



Department of Genetic Engineering and Biotechnology
Faculty of Health and Life Sciences
B. Sc. (Hons.) in Genetic Engineering and Biotechnology
Final Examination Spring 2026

Course Code: 0512-1203

Level and Term: L1, T2

Time: 2 Hour

Section: 253 (A+B)

Course Title: Basic Microbiology

Course Teacher Initials: DNH

Total Marks: 40

			Marks
1	(a) Compare batch and continuous culture systems of microorganisms in terms of growth phases, nutrient dynamics, and product formation.	[CLO1, PLO2, C1]	3
	(b) How do complex, selective, and enrichment media contribute to the isolation and growth of microorganisms? Explain with examples.	[CLO1, PLO2, C3]	3
	(c) Explain the methods used for microbial culture preservation.		2
2	(a) Describe the essential nutrients required by microorganisms for growth.	[CLO1, PLO1, C5]	3
	(b) Illustrate the growth curve of batch culture and explain how nutrient uptake and waste accumulation influence each growth phase.	[CLO2, PLO1, C4]	3
	(c) Explain the plate count method for direct measurement of microorganisms.		2
3	(a) State the principle of sterilization. Discuss the key factors that affect the efficacy of sterilization methods.	[CLO2, PLO1, C2]	3
	(b) Describe the chemical methods of sterilization.	[CLO1, PLO2, C1]	3
	(c) How do antimicrobial agents act on microorganisms?		2
4	(a) Diagrammatically present the pour plate and spread plate methods of microbial culture.	[CLO1, PLO1, C1]	3
	(b) Explain the serological and molecular techniques of microbial identification.	[CLO1, PLO2, C6]	3
	(c) Write short note on IMViC.		2
5	(a) Define microbial ecology. State and explain the biotic and abiotic factors influencing a microbial ecosystem	[CLO2, PLO2, C2]	4
	(b) Briefly discuss different types of microbial interactions with suitable examples.	[CLO2, PLO2, C6]	4