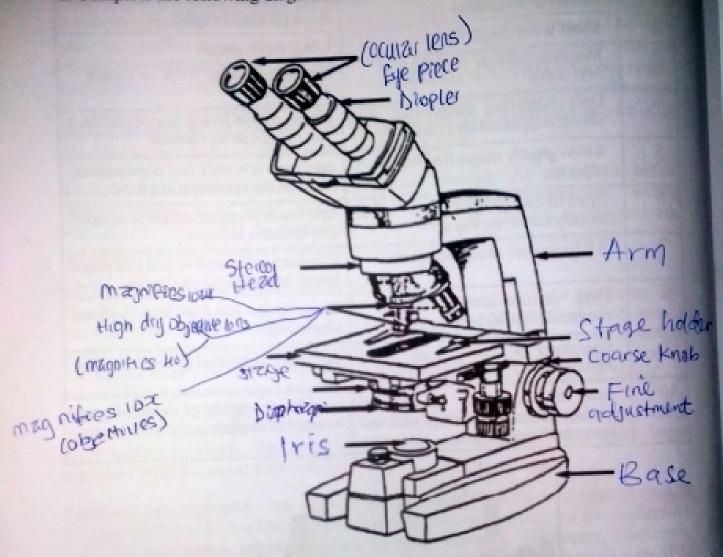
### Microscope Quiz Exercise

a. Complete the following diagram. Label the parts of the microscope



- Calculate the total magnification for each ocular/objective combination on your microscope.

- law	Objective	Total Magnification
Ocular	Hx	40x
10×	10x	100
10×	40.	400x
IOX.	102	1.000
104	1/2	
IOX		

d. List the magnification and numerical aperture for each objective on your microscope.

Magnification of Objective	Numerical Aperture (NA)
4×	4.45
10×	1.75
110 X	0.40
,noX	0.198

e.	What is the total magnification that you are using to examine a cell when you examine in ing it with the 40x objective The coular leps=10x The total magnification.	e?
	X. GX. G.	

- f. If 4 plant cells extend across the field of view when the observer is using 10X objective, what is the length of each cell?

  Lox Objective (Heelts Zaoss) length of the one cell.

  2 mon = 0.5 mm | Lox Objective 2 mm (1.76)
- If you were observing a live specimen under the 40x objective lens and it moved out of your field of view, explain how you could find it again quickly and easily. The observing area must be a white plan where you can easily. See and detect it but the observing area was a coloured easily. See and detect it but the observing area was a coloured easily. See and detect it but the observing area was a coloured easily. See and detect it but the observing area was a coloured easily. See and detect it but the observing area was a coloured easily.
- h. Microscopes that are \_\_\_\_ will remain nearly focused after the low-power objective lens is changed to the high-power objective lens. A), monocular. B), parcentered. C), parfocal. D), properly adjusted.
- i. At high power, always use which adjustment knob to focus the image?

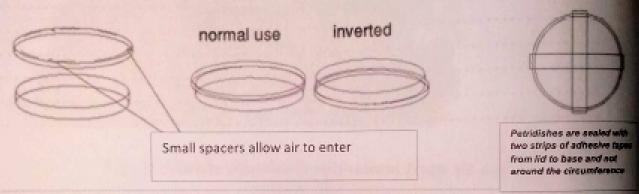
  At high power, always use the coarse Adjust ment knob.
- k. On the microscope stage, what is used to hold the glass slide in place and prevent it from moving. The part used to hold the glass. Sweet in place.

  Without moving is called a Chiff.

Which part of the microscope should you hold when lifting or moving the microscope? The only part one should hold and with when making ze
Transmitted light microscope has a light source below the stage while a reflected microscope has a light source that is located? microscope has a light source that is located? It is located just besde the objective lens where the auxiliary lensus. Can be attached
A stereo microscope is used to? Other the Durface of Solic Specimen of to Clary out work such as deserve, microsugen, watch making and Foreign engineering.  Which part of the microscope contains the shutter that regulates the amount of light entering the lens system? The Part which controls or regulates the amount of Plagnt is known as the Condenser lens.
A microscope is set to 10x eyepiece and 40x objective. What is the total magnification? The total magnification of 10x eye prece and Hox.  Objective as equal to Hook.
To obtain total magnification, you should the magnification of the eyepiece to the objective lens power: (a) Add (b) Multiply (c) Divide (d) Subtract.
हेब देव देव हैं है के क्षेत्र के किया के किया है

# LABORATORY EXERCISE FIVE MICROBIOLOGICAL TECHNIQUES

It is a known fact that any source of organic food which provides carbon compount for energy of respiration provides a suitable environment for Bacteria to grow. It study the growth of bacteria, two normal media are used and are a clear soup-like liquid nutrient broth, usually in tubes, and nutrient agar, which is set into a jelly by the addition of a seaweed extract called agar. When they are melted they are poured into glass or plastic Petri dishes(known as plates) as shown below.



Petri Dishes

#### 5.1 Microbiological Techniques

These consist of sterilization, aseptic techniques, inoculation, and incubation. The media must be sterilized by heating in an autoclave (like a pressure cooker) at 121°C (pressure 1 bar or 15 lb/sq. in.) for 15 minutes, which kills all living organisms, including spores. All apparatus used from this point onwards must be sterilized by heat (glassware - 160 °C for 2 hrs) or exposure to radiation. Aseptic techniques must be used to reduce the likelihood of bacterial contamination. This usually involves disinfection of working areas, minimizing possible access by bacteria from the air to exposed media, and use of flames to kill bacteria which might enter vessels as they are opened.

Cultures are usually examined after 24 hrs incubation. Liquid media such as broth become cloudy if bacteria are present. This could be the result of only one bacterial cell originally entering the medium, then dividing repeatedly to produce millions. Bacteria on agar "plates" become visible as distinct circular colonies; each colony should represent an individual bacterial cell (or group) which has divided repeatedly but, being kept in one place, the resulting cells have accumulated to form a visible patch.

7.	Record your observations in your practical book indicating the stain and organism used. Pay particular attention to the arrangements, shapes and relative sizes of the cells.
5.3	Review Questions
1.	Give three reasons why microorganisms are stained TO TO entrance wisualize from of the cell under a microscope b) To highlight dufferences of the more organisms C) To entrance or highlight cell components
2.	On what type of cells can staining be conducted
	Staming can be conducted on fixed or non-lawing cells
3.	Give two examples of stains. Simple Staining
4.	What is the result of staining with three dyes? It results in seperation.
5.	Why do we prefer stains with positively charged chromogen? H. (s. Preferred because bacterial muchas and cerain cell wall components carry a negative charge that strongly attracts and binds to the Cationic Chromagen
6.	Under what magnification do you examine all the stained slides?
7. cells	What stains are used to show the morphology and arrangement of bacterial
	***************************************

### LABORATORY EXERCISES SEVEN ECOLOGICAL TECHNIQUES

- 7.1 Ecological Terms
- i. Terrestrial organisms = organisms that live on land.
- ii. Aquatic organisms organisms that live in water. e.g fish, crayfish, octopus, etc
- iii. Arboreal organisms = organisms that live in the trees. e.g. Squirrel, monkey, pangolin, chameleon, etc
- iv. Planktonic organisms = organisms that float on water surface e.g copepods (and other microcrusteaceans), rotifer, algae, eggs and larvae of aquatic organisms in the water.
- Nektonic organisms = organisms that swim well in water, e.g fish, sea turtle, Blue crab, Octopus, etc
- vi. Benthos / Benthic organisms = organisms that dwell on sea or river bed (bottom-dwellers of a water body).
- Sessile or sedentary organisms live permanently attached to hard substrates (rocks, mangrove, ship-wreck, piers, in water). e.g. Oyster, sponge, barnacle, hydroid, tunicate, bryozoan, etc.
- viii. Semi-sessile organisms = Slow-moving animals like periwinkles, earthworms, garden snails, millipede, woodlice, etc that can easily be collected or captured.
- ix. Amphibious shoreline animals that spend some time in water and on land e.g. crocodile, River Otter, Nile monitor, sea turtle, terrapins, Hippopotamus, Seal, Sealion, etc.
- x. Fossorial = Burrowing animals e.g. Ocypod crabs, earthworm, crickets, giant rat (rabbit), etc.
- Cursorial ground-running animals like antelopes, duikers, bush-pig, grasscutter, etc.
- xii. Saltorial = leaping animals e.g kangaroo
- xiii. Authotrophs= Organisms that carry out the food-manufacturing process of photosynthesis, namely- green plants, algae, seagrass, etc.
- xiv. Herbivores = animals that feed on plants and plant parts.

pecies	EST. PRINCIPAL STREET	Count of An	imal Species		The second secon
-	Stone	Dead wood	Panel	Plastic	Total
Earthroom		0	6	0	10
Contipede		5 2	2		9
Cricket			8	0	12
Termiter		15	0	0	15
Ants	10	12	9	6	35
Spider	0	2	7	and all the	to
Milliferde	0	2	0	.0	2
Frog	0	0	5	6	11
Cockmach		2	3		13
OTAL	32	40	38	15	125
		mmonly found and B shide in these h be combe			ditions le mither
How do	Through.	these habitats?	60 n		T
Makea	list of other h	abitat types you	u can find on c	ampus, other	than these.
			nall are		

plastic container. Introduce some straw or dry leaves into it and let SHOTE ON POALS days. During the period, protozoan, rotifers, nematodes and algae etc in the culture will multiply.

Itel DUAL CAMERA

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## Laboratory Exercise

measurements? A Secchi disc is a metalic disc about soon across afternating black and white quadrant used to measure transparency of water body.
2 How are earthworms extracted from soil? Earthworm are extracted by pairing 5ml of 4% to formaldehyde into 5ml of water sprinkling over a square meter of glassfand, they will be driven out of their burrows.
3. What are the number of methods for extracting ectoparasites from their host?  There are two methods  Detergent washing methods.
4. What are the methods for extracting nematodes, small arthropods from soil and organic debris? Easymann fungel, lurgen funnel
5. Explain the methods for catching small mammals and birds. (b) Long nets with mammal trap is used for (atching small mammals like rate (B) Most net is used to trap birds.
6. What are the following used for in ecological studies?  i) Pooler Collecting minute and delicate organisms without ii) Pit trap collecting animals including Hweats, myriapods, Collecting animals including Hweats, myriapods, iii) Basket trap Capture fishes for studies
7. List the types of equipment used in collecting soil samples for analysis of microorganisms

8. What is the Pour Plate Method? A laboratory feahing Isolating and counting the viable mi 9. What are the conditions to ensure results that are accurate in the Pour Plate Method? Fixed amount of omensum from placed in the centre of stende, 15 inverted at 32°C for 24-48 hours. Preparing the Skeleton of a Small Mammal (e.g. a Giant Rat) 7.7.1 Oxidation Method (a) Procedure Sacrifice the animal humanely by putting it in a desiccator containing cotton wool soaked in chloroform or ether for 15 min. Dissect the animal to remove the visceral organs. Remove the skin. · Carefully chop off the major muscles of the body with scissors, making sure no bone is cut or cracked. Boil for 2 hrs. Put the animal in a transparent cellophane bag and fill it with manure. Open the earth and bury the cellophane and its content. Mark the spot with small pegs or caution tape Allow it to decay for 1 month. At the end of one month, exhume the animal and put it in a basin of water and wash off the soil. Immerse it in a detergent solution for 24 hrs to macerate any remaining tissue attached to the bones. Rinse Soak in 50% Ammonia solution Immerse it in container of hydrogen peroxide for 24 hrs. Rinse and air dry. clean, white, degreased set of skeleton, without any objectionable