

## **Chpt 5: the Cell Cycle**

the life of a cell

you started out life as a single cell

a fertilized egg

1 cell--->2 cells---->4 cells---->8 cells----->

16 cells----->ball of cells

this was rapid cell division w/no growth

then cell specialization began

nerve cells, muscle cells, blood cells, etc

every cell in body has same genetic material  
in nucleus regardless of type of cell

at birth--->complex multicellular organism

but you still needed to grow & develop

-growth involves cell division

-development involves turning on specific  
genes that regulate specific  
processes

## How do cells divide & how do they know when to divide?

it is programmed into genetic material

You are made up of trillions of cells

why not have less cells, but larger cells?

cells are limited by **SA to volume ratio**

cells must get O<sub>2</sub> & nutrients, get rid of CO<sub>2</sub> & wastes, all this has to move across surface of cell membrane

larger cells have larger volume, smaller

cells have better SA to volume ratio

**as cells grow, SA doubles as volume triples**, so you don't have enough surface for necessary items that are needed by the cell

also nucleus must control entire cell, if

contents get too big, nucleus has trouble directing cell

## **Cell cycle: phases in life of cell**

cell spends majority of time in growth & development, little time dividing

## **Purposes of cell division**

- 1-growth
- 2-replace worn out cells
- 3-repair tissues

## **Cell cycle: IPMAT**

interphase  
prophase  
metaphase  
anaphase  
telophase

**mitosis** = division of genetic material

prophase, metaphase, anaphase, telophase

**cytokinesis** = division of cytoplasm

this happens by the end of telophase

every cell must have complete copy of genetic material, thus **must copy DNA prior to mitosis**

**DNA** is found in several forms in cell depending on phase cell is in  
prior to division, DNA is in long strands called **chromatin**

after copying, chromatin condenses (shortens & thickens) into **double-stranded chromosomes** (proteins help do this)

each strand is copy of other called **sister chromatids**

chromatids are connected by **centromeres**

## **Cell Cycle**

**Interphase:** longest stage of cycle

cells spend approx 75-90% of life here  
used to be called "resting phase"

bad name, cell is carrying out all life processes  
growth & development, cell respiration,  
biosynthesis, active transport, passive  
transport, protein synthesis, etc.

divided into 3 parts

**1-G1** Gap1 or Growth

this is the first phase in the life of a  
newborn cell

the cell grows by increasing amount of  
cytoplasm, increasing # cellular  
organelles, the cell makes proteins,  
the cell obtains nutrients & O<sub>2</sub>, gets  
rid of wastes & CO<sub>2</sub>, these 2  
processes occur by passive &  
active transport

cells can stop in this phase called G<sub>0</sub>  
(zero)

your liver cells and nerve cells  
spend their lives in this phase  
most cells in adult multicellular  
organisms are in G<sub>0</sub>, these are  
metabolically active cells,  
maintaining the life of the cell  
and the organism

when cells get signal to divide, they  
move into next phase

**2-S phase:** Synthesis

copy genetic material, DNA  
copy centrioles

**3-G2 phase:** Gap2

cells get ready for mitosis  
make necessary proteins & RNA

Growth occurs in all subphases of interphase  
growth includes making proteins that are  
necessary for each stage

**Mitosis:** division of nucleus, series of steps ensuring that each daughter cell gets copy of genetic material

**Cytokinesis:** division of cytoplasm  
most cells exhibit cytokinesis, but muscle cells are an exception, this results in multinucleated cells that are very long

What happens in each stage of cell cycle? What do cells look like?

**Interphase:** nucleus present, nucleolus present  
DNA in form of chromatin, long strands

**Prophase:** DNA shortens and condenses into double-stranded chromosomes  
nucleus disappears, nucleolus disappears  
centrioles appear and begin migrating to opposite poles of cell, as they migrate, they set up the mitotic spindle  
these are microtubules that will attach to the chromosomes

**Metaphase:** centrioles arrive at opposite poles of the cell with the spindle fibers stretching out between them  
chromosomes attach to the mitotic spindle at the centromere of the double-stranded chromosomes  
the double-stranded chromosomes line up in center of cell, middle plate

**Anaphase:** centromeres divide resulting in sister chromatids separating  
sister chromatids are pulled to opposite poles by spindle fibers  
this is the shortest phase of mitosis

**Telophase:** chromatids reach opposite poles, now called single-stranded chromosomes  
spindle disassembles, nuclear membrane begins reappearing

**Cytokinesis** occurs  
in animal cells this is done by cleavage  
cells pinch in and pull apart  
in plant cells a cell plate forms

## **Differences in Mitosis: Plant vs Animal Cells**

plant cells do not have centrioles, the mitotic spindle is set by centrosomes and asters

### **cytokinesis**

**animal cells pinch in & pull apart**

**plants must begin making cell wall**

vesicles containing cellulose appear between 2 nuclei

vesicles then fuse together---->forms new cell membrane

cellulose inside vesicles forms new cell wall

forms from edges to inside



## **What controls mitosis?**

mitosis is a highly regulated process, things could go wrong at many places  
cells could duplicate chromatin twice  
cells could divide before genetic material is copied  
cells could divide when they are too small  
spindle fibers could fail to attach to chromo

Cell Fusion Experiments have been done to figure out what controls mitosis  
in cell fusion, you fuse 2 cells together  
removing the nucleus of one cell, then see what the cell does  
remember the nucleus is the control center, it tells the cell what to do

for example, if you take cells in S phase and remove the nucleus, then fuse them with cells in G1 (these cells have their nucleus intact)---->the DNA immediately begins to replicate  
thus, there is a factor in S phase cells that enters the G1 nucleus & initiates DNA replication

if you take G2 cells, remove nucleus, then fuse w/ early S phase cells----->these cells immediately enter G2 without DNA replication  
thus a factor from G2 cells is sending these cells into mitosis immediately

so what is going on? cyclins are regulating

**Cyclins** are proteins that regulate the cell cycle  
these are proteins are made just prior to  
each stage of the cycle and they tell the  
cell what to do  
the instructions for the proteins are in the  
DNA in the nucleus, but then proteins  
are made in the cytoplasm of the cell

G1 cyclins are made and begin accumulating  
in late G1---->these tell cell to begin S phase  
these reach peak during S phase, then drop  
mitotic cyclins are made and begin  
accumulating at end of S phase----->these  
tell cell to begin mitosis  
these peak at metaphase, then drop

cyclins stimulate every part of mitosis  
some cyclins turn on spindle formation  
some cyclins activate enzymes that break  
down centromeres so sister  
chromatids can separate at anaphase

So what if something still goes wrong?

UV radiation can damage DNA

what if sister chromatids don't attach to spindle?

the cell has **checkpoints** where the cell can be stopped, can make repairs, then proceed

there are checkpoints in G1, S, G2, & M

at these spots, proteins can detect if there is a problem, can stop cell cycle, put cell into **Cell-Cycle Arrest**, correct mistake & then restart cycle

for example, p53 is protein that detects DNA damage

when p53 detects damage, it activates inhibitors which prevent G1 cyclins from working

when the damage is repaired, then p53 is inactivated and the cell cycle proceeds

**Cancer** is cell division out of control

"cells gone wild"

cell cycle regulators prevent cells from leaving G0, but if regulators are inactivated, cells can divide at wrong time

when checkpoint controls are damaged, cells repeatedly divide forming mass of cells--> tumor

benign tumor: cells stay together, cause little harm, can be removed surgically

malignant tumor: cells break free & migrate to new areas called metastasis  
these cells can stimulate other cells to divide out of control

Everyone has the potential to have cancerous cells because the root lies within our genes

We all have genes in our cells called **proto-oncogenes**

these genes promote or turn on cell division  
there are also genes called **tumor-suppressor genes**: these genes inhibit cell division

in normal life, an external signal arrives at the cell to say it is time to grow, this turns on the proto-oncogene

this gene tells the cell to make cyclins to regulate cell cycle and send the cell into mitosis

then tumor suppressor genes turn on, which says stop after division, may put cell into G0

in cancerous cells, one or both of these processes may be interrupted

a mutation can change a gene from a proto-oncogene to an oncogene

oncogenes turn on cell division & don't listen to signals saying to turn off division

a mutation in tumor suppressor genes can result in them not making proteins that tell division to stop

both of these allow uncontrolled growth