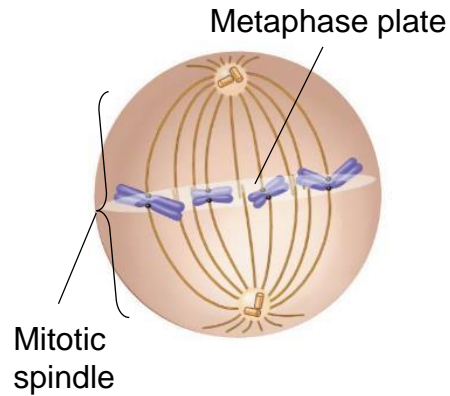
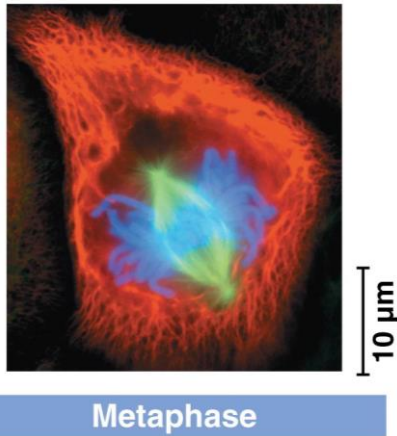


- 4). **metaphase**: 中期
 chromosomes line up on **metaphase plate**
 (**equatorial plane**) due to the pulling force from
 the opposite poles and cohesion



1

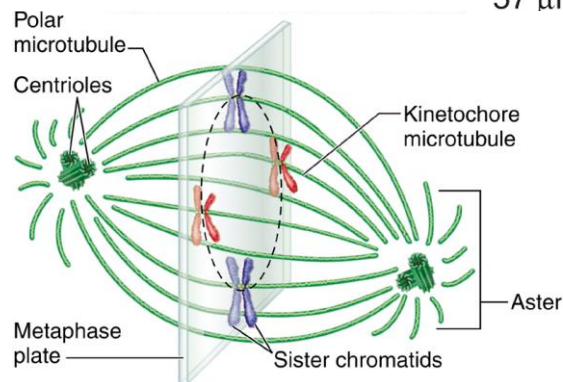
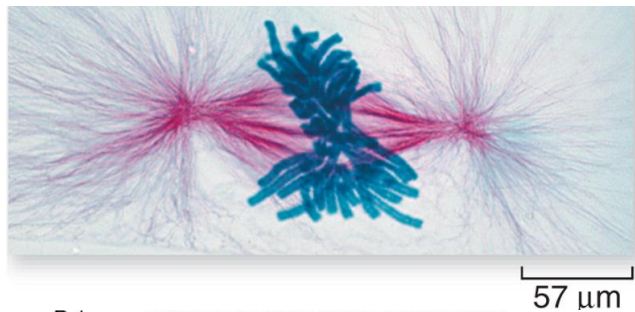
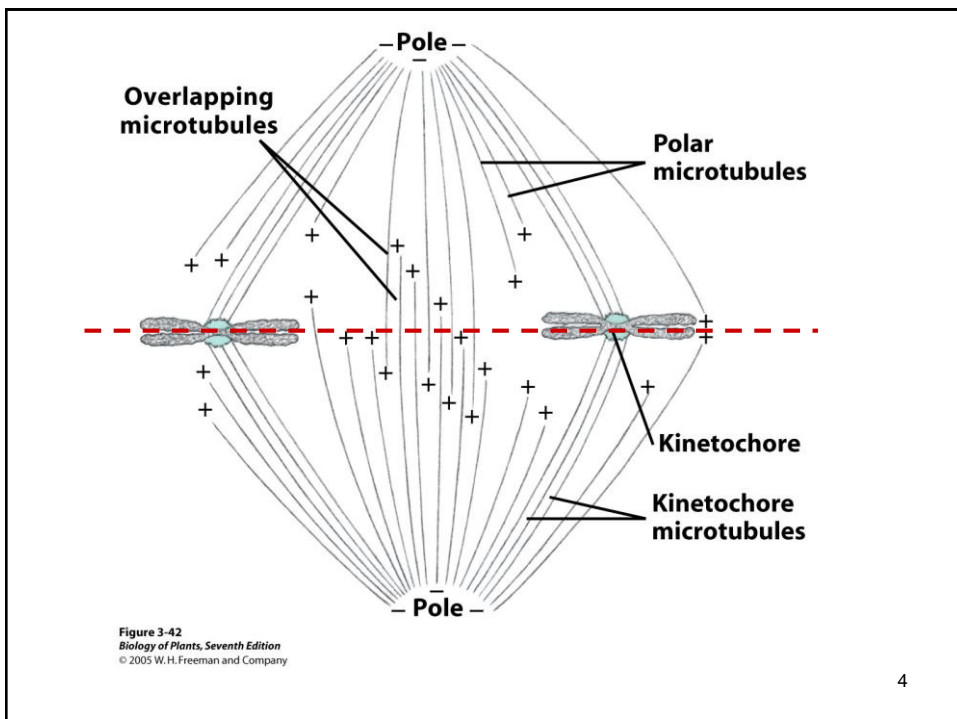
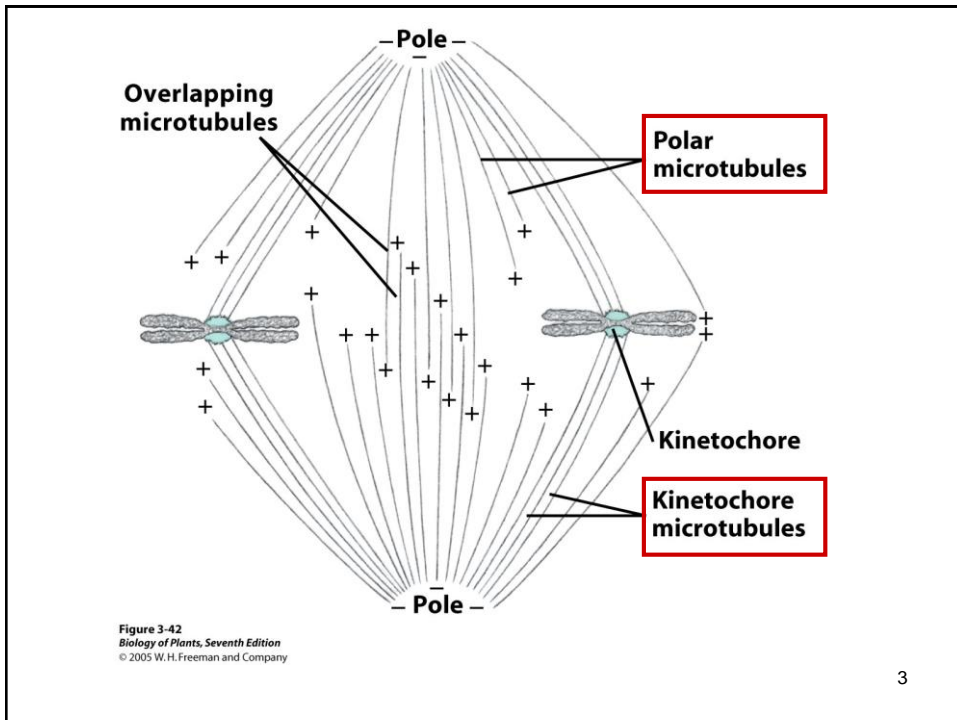


Fig.10.12

2



§ in mitosis and meiosis

chromosomes line up on equatorial plane
(metaphase plate), due to **bipolar attachments**

-> **bilateral symmetry (to each separation unit)**

-> to ensure proper separation of sister chromatids

do not forget cohesion

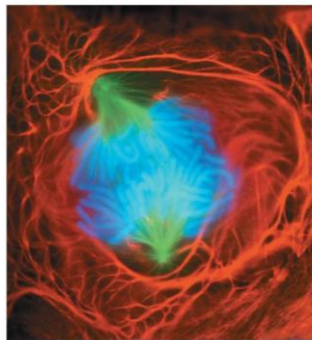
5

5). **anaphase**:後期

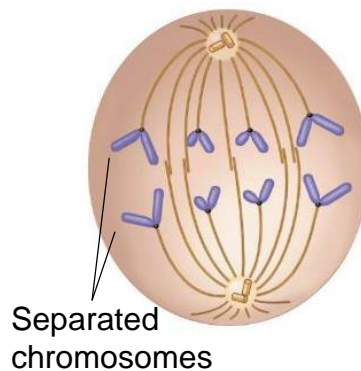
- sister chromatids separate

-> daughter chromosomes

-> move toward opposite poles



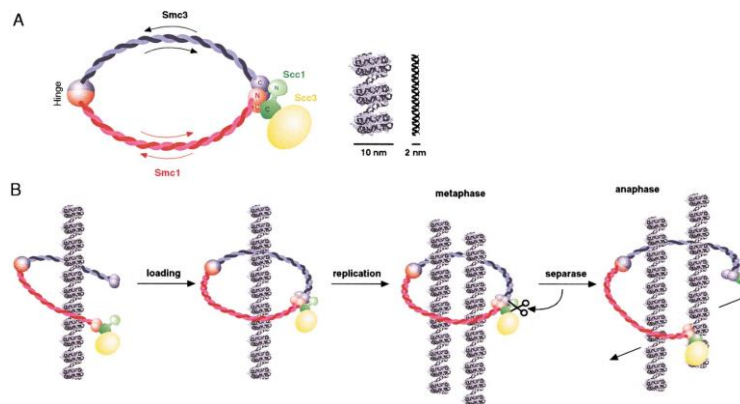
Anaphase



6

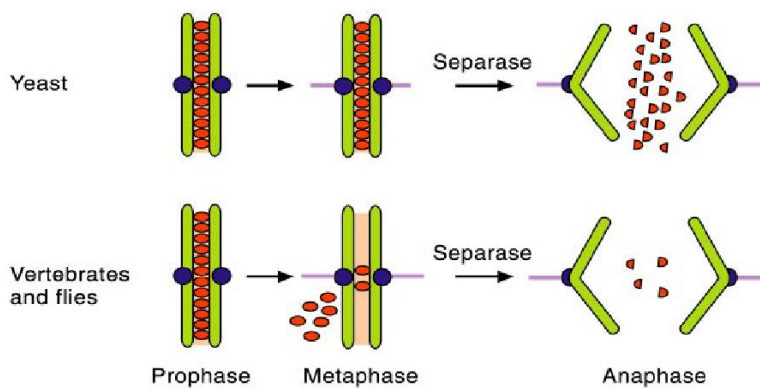
5). anaphase:

- sister chromatids separate
(proteolysis of cohesin complex)
- > daughter chromosomes
- > move toward opposite poles



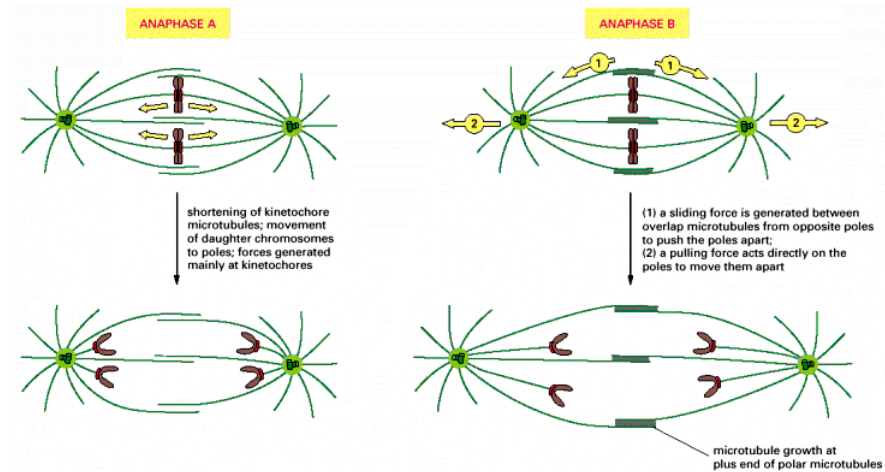
7

A. Mitosis



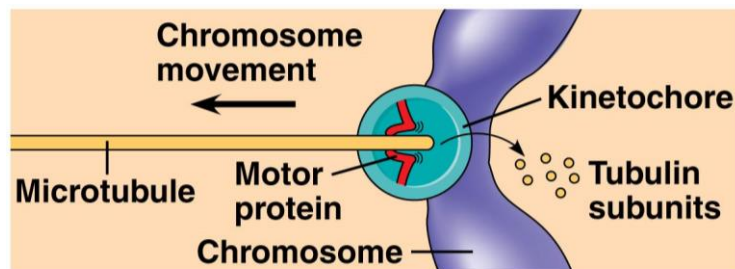
8

anaphase A: poleward movement of chromosomes
anaphase B: poles move farther apart



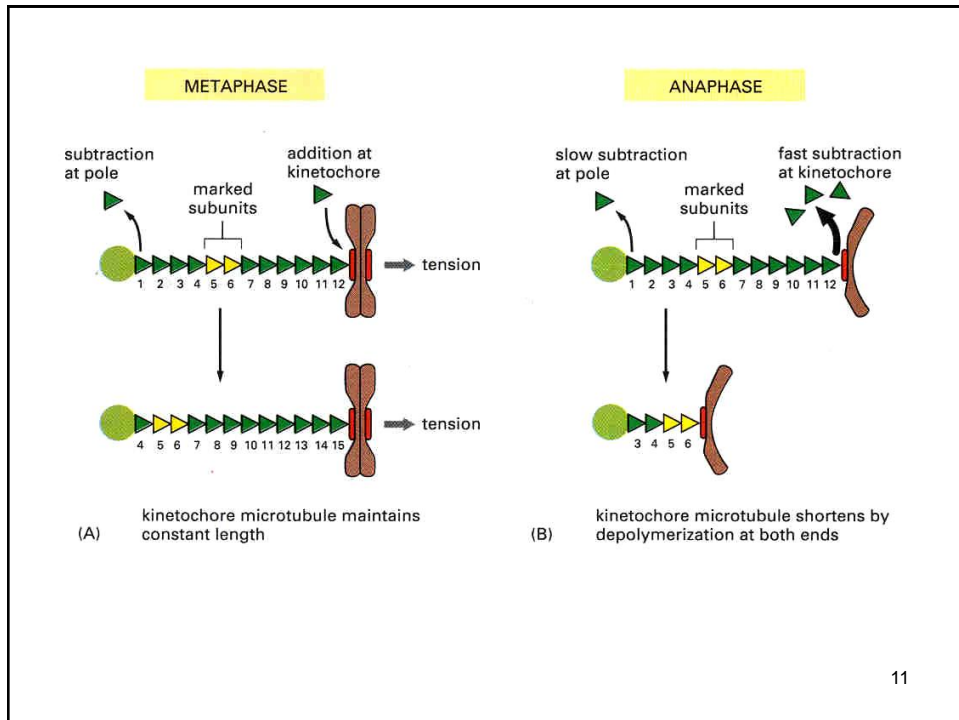
9

anaphase A: poleward movement of chromosomes
 two mechanisms:
 shortening of kinetochore microtubules
 kinetochore "walking" on microtubules

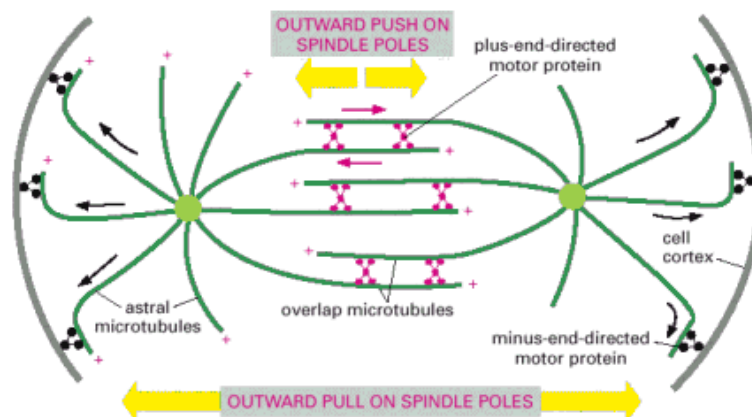


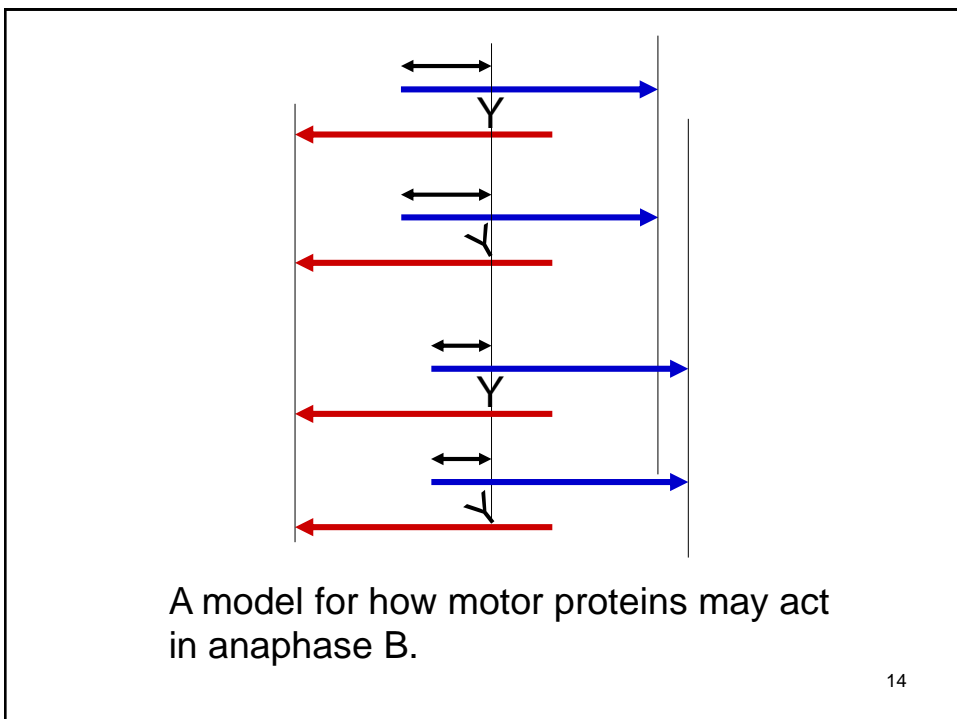
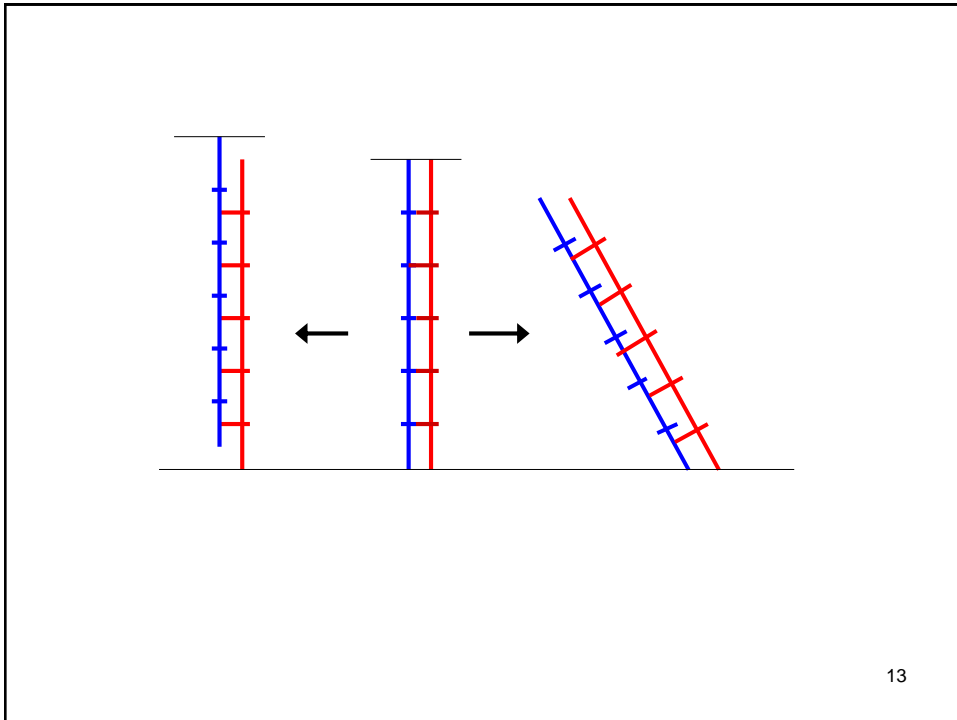
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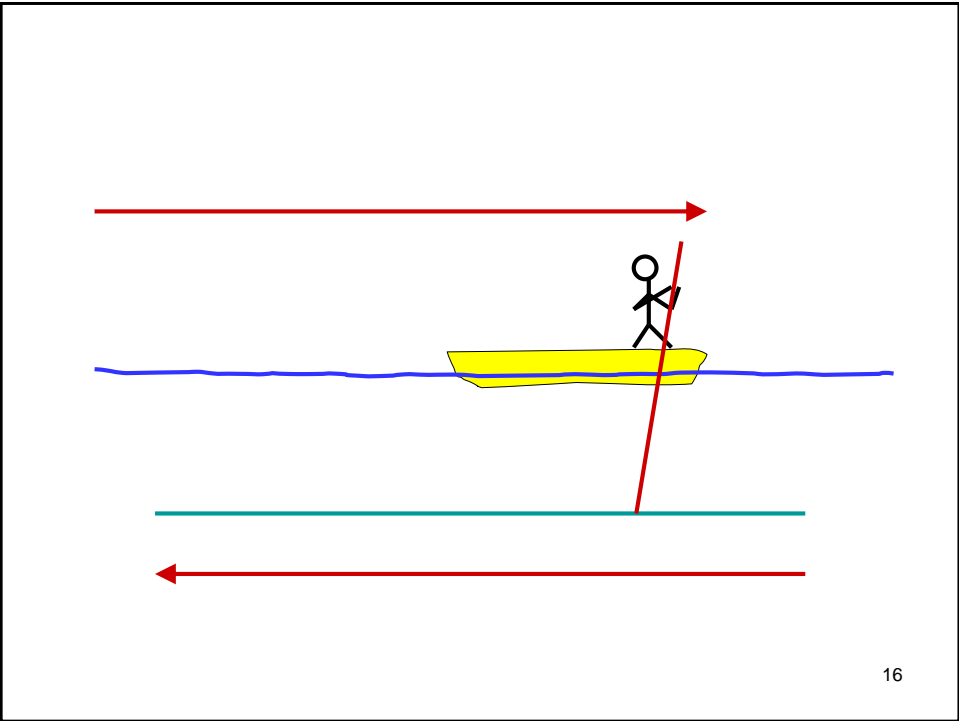
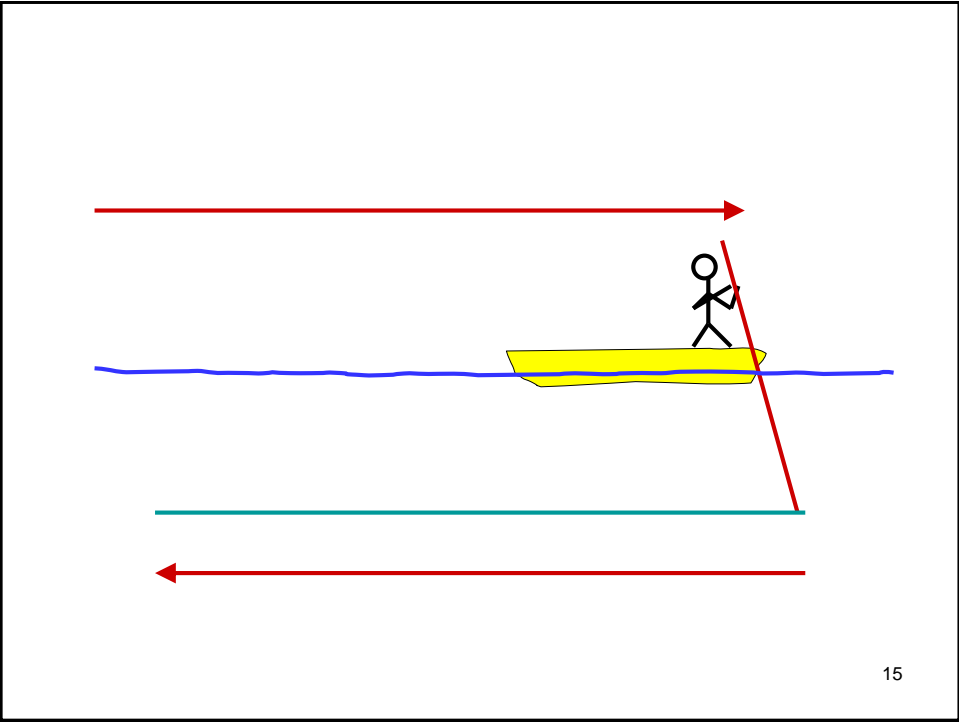
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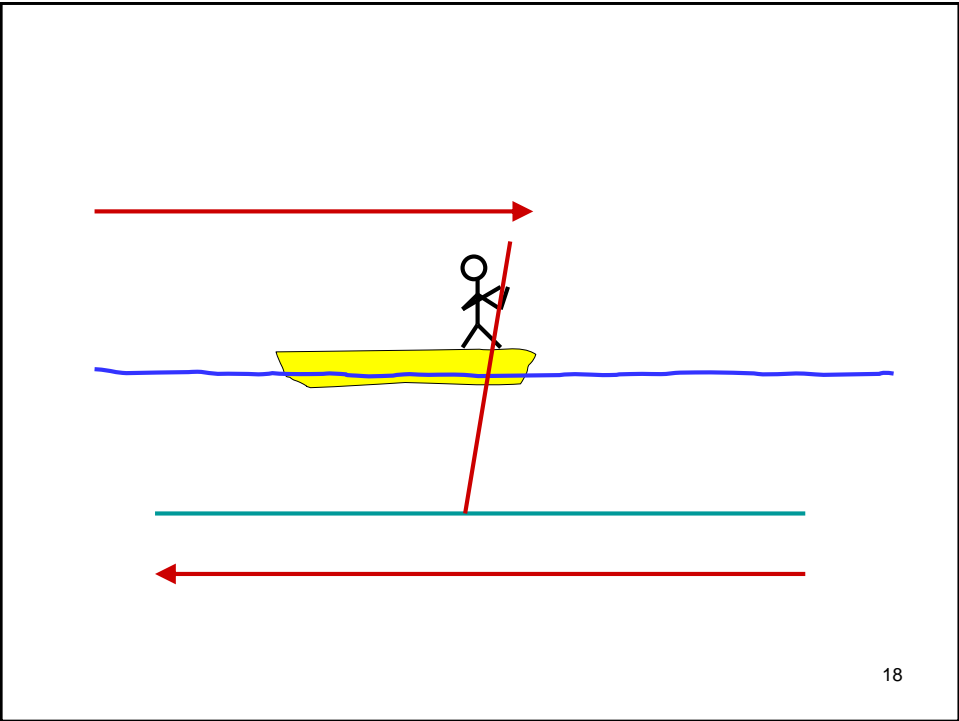
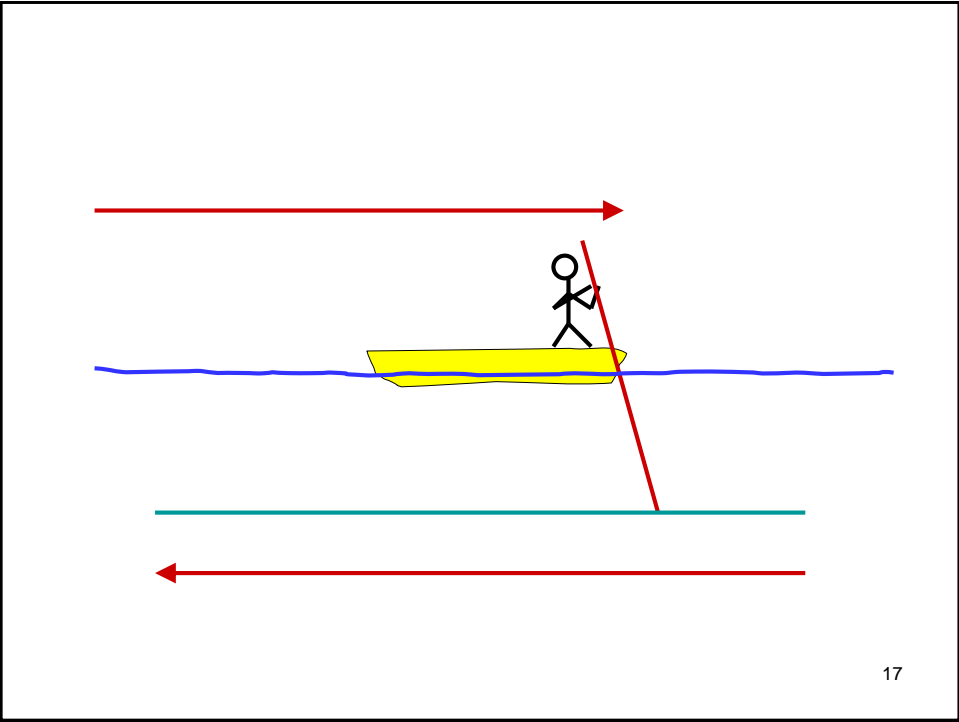


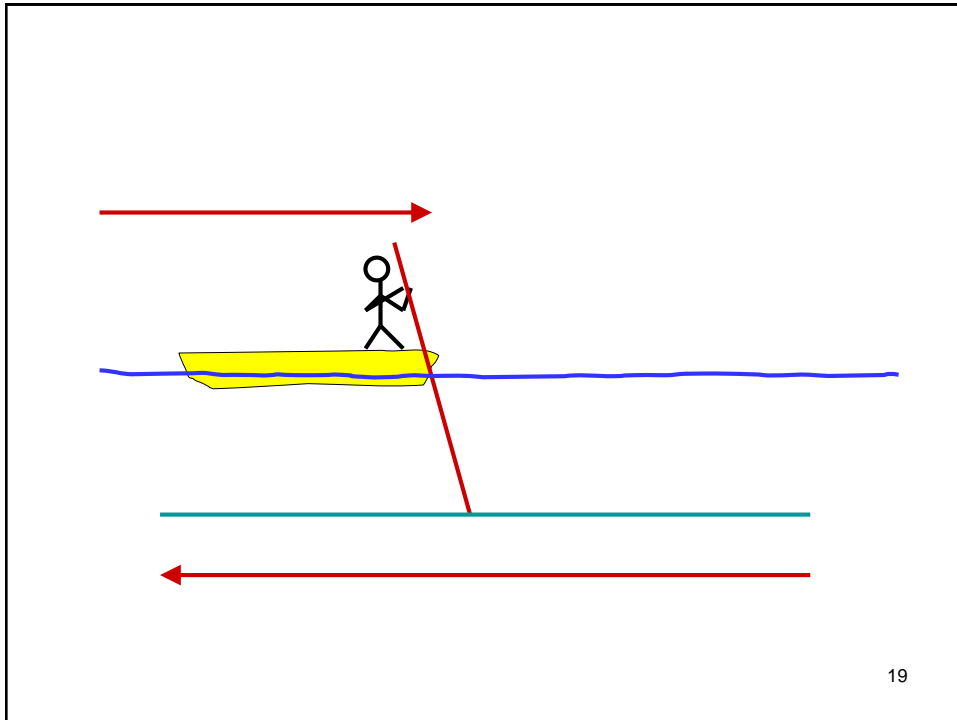
anaphase B: poles move farther apart
 by a **pushing force** between the polar microtubules
 from the opposite poles and **pulling force** to the
 poles by astral microtubules
 -> daughter chromosomes move further apart







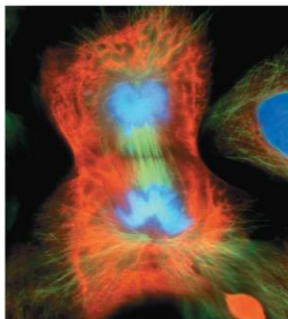




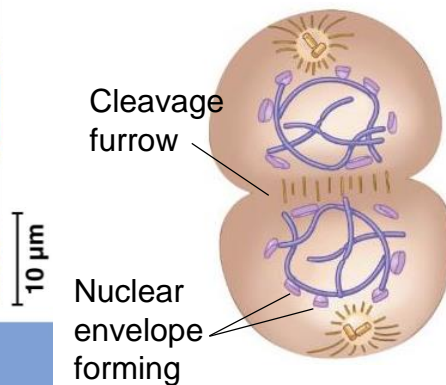
19

6). **telophase** 末期

- spindle disassembles
- nuclear envelopes form -> daughter nuclei appear
- nucleolus appears
- chromosomes decondense



Telophase and
Cytokinesis



20

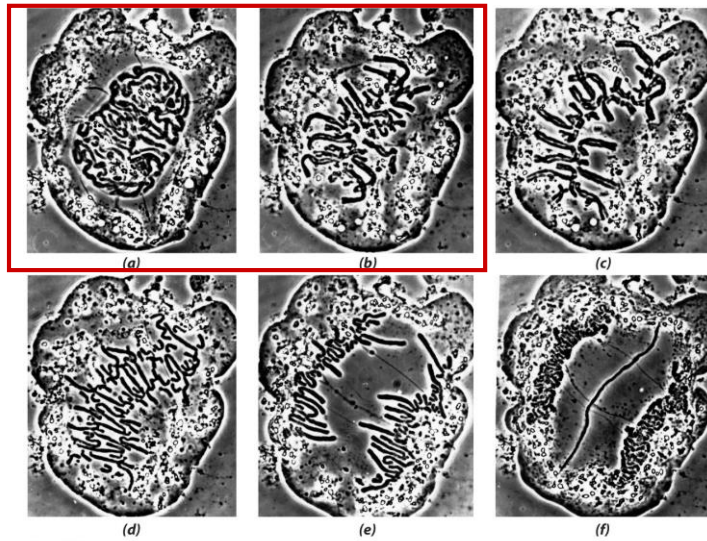


Figure 3-39
Biology of Plants, Seventh Edition
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a: prophase; b: prometaphase

21

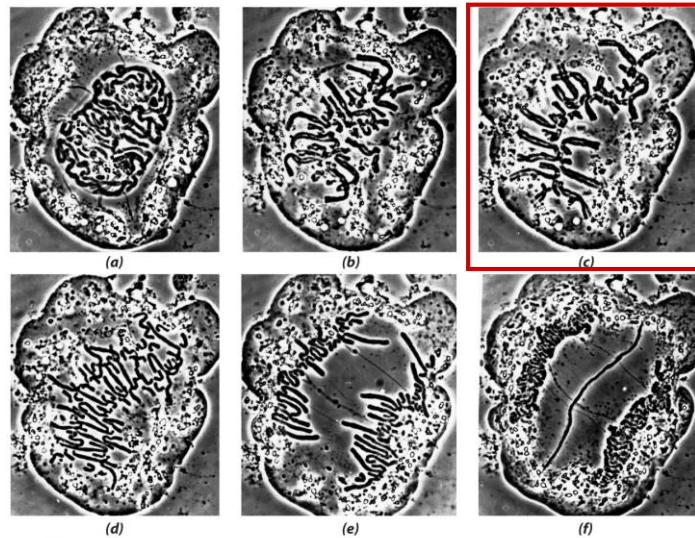


Figure 3-39
Biology of Plants, Seventh Edition
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c: metaphase

22

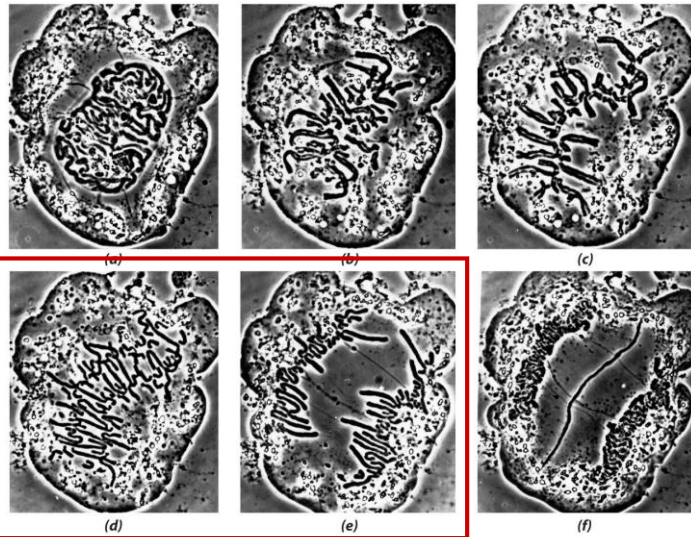


Figure 3-39
Biology of Plants, Seventh Edition
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d,e: anaphase

23

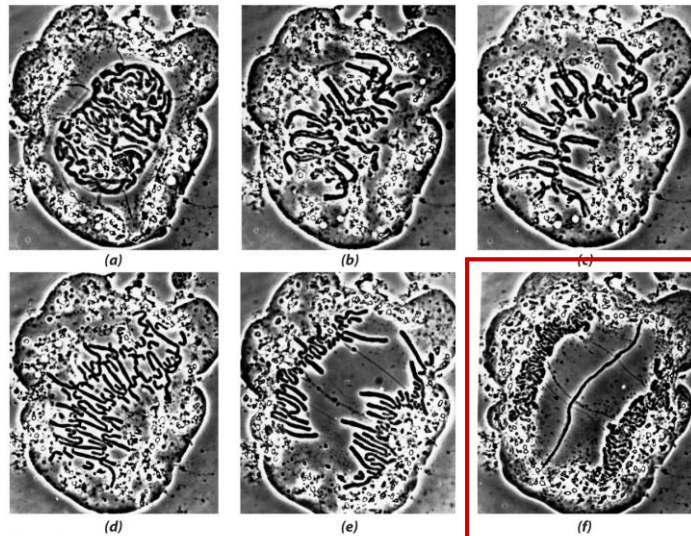
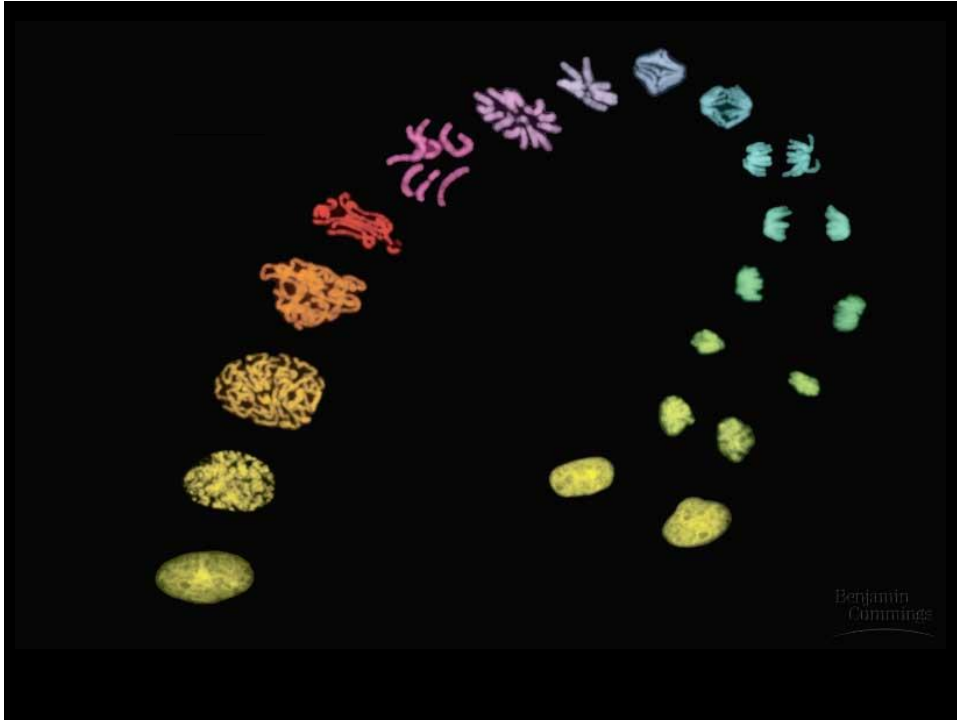


Figure 3-39
Biology of Plants, Seventh Edition
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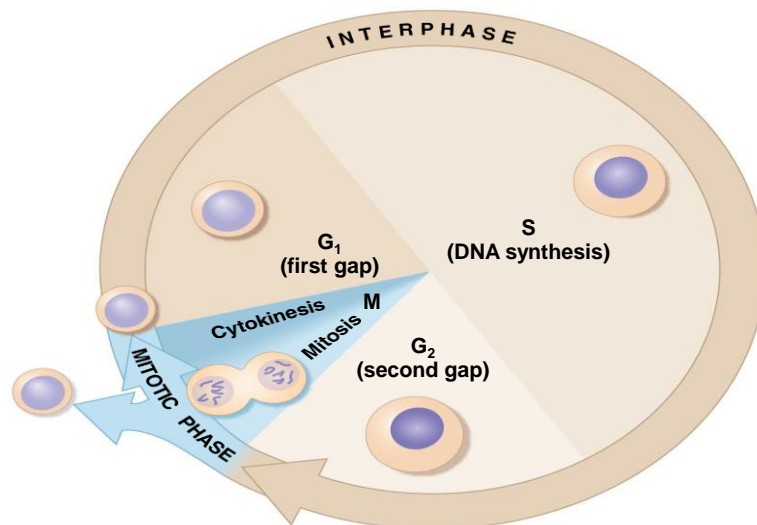
f: telophase

24



2). M (mitotic) phase

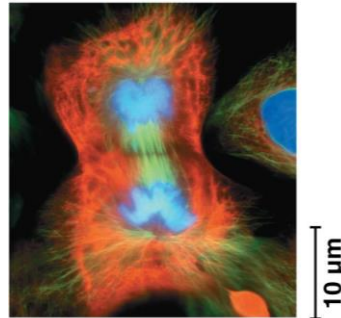
- a. mitosis - chromosome separation
- b. **cytokinesis** - division of cytoplasm



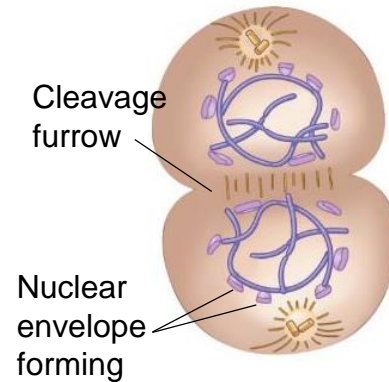
26

7). cytokinesis – divide the cell into two

- replication of organelles in S or G2
- > partition into two daughter cells
- usually occurs along with telophase



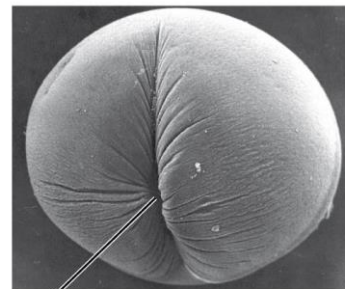
Telophase and Cytokinesis



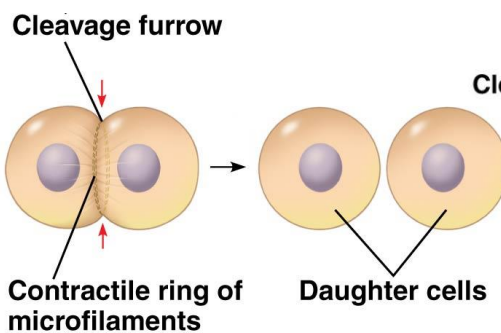
27

- animal cells:
- ring of microfilament
- > contracts
- > cleavage furrow 分裂溝
- > two cells

(a) Cleavage of an animal cell (SEM)



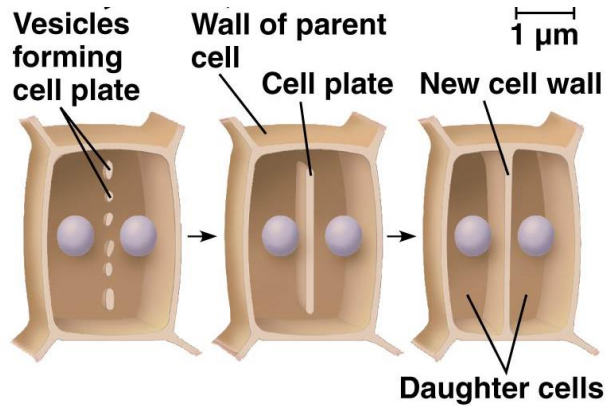
100 μm



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28

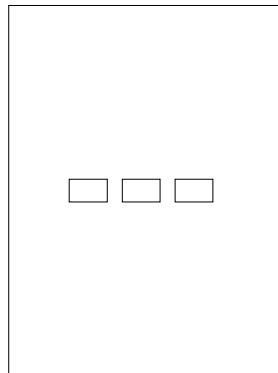
- plant cells:
membrane-enclosed vesicles (containing cell wall materials) from the Golgi apparatus move to the middle of the parent cell
- > fuse to form **cell plate** 細胞板



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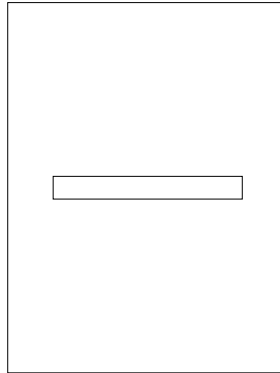
29

- > growing **cell plate** reaches and fuses with plasma membrane



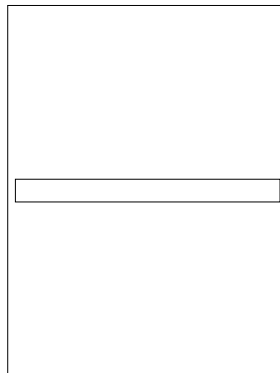
30

-> growing **cell**
plate reaches
and fuses
with plasma
membrane



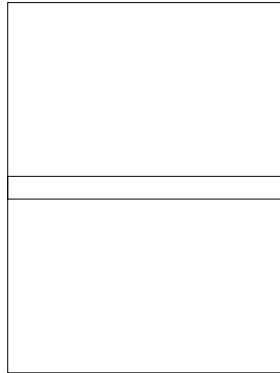
31

-> growing **cell**
plate reaches
and fuses
with plasma
membrane



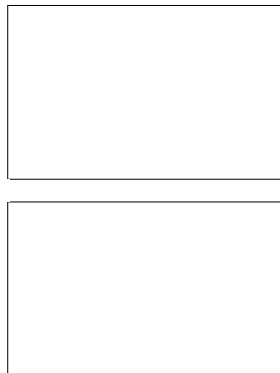
32

-> growing **cell**
plate reaches
and fuses
with plasma
membrane



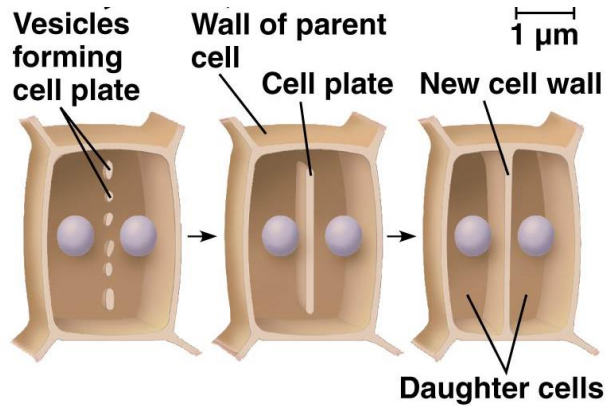
33

-> growing **cell**
plate reaches
and fuses
with plasma
membrane



34

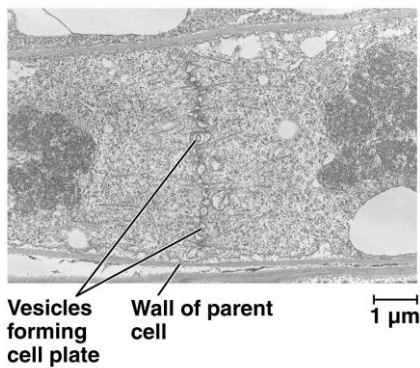
- > cell plate grows and fuses with plasma membrane
- > cell wall formation
- > two cells



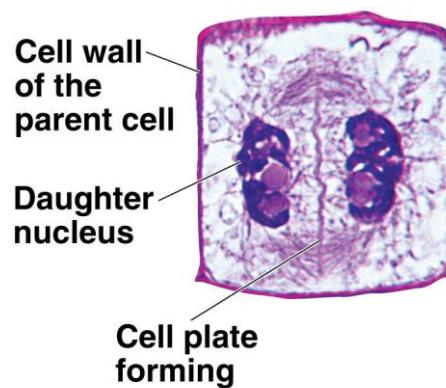
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35

- plant cells:
membrane-enclosed vesicles (containing cell wall materials) from the Golgi apparatus move to the middle of the parent cell



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36

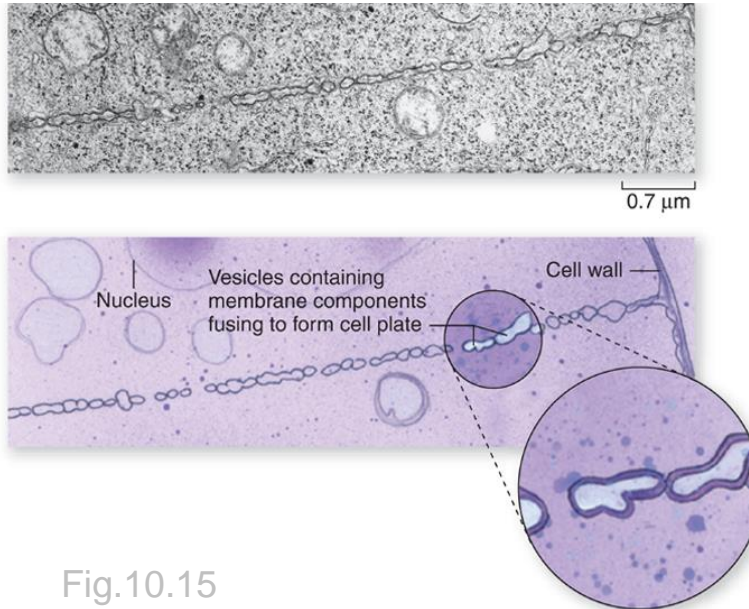
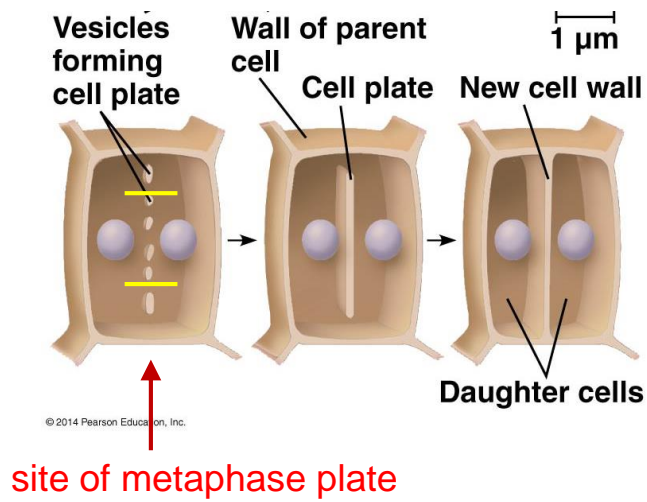
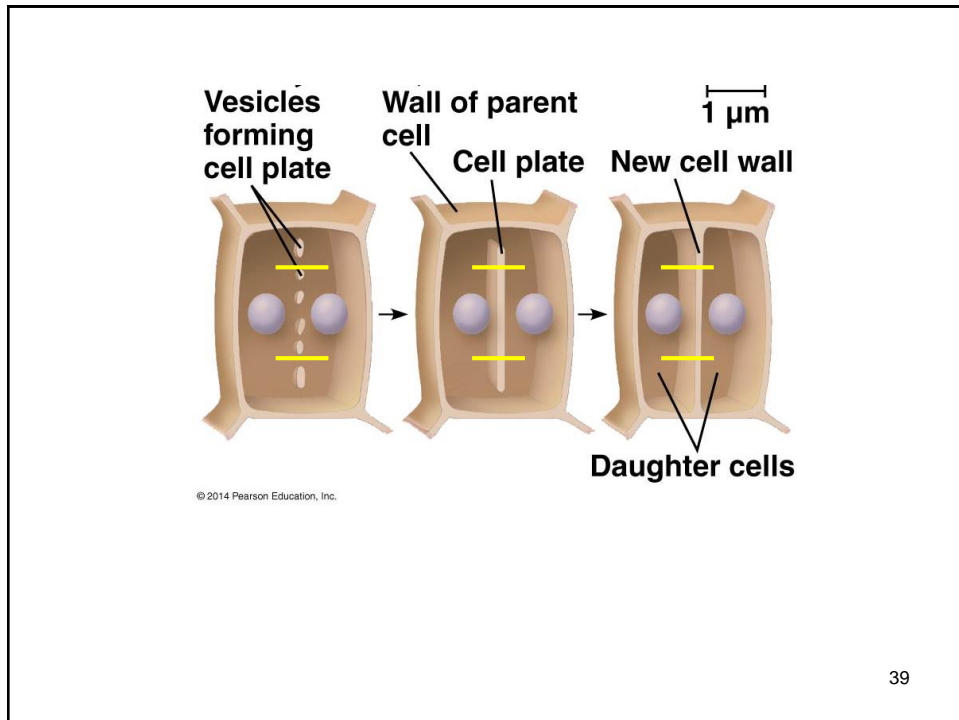


Fig.10.15

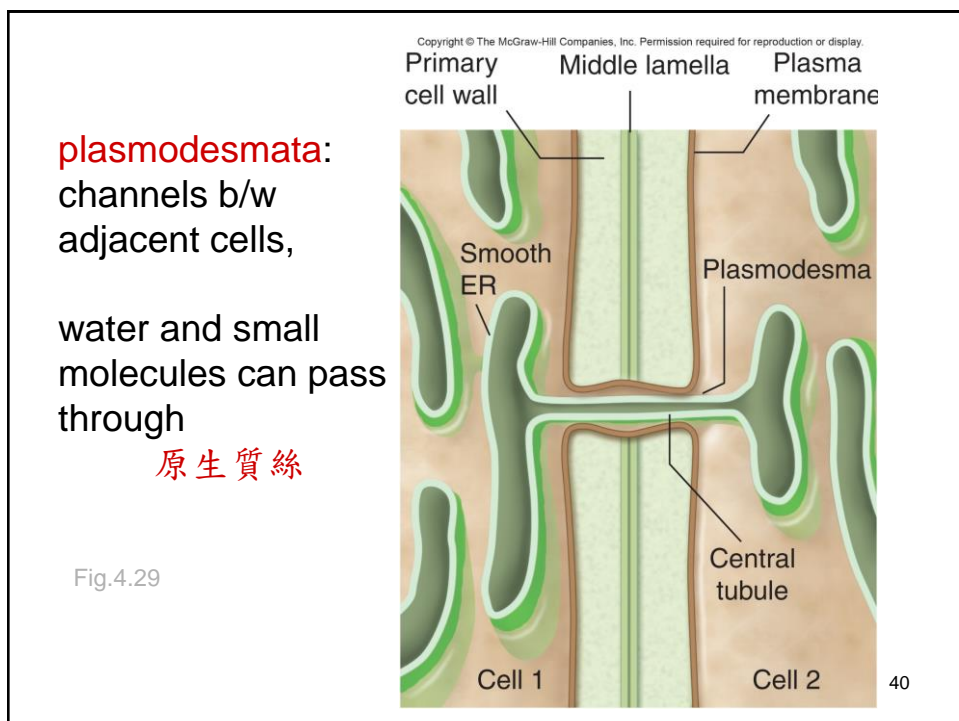
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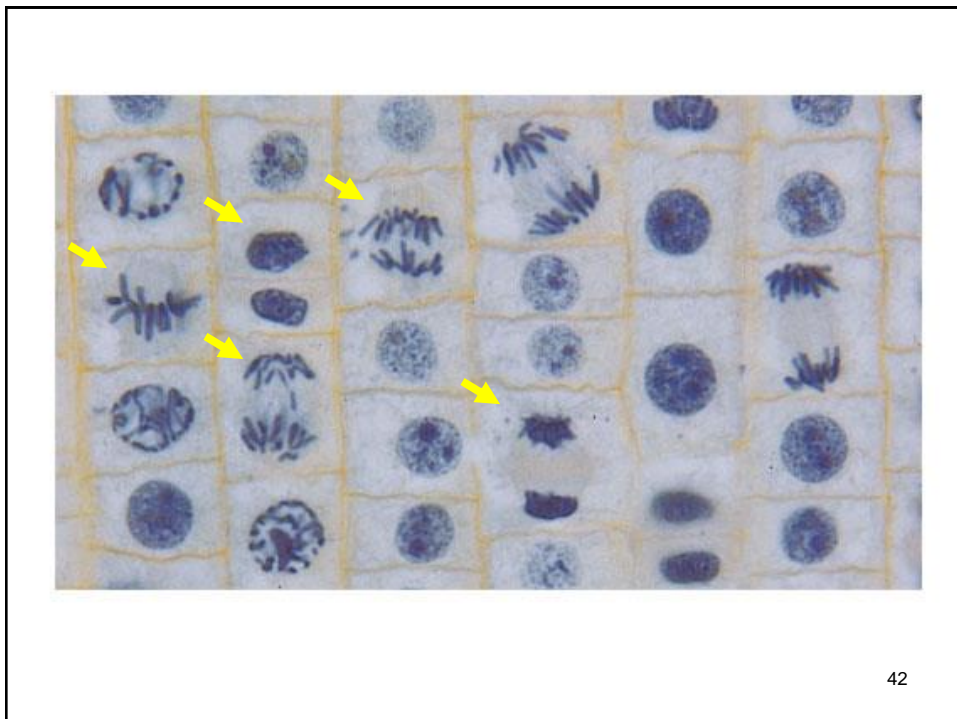
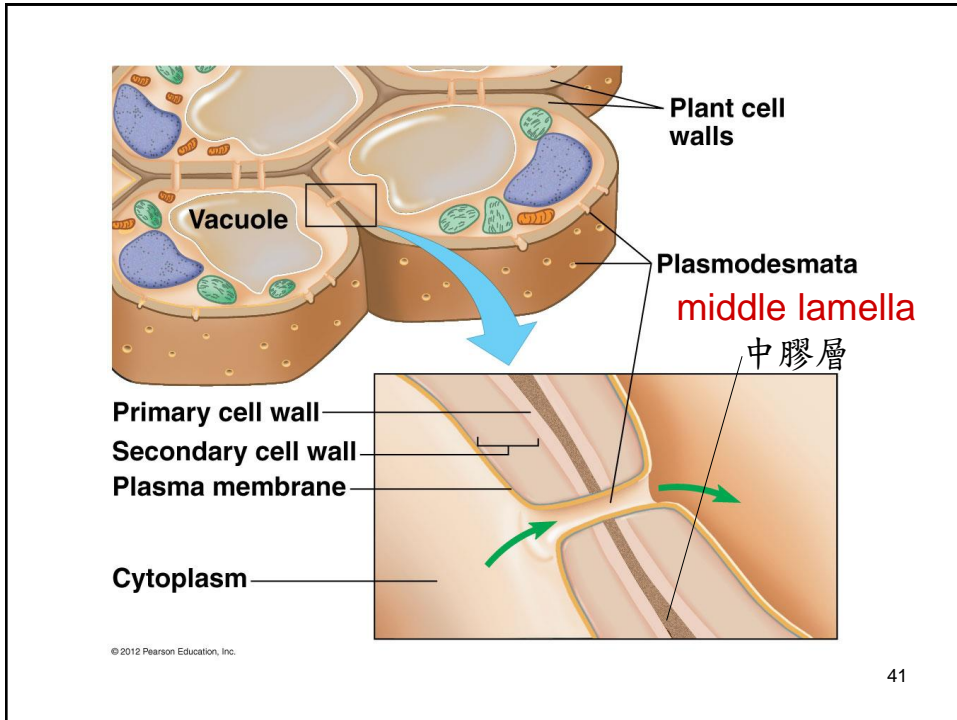


38



39





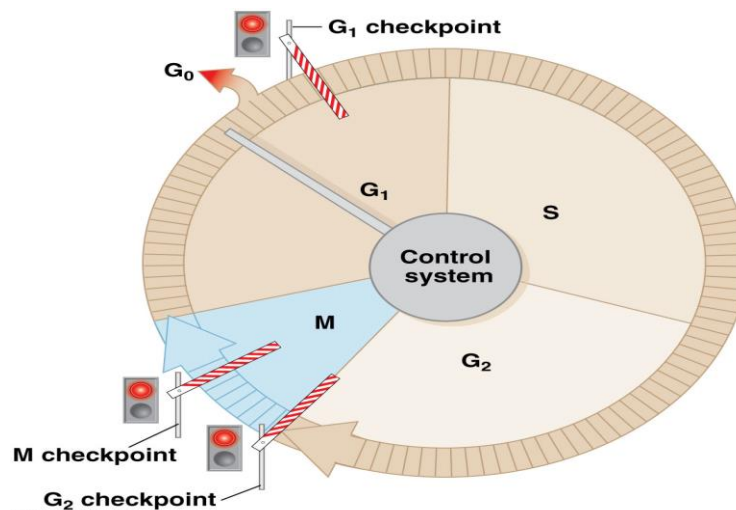
VI. Cell cycle control

- 1) Cell division is a complicated process and needs a good control
 - **cell-cycle control system** triggers and coordinates cell cycle events
- a. irreversible points:
- DNA replication, separation of sister chromatids
- b. can be arrest (put on hold) at specific points
 - > **checkpoints**

43

2) **checkpoints**

cell-cycle control works through checkpoints to determine the initiation or continuation of cell cycle:



44

2) checkpoints

- **sense and execute**

A. cell-cycle control system responds to signals

- outside:

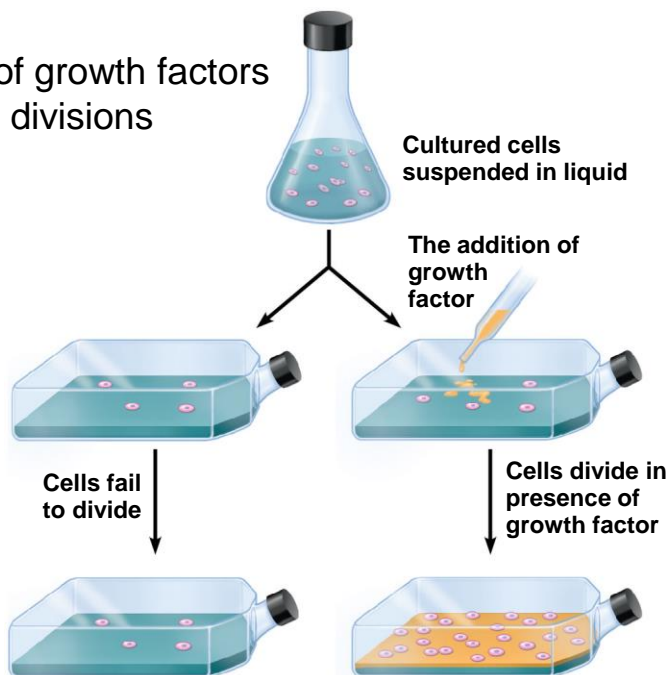
- nutrients,
 - growth factors,
 - anchorage dependence,
 - density-dependent inhibition

- inside:

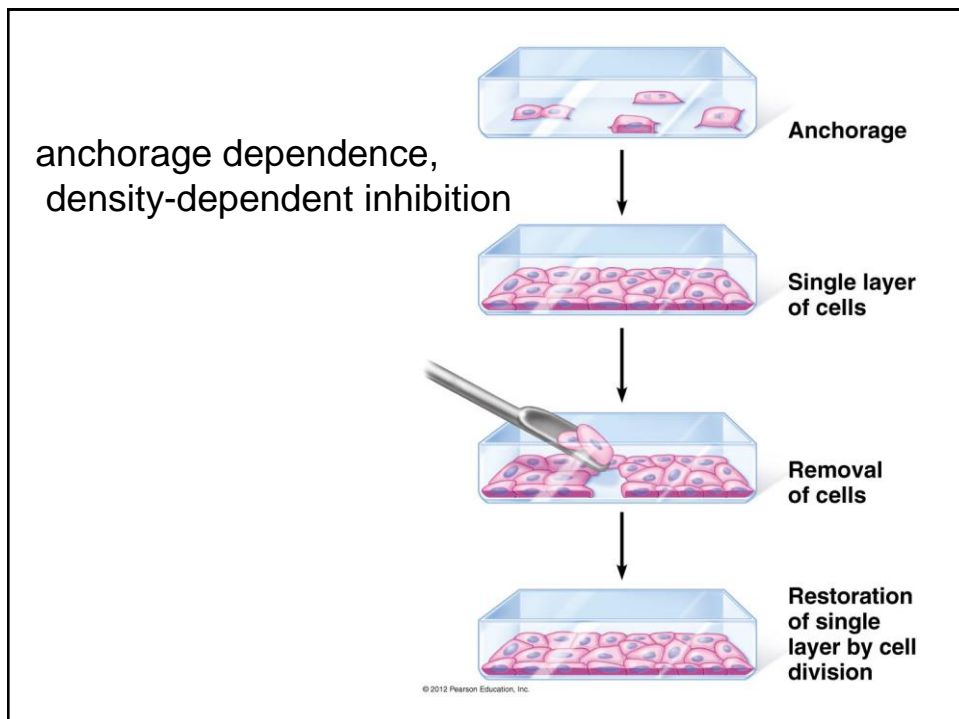
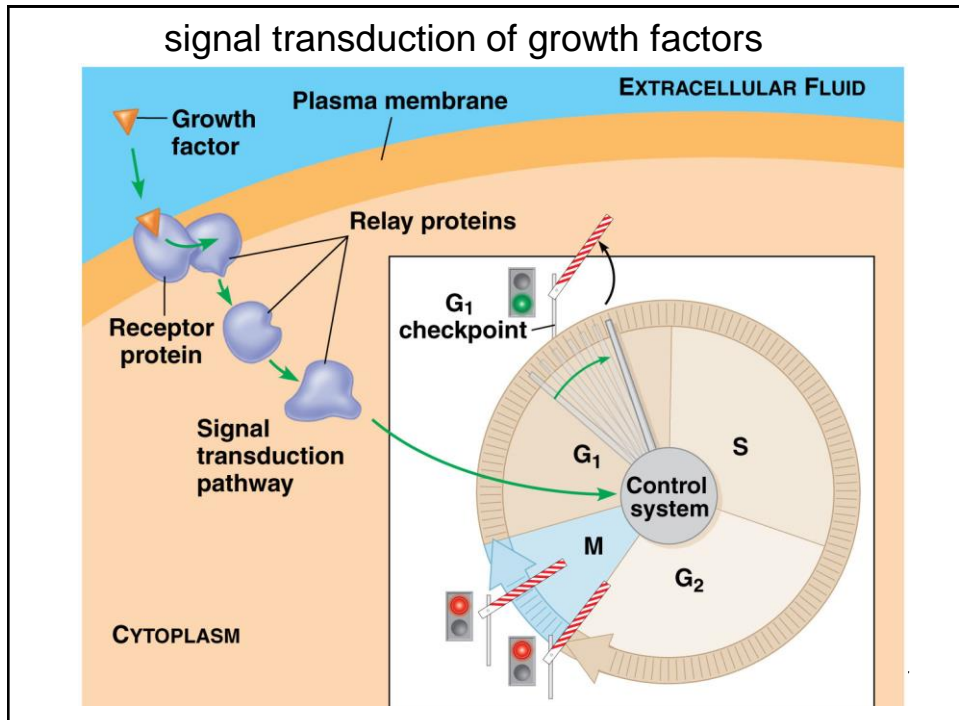
- cell size,
 - events of the cycle

45

effect of growth factors on cell divisions



46



2) checkpoints

- **sense and execute**

A. cell-cycle control system responds to signals

- outside:

nutrients,
growth factors,
anchorage dependence,
density-dependent inhibition

- inside:

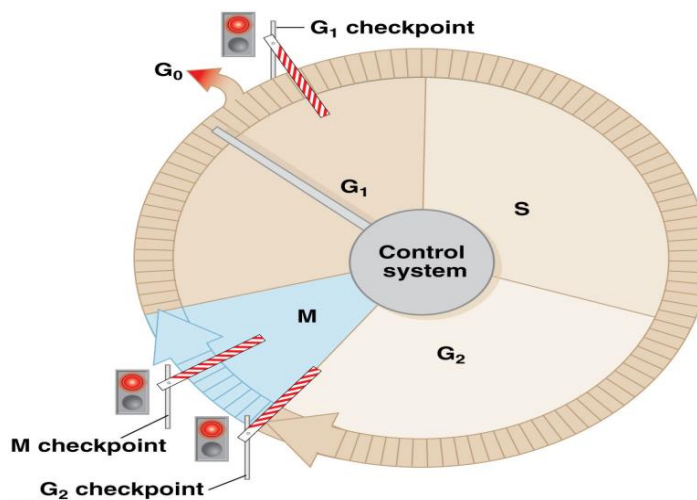
cell size,
events of the cycle

49

B. where and how to control

mostly, **built-in stop signal**, negative regulation

-> block, until overridden by go-ahead signal



50

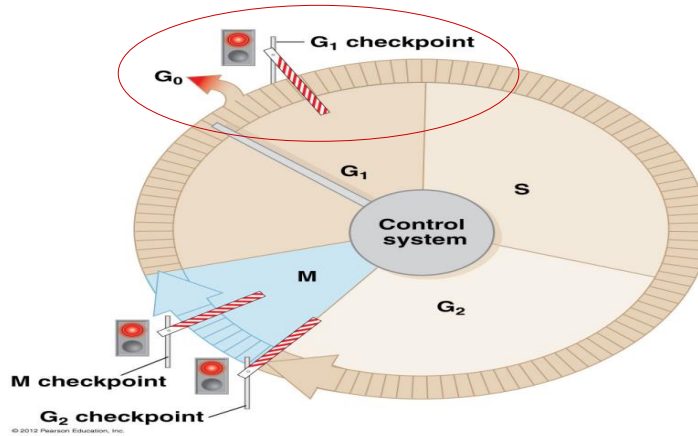
- major checkpoint:

a. **G1/S** start cell cycle?

START in yeast,

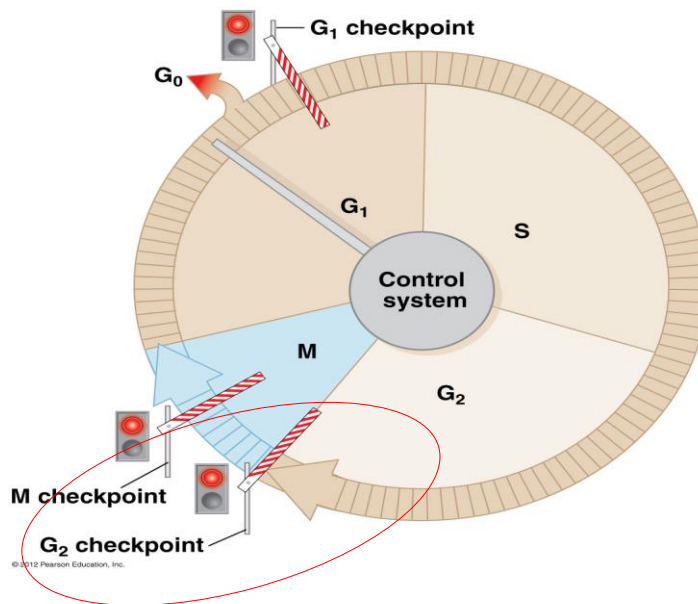
restriction point (**R point**) in animals

G0: exit from cell cycle, e.g., nerve cells



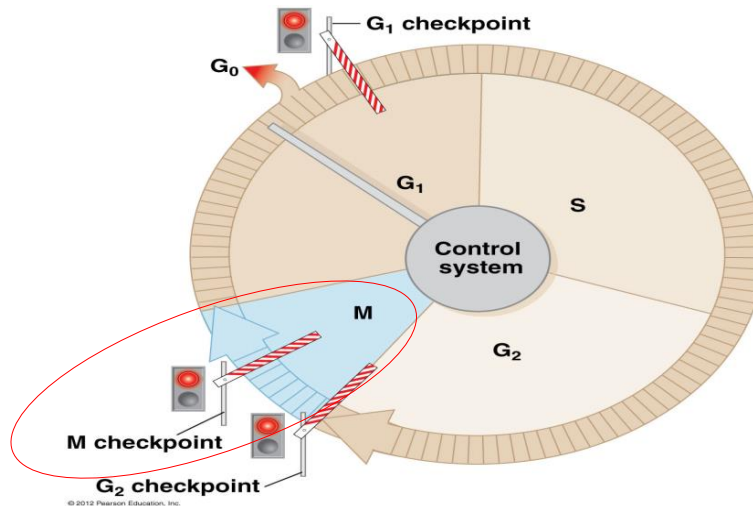
51

b. **G2/M** DNA replication complete?



52

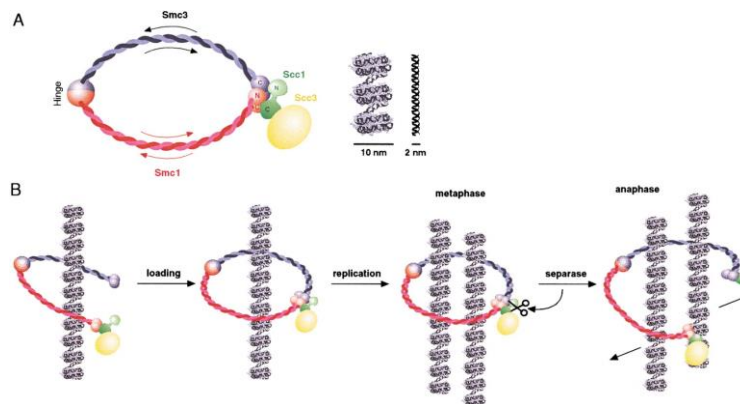
c. **metaphase/anaphase (spindle checkpoint)**
kinetochores attach to spindle microtubule?



53

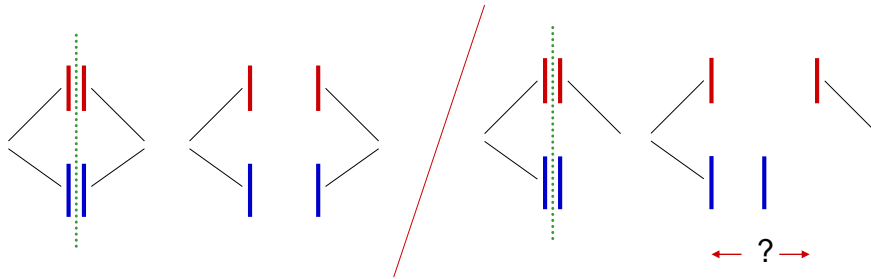
anaphase:

- sister chromatids separate
(**proteolysis of cohesin complex**)
- > daughter chromosomes
- > move toward opposite poles



54

- c. metaphase -> anaphase checkpoint,
 - if anaphase is initiated before kinetochores become properly attached to microtubules
 -> **nondisjunction**, daughter cells have missing or extra chromosomes



55

Checkpoints ensure the proper order of events in mitotic cell cycle by arresting or delaying in response to defects in cellular process.

Therefore, the integrity of genetic information is maintained

-> **to prevent failure and inviability in cell division**

56

- 3). cancer, a cell cycle disease
- single cell -> transformation (mutation)
 - > cell cycle out of control
 - > division
 - > excessive growth of cells
 - > tumor

57



The Nobel Prize in Physiology or Medicine 2001

for their discoveries of key regulators of the cell cycle



Leland H. Hartwell

Fred Hutchinson
Cancer Research
Center
Seattle, WA, USA
b. 1939



Sir R. Timothy Hunt

Imperial Cancer
Research Fund
London, United
Kingdom
b. 1943



Sir Paul M. Nurse

Imperial Cancer
Research Fund
London, United
Kingdom
b. 1949

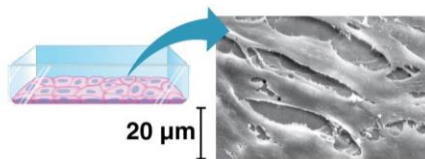
58

3). cancer, a cell cycle disease

- single cell -> transformation (mutation)
 - > cell cycle out of control
 - > division
 - > excessive growth of cells
 - > tumor

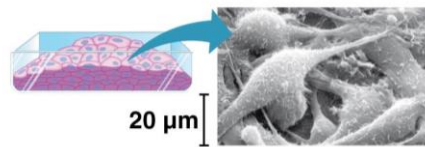
59

- immortal, don't need growth factor,
no density inhibition, no anchorage dependence,
no **apoptosis** 細胞凋亡



(a) Normal mammalian cells

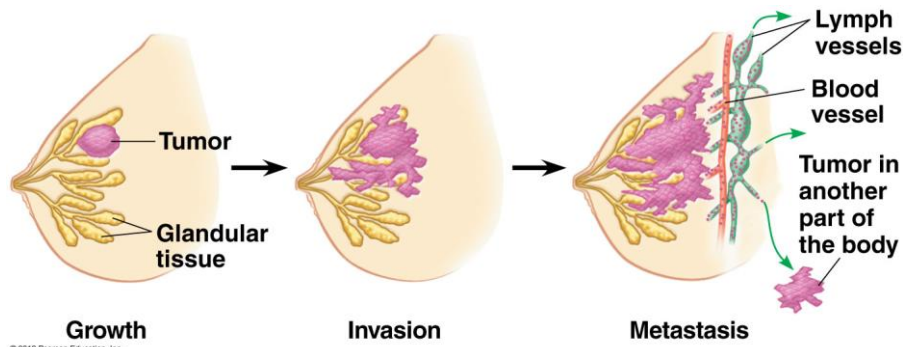
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(b) Cancer cells

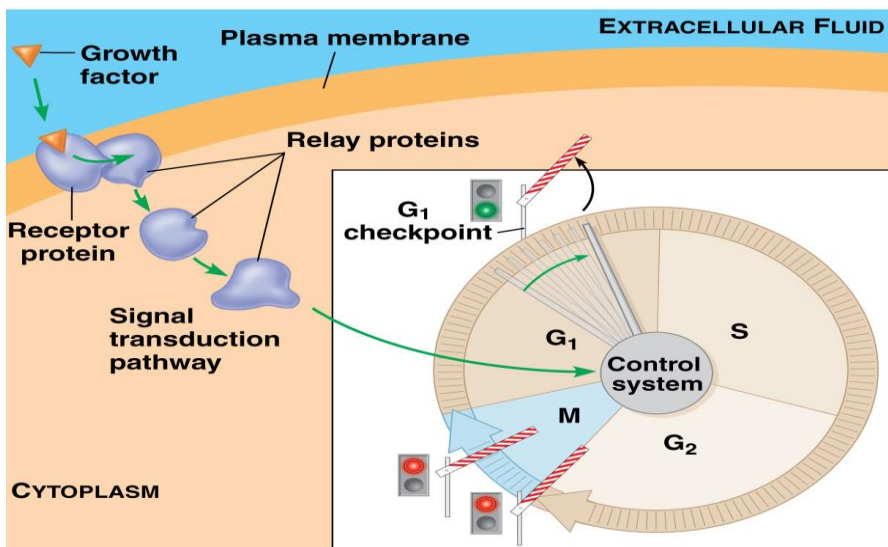
60

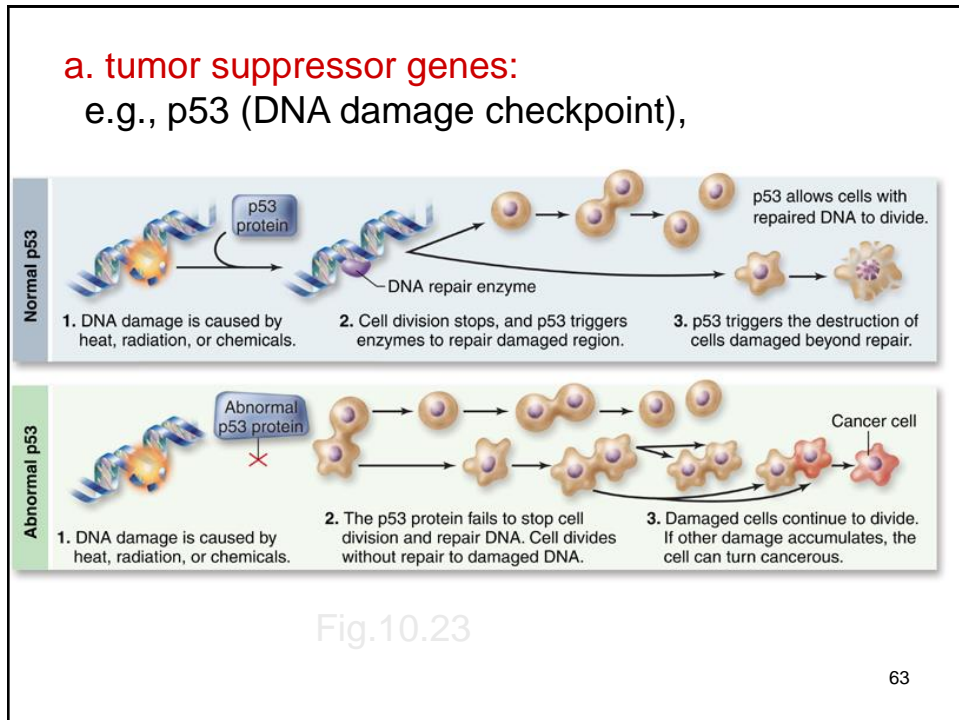
- a. benign tumor: remain at the original site
- b. malignant tumor:
 - > spreading into neighboring and other tissue
 - > displace normal cells -> cancer



61

- causes: defects (**mutations**) in the cell-cycle control system, e.g., receptor, messenger, checkpoints





- treatments

a. localized tumor:

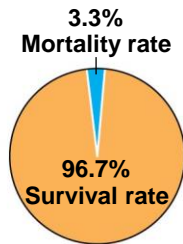
- surgery or radiation

b. spreaded tumor:

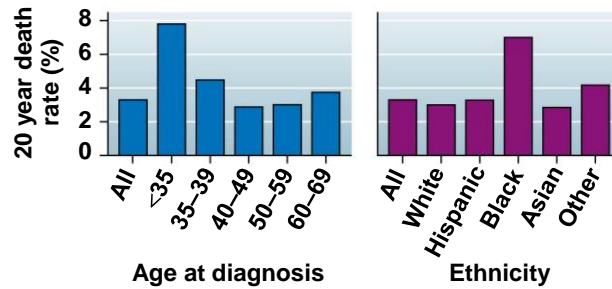
- chemotherapy
 - > disrupt specific steps in cell cycle,
e.g., Taxol -> freeze spindle
 - > side effect, also target on normal cells

the breast cancer treatment may vary by individual
 -> **personalized therapy**

Risk of death from
breast cancer 20 years
after DCIS diagnosis



Breast cancer death rates
for subsets of DCIS patients



DCIS: ductal carcinoma in situ 乳腺管原位癌