#### FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY

FIRST SEMESTER EXAMINATION 2016/2017 SESSION COURSE CODE: MCB 211 COURSE TITLE: INTRODUCTION TO MICROBIOLOGY I (3 UNITS) CLASS: 200 LEVEL TIME: 1hr 30 min Instruction: Answer two (2) questions from section A and one question from section B.

#### Section A

- (a) Define microbiology and state the branches of microbiology (b) Enumerate the benefits of microorganisms
- 2. (a) Give the major achievement of each of the following microbiologists:
  - (i) Louis Pasteur (ii) Robert Hooke
  - (iii) Alexander Fleming (iv) Antonni Van Leeuwenhoek
  - (v) Robert Koch

(b)Give two named examples of each of the following:

- (i) Bacteria (ii) Fungi
- (iii) Parasites (iv) Algae
- (v) Viruses
- 3. Describe what you understand by the term microbial taxonomy and give the taxonomic rank of a named bacterium.

#### Section B

- 4. Discuss why you consider microbiology as an important course.
- 5. (a) List some of the ways we can study microorganisms.(b) Describe the general morphology of bacteria with named examples.

## FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY FIRST SEMESTER EXAMINATION, 2016/2017 SESSION COURSE CODE: MCB 311- GENERAL MICROBIOLOGY (3 UNITS)

**INSTRUCTION:** Answer ALL Questions in Section A and Any 3 Questions in Section B

TIME ALLOWED: 2 hours only

## SECTION A

(i) Write the colour taken up by the underlisted cells and their Gram stain reactions.

Bacterial species		Cell Colour	Gram Positive/Negative	
(a) <i>Es</i>	cherichia coli			
(b) Sta	aphylococcus epidermidis			
(c) Sta	aphylococcus aureus			
. ,	eudomonas aeruginosa			
. ,	cillus subtilis			
(ii)	Explain the fundamental di	fference between a simi	ble and a differential staining.	
(iii)	Describe the function of the	-		
(111)	(a) Primary stain			
	(b) Decolourizer			
	(c) Counterstain			
(iv)	What culture will you advis	se individuals to use in (	Gram staining?	
(v)	Briefly describe the mecha		C	
(vi)	Name the primary stain use	ed in acid fast staining p	rocedure	
(vii)	What is the aim of heating/	steaming during the acie	d fast staining?	
(viii)	In clinical microbiology lab	boratory, the acid fast st	aining would be used for diagn	
	of			
(ix)	What makes a microorgani			
(x)		1	*	
(xi)	-	•	t unstained, we refer to the stai	
	as			
(xii)	~	-	ect staining	
(xiii)	Commonly used simple sta			
	(a)			
	(b)			
	(c)		1 11	
(xiv)	Presence or absence of met	achromatic granules is o	demonstrated by	
()	staining	ant is used in consult at	aining	
(xv)	reag	-	-	
(xvi)	is us	ed as primary stain in er	ndospore staining	

## **SECTION B**

1a) Write short notes on the following: autoclave, mycorrhizae, eutrophication, chemotrophs, selective medium.

1b) With a relevant diagram, describe the life cycle of a named bacterium?

2a) Describe the lytic cycle of a t4 bacteriophage?

2b) If the magnification of the objective lens of a binocular microscope is 100x, what is its total magnification?

3a) In a tabular form, give 5 differences between Gram positive and Gram negative bacteria giving an example of each?

3b) What is the generation time of a bacterial population that increases from  $10^4$  cells to  $10^7$  cells after four hours of growth (Note: Log  $10^7/10^4$  = 3)

4a) Write 5 distinguishing features between a prokaryote and a eukaryote?

4b) Write a short note on the metabolism of t4 bacteriophage?

5a Discuss DNA Homology experiment?

5b) Explain the interrelated parts of microbial taxanomy?

5c) Explain the two quality of bacteria classification and list the general methods of classification?

#### FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY

FIRST SEMESTER EXAMINATION 2016/2017 SESSION COURSE CODE: MCB 312 COURSE TITLE: PRINCIPLE OF STERILIZATION & DISINFECTION (3 UNITS) CLASS: 300 LEVEL TIME: 2 HOURS INSTRUCTION: Answer Two Questions Only from Section A and answer All Questions in Section B by filling in the correct answers in the blank spaces provided.

## Section A

- 1. (a) What factors must a physician consider when determining the appropriateness of antibiotics for therapeutic use?
- (b) Discuss the difference between broad and narrow spectrum antibiotics
- 2. (a) What is MIC? Why is it a common test in clinical microbiology laboratory?(b) Why is it so difficult to find antimicrobial agents for treating viral diseases?
- 3. (a) What causes drug resistance in microorganisms?
  (b) Why is penicillin ineffective against bacteria that produce β-lactamase?
  Section B

## Matriculation Number: \_\_\_\_\_

- 1. In microbiology, contaminants are (1) \_\_\_\_\_\_ present at a given place and time that are (2) \_\_\_\_\_\_ or (3) \_\_\_\_\_\_.
- Most decontamination methods employ either (4) \_\_\_\_\_\_, (5) such as (6) \_\_\_\_\_\_, or (8) \_\_\_\_\_\_, or (8) \_\_\_\_\_\_ such as (9) \_\_\_\_\_\_, and (10) \_\_\_\_\_.

- 3. A sterile object is totally free of (11) \_\_\_\_\_, (12) \_\_\_\_\_, (12)
- 4. When sterilization is achieved by a chemical agent, the chemical is called a (14)
- 5. Sterilization is a process that (15) \_\_\_\_\_ or (16) \_\_\_\_\_ all (17) \_\_\_\_\_ all (17)
- 6. Sterilization is also the process in which all (19) \_\_\_\_\_\_, including (20) \_\_\_\_\_\_, are (21) \_\_\_\_\_.

 7. Disinfection is the (22) \_\_\_\_\_\_, (23) \_\_\_\_\_\_, or (24)

 \_\_\_\_\_\_\_ of (25) \_\_\_\_\_\_ that may (26)

8. Disinfectants are agents, usually (27) \_\_\_\_\_\_.

 Any process that destroys the non-spore forming contaminants on inanimate objects is called (28) \_\_\_\_\_\_.

10. The primary mode of action of non-ionizing radiation is to (29) \_\_\_\_\_\_.

## 11. The most versatile method of sterilizing heat-sensitive liquids is (30) \_\_\_\_\_\_. FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY FIRST SEMESTER EXAMINATION 2016/2017 ACADEMIC SESSION COURSE CODE: MCB 313 - MICROBIAL ECOLOGY (3 UNITS)

**Instruction**: Answer any **FOUR** questions **Time allowed: 2 Hours** 

- 1a) Write short notes on the following terms as they relate to microbial ecology (i) rhizosphere,
- (ii) endogenous diseases, (iii) plankton, (iv) phyllosphere, (v) ecological niche.
- 1b) Discuss 5 impacts of microorganisms on the environment and human activities
- 2a) Discuss 5 ecological relationships that can be found in the rhizosphere
- b) Give five (5) reasons why air is a non-hospitable environment for microbial growth.

3a) In a healthy animal, the internal tissues are normally free of microorganisms however, many other sites are readily colonized by various microbial species, most of which are beneficial to human health, discuss with relevant examples.

3b) List five factors influencing microorganisms in the rhizosphere.

4a) Describe 5 ways Pyrolobus fumarii adapt to their environment

4b. Give 5 reasons why the normal microbiota may be harmful to their host

5. Write short notes on mineralization, nitrification and denitrification in nitrogen cycle by microorganisms highlighting factors that affect each of them.

6. With relevant examples and distinguishing characteristics, discuss the following microbial interactions in soil environment

i. Cometabolism

ii. Protocooperation

#### FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY

#### FIRST SEMESTER EXAMINATION 2016/2017 SESSION COURSE CODE: MCB 411 COURSE TITLE: IMMUNOLOGY AND IMMUNOCHEMISTRY (3 UNITS) CLASS: 400 LEVEL TIME: 2 HOURS Instruction: Answer question number ONE and any other three questions

**Instruction**: Answer question number **ONE** and any other three questions

- 1. Describe in details with adequate precautionary measures, how widal test can be conducted in the laboratory?
- 2. Explain the biological functions of immunoglobulins
- 3. Write an essay on types II and III hypersensitivity
- 4. Discuss all the pathways of complement activation
- 5. Describe in details antigens and antibodies reactions

6. (a) Explain primary and secondary immune responses

#### (b) State the advantages and disadvantages of each FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY

FIRST SEMESTER EXAMINATION 2016/2017 SESSION COURSE CODE: MCB 412 COURSE TITLE: FOOD MICROBIOLOGY (3 UNITS) CLASS: 400 LEVEL TIME: 2 HOURS Instruction: Answer any 4 questions

1. Study the food specific attack rate table below, fill the blanks and answer the questions that follow.

Food	No of people who ate and were ill	No of people who ate and were not ill	Total no of people who ate	% of people who ate and were ill	No of people who did not eat and were ill	No of people who did not eat and were not ill	Total no of people who did not eat	% of people who did not eat and were ill
Kwose	60		85		58		71	
Moi-Moi		17	70			11	74	
Massa	64			80	48			72
Custard	51	10			17	68		

The symptom of food borne diseases occurred two hours after consumption of the food. It included dizziness, vomiting and diarrhoea

- (a) What type of food borne disease is this?
- (b) What is the incriminating food?
- (c) What is the likely organism?
- (d) List the serological type of the causative agent based on their morbidity
- (e) How can future occurrence be prevented?
- 2. (a) What properties make bacteria important in food?
  - (b) Certain factors favour production of botulism toxins. Enumerate them
  - (c) What is histamine poisoning?
- 3. (a) Discuss microbial growth in succession
  - (b) Distinguish between food poisoning and food infection
  - (c) Describe the sequence of events that take place for food to get spoiled.
- 4. How would the following factors affect microbial activities in food
  - (a) Water activity
  - (b) Biological structure of food
  - (c) Antimicrobial constituents of food
- 5. (a) What is single cell protein?(b) How does food fermentation provide food security and health benefit?
- 6. Write short notes on the following:

- (a) Molecular method of detecting microorganisms in food
- (b) Microbial enumeration by membrane filter technique
- (c) Biological method of food preservation

# FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY

#### FIRST SEMESTER EXAMINATION 2016/2017 SESSION

#### **COURSE CODE: MCB 413**

#### COURSE TITLE: INDUSTRIAL MICROBIOLOGY (3 UNITS)

#### CLASS: 400 LEVEL

#### **TIME: 2 HOURS**

**Instruction:** Answer Two Questions Only from Section A and answer All Questions in Section B by filling in the correct answers in the blank spaces provided

#### **SECTION A**

1. (a) Outline the procedures for the isolation of amylolytic bacteria from a natural environment.

(b) Discuss any two (2) methods of culture preservation in a dehydrated form.

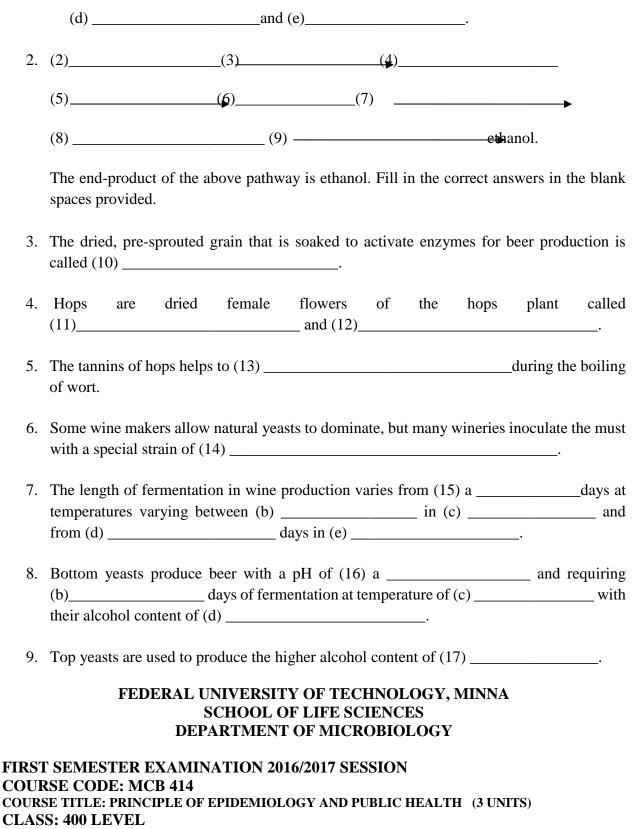
- 2. (a) What is strain improvement?
  - (b) How could an industrial microbiologist maximize its profit through strain improvement?
- 3. Write a comprehensive essay on how microorganisms could be used to enhance the economy growth of a nation.

#### **SECTION B**

#### Matriculation Number: \_\_\_\_\_

1. The major brewing materials used in the brewery industry include:

(1) a \_\_\_\_\_ (b) \_\_\_\_\_ (c) \_\_\_\_\_



TIME: 2 HOURS

**Instruction: Answer any 4 questions** 

- 1. (a) Differentiate between the following pairs as they relate to epidemiological study:
  - (i) Epidemic and endemic
  - (ii) Pathogenicity and infectivity
  - (iii) Incidence rate and prevalence rate
  - (iv) Herd immunity and risk factors

(b) How would you control an epidemic of a reported case of typhoid fever in your locality?

- 2. (a) Describe the various steps involved in epidemiological investigation.
  (b) In 2016, a study was carried out regarding an outbreak of Flu in Bosso with a population of 150, 800. Using 1000 as a constant, determine the incident rate if the number of new cases is given as 100 and if 560 are already diagnosed with Flu before 2016.
- 3. Explain in details the strategies you can use to effectively control the spread of communicable disease from one state to another in Nigeria.
- 4. Discuss Nosocomial infection
- 5. Write short note on antigenic shift and antigenic drift and explain their effect on the efficacy of vaccines against specific pathogens.
- 6. Explain the spectrum of disease

## FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA

## SCHOOL OF LIFE SCIENCES

## **DEPARTMENT OF MICROBIOLOGY**

## FIRST SEMESTER EXAMINATION 2016/2017 SESSION

## COURSE CODE: MCB 415

## COURSE TITLE: MICROBIAL GENETICS AND MOLECULAR BIOLOGY (3 UNITS)

## CLASS: 400 LEVEL

## TIME: 2 HOURS

**Instructions**: Answer **Four Questions** in All; **Two** from each Section. Question one (1) is compulsory from section A.

AUG CUG AAA CCC AUG.....UAC CGA ACG UAA ACG GGG CUG ACC UUU

In the hypothetical gene shown above:

- 1. (a)(i) Illuminate what will happen when there is a deletion of C from the codon CGA?
  - (ii). Produce the successive downstream codons.
  - (iii). What is the name of the mutation in ii?
  - (iv). State the two (2) new codons formed as a result of splitting the bases of the stop codon in the hypothetical gene above?

(v). State the new stop codon of the mutation in ii and make a distinction between the stop codon in the hypothetical gene above.



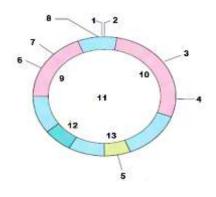


Figure 1

i. As a student of microbial genetics and molecular biology, provide a suitable title for the above Figures 1.

ii. Name 1-13 in Figure 1.

- iii. Divulge the universal significance of Figure 1 in microbial genetics and molecular biology. (To be answered in 15 words).
- 2. (a) When mutations are established, what will happen to the offspring of chloramphenicol sensitive cells that mutate to chloramphenicol resistance? On rare occasion, however, the nucleotide will reappear to its previous state, resulting in the chloramphenicol resistant cells becoming chloramphenicol sensitive. Explicate briefly the phenomenon.
  - (b) How many base pairs would have to be deleted in a mutational event to eliminate a single amino acid from a protein and not change the rest of the protein?

3. (a) Assume the following base sequence was found in a 20 base DNA strand:

## $3^1\,\text{ATT}$ CGA CCT TAT TAC TGC AC $5^1$

- i. What would be the 10 bases in the 3<sup>1</sup> end of the complementary strand?
  ii. What would be the 10 bases in the 5<sup>1</sup> end of the complementary strand?
- (b) How can you artificially stimulate a bacterium in the laboratory to take up DNA fragments from the environment? (To be answered in 15 words).

#### Section **B**

- 4. Explain the major roles of RNA in the metabolism of prokaryotes.
- 5. DNA is an amphipathic molecule. Discuss.
- 6. Discuss the synthesis of ribosomal RNA.

#### FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY

## FIRST SEMESTER EXAMINATION 2016/2017 SESSION COURSE CODE: MCB 511 COURSE TITLE: PATHOGENIC BACTERIOLOGY (3 UNITS) CLASS: 500 LEVEL TIME: 2 HOURS Instruction: Answer any four (4) questions

- 1. Discuss bacterial toxin. Compare and contrast the different types of toxin and give specific examples.
- 2. (i) Write short notes on bacterial virulence(ii) Explain three mechanisms by which bacteria demonstrate it.
- 3. (i) Describe bacterial morphology(ii) Explain the roles of at least three bacterial organelles
- 4. Give at least ten bacterial pathogens and highlight the mechanisms of pathogenesis of each of them.
- 5. Using standard laboratory procedures, explain how you will show that a given sample of bacterial culture contains *Staphylococcus aureus*.

6. (a) Describe the procedures for carrying out the following laboratory tests: lactose fermentation, methyl red, indole, gas formation, and sugar fermentation.

(b) How will you quickly differentiate the following bacteria in the laboratory? *Pseudomonas aeruginosa, Escherichia coli, Staphylococcus aureus, Salmonella typhi,* and *Streptococcus pneumoniae* 

#### FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY

#### FIRST SEMESTER EXAMINATION 2016/2017 SESSION COURSE CODE: MCB 512 COURSE TITLE: FERMENTATION TECHNOLOGY (3 UNITS) CLASS: 500 LEVEL TIME: 2 HOURS

#### Section A

#### Instruction: Answer only two (2) questions in this section. Question one is compulsory

1. (a) The maximum specific growth rate of an organism used in a chemostat for the production of microbial biomass is 0.40 h<sup>-1</sup> and the specific growth rate dictated by the dilution rate was 0.30 h<sup>-1</sup>, what was the substrate concentration at the steady state and the flow rate if the volume of the reaction vessel was 1.5 L? Assume the substrate utilization constant is 20mgL<sup>-1</sup>.

(b) A strain of *Lactobacillus bulgaricus* used in the production of yoghurt was known to divide every 30 minutes, if  $1 \times 10^3$  cells were used to inoculate 100 mL liquid milk, how many bacterial cells would be present after 6 h of fermentation? What is the specific growth rate and the generation time of the organism?

(c) Outline the sequence of events that occur when the substrate is reduced below the level that support the growth rate controlled by the dilution rate in a continuous culture system of fermentation.

2. (a) Describe the downstream processing for the purification of an intracellular microbial enzyme.

(b) Enumerate five (5) major groups of industrial fermentation processes with their relevant examples.

- 3. (a) Explain briefly how industrial fermentation could contribute to the economy growth of a nation.
  - (b) Distinguish between fed-batch and continuous system of fermentation.
  - (c) Why is fed-batch system more preferred for the cultivation of baker's yeast?

#### Section **B**

#### Instruction: Attempt any two questions in this section

- 4. (a) Describe food fermentation with appropriate examples(b) List the two types of fermentation and explain the principles behind any of the listed types with production technique
- 5. (a) Explain the advantages and disadvantages of batch fermentation process. What is the role played by head space?

(b) Give two examples of the following fermented foods classified by substrate type:

(ii) Fruit based

- (i) Bean based
- (iii) Dairy based (iv) Grain based
- (v) Vegetable based

## 6. (a) Enumerate the basic functions of a fermenter

- (b) Write short notes on the following:
- (i) Baffles

- (ii) Sparger
- (iii) Chemostat (iv) Foam control

#### FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY

FIRST SEMESTER EXAMINATION 2016/2017 SESSION COURSE CODE: MCB 513 COURSE TITLE: GENERAL TOXICOLOGY (3 UNITS) CLASS: 500 LEVEL TIME: 2 HOURS Instruction: Answer four questions, at least two from each section

#### SECTION A

- 1. Write short notes on the following: Pyrrophyceae, Cyanophyceae and Chrysophyceae, carcinogens and lethal dose
- 2. Describe the following: Aflatoxins, Sterigmatocystin, Trichothecenes, *Corynebacterium diphteriae*, and *Clostridium*
- 3. a. Explain the mechanism of drug resistanceb. The techniques of removal of toxic substances from patients

#### **SECTION B**

- 4. a. Highlight the factors that can predispose an individual to drug toxicityb. write short notes on any two mechanisms of drug toxicity
- 5. With relevant examples discuss three natural toxicants and their effects on human body
- 6. a. Write short notes on different techniques that can be employed to manage acute drug toxicity

b. What are the factors that modulate effects of toxicant in human body? FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY

#### FIRST SEMESTER EXAMINATION 2016/2017 SESSION COURSE CODE: MCB 512 COURSE TITLE: FERMENTATION TECHNOLOGY (3 UNITS) CLASS: 500 LEVEL TIME: 2 HOURS

#### Section A

#### Instruction: Answer only two (2) questions in this section. Question one is compulsory

1. (a) The maximum specific growth rate of an organism used in a chemostat for the production of microbial biomass is 0.40 h<sup>-1</sup> and the specific growth rate dictated by the dilution rate was 0.30 h<sup>-1</sup>, what was the substrate concentration at the steady state and the flow rate if the volume of the reaction vessel was 1.5 L? Assume the substrate utilization constant is 20mgL<sup>-1</sup>.

(b) A strain of *Lactobacillus bulgaricus* used in the production of yoghurt was known to divide every 30 minutes, if  $1 \times 10^3$  cells were used to inoculate 100 mL liquid milk, how many bacterial cells would be present after 6 h of fermentation? What is the specific growth rate and the generation time of the organism?

(c) Outline the sequence of events that occur when the substrate is reduced below the level that support the growth rate controlled by the dilution rate in a continuous culture system of fermentation.

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- 3. (a) Explain briefly how industrial fermentation could contribute to the economy growth of a nation.
  - (b) Distinguish between fed-batch and continuous system of fermentation.
  - (c) Why is fed-batch system more preferred for the cultivation of baker's yeast?

## Section B

#### Instruction: Attempt any two questions in this section

- 4. (a) Describe food fermentation with appropriate examples
  - (b) List the two types of fermentation and explain the principles behind any of the listed types with production technique
- 5. (a) Explain the advantages and disadvantages of batch fermentation process. What is the role played by head space?

(b) Give two examples of the following fermented foods classified by substrate type:

- (i) Bean based (ii) Fruit based
- (iii) Dairy based (iv) Grain based
- (v) Vegetable based
- 6. (a) Enumerate the basic functions of a fermenter
  - (b) Write short notes on the following:
  - (i) Baffles (ii) Sparger
  - (iii) Chemostat (iv) Foam control

## FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA

## SCHOOL OF LIFE SCIENCES

## DEPARTMENT OF MICROBIOLOGY

#### FIRST SEMESTER EXAMINATION 2016/2017 SESSION

#### COURSE CODE: MCB 516

#### COURSE TITLE: INTRODUCTION TO BIOTECHNOLOGY (3 UNITS)

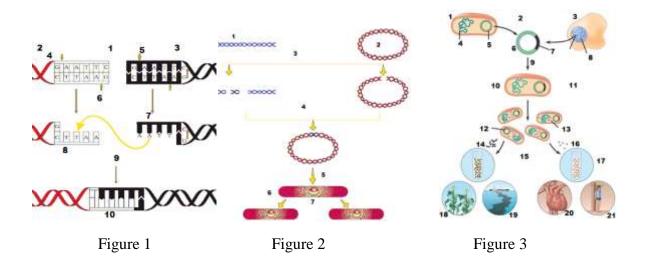
#### CLASS: 500 LEVEL

#### **TIME: 2 HOURS**

**INSTRUCTIONS:** Answer **Four Questions** in All; **Two** from each Section. Question one (1) is compulsory from section A.

**Time Allowed:** 2 Hours

Section A



- 1. (a). As a student of biotechnology, provide a suitable title for the above Figures 1-3 in biotechnology and genetic engineering.
  - (b). Name 1-10, 1-7 and 1-21 in Figures 1-3 respectively.
  - (c). Mention the salient feature universal to the three (3) figures and its significance in biotechnology and genetic engineering.
- 2. A case of defective coagulation factor gene was reported to the school clinic from a student. Observable signs and symptoms were detected by the doctor.
  - (a) As a student of biotechnology, kindly explicate to the doctor on how to treat or cure the disease condition using genetic engineering and biotechnology techniques.
  - (b) What is the name of this genetic defect?

3. Write a compendium of historical advancement and important milestones of biotechnology with regards to time period and major break-through. (To be answered in a tabular form).

#### Section B

- 4. (a) What is Enzyme Immobilization?
  - (b) List the three major groups of supports for Enzyme Immobilization and three examples for each support.
  - (c) List five methods of Enzyme Immobilization.
- 5. (a) Give three examples and three characteristics of secondary metabolites
  - (b) Discuss any three of the following terms: (i) Bioconversions (ii) Induction (iii) Phosphate regulation (iv) End product regulation (v) Catabolite regulation.
- 6. (a) What are the necessary steps needed for the purification of a product?
  - (b) List three types of each of the following product purification operation: (i) Chromatography (ii) Centrifuges.

#### FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA SCHOOL OF LIFE SCIENCES DEPARTMENT OF MICROBIOLOGY

#### FIRST SEMESTER EXAMINATION 2016/2017 SESSION COURSE CODE: MCB 517 COURSE TITLE: ENTREPRENEURSHIP (2 UNITS) CLASS: 500 LEVEL TIME: 1hr 30 min Instruction: Answer only three (3) questions at least one question from each section

#### Section A

- (a) What factors motivate entrepreneurship?
   (b) Enumerate six (6) sources of entrepreneurship ideas
- 2. Taking into cognizance the raw materials, discuss the production of the following home made products
  - (i) Ginger ale drink
  - (ii) Bread loaf
- 3. Ability to make good decisions is essential to success in business. Discuss

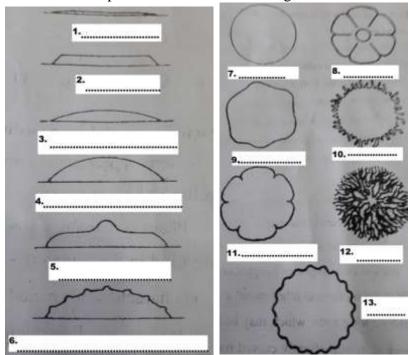
## Section B

4. (a) What is entrepreneurial microbiology?(b) Discuss the prospects of entrepreneurial microbiology in a developing country.

- 5. How would you use effective microorganisms to boost food and animal production in a country faced with economic recession?
- 6. Biogass technology is more sustainable in Nigeria than Cotonou. Discuss

FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA DEPARTMENT OF MICROBIOLOGY SECOND SEMESTER EXAMINATION 2016/2017 SESSION INTRODUCTION TO MICROBIOLOGY II (MCB 221) 3 UNITS INSTRUCTIONS: Answer All Questions. Time Allowed: 30 Minutes Matriculation Number:

#### Section A



Name 1-13 and provide a suitable title to Figure 1 and 2.

Figure 1 (14)..... Figure 2 (15).....

Fill in the correct answers in the blank spaces provided (16-20)

Some varieties of culture media use in Microbiology Laboratory include (16) \_\_\_\_\_ (17) \_\_\_\_\_ (18) \_\_\_\_\_

FEDERAL UNIVERSITY OF TECHNOLOGY DEPARTMENT OF MICROBIOLOGY INTRODUCTION TO MICROBIOLOGY 2 (MCB 221) SECOND SEMESTER EXAMINATIONS 2016/2017 ACADEMIC SESSION SECTION B Name; ...... Mat No.; ..... INSTRUCTIONS; Answer all questions Time Allowed: 45 Minutes Column A Column B 1. Dye repelled by bacteria (a) Gram stain technique 2. Iodine used (b) Acid-fast technique 3. Developed in 1880s (c) Simple stain technique 4. Carbon fuchsin used 5. Uses alcohol decolorizer 6. Microorganisms outlined in stain 7. Basic dye used 8. Acid – alcohol decolorizer 9. All bacteria divided to two groups 10. Mycobacterium detected Completion; Add the word or words that best complete each of the following statements 11. The unit most often used to measure the diameter of viruses is the \_\_\_\_\_ 12. Whereas a simple microscope has a single lens, a compound microscope has 13. The ability of a microscope to distinguish two closely related points is called 14. A meter is equivalent to micrometers that number\_\_\_\_\_ 15. The part of the microscope known as the condenser consist of a series of 16. Among the microorganisms most often studied with the phase contrast microscope are the 17. Two types of electron microscopes are the transfusion electron microscope and the 18. Bacteria and microorganisms must be stained before microscopy because their cytoplasm is usually 19. Those stains that carry a positive charge are referred to as \_\_\_\_\_

20. Among the smallest bacteria are three groups known as rickettsiiae, chlamydiae and

(19) \_\_\_\_\_\_and (20) \_\_\_\_\_\_

Match the choices from column B with the appropriate statements in column A

- (d) Negative stain technique

#### FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA DEPARTMENT OF MICROBIOLOGY SECOND SEMESTER EXAMINATION, 2016/2017 SESSION INTRODUCTION TO MICROBIOLOGY II (MCB221)

#### SECTION C

INSTRUCTION: Answer all Questions MATRIC. NO.:

#### Time Allowed: 1Hr

Define the following: 1.Ecology:				
2. Niche: 3. Habitat: 4. Limiting factors				
5. Extremophiles species:				
What is the role of the following organisms in 6.Producer:	an ecosystem:			
7.herbivore:	8.carnivore:			
9.omnivore:				
11.How many trophic levels has this food chain:		<del>Grass →</del>	Goat ——	— <b>M</b> an
<ul><li>List three elements of aseptic techniques:</li><li>(13)</li></ul>	(12) (14)			
<ul><li>15. When several food chains overlap and intersec</li><li>&gt; Briefly describe the following relationships</li></ul>	t is called			
16. predator-prey:				
17. parasite-host:				
18. mutualism:				
19. commensalism:				

20. cooperation: -----

## FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA DEPARTMENT OF MICROBIOLOGY COURSE TITLE: MEDICAL MYCOLOGY COURSE CODE: MCB 322 (3 units) SECOND SEMESTER EXAMINATION, 2016/2017 ACADEMIC SESSION

**Instruction:** Answer all questions in this Section **Time allowed:** 1HR

#### Section A

- 1. Describe the various clinical manifestitations of candidiasis
- 2. Name the dimorphic fungi and describe the morphology and pathology of any one of them
- 3a. What are the unique features of dermatophytes?
- b. Define the following terms and give three examples for each:
- (i) Anthropophilic dermatophyte (ii) Zoophilic dermatophyte
- (iii) Geophilic dermatophyte
- 4 Outline the detail procedure in carrying out diagnosis of dermatomycosis in a clinical laboratory
- 5 Name the various systemic mycoses and discuss any one of them

## FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA DEPARTMENT OF MICROBIOLOGY MYCOLOGY (MCB 322) SECOND SEMESTER EXAMINATION, 2016/2017 SESSION SECTION B

Mat No.;Time: 1HRINSTRUCTIONS; Answer all questionsTime: 1HRMatching; select from Column B the class of fungi that fits the description in Column AColumn AColumn AColumn B1. Form ascospores(a) Chytridiomycetes2. Common mushroom(b) Zygomycetes3. "Imperfecti fungi"(c) Ascomycetes

- 4. Zygospore forms
- 5. Form oospore
- 6. Include the fermentation yeast
- 7. No known sexual cycle
- 8. Spore-beaning sexual structures
- 9. <u>Candida albicans</u> a member
- 10. Species causes ergot disease
- 11. Non septate hypae with sporangiospores
- 12. Rhizoids for anchorage
- 13. Include Rhizopus
- 14. Include rust & smut fungi

Completion; Add the word or words that best complete each of the following statements

- 15. Those fungi which are biphasic occur as yeast or as \_\_\_\_\_
- 16. Fungi tolerates environments that are high in sugar and high in \_\_\_\_\_
- 17. Yeasts are notable for their ability to live in the presence or absence of \_\_\_\_\_\_
- 18. Where the bacterial spore is used for resistance, fungal spore is used for \_\_\_\_\_
- 19. Asexual spore formed within a sporangium are known as \_\_\_\_\_
- 20. Fungi display a mode of nutrition that is \_\_\_\_\_
- 21. The cell wall of fungi contains cellulose, chitin and polymers of glucose known as
- 22. Those hyphae that form the reproductive structures of the fungus are called
- 23. The typical shape observed in a common yeast cell is called \_\_\_\_\_\_
- 24. The metabolic process performed by yeast and used in the beer and wine industries is called
- 25. The sexual reproductive process of a fungus, the terms "male" and "female" are used interchangeably with the term \_\_\_\_\_
- 26. Fungi are separated into two major divisions, with the true fungi placed into the division
- 27. Budding is the type of reproduction used by \_\_\_\_\_
- 28. The fungus used to ferment soybeans to soy sauce, to modify steroids and in the production of organic acids is \_\_\_\_\_\_
- 29. The mushroom agaricus is safe to eat, but the poisonous mushroom belongs to the genus
- 30. All of the following are considered fungi except
- 31. Mushrooms b. Yeasts c. Molds d. Amoebas
- 32. Those fungi which are biphasic occur as yeast or as \_\_\_\_\_
- 33. Fungi tolerates environments that are high in sugar and high in \_\_\_\_\_
- 34. Yeasts are notable for their ability to live in the presence or absence of \_\_\_\_\_\_
- 35. Where the bacterial spore is used for resistance, fungal spore is used for \_\_\_\_\_
- 36. Asexual spore formed within a sporangium are known as \_\_\_\_\_
- 37. Fungi display a mode of nutrition that is \_\_\_\_\_
- 38. Yeasts are notable for their ability to live in the presence or absence of \_\_\_\_\_\_
- 39. The important feature in the reproduction of fungi is the

(d) Basidiomycetes

\_\_\_\_\_

(e) Deuteromycetes

a. Septa b. Spore c. Structure of the cell wall d. Type of Nutrition 40. Both blastospores and arthrospores are

- a. Types of sexual spores c. Produced by yeasts d. similar to bacteria
- b. Various kinds of asexual spores

# FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA MICROBIOLOGY DEPARTMENT

#### SECOND SEMESTER EXAMINATION, 2016/2017 SESSION

#### COURSE: Microbial Physiology and Metabolism (MCB323, 3Units)

#### Instructions: Answer FIVE Questions in All: AT LEAST ONE from each Section.

**Time Allowed**:  $2^{1}/_{2}$  Hours

#### **SECTION A**

- 1. (a) What is bacterial growth?
  - (b) How can cells multiply when they have exhausted their supply of nutrients?
- 2. (a) What happens when a stock culture stored in the refrigerator for several weeks is inoculated into fresh medium? Provide a suitable reason for your answer.
  - (b) Briefly explain what happens if young cells are transferred to a medium similar in composition?

#### **SECTION B**

- 3. (a) (i)What are the major pathways for energy generation in microorganisms?
  - (ii) Enumerate the steps in protein synthesis.
  - (b) Describe the two sub processes involved in catabolism.
  - (c) Distinguish between oxygenic and anoxygenic photosynthesis.
- 4. (a) (i)What are the major components of electron transport chain?
  - (ii) Describe purine synthesis in bacteria.
  - (b) Explain the term "anaplerotic pathway."
  - (c) Discuss the functions of pentose phosphate pathway in heterotrophs.

- 5. (a) (i).Describe the type 2 reaction center in photosynthetic microorganisms.
  - (ii)What are non-sense codes? List them.
  - (b) What is the primary purpose of glyoxylate cycle in metabolism.
  - (c) With examples distinguish between homofermenters and heterofermenters.

#### **SECTION C**

6. (a) What is water activity?

(b) How do hypotonic and hypertonic solutions affect microbial activities? Explain how microorganisms adapt to these environmental conditions?

- (c) Describe microaerophiles and facultative anaerobes.
- 7. (a) How does a higher temperature above the optimum growth temperature of an organism affect its activities?
  - (b) Describe how Coulter counter method is used to measure microbial growth.
  - (c) Outline how anaerobic microorganisms can be cultured in the laboratory.

#### FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA

#### DEPARTMENT OF MICROBIOLOGY

#### COURSE TITLE: MICROBIOLOGICAL TECHNIQUES

#### COURSE CODE: MCB 324 (3 units)

#### SECOND SEMESTER EXAMINATION, 2016/2017 ACADEMIC SESSION

# Instruction: Answer 5 questions in all (attempt at least 2 question in each section) Time : $2^{1}/_{2}$ Hours

#### Section A

- 1. a). i, Enumerate the characteristics often used as additional aid to identification in bacteriology
  - ii, With named examples, distinguish between selective and differential medium
  - b). Enumerate the precautions to be taken in the course of pour plating in pour plate technique
- a). Calculate the CFU/ml of fruit juice samples submitted to the laboratory for analysis if 207 and 209 colonies were counted in a duplicate plating sample after dispensing 0.1ml of 10<sup>-5</sup> dilution.
  - b). i, Distinguish between synthetic and non-synthetic medium
  - ii, How are cultures preserved using soil?

3. a).What are the components of a good "Materials and Methods" in good scientific writing?

b). i, Describe the procedure for nutrient agar preparation if 200ml quantities is required. Nutrient gar is usually weighed 28g/L

ii, Describe the Gram staining procedure

4. a).Describe the operation of an autoclave

b). Discuss the Bacitracin and Bile solubility tests in biochemical identification of bacteria.

#### Section B

- 5. (a) Describe catalase test and explain how this test is used to distinguish bacteria(b) Discuss the morphology of bacteria
- 6. (a) Explain classification and identification of bacteria(b) What are vitamins? Describe the B vitamins giving their specific names
- 7. (a) List 6 laboratory tests used in identification of bacteria(b) Describe serology and briefly explain its uses

## FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA DEPARTMENT OF MICROBIOLOGY COURSE TITLE: VIROLOGY COURSE CODE: MCB 521 (3 units) SECOND SEMESTER EXAMINATION, 2016/2017 ACADEMIC SESSION

**Instruction:** Answer any five (5) questions **Time allowed:**  $2^{1}/_{2}$  **Hours** 

- 1a. Explain in details the lytic phase of viral replication
- b. Enumerate the properties of a viral particle

- 2. Given a blood sample that is suspected to contain Hepatitis C virus particle, describe details of analysis you will carry out on the blood sample to confirm the presence of the virus in the given sample.
- 3. Write an essay on Monkey pox virus
- 4. Virilogists have proposed that lysogenic phase of viral replication should be considered as an alternative to cloning. Discuss the pros and cons of this proposal?
- 5. In a tabular form clearly differentiate between the underlisted Hepatitis viruses
  - (i) Hepatitis A
  - (II) Hepatitis B
  - (III) Hepatitis C
- 6. Describe in details how Rubella can be diagnosed and prevented
- 7. Briefly explain the important medical considerations in viral diseases

## FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA DEPARTMENT OF MICROBIOLOGY COURSE TITLE: PETROLEUM MICROBIOLOGY COURSE CODE: MCB 522 (2 units) SECOND SEMESTER EXAMINATION, 2016/2017 ACADEMIC SESSION

## Instruction: Answer any 3 questions Time allowed: $1^{1}/_{2}$ Hours

1a) Write a short note on the following terms: (i) Biocompetitive exclusion(ii) Biodeterioration (iii) Biofilm (iv) Biogas composition (v) Dissimilatory sulphate reduction

1b) Outline the mechanism of biodegradation of a named hydrocarbon.

1c) Differentiate between detention time and retention time as it relates to anaerobic digestion.

2a) Describe the microbiology of methanogenesis.2b) Outline the mechanism of corrosion by methanogenesis

2c) List five impacts of sulphate reducing bacteria in the environment

3a) Define Microbial Enhanced Oil Recovery (MEOR)

3b) Discuss the different mechanisms of Microbial Enhanced Oil Recovery

4a) Enumerate five characteristics of microorganisms that enable them to degrade petroleum products

4b) Discuss the chemical composition of petroleum

5) In the petroleum industry, surface-active agents from microorganisms are preferred to synthetic surfactants. Discuss

## FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA DEPARTMENT OF MICROBIOLOGY COURSE TITLE: ENVIRONMENTAL MICROBIOLOGY COURSE CODE: MCB 523 (3 units) SECOND SEMESTER EXAMINATION, 2016/2017 ACADEMIC SESSION

## Instruction: Answer any 5 questions

## Time allowed: $2^1/_2$ Hours

1a) Write short notes on the following: (i) Plankton, (ii) eutrophication, (iii) biosorption, (iv) bioaccumulation, (v) bioconcentration.

1b) Describe two methods by which elements are made readily available by microorganisms?

2a) Differentiate between bioamplification and bioactivation?

2b) List 5 factors influencing the bioaccumulation and biodegradation of pesticides in the environment?

3a) Write briefly on 5 impacts of pesticides contamination on the environment?

3b) Outline 5 ways you would minimise the negative effects of pesticides in the environment?

4a) Discuss the different categories of contaminants present in wastewater and how it can be treated biologically.

4b) Highlight the objectives of wastewater treatment.

5a) What are the dangers associated with the presence of microorganisms in the air?5b) Discuss three mechanisms used by microorganisms in the air to adapt to unfavourable environment.

6a) Enumerate the differences between exotoxins and endotoxins

6b) Write short notes on the following: (i) Ergot, (ii) Aflatoxins, (iii) *Fusarium* toxins, (iv) Bacillosis, and (v) *Cyanobacteria diphtheriae* 

7a) Write short notes on any three of the following

- i) Biopesticides
- ii) Bioleaching
- iii) Organophosphate
- iv) Alkanes in petroleum

b) Explain the detrimental effects of Sulphur Cycle

#### FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA DEPARTMENT OF MICROBIOLOGY COURSE TITLE: PHARMACEUTICAL MICROBIOLOGY COURSE CODE: MCB 525 (3 units) SECOND SEMESTER EXAMINATION, 2016/2017 ACADEMIC SESSION

#### **Time Allowed : 2<sup>1</sup>/<sub>2</sub> Hours**

Instruction: Answer 5 questions in all (attempt at least 2 questions in each section)

#### Section A

1. (a) What is pharmaceutical microbiology?

(b) List 10 secondary metabolites of plants origin that are useful in pharmaceutical microbiology (c) State the traditional Welsh and American rhymes and explain their relevance in medicine

- 2. (a) Explain the following terms as they are used in pharmaceutical microbiology
  - i. Chemotherapy
  - ii. Prophylaxis
  - iii. Drug
  - iv. Antibiotic
  - v. Vaccine
  - vi. Antimicrobial agent
  - vii. Synthetic, naturally occurring and semi synthetic agents
  - (b) Give a diagramatical representation of the core structure of cephalosporin
- 3. (a) List the pharmaceutical ingredients that make pharmaceutical products susceptible to microbial attack
  - (b) What are the observable effects of microbial attack on pharmaceutical products?
  - (c) Enumerate the parameters to be controlled in order to mitigate spoilage of drugs and other pharmaceuticals.
- 4. (a) Discuss the role of NAFDAC in the quality control of pharmaceuticals in Nigeria
  - (b) Explain control of the phased production and control of final products of pharmaceuticals
  - (c) Describe the following:
  - i. Validation activities and monitoring
  - ii. Analytical development of new products and stability

#### Section B

- 5. (a) Giving specific examples, describe the possible effects of mutational resistance in microorganisms.
  (b) Explain how a microbiologist may contribute to incidence of microbial resistance
- 6. Explain how you would determine the susceptibility of an organism to a plant extract by poison food techniques
- 7. Discuss the general mode of action of antiseptics and disinfectants.

## FEDERAL UNIVERSITY OF TECHNOLOGY, MINNA DEPARTMENT OF MICROBIOLOGY COURSE TITLE: MEDICAL PARASITOLOGY COURSE CODE: MCB 526 (3 units) SECOND SEMESTER EXAMINATION, 2016/2017 ACADEMIC SESSION

**Instruction:** Answer five (5) questions with at least two questions from each section **Time allowed:**  $2^{1}/_{2}$  **Hours** 

## SECTION A

1a) Identify and discuss at least three factors that promote the spread of helminthic diseases in Nigeria1b) for each factor in (1a) above suggest public health measures for overcoming them

2. Discuss the importance of each of the underlisted concepts to the study of parasites(a) Morphology (b) Geographical distribution (c) Diagnosis (d) Life cycle(e) Host-Parasite interaction

3) Identify a parasitic disease for which haematuria is a key symptom and discuss it under the following headings

(a) Transmission (b) Pathology (c) Prevention (d) Control

## SECTION B

4) In a tabular form under the following headings compare Bancroft filariasis and River blindness:

(i) Parasites(ii) Symptoms(iii) Transmission(iv) Prevention(v) Treatment

- 5) Discuss malaria and its transmission in an Urban Slum
- 6) Describe the various classical methods used in the diagnosis of parasitic infections
- 7a) Name two examples each of the important groups of parasitic protozoa
- b) In what ways is the environment important in the transmission of parasitic disease?