

4-Year Bachelor of Pharmacy (B.Pharm) Curriculum and Syllabus <u>Course Structure</u> Third Semester

Course Code	Name of the Course	No.of hours	Tutorial	Credit points
TIU-UBP-301T	Pharmaceutical Organic Chemistry II - Theory	3	1	4
TIU-UBP-302T	Physical Pharmaceutics I-Theory	3	1	4
TIU-UBP-303T	Pharmaceutical Microbiology -Theory	3	1	4
TIU-UBP-304T	Pharmaceutical Engineering-Theory	3	1	4
TIU-UBP-305P	Pharmaceutical Organic Chemistry II- Practical	4	-	2
TIU-UBP-306P	Physical Pharmaceutics I -Practical	4	-	2
TIU-UBP-307P	Pharmaceutical Microbiology– Practical	4	-	2
TIU-UBP-308P	Pharmaceutical Engineering– Practical	4	-	2
	Total	28	4	24



B.PHARM SYLLABUS

SEMESTER – III

Pharmaceutical Organic Chemistry II – Theory Contact hours: 45 hr

Course Objectives

1. To understand the basic structure of Benzene, Polyneuclear hydrocarbons, Cycloalkanes, aromatic amines, acids, fats and oils.

2. To know the mechanism of action of the different chemical reaction of organic compounds, their resonance hybridization and their orientation reactions

3. To gain knowledge of the basic preparation, physicochemical properties, use and preparation of different organic compounds.

Course Outcomes

Upon completion of the course, the student shall be able

CO1. Write the structure, name and the type of isomerism of the organic compound

CO2. Write the reaction, name the reaction and orientation of reactions

CO3. Understand the resonance hybridization of different organic compounds.

CO4. Account for reactivity/stability of compounds,

CO5. Preparation chemical reaction and use of different organic compounds

Course Content

45 Hours

10 Hours

UNIT I

Benzene and its derivatives•

A. Analytical, synthetic and other evidences in the derivation of structure of benzene, Orbital picture, resonance in benzene, aromatic characters, Huckel's rule

B. Reactions of benzene - nitration, sulphonation, halogenationreactivity, Friedelcrafts alkylation- reactivity, limitations, Friedelcrafts acylation.

C. Substituents, effect of substituents on reactivity and orientation of mono substituted benzene compounds towards electrophilic substitution reaction D. Structure and uses of DDT, Saccharin, BHC and Chloramine

UNIT II

• **Phenols*** - Acidity of phenols, effect of substituents on acidity, qualitative tests, Structure and uses of phenol, cresols, resorcinol, naphthols

•Aromatic Amines* -Basicity of amines, effect of substituents onbasicity, and synthetic uses of aryl diazonium salts

•Aromatic Acids* –Acidity, effect of substituents on acidity and important reactions of benzoic acid.

10 Hours



UNIT III

• Fats and Oils

a. Fatty acids – reactions.

b. Hydrolysis, Hydrogenation, Saponification and Rancidity of oils, Drying oils.

c. Analytical constants – Acid value, Saponification value, Ester value, Iodine value, Acetyl value, Reichert Meissl (RM) value – significance and principle involved in their determination.

UNIT IV

• Polynuclear hydrocarbons:

a. Synthesis, reactions

b. Structure and medicinal uses of Naphthalene, Phenanthrene, Anthracene,

Diphenylmethane, Triphenylmethane and their derivatives

UNIT V

•Cyclo alkanes*

Stabilities – Bae yer's strain theory, limitation of Baeyer's strain theory Coulson and Moffitt's modification, Sachse Mohr's theory (Theory of strainless rings), reactions of cyclopropane and cyclobutane only

Reference Books(Latest Editions)

- 1. Organic Chemistry by Morrison and Boyd
- 2. Organic Chemistry by I.L. Finar , Volume-I
- 3. Textbook of Organic Chemistry by B.S. Bahl& Arun Bahl.
- 4. Organic Chemistry by P.L.Soni
- 5. Practical Organic Chemistry by Mann and Saunders.
- 6. Vogel's text book of Practical Organic Chemistry
- 7. Advanced Practical organic chemistry by N.K.Vishnoi.
- 8. Introduction to Organic Laboratory techniques by Pavia, Lampman and Kriz.

Relationship between the Course Outcomes (COs) and Program Outcomes (POs)

	Mapping between COs and POs									
	Course Outcomes (COs)	Mapped Program Outcomes								
CO1	Write the structure, name and the type of isomerism of the	PO1, PO11, PO12,								
	organic compound	PS01								
CO2	Write the reaction, name the reaction and orientation of	PO1, PO5, PO11,								
02	reactions	PO12 PEO2,								
	Understand the resonance hybridization of different organic	PO1,PO3,PO5, PO8,								
CO3	compounds.	PO11, PO12,								
		PEO1,PEO2								

10 Hours

08 Hours

07 Hours



CO4	Account for reactivity/stability of compounds,	PO1,PO11,PO12,PEO1
CO5	Preparation chemical reaction and use of different organic	PO1,PO5,PO11,
COS	compounds	PO12,PEO1,PEO2

Course Code	Course Title	Pharmaceutical Knowledge	Problem solving	Conduct, analyze and interpret data	A Ability to design and formulating a process	A bility to understand mechanism	Demonstrate skills in problem solving	Professional and ethical responsibilities	Communication to present a technical report	Impact on society and responsibilities	Leadership qualities	Self educating and Life-long Learning	Preparation for competitive examinations	Building a theoretical knowledge base along with necessary practical skills	To build a strong foundation as per the requirements of pharmaceutical Industries, Community and Hospital	Training students to achieve expertise
TIU- UBP- 301T	PHARMACEUTICAL ORGANIC CHEMISTRY – II Theory	3	-	2	-	3	-	-	2	-	-	3	3	3	3	2

1=weakly mapped

2= moderately mapped

3=strongly mapped

Pharmaceutical Organic Chemistry II- Practical (TIU-UBP-305P) Contact hours: 4hrs/wk

Course Objectives

- 1.To know the systematic qualitative analysis of unknown organic compounds
- 2.To prepare solid derivatives of organic compounds.
- 3.To understand the molecular models.

Course Outcomes:

After successful completion of this course, students will be able to:

- CO1. **Demonstrate** the systematic qualitative analysis of unknown organic compounds
- CO2. Evaluate the Functional group and melting point of organic compounds
- CO3. Understand the suitable solid derivatives from organic compounds

CO4. Demonstrate the molecular models



Course Content Hours/Week

1. Systematic qualitative analysis of unknown organic compounds like

- a. Preliminary test: Color, odour, aliphatic/aromatic compounds, saturation and unsaturation, etc.
- b. Detection of elements like Nitrogen, Sulphur and Halogen byLassaigne's test
- c. Solubility test
- d. Functional group test like Phenols, Amides/ Urea, Carbohydrates, Amines, Carboxylic acids, Aldehydes and Ketones, Alcohols, Esters, Aromatic and Halogenated Hydrocarbons, Nitro compounds and Anilides.
- e. Melting point/Boiling point of organic compounds
- f. Identification of the unknown compound from the literature using melting point/ boiling point.
- g. Preparation of the derivatives and confirmation of the unknown compound bymelting point/ boiling point.
- h. Minimum 5 unknown organic compounds to be analysed systematically.
- 2. Preparation of suitable solid derivatives from organic compounds
- 3. Construction of molecular models

Recommended Books (Latest Editions)

- 1. Organic Chemistry by Morrison and Boyd
- 2. Organic Chemistry by I.L. Finar , Volume-I
- 3. Textbook of Organic Chemistry by B.S. Bahl& Arun Bahl.
- 4. Organic Chemistry by P.L.Soni
- 5. Practical Organic Chemistry by Mann and Saunders.
- 6. Vogel's text book of Practical Organic Chemistry
- 7. Advanced Practical organic chemistry by N.K.Vishnoi.
- 8. Introduction to Organic Laboratory techniques by Pavia, Lampman and Kriz.
- 9. Reaction and reaction mechanism by Ahluwaliah/Chatwal.

Relationship between the Course Outcomes (COs) and Program Outcomes (POs)

Mapping between COs and POs									
	Course Outcomes (COs)	Mapped Program Outcomes							
CO1	Demonstrate the systematic qualitative analysis of unknown organic compounds	PO1,PO3,PO12,PEO1							
CO2	Evaluate the Functional group and melting point of organic compounds	PO1, PO3, PO12,PEO1,PEO2, PEO3							



CO3	Understand the suitable solid derivatives from organic compounds	PO1,PO12
CO4	Demonstrate the molecular models	PO1,PO4,PO5,PO12,
CU4		PEO1,PEO2, PEO3

		Pharmaceutical Knowledge	Problem solving	Conduct, analyze and interpret data	Ability to design and formulating a process	Ability to understand mechanism	Demonstrate skills in problem solving	Professional and ethical responsibilities	Communication to present a technical report	Impact on society and responsibilities	Leadership qualities	Self educating and Life-long Learning	Preparation for competitive examinations	Building a theoretical knowledge base along with necessary practical skills	To build a strong foundation as per the requirements of pharmaceutical Industries, Community and Hospital	Training students to achieve expertise
Course Code	Course Title	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PEO1	PEO2	PEO3
TIU- UBP- 305P	Pharmaceutical Organic Chemistry II– Practical	3	-	2	2	2	-	-	-	-	-	-	2	3	2	2

1=weakly mapped 2= moderately mapped 3=strongly mapped

Physical Pharmaceutics I-Theory (TIU-UBP-302T) Contact hours: 45 hrs

Course Objectives

1.To understand various physicochemical properties of drug molecules in the designing the dosage forms

2.To know the principles of chemical kinetics & to use them for stability testing and determination of expiry date of formulations



3.To demonstrate use of physicochemical properties in the formulation development and evaluation of dosage forms.

4. To understand various chemical and physical properties of different drug molecules.

Course Outcomes

Upon completion of the course, the student shall be able

CO-1.**Demonstrate** the various physicochemical properties of drug molecules in the designing of the formulation.

CO-2. **Identify** the different states of matter and their properties.

CO-3.**Discuss** the different surface and interfacial properties related to pharmaceutical formulations..

CO-4.**Identify** the role of pH and buffers in pharmaceutical formulations.

CO-5. **Discuss** the role of protein binding and complexation in pharmaceutical formulations.

Course Content	45 Hours

Unit-

10h

Solubility of drugs: Solubility expressions, mechanisms of solute solvent interactions, ideal solubility parameters, solvation & association, quantitative approach to the factors influencing solubility of drugs, diffusion principles in biological systems. Solubility of gas in liquids, solubility of liquids in liquids, (Binary solutions, ideal solutions)Raoult'slaw, real solutions. Partially miscible liquids, Critical solution temperature and applications.Distribution law, its limitations and applications.

U	nit-
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10h

Describes of Matter and properties of matter:Describe of matter, changes in the Describe of matter, latent heats, vapour pressure, sublimation critical point, eutectic mixtures, gases, aerosols – inhalers, relative humidity, liquid complexes, liquid crystals, glassy Describes, solid- crystalline, amorphous &polymorphism.Physicochemical properties of drug molecules: Refractive index, optical rotation,dielectric constant, dipole moment, dissociation constant, determinations and applications.

Unit- III

Surface and interfacial phenomenon: Liquid interface, surface & interfacial tensions, surface free energy, measurement of surface & interfacial tensions, spreading coefficient, adsorption at liquid interfaces, surface active agents, HLB Scale, solubilization, detergency, adsorption at solid interface.

09h

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Unit-

08h

Complexation and protein binding: Introduction, Classification of Complexation, Applications, methods of analysis, protein binding, Complexation and drug action, crystalline structures of complexes and thermodynamic treatment of stability constants.

Unit

08h

pH, buffers and Isotonic solutions: Sorensen's pH scale, pH determination (electrometric and calorimetric), applications of buffers, buffer equation, buffer capacity, buffers in pharmaceutical and biological systems, buffered isotonic solutions.

Recommended Books: (Latest Editions)

1.Physical Pharmacy by Alfred Martin

2. Experimental Pharmaceutics by Eugene, Parott.

3. Tutorial Pharmacy by Cooper and Gunn.

4. Stocklosam J. Pharmaceutical Calculations, Lea & Febiger, Philadelphia.

5.Liberman H.A, Lachman C., Pharmaceutical Dosage forms, Tablets, Volume-1 to 3, MarcelDekkar Inc.

6.Liberman H.A, Lachman C, Pharmaceutical Dosage forms. Disperse systems, volume 1, 2, 3. Marcel Dekkar Inc.

7. Physical Pharmaceutics by Ramasamy C and ManavalanR.

8. Laboratory Manual of Physical Pharmaceutics, C.V.S. Subramanyam, J. Thimma settee

9. Physical Pharmaceutics by C.V.S. Subramanyam

10. Test book of Physical Phramacy, by Gaurav Jain & Roop K. Khar

Relationship between the Course Outcomes (COs) and Program Outcomes (POs)

	Mapping between COs and Pos									
	Course Outcomes (COs)	Mapped Program Outcomes								
CO1	Recognize the various physicochemical properties of drug molecules in the designing formulation Know the thermodynamic principles to use them to study stability of dosage form and demonstrate knowledge of pharmaceutical science & basic science.	PO1, PO4, PO5, PEO2,PO12								
CO2	Identify various use of physicochemical properties in evaluation of dosage forms.	PO1, PEO2								

IV

V



	CO3	Demonstrate and apply different formulate and solve the problems associated with pharmaceutical industries.PO1, PO4 PO1										, PO4, PO11	PO6,			
	CO4	releva	nt to p of phy	harma sicoch	ceutica emical	l indus prope	stries a	& syn as well of dru	as int	erpret	the		O2,PO2 PEO2			
		Pharmaceutical Knowledge	Problem solving	Conduct, analyze and interpret data	Ability to design and formulating a process	Ability to understand mechanism	Demonstrate skills in problem solving	Professional and ethical responsibilities	Communication to present a technical report	Impact on society and responsibilities	Leadership qualities	Self educating and Life-long Learning	Preparation for competitive examinations	Building a theoretical knowledge base along with necessary practical skills	To build a strong foundation as per the requirements of pharmaceutical Industries, Community and Hospital	Training students to achieve expertise
Course Code	Course Title	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PEO1	PEO2	PEO3
TIU- UBP- 302T	Physical Pharmaceutics- I- Theory	3	3	2	3	2	3	-	-	_	-	3	2	3	3	2

1=weakly mapped 2= moderately mapped 3=strongly mapped

Physical Pharmaceutics I – Practical (TIU-UBP-306P)

Contact hours: 4hrs/wk

Course Objectives:-

1.To understand physical pharmacy and able to design as per pharmaceutical industry

2.To know the principles and acquire basic knowledge about identifying the basic properties and the ADME of dosage form

3.To demonstrate various kinds of solubilization, absorption technique in lab scale as well as industry scale.

EM-4,Sector-V, SaltLake,Kolkata-700091,India



Course Outcomes:

After successful completion of this course, students will be able to: CO1. **Demonstrate** the role of various physicochemical properties related to pharmaceutical formulation design. CO2. **Identify** the role of pKa and surface tension in design of pharmaceutical formulation. CO3. **Evaluate** the effect of partition coefficient andstability constant related to pharmaceutical formulation design.

CO4. Explain suitable graphical plots for presentation of results.

Course Content Hours/Week

4

1.Determination the solubility of drug at room temperature

2. Determination of pKa value by Half Neutralization/ Henderson Hasselbalch equation.

3.Determination of Partition co- efficient of benzoic acid in benzene and water

4. Determination of Partition co- efficient of Iodine in CCl4 and water

5.Determination of % composition of NaCl in a solution using phenol-water system by CST method

6.Determination of surface tension of given liquids by drop count and drop weight method

7.Determination of HLB number of a surfactant by saponification method

8. Determination of Freundlich and Langmuir constants using activated char coal

9.Determination of critical micellar concentration of surfactants

10.Determination of stability constant and donor acceptor ratio of PABA-Caffeine complex by solubility method

11.Determination of stability constant and donor acceptor ratio of Cupric-Glycine complex bypH titration method

Reference Books (Latest Editions)

1. Physical Pharmacy by Alfred Martin

2. Experimental Pharmaceutics by Eugene, Parott.

3. Tutorial Pharmacy by Cooper and Gunn.

4. Stocklosam J. Pharmaceutical Calculations, Lea & Febiger, Philadelphia.

5.Liberman H.A, Lachman C., Pharmaceutical Dosage forms, Tablets, Volume-1 to 3, MarcelDekkar Inc.

6.Liberman H.A, Lachman C, Pharmaceutical Dosage forms. Disperse systems, volume 1, 2,

3. Marcel Dekkar Inc.

7. Physical Pharmaceutics by Ramasamy C and ManavalanR.

8. Laboratory Manual of Physical Pharmaceutics, C.V.S. Subramanyam, J. Thimma settee

9. Physical Pharmaceutics by C.V.S. Subramanyam

10. Test book of Physical Phramacy, by Gaurav Jain & Roop K. Khar



Relationship between the Course Outcomes (COs) and Program Outcomes (POs)

	Mapping between COs and Pos									
	Course Outcomes (COs)	Mapped Program Outcomes								
CO1	Demonstrate various formulation Describeed in the pharmaceutical industry	PO1,PEO1, PEO2, PEO3								
CO2	Identify various use of physicochemical properties in evaluation of dosage forms.	PO1,PO5, PEO1, PEO2, PEO3								
CO3	Evaluate the effect of various factors related to different physicochemical properties of drugs	PO1, PO3, PEO1, PEO2, PEO3								
CO4	Explain suitable graphical plots for presentation of results	PO1, PO3, PEO1,PEO2, PEO3								

		Pharmaceutical Knowledge	Problem solving	Conduct, analyze and interpret data	Ability to design and formulating a process	Ability to understand mechanism	Demonstrate skills in problem solving	Professional and ethical responsibilities	Communication to present a technical report	Impact on society and responsibilities	Leadership qualities	Self educating and Life-long Learning	Preparation for competitive examinations	Building a theoretical knowledge base along with necessary practical skills	To build a strong foundation as per the requirements of pharmaceutical Industries, Community and Hospital Pharmacv	Training students to achieve expertise
Course Code	Course Title	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PEO1	PEO2	PEO3
TIU-UBP- 306P	Physical Pharmaceutics I - Practical	3	2	3	2	2	2	-	-	-	-	-	-	3	2	3

1=weakly mapped 2= moderately mapped

3=strongly mapped

Pharmaceutical Microbiology –Theory (TIU-UBP-303T) Contact hours: 45 hrs

Course Objectives

- 1. To understand the methods of identification, cultivation and preservation of various microorganisms
- 2. To have an idea about the importance and implementation of sterilization in pharmaceuticalprocessing and industry.

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- 3. To know sterility testing of pharmaceutical products.
- 4. To learn microbiological standardization of Pharmaceuticals.
- 5. To gain knowledge about the cell culture technology and its applications in pharmaceutical industries.

Course Outcomes

Upon completion of the course, the student shall be able

CO1.**Summarize** the various nutritional requirements, physical parameters required for the growth and preservation of bacterial culture, together with bacterial count.

CO2. Classify the different sterilization techniques along with their merits, demerits and efficiency.

CO3.Differentiate between antiseptics and disinfectants.

CO4.**Demonstrate** the types of microorganisms, disinfectants as well as principles and methods of different microbiological assay.

CO5.**Identify** sources and types of microbial contaminants, assessment of microbial contamination and spoilage and method for prevention of pharmaceutical contamination.

Course Content	45 Hours
Unit I	10 Hours
Introduction, history of microbiology, its branches, scope and its	
importance.	
Introduction to Prokaryotes and Eukaryotes	
Study of ultra-structure and morphological classification of bacteria,	
nutritional requirements, raw materials used for culture media and physical	
parameters for growth, growth curve, isolation and preservation methods	
for pure cultures, cultivation of anaerobes, quantitative measurement of	
bacterial growth (total & viable count). Study of different types of phase contrast microscopy, dark field	
microscopy and electron microscopy.	
incroscopy and electron incroscopy.	
Unit II	10 Hours
Identification of bacteria using staining techniques (Simple, Gram's &Acid fast staining) and biochemical tests (IMViC).	
Study of principle, procedure, merits, demerits and applications of physical,	
chemical gaseous, radiation and mechanical method of sterilization.	
Evaluation of the efficiency of sterilization methods.	
Equipments employed in large scale sterilization.	
Sterility indicators.	
Unit III	10 Hours
Study of morphology, classification, reproduction/replication and	10 110415



cultivation of Fungi and Viruses. Classification and mode of action of disinfectants Factors influencing disinfection, antiseptics and their evaluation. For bacteriostatic and bactericidal actions Evaluation of bactericidal & Bacteriostatic. Sterility testing of products (solids, liquids, ophthalmic and other sterile products) according to IP, BP and USP.

Unit IV

08 Hours

Designing of aseptic area, laminar flow equipments; study of different sources of contamination in an aseptic area and methods of prevention, clean area classification. Principles and methods of different microbiological assay. Methods for

Principles and methods of different microbiological assay. Methods for standardization of antibiotics, vitamins and amino acids. Assessment of a new antibiotic.

Unit V

07 Hours

Types of spoilage, factors affecting the microbial spoilage of pharmaceutical products, sources and types of microbial contaminants, assessment of microbial contamination and spoilage.

Preservation of pharmaceutical products using antimicrobial agents,

evaluation of microbial stability of formulations.

Growth of animal cells in culture, general procedure for cell culture,

Primary, established and transformed cell cultures.

Application of cell cultures in pharmaceutical industry and research.

Reference Books

1. W.B. Hugo and A.D. Russel: Pharmaceutical Microbiology, Blackwell Scientific publications, Oxford London.

2. Prescott and Dunn., Industrial Microbiology, 4th edition, CBS Publishers & Distributors, Delhi.

- 3. Pelczar, Chan Kreig, Microbiology, Tata McGraw Hill edn.
- 4. Malcolm Harris, Balliere Tindall and Cox: Pharmaceutical Microbiology.
- 5. Rose: Industrial Microbiology.
- 6. Probisher, Hinsdill et al: Fundamentals of Microbiology, 9th ed. Japan
- 7. Cooper and Gunn's: Tutorial Pharmacy, CBS Publisher and Distribution.
- 8. Peppler: Microbial Technology.
- 9. I.P., B.P., U.S.P.- latest editions.
- 10. Ananthnarayan: Text Book of Microbiology, Orient-Longman, Chennai
- 11. Edward: Fundamentals of Microbiology.
- 12. N.KJain: Pharmaceutical Microbiology, Vallabh Prakashan, Delhi
- 13. Bergeys manual of systematic bacteriology, Williams and Wilkins- A Waverly company

Relationship between the Course Outcomes (COs) and Program Outcomes (POs)

Mapping between COs and Pos



	Course Outcomes (COs)	Mapped Program Outcomes
CO1	Summarize the various nutritional requirements, physical parameters required for the growth and preservation of bacterial culture, together with bacterial count.	PO1, PO3, PO12, PEO1
CO2	Classify the different sterilization techniques along with their merits, demerits and efficiency.	PO1, PO5, PO11, PO12, PEO1, PEO2
CO3	Compare between antiseptics and disinfectants.	PO1,PO11, PO12,PEO2
CO4	Demonstrate the principles and methods of different microbiological assay.	PO1,PO12,PEO1, PEO3
CO5	Identify sources and types of microbial contaminants,assessment of microbial contamination and spoilage and method for prevention of pharmaceutical contamination.	PO1,PO12,PEO1,PEO2

		Pharmaceutical Knowledge	Problem solving	Conduct, analyze and interpret data	Ability to design and formulating a process	Ability to understand mechanism	Demonstrate skills in problem solving	Professional and ethical responsibilities	Communication to present a technical report	Impact on society and responsibilities	Leadership qualities	Self educating and Life-long Learning	Preparation for competitive examinations	Building a theoretical knowledge base along with necessary practical skills	To build a strong foundation as per the requirements of pharmaceutical Industries, Community and Hosnital Pharmacy	
Course Code	Course Title	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PEO1	PEO2	PEO3
TIU- UBP- 303T	Pharmaceutical Microbiology - Theory	3	-	2	-	2	-	-	-	-	-	2	3	3	2	2

1=weakly mapped 2= moderately mapped 3=strongly mapped

Pharmaceutical Microbiology– Practical (TIU-UBP-307P) Contact hours: 4hrs/wk

Course Objectives

- 1. To know the fundamentals of sterilization and its industrial utility.
- 2. To administer the knowledge and techniques required for sub culturing of microbes.
- 3. To understand the concept of microbiological assay.

Course Outcomes:

After successful completion of this course, students will be able to:



CO1.**Operate** the various sterilizing equipments.

CO2.Prepare the sub-cultures of bacteria and fungus.

CO3.**Describe** the various staining techniques.

CO4.**Describe** the microbiological assay and bacteriological analysis of various pharmaceutical products.

Course Content

4 Hrs/week

1. Introduction and study of different equipments and processing, e.g., B.O.D. incubator, laminar flow, aseptic hood, autoclave, hot air sterilizer, deep freezer, refrigerator, microscopes used in experimental microbiology.

2. Sterilization of glassware, preparation and sterilization of media.

3. Sub culturing of bacteria and fungus. Nutrient stabs and slants preparations.

4. Staining methods- Simple, Grams staining and acid-fast staining (Demonstration with practical).

5. Isolation of pure culture of micro-organisms by multiple streak plate technique and other techniques.

6. Microbiological assay of antibiotics by cup plate method and other methods

7. Motility determination by Hanging drop method.

- 8. Sterility testing of pharmaceuticals.
- 9. Bacteriological analysis of water

10. Biochemical test.

Reference Books

1. W.B. Hugo and A.D. Russel: Pharmaceutical Microbiology, Blackwell Scientific publications, Oxford London.

2. Prescott and Dunn., Industrial Microbiology, 4th edition, CBS Publishers & Distributors, Delhi.

3. Pelczar, Chan Kreig, Microbiology, Tata McGraw Hill edn.

- 4. Malcolm Harris, Balliere Tindall and Cox: Pharmaceutical Microbiology.
- 5. Rose: Industrial Microbiology.
- 6. Probisher, Hinsdill et al: Fundamentals of Microbiology, 9th ed. Japan
- 7. Cooper and Gunn's: Tutorial Pharmacy, CBS Publisher and Distribution.
- 8. Peppler: Microbial Technology.
- 9. I.P., B.P., U.S.P.- latest editions.
- 10. Ananthnarayan: Text Book of Microbiology, Orient-Longman, Chennai
- 11. Edward: Fundamentals of Microbiology.
- 12. N.KJain: Pharmaceutical Microbiology, Vallabh Prakashan, Delhi
- 13. Bergeys manual of systematic bacteriology, Williams and Wilkins- A Waverly company

Relationship between the Course Outcomes (COs) and Program Outcomes (POs)

	Mapping between COs and POs	
	Course Outcomes (COs)	Mapped Program Outcomes
CO1	Operate the various sterilizing equipments.	PO1, PO4, PO11, PO12, PEO1, PEO2,



		PEO3
CO2	Prepare the subcultures of bacteria and fungus.	PO1, PO4, PO12, PEO1, PEO2, PEO3
CO3	Describe the various staining techniques.	PO1, PO5, PO12, PEO1,PEO2, PEO3
CO4	Describe the microbiological assay and bacteriological analysis of various pharmaceutical products.	PO1, PO2, PO5, PO12, PEO1, PEO2, PEO3

Course	Course	Pharmaceutical Knowledge	Problem solving	Conduct, analyze and interpret data	Ability to design and formulating a process	Ability to understand mechanism	Demonstrate skills in problem solving	Professional and ethical responsibilities	Communication to present a technical report	Impact on society and responsibilities	Leadership qualities	Self educating and Life-long Learning	Preparation for competitive examinations	Building a theoretical knowledge base along with necessary practical skills	To build a strong foundation as per the requirements of pharmaceutical Industries, Community and Hospital	Training students to achieve expertise
Code	Title															
TIU- UBP- 307P	Pharmaceutical Microbiology - Practical	3	2	_	2	2	_	-	-	-	_	2	3	3	3	3

1=weakly mapped

2= moderately mapped

3=strongly mapped

Pharmaceutical Engineering-Theory (TIU-UBP-304T) Contact hours: 45 hrs

Course Objectives

1. To know various unit operations used in Pharmaceutical industries.

2. To understand the material handling techniques.

3. To carry out various test to prevent environmental pollution.

4. To appreciate and comprehend significance of plant lay out design for optimumuse of resources.

5. To appreciate the various preventive methods used for corrosion control inPharmaceutical industries.

Course Outcomes



Upon completion of the course, the student shall be able

CO1. **Recognize** the importance of the various unit operations used in pharmaceutical

Industries like filtration, centrifugation, mixing, size separation, size reduction, flow of fluids, heat transfer, distillation, evaporation, drying, etc

CO2. **Identify** the various the various preventive methods used for hazards and corrosion control in Pharmaceutical Industries.

CO3. **Demonstrate** the applications of laws in the various unit operations for optimization of the process.

CO4. **Interpret** the results for problem solving in various unit operations.

CO5. **Classify** the equipments according to their mechanism of operation and identify their use for production of various pharmaceuticals.

Course Content

UNIT-I

Flow of fluids: Types of manometers, Reynolds number and its significance, Bernoulli's theorem and its applications, Energy losses, Orifice meter, Venturimeter, Pitot tube and Rotometer.

Size Reduction: Objectives, Mechanisms & Laws governing size reduction, factors affecting size reduction, principles, construction, working, uses, merits and demerits of Hammer mill, ball mill, fluid energy mill, Edge runner mill & end runner mill.

Size Separation: Objectives, applications & mechanism of size separation, official standards of powders, sieves, size separation Principles, construction, working, uses, merits and demerits of Sieve shaker, cyclone separator, Air separator, Bag filter & elutriation tank.

UNIT-II

Heat Transfer: Objectives, applications & Heat transfer mechanisms. Fourier's law, Heat transfer by conduction, convection & radiation. Heat interchangers & heat exchangers.

Evaporation: Objectives, applications and factors influencing evaporation, differences between evaporation and other heat process. principles, construction, working, uses, merits and demerits of Steam jacketed kettle, horizontal tube evaporator, climbing film evaporator, forced circulation evaporator, multiple effect evaporator& Economy of multiple effect evaporator.

Distillation: Basic Principles and methodology of simple distillation, flash distillation, fractional distillation, distillation under reduced pressure, steam distillation & molecular distillation

UNIT-III

Hours

Drying: Objectives, applications & mechanism of drying process, measurements & applications of Equilibrium Moisture content, rate of drying curve. principles, construction,

12 Hours

10 Hours

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working, uses, merits and demerits of Tray dryer, drum dryer spray dryer, fluidized bed dryer, vacuum dryer, freeze dryer.

Mixing: Objectives, applications & factors affecting mixing, Difference between solid and liquid mixing, mechanism of solid mixing, liquids mixing and semisolids mixing. Principles, Construction, Working, uses, Merits and Demerits of Double cone blender, twin shell blender, ribbon blender, Sigma blade mixer, planetary mixers, Propellers, Turbines, Paddles&Silverson Emulsifier.

UNIT IV

Hours

Filtration: Objectives, applications, Theories & Factors influencing filtration, filter aids, filter medias. Principle, Construction, Working, Uses, Merits and demerits of plate & frame filter, filter leaf, rotary drum filter, Meta filter & Cartridge filter, membrane filters and Seidtz filter.

Centrifugation: Objectives, principle & applications of Centrifugation, principles, construction, working, uses, merits and demerits of Perforated basket centrifuge, Non-perforated basket centrifuge, semi continuous centrifuge & super centrifuge.

UNIT V

Hours

Materials of pharmaceutical plant construction, Corrosion and its prevention: Factors affecting during materials selected for Pharmaceutical plant construction, Theories of corrosion, types of corrosion and there prevention. Ferrous and nonferrous metals, inorganic and organic non metals, basic of material handling systems.

Reference Books (Latest Editions)

1. Introduction to chemical engineering . Walter L Badger & Julius Banchero, Latest edition

2. Unit operation of chemical engineering .Mcabe Smith, Latest edition,

3. Pharmaceutical Engineering principles and practices .C.V.S.Subrahmanyam et al., Latest edition

4. Physical Pharmacy-Martin et al., Latest edition

5. Bentleys Pharmaceutics. Davis, Latest edition

6.Physical Pharmaceutics.Shotton, Latest edition

7.Remington Practice of Pharmacy.Martin, Latest edition

8. Cooper and Gunn.s Tutorial Pharmacy, S.J. Carter., Latest edition

9. Theory and practice of Industrial Pharmacy by Lachman., Latest edition

10.Refrigeration and Air conditioning by L. Ballaney., Latest edition.

11."Coulson and Richardson's Chemical Engineering", Vol. I, 3rd Edition,

Butterworth Heinemann Publishers, 2004. Coulson, J.M., and Richardson, J.F.

12. Pharmaceutical Engineering, K. Sambhamurthy, New Age International publishers

08



Relationship between the Course Outcomes (COs) and Program Outcomes (POs)

	Mapping between COs and POs	
	Course Outcomes (COs)	Mapped Program Outcomes
CO1	Recognize the importance of the various unit operations used in pharmaceutical Industries like filtration, centrifugation, mixing, size separation, size reduction, flow of fluids, heat transfer, distillation, evaporation, drying, etc	PO1, PO12, PEO1, PEO2
CO2	Identify the various the various preventive methods used for hazards and corrosion control in Pharmaceutical Industries.	PO1, PEO2
CO3	Demonstrate the applications of laws in the various unit operations for optimization of the process.	PO1, PO4, PO11, PO12,PEO1,PEO2, PEO3
CO4	Interpret the results for problem solving in various unit operations.	PO1,PO2,PO3, PO6, PEO1, PEO2, PEO3
CO5	Classify the equipments according to their mechanism of operation and identify their use for production of various pharmaceuticals.	PO1,PO5,PO11, PO12,PEO1,PEO2

	Gumm	Pharmaceutical Knowledge	Problem solving	Conduct, analyze and interpret data	Ability to design and formulating a process	Ability to understand mechanism	Demonstrate skills in problem solving	Professional and ethical responsibilities	Communication to present a technical report	Impact on society and responsibilities	Leadership qualities	Self educating and Life-long Learning	Preparation for competitive examinations	Building a theoretical knowledge base along with necessary practical skills	To build a strong foundation as per the requirements of pharmaceutical Industries, Community and Hospital	Training students to achieve expertise
Course Code	Course Title	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PEO1	PEO2	PEO3
TIU- UBP- 304T	Pharmaceutical Engineering- Theory	3	3	3	2	2	3	-	-	-	-	3	3	3	3	2

1=weakly mapped 2= moderately mapped

3=strongly mapped

Pharmaceutical Engineering– Practical (TIU-UBP-308P)

EM-4,Sector-V, SaltLake,Kolkata-700091,India



Contact hours: 4hrs/wk

Course Objectives

- 1.ToDescribe various processes involved in pharmaceutical manufacturing process.
- 2.To interpret the results with the help of suitable calculations
- 3.To apply the knowledge in problem solving in manufacturing.

Course Outcomes:

After successful completion of this course, students will be able to:

CO1. Demonstrate various unit operations Describeed in the pharmaceutical industry

- CO2. Relate various laws and related equations governing the various unit operations
- CO3. Evaluate the effect of various factors on the unit operations.
- CO4. Use suitable graphical plots for presentation of results

Course Content

4

Hours/Experiment

- 1. Determination of radiation constant of brass, iron, unpainted and painted glass.
- 2. Steam distillation To calculate the efficiency of steam distillation.
- 3. To determine the overall heat transfer coefficient by heat exchanger.
- 4. Construction of drying curves (for calcium carbonate and starch).
- 5. Determination of moisture content and loss on drying.
- 6. Determination of humidity of air i) From wet and dry bulb temperatures –use of Dew point method.
- 7. Description of Construction working and application of Pharmaceutical Machinery such as rotary tablet machine, fluidized bed coater, fluid energy mill, de humidifier
- Size analysis by sieving To evaluate size distribution of tablet granulations Construction of various size frequency curves including arithmetic andlogarithmic probability plots.
- 9. Size reduction: To verify the laws of size reduction using ball mill and determining Kicks, Rittinger's, Bond's coefficients, power requirement and critical speed of Ball Mill.
- 10.Demonstration of colloid mill, planetary mixer, fluidized bed dryer, freeze dryer and such other major equipment.
- 11.Factors affecting rate of filtration and evaporation (surface area, concentration and thickness/viscosity)
- 12. To study the effect of time on the Rate of Crystallization.
- 13. To calculate the uniformity Index for given sample by using Double Cone

Blender.

Reference Books (Latest Editions)

1.Introduction to chemical engineering . Walter L Badger & Julius Banchero, Latest edition

- 2. Unit operation of chemical engineering .Mcabe Smith, Latest edition,
- 3. Pharmaceutical Engineering principles and practices .C.V.S.Subrahmanyam et al., Latest edition
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- 5. Bentleys Pharmaceutics. Davis, Latest edition
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Relationship between the Course Outcomes (COs) and Program Outcomes (POs)

	Mapping between COs and POs	
	Course Outcomes (COs)	Mapped Program Outcomes
CO1	Demonstrate various unit operations Describeed in the pharmaceutical industry	PO1,PEO1, PEO2, PEO3
CO2	Relate various laws and related equations governing the various unit operations	PO1,PO5, PEO1, PEO2, PEO3
CO3	Evaluate the effect of various factors on the unit operations.	PO1, PO3, PEO1, PEO2, PEO3
CO4	Use suitable graphical plots for presentation of results	PO1, PO3, PEO1,PEO2, PEO3

Course Code	Course Title	Pharmaceutical Knowledge	Problem solving	Conduct, analyze and interpret data	Ability to design and formulating a process	Ability to understand mechanism	Demonstrate skills in problem solving	Professional and ethical responsibilities	Communication to present a technical report	Impact on society and responsibilities	Leadership qualities	Self educating and Life-long Learning	Preparation for competitive examinations	Building a theoretical knowledge base along with necessary practical skills	To build a strong foundation as per the requirements of pharmaceutical Industries, Community and Hospital	Training students to achieve expertise
TIU- UBP- 308P	Pharmaceutical Engineering - Practical	3	2	3	2	2	2	-	-	-	-	-	-	3	2	3

1=weakly mapped

2= moderately mapped

3=strongly mapped