spindles radiating out from centrosome (asters)
- Prometaphase: nuclear envelope disappears, kinetochore protein at centromere, opposite spindle microtubules attach to kinetochore
- Metaphase: centrosomes at opposite poles, chromosomes on metaphase plate, alignment of chromosomes on the equator

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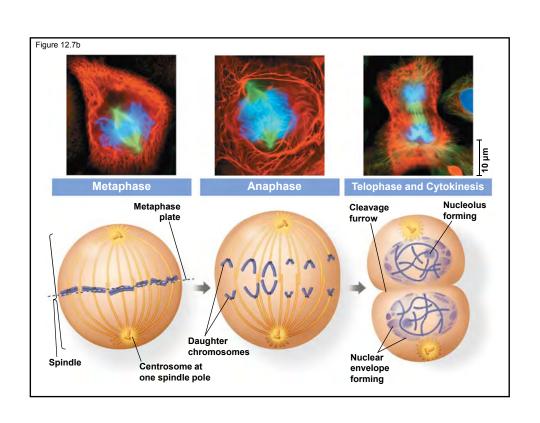


Substages of Mitosis.

- Anaphase: short, cohesins cleaved allowing sister chromatids of each pair to part, kinetochore microtubules attached shorten
- Telophase: 2 nuclei form from ER and old envelope, chromosomes decondense
- Cytokinesis: cleavage furrow pinches cell into two, two daughter cells

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A closer look at centrosomes.

- Microtubules consists of tubulin units
- Centrosome organizes microtubules
- Centriole pair inside centrosome nonessential
- Centrosome divides and move to opposite sides of cell by prometaphase
- Aster microtubule array extends out
- Opposite facing kinetochores attach to each sister chromatid, tugged into metaphase plate

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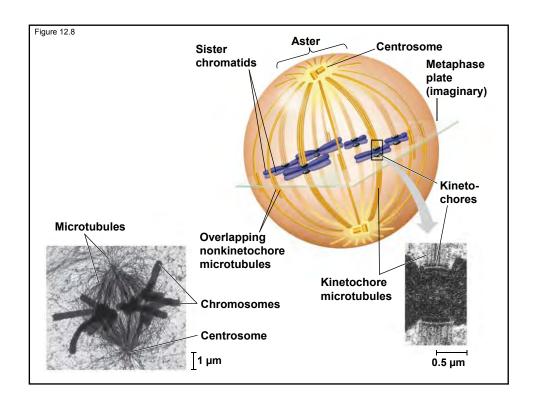
A closer look at centrosomes.

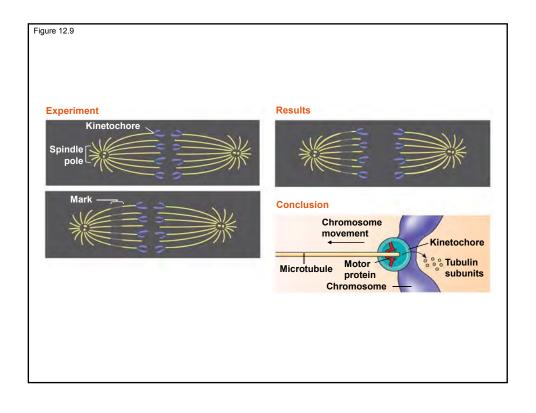
- Separase enzyme cleaves cohesins allowing chromatids to move to opposite sides
- Motor proteins pull at spindle pole? or Pacman: kinetochore push walk proteins along microtubules which shorten at chromatid end
- Nonkinetochore microtubules from opposite poles join and lengthen, elongating cell

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A closer look at cytokinesis.

- Animals: cleavage furrow at site of metaphase place formed by actin-myosin contractile ring pinching cell into two.
- Plants: vesicles from Golgi move along microtubules to center of cell to form cell plate, membranes fuse new cell wall.

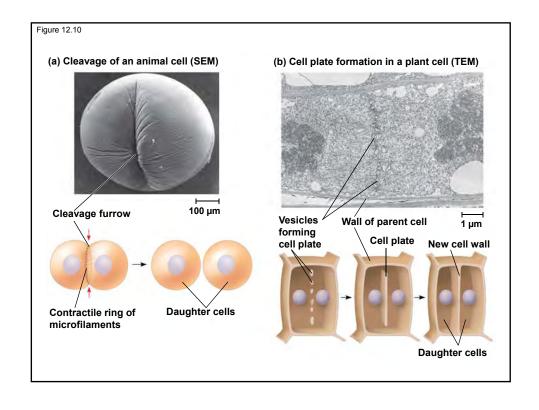
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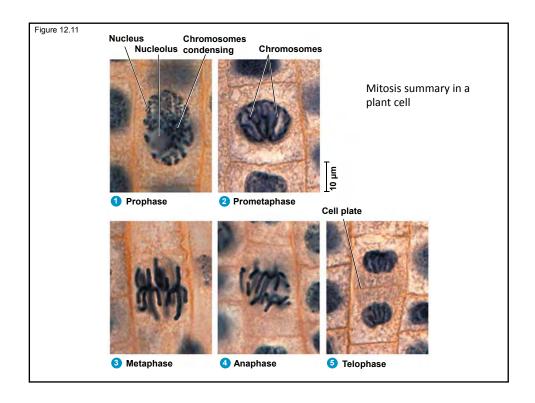
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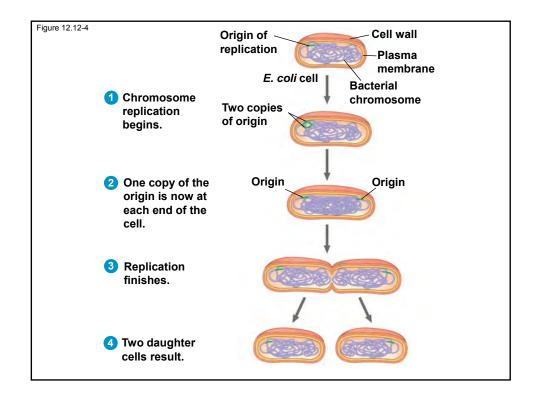
Prokaryotic cell division.

- Bacteria binary fission (no mitosis)
- Single circular DNA: replication start at origin to make two forks to make DNA copies
- No spindles, actin moves, tubulin pinches membrane inward to get two cells
- Evolution: Prokaryotes -> intact nuclei division in dinoflagellates and diatoms -> mitosis

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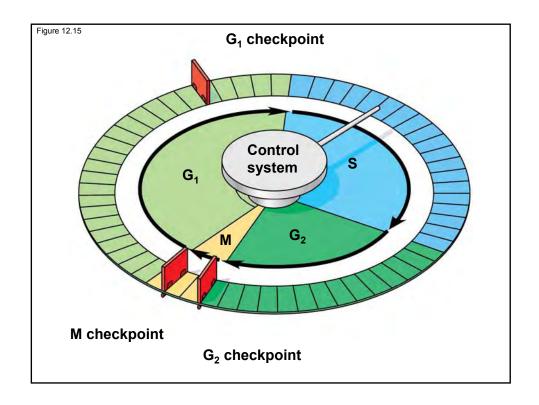
Cell cycle control by cylin dependent kinase proteins.

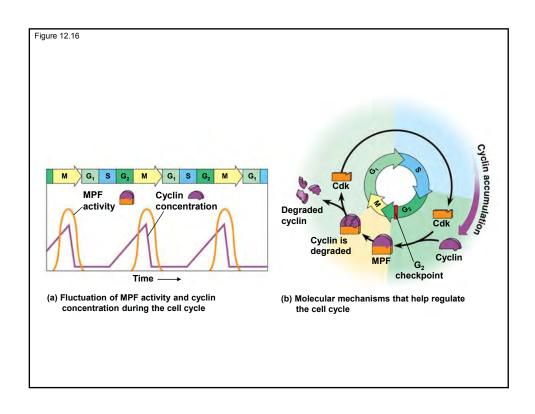
- Fused one cell with two nuclei: if any was in M or S, the other nuclei goes to M or S.
- Cell cycle control as washing machine: built-in clock plus external control, G1 G2 M checkpts.
- Protein kinases phosphorylate proteins, activating or inactivating them.
- Cyclin-dep kinases CDKs activity fluctuates with cyclin concentration (rise in S G2, fall M).
- Ex MPF triggers mitosis, phosphorylate nuclear lamin -> fragment, anaphase destroys own cyclin.

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Cell Cycle control system.

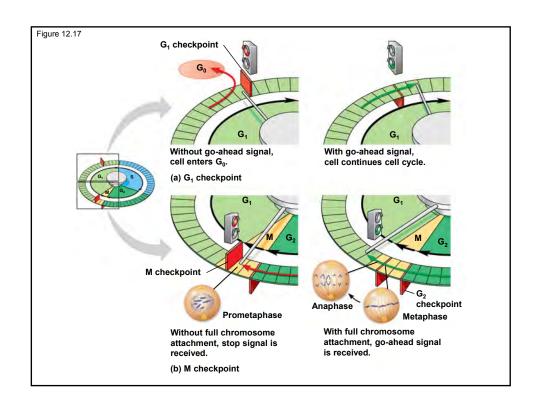
- G1 checkpoint restriction point: go -> S G2 M, if no signal -> nondividing G0 (neurons, liver)
- Ex. anaphase only when ALL kinetochores are attached will cohesins be cleaved chromatids
- Growth factor + essential nutrients -> divide,
 ex. Injuries -> release PDGF -> platelets up
- Cell cycle control of excessive cells: density dep inhibition, anchorage dependence

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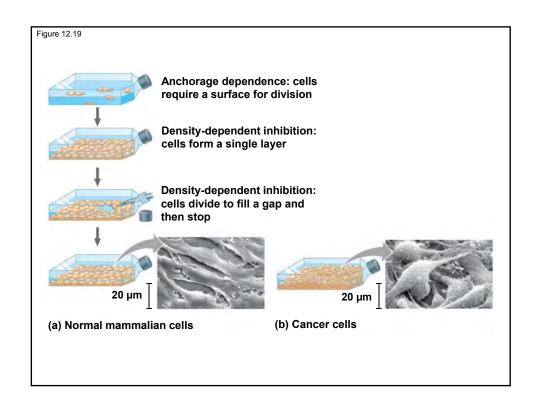


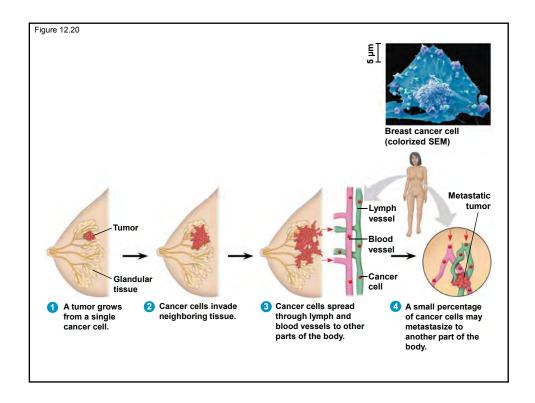
Cancer cells result from failure in cell cycle control.

- Cancerous cells divide without growth factors, stop at random points in cycle, HeLa cell line.
- Usually 20-50 cycles; cancer evades apoptosis
- Benign tumor: can't survive in diff site
- Malignant tumor: spread to new tissue organs
- Metastasis: spread of cancer e.g. via blood ves
- Radiation of cancers (can't repair), side effect on cells that divide (intestine, hair, immune).

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Team activity: answer these questions in teams of 3 or 4 (and justify): 3. How is plant cell cytokinesis different from animal cell cytokinesis?

1. If there are 20 duplicated chromosomes in a cell, how many centromeres are there?

- A) 10
- B) 20
- C) 30

2. Some cells have several nuclei per cell. How could such multinucleated cells be explained?

- A) The cell underwent repeated mitosis, but cytokinesis did not occur.
- B) The cell had multiple S phases before it entered mitosis.
- C) The cell underwent repeated cytokinesis but
- D) The cell underwent repeated mitosis with simultaneous cytokinesis.

- A) Plant cells divide after metaphase but before anaphase; animal cells divide after anaphase.
- B) The structural proteins of plant cells separate the two cells; in animal cells, a cell membrane separates the two daughter cells.
- C) Plant cells deposit vesicles containing cell-wall building blocks on the metaphase plate; animal cells form a cleavage furrow.
- D) The contractile filaments found in plant cells are structures composed of carbohydrates; the cleavage furrow in animal cells is composed of contractile phospholipids.
- 4. Besides the ability of some cancer cells to overproliferate, what else could logically result in a
- A) changes in the order of cell cycle stages
- B) inability to form spindles
- C) inability of chromosomes to meet at the metaphase plate
- D) lack of appropriate cell death

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