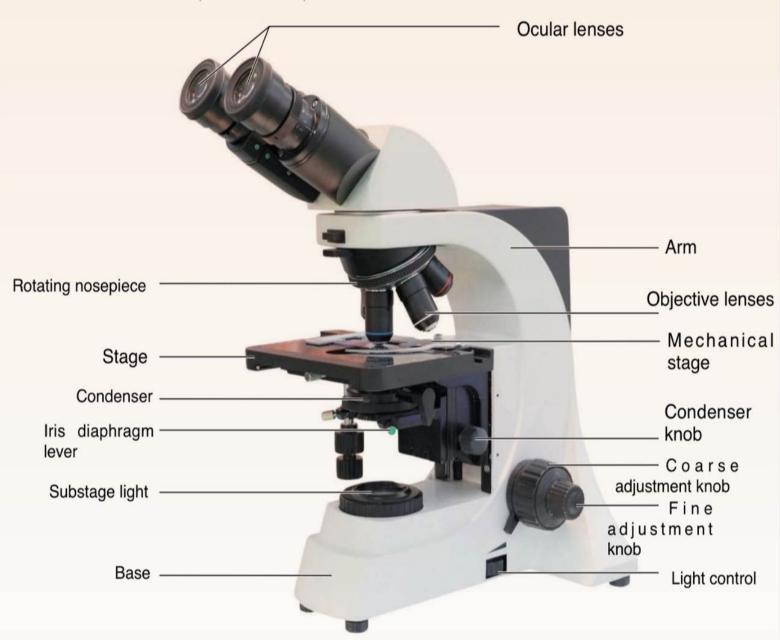


REVIEW SHEET The Microscope

Name Elvana Lleshaj-Gjoka Lab Time/Date Tuesday's 2:30-5:00 pm

Care and Structure of the Compound Microscope

1. Label all indicated parts of the microscope.



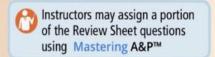
2. Explain the proper technique for transporting the microscope.

When transporting the microscope, should be held in an upright position, with one hand on its arm and the other supporting its base. The instrument should not be swinged during its transport or jar the instrument when setting it down.

| 3. | | | ents is either true or false. If true, write T on the anser word or phrase to replace the one that is underline | | nk. If false, correct the statement by | | | | |
|-----|---|---------------------|---|--|--|--|--|--|--|
| S | Special grit-free | lens pa | per 1. The microscope lens may be cleaned with | any sof | t tissue. | | | | |
| | Scanning obje | ctive le | ns 2. The microscope should be stored with the | oil imm | ersion lens in position over the stage. | | | | |
| | T | | 3. When beginning to focus, use the scanning | | | | | | |
| | Т | 1 | | | | | | | |
| | Т | | 4. When focusing on high power, always use | e the <u>co</u> | <u>arse</u> adjustment knob to focus. | | | | |
| | | | 5. A coverslip should always be used with w | | | | | | |
| 4. | Match the microscop Column A | oe structu | res in column B with the statements in column A tha | n column B with the statements in column A that identify or describe them. Column B | | | | | |
| | T T | | | Col | | | | | |
| | 1. | platform | on which the slide rests for viewing | a. -b . | coarse adjustment knob condenser | | | | |
| | <u> </u> | used to the spec | adjust the amount of light passing through men | 4. | fine adjustment knob iris diaphragm lever mechanical stage | | | | |
| | <u>E</u> 3. | controls | the movement of the slide on the stage | ≱. g. | nosepiece objective lenses | | | | |
| | 4. | delivers a | concentrated beam of light to the specimen | h. | ocular lens stage | | | | |
| | <u> </u> | used for been do | or precise focusing once initial focusing has | | | | | | |
| | <u> </u> | | the objective lenses; rotates so that the differ- ective lenses can be brought into position over cimena. | | | | | | |
| 5. | Define the following total magnification: | The to | tal magnification of any specimen bein ular lens multiplied by the power of the | | | | | | |
| | resolution: is t | the abil | ity to discriminate two close objects as | sepa | rate. | | | | |
| Vie | ewing Objects | Throug | h the Microscope | | | | | | |
| 6. | Complete, or respon | d to, the | following statements: | | | | | | |
| | Working distand | <u>ce</u> 1. | The distance from the bottom of the objective | lens to | the surface of the slide is called | | | | |
| | Diabt | | the | | | | | | |
| | Right 2. Ass | | Assume there is an object on the left side of the | field th | nat you want to bring to the center | | | | |
| | | | (that is, toward the apparent right). In what direction | n would | you move your slide? | | | | |
| | Field | 3. | | | | | | | |
| | 25, | 4. | If a microscope has a $10\times$ ocular lens and the total | l magnit | fication is 950 $	imes$, the objective lens in | | | | |
| | | | use at that time is $___$ $\times.$ | | | | | | |

| To provide more of | contrast for viewing the cells 5. | Why should the light be dim | med when looking at living | nearly transparent) cells? | |
|--|--|---------------------------------|--|---|--|
| | Parfocal | | | | |
| | 6. | if, after focusing in low pow | er, you need to use only the | fine adjustment to focus the specimen | |
| | 075 | at the higher powers, the m | croscope is said to be | <u>-</u> | |
| | 0,75 mm 7. | You are using a 10× ocular | and a 15 \times objective, and | the field diameter is 1.5 mm. The ap- | |
| | | proximate field size with a 3 | 0× objective is | _ mm. | |
| |).5 mm 8. | If the diameter of the low-p | ower field is 1.5 mm, an obj | ect that occupies approximately a third | |
| | | of that field has an estimate | d diameter of | _mm. | |
| 7. | You have been asked to prepa low-power field. | re a slide with the letter F on | it (as shown below). In the | circle below, draw the F as seen in the | |
| | | | | | |
| | | F | F | | |
| | | | | | |
| 8. | Estimate the length (longest d | mension) of the object in μm | | | |
| | | Text | | | |
| | | | | | |
| | Total magnification = 100× | | | | |
| | Field diameter = 1.6 mm | | | | |
| | Length of object = $\frac{16}{100}$ | μm | | | |
| 9. | Say you are observing an object | t in the low-power field. Wh | en you switch to high power | it is no longer in your field of view. | |
| | Why might this occur? Cha | | er lens narrows the out of focus, not ce | field view and the object wintered. | |
| | What should you do initially to | | D-1 | Salara Blakarana ara- | |
| | into a higher power lens d focus the object. | | | | |
| 10. Do the following factors increase or decrease as one moves to higher magnifications with the microscope? | | | | | |
| | resolution: Increa | ise a | mount of light needed: | Increase | |
| | working distance: | crease | depth of field: | Decrease | |
| 11. | | lens in position and appears t | | pecimen. The instructor, noting a work- | |
| | How so? H | igh-power lenses are used a s | horter working distances than | 1 1cm. | |
| | | | | | |

| 12. | Place a drop of saline in the center of a clean slide. After that, we place the object. Hold the coverslip at a 45 degree angle with fingertips and slowly lower it. | | | | | |
|-----|---|--|--|--|--|--|
| 13. | Indicate the probable cause of the following situations during use of a microscope. | | | | | |
| | a. Only half of the field is illuminated: The light path can be blocked and does not illuminate the field completely. | | | | | |
| | b. The visible field does not change as the mechanical stage is moved: | | | | | |
| | When something like this happens it can be a problem with the lens | | | | | |
| 14. | A blood smear is used to diagnose malaria. In patients with malaria, the protozoa can be found near and inside red blood cells. Explain why a microscope capable of high magnification and high resolution would be needed to diagnose malaria. | | | | | |
| | Because only under a microscope with the appropriate lenses can be | | | | | |
| | distinguished the Protozoa which can be found on the red blood cells. | | | | | |
| 15. | Histopathology is the use of microscopes to view tissues to diagnose and track the progression of diseases. Why are thin | | | | | |
| | slices of tissue ideal for this procedure? Thin slices of tissues are ideal because can allow that on the microscope can be seen the tissue and its components meticulously | | | | | |

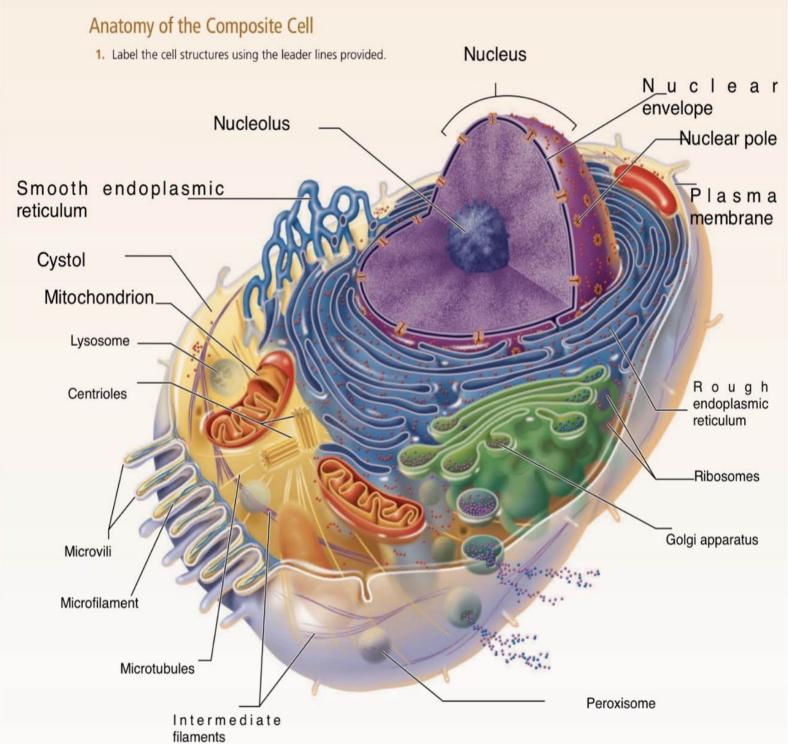




REVIEW SHEET

The Cell: Anatomy and Division

Name Elvana Lleshaj-Gjoka Lab Time/Date Tuesday's 2:30-5:00 pm.



| 2. | Match each | cell structure | listed on | the left v | with the correct | description on | the right. |
|----|------------|----------------|-----------|------------|------------------|----------------|------------|

| F_ | 1. | ribosome | <u>a</u> . | main site of ATP synthesis |
|----------------|----|-----------------|------------|---|
| _H_ | 2. | smooth ER | ф. | encloses the chromatin |
| _A_ | 3. | mitochondrion | pe. | sac of digestive enzymes |
| B | 4. | nucleus | N. | examples include glycogen granules and ingested foreign materials |
| | 5. | Golgi apparatus | بھ | forms basal bodies and helps direct mitotic spindle formation |
| | 6. | lysosome | 4 | site of protein synthesis |
| E, R | 7. | centriole | 18. | forms the external boundary of the cell |
| E | 8. | cytoskeleton | X | site of lipid synthesis |
| \overline{P} | 9. | inclusion | 1 | packaging site for ribosomes |
| <u> </u> | 0. | plasma membrane | / | packages proteins for transportation |
| I 1 | 1. | nucleolus | + | internal cellular network of rodlike structures |

Differences and Similarities in Cell Structure

| 3. | Choose the specimen | observed in | Activity 5 | (squamous | epithelium, | sperm cell | s, smooth | muscle, | or human | red | blood | cells) |
|----|---------------------------|-------------|------------|-----------|-------------|------------|-----------|---------|----------|-----|-------|--------|
| | that fits the description | helow | | | | | | | | | | |

| 1. | Sperm | cell has a flagellum for movement |
|----|-----------------------|---|
| 2. | Smooth | cells have an elongated shape (tapered at each end) |
| 3. | Squamous epithelium | cells are close together |
| 4. | Human red blood | cells are circular |
| 5. | Squamous | cells are thin and flat, with irregular borders |
| 6. | Human red blood cells | cells are anucleate (without a nucleus) |
| 7. | Smooth muscle | longest cell |

Cell Division

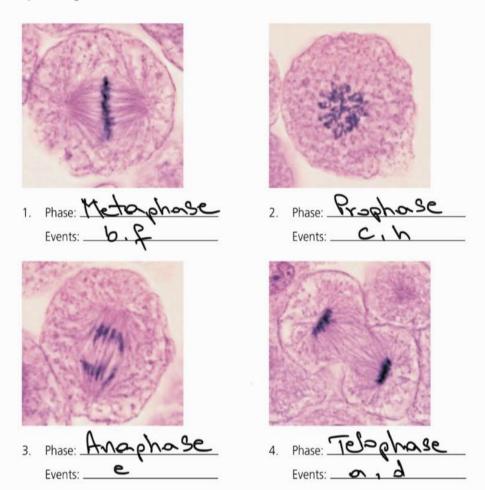
| 4 | What is the function of mitotic cell division? | |
|---|---|--|
| | What is the falletion of filltotic cell division: | |

Cell division is essential for growth and repair. Mitotic cell division is the period when the cell reproduces itself by dividing.

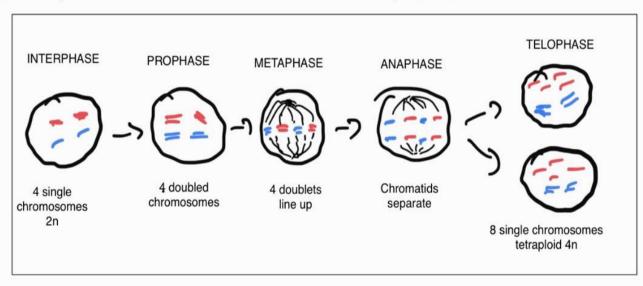
5. Identify the four phases of mitosis shown in the following photomicrographs, and select the events from the key that correctly identify each phase. On the appropriate answer line, write the letters that correspond to these events.

Key:

- a. The nuclear envelope re-forms.
- ර. Chromosomes line up in the center of the cell.
- Chromatin coils and condenses, forming chromosomes.
- d. Chromosomes stop moving toward the poles.
- F The chromosomes are V shaped.
- The nuclear envelope breaks down.
- Chromosomes attach to the spindle fibers.
- The mitotic spindle begins to form.



6. Draw the phases of mitosis for a cell that contains four chromosomes as its diploid, or 2n, number.



7. Describe the events that occur during interphase.

Interphase is the period when the cell carries out its normal metabolic activities and grows. The DNA- containing material is in the form of chromatin. The nuclear envelope and one or more nucleoli are intact and visible.

| | The DNA- containing material is in the form of chro | |
|-----|--|--|
| 8. | 8. Complete or respond to the following statements: | |
| | Division of the1 is referred to as mitosis. Cytokinesis is division of the2 The major structural difference between chromatin and chromosomes is that the latter are3 Chromosomes attach to the spindle fibers by undivided structures called4 If a cell undergoes mitosis but not cytokinesis, the product is5 The structure that acts as a scaffolding for chromosomal attachment and movement is called the67_ is the period of cell life when the cell is not involved in division. Three cell populations in the body that do not routinely undergo cell division are8_,9_, and10 | Cytoplasm Cytoplasm Condensed Centromeres Binucleate cell Spindle Spindle Therphase Steletal muscle Cordina muscle |
| | | Neurong |
| 9. | 9. Plasma cells are key to the immune response because they secrete | antibodies. Given that antibodies are made of protein, |
| | which membrane-enclosed cell organelle would you expect the plasma | cells to have in abundance? Why? |
| | Ribosomes, because they are particularly abundant in ce | ells that synthesize large amounts of protein. |
| 10. | 10. Hame which organelle you would expect to play the largest role in | decomposition of the human body. Why? |
| | Lysosomes, because they degrae | de. |
| 11. | 11. Some antifungal medications work by blocking DNA synthesis in the | fungal cell. Describe where in the cell cycle such a medi- |
| | cation would halt the fungal cell and the consequences of this early term | nination of the cycle |
| | This may occur during interphase, in the S phase which | is responsible for the synthesis of DNA. |
| | | |