Name:	Date:
-	

Genetic material duplicates.

Guided Notes

Unit 4: Cellular Reproduction

Chapter 5: Cell Growth and Division

Ι. Background

- a. "Where a cell exists, there must have been a preexisting cell...." Rudolf Virchow
- b. The division of cells into more cells enables living things:
 - i. To repair damage
 - ii. To grow
 - iii. To produce offspring
- c. Cell Division
 - i. Two Types

1. _____

- a. allows for cell reproduction leading to growth and repair
 - i. Example: Cell reproduction through mitosis enables your body to produce new skin cells that replace dead cells at your skin's surface.
- 2. ____
 - a. allows for sexual reproduction
 - i. Example: Cell reproduction through meiosis enables multicellular organisms to produce cells that are necessary for sexual reproduction, such as sperm and egg cells in animals.

II. Concept 5.1: The Cell Cycle

a. Cell Cycle



- b. Interphase
 - i. 90% of a cell's life is spent in interphase.
 - ii. Interphase: the stage during which a cell _____
 - iii. $G_1 \rightarrow S \rightarrow G_2$
 - 1. Gap 1 (G₁) phase: _____
 - 2. DNA Synthesis (S) phase: _____
 - 3. Gap 2 (G₂) phase: additional growth each duplicated chromosome remains loosely packed as chromatin fibers

c. Mitotic Phase

- i. Mitotic phase (M phase): stage of the cell cycle when the cell is actually dividing; _____
 - 1. Mitosis: process where the nucleus and the duplicated chromosomes divide and are evenly distributed, forming two "daughter" nuclei
 - Cytokinesis: process by which the cytoplasm is divided in two; usually begins before mitosis is completed



- ii. The combination of mitosis and cytokinesis produces _____
- iii. Mitosis is unique to eukaryotes.
- iv. The rate of cell division varies with the need for those types of cells.
- v. Some cells are unlikely to divide or never divide and remain in a stage called ______.

FIGURE 5.2 CELL DIVISION			
CELL TYPE	APPROXIMATE LIFE SPAN		
Skin cell	2 weeks		
Red blood cell	4 months		
Liver cell	300–500 days		
Intestine—internal lining	4–5 days		
Intestine—muscle and other tissues	16 years		

III. Concept 5.2: Mitosis and Cytokinesis

- a. Overview
 - i. Interphase: chromatin of each chromosome doubles, normal growth and cell functions occur (90% of cell's life)
 - ii. Mitotic phase: takes place rapidly, distributing the duplicate sets of chromosomes to two daughter nuclei
 - iii. Cytokinesis: divides the cytoplasm, producing two daughter cells
- b. Chromosomes
 - i. Chromatin:
 - ii. Centromere: _____
 - iii. Telomere: _____
 - iv. Chromosomes: _____

1. (Each chromosome may contain many hundreds of genes.)

v. Sister chromatids:

1. (Before cell division begins, a cell duplicates all of its chromosomes to make these.)



vi. Once separated from its sister, each chromatid is considered a full-fledged chromosome.



- c. Human Cells
 - i. Humans have _____ pairs of chromosomes in their cells. (So you have _____ chromosomes in total.)
- d. The Mitotic Phase
 - i. Spindle: _____
 - ii. Centrosome: ____
- e. Mitosis occurs in four stages. Your textbook with reference these by name. However, in this class you do not need to memorize stage names, just what is occurring within mitosis as a whole.









- STAGE 2 (METAPHASE)

 Sister chromatids gather in a plane across the middle of the cell.
- The mitotic spindle is now fully formed.
- All the sister chromatids are attached to the spindle microtubules, with their centromeres lined up about halfway between the two ends, or poles, of the spindle.



STAGE 4 (TELOPHASE)

- The chromosomes reach the poles of the spindle.
- The spindle disappears.
- Two nuclear envelopes reform (one around each set of daughter chromosomes).
- The chromosomes uncoil and lengthen.
- The nucleoli reappear.

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f. Cytokinesis

i. Cytokinesis:

_, typically occurs during the fourth stage of mitosis (telophase)

- ii. In animal cells, the first sign of cytokinesis is the appearance of an indentation around the middle of the cell.
- iii. In plant cells, a disk containing cell wall material called a ______ forms inside the cell and grows outward.



IV. Concept 5.3: Regulation of the Cell Cycle

- a. The timing of cell division is critical to normal growth and development. When the "control system" for cell growth malfunctions, cells may reproduce at the wrong time or in the wrong place.
- b. Cell Cycle Regulation
 - i. External Factor
 - 1. Growth factors: _____
 - ii. Apoptosis
 - 1. Apoptosis: _____
 - The timing of cell division is critical to normal growth and development. When the "control system" for cell growth malfunctions, cells may reproduce at the wrong time or in the wrong place.

c. Tumors

i. Tumor: ______

(fast reproduction rate and irregular appearance)

- - be completely removed by surgery)
- 2. Malignant tumors: _____

_____ (more problematic than benign – they spread

to surrounding tissues)

d. Cancer

- Cancer: a disease caused by the severe disruption of the mechanisms that normally control the cell cycle; ______, which if unchecked can result in death
- ii. Metastasis:

______ - allowing for new tumors to form elsewhere (This is what makes

cancer so dangerous.)



- iii. How does cancer happen?
 - 1. Gene Mutations caused by: _____
 - a. Genetics: some gene mutations are inherited
 - b. Carcinogens: _____
 - 2. When genes are mutated, the cell cycle may speed up, causing a normal cell to no longer perform its normal functions and to divide and spread quickly – labeling it as a "cancer cell."

e. Cancer Treatment

- i. There is no single "cure" just multiple approaches that can help control or halt the progress of the disease.
- ii. Malignant tumors can be:
- iii. Radiation Therapy: exposing the parts of the body with cancerous tumors to high-energy radiation, which disrupts cell division
 - 1. Side effects: can damage cells of the ovaries or testes, causing sterility
- iv. Chemotherapy: treating the patient with anti-mitotic drugs that disrupt cell division
 - 1. Side effects: nausea or hair loss

V. Concept 5.4: Asexual Reproduction

- a. Types of Reproduction
 - i. Asexual Reproduction
 - 1. Usually unicellular organisms
 - 2. A single cell or group of cells each duplicates its genetic material and then splits into _____

_ (offspring inherit all

their genetic material from just one parent)

- 3. Example: Paramecium, some sea stars
- ii. Sexual Reproduction
 - 1. Usually multicellular organisms
 - 2. When two parents are involved in the production of offspring _____

(involves

the union of sex cells, such as an egg and a sperm)

- 3. Example: plants, animals
- b. Asexual Reproduction
 - i. Asexual reproduction: creation of offspring from a single parent
 - ii. Binary fission:

_____- occurs in prokaryotes

 Binary fission is very similar to mitosis. However, prokaryotes do not have a nucleus, so the division is distinctly different.



- c. Advantages and Disadvantages of Asexual Reproduction
 - i. Advantages

 ______ (often requires less time and energy)
 ______ (new traits could be harmful in some environments)

- ii. Disadvantages
 - 1. _____ (which

may be needed if environment changes)

Chapter 8: From DNA to Proteins

I. Concept 8.1: Identifying DNA as the Genetic Material

- a. The Scientists Who Discovered DNA
 - i. Watson & Crick
 - 1. James Watson and Francis Crick modeled DNA's structure with tin and wire.
 - Using the clues provided by Franklin's work, Watson and Crick created a model in which two strands of nucleotides wound about each other.
 - 3. This formed a twisting shape called a
 - a. This new model successfully represented DNA's structure.

II. Concept 8.2: Structure of DNA

- a. The Building Blocks of DNA
 - i. Deoxyribonucleic acid (DNA):
 - ii. Nucleotide: _______ (Nucleic acids are a polymer.)
 - 1. DNA uses four different types of nucleotides but the chemical structure of each type is very similar.
 - 2. Nucleotides have three parts:
 - a. A ring-shaped sugar called _____
 - b. A _____ group
 - c. A _____: a single or double ring of carbon and

nitrogen atoms with functional groups (nitrogenous means "nitrogen-containing")





- b. Nitrogenous Bases
 - i. The four nucleotides found in DNA differ only in their nitrogenous bases.
 - 1. single-ring structures:



Name of Base	Structural Formula	Model	Name of Base	Structural Formula	Model
thymine		T	adenine		A
cytosine		C	guanine		G

c. DNA Strands

- i. Nucleotide monomers join together by covalent bonds between the sugar of one nucleotide and the phosphate of the next, forming a _____.
 - 1. The nucleotides can combine in many different sequences.
 - The part of DNA shown has 9 nucleotides arranged in the order CTGCTATCG. This arrangement is only one of many possible.



d. DNA's Structure

- i. The Double Helix
 - Remember: James Watson and Francis Crick modeled DNA's 3-D structure with tin and wire that formed a twisting shape called a double helix.
 - 2. The _____



_____, forming the double helix.

- 3. Watson and Crick's discovery built on the work of Rosalind Franklin and Erwin Chargaff.
- ii. Complementary Base Pairs
 - Individual structures of the nitrogenous bases determine very specific pairings between the nucleotides of the two strands of the double helix. These pairings are due to the sizes of the bases and their abilities to form hydrogen bonds with each other.
 - a. The ______ pairs with the ______.
 - b. The ______ pairs with the ______.
 - i. (A is said to be "complementary" to T, and G is complementary to C.)
 - 2. Each base must pair up with its complementary base.



III. Concept 8.3: DNA Replication

- a. Template Mechanism
 - i. Remember: Cells duplicate their DNA in Interphase before Mitosis/Meiosis.
 - ii. How do cells duplicate DNA?
 - 1. DNA-copying uses a _____
 - a. This means it uses one part of the DNA to make the opposite side.
 - iii. How can cells do this?
 - 2. (Remember: A pairs with T, and G pairs with C.)
 - Know the sequence of bases on one strand of DNA → Determine the sequence on the other
 - iv. DNA replication: _____

1.

v. Daughter strands: _____



- b. Replication of the Double Helix
 - i. DNA replication is important when a cell wants to start mitosis.
 - ii. Without DNA replication, your DNA wouldn't be in every cell!

1