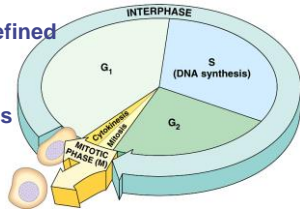




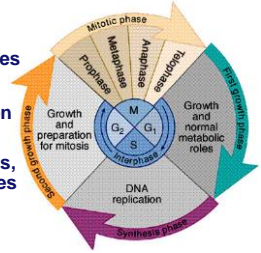
### Interphase

- 90+% of cell life cycle
  - cell doing its “everyday job”
    - produce RNA, synthesize proteins
  - prepares for duplication if triggered
- Characteristics
  - nucleus well-defined
  - DNA loosely packed in long chromatin fibers



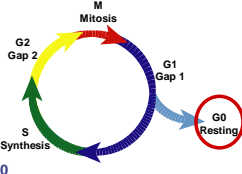
### Interphase

- Divided into 3 phases:
  - $G_1 = 1^{st}$  Gap
    - cell doing its “everyday job”
    - cell grows
  - S = DNA Synthesis**
    - copies chromosomes
  - $G_2 = 2^{nd}$  Gap
    - prepares for division
    - cell grows
    - produces organelles, proteins, membranes



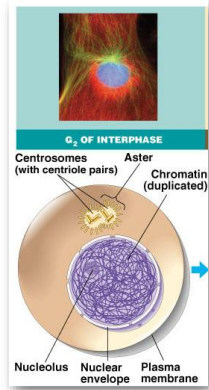
### $G_0$ phase

- $G_0$  phase
  - non-dividing, differentiated state
  - most human cells in  $G_0$  phase
    - nerve & muscle cells
      - highly specialized; arrested in  $G_0$  and can never divide!
    - liver cells
      - in  $G_0$ , but can be “called back” to cell cycle by external cues



### Interphase $G_2$

- Nucleus well-defined
  - chromosome duplication complete
  - DNA loosely packed (more or less) in long chromatin fibers
- Prepares for mitosis
  - produces proteins & organelles



### Cell Cycle

**Gap 2** - the cell will continue to grow and produce new proteins required for cell division. At the end of this gap is another control checkpoint (G2 Checkpoint) to determine if the cell can now proceed to enter M (mitosis) and divide.

**Gap 1** - Cells increase in size, produce RNA and synthesize proteins. An important cell cycle control mechanism activated during this period (G1 Checkpoint) ensures that everything is ready for DNA synthesis.

**Gap 0** - There are times when a cell will leave the cycle and quit dividing. This may be a temporary resting period (i.e. liver cell, or more permanent, i.e. a cell that has reached an end stage of development and will no longer divide (e.g. nerve cells in the brain).

**S phase (DNA synthesis)**

**Synthesis phase** - DNA replication takes place

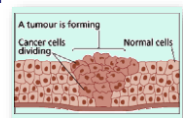
**Interphase =  $G_1 + S + G_2$**

**M (mitosis)**

Cells that ceases division

### Coordination of Cell Cycle

- Multicellular organism
  - need to coordinate across different parts of organism
    - timing of cell division
    - rates of cell division
  - crucial for normal growth, development & maintenance
    - do all cells have same cell cycle?

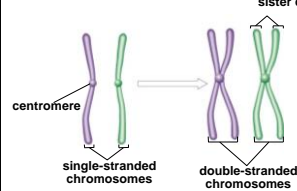


### Frequency of Cell Cycle

- Frequency of cell division varies with cell type
  - ◆ skin cells
    - divide frequently throughout life
  - ◆ liver cells
    - retain ability to divide, but keep it in reserve
  - ◆ mature nerve cells & muscle cells
    - do not divide at all after maturity

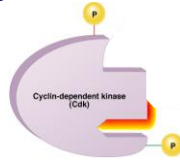
### Cell Cycle Control

- Cell cycle can be put on hold at specific **checkpoints**
- **Irreversible** points in cell cycle
  - ◆ replication of genetic material
  - ◆ separation of sister chromatids



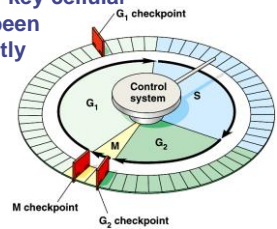
### “Go-ahead” signals

- Signals that promote cell growth & division
  - ◆ **intracellular** signals
    - “promoting factors”
  - ◆ **extracellular** signals
    - “growth factors”
- Primary mechanism of control
  - ◆ phosphorylation
    - kinase enzymes



### Checkpoint control system

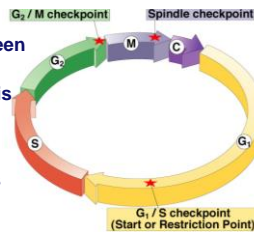
- Checkpoints
  - ◆ cell cycle controlled by **STOP & GO** chemical signals at critical points
  - ◆ signals indicate if key cellular processes have been completed correctly



### Checkpoint control system

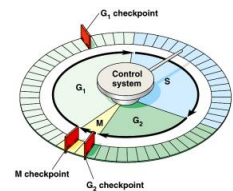
- 3 major checkpoints:

- ◆ G<sub>1</sub>
  - can DNA synthesis begin?
- ◆ G<sub>2</sub>
  - has DNA synthesis been completed correctly?
  - commitment to mitosis
- ◆ M phases
  - spindle checkpoint
  - can sister chromatids separate correctly?



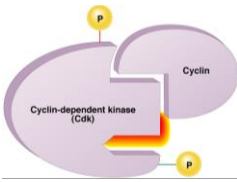
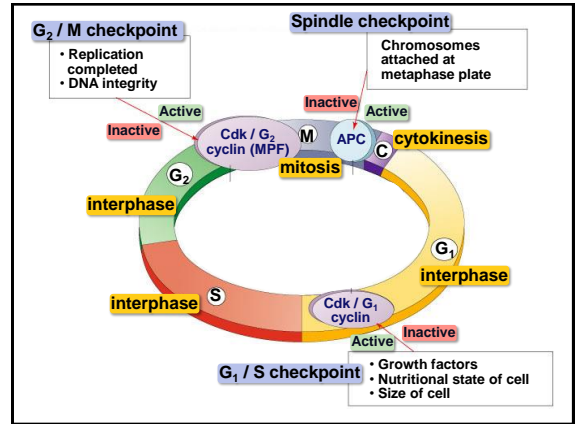
### G<sub>1</sub> checkpoint

- G<sub>1</sub> checkpoint is critical
  - ◆ primary decision point
    - “restriction point”
  - ◆ if cell receives “go” **signal**, it continues on...
  - ◆ if does **not** receive “go” signal, cell exits cycle & switches to **G<sub>0</sub> phase**
    - non-dividing state



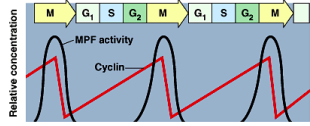
### Intracellular signals


- Promoting factors**
  - Cyclins**
    - regulatory proteins
    - levels cycle in the cell
  - Cdks**
    - cyclin-dependent kinases
    - enzyme activates cellular proteins
- MPF (for G<sub>2</sub> checkpoint):**  
maturation/mitosis promoting factor
- APC (for M checkpoint):**  
anaphase promoting complex


### Cyclins & Cdks 1970s-'80s | 2001

- Interaction of Cdks & different Cyclins triggers the stages of the cell cycle.**






Leland H. Hartwell  
checkpoints



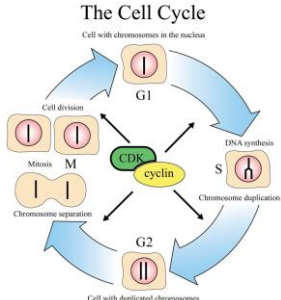
Tim Hunt  
Cdks



Sir Paul Nurse  
cyclins

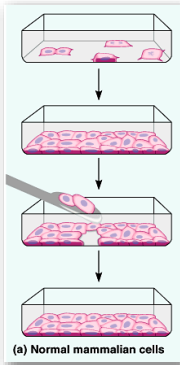
### Intracellular Signals

- CDKs & cyclin drive cell from one phase to next in cell cycle**
  - proper regulation of cell cycle is so key to life that the genes for these regulatory proteins have been **highly conserved through evolution**
  - the genes are basically the same in yeast, insects, plants & animals (including humans)



### Extracellular Signals

- Growth factors**
  - external signals
  - protein signals released by body cells that stimulate other cells to divide
  - density-dependent inhibition**
    - crowded cells stop dividing
    - mass of cells use up growth factors
    - not enough left to trigger cell division

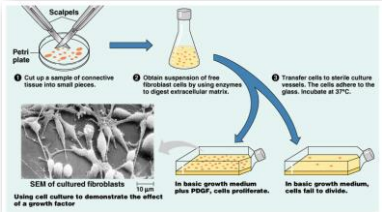


(a) Normal mammalian cells

### Example of a Growth Factor

- Platelet Derived Growth Factor (PDGF)**
  - made by platelets (blood cells)
  - binding of PDGF to cell receptors stimulates fibroblast cell division

**Growth of fibroblast cells (connective tissue cells) helps heal wounds!**



1 Cut up a sample of connective tissue into small pieces.

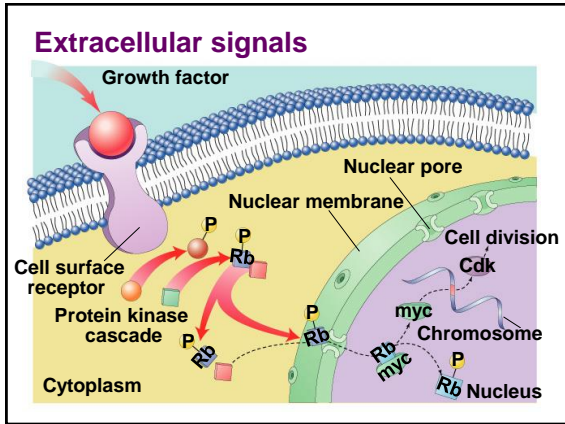
2 Obtain suspension of free fibroblast cells by using enzymes to digest extracellular matrix.

3 Transfer cells to sterile culture vessels. The cells adhere to the glass. Incubate at 37°C.

SEM of cultured fibroblasts. Using cell culture to demonstrate the effect of a growth factor.

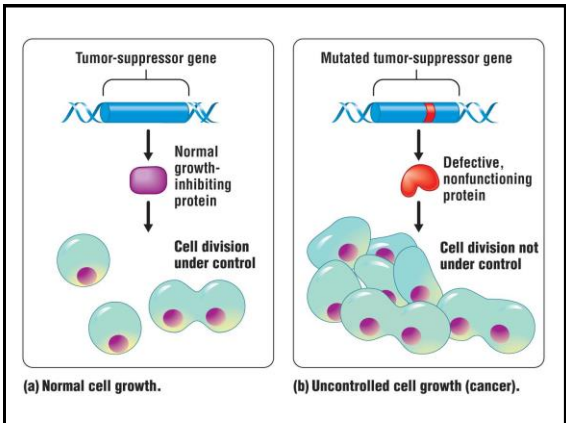
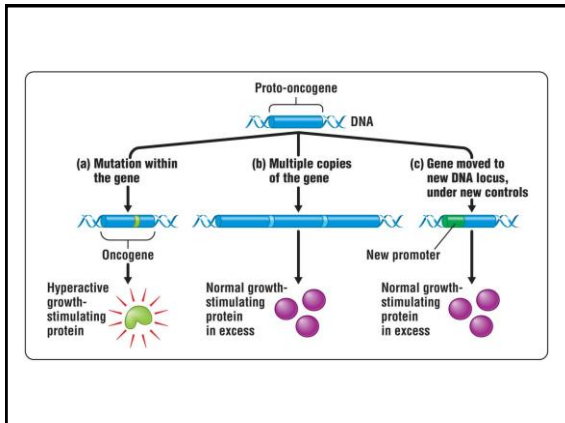
In basic growth medium, cells fail to divide.

In basic growth medium plus PDGF, cells proliferate.



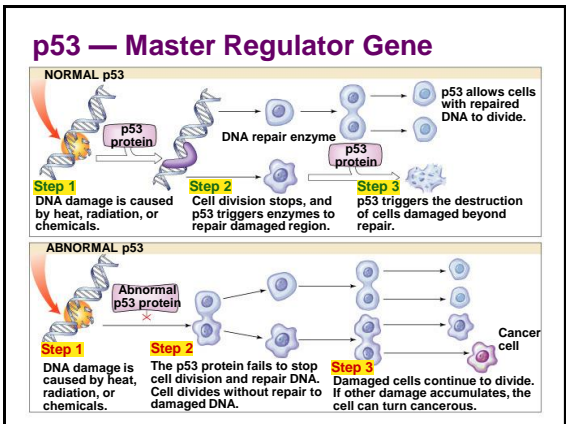
### Growth Factors, Genes, and Cancer

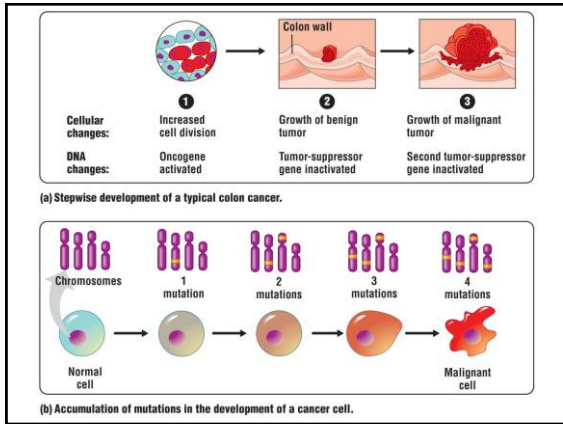
- Cancer is a “genetic” disease...
  - ♦ **proto-oncogenes**
    - normal genes that become oncogenes (cancer-causing) when mutated
    - stimulates cell growth
    - if switched **on** or **increased expression** can cause cancer
    - example: RAS (activates cyclin production)
  - ♦ **tumor-suppressor genes**
    - inhibits cell division
    - if switched **off** can cause cancer
    - example: p53



### Cancer & Cell Growth

- Cancer is essentially a failure of cell division control
  - ♦ unrestrained, uncontrolled cell growth
- What control is lost?
  - ♦ **checkpoint stops**
  - ♦ gene **p53** plays a key role in G<sub>1</sub> checkpoint
    - p53 protein halts cell division if it detects damaged DNA
      - ♦ stimulates repair enzymes to fix DNA
      - ♦ forces and keeps cell in G<sub>0</sub> resting stage
      - ♦ causes apoptosis of severely damaged cell
    - **MOST** cancers have to shut down p53 activity





### Development of Cancer

- Cancer develops only after a cell line experiences ~6 key mutations (“hits”)**
  - unlimited growth
    - turn **on** oncogenes
  - ignore checkpoints
    - turn **off** tumor suppressor genes
  - escape apoptosis
    - turn **off** programmed cell death genes
  - immortality = unlimited divisions
    - turn **on** chromosome maintenance genes
  - promotes blood vessel growth
    - turn **on** blood vessel growth genes
  - overcome anchor & density dependence
    - turn **off** “touch sensor” gene

**(b) Cancer cells**

Cancer cells do not exhibit anchorage dependence or density-dependent inhibition.

### What causes these “hits”?

- Mutations in cells can be triggered by:**
  - UV radiation
  - chemical exposure
  - radiation exposure
  - heat
  - cigarette smoke
  - pollution
  - age
  - genetics

**1** A tumor grows from a single cancer cell. **2** Cancer cells invade neighboring tissue. **3** Cancer cells spread through lymph and blood vessels to other parts of the body.

### Tumors

- Mass of abnormal cells**
  - Benign tumor (not totally safe...)**
    - abnormal cells remain at original site as a lump
      - p53 has halted cell divisions
    - still have properties of ‘original’ tissue
    - most do not cause serious problems & can be removed by surgery
  - Malignant tumors**
    - cells leave original site
      - lose attachment to nearby cells
      - carried by blood & lymph system to other tissues
      - start more tumors = metastasis
    - no longer resembles ‘original’ tissue
    - impair functions of organs throughout body

### Traditional treatments for cancers

- Treatments target rapidly dividing cells**
  - high-energy radiation & chemotherapy with toxic drugs
    - kill rapidly dividing cells at expense of healthy cells

### New “miracle drugs”

- Drugs targeting proteins (enzymes) found only in SPECIFIC tumor cells**
  - Gleevec**
    - treatment for adult leukemia (CML) & stomach cancer (GIST)
    - 1st successful targeted drug

**Gleevec: HOW IT WORKS**