GOA UNIVERSITY Taleigao Plateau, Goa 403 206

# **REVISED MINUTES**

of the 5<sup>th</sup> Meeting of the Standing Committee of

X ACADEMIC COUNCIL

Day & Date

Tuesday, 14<sup>th</sup> February, 2023 & Thursday, 23<sup>rd</sup> February, 2023

<u>Time</u>

10.00 a.m.

Venue Council Hall, Administrative Block Goa University

D 3.12	Minutes of the Board of Studies in Portuguese meeting held on 18.10.2022.
	The Standing Committee of the Academic Council approved the minutes of the Board of Studies in Portuguese meeting held on 18.10.2022 with the suggestion to thoroughly verify the translation of titles/font in languages before uploading the Syllabus on the website.
	(Action: Assistant Registrar Academic-PG)
D 3.13	Minutes of the Board of Studies in Biochemistry meeting held on 22.10.2022.
	The Standing Committee of the Academic Council approved the minutes of the Board of Studies in Biochemistry meeting held on 22.10.2022 with the suggestion to replace terminology 'Text Books/References/Readings' with 'References/Readings.'
	(Action: Assistant Registrar Academic-PG)
D 3.14	Minutes of the Board of Studies in Philosophy meeting (by circulation)
	The Standing Committee of the Academic Council approved the minutes of the Board of Studies in Philosophy meeting held by circulation with the following suggestions:
	<ol> <li>Prerequisites for the Course PYTC 601 – Research Methodology to be added.</li> <li>Pedagogy for the Course to be included.</li> </ol>
	(Action: Assistant Registrar Academic-PG)
D 3.15	Minutes of the Board of Studies in Biotechnology meeting held on 13.10.2022. The Standing Committee of the Academic Council approved the minutes of the Board of Studies in Biotechnology meeting held on 13.10.2022 with the following suggestions:
	1. Heading 'Elective Generic Course' mentioned under Course Structure to be replaced with 'Generic Elective Course'.
	2. Course level listed below the Course structure to be deleted.
	<ol> <li>The synabus to be submitted as per prescribed synabus template.</li> <li>Number of hours of the Course Code GBTR-502, Bioprocess Technology to be corrected.</li> </ol>
	5. Course Code of the Course Scuba Diving to be verified.
	(Action: Assistant Registrar Academic-PG)
D 3.16	Minutes of the Board of Studies in Marathi meeting held on 19 and 20.10.2022. The Standing Committee of the Academic Council approved the minutes of the Board of Studies in Marathi meeting held on 19.10.2022 and 20.10.2022 with the following suggestions:
	<ol> <li>Title of the Courses to be indicated in both Marathi and English languages.</li> <li>Translation of titles/font in languages to be thoroughly verified before uploading on the website.</li> </ol>
	Hon'ble Vice-Chancellor (Chairperson) thanked Prof. K. S. Bhat on behalf of the Academic Council for his cooperation and contribution to the academic progress of

### GOA UNIVERSITY Taleigao Plateau, Goa 403 206

### FINAL AGENDA

For the 5<sup>th</sup> Meeting of the Standing Committee of

X ACADEMIC COUNCIL

Day & Date

Tuesday, 14<sup>th</sup> February, 2023

<u>Time</u>

10.00 a.m.

Venue Conference Hall Administrative Block Goa University

		Std. Com. X AC-5
		<u>14.02.2023</u>
	The meeting ended with closing remarks by Prof. K. Shripad Bhat, D School of Languages & Literature and Chairman, BoS Departmen Lusophone Studies.	: NIL ean, Shenoi Goembab nt of Portuguese and
	Date: 18.10.2022. Signature of	d/- the Chairman
	<ul> <li>Part G:</li> <li>Remarks of the Dean, Faculty of Languages and Literature: <ul> <li>i) The minutes are in order.</li> <li>ii) The minutes may be placed before the Academic Council.</li> <li>iii) May be recommended for approval by the Academic Council.</li> </ul> </li> </ul>	
	Date: 18.10.2022. Sd/- Dean, Shenoi Goembab School of Lan	guages and Literature
D 3.13	<ul> <li>Minutes of the Board of Studies in Biochemistry meeting held on 2</li> <li>Part A         <ol> <li>Recommendations regarding courses of study in the subject of the undergraduate level:</li> <li>NIL</li> <li>Recommendations regarding courses of study in the subject of the postgraduate level: (Detailed minutes of the BOS meeting</li> </ol> </li> <li>To discuss the revision of semester III &amp; IV syllabus of M.Sc. Biochemistry program         <ul> <li>After due deliberations and incorporating the sugge members, the BOS unanimously resolved to approve semester III and IV of M.Sc.Biochemistry program. The app III &amp; IV of M.Sc. Biochemistry program is attached as [Anne 496]</li> </ul> </li> </ul>	22.10.2022. or group of subjects at may please be seen). emistry stions made by the the draft syllabus of roved syllabus of SEM. exure I Refer page No.
	<ul> <li>Approval of syllabus for the Ph.D. Coursework paper in Research I (Paper-I; 4 Credits) for Research Scholars         <ul> <li>After due deliberations and incorporating the suggestions r the BOS unanimously resolved to approve the draft syllabutitled "Research methodology" for the Ph.D. Coursework The BOS approved syllabus of Ph.D. coursework in Researched as [Annexure II] Refer page No. 530]</li> </ul> </li> <li>Approval of around 16 PG level Swayam courses in Biochemistry a subjects that M Sc. students can offer</li> </ul>	Methodology, nade by the members, ous of 4 credit Paper-I for research scholars. earch Methodology is nd allied
	After going through the syllabus of total 22 different co portal, 16 were selected which Biochemistry PG students attached as [Annexure III] Refer page No. 533] Part B i. Scheme of Examinations at undergraduate level: NIL	ourses from SWAYAM s can offer. The list is

- iii. Scheme of Examinations at postgraduate level: NIL
- iv. Panel of examiners for different examinations at post-graduate level: NIL

#### Part C

i. Recommendations regarding preparation and publication of selection of reading material in the subject or group of subjects and the names of the persons recommended for appointment to make the selection:NIL

#### Part D

- i) Recommendations regarding general academic requirements in the Departments of University or affiliated colleges:NIL
- ii) Recommendations of the Academic Audit Committee and status thereof:NIL

#### Part E

- i. Recommendations of the text books for the course of study at undergraduate level:NIL
- ii. Recommendations of the text books for the course of study at post graduate level: NIL

#### Part F

Important points for consideration/approval of Academic Council

i. The important points/recommendations of BoS that require consideration/approval of Academic Council as mentioned below

### a) **PART-A (ii)**

ii. The declaration by the Chairperson:

Hereby, it is declared that the minutes were readout by the Chairperson at the meeting itself

Date: 22<sup>nd</sup> October 2022 Place: Taleigao Plateau Sd/-Signature of the Chairperson

**Part G** The Remarks of the Dean of the Faculty

- i. The minutes are in order.
- ii. The following important points / recommendations of BOS may be considered /approved by the Academic Council.

Attention of the Academic Council is drawn to item Nos. PART-A (ii)

- iii. May be recommended for approval of Academic Council.
- iv. Special remarks if any.

Sd/-

Date: 22.10.2022 Place: Taleigao Plateau Prof. V. M. S. Verenkar Dean, School of Chemical Sciences (Back to Index)

# D 3.13 Minutes of the Board of Studies in Biochemistry meeting held on 22.10.2022.

Annexure I

SI.	Subject	Paper title	Credits
No.	code		
	11	Research Specific	•
1.	CHBR-511	Practical Course in Biochemistry-III	4
2.	CHBR-512	Practical Course in Biochemistry-IV	4
3.	CHBR-513	Medical Biochemistry	4
4.	CHBR-514	Nanobiotechnology	4
5.	CHBR-515	Concepts in Genetic Engineering	4
6.	CHBR-516	Research methodology, Biostatistics and Bioethics	4
		Generic Elective	
1.	CHBG-511	Hormones and Neurochemistry	4
2.	CHBG-512	Clinical Microbiology and Food Biochemistry	4
3.	CHBG-513	Drug metabolism and Pharmaceutics	4
4.	CHBG-514	Bioprospecting and Bioremediation	4
5.	CHIG-515	Bioinorganic Chemistry	4
6.	CHAG-511	Fundamentals of Crystallography	4
		Dissertation	
1.	CHBD-511	Discipline Specific Dissertation	16

# M.Sc. Biochemistry Part-II revised syllabus (SEM III and SEM IV) for AY 2023-24 based on NEP-2020

### M.Sc. Biochemistry Part-II revised syllabus (SEM III and SEM IV)

SEM-III					
Sr. No.	Subject code	Paper title	Credits		
	Research Specific				
1.	CHBR-511	Practical Course in Biochemistry-III	4		
2.	CHBR-512	Practical Course in Biochemistry-IV	4		
3.	CHBR-515	Concepts in Genetic Engineering	4		
4.	CHBR-516	Research methodology, Biostatistics and Bioethics	4		
Generic Elective					

			<u>Std. C</u> 14	Com. X AC-5	
5.	CHBG-511	Hormones and Neurochemistry		4	
6.	CHBG-512	Clinical Microbiology and Food Biochemistry		4	
7.	CHBG-513	Drug metabolism and Pharmaceutics		4	
8.	CHBG-514	Bioprospecting and Bioremediation		4	
5.	CHIG-511	Bioinorganic Chemistry		4	
6.	CHAG-511	Fundamentals of Crystallography		4	
	SEM-IV				
Research Specific					
1.	CHBR-513	Medical Biochemistry		4	
2.	CHBR-514	Nanobiotechnology		4	
3.	CHBD-511	Discipline Specific Dissertation		16	

# Programme: M.Sc. Part-II (Biochemistry)

Course Code: CHBR-511

Title of the course: Practical course in Biochemistry-III

Number of Credits: 04 Total Hours: 120 Effective from AY: 2023-24

Prerequisites for the course:	Students should have studied biochemistry courses at MSc. part I level.
Course Objective:	<ol> <li>To acquaint the students with various methods of analyses of clinical samples for metabolic diseases/ disorders essential in pathological laboratories.</li> <li>To develop skills in the analysis of water samples according to critical parameters.</li> <li>To impart an understanding of various statistical operations needed to process biological data and improve technical writing skills.</li> <li>To develop techniques for handling, identification, and culturing of microorganisms.</li> </ol>
Course Outcome:	<ol> <li>Students will be able to analyse clinical samples for metabolic diseases/ disorders essential in pathological laboratories and further will be able to design various techniques in clinical biochemistry research.</li> <li>Students will be able to evaluate water samples and assess its suitability</li> <li>Students will be able to apply various statistical operations needed to process any biological data and have good technical writing skills.</li> <li>Students will be in a position to handle, culture, and identify microorganisms</li> </ol>

	Std.	. Com. X AC-5
Contract	<u> </u>	4.02.2023
Content		Hrs
A. Medical Biochemistry		30
Introduction to use of autoanalyzer and Rapid test for various cl	nical	
samples		
1. Analysis of blood sample: (ANY THREE)	L	
a. Examination of Haemoglobin (Hb) content of blood by copper sup	nate	
(FSR) of blood by Westergren method and ABO Blood grouping	for	
determination of blood group	, 101	
b. Examination of clotting time of blood by capillary tube method	and	
examination of total cell and differential cell (TC/DC) counts of blood sar	nple.	
c. Examination of blood glucose by glucose oxidase method or Folin-Wu me	thod	
or HbA1c rapid test		
d. Examination of blood cholesterol level by Zak's method.		
e. Rapid test for drug abuse		
f. Rapid test for pregnancy		
2. Liver function tests: (ANY ONE)		
a. Estimation of serum alarine transaminase (SGPT) and aspartate transam	nase	
b Estimation of serum bilirubin level by Malloy and Evelyn method		
3. Renal function tests:		
a. Physical examination of urine: assessment of volume, appearance, or	dour,	
color, pH and specific gravity and microscopic examination of u	rine:	
assessment of crystals, casts, cells in urine sample.		
b. Chemical examination of urine: (ANY ONE)		
i. Estimation of glucose in urine sample by Benedict's method and estim	ation	
of albumin content in urine sample by Sulfosalicylic acid method.		
ii. Estimation of blood urea by Diacetyl-monoxime method.		
B. Bioprospecting and Bioremediation (ANY FIVE)		25
1. Estimation of Dissolved oxygen (DO) and Biochemical Oxygen Dem	ands	
(BOD) of given water sample using Winkler method.		
2. Estimation of Chemical Oxygen Demands (COD) of water sample	and	
assessment of water quality using observed BOD and COD values.	ch ac	
5. Detection of sewage politition by screening for indicator organisms su		
4 Biotransformation of xenobiotics		
5. Bioassay: Antibiotic assays		
6. Techniques of strain improvement:		
a. Using UV radiations		
b. Using a Chemical mutagen		
7. Production of protoplast:		
a. Using lytic enzymes		
b. Using antibiotics.		
8. Immobilization of enzymes and determination of its activity.		

<u>Std.</u>	Com.	Х	AC-5
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	9.	Separa extrac	ation and purification of secondary metabolites from microbial ts using preparative HPLC.	
С.	<ul> <li>Bit</li> <li>1.</li> <li>2.</li> <li>3.</li> <li>4.</li> <li>5.</li> <li>6.</li> <li>7.</li> <li>8.</li> <li>9.</li> <li>10.</li> <li>11.</li> </ul>	ostatist Use of Develo To stu To car To dev To dev Forma measu error Study To stu ANOV . To est	<b>tics and technical writing (ANY FIVE)</b> graphical modes to represent biological data oping understanding for linear equation analysis (regression analysis). dy normal distribution curve ry out Hypothesis testing using Z-test and t-test velop scientific abstract writing skills. velop scientific reports writing skill atton of frequency distribution and calculation of descriptive ares-mean, median, mode, variance, standard deviation and standard application of biostatistics in pharmacology and medical research dy one way and two-way classification of biological data using A. timate genetic components and heritability using ANOVA data. alyse Paired t-test with an example.	25
D.	Clir 1. 2. 3. 4. 5. 6. 7. 8.	hical M Study Micros parasi Study Antibi Study Exami Study Study	icrobiology and food biochemistry (ANY FIVE) of the bacterial growth curve. scopic examination of blood films for identification of malarial tes/ Rapid test for malaria. and identification of bacterial pathogens. otic susceptibility testing for bacterial pathogens. and identification of fungi. nation of foods and determination of food spoilage microorganisms of Enzymatic browning of fruits of Auto Oxidation and Rancidity of fats.	25
Ε.	<b>QA</b> ; 1. 2. 3. 4. 5. 6.	and QC Qualit Mono To stu tablet To de perfor To ide with t Titrim /Furos UV Sp forms	<b>in pharmaceuticals (ANY THREE)</b> ative and Quantitative tests of Paracetamol/Aspirin as per IP graph dy the dissolution rate of sustained release Diclofenac/Theophylline s IP. velop and validate the analytical method of any one drug using high mance liquid chromatography. ntify the given drug amongst paracetamol, aspirin, and caffeine citrate he help of thin layer chromatography and calculate its Rf value. etric Assay of the following bulk drugs: Chloramphenicol capsules IP semide injection IP/Ketoprofen/ Phenytoin (Any 1) rectrophotometric Assay of the following drugs (in different dosage ): Mefenamic acid/ Furosemide/ Chloramphenicol (Any 1)	15
Pe	dag	ogy	Prelab exercises / assignments / presentations / lab hand-out or a com some of these. Sessions shall be interactive in nature to enable learning.	nbination of peer group

Text Books/ References / Readings	<ol> <li>Damodaran, G.; Practical Biochemistry; Jaypee Brothers Medical Publishers (P) Ltd., 2011.</li> <li>Mohanty, S.; Practical clinical Biochemistry; Jaypee Brothers Medical</li> </ol>			
	Publishers (P) Ltd., 2013, 1 <sup>st</sup> Edition.			
	3. Glasman-Deal H.; Science Research Writing, Imperial College Press, 2010.			
	<ol> <li>Vogel's Text book of Quantitative Inorganic Analysis, Pearson Education, Asia, 2000, 6<sup>th</sup> Edition.</li> </ol>			
	<ol> <li>Wilson K, Walker J; Principles and Techniques of Practical Biochemistry; Cambridge University Press; 2010; 7<sup>th</sup> Edition.</li> </ol>			
	<ol> <li>Sawhney, S. K., Singh, R.; Introductory Practical Biochemistry; Narosa Publishing House; 2005.</li> </ol>			
	<ol> <li>Poornima B., Food Science &amp; Quality Control, Centrum Press First, 2014, 1<sup>st</sup> Edition.</li> </ol>			
	<ol> <li>Sathe, A.Y., A first course in Food Analysis, New Age International Pvt. Ltd., 1999, 1<sup>st</sup> Edition.</li> </ol>			
	<ol> <li>Prescott, H. Laboratory exercise in Microbiology. MacGraw-Hill Companies. 2002, 5<sup>th</sup> edition.</li> </ol>			
	10. K. A. Connors, Text book of Pharmaceutical analysis, 3rd Edition, Wiley Interscience Publication, 1990.			
	<ol> <li>J. Moini, Pharmaceutical Laboratory Procedures, 1st Edition, Cengage Learning India Pvt. Ltd., New Delhi, 2010.</li> </ol>			

# Programme: M.Sc. Part-II (Biochemistry)

Course Code: CHBR- 512

Title of the course: **Practical course in Biochemistry-IV** 

Number of Credits: 04 Total Hours: 120 Effective from AY: 2023-24

Prerequisites for the course:	Students should have studied biochemistry courses at MSc. part I level.		
Course Objective:	<ol> <li>To develop hands-on experience of skills in various instruments and techniques in animal cell and tissue culture and microbial cells.</li> <li>To develop skills in genomics and proteomics</li> <li>To gain experience in bioprospecting of microbes for industrial purpose</li> <li>To study advanced analytical techniques in the separation and characterization of biomolecules.</li> </ol>		
Course Outcome:	<ol> <li>Students will be able to use various instruments and techniques in tissue culture and microbial culture.</li> <li>Students will be able to have skills in genomics and proteomics.</li> <li>Students will be able to apply the techniques in bioprospecting of microbes for industrial purposes</li> <li>Students will be able to use advanced analytical techniques in the separation and characterization of biomolecules.</li> </ol>		
A. Animal and p	ant tissue culture techniques and Microbial techniques (any nine) 45		

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<ol> <li>Animal tissue culture techniques:         <ul> <li>Laboratory safety protocols, Preparation of media and sterilizatitechniques.</li> <li>Primary cell culture</li> <li>Establishing cell lines</li> <li>Cell counting and viability techniques.</li> <li>Preservation of cell lines.</li> </ul> </li> <li>Plant tissue culture techniques:         <ul> <li>Laboratory safety protocols and Preparation of media and sterilizatitechniques.</li> <li>Germination of seeds in vitro.</li> <li>Establishment of primary culture and Micropropagation.</li> <li>Low cost strategies in plant tissue culture.</li> </ul> </li> <li>Microbial culture techniques:         <ul> <li>Laboratory safety protocols and Preparation of media and sterilizatitechniques.</li> <li>Setablishment of primary culture and Micropropagation.</li> <li>Low cost strategies in plant tissue culture.</li> </ul> </li> <li>Microbial culture techniques:         <ul> <li>Laboratory safety protocols and Preparation of media and sterilizatitechniques.</li> <li>Isolation and enumeration of bacterial and fungal cultures from varienvironmental samples.</li> <li>Identification of microbial isolates: Morphological and biochemidentification techniques.</li> </ul> </li></ol>	ion ion ous ical
<ul> <li>B. Genomics and proteomics (any six) <ol> <li>Isolation of genomic DNA from Prokaryotic cells.</li> <li>Isolation of genomic DNA from Eukaryotic cells.</li> <li>Isolation of RNA from prokaryotic cells</li> <li>Isolation of plasmid DNA using Rapid boiling and Alkaline lysis method.</li> <li>Isolation of protease degraders from soil and estimation of protease activ</li> <li>Quantitative Estimation of DNA and RNA</li> <li>Electrophoretic techniques and various gel staining techniques.</li> <li>DNA: PCR amplification, electrophoresis and purification.</li> <li>Molecular identification techniques for microbial isolates: understanding 16s and 18s rRNA sequencing, BLAST analysis and construction of phylogen trees.</li> <li>Protein identification techniques: understanding of protein sequence Protein BLAST, Protein Data bank (PDB) studies.</li> </ol> </li> </ul>	30 ity. g of etic ing,
<ul> <li>C. Advanced Analytical techniques in industry and research (any nine) <ol> <li>Extraction, purification and quantification of bioactive components fr different source</li> <li>Gas chromatographic analysis of volatile organic impurities in differ samples</li> <li>Purification of various analytes using advance chromatographic techniq such as size exclusion and ion exchange chromatography</li> <li>Fluorometric analysis of the vitamins and drug molecules</li> <li>Removal of impurity from commercial food products using adsorption column and analysis by potentiometry.</li> </ol> </li> </ul>	45 om ent ues on

		Std. Com. X AC-5
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7. Determi 8. Determi 9.Determi 10. Extrac chromatog 11. Qualita 12. Structu various spo 13.Decolor charcoal using gravi 14. Estima 15. Estima 16. Struc spectrosco	nation of potassium in plants by Flame Emission Spectroscopy nation of Caffeine in tablets by UV- visible spectroscopy nation of Aspirin in tablets by UV- visible spectroscopy tion and Separation of microbial pigments using TLC and pa graphy ative and quantitative analysis of given sample using HPLC ural elucidation of amino acids (proline/tryptophan/cysteine) us ectroscopic techniques. rization and crystallization of brown sugar (sucrose) with ani ity filtration. tion of lead/cadmium in water sample by AES/AAS/ICP. tion of iron/ manganese in water sample by AES/AAS/ICP. tural elucidation of carbohydrates (glucose) using vari opic techniques.	ous
Pedagogy	Prelab exercises / assignments / presentations / lab hand-out or some of these. Sessions shall be interactive in nature to en learning.	a combination of able peer group
Text Books/ References / Readings	<ol> <li>Freshney, R. I., Masters, J. R. W.; Animal Cell Culture: A Pr No. 232; Oxford University Press; 2002, 3<sup>rd</sup> Edition.</li> <li>Freshney, R. I., Culture of Animal Cells: A Manual of Bas Specialized Applications; Wiley-Blackwell; 2016, 7<sup>th</sup> Editio</li> <li>Smith, R. H.; Plant Tissue Culture: Techniques and Experi Press; 2012, 3<sup>rd</sup> Edition.</li> <li>Vogel's Text book of Quantitative Inorganic Analysis, Pe Asia, 2000, 6<sup>th</sup> Edition.</li> <li>Wilson K, Walker J; Principles and Techniques of Practi Cambridge University Press; 2010; 7<sup>th</sup> Edition.</li> <li>Sawhney, S. K., Singh, R.; Introductory Practical Bioch Publishing House; 2005</li> <li>Poornima B., Eood Science &amp; Quality Control. Centrum Press</li> </ol>	actical Approach: ic Technique and on. ments, Academic varson Education, cal Biochemistry; nemistry; Narosa
	<ol> <li>Poornima B., Food Science &amp; Quality Control, Centrum Pro Edition.</li> <li>Sathe, A.Y., A first course in Food Analysis, New Age Inter 1999, 1<sup>st</sup> Edition.</li> </ol>	ess First ,2014, 1 <sup>st</sup> national Pvt. Ltd.,

# Programme: M.Sc. Part-II (Biochemistry)

#### Course Code: CHBR-513 Title of the course: Medical Biochemistry Number of Credits: 04 Total Hours: 60 Effective from AY: 2023-24 Prerequisites Students should have studied biochemistry courses at MSc. part I level. for the course: Course To understand the biochemistry of metabolic diseases/disorders of the 1. Objective: human body.

	<ol> <li>To introduce knowledge on clinical investigations and analyse samples.</li> <li>To provide insights on biochemistry of cancer and ageing.</li> </ol>	s of clinical	
Course Outcome:	<ol> <li>Students will be able to explain the biochemistry of metabolic disorders/diseases caused due to imbalances and metabolic errors.</li> <li>Students will be able to illustrate the mechanisms of cancer and aging in the human body.</li> <li>Students will be able to employ technical knowledge for assessment of various clinical samples.</li> <li>The students will be able to devise strategies in designing experiments based on their understanding about physiological processes.</li> </ol>		
	Content	Hrs	
1. Analysis a. Blood sa i. ii. ii. iiv.	of Clinical sample mple Collection and safety measures involved. Composition and function: Composition of blood, RBCs, Erythropoiesis, Hemoglobin, gas transport by hemoglobin, Blood buffer system: acid- base balance and imbalance. Analysis: Haemoglobin, total cell and differential cell (TC/DC) counts, Erythrocyte sedimentation Rate (ESR); Bleeding time and Clotting time, glucose; lipid profile; urea; gases: oxygen and carbon dioxide levels; pH. Immunohaematology: Blood group systems – MN, Rh, ABO; hemolytic disease of newborn.	8	
b. Serum sa i. ii. iii.	Collection and safety measures involved. Analysis: Proteins, albumin/globulin ratio; bilirubin; creatinine; uric acid; electrolytes; Thyroid function tests (serum free and total T3 & T4 and serum TSH) Enzymes of clinical and diagnostic importance: Enzymes as markers in the diagnosis of diseases; clinical significance of cholinesterase, alkaline and acid phosphatase, lactate dehydrogenase (LDH), creatine phosphokinase (CPK), aspartate aminotransferase (AST/SGOT), alanine aminotransferase (ALT/SGPT).	7	
<b>c. Liver fun</b> i. ii. iii.	<ul> <li>c. Liver function tests (LFTs)</li> <li>i. Functions of the liver and liver profile in health and disease</li> <li>ii. Bilirubin metabolism and clinical significance</li> <li>iii. Classification of LFTs and their clinical significance in the diagnosis of liver diseases.</li> </ul>		

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<ul> <li>d. Renal function test (RFTs)         <ol> <li>Urine: Composition of urine, collection and safety measures,</li> <li>Kidney functions: Urine formation, glomerular and tubular functionate electrolyte balance.</li> <li>Analysis of urine/RFTs: Physical, chemical and micros examination.</li> </ol> </li> </ul>	tions, copic	4
e. Gastric and Pancreatic Function tests Gastric function tests (gastric analysis), hypo (achlorhydria) and hyper ac tests to confirm pancreatic involvement in disease.	idity,	2
<ul> <li>2. Metabolic disorders <ul> <li>a. Disorders in metabolism</li> <li>i. Carbohydrates: Regulation of blood glucose, insulin and diabetes metabolish (classification, stages and diagnosis); Hypoglycaemia; Diabetic ketoacidos hypercholesteremia,</li> <li>iii. Heart: Cardiovascular disease (Atherosclerosis and Coronary artery disethypertension</li> <li>iv. Proteins: Kwashiorkor, Marasmus Protein misfolding, Creutzfeldt-Jakob disease, mad cow disethyperchologithy</li> <li>v. Blood Anaemia: Iron deficiency anemia, Megaloblastic anemia, Pernianemia, Sickle cell disease, hemolytic anemia</li> <li>vi. Liver: Jaundice, cirrhosis</li> <li>vii. Kidney: Diabetes insipidus, Renal calculi.</li> </ul> </li> </ul>	ellitus sis. terol, ease), ease, icious	15
<ul> <li>b. Inborn errors of metabolism</li> <li>i. Prenatal diagnosis, newborn screening, laboratory investigations to diag metabolic disorders.</li> <li>ii. Carbohydrate: Lactose intolerance, galactosemia, Glycogen storage disea</li> <li>iii. Lipids: Lysosomal storage disorders: Tay-Sach'sdisease; Gaucher's dis Niemann Pick disease; Fabry's disease.</li> <li>iv. Amino acids: Phenylketonuria, Albinism</li> <li>v. Purine/pyrimidine: Lesch-Nyhan Syndrome, Gout.</li> <li>vi. Blood: Thalassemia</li> <li>vii. Thyroid hormone: hyperthyroidism and hypothyroidism</li> <li>viii. Skin: Xeroderma Pigmentosum</li> </ul>	;nose ise. ease;	7
<ul> <li>3. Biochemistry of cancer</li> <li>a. Properties of cancer cells</li> <li>b. Biochemistry of cancerous growth</li> <li>c. Etiology of cancer cells</li> <li>d. Apoptosis in carcinogenesis</li> <li>e. Metastasis</li> <li>f. Mutagens and carcinogens</li> </ul>		8

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g. Oncoge h. RNA vi i. Tumor j. Anticar	enic viruses: DNA viruses (Hepatitis B virus and Epstein-Barr virus ruses (Rous sarcoma virus and Human T-cell lymphotropic virus-1 markers ncer drugs	;) 1)
<ul><li>4. Biochemisti a. Definit</li><li>b. Ageing</li></ul>	<b>ry of ageing</b> ion and symptoms theories: Programmed theories and Error theories	4
Pedagogy Mainly lectures and tutorials. Seminars / term papers /assignments / presentations / self-study or a combination of some of these can also be used. ICT mode should be preferred. Sessions should be interactive in nature to enable peer group learning.		
Text Books/ References / Readings	<ol> <li>Vasudevan, D. M.; Sreekumari, S., Vaidyanathan, K. Biochemistry for Medical students, Jaypee brothers Me 2011, 6<sup>th</sup> Edition.</li> <li>Chattergee, M. N; Shinde, R.; Textbook of Medical Bioch brothers Medical publishers Ltd., 2012, 8<sup>th</sup> Edition.</li> <li>Smith, C.; Mark, A. D; Lieberman, M.; Marks' Basic Medica Clinical Approach; Lippincott's William and Wilkins; 2004, 2<sup>th</sup></li> <li>Gaw, A.; Cowan, R. A.; Murphy, M. J.; O'Relly, D. S. J.; Sriva Biochemistry, Elsevier; 2013, 5<sup>th</sup> Edition.</li> </ol>	, Textbook of dical publishers; hemistry, Jaypee I Biochemistry: A <sup>nd</sup> Edition. Istava, R.; Clinical

# Programme: M.Sc. Part-II (Biochemistry)

Course Code: CHBR-514

# Title of the course: Nanobiotechnology

Number of Credits: 04 Total Hours: 60

Effective from AY: 2023-24

Prerequisites for the course:	Students should have studied biochemistry courses at MSc. part I level.	
Course Objective:	<ol> <li>To introduce the concept of nanoparticles and nanomaterials.</li> <li>To understand methods to develop nanoparticles from plants and microbes.</li> <li>To familiarize students with different characterization tools, to identify bio-nanoparticle</li> <li>To develop an understanding of applications of Bio-nanomaterials in Health, Food, and the Environment.</li> </ol>	
Course Outcome:1. Students will be able to biosynthesize, and characterize nanoparticles 2. Students will be able to apply concepts of Nano-biotechnology in Healthcare, Environment and Food Industry.		
Content Hrs		
1. Introduction to biological cellular nanostructure and nanomaterials20		

	a. Introduction to nanobiotechnology: definition; historical background;	
	b. Basics of biology for nanobiotechnology: cell, organelles and nucleic acids as genetic material.	
	<ul> <li>c. Biological cellular nanostructures:</li> <li>i. Protein and Peptide based: Proteins, Bilayers and membrane arrays: ATPase</li> <li>Archaeal S-layers, bacteriorhodopsin</li> <li>ii. Eubacterial magnetosomes – greigite, magnetite.</li> </ul>	
	iii. DNA based: DNA molecule; self-assembled DNA nanotubes iv. Virus particles v. Diatoms	
2.	<ul> <li>Application of nanobiotechnology to biomineralization</li> <li>Nanomaterials</li> </ul>	
	<ul> <li>a. Shapes, size and properties: spherical, triangular, prisms, rods, cubes.</li> <li>Nanoparticles, nanocrystals, quantum dots, nanotubes and nanowires.</li> <li>b. Miniaturized devices in nanobiotechnology - types and applications</li> <li>c. Introduction to lab-on-a-chip (LOC).</li> </ul>	
3.	Biosynthesis of nanomaterials and characterization	15
	i. Concept of top-down versus bottom-up approach.	
	ii. Uniformity and heterogeneity.	
	collision efficiencies. <b>b. Green technologies:</b> nanoparticle biosynthesis using microbes, plant extracts,	
	reductases.	
	c. Detection and characterization of nanoparticles: Optical: i. Visual colour change; UV-Vis spectrum; Fluorescence, single molecule spectroscopy	
	ii. Size imaging: Electron microscopy (SEM, TEM), light scattering, FRET	
	iii. Zeta Potential surface and composition: FT-IR, Raman spectroscopy, EDAX, AFM, XRD, <sup>1</sup> H NMR, <sup>13</sup> C-NMR.	
4.	Nanobiotechnological applications in health and disease - infectious and chronic.	15
	A. Introduction to Biosensors:	
	i. Different classes -molecular recognition elements and transducing elements. ii. Applications of molecular recognition elements in nanosensing of different analytes	
	<ul> <li>iii. Various transducing elements as part of nanobiosensors.</li> <li>iv. Miniaturized devices in nanobiotechnology - types and applications,</li> <li>v. Lab on a chip concept (discussion with example)</li> <li>B. Medical Applications</li> </ul>	
	i. Drug development – Drug discovery; toxicity evaluation: cyto-toxicity, geno- toxicity.	

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ii. Diag stage ca iii. Nand iv. Drug v. Antii viruses. vi. Ther and Pa delivery	nostics – LOC technology; Imaging agents: MRI; nanosensors for early- ancer detection p-optics and fluorescence-based assays g delivery systems –Lipid and inorganic nanoparticles microbials – Metal/metal oxide nanoparticles against bacteria, fungi, rapeutics – Cardiovascular diseases; neurological disorders (Alzheimer's, rkinson's disease). Cancer therapy – quantum dots for targeted drug /.	
5. Nanobi mitigat a. Envir i. Nanol ii. Wate iii. Biore b. Food i. Magn ii. Nano iii. Nano	otechnological applications in Environment and food - detection and ion ronment analysis and remediation biosensors for pollution detection er purification – Nanoadsorbents and magnetic nanoparticles emediation –nanoparticles for degradation of biological pollutants i industry etosomes for detection of pathogens obiosensors for food quality monitoring.	10
PedagogyMainly lectures and tutorials. Seminars / term papers /assignm presentations / self-study or a combination of some of these can also be ICT mode should be preferred. Sessions should be interactive in nature to peer group learning.		ents / e used. enable
Text Books/ References / Readings	<ul> <li>Fext Books/</li> <li>1. Nicolini, C.; Nanobiotechnology &amp;Nanobiosciences, Jenny Stanfor References /</li> <li>Publishing., 2008,1<sup>st</sup> Edition.</li> <li>2. Niemeyer C.M., and Mirkin, C.A.; Nanobiotechnology, Concepts, Application and perspectives, Wiley- Verlag GmbH &amp; Co., 2004.</li> <li>3. DeVilliers, M.M., Aramwit, P., and Kwon, G.S.; Nanotechnology in Dru Delivery; Springer-American Association of Pharmaceutical Scientists Press 2009</li> <li>4. Yao, N. and Wang, Z.L.; Handbook of Microscopy for Nanotechnology. Kluw Academic Publishers., 2005</li> <li>5. Pradeep T.; Nano: The Essentials, Understanding Nanoscience ar Nanotechnology, Tata McGraw-Hill Publishing Company Limited., 2007, 2 Edition.</li> <li>6. Mirkin, C.A. and Niemeyer, C.M. Nanobiotechnology- II, More Concepts ar Applications, Wiley, Verlag GmbH &amp;Co., 2007</li> <li>7. Bulte, J.W.M. and Modo, M.M.J.; Design and Applications of Nanoparticles Biomedical Imaging, Springer International Publishing, 2016</li> <li>8. Shoseyov, O. and Levy, I.; Nanobiotechnology-Bio Inspired Devices ar Materials of the Future, Humana Press Inc., 2008.</li> </ul>	

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# Course Code: CHBR-515

Title of the course: Concepts in Genetic Engineering

Number of Credits:	04 Total Hours: 60 Effective from AY: 2023-24			
Prerequisites for the course:	Students should have studied biochemistry courses at MSc. part I le	vel.		
Course Objective:	<ol> <li>To introduce fundamental tools and techniques in Genetic engir</li> <li>To understand the mechanisms of recombinant DNA technology</li> <li>To familiarize the students with the applications of genetic engir agriculture, therapeutics, environment and industry.</li> </ol>	neering. / neering in		
Course Outcome:	<ol> <li>The students will be able to explain the tools and techniques i Genetic Engineering.</li> <li>The students will be able to apply the techniques learnt in red DNA technology.</li> <li>They will be able to explain the significance of transgenic org various sectors of human development.</li> <li>Students will be able to understand the risks and benefits of g modified organisms.</li> </ol>	nvolved in combinant ganisms in genetically		
	Content	Hrs		
<ol> <li>Introduction         <ol> <li>Concept of</li> <li>History and</li> <li>Introductio</li> <li>transforma</li> </ol> </li> </ol>	1. Introduction       5         a. Concept of genetic engineering       5         b. History and milestones       5         c. Introduction to gene manipulation tools (enzymes, hosts, vectors and transformation techniques).       5			
<ul> <li>2. Tools in Recombinant DNA technology         <ul> <li>a. DNA modifying enzymes: restriction endonucleases, exonucleases, DNA ligases, terminal DNA transferase, DNA polymerases, reverse transcriptase, T4 polynucleotide kinases, alkaline phosphatase, S-1 Nuclease, mung bean nuclease, RNases.</li> <li>b. Gene cloning systems/Hosts: Gene cloning in <i>E. coli, Saccharomyces cerevisiae</i>.</li> <li>c. Vectors: Plasmid (pUC19, pBR 322), λ phage-based vectors, cosmid vectors, phasmid vectors, shuttle vectors, high capacity cloning vectors.</li> <li>d. Gene transfer techniques: Transformation, electroporation, transfection, gene gun.</li> </ul> </li> </ul>				
3. Recombinant D a. Preparation b. Principles & c. Restriction d. Polymerase e. DNA Micro f. DNA sequ automated	NA techniques: n of probes & applications of nucleic acid hybridization, mapping, RFLP, e chain reaction: PCR, RT- PCR, real time PCR, parray encing using Sanger's dideoxy chain termination method and sequencer	10		
g. Gene editir	ng: Introduction to CRISPR/cas9 gene editing system.			

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4. Genetic Engine a. Screening b. Productio c. DNA vacc d. Edible vac e. Applicatio criminal c	eering in Biology, forensics and medicine of genetic diseases using DNA probes (DNA diagnostics). n of recombinant proteins and drugs (insulin, Antibodies), ines: merits and demerits ccines- merits and demerits on of recombinant DNA technology in paternity disputes and ases (DNA fingerprinting).	solving	10
5. Genetic Engine a. Importan b. Transgeni c. Significan d. Biofortific	eering in Agriculture ce of Agrobacterium tumefaciens c plants ce of Bacillus thuringiensis (Bt genes) cation of foods using genetic engineering.		8
6. Genetic Engine a. Developm b. Developm c. Animal cle	eering in Animal Husbandry and Aquaculture nent of transgenic animals nent of transgenic fish oning		6
<ul> <li><b>7. Genetically en</b></li> <li>a. Application</li> <li>b. Bioremedic. Safety and</li> </ul>	<b>gineered microbes in industries and the environment.</b> on of genetic engineering for enzyme production. iation using genetically modified microbes. d bioethics of genetically modified organisms.		5
PedagogyMainly lectures and tutorials. Seminars /term papers /assignments presentations / self-study or a combination of some of these can also be used ICT mode should be preferred. Sessions should be interactive in nature to enable peer group learning.			gnments / so be used. re to enable
<ul> <li>Text Books/ References /</li> <li>1. Old, R. W. and Primrose, S. B., Principles of Gene Manipulation: An introduction to Genetic Engineering, University of California Press 1981, 2<sup>nd</sup> edition.</li> <li>2. Glick, B. R., Pasternak, J. J. and Patten, C. L., Molecular Biotechnology: Principles and Applications of Recombinant DNA, ASM Press, 2010, 4<sup>th</sup> edition.</li> <li>3. Williamson, R., Genetic Engineering, Volumes 4-7, Academic Press 1981, 1<sup>st</sup>edition.</li> <li>4. Glover, D. M., Gene cloning: The Mechanics of DNA Manipulation, Springer, 1984.</li> <li>5. Green, M. R. and Sambrook, J., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory, New York (2014), 4<sup>th</sup> edition.</li> <li>6. Davis, L. G., Dibner, M. D. and Battey, J. F., Basic Methods in Molecular Biology, Elsevier, 1986, 1<sup>st</sup> edition.</li> <li>7. Gerhardt, P., Methods for General and Molecular Bacteriology, Elsevier 1994.</li> <li>8. Brown, T.A., Gene Cloning and DNA analysis: An introduction, John Wiley and Sons, UK, 2021, 8<sup>th</sup> edition.</li> </ul>			
	<u>(Back to Ind)</u> Programme: <b>M.Sc. Part-II (Biochemistry)</b>	<u>ex) (Back</u>	<u>k to Agenda)</u>

Course Code: CHBR-516

Title of the course: Research methodology, Biostatistics and Bioethics

Number of Credits: **04** Total Hours: **60** 

Effective from AY: 2023-24

Prerequisites for the course:	Students should have studied Biochemistry courses at MSc. part I le	vel.	
Course Objective:	<ol> <li>To develop a basic understanding of various types of biological data, its handling and processing.</li> <li>To introduce various technical writing skills.</li> <li>To understand various ethical considerations while studying biological data.</li> </ol>		
Course Outcome:	<ol> <li>Students will be able to collect, handle, process and present the biological data.</li> <li>Students will be able to apply statistics to biological data.</li> <li>Students will be able to develop the skills needed to successfully communicate through technical writing skills.</li> </ol>		
	Content	Hrs	
<ol> <li>Introduction to a. Basics of re i. Definition a scientific meth ii. Types of re researchers in iii. Research ap iv. Basic princi plan for data of analysis.</li> <li>Literature i. Primary and ii. Web source iii. Hypothesis null hypothesis iv. Research Monthesis do inductive log</li> </ol>	<ul> <li><b>D</b> Research, Research Design &amp; literature review</li> <li><b>esearch</b></li> <li>nd meaning of research, the significance of research, research &amp; nod.</li> <li>research, criteria for good research, problems encountered by India, selecting &amp; defining a research problem.</li> <li>Dproaches: research methods vs methodology.</li> <li>ples of experimental designs, sampling, sample size determination, collection, methods of data collection, plan for data processing and</li> <li><b>Review</b></li> <li>secondary Sources</li> <li>s –critical literature review</li> <li>Different types, significance, development of working hypothesis, esis</li> <li>Methods: <u>S</u>cientific method vs arbitrary method, logical scientific leductive, inductive, deductive-inductive, pattern of deductive – gical process, different types of inductive logical methods.</li> </ul>		
<ul> <li>2. Technical writing <ul> <li>a. Different forms of technical writing: articles, research notes and reports in journals, review articles, monographs, dissertations, bibliographies.</li> <li>b. How to formulate outlines: The reasons for preparing outlines, guide for plan of writing, skeleton for the manuscript, drafting titles, subtitles, tables, illustrations.</li> <li>c. Parts of dissertation/research report/article: introduction, review of literature, method, results and discussion.</li> </ul> </li> </ul>		5	

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<ul> <li>d. Significant subtopics related to scientific writing such as content, its contellarity, validity, internal consistency and objectivity</li> <li>e. Basic attributes for writing for grants.</li> </ul>	tinuity,	
<ul> <li>3. Introduction to Biological data <ul> <li>a. Basic characteristics of biological data</li> <li>i. Variables and constants, discrete and continuous variables, relationsh prediction, variables in biology (measurement, ranked, attributes), d variables (ratio, index, rates).</li> <li>b. Types of measurements in biological data</li> <li>i. Interval scale, ratio scale, ordinal scale, nominal scale, discrete and cont data, exact and approximate numbers.</li> <li>ii. Classification of errors, decimal notation and rounding off numbers, at and relative errors, valid significant digits, relationship between number or digit and error, the error of sum, difference, product, quotient, power an and rules of calculating digits.</li> </ul> </li> </ul>	ip and lerived inuous osolute of valid id root	10
<ul> <li>4. Data handling <ul> <li>a. Population and Sampling <ul> <li>i. Random samples, parameter and statistics, accuracy and precision, accin observations</li> <li>ii. Tabulation and types of frequency distribution: relative &amp; cumulative</li> <li>iii. Graphical representation: types of graphs, preparation and applications.</li> <li>b. Measures of central tendency: <ul> <li>i. Characteristics of ideal measure, arithmetic mean – simple, weicombined, and corrected mean, limitations of arithmetic mean;</li> <li>ii. Median – calculation for raw data, for grouped data, for continuous limitations of median;</li> <li>iii. Mode – computation of mode for individual series, by grouping methal a continuous frequency distribution, limitations of modes</li> <li>iv. Relationship between mean, median and mode</li> <li>c. Measures of dispersion:</li> <li>i. Variability, Range, mean deviation, coefficient of mean deviation, stateviation (individual observations, grouped data, continuous series)</li> <li>ii. Skewness – definition, positive, negative, purpose, measure, remeasure, iv. Karl Pearson's coefficient, Bowley's coefficient, Kelly's memoments.</li> </ul> </li> </ul></li></ul></li></ul>	curacy their ighted, series, hod, in andard relative easure,	15
<ul> <li>5. Correlation analysis, Population Biostatistics and Hypothesis testing         <ul> <li>a. Covariance, correlation coefficient for ungrouped and grouped scatter and dot diagram (graphical method).</li> <li>i. Regression analysis - linear and exponential function</li> <li>ii. Examples: DNSA conversion by reducing sugar, survival/growth of bar regression coefficients, regression analysis for linear equations.</li> <li>b. Population Biostatistics</li> </ul> </li> </ul>	l <b>data,</b> acteria,	15

		<u>Std. C</u> 14.	om. X AC-5 02.2023
i. Concep theory ii. Probab <b>c Hypoth</b> i. Hypoth ii. Level o single pro iii. Para t-test, Z- Introduc	bt of probability, theories of probability- additive and multipli pility distributions: binomial, poisson and normal mesis testing. Tesis and its types: Null and Alternative of significance, one tailed and two tailed test, test for single mea opportion, critical region, level of confidence, level of significance metric and Non- Parametric test test. F-test and ANOVA tion to Chi-square test	icative an and e,	
<ul> <li>6. Bioethics <ul> <li>a. Bioethics</li> <li>basic app</li> <li>b. Legal and</li> <li>c. Bioethics</li> <li>rights/PE</li> <li>commerce</li> <li>d. Bioethics</li> <li>GMOs, be</li> <li>of GMOs</li> <li>e. Past and</li> <li>f. Biopiracy</li> <li>plagiarising</li> <li>g. Bio-wast</li> </ul> </li> </ul>	E: Definition, ethics in biology, role and importance of ethics in bioroaches to ethics. If regulatory values related to bioethics. If in Healthcare, agriculture, biotechnology, animal welfare TA in research, wildlife conservation and manage cialization in scientific research. If related to genetically modified organisms (GMOs): concerns enefit and risk of GMOs, reasoning behind acceptance and reju- present bioethical conflicts in life sciences. If thical committees, copyright, royalty, IPR and patent m, citation and acknowledgement. If disposal: Types of biowaste, ways to dispose of biowaste.	iology, e and ement, about ection t law,	5
Pedagogy	Mainly lectures and tutorials. Seminars / term paper presentations / self-study or a combination of some of these ca mode should be preferred. Sessions should be interactive in na group learning	rs /assi an also ature to	ignments / be used. ICT enable peer
Text Books/ References / Readings	<ol> <li>Daniel W.W.; Biostatistics: Basic Concepts and Methodo Sciences, Wiley publishers, 2014, 10<sup>th</sup> Edition</li> <li>Antonisamy B.; Premkumar P.S.; Christopher S.; Princip Biostatistics, Elsevier India, 2017, 1<sup>st</sup> Edition</li> <li>Glasman-Deal H.; Science Research Writing, Imperial C</li> <li>Kothari, C. R.; Quantitative Techniques, Vikas Publishin Edition</li> <li>Arora, P. N. and Malhan, P. K.; Biostatistics, Himalaya 2006, 9<sup>th</sup> Edition</li> <li>Surya, R. K.; Biostatistics for health and life sciences, H House., 2010, 1<sup>st</sup> Edition</li> <li>Annadurai A.; A Textbook of Biostatistics, New Age Pu Edition.</li> </ol>	ology fo ples and College F ng House a Publis Himalay ublicatic	or the Health d Practice of Press, 2010. e., 2013, 3 <sup>rd</sup> hing House. a Publishing on, 2017, 1 <sup>st</sup>

Programme: M.Sc. Part-II (Biochemistry)

Prerequisites for the course:	Students should have studied life sciences at M.Sc. Part I Level	
Course1. To develop knowledge on the human endocrine system human physiology.2. To acquaint students with the mechanism of hormor regulation and clinical disorders associated with them.3. To develop insights into the structure and organization system, sensory organs and their functions.4. To develop a basic understanding of the significance of neurotransmitters.5. To introduce the biochemistryof mental disorders.		em and its role in none action, their tion of a nervous
Course Outcome:	<ol> <li>Students will be able to apply the knowledge of the sig mechanisms of different hormones in the human syste</li> <li>The students will also be able to correlate the diseases hormonal imbalance and the biochemistry behind then</li> <li>Students will be able to explain the significance of the for the normal functioning of the human body.</li> <li>Students will be able to illustrate the role of neurotrans generation and the biochemistryof mental disorders in</li> </ol>	nalling m. associated with n. nervous system smitters in signal the human body.
	Content	Hrs
	Hormones	
<ol> <li>Introduct</li> <li>a. Defin of h</li> <li>b. Unde</li> <li>c. Signa signa</li> <li>d. Horm</li> <li>throu</li> <li>Thyro</li> <li>e. Mech</li> </ol>	tion to hormones ition, history, classification, and mechanism of action, History ormones, Classification of hormones. Instanding of endocrine system, Pathways of hormone release, I transduction pathways, second messengers, regulation of ling Pathways. Nones and their receptors: cell surface receptor, signaling righ G-protein coupled receptors, Steroid hormone receptors, bid hormone receptors nanism of sensitization & desensitization of hormone receptors	6
<ul> <li>2. Stimulus,</li> <li>a. Hypo</li> <li>b. Anter</li> <li>pepti</li> <li>c. Poste</li> <li>d. Adrer</li> <li>&amp; cor</li> </ul>	, regulation of biosynthesis and release of hormones thalamic Hormones: CRH, TRH, GnRH, PRL/PRIH, GHRH/GHRIH for Pituitary hormones: Growth hormone, Prolactin, POMC de family, LH, FSH, TSH erior Pituitary Hormones: Vasopressin, Oxytocin nal Cortex Hormones: Aldosterone (renin angiotensin system) tisol	9

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<ul> <li>e. Hormones of Adrenal Medulla: Epinephrine and norepinephrine Hormones regulating Ca2+ Homeostasis: PTH, Vitamin D, Calcitonin</li> <li>f. Pancreatic Hormones: Insulin, Glucagon.</li> <li>g. Gl tract Hormones: Gastrin, Secretin, CCK, GIP, Ghrelin.</li> </ul>	
<ul> <li>3. Reproductive hormones and hormones by organs with endocrine function: <ul> <li>a. Reproductive Hormones: Male and female Sex hormones, interplay of hormones during reproductive cycle, pregnancy, parturition and lactation. Introduction to rapid test for pregnancy.</li> <li>b. Role of oral contraceptives.</li> <li>c. Other organs with endocrine function: Heart (ANP), Kidney (erythropoietin), Liver (angiotensinogen, IGF-1), adipose tissue (leptin, adiponectin); growth factors: PDGF, EGF, IGF-I, II</li> </ul> </li> </ul>	6
<ul> <li>4. Biochemistry and diseases associated with hyper or hypo secretion: <ul> <li>a. Hypothalamus and pituitary associated hormonal conditions: Goiter, Graves' disease, Cretinism, Myxedema, Hashimoto's disease, Gigantism, Acromegaly, dwarfism.</li> <li>b. Adrenal cortex-associated hormonal conditions: Addison's disease, Conn's syndrome, Cushing's syndrome,</li> <li>c. Calcium homeostasis-related hormonal conditions: Rickets, Osteomalacia, Osteoporosis.</li> <li>d. Pancreatic hormone-associated hormonal conditions: Diabetes insipidus.</li> </ul> </li> </ul>	9
Neurochemistry	
<ol> <li>Organization of Nervous system: Definition, parts and anatomy         <ul> <li>Central Nervous system and Peripheral nervous system; Blood Brair Barrier.</li> <li>Cerebrospinal fluid: composition, function and circulation.</li> <li>Cellular components of nervous system: Nerve, neuron, neuroglial cells</li> </ul> </li> </ol>	4
<ul> <li>2. Nerve cell Membranes:</li> <li>a. Structures and Functions of nerve cells and membrane transport:         <ol> <li>Phospholipid bilayer, membrane proteins, Biological membrane</li> <li>Membrane transport: Primary ion transporters, Ca2+ pumps, V-ATPase pump, secondary active transport, cation antiporters, facilitators</li> </ol> </li> </ul>	3
<ul> <li>b. Energy metabolism in brain:</li> <li>Substrates for cerebral energy metabolism, regulation of the cerebral metabolic rate, glycolysis, glycogen metabolism, Pentose, phosphate shunt, Malate–aspartate shuttle, lactate metabolism, TCA, Glutamate/glutamine metabolism.</li> </ul>	3

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<b>3. Synaptic Tra</b> a. Synapse in stead post syn	nsmission: structure, Chemical and Electrical synapses, membrane potential ly state, Action potential generation and propagation, pre and naptic events.	4
b. Neurotra metabol Acetylcholi Inhibitory Glycine, Se (Cannabino c. Sensory	ansmitters and neuromodulators: Structure, functions, ism, receptors: ine, Excitatory Amino Acids (EAAs): Glutamic Acid, Amino Acids (IAAs): g-Aminobutyric Acid and rotonin (5-HT), Catecholamine, Purines bids), Neuropeptides and Nitric oxide.	4
balance	e, touch.	
d. <b>Bioche</b> i. Biochen (Qualita and mer ii. Mental iii. Neuro disease	mistry of memory; mental and neurodegenerative disease: nistry of memory: Learning and memory; Divisions of memory tive and Quantitative categories); Synaptic signalling in learning mory illness: Depression, Schizophrenia degenerative diseases: Alzheimer's disease, Parkinson's , Huntington's disease, Dementia	6
e. <b>CNS activ</b> Drugs of receptors	e drugs and drugs of abuse: classification and mode of action abuse: Opiates, Nicotine, alcohol: Molecular mechanisms, and signalling	3
Pedagogy	Mainly lectures and tutorials. Seminars / term papers /assignme / self-study or a combination of some of these can also be used be preferred. Sessions should be interactive in nature to e learning.	ents / presentations d. ICT mode should enable peer group
Text Books/ References / Readings	<ol> <li>Kline B and Rossmanith W.G, Hormones and the endocrin 2016</li> <li>Ilie I.R. Introduction to endocrinology, Springer, 2020.</li> <li>Berg, J.M., Stryer, L., Tymoczko, J., Gatto, G., Biochemi 2019, 9<sup>th</sup> edition.</li> <li>Mathews, C.K., van Holde, K.E. &amp; Ahern, K.G. Bioc Publishers, 1999, 3<sup>rd</sup> edition</li> <li>Nelson, D. L. and Cox, M. M. Lehninger Principles of Freeman, 2017, 7<sup>th</sup> edition.</li> <li>Norman A. W., Gerald Litwack. Hormones, Elsevier, 1997</li> <li>David, G. &amp; Dolores, S., Greenspan's Basic and Clinical Graw Hill Education, (2018), 10th edition</li> <li>Belfiore A and Leroith D, Principles of Endocrinology an Springer, 2018.</li> <li>Albers, R.W., Brady, S.T., Price, D. L., Basic neuroche cellular and medical aspects., Elsevier Academic Press p Edition.</li> </ol>	e system, Springer, stry, WH Freeman, chemistry, pearson Biochemistry, WH 7, 2 <sup>nd</sup> edition Endocrinology, Mc ad hormone action, emistry: Molecular, ublishers; 2006, 7 <sup>th</sup>

E

- 10. Smith, C.U.M., Elements of Molecular Neurobiology., John Wiley & Sons Ltd., 2002, 3<sup>rd</sup> edition.
- 11. Kandel, E.R., Swartchz, J.H., Jesselle, T.M., Principles of Neural science, McGraw-Hill, New York publishers., 2000, 4<sup>th</sup> edition.
- 12. Mathew, B., Parambi, T., Principles of Neurochemistry: Fundamentals and Applications. Springer, Singapore., 2020.

### Programme: M.Sc. Part-II (Biochemistry)

#### Course Code: CHBG-512

Title of the course: (	<b>Clinical Microbiology and</b>	<b>Food Biochemistry</b>
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Number of Credits: 04	Total Hours: 60	Effective from AY: 2023-24

Prerequisites for the Course:	Students should have studied life sciences at M.Sc Part I Level	
Course Objective:	<ol> <li>To develop an understanding of the diseases caused by microorg and their biochemistry.</li> <li>To develop a basic understanding on significance of commensa normal microflora for human health.</li> <li>Introduction of the fundamental concepts of food spoilage and preservation.</li> <li>To provide insights on quality control and good practices in the industry.</li> </ol>	anisms II and food food
Course Outcome:	<ol> <li>Students will be able to explain the significance of normal microbiota and the biochemistry of infectious diseases in the human body.</li> <li>Students will be able to explain the importance of antimicrobial agents in antibiotic therapy.</li> <li>They will be able to apply the concepts of food spoilage and food preservation in maintaining food safety.</li> <li>The student will be able to implement the Good Laboratory Practices and Good Manufacturing Practices used in industries to maintain food hygiene.</li> </ol>	
	Content	Hrs
<ol> <li>Introduct         <ol> <li>Introduct</li> <li>Introduct</li> <li>Sterilization</li> <li>its applicits</li> </ol> </li> </ol>	ion to Microbiology ction to bacteriology, mycology, virology and parasitology. tion and Disinfection: Introduction and its types, principle, procedure and cation, biosafety in microbiology lab, biowaste management.	3

	Std. Com. X AC-5
	<u>14.02.2023</u>
<ul> <li>2. Normal microbial flora and pathogenic microorganisms <ul> <li>a. Introduction: Distribution of the normal microbiota; Commensals; between normal microbiota and host; collection and transport of processing of clinical specimens for microbiological examination.</li> <li>b. Human microbiota in health: functions, microbe-host interaction, heal Skin microbiota, Gut microbiota, Normal microbiota of oral cavity, Norma of genitourinary tract</li> <li>c. Human microbiota in disease <ul> <li>i. Human microbiota and infectious disease: Opportunistic Nosocomial infections; bacterial Infections: Gastroenteric (<i>Clostridi Helicobacter pylori; E. coli</i>); Skin (<i>Staphylococcal</i>); Respiratory (<i>St. Pneumococcal, tuberculosis</i>); Urogenital tract (UTIs, Bacterial vagi cavity (Dental caries, Periodontitis).</li> <li>ii. Human microbiota and metabolic disorders: Irritable bowel disea Type 2 diabetes mellitus; Allergic diseases; Liver diseases.</li> <li>iii. Secondary infections: Infections associated with HIV; Influenza</li> </ul> </li> </ul></li></ul>	12 relationship specimens, th benefits: I microbiota infections; <i>ium difficile</i> ; <i>reptococcal</i> , inosis); Oral se; Obesity;
<ul> <li>3. Fungal and parasitic infections         <ul> <li>a. Fungal infections/mycoses: Cutaneous, Sub-cutaneous, systemic and or mycoses</li> <li>b. Parasitic infectious:</li></ul></li></ul>	pportunistic
<ul> <li>4. Viral infections:</li> <li>HIV, Influenza, Poliomyelitis, Dengue fever, Chikungunya, Hepatiti</li> <li>Coronavirus disease (COVID-19)</li> </ul>	s, Rabies.
<ul> <li>5. Antimicrobial agents and drug resistance:</li> <li>a. Classification, mechanism of action of antibacterial agents; antifung antiviral agents and their resistance</li> <li>b. Antibiotic sensitivity tests and its medical importance</li> </ul>	gal agents;
<ul> <li>4. Food Spoilage and Food Preservation <ul> <li>a. Forms of food spoilage: physical, chemical, microbiological parameters.</li> <li>b. Factors affecting the growth and survival of microorganisms in foods: In extrinsic factors</li> <li>c. Predictive food spoilage microbiology of milk, meat, poultry, vegetables grains and legumes.</li> <li>d. Food preservation technologies: Traditional methods of food preservat processing, low temperature storage, control of water activity, irradia pressure processing, modified atmospheres, preservatives (chemical organic molecules (nisin) and enzymes).</li> </ul> </li> </ul>	12 Itrinsic and and fruits, tion, Heat Ition, high s, natural
<ul> <li>5. Vitamins and minerals in health</li> <li>a. Fat soluble vitamins: physiological role, deficiency disorders, toxicity.</li> <li>b. Water soluble vitamins: physiological role, deficiency disorders, toxicity</li> <li>c. Mineral metabolism, physiologic role and deficiency disorders: calcondered</li> </ul>	cium, iron,

<u>Std. Co</u> <u>14.0</u>	<u>m. X AC-5</u> 2.2023
magnesium, sodium, zinc, manganese, potassium, phosphorus, sulphur ar chlorine.	nd
<ul> <li>6. Quality control and Quality Assurance in Food industries <ul> <li>a. Microbiological examination of food, air and water in industries.</li> <li>b. Plant sanitation</li> <li>c. Hazard analysis and critical control point concept</li> <li>d. Good lab practices (GLP) Good Manufacturing Practice (GMP) and Quality Systee in the food industry.</li> </ul> </li> </ul>	8 ms
PedagogyMainly lectures and tutorials. Seminars / term papers /assignments / pro / self-study or a combination of some of these can also be used. ICT m be preferred. Sessions should be interactive in nature to enable p learning.	esentations ode should peer group
<ol> <li>Text Books/ References / Readings</li> <li>Tortora, G. J., Funke, B. R., Case, C. L., Microbiology: An Intr Pearson Benjamin Cummings publishers; 2010, 10<sup>th</sup> Edition.</li> <li>Willey, J., Sandman, K., Wood, D.; Prescott's Microbiology., Mc G 2020, 11<sup>th</sup> Edition.</li> <li>Harvey, R. A., Cornelissen, C. N., Fisher, B. D., Lippincott's Illustrat Microbiology., Lippincott's William and Wilkins; 2007, 3<sup>rd</sup> Edition</li> <li>Chauhan, N. S. Introductory Chapter: Human and Microbes in F Diseases. In Role of Microbes in Human Health and Diseases. Int 2019.</li> <li>Feng, Q., Chen, W. D., &amp; Wang, Y. D. (2018). Gut microbiota: a moderator in health and disease. Frontiers in microbiology. 9, 15</li> <li>Frazier, W. C &amp;Westhoff, C.W. Food Microbiology. Graw-Hill C Inc., New York (2017), 5<sup>th</sup> edition.</li> <li>Hayes, P. R. Food Microbiology and Hygiene. Springer, 1995, 2<sup>nd</sup></li> <li>Kniel, K. E., Montville, T. J., Matthews, K. R, Food Microbiolo Press, NW Washington, USA., 2017, 4<sup>th</sup> edition</li> <li>Jay, J. M., Loessner, M.J., Golden, D.A., Mode Microbiology. Springer Science, New York, 2005, 7<sup>th</sup> edition</li> <li>Adams, M. R. &amp; Moss, M. O. Food Microbiology. Royal Sc Chemistry, 2015, 4<sup>th</sup> edition</li> <li>Mudambi, R. Sumathi, Rajagpal M.V, Fundamentals of Food, Nu diet therapy, New age International Publishers, 1983, 6<sup>th</sup> edition</li> </ol>	oduction., raw Hill., ed review: ealth and echOpen., n integral 1. ompanies, edition. gy., ASM m Food ociety of trition and

Programme: M.Sc. Part-II (Biochemistry)

Course Code: CHBG-513Title of the course: Drug metabolism and Pharmaceutics (GE)Number of Credits: 04Total Hours: 60Effective from AY: 2023-24

<u>Std. Com. X AC-5</u> <u>14.02.2023</u>

Prerequisites for the course:	Students should have studied natural and life sciences at M.Sc Part I Lev	vel
Course Objective:	<ol> <li>To introduce concepts of drug administration, distribution, metabolism and excretion.</li> <li>To introduce the students to pharmacopoeia, and types of drug formulations.</li> <li>To acquaint the students with GMP and quality control practices in a pharmaceutical set-up.</li> </ol>	
Course Outcome:	<ol> <li>Students will be able to explain the basic pathways of drug dis metabolism and excretion in the body.</li> <li>Students will be able to categorize different types of drug formula their contents.</li> <li>They will be able to implement quality assurance and quality procedures for drug formulations.</li> </ol>	tribution, tions and y control
	Content	Hrs
<ol> <li>Drugs A         <ol> <li>Definition</li> <li>Introduct</li> <li>Introduct</li> <li>Routes</li> <li>metabo</li> <li>Absorpt</li> <li>Factors</li> <li>organ/ti</li> <li>distribut</li> <li>Protein/significa</li> </ol> </li> </ol>	bsorption and distribution in human body: on and types of drugs (therapeutic, drugs of abuse, poisons). ction to pharmacokinetics and pharmacodynamics. of drug administration, introduction to absorption, distribution, lism, and excretion (ADME) of drug. ion and distribution of drug through organ /tissue. affecting drug distribution: Physicochemical properties of drugs, issue size, blood flow to the organ, physiological barriers to the tion of drugs, drug binding blood/ tissue/ macromolecules. /tissue binding of drugs – factors affecting protein binding of drugs, ince and kinetics, tissue binding of drugs.	6
<ul> <li><b>2.</b> Drug Ma</li> <li>a. Biotran</li> <li>drug me</li> <li>b. Mechan</li> <li>interme</li> <li>c. Phase 1</li> <li>Dealkyla</li> <li>Non-CYI</li> <li>Contain</li> <li>Reducta</li> <li>d. Phase 2</li> <li>(minor),</li> <li>e. Significa</li> </ul>	etabolism sformation of drugs and factors affecting biotransformation. Organs of etabolism: hepatic and extrahepatic metabolism. iisms of drug metabolism — inactivation, bioactivation, reactive diates. reactions - CYP-Catalyzed: Hydroxylation (Primarily at C, N, some at S), ation (N- and O-dealkylation), Deamination, Epoxidation, Reduction. P-Catalyzed: Oxidation (Alcohol and Aldehyde Dehydrogenase, Flavin- ing Monooxygenase, Monoamine Oxidase), Reductase (Quinone use), Hydrolysis (Esterases, Amidases, Epoxide Hydrolase). reactions -Glucuronidation, Sulfation, Acetylation, Glycine conjugation of drug metabolism (paracetamol/aspirin/ ibuprofen/ antibiotics).	7
<b>3. Excretic</b> a. Renal ex b. Non rer circulati	on of drugs Accretion, factors affecting renal excretion. Thal routes of excretion, factors affecting excretion and enterohepatic on.	2

			<u>Std. Co</u>	<u>m. X AC-5</u> 2 2023
	<b>4.</b> a. b.	Posology Determination of doses; dose response relationship, dosage form biopharmaceutical consideration. Drug antagonism and drug–drug interaction.	design,	2
	<b>5.</b> a. b.	<b>Drug Extraction</b> Solvents used in extraction of drugs, processes used for extraction (in decoction, maceration, percolation, hot extraction). Water as a universal pharmaceutical vehicle.	nfusion,	5
	<b>6.</b> a.	<b>Types of formulations:</b> Tablets: advantages of tablets; types of tablets: effervescent, lo chewable, buccal and sublingual, dispersible, orodispersible, soluble; ex in tableting, coating in tablets.	ozenges, cipients	
	b. c.	Granulation: methods and equipment, direct compression. Sustained release: Delayed absorption and/or a mixture of slow- and release particles to produce rapid and sustained absorption in the same	nd fast- dose.	
	d.	Capsules: hard gelatin and soft gelatin capsules- differences and comp advantages and limitations, Excipients in capsule.	oosition,	15
	e.	Liquids and Gels: Types of liquid formulations, excipients including solu stabilizers, buffers, tonicity modifiers, bulking agents, w enhancers/reducers, surfactants, chelating agents and adjuvants, hydr	bilizers, /iscosity rophilic-	
	f.	Parenterals: Intravenous, subcutaneous, intramuscular or intra a administration, stored in liquid form, or in lyophilized form if unstable.	articular	
-	б·			
	7. a.	Quality assurance/ Quality control Introduction to GLP, GMP and SOPs Raw material analysis (RMA), control of pharmaceutical excipients.	Quality	
	b.	Packaging material testing (PMT): Permeability of plastic; testing of foil, carrions. Limit tests – chloride, sulphate, arsenic, lead, iron, nitrate, al alkaline earth metals Limits of insoluble matter, soluble matter, nor matter, volatile matter, residue on ignition and ash value.	bottles, kali and wolatile	15
	C.	Sources of contamination in pharmaceutical compounds (a Pharmacopoeia).	as per	
	d. e.	Physico-chemical and microbiological analyses of formulations. Types of errors, selection of sample, precision and accuracy.		
	8.	Drug Stability		
	a.	Solid state, solution phase physical stability testing, Stability testing	general	
	b.	Kinetic principles applied for stability evaluation and their application predicting shelf life, accelerated stability study and shelf life assignment	tions in	5
	с.	roited degradation studies.		
	<b>9.</b> a.	Research and Development Introduction to drug design		3

b. Drug o c. Clinica	<ul><li>b. Drug discovery and development</li><li>c. Clinical trials.</li></ul>		
Pedagogy	Mainly lectures and tutorials. Seminars / term papers /assignments / pre / self-study or a combination of some of these can also be used. ICT mode preferred. Sessions should be interactive in nature to enable peer group I	rs / term papers /assignments / presentations of these can also be used. ICT mode should be tive in nature to enable peer group learning.	
Text Books/ References / Readings	<ol> <li>Brunton, L. L., Hilal-Dandan, R., Knollmann, B. C.; Goodman &amp; Gilr Pharmacological Basis of Therapeutics, McGrawHill Education, 2018, 13<sup>th</sup></li> <li>Mahato R. I., Narang A. S., Pharmaceutical Dosage Forms and Drug Revised and Expanded, CRC Press, 2017, 3<sup>rd</sup> Edition.</li> <li>Aulton, M. E., Pharmaceutics: The Science of Dosage Form Design Livingstone; 1988, 7<sup>th</sup> edition.</li> <li>Aulton, M. E., Taylor, K.; Aulton's Pharmaceutics: The Design and Manu Medicines, Elsevier, 2017, 5<sup>th</sup> Edition.</li> <li>Allen, L., Popovich, N. G., Ansel, H.; Ansel's Pharmaceutical Dosage Form Delivery Systems, Lippincott Willimas &amp; Wilkins, 2018, 11<sup>th</sup> Edition</li> </ol>	<ul> <li>J., L. L., Hilal-Dandan, R., Knollmann, B. C.; Goodman &amp; Gilman's: The ogical Basis of Therapeutics, McGrawHill Education, 2018, 13<sup>th</sup> Edition.</li> <li>R. I., Narang A. S., Pharmaceutical Dosage Forms and Drug Delivery: d Expanded, CRC Press, 2017, 3<sup>rd</sup> Edition.</li> <li>M. E., Pharmaceutics: The Science of Dosage Form Design, Churchill e; 1988, 7<sup>th</sup> edition.</li> <li>M. E., Taylor, K.; Aulton's Pharmaceutics: The Design and Manufacture of Elsevier, 2017, 5<sup>th</sup> Edition.</li> <li>Popovich, N. G., Ansel, H.; Ansel's Pharmaceutical Dosage Forms and Drug <i>r</i>stems, Lippincott Willimas &amp; Wilkins, 2018, 11<sup>th</sup> Edition</li> </ul>	
	(Back to Index) (Back	to Agenda)	

## Programme: M.Sc. Part-II (Biochemistry)

### Course Code: CHBG-514

# Title of the course: Bioprospecting and Bioremediation

Number of Credits: 04Total Hours: 60Effective from AY: 2023-24

Prerequisites for the course:	Students should have studied natural and life sciences at M.Sc. Part I Level		
Course Objective:	<ol> <li>To introduce the concept of bioprospecting of bioactive compounds plant and microbial sources.</li> <li>To impart knowledge on purification and characterization of novel metabolites from biological sources using analytical techniques.</li> <li>To develop concepts in environmental pollution and role of microorganisms in biogeochemical cycles and bioremediation of po</li> </ol>	s from Ilutants	
<ol> <li>Students will be able to screen, isolate and identify novel biomolecules from plant and microbial sources.</li> <li>The students will be able to correlate the role of microorganisms in biogeochemical cycles and design strategies for bioremediation of pollutants.</li> </ol>		ules	
	Content Hrs		

	Std. Com.	<u>X AC-5</u>
	<u>14.02.2</u>	023
1. Sources and Sampling of potential microbes and plants sources		6
a. Sources: microbes and plants		-
i. Marine and other coastal ecosystems: Water and sediment	samples,	
, microorganisms from mangroves, sand dunes and salterns.	1 /	
ii. Terrestrial: Forest/Ghats		
iii. Microbes in Extreme environments: thermophilic, psychrophilic,	halophilic,	
alkaliphilic, barophilic	·	
b. Sampling microorganisms		
i. Niskin water sampler		
ii. Van Veen Grab sediment sampler		
c. Aseptic collection of samples		
i. Sampling of plants: Selection criteria: Type, physical condition, stage	of growth,	
plant part.	C ,	
ii. Sample treatment: surface sterilization, excision of desired plant c	omponent,	
extraction.		
2. Industrially and medically important biomolecules from plants and micro	organisms:	
Screening, detection, purification and characterization using analytical tools		24
a. Enzymes: extremozymes; food additives/ quality enhancers,	medicine,	
antioxidants and antitumor agents		
b. Pigments: food colorants, fabric dyes		
c. Biocontrol agents:herbicides, pesticides		
d. Nanoparticles : medicine, drug carriers.		
e. Biofuels: microbially produced; plant based		
f. Optical and electronic devices: archaeal metabolites (bacteriorhodops	sin and cell	
wall S-layer as membrane for ultrafiltration)		
g. Biopolymers – biodegradable plastics: PHAs, blended plastic poly	mers, EPS,	
biosurfactants and bioemulsifiers		
h. Plant growth promoters- gibberellins, auxins, cytokinins		
i. Pharmaceuticals: Antimicrobials, Antitumour agents, drug carriers.		
j. Nutraceuticals: PUFAs, β-carotenes, antioxidants		
k. Cosmeceuticals: humectants (polyols).		
I. Drugs from Sea		
3. Pollutants in the environment and their impact:		
a. Environment and pollutants		10
i. Classification of pollutants		
ii. Toxicity, synergistic or antagonistic action.		
iii. Eco-toxicology: concept of permissible limits. ED50 & LD50		
iv. Acute and chronic exposures: biochemical effects and genotoxic	city.	
b. Significant environmental pollutants: source. effect and impact		
i. Soil Xenobiotics		
ii. Agricultural chemicals		
iii. Pesticides		
iv. lead and other heavy metals		
v. Marine pollutants		
c. Monitoring of pollution		

		<u>sta. com.</u> <u>14.02.2</u>	<u>X AC-5</u> 2023
i. Using ir ii. Biosens <b>d. Significant envi</b> i. Dissolved ii. Biochem iii. Chemica iv. Environn v. Environn	dicator microorganisms ors: genetically modified organisms and enzymes <b>onmental monitoring parameters</b> oxygen cal Oxygen Demand Oxygen demand . eent protection regulations, impact assessment and s eental pollutants, improper waste disposal	standards.	
<ul> <li>4. Remediation of <ul> <li>a. Treatment of wa</li> <li>b. Introduction to w</li> <li>i. Wastewater/sew</li> <li>ii. Solid waste mana</li> <li>iii. Hospital waste r</li> <li>c. Biological system</li> <li>d. Microbial consor</li> <li>i. Enzymatic transf</li> <li>ii. Co-metabolism</li> <li>iii. Microbial adhes</li> <li>iv. Biofilms</li> <li>v. Production of ex</li> <li>e. Other pollutant r</li> <li>i. Sorption</li> <li>iii. Precipitation</li> <li>iv. Speciation conve</li> <li>f. Emerging eco-friet</li> <li>Technology</li> </ul></li></ul>	waste ste : Concepts of Reuse, Recycle, Recovery. "aste treatment ge treatment gement nanagement. s for remediation: plants, bacteria and fungi tia and related microbial processes ormations on tracellular polymers and emulsifiers. emoval techniques	stry and Green	10
<ul> <li><b>5. Biotechnological</b> <ul> <li>a. Bioremediation</li> <li>i. In situ and Ex-siti</li> <li>ii. Factors affecting</li> <li>iii. Methods in deteriv. Use of microbes</li> <li>v. Bioremediation</li> <li>vi. Evaluating Biore</li> <li>b. Biofilters</li> <li>c. Biotransformation</li> <li>d. Phytoremediation</li> </ul> </li> <li>Pedagogy Mainly I self-stude</li> </ul>	methods to control pollution u bioremediation process of bioremediation rmining Biodegradability (bacteria and fungi) bioremediation of common environmental pollutant mediation ion on ectures and tutorials. Seminars / term papers /assign ly or a combination of some of these can also be us d. Sessions should be interactive in nature to enable	ments / presen ed. ICT mode si	10 tations / hould be

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Text Books/ References / Readings	<ol> <li>Manahan S. E; Environmental Chemistry; Lewis Publishers, 2000/7 <sup>th</sup> edition.</li> <li>Salker A. V; Environmental Chemistry; NarosaNublishing; 2017, 1<sup>st</sup> edition</li> <li>De A. k; Environmental Chemistry; New Age International Publishers; 2005, 3 <sup>rd</sup> Ed</li> <li>Dara, S.S., Mishra D. D; A text book of Environmental Chemistry and Pollution Control; S. Chand Publishers; 2004.</li> <li>Enger E. D., Smith B. E.; Environmental Science: A study of Interrelationships; WCB Publication, McGraw-Hill Higher Education.; 2019, 15 <sup>th</sup> edition.</li> <li>Khopkar S. M., Environmental Pollution Analysis. New Age International Pvt. Ltd.; 2005, 1 <sup>st</sup> edition.</li> <li>Mitchell R., Cu J. D.; Environmental Microbiology; Wiley-Blackwell Publication; 2009.</li> <li>Moore J. W., Moore, E. A.; Environmental Chemistry; Academic Press; 1976, 1 <sup>st</sup> edition</li> <li>Moore J. W., Moore, E. A.; Environmental Chemistry; Academic Press; 1976, 1 <sup>st</sup> edition</li> <li>Jogdand, S.N., Gene Biotechnology. Himalaya publishing house, 2016, 4<sup>th</sup> edition.</li> <li>I. Jogdand, S.N., Advances in Biotechnology. Himalaya publishing house. 2007, 2<sup>nd</sup> edition.</li> <li>Ravi, I., Baunthiyal, M. &amp; Saxena, J. Advances in Biotechnology. Springer, 2014, 1<sup>st</sup> edition.</li> <li>Satyanarayana, U. and Chakrapani U. Biotechnology and Agriculture, Elsevier 2011, 1<sup>st</sup> edition</li> <li>Clark, D. &amp; Pazdernik, N. Biotechnology, Academic Press cell, 2015, 2<sup>nd</sup> edition.</li> <li>Bielecki, S., Tramper, J., &amp;Polak, J. Food Biotechnology. Churchill Livingstone, 2009.</li> <li>Pongracz, J. &amp; Keen, M., Medical Biotechnology. Churchill Livingstone, 2009.</li> </ol>
	<ol> <li>Bielecki, S., Tramper, J., &amp;Polak, J. Food Biotechnology. Elsevier, 2000.</li> <li>Pongracz, J. &amp; Keen, M., Medical Biotechnology. Churchill Livingstone, 2009</li> <li>Fletcher, G. L. &amp; Rise, M. L. Aquaculture Biotechnology. Wiley, 2011.</li> <li>Verma, A. &amp; Singh, A. Animal Biotechnology Models in Discovery and Translation. Academic press, 2020, 2<sup>nd</sup> edition.</li> </ol>
I	

### Programme: M.Sc. Part-II (Biochemistry) Course Code: CHBD-511 Title of the course: Discipline Specific Dissertation Number of Credits: 16Total Hours: 480

Effective from AY: 2023-24

Prerequisites f	for	or Students should have studied Biochemistry courses at MSc-I level.		
the course:				
Course		To develop the skills of preparing and conducting independent re	esearch.	
Objective:				
Course		Students will be able to understand and apply the tools and techniques of		
Outcome:		Biochemistry in conducting independent research.		
<i>Content</i> Hours				
As per OA-35 480			480	
Pedagogy	Dissertation carried out individually by each student throughout the academic year.			
Textbooks/	As required for the development of review and methodology			
References /				
Readings				

### Programme: M.Sc. Part-II (Chemistry/Biochemistry)

Course Code: CHIG-511

Title of the course: **Bioinorganic Chemistry** 

Number of Credits: **04** 

Total Hours: 60

Effective from AY: 2023-24

Prerequisites	Students have studied chemistry/biochemistry courses at M.Sc. Part-I			
for the course:				
Course Objective:	<ol> <li>To understand the role of inorganic elements especially metal biology.</li> <li>To introduce metallobiolecules, metalloproteins &amp; metalloenymes.</li> <li>To understand the role of small molecule model compounds.</li> <li>To introduce the concept of Biomimetic chemistry.</li> </ol>	ions in		
	1. Students will be in a position to clarify the significance of essential in biology.	elements		
Course	2. Students will be able to explain the role played by metal ion	s in vital		
Outcome:	processes like i) oxygen storage and transport and ii) electron transf	er.		
	3. Students will be able to explain basic concepts in Biomimetic chemistry.			
4. The students will be able use different techniques in Bioinorganic Chemistry				
	Content	Hours		
1. Essential eler	nents in biology	12		
Periodicity of	elements, distribution of elements in biosphere, bio-availability, bio-			
stability, bui	stability, building blocks of the biosphere; carbohydrates, nucleic acids and			
proteins, bio	proteins, biological importance of water, and brief review of the chemistry of			
biopolymers.	biopolymers. Metallobiomolecules: classification, metalloproteins (enzymes),			
metal activated proteins (enzymes), metal functions in metalloproteins, Principles				
of coordinati	on chemistry related to bioinorganic research, physical methods in			
bioinorganic	chemistry.			
2. Alkali and alk	aline earth metals in biology	12		
Introduction,	biological importance of the alkali and the alkaline earth cations,			

		Std. Com.	X AC-5
		<u>14.02.2</u>	023
Cation transport through membranes (ion pumps). Photosynthesis, Hill reaction,			
Chlorin mac	rocycle and chlorophyll, Absorption of light by chlorophy	ll, role of	
metals in ph	otosynthesis, in vitro photosynthesis.		
3. Non-redox r	netalloenzymes		12
Zinc metalle	penzymes like carboxypeptidase, carbonic anhydrase an	d alcohol	
dehydrogen	ase, Bio-functions of zinc enzymes, active site structure a	ind model	
complexes.			
4. Biochemistr	y of a few transition metals		12
Role of Fe,	Mo, Cu and Ni. Oxygen carriers and oxygen transport pro	teins, iron	
porphyrins	(Haemoglobin and myoglobin). Haemocyanins and Haem	erythrins,	
Synthetic m	odels for oxygen binding haemproteins. Cytochrome C,	catalase,	
peroxidase,	and superoxide dismutase, blue copper proteins, vi	tamin B <sub>12</sub>	
coenzymes,	nitrogen fixation and iron-sulfur proteins, biological nitroge	n fixation,	
nitrogenase	and dinitrogen complexes, iron-sulfur proteins, synthetic	analogues	
for Fe-S pro	teins, core extrusion reactions. Metal transport and storage	ge: A brief	
review of i	ron transport. transferrin, ferritin, hemosiderin, sideroph	ores, iron	
biomineraliza	ation		10
5. Biomimetic I	norganic Chemistry		12
Fundamenta	is of biomimetic chemistry, metal – oxygen intermediates, tech	niques used	
to probe the	active sites of oxygen carriers, redox chemistry of free molecula	ar dioxygen,	
spectroscopy	of Fe-U-Fe molety, geometry and electronic structure of a		
dioxygen, ot	ner ligands for biological oxygen carriers, reactions of me	et al-oxygen	
compounds, oxygenases, Cytochrome P-450, synthetic procedures of simple			
	stallization of carboxylic acids. Non-Heme and heme ligand	s s s s s s s s s s s s s s s s s s s	
Pedaaoay	Mainly lectures / tutorials / assignments /group dis	scussion /	self-study
, cuugogy	/presentations or a combination of some of these could	also be used	d to some
	extent.		
Textbooks /	1. S. L. Lippard & L. M. Berg. Principles of Bioinorgani	c chemistrv	Panima
Reference /	Publishing Corporation	<i>e enemier</i> ,	, i annia
Readinas	2. I. Britini, H. B. Grav. S. J. Lippard & J. S. Valentine, B.	ioioraanic cl	hemistrv.
	University Science books, Mill Valey, CA, 1994.	<u> </u>	, ,
	3. E. Fenton, <i>Biocoordination Chemistry</i> , Oxford Chemist	v Printers, 2	5 Oxford
	University Press, 1995		
	4. E. Conn, P.K. Stumpf, G. Bruening & R. H. Doi, Out	lines of Bioi	inorganic
	Chemistry, 5 <sup>th</sup> Ed.; Wiley Eastern, 1983.	-	-
	5. F.A. Cotton, G. Wilkinson, P.L. Gaus, Basic Inorgania	c Chemistry	, 3 <sup>rd</sup> Ed.
	(Chapter 31); WileyIndia, 2007.		
	6. M. Weller, T. Overton, J. Rourke & F. Armstrong Inor	ganic Chem	<i>istry,</i> Int.
	Ed. (Chapter 25); Oxford University Press, 2018.		
	7. P Atkins, T Overton, J Rourke, M Weller & F Armstro	ng, Shriver	& Atkins'
	Inorganic Chemistry, 5 <sup>th</sup> Ed. (Chapter 27); Oxford Unive	ersity Press,	2010.
	8. J. E. Huheey, E. A. Keiter, R. L. Keiter, Inorganic Che	emistry: Prin	nciples of
	Structure and Reactivity, 5 <sup>th</sup> Ed. (Chapter 19); Addison	Wesley Pub	lishing.
	9. R. W. Hay, Bioinorganic chemistry, Ellis Horwood Chiche	ster, 1984.	-
	10.M.N. Hughes, The Inorganic Chemistry of Biological pro	cesses, 2 <sup>nd</sup>	Ed.; Wiley
	(Interscience), 1984.		-

11.R. R. Crichton, <i>Biological Inorganic Chemistry</i> , Elsevier, 2012.
12.R. Breslow, Biomimetic Chemistry: Biology as an Inspiration, The Journal of
Biological Chemistry, vol. 284, no. 3, pp. 1337–1342, 2009.
13.C. Housecroft, A. G. Sharpe, <i>Inorganic Chemistry</i> , 4 <sup>th</sup> Ed; Pearson Publishing, 2012.

### Programme: M.Sc. Part-II (Analytical Chemistry)

# Course Code: CHAG- 511

Title of the course: Fundamentals of Crystallography

Number of Credits: <b>04</b>	Total Hours: 60	Effective from AY: 2023-24

Prerequisites	Students should have studied M.Sc. Part-I.		
for the course:			
	1. To introduce basic concepts of crystallography.		
Course	2. To impart knowledge of single crystal and powder X-ray diffraction		
Objective:	methods.		
	3. To analyse Materials and understand Structure.		
	1. Student will acquire fundamental concepts of crystallography.		
Course	2. Students will gain insights into single crystal and powder X-ray		
Course	diffraction methods.		
Outcome:	3. Students will be able to use X-ray diffraction methods for materials		
	characterization.		
	Content	Hours	
1. Basics of Crys	tallography	10	
a. The Crystal	line state, symmetry elements.		
b. Lattices, ur	nit cell, crystallographic directions, planes, point groups and		
symmetry	classes.		
c. The Laue classes, the seven crystal systems, Bravais lattices, space groups			
and International Tables.			
d. Description of crystal structures, unit cell projections and atomic			
coordinate	s, unit cell content.		
e. Ionic crysta	als, molecules and molecular crystals, protein crystals, physical		
properties	of crystals.		
2. Diffraction of	X-rays by Crystals:	10	
a. Interaction of X-rays with matter.			
b. Scattering of X-rays by an electron, atom, atomic scattering factor,			
temperature	temperature factor, scattering by molecule or unit cell.		
c. Diffraction by crystals, structure factor, Bragg's law, the reflection and the			
limiting spheres, symmetry in reciprocal space, systematic absences,			
diffraction intensities.			
d. Experiment	al methods in X-ray crystallography: X-ray sources,		
monochrom	atization, collimation, and focusing of X-rays.		
3. Single Crystal X-ray Diffraction:		10	
a. Crystals an	d their properties: crystallization, growing and choosing		
crystals, mi	croscopic observation		
b. Data collec	tion techniques for single crystals, diffractometer geometry,		
measurem	ent of the integrated intensities, data collection with area		

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detecto	rs,		
c. Data rec	luction: Lorentz correction, polarization correction, absorption		
correcti	ons, radiation damage corrections, relative scaling.		
d. Solutior	and refinement of crystal structures: Wilson plot, the heavy ato	m	
metho	d, Direct methods, phase determination procedures, figures of		
merit,			
e. Comple	ting and refining the structure: difference Fourier method, least-		
square	s method, absolute configuration.		
f. Introduc	tion to crystallographic software's (e.g. APEX 4, Olex2 etc) and		
IUCr val	idation of the data (CIF)		
4. Powder X-	ray Diffraction:		10
a. Origin o	f powder diffraction pattern, position, shape, and intensity of		
powder	diffraction peaks.		
b. Powder	diffractometry: beam conditioning, goniometer design,		
nonamb	ient powder diffractometry.		
c. Collectir	ng quality powder diffraction data: sample preparation, data		
acquisiti	on, quality of data, data processing.		
d. Determ	ination of unit cell: indexing methods.		
e. Introdu	ction to the Rietveld method.		
d. Introdu	ction to powder diffraction software's for indexing, unit cell		
refinem	ent (e.g. Winplotr, UnitCell).		
5. Applicatio	ns of Crystallography:		10
a. Chemist	ry and Materials science: understanding crystal structures of		
compou	nds, alloys, metals, polymers, phase transitions etc.		
b. Geology	v, mineralogy, gemology.		
c. Pharma	ceuticals: polymorphs, excipient analysis, active pharmaceutical		
ingredie	nts.		
d. Forensio	cs and environmental analysis.		
e. Nano m	aterials characterization.		
f. Biomole	cules: determination of structures of proteins, nucleic acids and		
other bio	ological macromolecules.		
g. Other d	iffraction techniques: neutron diffraction, thin film, microstructu	re	
properti	es, pair distribution function analysis, etc.		
6. Analysis of	f Materials and Structural Understanding:		10
a. Characte	erisation of Solids using diffraction techniques.		
b. Introduc	tion to databases: powder diffraction files, inorganic and organic	;	
crystal st	ructure database, protein data bank etc.		
c. Inspectio	on of crystals/powders with light microscope.		
d. Visualiza	ition of crystal structures using softwares (e.g. Diamond,		
VESTA).			
e. Beyond	ideal crystals: crystal twins, modulated structures, quasicrystals		
Pedagogy	Mainly lectures and tutorials. Seminars / term papers /assignm	ients / pres	sentations /
	self-study or a combination of some of these can also be used	I. ICT mode	e should be
	preterred. Sessions should be interactive in nature to enable pe	er group le	arning.
Textbooks/	1. M. Milanesio, G. Zanotti, G. Gilli, M. Catti, H. Monaco, G. Fe	erraris, G. A	Artioli, P.
References	Gilli, D. Viterbo, C. Giacovazzo - Fundamentals of Crystallog	raphy, 3 <sup>rd</sup>	Ed., Oxford
/ Readings	University Press, 2015.		

2. C. Hammond - The Basics of Crystallography and Diffraction (International Union
of Crystallography Texts on Crystallography) 4 <sup>th</sup> Ed., Oxford University Press,
2015.
3. R. West, Solid State Chemistry and Its Applications, 2 <sup>nd</sup> Ed.; Wiley, 2022.
4. F. Hoffmann, Introduction to Crystallography, 1 <sup>st</sup> Ed. Springer, 2020.
5. D. Sherwood, Crystals, X-rays and Proteins: Comprehensive Protein
Crystallography, 1st Ed. Oxford University Press, 2015.
6. A. Hofmann, S. Clokie, Wilson and Walkers Principles and Techniques of
Biochemistry and Molecular Biology, 8 <sup>th</sup> Ed.; Cambridge University Press, 2018.
7. V. Pecharsky and P. Zavalij, Fundamentals of Powder Diffraction and Structural
Characterization of Materials, 2 <sup>nd</sup> Ed.; Springer, 2009.
8. R. Young, The Rietveld Method, 1 <sup>st</sup> Ed., Oxford University Press, 1995
9. W. David, K. Shankland, L. McCusker, C. Bärlocher, Structure Determination from
<i>Powder Diffraction Data</i> , 1 <sup>st</sup> Ed., Oxford University Press, 2006.
10. B. He, Two-dimensional X-ray Diffraction, 1 <sup>st</sup> Ed., Wiley, 2009.
11. W. Massa, Crystal Structure Determination, 2 <sup>nd</sup> Ed., Springer, 2010.
12. R. Dinnebier, S. Billinge, <i>Powder Diffraction: Theory and Practice</i> , 1 <sup>st</sup> Ed., Royal
Society of Chemistry, 2008.

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# Programme: Ph.D. Biochemistry Paper-I Syllabus Title of the course: Research Methodology

Number of Credits: 04

Total Hours: 60 Effective from AY: 2022-23

Prerequisites f	tes for Provisional registration for PhD in Biochemistry		
the course:			
	1. To introduce research students to various aspects of	research	
	methodology		
	2. To provide an understanding of various databases used in biod	chemistry	
Course	3. To introduce the fundamental roles of computers in bio		
Objective:	research.		
	4. To provide understanding and importance of lab safety	to make	
	students aware of the statistical methods used in biochemical	research	
	5. To understand the usefulness of various techniques	in the	
	Characterization and purification of biochemical compounds		
	1. Students will be tamiliar with research methodology concepts	nrohlomo	
	in biochemictry	problems	
Course	2 Students will know in advance the safety precautions to be tal	on in the	
Outcome:	biochemistry lab		
outcome.	4 Students will apply statistical methods of data handling	in their	
	research. Students will gain fundamental knowle	edge of	
	characterization and purification techniques.		
	Content	Hours	
1. Introduction	to Research Methodology		
a) Rese	arch- meaning, objectives, motivation, types and methodology.		
b) Proc	ess- formulating the research problem; literature survey;		
deve	loping the hypothesis and the research design; sample design and		
colle	ction of the data; execution of the project; analysis of data; testing		
of h	ypothesis; generalizations and interpretation, and preparation of	10	
the	eport or presentation of the results & conclusions.		
c) Natu	re of scientific information- types of books, types of presentations		
pub	ished in journals, standard format for reporting original research,		
intro	duction to various scientific (chemistry) databases & sources from		
the	nternet.		
2. Role of Com	puters in Research and Biochemistry	10	
a) Role	of Computers in Research and chemistry		
b) Appl	Applications of computers in research.		
c) Appl	ications of computer in Chemistry - Need of computers in		
cnen	ramming languages: Solving a problem with computers		
prog	ithm flowchart and program: Use of software for data handling		
	ing graphs and drawing molecular structures visualisation of 2		
D da	ta: Software for literature survey, software for reference citing		

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d)	Optimisation techniques and applications in molecular geometric	ry		]			
	optimisation						
3. Saf	. Safety in biochemistry						
a)	Introduction to lab safety.						
b)	Handling of various chemicals, solvents and glassware						
c)	Fires and fighting with fires						
d)	Hazardous substances and their classification and handling						
e)	Biosafety procedures for handling of biological samples						
(†)	Preventing laboratory acquired infections			4			
4. Inti	oduction to Statistical Methods		10				
a)	Errors & their types, precision & accuracy in chemical analysis.						
b)	Application of statistical methods to data treatment & evaluation.						
c)	Confidence limits; hypothesis testing.						
(b)	F-tests, Chi square test, correlation and linear regression.						
e)	Use of software for statistical analysis.			_			
5. A. Int	roduction to Basic concepts		22				
	a) Sterilization and disinfection						
	b) Sample collection						
	c) Biological sample maintenance						
B. Pu	rification and characterization techniques in biochemistry research	n al					
a)	Purification techniques: agarose and polyacrylamide gi	ei					
	electrophoresis, gel filtration, ion exchange; affinity chromatography						
h)	Lechniques, HPLC, GC	ie					
(0	Methods of DNA analysis: Extraction and purification of nucle	IC					
	acids, PCR, DNA sequencing and cioning techniques						
()	Spectroscopic techniques: Oltraviolet-visible (OV-vis), basics of Infi	rared					
d)	(IR), NUClear magnetic resonance NVIR and Mass spectroscopy (Ms	5).					
() ()	Microscopic tochniques: Optical Microscopy (light dark phase con	tract					
(	Eluoroscopic techniques. Optical Microscopy (light, dark, phase con	trast,					
Padagogy	Loctures Discussions cominars internal exame assignments		tudy or a	-			
Peuuyoyy	combination of some of these can be used. ICT mode show	/sell-si	noferred				
	sessions should be interactive in nature to enable near group l	la de l	σ				
Text Boo	ks/ 1 Research Methodology: Methods and Techniques by C. R. k	(othari	<u>s.</u> New	-			
Reference	Δge International Pvt 1td 2004	Cothair	. INC W				
/ Reading	2 The ACS Style Guide: Effective Communication of	of Scie	ontific				
/ neuung	Information Edited by Anne M Coghill and Lorrin B Garso	$n \Delta m \epsilon$	erican				
	Chemical Society Washington DC and Oxford University	nress	New				
	York Oxford 2006	, press					
	2 Eurodamontals of Posoarch Mothodology and Statistics by V.K. Singh Now						
	3. Fundamentals of Research Methodology and Statistics by Y K Singh, Nev						
	A Prudent practices in the laboratory handling and my	anagar	nent of				
	chemical bazards The National Academies Press USA 201	11 11					
	5 Spectrometric Identification of Organic Compounds (5th Ed) by P M						
	Silverstein, G.C. Bassler and T.C. Morrill, John Wiley, Singan	ore	,				
	enversion, e e bassier and re mornin sonn whey, singape	5.0.		1			

I

<ol> <li>Principles of Biochemistry (7th Ed). D L Nelson, M M Cox, Lehninger. WH Freeman, 2017.</li> </ol>
<ol> <li>An introduction to practical biochemistry (3rd Ed). D T Plummer. Tata McGraw Hill, 2006</li> </ol>
8. Principles and Techniques of Practical Biochemistry (7 th Ed) Wilson K, Walker J. Cambridge University Press, 2010
9. Analytical Chemistry (7th Ed) Christian G D, Dasgupta P K, Schug K A. John Wiley & Sons, 2013
10. Analytical Biochemistry, Homes D J, Peck H. Pearson education Limited, 1998.
<ol> <li>Principles of Instrumental Analysis, (7th Ed) Skoog D A, Holler F J, Crouch S R, Cengage Learning. 2016.</li> </ol>
<ol> <li>Molecular cloning: a laboratory manual (2nd edition) J Sambrook, E F Fritsch, T Maniatis, Cold Spring Harbor Laboratory Press, New York, 1989.</li> </ol>
<ol> <li>Modern Quantum Chemistry Introduction to Advanced Electronic Structure Theory, Attila Szabo, Neil S. Ostlund, Dover Publications, Inc. Mineola, New York 1989</li> </ol>
14. Introduction to Organic Spectroscopy Fifth Edition, D. Pavia, G.
15. Computer Programming in Fortran 90 And 95, V. Rajaraman, PHI Learning Pvt. Ltd., 2013

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# SWAYAM COURSES FOR BIOCHEMISTRY STUDENTS

Courses of UG/PG level which could be opted over and above existing PG courses from the curriculum:

Sr.	Name of the	Credits	Level as per	Intended audience	Whether
NO.	course		SWAYAM		Included in
1.	Animal Physiology	3	Undergraduate /Postgraduate	UG and PG students pursuing biology, biotechnology, zoology and bio-engineering	Yes
2.	Biomedical Nanotechnology	1	Undergraduate /Postgraduate	UG/PG students of Biotechnology/ Nanotechnology	No
3.	Environmental Biotechnology	3	Undergraduate /Postgraduate	B Sc. (Biosciences, Microbiology, Biotechnology, Botany, Any other branches of Life Sciences); M Sc. (Biotechnology / Life Sciences/ Environmental Sciences); B Tech (Biotechnology/Bioengineering )	Yes (partly)
4.	Fundamentals of Protein Chemistry	3	Undergraduate /Postgraduate	Undergraduate and Graduate students	Yes
5.	Genetic Engineering	3	Undergraduate /Postgraduate	-	Yes (partly)
6.	Genome Editing and Engineering	3	Undergraduate /Postgraduate	UG/PG/PhD/Scientist in industry	No
7.	Introduction to Proteogenomics	2	Undergraduate /Postgraduate	Biotechnology or Engineering background students having interest in latest technologies, (BE/B.Tech) Biotechnology. Students with science or engineering background but course is open to all.	Yes
8.	Introduction to Proteomics	2	Undergraduate /Postgraduate	It would be applied to B.Sc., M.Sc. and MS.	Yes
9.	Human Molecular Genetics	1	Undergraduate /Postgraduate	Students interested in pursuing research in human molecular genetics. Medical students and practicing clinicians interested in understanding the principles	Yes (partly)

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				and complexities of hur genetics. Students interested in care in genetic counselling and I diagnostics. Scientists working in pu health service, counsel centres, and diagno laboratories.	man eers DNA ublic Iling ostic
10	Cell Culture Technologies	2	Postgraduate	UG and PG students pursu biology, biotechnology, zool and bio-engineering.	uing No logy
11	Tissue Engineering	2	Postgraduate	Masters/Doctoral students faculty interested in teach Tissue Engineering	and No hing
12	Bioreactors	1	Undergraduate	-	Yes (partly)
13	Functional Genomics	1	Undergraduate	<ul> <li>Students interested pursuing research in genor and biotechnology field.</li> <li>Medical students practicing clinicians interest in genomics and molec medicine.</li> <li>Students interested in card in biotechnology biopharma fields.</li> <li>Technical staff /scient working in biotechnology biopharma and molec diagnostic laboratories/industry</li> </ul>	in No mics and sted cular eers and tists ogy, cular
14	Advanced Microbiology	4	Undergraduate	-	No

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### Courses of PG level which could be opted in lieu of existing PG optional courses of Biochemistry Part-II from the curriculum:

Sr. No.	Name of the course	Credits	Level as per SWAYAM	Intended audience	Whether included
					in NET syllabus
1.	Cell Culture Technologies	2	Postgraduate	UG and PG students pursuing biology, biotechnology, zoology and bio-engineering.	No
2.	Tissue Engineering	2	Postgraduate	Masters/Doctoral students and faculty interested in teaching Tissue Engineering	No

### Syllabus for online courses related to Biochemistry field at UG/PG level

#### <u>1-Animal Physiology</u> Credit=3

Level : Undergraduate/Postgraduate INTENDED AUDIENCE : UG and PG students pursuing biology, biotechnology, zoology and bioengineering PREREQUISITES : Biology at standard 10th (Secondary school examination)

#### Course layout

Week 1: Introduction Week 2: Skeletal system Week 3: Muscles Week 4: Neural system Week 5: Neural system Week 6: Neural system Week 7: Neural system Week 8: Neural system and Special senses Week 9: Neural system and Special senses Week 10: Cardiovascular system Week 11: Respiratory and Blood Week 12: Endocrine, Digestive, Blood, Kidney, and Reproductive system Books and references 1-Guyton and Hall Textbook of Medical Physiology 2-Ganong'S Rev of Med Physiology 3-Fundamentals of anatomy and physiology by Martini

### 2-Biomedical Nanotechnology

#### <u>Credit=1</u>

#### Level : Undergraduate/Postgraduate

INTENDED AUDIENCE : UG/PG students of Biotechnology/ Nanotechnology.

#### <u>Course layout</u>

Week 1: Introduction to nano, Nano-biomimicry, Synthesis of nanomaterials by physical and chemical methods, Synthesis of nanomaterials by biological methods, Characterisation of nanomaterials.

Week 2: DNA nanotechnology, Protein & glyco nanotechnology, Lipid nanotechnology, Bionanomachines, Carbon nanotube and its bio-applications.

Week 3: Nanomaterials for cancer diagnosis, Nanomaterials for cancer therapy, Nanotechnology in tissue engineering, Nano artificial cells, Nanotechnology in organ printing.

Week 4: Nanotechnology in point-of-care diagnostics, Nanopharmacology & drug targeting, Cellular uptake mechanisms of nanomaterials, In vitro methods to study antibacterial and anticancer properties of nanomaterials, Nanotoxicology.

### **Books and references**

1-Malsch NH; Biomedical Nanotechnology; CRC Press; 2005; 1st edition.

2-Mirkin CA, Niemeyer CM; Nanobiotechnology II: More Concepts and Applications; Wiley-VCH; 2007; 1st edition.

3-Kumar CSSR, Hormes J, Leuschner C; Nanofabrication Towards Biomedical Applications: Techniques, Tools, Applications, and Impact; Wiley-VCH Verlag GmbH & Co; 2005; 1st edition. 4-Lamprecht A; Nanotherapeutics: Drug delivery concepts in nanoscience; Pan Stanford; 2009; 1st edition.

5-Jain KK; The Handbook of Nanomedicine; Humana press; 2008; 1st edition.

### <u>3-Environmental Biotechnology</u> <u>Credit=3</u>

### Level : Undergraduate/Postgraduate

**INTENDED AUDIENCE**: B Sc. (Biosciences, Microbiology, Biotechnology, Botany, Any other branches of Life Sciences); M Sc. (Biotechnology / Life Sciences/ Environmental Sciences); B Tech (Biotechnology/Bioengineering)

Week 1: Environmental Biotechnology and Sustainability. Scope and applications of the subject. Basics of ecosystem structure and function

Week 2: Microbial Ecology and Environmental Biotechnology: Concepts and importance of microbial ecology in Environmental Biotechnology

Week 3: Microbiology of Environmental Engineering System: Microbial diversity, growth and decay. Stoichiometry of microbial energetics and kinetics.

Week 4: Resource Exploitation by Microorganisms: Functions of various microbial groups relevant to environmental systems, including waste treatment and resource recovery, implications in biogeochemistry.

Week 5: Methods in Microbial Ecology with relevant to Environmental Biotechnology: Culture dependent and - independent analyses of microbial communities; PCR based methods, Microarray, Environmental genomics

Week 6: Microbial Principles of Biodegradation, Biodetoxification and other processes relevant for Environmental Applications: Microbial engines, (metabolism), Requirements for biodegradation, acclimation, Common biotransformation mechanisms; Effect of organic contaminant structure on biodegradability; Cooperation between different microbial species for enhanced biodegradation; Applying biodegradation kinetics to fate and transport modelling

Week 7: Bioremediation Technologies: Concepts, methods and applications of natural attenuation and engineered bioremediation (e.g bioaugmentation and biostimulation)

Week 8: Microbial Interactions with Heavy Metals and Metalloids: Bioremediation, Biohydrometallurgy and other aspects of Environmental Biotechnology

Week 9: Aerobic and Anaerobic Degradation of Aliphatic and Aromatic Compounds. Microbial interaction with plastics, antibiotics and others emerging pollutants.

Week 10: Microbially Enhanced Phosphorus and Nitrogen Removal

Week 11: Microbially Enhanced Oil Recovery; Microbial role in Carbon Storage and Capture (sequestration, conversion to useful biopolymers, etc.).

Week 12: Case studies : Bioremediation, Carbon Storage and Capture, Bioenergy.

#### **Books and references**

1. Rittman B. E, McCarty P. L ; Environmental Biotechnology Principles and Applications; McGraw-Hill Higher education.

2. Jördening H J, Wint J; Environmental Biotechnology; WILEY-VCH Verlag Gmbh & Co.

3. Alvarage P J J, Illman W A; Bioremediation and Natural Attenuation; Wiley Interscience.

4. L K Wang et al; Environmental Biotechnology; Handbook of Environmental Engineering; Humana Press; Vol 10.

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### <u>4-Fundamentals of Protein Chemistry</u> <u>Credit=3</u>

Level : Undergraduate/Postgraduate

INTENDED AUDIENCE : Undergraduate and Graduate students

### <u>Course layout</u>

- Week 1: Amino acids and the Peptide Bond
- Week 2: Protein Architecture
- Week 3: Protein Folding and Denaturation
- Week 4: Protein isolation and characterization
- Week 5: Protein-ligand interactions
- Week 6: Membrane Proteins and Transport
- Week 7: Enzymes and Enzyme mechanisms
- Week 8: Enzyme kinetics and Enzyme inhibition
- Week 9: Metalloproteins and Motor Proteins
- Week 10: Protein-Protein interactions
- Week 11: Protein Structure analysis
- Week 12: Special Topics in Protein Chemistry

### **Books and references**

- 1- Branden C, Tooze J; Introduction to Protein Structure; 2nd Edition.
- 2-Creighton T E; Proteins: Structures and Molecular Properties.
- 3- Holde K.E.van , Johnson C , Ho S P; Physical Biochemistry.
- 4- Kyte J; Structure in Protein Chemistry.

5- Lehninger L Albert, Nelson L D , Cox M M; Principles of Biochemistry; 6th Edition

6- Kessel A, Tal Ben N; Introduction to Proteins: Structure, Function, and Motion; 2018;2nd Edition .

7- Stryer L, Berg J , Tymoczko J , Gatto G; Biochemistry; 2019;9th Edition.

# 5- Genetic Engineering

# <u>Credit=3</u>

# Level: Undergraduate/Postgraduate

# <u>Course layout</u>

Week 1: Introduction and Basics of Biological System.
Week 2: Basics of Biological System
Week 3: Basics of Cloning (Part I)
Week 4: Basics of Cloning (Part II)
Week 5: Recombinant DNA Technology (Part I)
Week 6: Recombinant DNA Technology (Part II)
Week 7: Product Recovery and Purification (Part I)
Week 8: Product Recovery and Purification (Part II)
Week 9: Characterization of Isolated Products (Part I)
Week 10: Characterization of Isolated Products (Part II)
Week 11: Biotechnology in Social Welfare
Week 12: Summary & Conclusions

# **Books and references**

Berg M J, Tymoczko L J,Stryer L; Biochemistry; W. H. Freeman and Company(New York);2006.
 Nelson L D, Cox M. M; Lehninger Principles of Biochemistry; Macmillan Worth; 2007;5th Edition.

3. Alberts B, Johnson A, Lewis J, Raff M, Roberts K, Walters P; Molecular Biology of Cell; Garland Publishing; 2007;5th Edition.

4. Prescott M L, Harley P J, Klein A D; Microbiology; McGraw Hill; 2005; 6th Edition.

5. Primrose SB, Twyman RM; Principles of Gene Manipulation; Blackwell Science; 2006.

6. Lewin B; Genes IX; International Edition; Pearson education; 2008.

# <u>6- Genome Editing and Engineering</u> <u>Credit=3</u>

Level : Undergraduate/Postgraduate

# INTENDED AUDIENCE: UG/PG/PhD/Scientist in industry

# <u>Course layout</u>

Week 1: Introduction to genetics and genetic engineering

Week 2: Breakage and Repair Of Genomic DNA

Week 3: Recombination

Week 4: Targeted genetic modification

Week 5: Zinc Finger Nuclease (ZFN) Technology

Week 6: Transcription activator-like effector nuclease (TALEN) Technology

Week 7: Clustered regularly interspaced short palindromic repeats (CRISPR)/Cas9 technology

Week 8: Applications of genome editing in treating human diseases

Week 9: Genome engineered Disease modeling

Week 10: Engineered immune cells for cancer therapy

Week 11: Personalized therapy; Challenges: safety and specificity

Week 12: Ethical concerns: Germ line gene editing

## **Books and references**

1-Harber , J. E., Genome Stability: DNA Repair and Recombination , Garland Science, 2013.

2-Yamamoto, T., Targeted Genome Editing Using Site-Specific Nucleases, Springer, 2015.

3-Zlatanova, J. and Holde, K. van, Molecular Biology: Structure and Dynamics of Genomes and Proteomes. Garland Science, 2015.

4-Yamamoto, T.(Ed.), Targeted Genome Editing Using Site-Specific Nucleases: ZFNs, TALENs, and the CRISPR/Cas9 System, Springer 2015.

5-Barrangou, R. and Oost, J. van der, CRISPR-Cas Systems: RNA-mediated Adaptive Immunity in Bacteria and Archaea, Springer, 2013.

6-Addgene, CRISPR 101:A Desktop Resource , January 2016

7-Alberts , B. , Johnson , A., Lewis , J., Morgan, D., Raff, M., Roberts, K.and Walter, P., Molecular Biology of the Cell, 6th Edn., Garland Science, 2014.

# 7-Introduction to Proteogenomics

### <u>Credit=2</u>

# Level : Undergraduate/Postgraduate

INTENDED AUDIENCE : Biotechnology or Engineering background students having interest in latest technologies, (BE/B.Tech) Biotechnology. Students with science or engineering background but course is open to all.

# <u>Course layout</u>

Week 1: Proteogenomics overview- Part I, Proteogenomics overview- Part II, Introduction to Genomics- Part I : Gene sequencing and mutations Introduction to Genomics-Part II : Sequence alignment, Introduction to Genomics-Part III :Transcriptome, SL1: Advancement in Cancer Genomics, SL2: Advancement in Cancer Genomics

Week 2: Introduction to Genomics IV : Epigenome, Introduction to Genomics : cBioPortal, Genotype, Gene expression & Phenotype - Part I, Genotype, Gene expression & Phenotype- Part II, An overview of NGS technology, SH1: NGS-Sequencing by synthesis, SH2: NGS- Sequencing by synthesis

Week 3: Introduction to Proteomics, Proteomics: Sample Prep & Protein Quantification, Proteomics: Sample Prep & Protein Quantification (Hands-on), Introduction to MS-based Proteomics- Part I, Introduction to MS-based Proteomics- Part II, SL 3: Applications of NGS – Ion Torrent, SL4: Applications of NGS – Ion Torrent

Week 4: Introduction to MS-based Proteomics- Part I (Hands-on), Introduction to MS-based Proteomics- Part II (Hands-on), Data analysis: Normalization, Data analysis: Batch Correction and Missing values, Data analysis: Statistical Tests, SH3: NGS- Ion Torrent, SH4: NGS- Ion Torrent

Week 5: Machine learning and Clustering, Hypothesis testing, ProTIGY- Part I, ProTIGY- Part II, Proteogenomics approach to unravel proteoforms, SL5: Genomic Analysis using Droplet PCR, SL6: Genomic Analysis using Droplet PCR

Week 6: Workflow to Automated Data Processing, Introduction to Fire Cloud, Fire Cloud and Data Model, Bioinformatics solutions for 'Big Data' Analysis- Part I, Bioinformatics solutions for 'Big Data' Analysis-Part II, SH5: Genomic Analysis using Droplet PCR, SH6: Genomic Analysis using Droplet PCR

Week 7: Data Science infrastructure management- Part I, Data Science infrastructure management- Part II, Data Science infrastructure management- Part III, DIA-SWATH Atlas-Part I, DIA-SWATH Atlas-Part II, SL7: Introduction to Targeted Proteomics, SH7: Data Analysis using Skyline

Week 8: Human Protein Atlas-Part I Clinical, Human Protein Atlas-Part II, Affinity based proteomics & HPA, Clinical Considerations for OMICS-Part I, Considerations for OMICS- Part II, SL8: Proteomics: PTMs, SL9: Clinical Proteomics

Week 9: ntroduction to Proteogenomics-Part I, Introduction to Proteogenomics-Part II, Sequence centric proteogenomics, Gene Variant Analysis, Proteomics in Clinical studies, SH8: ProTIGY

Week 10: Supervised Machine learning- Predictive Analysis Part I, Supervised Machine learning-Predictive Analysis Part II, Supervised Machine learning- Marker Selection, Gene Set Analysis using WebGestalt- Part I, Gene Set Analysis using WebGestalt- Part II, SH9: Supervised Machine Learning

Week 11: Biological Network Analysis- Part I, Biological Network Analysis- Part II, Mutation and Signaling - Part I, Mutation and Signaling- Part II, Pathway Enrichment, SH10: Pathway Enrichment and Network Analysis

Week 12: Gene Set Enrichment Analysis (GSEA), Pathway enrichment: GSEA, Linked Omics, Linked Omics (Hands-on), Proteogenomics Conclusions, SL10: Topics in Proteogenomics- Malaria and Cancer case study

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#### **Books and references**

Link A J, LaBaer J: Proteomics: A Cold Spring Harbor Laboratory Course Manual; Cold Spring Harbor Laboratory Press; 2009.

#### <u>8-Introduction to Proteomics</u> Credit=2

Level : Undergraduate/Postgraduate INTENDED AUDIENCE : It would be applied to B.Sc., M.Sc. and MS.

#### <u>Course layout</u>

Week 1 : Basics of Proteins and Proteomics

Lecture 1 : Introduction to amino acids

Lecture 2 : Introduction to Proteins

Lecture 3 : Protein folding & misfolding

Lecture 4 : Introduction to Proteomics

Lecture 5 : Lab session – Protein-protein interaction using label-free biosensors

Week 2: Gel-based proteomics

Lecture 6: Sample preparation and pre-analytical factors

Lecture 7 : Sample preparation: Pre-analytical factors (contd.)

Lecture 8 : Sample preparation: Protein extraction and quantification

Lecture 9 : One-dimensional electrophoresis

Lecture 10 : Introduction to 2-DE

- Week 3 : Two-dimensional gel electrophoresis (2-DE)
- Lecture 11 : 2-DE: Second dimension, staining & destaining

Lecture 12 : 2-DE: Gel analysis

Lecture 13 : 2-DE Applications

Lecture 14 : 2-DE Applications (contd.) & Challenges

Lecture 15: Lab session - Protein/peptide pre-fractionation using OFFGEL FRACTIONATOR & data analysis

- Week 4 : Difference in gel electrophoresis (DIGE) & Systems Biology
- Lecture 16 : 2D-DIGE: Basics
- Lecture 17 : 2D-DIGE: Data analysis
- Lecture 18 : 2D-DIGE: Applications
- Lecture 19 : Systems biology and proteomics I
- Lecture 20 : Systems biology and proteomics II

Week 5: Basics of mass spectrometry

- Lecture 21 : Fundamentals of mass spectrometry
- Lecture 22 : Chromatography technologies
- Lecture 23 : Liquid chromatography

Lecture 24 : Mass spectrometry: Ionization sources

Lecture 25 : Mass spectrometry: Mass analyzers

Week 6 : Basics of mass spectrometry and sample preparation

Lecture 26 : MALDI sample preparation and analysis

Lecture 27: Hybrid mass spectrometry configurations

Lecture 28 : Lab session - Demonstration of Q-TOF MS technology

- Lecture 29 : In-gel & in-solution digestion
- Lecture 30 : Lab session Sample preparation: tissue sample preservation technology
- Week 7 : Quantitative proteomics
- Lecture 31 : Introduction to quantitative proteomics
- Lecture 32 : SILAC: In vivo labelling
- Lecture 33 : iTRAQ: In vitro labelling
- Lecture 34 : TMT: In vitro labelling

Lecture 35 : Quantitative proteomics data analysis

### Week 8: Advancement in Proteomics

Lecture 36 : Proteomics applications

Lecture 37 : Challenges in proteomics

Lecture 38 : OMICS and translational research

Lecture 39 : Lab session – Targeted proteomics using triple quadrupole mass spectrometry

Lecture 40 : Lab session - Targeted proteomics: multiple reaction monitoring

### 9. Human Molecular Genetics

### <u> Credits : 1</u>

### Level: Undergraduate/Postgraduate

INTENDED AUDIENCE : Students interested in pursuing research in human molecular genetics.

Medical students and practicing clinicians interested in understanding the principles and complexities of human genetics.

Students interested in careers in genetic counselling and DNA diagnostics.

Scientists working in public health service, counselling centres, and diagnostic laboratories.

PRE- REQUISITES: Participants are expected to have at least class 12 level understanding in genetics.

### Course layout

Week 1: Fundamentals of central dogma (DNA, RNA and proteins; mutations), Chromosome structure and function (organization; structure-function relationship; chromosome abnormalities).

Week 2: Genes in pedigree (Mendelian pedigree patterns, complications to pedigree patterns), DNA cloning and hybridization techniques (vector based cloning; nuclei acid hybridizations; PCR-based DNA analyses)

Week 3: Mutation and instability of human DNA (mutation and polymorphism; pathogenic mutations, repeat expansions), Molecular pathology (types of mutations; animal models for human disease)

Week 4: Identifying human disease genes (functional cloning versus positional cloning; mutation screening), Complex diseases; The Human Genome and HapMap projects

#### **Books and references**

Strachan T, Read A; Human Molecular Genetics; Science/Taylor & Francis Group, 2011; 4 th Edition

### <u>10-Cell Culture Technologies</u> <u>Credit=2</u>

### Level : Postgraduate

INTENDED AUDIENCE : UG and PG students pursuing biology, biotechnology, zoology and bioengineering.

### <u>Course layout</u>

Week 1: Introduction & biology of cultured cells

Week 2: Equipments, aseptic techniques, safety protocols

Week 3: Culture vessels & media development

Week 4: Serum-free medium development & sterilization

Week 5: Primary culture, secondary culture, cloning & selection

Week 6: Cell separation, characterization, differentiation & transformation

Week 7: Contamination, cryo-preservation & cyto-toxicity

Week 8: Organo-typic culture & specialized cell culture techniques

### **Books and references**

1. Freshney I; Culture of Animal Cells; Wiley; 2016; 7th edition.

2. Bruserud Ø, Englezou A; Cell culture technology: Recent advances and future prospects Euroscicon Meeting Reports; Honnao publishing; 2012; 1st edition.

3. Bückmann A, Carrea G, et al; Vertebrate Cell Culture II and Enzyme Technology: Volume 39 Advances in Biochemical Engineering/ Biotechnology; Springer; 2013.

4. Butler M; Animal Cell Culture and Technology: The Basics; Garland Science, 2003, 2nd edition.5. Skloot R; The Immortal Life of Henrietta Lacks; Crown; 2011.

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# <u> 11- Tissue Engineering</u>

<u>Credit=2</u>

### Level : Postgraduate

**INTENDED AUDIENCE** : Masters/Doctoral students and faculty interested in teaching Tissue Engineering

### <u>Course layout</u>

Week 1: Introduction to tissue engineering

Week 2: Scaffolds: extracellular matrix, natural and synthetic polymers

Week 3: Hydrogels, bioceramics, scaffold fabrication

Week 4: Material characterization

Week 5: Cell source, isolation, growth, differentiation

Week 6: Cell adhesion, migration, signaling, bioreactors and challenges in tissue engineering

Week 7: Host integration, bioethics, Applications: Skin tissue engineering

Week 8: Applications: Bone tissue engineering, Vascular tissue engineering, and Corneal tissue engineering

### **Books and references**

1-Palsson O B, Bhatia S N; Tissue Engineering; Pearson; 2004

2-Brown R A; Extreme Tissue Engineering: Concepts and Strategies for Tissue Fabrication; Wiley Blackwell; 2013

3-Saltzman W M; Tissue Engineering: Engineering Principles for the Design of Replacement Organs and Tissues; Oxford University Press; 2004

4-Fisher J P, Mikos A G, Bronzino J D; Tissue Engineering; CRC Press; 2006

5-Lanza R, Langer R, Vacanti J; Principles of Tissue Engineering; Elsevier Academic Press; 2007; 3rd Edition

#### <u>12-Bioreactors</u> Credit=1

#### Level : Undergraduate/postgraduate

### <u>Course layout</u>

Week 1: Introduction

**Week 2:** Two important outcomes of a bioprocess: biomass (cells) and bio-products **Week 3:** Common bioreactor operation modes, factors that affect bioreactor performance **Week 4:** The cell view of a bioreactor

### **Books and references**

1. Shuler, M.L. and Kargi, F. 2002. Bioprocess Engineering: Basic Concepts, Prentice Hall, 2. Englewood Cliffs, NJ J. Bailey and D. Ollis, Biochemical Engineering Fundamentals; McGraw Hill, 1986.

In addition, the students would be directed to specific sources during the course – they will become available during the course on the course page.

# 13-Functional Genomics

### <u>Credit=1</u>

#### Level : Undergraduate/postgraduate INTENDED AUDIENCE :

- Students interested in pursuing research in genomics and biotechnology field.
- Medical students and practicing clinicians interested in genomics and molecular medicine.
- Students interested in careers in biotechnology and biopharma fields.
- Technical staff /scientists working in biotechnology, biopharma and molecular diagnostic laboratories/industry

### <u>COURSE LAYOUT</u>

Week 1: [2.5 hrs; 4 lectures]

Introduction to Functional Genomics:

Pre- and post-genomic era; major advancements in genomic approaches; epigenetics and metagenomics; forward versus reverse genetics

Week 2: [2.5 hrs; 4 lectures] Genome Analyses - Part 1 Genome editing approaches and their applications; gene expression analyses and applications

Week 3: [3 hrs: 4 lectures and 2 tutorial sessions] Genome Analyses - Part 2 Methods for DNA/RNA sequencing, sequence analysis and their applications

Week 4: [2.5 hrs: 3 lectures and 2 laboratory sessions] Comparative Genomics Genomic insight into evolution; power of comparative genomic analysis

### **Books and references**

1. Mostly publically available literature. Will be shared with the participants during the launch of the course.

<u>14- Advanced Microbiology</u> <u>Credit=4</u>

### Level : Undergraduate/postgraduate

Course layout Week1 Module 1: Morphology, Disease and Holmes Classification. Module 2: Classification of viruses-I Module 3: Classification of viruses II

Week 2

Module 4: Classification of fungi: Classical Concept -I Module 5: Classification of fungi: Classical Concept -II Module 6: Modern Classification of fungi: Modern Concept - I Week 3 Module 7:Classification of fungi: Modern Concept - II Module 8:Classification of fungi: Modern Concept - III Module 9:Classification of yeast-I

Week 4 Module 10:Classification of yeast-II Module 11: Application of microbes Module 12:Antibiotics: The secondary Metabolites

Week 5 Module 13:Role of Staphylococcus in infection Module 14:Nitrogen fixation & Role of Nif and Nod genes. Module15:Microbiology & Human Health

Week 6 Module 16: Bacterial Diseases Module 17.The pathogenesis of Vibrio cholerae Module 18: Introduction to Salmonella

Week 7

Module 19: Preparation of Culture Media: Liquid Medium Module 20: Preparation of Culture Media: Solid Medium Module 21: Sterilization Techniques (Practical)

Week8

Module 22: Isolation of single colony on solid media (Practical) Module 23: Enumeration of bacterial numbers by serial dilution & plating (Practical) Module 24: Measurement of fungal bio-mass

Week 9 Module 25: Animal Handling (Practical) Module 26: Staphylococcus aureus Module27: Isolation of protozoa from soil

Week 10

Module 28: Antibiotic sensitivity assay: natural products Module 29: Preparation of cultu re media for pathogenic bacteria – Part 1 Module 30: Preparation of culture media for pathogenic bacteria – Part 2

Week 11 Module 31: Albert and Giemsa Staining Module 32: Determination of Growth Phase of E. coli by Measurement of OD & cfu

Week 12 Module 33: Preparation of Buffer Solution

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