PH 501.4

Reg. No:	

St Aloysius College (Autonomous)

Mangaluru

Semester IV - P.G. Examination - M. Sc. Biotechnology

September - 2020

FOOD BIOTECHNOLOGY

Time: 3 Hours Max. Marks: 70

Note: Draw neat labeled diagrams/schematic sketches/structures wherever Necessary.

I Write short notes on any <u>FIVE</u> of the following:

(5x3=15)

- 1. Maillard reactions
- 2. Preservativation of volatiles
- 3. Factors affecting quality of foods
- 4. Smoking and pickling in preservation
- 5. SCP
- 6. Tempeh production
- 7. ISO 22000
- 8. Irradiation

II Write explanatory notes on any <u>FIVE</u> of the following:

(5x5=25)

- 9. Factors affecting growth of micro organisms in food
- 10. Macro nutrients in food. Explain each.
- 11. Principles of HACCP.
- 12. Production of cheese
- 13. Food processing using low temperature
- Preparation of soy sauce
- 15. Natural preservatives
- 16. Microbial food poisoning

III Answer any THREE of the following:

(3x10=30)

- 17. Discuss in detail on beer production
- 18. Write a detailed note on the types of food additives.
- 19. Discuss on biochemical changes in foods during processing and storage.
- 20. Describe the food spoilage mechanism in canned foods.
- 21. Explain in detail about the antinutritional factors and their effects with suitable examples

Reg. No:	

St Aloysius College (Autonomous)

Semester IV - P.G. Examination - M. Sc. Biotechnology Mangaluru

September - 2020

IMMUNOLOGY

ST. ALOYSIUS COLLEGE PG Library MANGALORE . 575 003

Max. Marks: 70

Time: 3 Hours

Note: Draw neat labeled diagrams/schematic sketches/structures wherever Necessary.

Write short notes on any FIVE of the following: I

(5x3=15)

- 1. Haptens
- 2. Type I hypersensitivity reaction
- 3. Immuno surveillance
- 4. Edible vaccines
- 5. Tumor antigens
- Cytokine antagonists
- 7. Characteristics of innate immunity
- 8. RIA

II Write explanatory notes on any <u>FIVE</u> of the following:

(5x5=25)

- 9. ELISA
- 10. Graft Vs Host disease
- 11. Antigen processing and presentation
- 12. Immunoglobulin structure and functions
- 13. Systematic lupus erythematosus
- 14. T cells activation and differentiation
- 15. Production of monoclonal antibodies
- 16. Antigen antibody reaction kinetics

III Answer any THREE of the following:

(3x10=30)

- 17. Discuss on antibody diversity and the mechanism of DNA rearrangements
- 18. Discuss on concepts in vaccine development and types of vaccines with suitable examples.
- 19. Discuss on a) Immune response to viral infections
 - b) Rheumatoid Arthritis
- 20. Describe the types of cytokines based on the functions with suitable examples.
- 21. Write a note on complement systems.

PS 505.4a

ST. ALOYSIUS COLLEGE
PG Library
MANGATORES

St Aloysius College (Autonomous) Mangaluru

Semester IV – P.G. Examination - M. Sc. Biotechnology September - 2020

IPR AND REGULATORY AFFAIRS

Time: 3 Hours Max. Marks: 70

Note: Draw neat labeled diagrams/schematic sketches/structures wherever Necessary.

I Write short notes on any <u>FIVE</u> of the following:

(5x3=15)

- 1. Copy right
- 2. Biopiracy
- 3. NOEL
- 4. Pharmacokinetics
- 5. Various regulatory bodies in clinical research
- 6. Role and responsibilities of IRB
- 7. Conditions for patenting inventions
- 8. Research integrity

II Write explanatory notes on any <u>FIVE</u> of the following:

(5x5=25)

- 9. Discuss the legislature regulating IPRs in India
- 10. Comment on CPCSEA guidelines for animal experimentation
- 11. What are clinical trials? Discuss the significance of various phases in clinical trials.
- 12. Explain patent revocation in India with an example.
- 13. Write about various animal models for preclinical research.
- 14. Geographical indications
- 15. Write a note on significance and ICH-GCP guidelines for preparing informed consent form.
- 16. Comment on various types of clinical research.

III Answer any THREE of the following:

(3x10=30)

- 17. Explain the regulatory requirements in filing a IND.
- 18. What are the requirements to be fulfilled for a patent grant? State the process from patent search till the grant of patent.
- 19. Discuss in details the good manufacturing practices in drug manufacturing.
- 20. Discuss the design and maintenance of case report form.
- 21. Describe in detail protection of plant varieties and the registration of a new plant variety.
