



BRAINWARE UNIVERSITY

Term End Examination 2024-2025 Programme - M.Sc.(MB)-2023 Course Name - Microbial Biotechnology Course Code - MMBE304 (Semester III)

Full Marks: 60

Time: 2:30 Hours

[The figure in the margin indicates full marks. Candidates are required to give their answers in their own words as far as practicable.]

Group-A

(Multiple Choice Type Question)

1 x 15=15

- 1. Choose the correct alternative from the following:
- (i) Which microorganism is commonly used in the fermentative production of beer?
 - a) Lactobacillus

b) Escherichia coli

c) Saccharomyces cerevisiae

- d) Streptococcus
- (ii) What is the primary function of malt adjuncts in beer production?
 - a) To enhance color and aroma
- b) To filter the beer
- c) To provide additional fermentable sugars
- d) To increase alcohol content
- (iii) In beer production, what is the purpose of wort boiling?
 - a) Enhance color

- b) Decrease alcohol content
- c) Sterilize and extract flavors from hops
- d) Increase sweetness
- (iv) Select the reason why Aspergillus oryzae is preferred over Bacillus subtilis in the industrial production of amylase.
 - a) Aspergillus oryzae produces higher yields
 - c) Aspergillus oryzae has more stable enzyme

 - activity at extreme pH
- b) Bacillus subtilis has slower growth rates
- d) Bacillus subtilis requires a more complex growth medium
- (v) Select the importance of inoculum preparation in microbial enzyme production.
 - a) It minimizes contamination risk during recovery
 - c) It regulates the pH of the fermentation
- b) It optimizes microbial growth for efficient enzyme production
- d) It reduces the cost of enzyme purification
- (vi) Select from the following that is NOT a biotechnological product.
 - a) A. Antibiotics

b) B. Smartphones

c) C. Biofuels

- d) D. Vaccines
- (vii) Select type of microorganism that is commonly used in the production of enzymes
 - a) A. Mammals

b) B. Bacteria

c) C. Birds

- d) D. Fungi
- (viii) Select the term "bioproducts" refer to in the context of biotechnology

| | STYLEST. | |
|--|---|---|
| C Constitution diffed examines | d) D. Padioactive materials | |
| c) C. Genetically modified organisms d) D. Radioactive materials (ix) Differentiate the role of carbon and nitrogen sources in the formulation of the | | |
| production medium for microbial enzymes. | ources in the formaldion of the | |
| a) A. Nitrogen sources improve enzyme yield, | b) B. Carbon sources regulate pH, nitro | gen |
| while carbon sources provide energy | sources enhance product stability | , |
| c) C. Both carbon and nitrogen sources | d) D. Carbon sources determine enzym | e |
| enhance microbial growth equally | specificity, nitrogen sources impact y | rield |
| (x) Compare different recovery methods for enzymes after microbial fermentation. | | |
| a) A. Centrifugation removes microbial cells | b) B. Crystallization offers higher enzym | ne |
| while filtration retains enzymes | purity than liquid-liquid extraction | |
| c) C. Filtration is slower but offers higher | d) D. Chromatography yields higher en | zyme |
| recovery than centrifugation | purity but is costlier than filtration | |
| (xi) Identify the first step in the wine making proce | ess after the fruit has been selected | |
| a) Crushing | b) Fermentation | |
| c) Bottling | d) Aging | |
| (xii) Vitamin B12 production primarily involves imp | fortant metal, select them from the | |
| following. | hard and the subsequence and man | 11、11、11、11、11、11、11、11、11、11、11、11、11、 |
| a) Iron | b) Zinc | |
| c) Cobalt(xiii) Complex media containing sugars and nitroger | d) Magnesium | |
| microorganism for vitamin B12 production, ide | | |
| | b) Clostridium botulinum | |
| a) Saccharomyces cerevisiae c) Propionibacterium freudenreichii | d) Fusarium oxysporum | |
| (xiv) Choose the method commonly used for the re | | |
| fermentation broth. | | |
| a) Centrifugation | b) Precipitation and extraction | |
| c) Chromatography | d) Filtration | |
| (xv) Choose the main application of tetracyclines in medicine. | | |
| a) Treatment of fungal infections | b) Treatment of viral infections | |
| c) Antibiotic for bacterial infections | d) Pain management | |
| marries todayle samuel Gre | NUMB RESIDENCE | |
| Group-B (Short Answer Type Questions) 3 > | | 3 x 5=15 |
| off planting and a second | Type Questions, | HOLE TO |
| 2 Differentiate between chlortetracycline oxy-tetr | acycline, and tetracycline. | (3) |
| 2. Differentiate between chlortetracycline, oxy-tetracycline, and tetracycline.3. Discuss the role of microalgae in the production of biofuels. | | (3) |
| 4. Identify the microorganisms involved in the fermentative production of tetracyclines. | | (3) |
| 5. Assess the benefits of using microbial enzymes in the detergent industry compared to | | (3) |
| synthetic chemical alternatives. | ince reduid internation to the residual and | 4=1 |
| 6. Explain the importance of regulatory compliance in microbial biotechnology research. | | (3) |
| | OR | (3) |
| Compare the inoculum preparation techniques uproteases. | used for the production of anylases and | (3) |
| Gro | oup-C | |
| | Type Questions) | 5 x 6=30 |
| all a see production of electronics | of maximum at your Alice (great state or with | |
| 7. Evaluate the impact of microbial biotechnology on sustainable agriculture. Provide (5) | | |
| examples of microbial products used in agriculture and their benefits. | | |

- 8. Discuss the applications of extremophiles in biotechnology. Provide examples of (5) extremophiles and their unique adaptations for industrial processes. 9. Analyze the impact of biotechnology on the production of biofuels. Provide examples of (5) microbial processes used to produce biofuels and their advantages over conventional fuel sources. 10. Discuss the fundamental principles of fermentation. (5) 11. Discuss the significance of recovery processes in the commercial production of benzyl (5) penicillin. 12. Design a microbial vaccine production strategy using recombinant technology, detailing the (5)
- choice of microorganism, antigen expression, and recovery.

Formulate a step-by-step process for insulin production using genetically modified bacteria, (5) highlighting the role of recombinant DNA technology.

LIBRARY Brainware University Lerasat, Kokata -700125