

Principal: Dr. S. B. Bari M.Pharm. Ph.D., D.I.M.F.J.C.

List of Tutorials (S. Y. B. Pharmacy)

Academic Year 2023-24

BP301T Pharmaceutical Organic Chemistry-II – Theory

- 1. Explain resonance in Benzene.
- 2. Explain the Huckle rule.
- 3. Give the structure and uses of cresol and resorcinol.
- 4. Explain reaction of benzene.
- 5. Explain analytical constat of acid value, iodine value, ester value, and saponification value.
- 6. Give the structure and uses of naphthalene, phenanthrene, and triphenylmethane.
- 7. Explain bayers strain theory.
- 8. Explain sachse mohrs theory.
- 9. Draw Haworth synthesis of naphthalene.
- 10. Draw Haworth synthesis of Phenanthrene.
- 11. Give the reaction of anthracene.
- 12. Give in detail about the reaction of cyclopropane.
- 13. Give in detail about the reaction of cyclobutane.
- 14. Explain Coulson-moffit modification.
- 15. Explain synthetic uses of aryl diazonium salt..



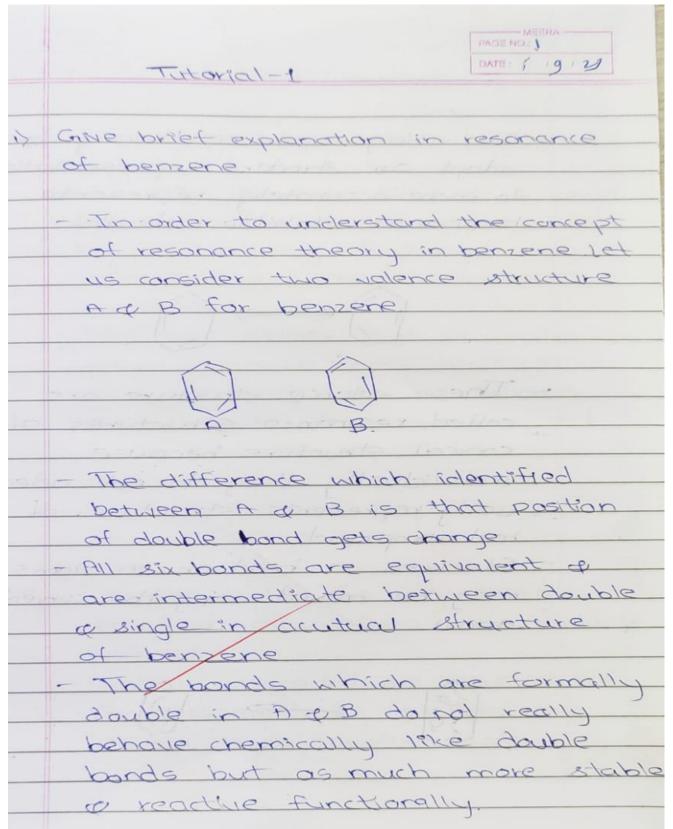


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PAGE NO. 2	MSE NO. 35 DATE 13 19 123
	Tutorial-Z
- In using resonance theory to adapt our standard representation	2) Explain Hullers Rule
to more occurately represents responding stabilise molecules	- In 1931, German chemist & physi- cist Frich Mckel proposed a rule
and protest generally to	by determining if planar ring molecules would have aromatic
	properties This rule states that if cyclic
- These valence structure are	planax molecule has 4n to 77 - A cyclic ring molecule follows Hudsel rule when number of
called resonance structures of	it is electrons equal 4n+2
conical structure because individually they do not adequetly represents structure of real compound	n = zero or any positive
- Electrons in bond or an atoms may be moved using the curved arrow	- Huckles rule was organizing based on calculations using Huckle's method
	- promotic compound are more stable than theoretically predicted by alkene hydrogenation data
The state of the stand	





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BP303T Pharmacognosy Microbiology – Theory

Prepare 10 multiple choice questions (MCQs) with four options (Underline correct option).

Sr. No	Topics	Sr. No	Topics
01	Introduction, history of microbiology, its branches, scope and its importance, Introduction to Prokaryotes and Eukaryotes	9	Designing of aseptic area, laminar flow equipments; study of different sources of contamination in an aseptic area and methods of prevention, clean area classification
02	Study of ultra-structure and morphological classification of bacteria, Nutritional requirements, raw materials used for culture media and physical parameters for growth, growth curve	10	Principles and methods of different microbiological assay. Methods for standardization of antibiotics, vitamins and amino acids
03	Isolation and preservation methods for pure cultures, cultivation of anaerobes, quantitative measurement of bacterial growth (total & viable count)	11	Assessment of a new antibiotic and testing of antimicrobial activity of a new substance. General aspects-environmental cleanliness
04	Study of different types of phase contrast microscopy, dark field microscopy and electron microscopy	12	Study of morphology, classification, reproduction/replication and cultivation of Virus
05	Identification of bacteria using staining techniques (Simple, Gram's & Acid fast staining) and biochemical tests (IMViC)	13	Types of spoilage, factors affecting the microbial spoilage of pharmaceutical products, sources and types of microbial contaminants, assessment of microbial contamination and spoilage
06	Study of principle, procedure, merits, demerits and applications of Physical, chemical and mechanical method of sterilization. Evaluation of the efficiency of sterilization methods, Sterility indicators	14	Preservation of pharmaceutical products using antimicrobial agents, evaluation of microbial stability of formulations
07	Study of morphology, classification, reproduction/replication and cultivation of Fungi. Classification and mode of action of disinfectants	15	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
08	Factors influencing disinfection, antiseptics and their evaluation. For bacteriostatic and bactericidal actions. Evaluation of bactericidal & bacteriostatic		

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	Tutorial work	PAGE NO.: DATE: 30 /12/23
4	Topici- History & branch biology, scope & imported biology.	ince of misso-
1	T+ deals with the students of infections	by of rausa-
2)	a) Soil Microbiology b) Ai c) Medical microbiology a) foo	n microbiology.
- 2	of RBC cells.	human shape
	b) Lours parsteur c) Robert Koch d) Lord joseph 195ter.	
	Methods is use to killi microorganisms.	ng of
	Jonembrane Filtration of All	1 of this

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List of Tutorials (S. Y. B. Pharmacy)

Academic Year 2023-24

BP 304T Pharmaceutical Engineering – Theory

Sr No	Tutorials	
1.	Write a principle & draw a labelled diagram of FBD.	
2.	Write a principle & draw a labelled diagram of lyophilizer.	
3.	Write a principle & draw a labelled diagram of plate & frame	
	filter.	
4.	Write a principle & draw a labelled diagram of rotary drum filter.	
5.	Write a principle & draw a labelled diagram of perforated basket	
	Centrifuge	
6.	Write a principle & draw a labelled diagram of fluid energy mill.	
7.	Write a principle & draw a labelled diagram of cyclone separator.	
8.	Write a principle & draw a labelled diagram of ball mill.	
9.	Write a principle & draw a labelled diagram of orifice meter.	
10.	Write a principle & draw a labelled diagram of multiple effect	
	evaporator.	
11.	Write a principle & draw a labelled diagram of fractional	
	distillation.	
12.	Write a principle & draw a labelled diagram of steam distillation.	
1.0		
13.	Write a principle & draw a labelled diagram of twin shell blender.	
14.	Write a principle & draw a labelled diagram of planetary mixers.	
15.	Write a principle & draw a labelled diagram of Silverson	
	Emulsifier.	







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	Tulorial - 1	PAGE NO.: 1 DATE: 5 19 123
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0	diagram of FBD.	AN -
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200000	to prepare granules of a Content in the production	of the tablets.
Conto I		
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	Principle: In fluidized bed de is passed through a perfe of the Container Containing	ng the wet golids
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	and become Suspended	in the air/gas
	Stream. At this stage So	olid bed looks
	like the boiling liquid a as fluidized. The hot ai	and hence Called
	as fluidized. The hot ai	r / gas Surrounds
	each fluidized particle.	leat transfer
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	each fluidized particle. It is accomplished by direct the out solid & hot as	I/gases. The
	Vapourised liquid is Carrie	ed away by the
	drying air/ gasses.	0

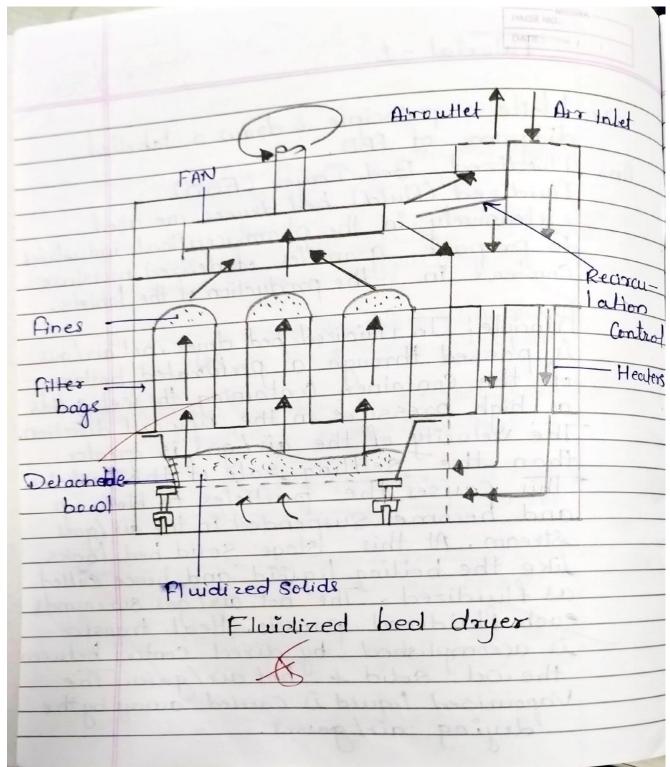
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Academic Year 2023-24

BP302T Physical Pharmaceutics I – Theory

Tutorial 1

Q. Define solubility and explain the ways of expressing solubility.

Tutorial 2

Q. What is the supersaturated solution? Add a note on mechanism of solute solvent interaction.

Tutorial 3

Q. Write in brief about Raoults law and its significance.

Tutorial 4

Q. What do you mean by states of matter? Add a note on sublimation critical point.

Tutorial 5

Q. Define aerosol. Give principle, construction and working of it.

Tutorial 6

Q. Explain in detail about refractive index and optical rotation as a physicochemical property of drug molecule.

Tutorial 7

Q. Write in detail about surface tension and interfacial tension.

Tutorial 8

Q. Give methods of surface tension determination.

Tutorial 9

Q. What do you mean by HLB scale? Give its its significances.

Tutorial 10

Q. Define complexation and give detail classification of it.

Tutorial 11

Q. Prepare 15 MCQs on complexation and protein binding.

Tutorial 12

Q. What is crystalline structure of complexes? Write its significances.

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Tutorial 13

Q. Define pH and add a note on pH scale and application of it.

Tutorial 14

Q. Give detail account of pH determination methods.

Tutorial 15

Q. What do you mean by buffer & buffer capacity? Write the importance of buffer in

Pharmaceutical and biological system.

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		MEERA
		PAGE NO.:
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	litee of Solution.	
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(0)	N = No of geam equival	ent of solute
9	Litee of Solu	ition

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	1-lorgo DATE:
- 10	Geam equivalent = Geam of solute 4
2	Molarity - is defined as the molarity of solute per liters of a solution of a substitution of a certain volume of solution. Molarity is also known as the molarity of a solution.
and and	Molarity = Number of moles Solute (n) x1000 Volume of Solution in L.
3	Molality (m) or molal concentration is the amount of a substance dissolved in a certain mass of solvent.
ano ni	of solute per kilogeam of solvent
skilos h	m = Moles of abolite. Mass of Solvent (inkg)

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	PAGE NO.: DATE: / /
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	the components, moles representing of
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	Total Moles of elolution.
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	Percentage by weight. Percentage mass (Percentage by weight) is the percent of the fotal mass of the Solution that is one component.
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	× 100
6.	Percentage by Volume :-
	Volume / Volume Percentage is
	Volume / Volume l'excentage is a measure of concentration of a
	substance in a solution.
	It is expressed as the
	substance in a Solution. It is expressed as the ratio of the solute

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	PAGE NO.: DATE:
	male fraction-
90 300 Oc.	to total volume of the solution multiplied by 100 ml.
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To post	Vol. / = Valume of Solute X 100
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	mililities of solution.
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BP405T Pharmacognosy and Phytochemistry I – Theory

Prepare 10 separate MCQs with correct option underlined on the following topics by individual student. (Avoid repetition of MCQs)

Tutorial No.	Topic	
01	Definition, history and scope of Pharmacognosy	
02	Sources of drugs, Organized and unorganized drugs	
03	Classification of drugs	
04	Adulteration of drugs	
05	Evaluation of drugs by organoleptic and microscopic methods	
06	Evaluation of drugs by chemical and biological methods	
07	Factors affecting cultivation of medicinal plants	
08	Plant hormones, polyploidy, mutation, hybridization	
09	Plant tissue culture	
10	Pharmacognosy in various system of medicine	
11	Secondary metabolites	
12	Plant fibres, Hallucinogens, Teratogens, Natural allergens	
13	Carbohydrates	
14	Proteins and enzymes	
15	Lipids, Marine drugs	







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	CH CO.	PAGE NO.:
	Tutorial - 1	DATE: / /
W)	Define history & scope	of Planting
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Academic Year 2023-24

BP401T Pharmaceutical Organic Chemistry III- Theory

Tutorial 1: Define & classify stereoisomer with example

Tutorial 2: What do you mean by racemic mixture & racemic modification? Explain various methods of resolution of a racemic mixture.

Tutorial 3: Give the various method for determination of geometrical isomers

Tutorial 4: Explain in detail the RS system of nomenclature of optical isomers.

Tutorial 5: What are stereoselectivity and stereospecificity? Explain it with a suitable example.

Tutorial 6: Explain the reaction & mechanism involved in Backmanns rearrangements.

Tutorial 7: Explain the reaction & mechanism involved in the Schmidt reaction.

Tutorial 8: Explain synthesis, reactions & medicinal uses of furan

Tutorial 9: Explain synthesis, reactions & medicinal uses of thiophene

Tutorial 10: Explain the synthetic method & two characteristic reactions for Imidazole & pyrrole

Tutorial 11: Give any two methods of preparation & chemical reaction of pyridine.

Tutorial 12: Explain the chemical reactions of Quinoline and Isoquinoline

Tutorial 13: Write a note on reactions and synthesis of Indole.

Tutorial 14: Write synthesis and medicinal uses of Pyrimidine.

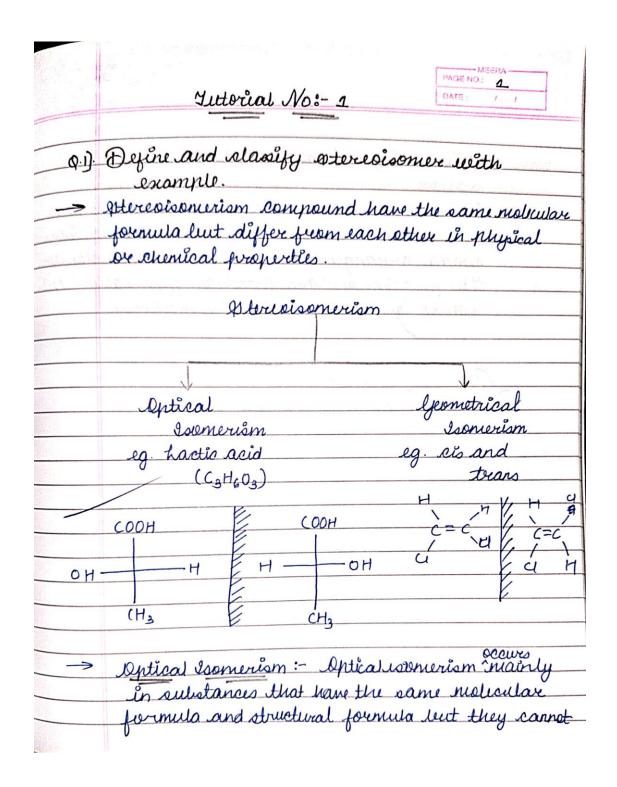
Tutorial 15: Comment on electrophilic substitution in five-membered heterocycles.







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	PAGE NO.: 3
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0.2].	What do you mean by reacenic mixture and
	seacenic modification? Explain various metrodo
Renig	What do you mean ley reacente mixture and reacente modification? Explain various metrods of resolution of a reacente mixture.
	The state of the s
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	A mixture having equal amounts of
	enantionier is called recenile mixture.
	Racenic modification:
	It is the process of separation of a racenic modification ento enartioners sonstituents.
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	have different shapes from each others. So they
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List of Tutorials (S. Y. B. Pharmacy)

Academic Year 2023-24

BP403T Physical Pharmaceutics II – Theory

Tutorial 1

Q. What do you mean by colloidal dispersion? Give detail account of their general characteristics.

Tutorial 2

Q. Explain in detail optical and electrical properties of colloids..

Tutorial 3

Q. Frame at least 15 MCQs on colloidal dispersion.

Tutorial 4

Q. What do you mean by newtonian system? Explain in detail.

Tutorial 5

Q. What are the non-newtonian system? Explain in brief dilatant and pseudoplastic flow.

Tutorial 6

Q. Define viscosity; enlist different viscometers and explain in brief cup and bob viscometer.

Tutorial 7

Q. What do you mean by suspension? Comment on interfacial properties of suspended particles.

Tutorial 8

Q. Define emulsion and explain theory of emulsification.

Tutorial 9

Q. Write detail note on emulsion formulation by HLB method.

Tutorial 10

Q. Define micromeritics; add a note on particle size and its distribution.

Tutorial 11

Q. Enlist and explain any one method for particle size determination.

Tutorial 12

Q. What are the derived properties of powder explain in detail.

Tutorial 13

Q. What do you mean by drug stability? Explain in brief second order kinetics.

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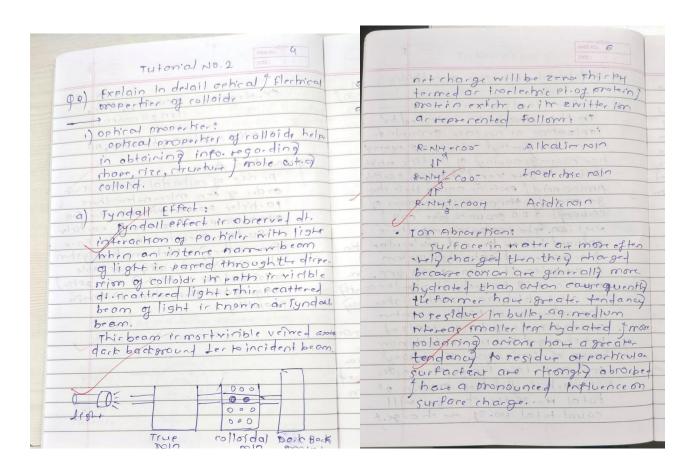
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Tutorial 14

Q. Enlist and explain physical and chemical factors influencing stability of pharmaceutical product

Tutorial 15

Q. Write a note on accerated stability study.





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Electrical properties of collections

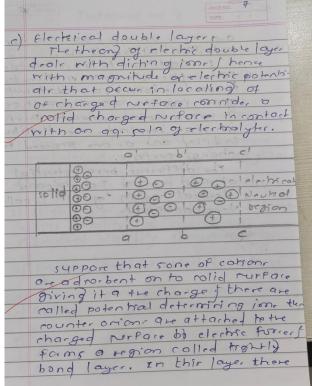
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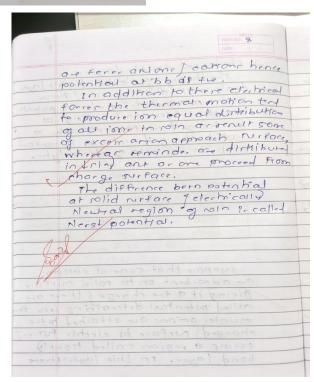
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Academic Year 2023-24

BP402T Medicinal Chemistry – Theory

- 1. Define Lead. Give its sources of Optimization.
- 2. Explain Bioisosterism and its application on biological acton
- 3. Explain Ionization and Partition Coefficient.
- 4. Discuss about neurotransmitters in body
- 5. Write in brief about biosynthesis of catecholamine
- 6. I. Write a classification of sympathomimetic agents
 - II. Write SAR of Adrenergic agents
- 7. Discuss SAR of β blockers.
- 8. Write in brief about chemistry of acetylcholine.
- 9. Write the short note on coline esterase reactivator. Write SAR of cholinolytic.
- 10. Discuss SAR of Barbiturate, Phemnothiazine and Benzodiazepines.
- 11. Write 5 MCQ's on Sympathomimetic agents
- 12. Write 5 MCQ's on Parasympathomimetic agents
- 13. Write the classification and stages of G.A.
- 14. Give the MOA of NSAID.
- 15. Give SAR of mepiride series ansd classification of Opoid analgesics







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BP701T Instrumental Methods Of Analysis

(Tutorial)

Sem VII A Y 2023-24

- 1. Comment on type of electrons & electronic transitions involved in UV-spectroscopy.
- 2.Add a note on Beer's & Lambert's law.
- 3. What are the factors affecting fluorescence intensity.
- 4.Describe in details about vibrations in IR Spectroscopy.
- 5. Explain instrumentation of Flame Photometry give its applications.
- 6.Discuss the principle and applications of Atomic absorption spectroscopy.
- 7. Explain the instrumentation of Nephelo-turbidimetry.
- 8. Write principle, advantages, disadvantages & applications of TLC.
- 9. Give principle and explain various steps involved in Paper chromatography.
- 10. Explain the factors offering electrophoretic mobility.
- 11. Explain the types of detectors used in Gas chromatography.
- 12. Explain applications of HPLC with a schematic diagram.
- 13. Explain the ion exchange resins for Ion exchange Chromatography.
- 14. Write a note on Gel chromatography.
- 15. Explain instrumentation of Affinity chromatography

PAGE NO .: DATE: Tatorial No:1 comment on types of electrons of electronic transitions involved in UV-spectroscopy. Types of electrons: 1) o (sigma) electrons . Trivolved in single bonds 2) IT (pi) electrons: Involved in multiple bonds, such as double of triple bonds 3) n(non bonding electrons? Lucalized on heteroatoms (eg. 0, N) not involved in bonding Electronic transitions ot (antibonding) TI+ (antibording) D->+ THOP Trot n (non-bonding) T (bonding) 11+)11* J-IT T (bonding)

	PAGE NO.:	
	DATE: / /	
::::::::::::::::::::::::::::::::::::::		1
	σ→σ* transition →	<i>.</i>
•	or electron from orbital is excited to	
	corresponding anti-bonding orbitals	
^	The energy required is large for	
	this transition.	
2]	Π→ Π ^{tr} transition →	
-	compounds like alkenes, alkyny	1
	carbonye initriles , aromatic compound	1
	containing multiple bonds undergo	1
	IT → IT+ transition.	*
		1
3)	n→ σ* Transition :-	1
	sourated compounds contaîning	
	atoms with lone pair of electrons like	爾
	O, NI, S & halogens are capable of mot	
	fransition.	
-	requires 1855 energy than or of Hanse,	
	4) n -> TT# transition !-	
	compound contening double bond	
	involving heteroatoms (=0.1(ZN, N=0)	100
	undergo such transitions.	

		Duggue	
		PAGE NO.: DATE: / /	
		·	
_	requires minimum ene	2794	
5_) r -> 77# & T -> 6# +	ransition ->	
	These electronic tr	ansition are	
	Forbidden transitions	4 are only	
	theoretically possible.		
	thoughand vestible.		
	·		

•	
	PAGE NO.:
	Tutorial NO:2
9	Add a note on Beer's & lamberts
	(aw -)
9)	Lampelis law:
	It states that when a beam
	of monochromatic radiation passes
	through a homogeneous absorbing
	medium, the rate of decrease of
	intensity of radiation with thickness
	of absorbing medium is proportium
	to intensity of incident radiation.
	Mathematically, the law is
	expressed as - d? = KI
	चेंद्र
	Let Io be the intensity of radiation
	before entering into absorbing
	medium (x=0)
	Then 2, the intensity of radiation
	after passing through any thickness
	SCIU IT
	(dz = x=2 (kdx
	J TP J
	20 + 20
	- In I KX
	T

PAGE NO .: DATE: 2.303 log 2 = -kz A = KX A= ER 2.303 Absorption of Thickness b) Beer's low: This law states that when a beam of monochromatic radiation is passed through a sul of absorbing sub. the rate of decrease of intensity of radiation which thickness or absorbing soin is propertioned to intensity of incident rediction as which thickness of absorbing sol is propertional to intensity -dI - KIC suppose to be the intensity of radiation before entering into

PAGE NO .: DATE: absorbing medium when thickness 2.303 log I = -kcte 109 I = - KCX 2,303 K' CX 2.303 A = GiCR absorption of Thickness & concentration



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List of Tutorials (Final Year B. Pharmacy)

Academic Year 2023-24

BP 702 T Industrial Pharmacy - II - Theory

Sr. No.	Tutorials
1.	What is pilot plant scale up?
2.	Give the pilot plant consideration for solid formulations
3.	What are SUPAC guidelines? Discuss in detail
4.	Write in brief about the technology transfer protocol
5.	Write in brief about the quality risk management
6.	Management of Clinical Studies
7.	Contents of the Investigational New Drug (IND) Application
8.	Investigator's Brochure
9.	Explain the principles of Total Quality Management
10.	Define the various concepts in the Quality by Design (QbD)
11.	Write in brief about the Out of Specifications (OOS)
12.	Write in brief about NABL
13.	Central Drug Standard Control Organization (CDSCO)
14.	Certificate of Pharmaceutical Product (COPP),
15.	Regulatory requirements and approval procedures for New Drugs

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102.004		
PAGRIO.		Tutorial :4. BAGE NO.
Automorphism of the control of		Control of the second of the s
	1	
	Q.	Write in brief about tech. transfer
		protocol.
on	1)	大大学 1000 1000 1000 1000 1000 1000 1000 1
		The transfer process should be
fg.		managed by SUGRU. If
		organised, can additional agency
		in which proper direction and
		approval are provided. These
		should be a proper managemen
		plant and format agreement
		for fr.
		/ Following Steps would be follow
Company of the Compan		os per er protocol.
	(1)	pumpose and objective of transfer
	カ	sope of transfor
	3)	skilled personnel.
Market Control of Cont	4)	Companision of materials.
	5)	Documented evidence
	. 6)	The transfer of documented
	7)	assessment of ccp.
	8)	processment of expt.
	9)	Information of different butches.

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	PASE NO: DATE:		
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List of Tutorials (Final Y. B. Pharmacy) Academic Year 2023-24 BP704T_Novel Drug Delivery System- Theory

Tutorial	Tutorial Questions			
number	A Division (VKC)			
1	Give classification of polymers			
2	Explain the concept suntained & CRDDS ?			
3	Enlist different approaches designing CRDDS.			
4	Enlist Factors affecting Formulation For TDDRS.			
5	Enlist approaches for retentive DDS			
6	Give principle & gastro-mucoadhesion.and bio adhesion			
7	Give advantage of mucosal drug delivery system			
8	Enlist method of encapsulation (micro)			
9	Write the concept of implant & armatic pump. 10. 4/9/23 Explain the concept of			
	navopulmonary drug delivery system			
10	Write method to overcome challenge occur in ocular drug delivery system.			
11	Write method to overcome challenge occur in ocular drug delivery system.			
12	Enlist occular formulation & write in detail about ocuserts.			
13	Give advantages and disad. vantages of intrauterine drug delivery system.			
14	Explain the concept of drug delivery system.			
15	Enlist approaches for targeted drugs			



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Tetorial: 1

Q d	Give classification of polymers.
reas parties and his factories and the factories	Classification:
	Based on origin / source
	. Natural polymers
ا ا	Synthetic polymers
	semisynthetic polymers.
	Bared on structure
Q.	Linear polymer
6.	Branched chain polymer Coors-linked polymer
С.	Coors - linked polymer
	Bared on molecular forces.
and the first of the second se	elastomers
	fibres
	Themplastic
d.	Thermosetting polymers.
4	Based in mode of polymensation.
Q.	Based on mode of polymerisation. Addition polymer Condensation polymer.
5.	condensation polymer.



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Tutorial: 2

malerille

9. Explain the concept sustained selease and controlled selease drug delivery system

> Sustained seleased drug delivery system:

Many provide an immediate dose sequised for the normal therapeutic sesponse followed by graphical seleased of drug in amounts of sufficient to maintain the therapeutic sesponse for specific extended period of time usually 8-12 hours

immediate relevence

a pre-determined ratio for locally

t (time)



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List of Tutorials (Final Year B. Pharmacy)

Academic Year 2023-24

BP804 ET Pharmaceutical Regulatory System-Theory

Sr. No.	Tutorials
1.	Flow chart of pre-clinical trials.
2.	Flow chart of stages of drug discovery process.
3.	Flow chart stages of clinical trials.
4.	Flow chart of NDA review process.
5.	Flow chart of ANDA review process.
6.	Summary of CDSCO.
7.	Summary of TGA.
8.	Summary of USFDA.
9.	Summary of MHLW.
10.	Summary of Belmont Report.
11.	Summary of Helsinki Declaration.
12.	Duties of Principle investigator.
13.	Duties of Sponsor.
14.	Outline of Clinical trial protocol.
15.	Flow chart of procedure for export of pharmaceutical products.

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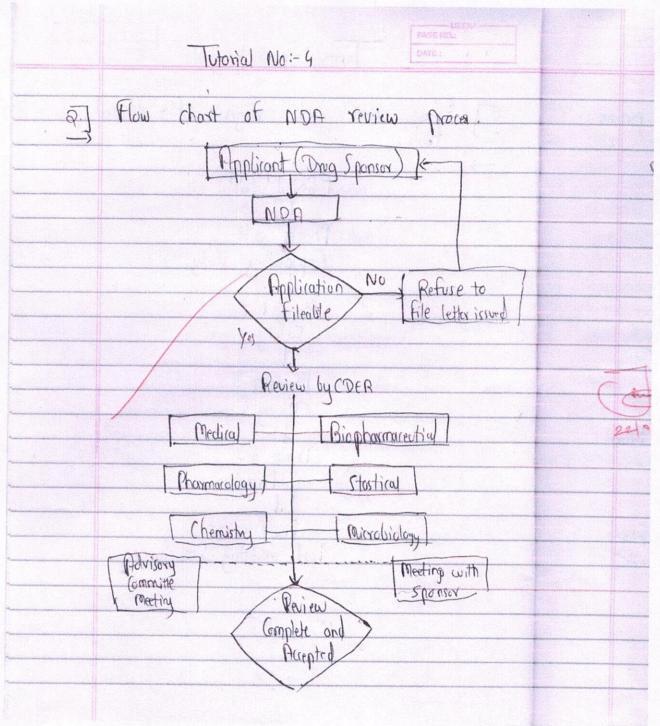
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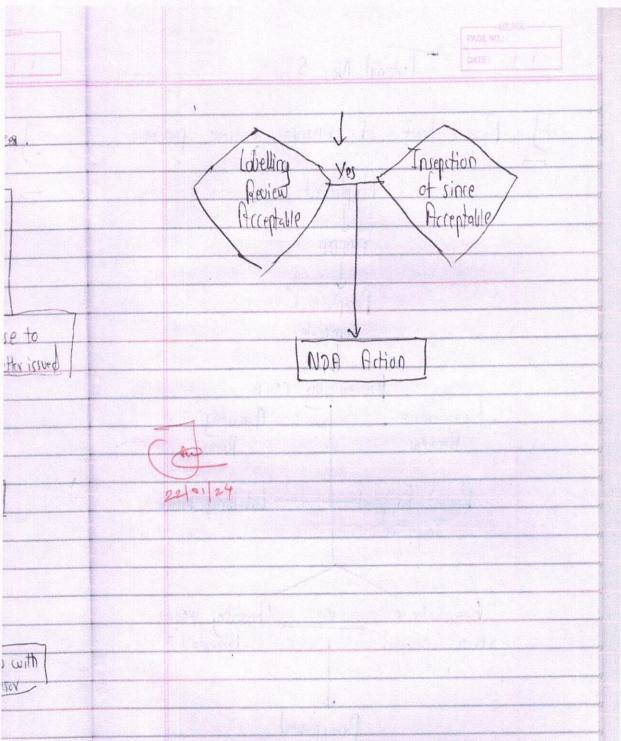
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List of Tutorials (Final Year B. Pharmacy)

Academic Year 2023-24

BP802T Social and Preventive Pharmacy-Theory

- 1. Prepare MCQ on with answer on topic Concept of health and disease.
- 2. Prepare MCQ on with answer on topic Social and health education.
- 3. Prepare MCQ on with answer on topic Sociology and health.
- 4. Prepare MCQ on with answer on topic Hygiene and health.
- 5. Write note on prevention and control of cholera, influenza and SARS.
- 6. Write note on Prevention and control on diabetes and cancer.
- 7. Explain in details HIV and AIDS programme.
- 8. Elaborate national mental health programme.
- 9. Explain pulse polio programme.
- 10. Discuss the role of WHO Indian national programme.
- 11. Give details about intervention programme for mother and child.
- 12. Describe tuberculosis programme mission and objective.
- 13. Write down function of PHC.
- 14. Elaborate tobacco control programme.
- 15. Describe National urban health mission.

Mr S.K. Amale

Subject in-Charge

a write note on constation and continuous Tutorial-5 of cholera, influence and SAP. -> * cholera -1) safe water and sanitation-Ensuring access to clean water and proper sanitation facilities is crucial in preventing cholero transmission. Impris water quality and promoting hygiene practices such as handwashing can reduce the risk of contamination a) vaccination - vaccination against cholers can provide protection for individual for endemic areas or during outer and cholero vaccines are available and recommended for travelers to regions with known cholera authorit 3) public Health education- Educating communities about cholery transmis Symptoms and preventive measures is essential for early detection and response promoting safe food hand

practices and encouraging proper

to cholero prevention.

waste disposal can also contribute

* SARS (Severe Acute Respiration) Syndrome) :-1) Infection control measures; Implementing strict infection measures in healthcase settings. Encluding proper use of prince protective equipment (PPE) Suc. 1 marks, gloves and gowns, can he prevent nosocomical transmission of e) Isolation and avarantine: Identific and Isolating suspected SARS (0) along with quarantining class conhelp contains the spread of the visur withing Communities and prevent further transmission. 3) Travell restrictions Implementing travel advisories and restrictions regions affected by SARS Outbreet can reduce the spread of the virus across borders and limit !! global impact

Influenta:

Naccination - Annual influency vaccination

Ps recommended, especially for high

nist Individuals Such as the elderly,

young children, pregnent women, and
individuals with undelying health

condition, vaccination helps reduce

the severity of illness and prevent

complications.

2) personal hygiene - practice good respiration

hygiene, Such as covering cough and

personal hygiene - practice good respiration hygiene, such as covering cough and sneezes with a tissue are elbow, and regular handwalking with soap and water can help prevent influenza transmission.

3) Social distancing measures: During Influence outbreak, implementing Social distancing measures such as awording Crowded places, staying home when sick, and maintaining physical dist from individual with flu-like symptoms can help reduce transmission mate

		,
*	Mca	

- 1) what is health
 - A Absence of disease
- 18. complete physical well-being
- c. presence of mental itt-near
 - D. Lack Of social Connections
- 2) which model of health emphasizes the interaction of biological, psychological. and social factors
 - A. Biomedical model
 - B. Ecological model
 - C. social model
- L. Biopsychorocial model
- 3) what is the primary focus of preventire medicine
- A treating existing disease

 B Early detection and intervention
 - c. Rehabilitation.
 - d. palliative case.

- 9) which of the following is a non-rommy disease
 - A. Influenza
 - B. Tuberculoric
 - Le Diabetes
 - d) Malaria.
- 5) what is the term for the spread of a dyeare within a specific population or geographic asea.
 - A pandemic
- & Epidemic
- C outbrek
- D Endemic
- 6 which lifestyle factor is a major contributor to cardiovarcular disease
 - A Regular exercise
 - B. Adequale sleep
- D vegetarian diet

what is	the	Dux pose	20	the	who
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- A. provide medical insurance
- 8. Combat infections disease
 - c. conduct scientific meseasch
 - o. promote specific diety
- 8) which level of prevention focused on minimizing the impact of established disease
 - A. primary powention
 - B. Secondary prevention
 - E. Tertiary prevention
 - D. quaternooy prevention
- 9) what is the term for the ability of an organism to resist infection or diseases
- LA. Immunity
- B. Antibody response
- c. pathogenesis
- O vaccination.

10	which	of the	Following	es a	٠
	borne	disease	following	T-W-V	8(+
				CONTRACTOR	

- A Measles
- B Typhoid.
- LC. Denque fever





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List of Tutorials (Final. Y. B. Pharmacy)

Academic Year 2023-24

BP804ET Pharmaceutical Regulatory Science - Theory

- 1. Write a short note on drug master file (DMF).
- 2. What is CFR. Explain detail type of CFR.
- 3. Write a short note on CTD.
- 4. Write a short note on IND.
- 5. Write a short note on NDA.
- 6. Write a short note on ANDA.
- 7. Give difference between the submission process of IND, NDA & ANDA.
- 8. Enlist the parts of clinical trial protocols.
- 9. Draw organization structure of CDSCO.
- 10. Enllist the stage of drug development process.
- 11. Define bioequivalence, pharmacokinetic, pharmacodynamic, bioavailability & partition coefficient.
- 12. Write a short note on IRB board.
- 13. MCQ on drug approval process.
- 14. Draw a flow chart of IND application.
- 15. MCQ on clinical protocols.



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	Tutorial - 1 Page No. Date
0	Drucy Master File (DMF)
→	FDA of information was submirror to the
	of drucy product or a consent of
	of drung product or a component of drug product
	Type of OMF's -
- 1	Type I - Plant information
	Type I - Plant information
	Type II - Drug substance, drug produt tis
	Type III - Package
	Type IV - Excipient
	Type II - Packaging Type IV - Excepted Type V - Other information
	Current types of DMFs
	Now from typs
	Now from typs Type I DMF muthdrauer
	Taype II drug substance, drug product 4 its
	Type III Packaging
	Type It Excipins
	Type I oth information.
	DMF is submitted to the FDA for the info which is confidential.
	is confidential.



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	Type 3 = 46 (11.)
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	Type 5 = 88 (14 1.)
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Term-I

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F. Y. B. Pharmacy

List of Tutorials

Academic Year 2023-24

BP101T Human Anatomy and Physiology I– Theory

Tutorial	Tutorial Questions		
number	A Division (PSB)	B Division (PSB)	
1	Explain the structural organization of	Explain the structural organization of	
	human body	human body	
2	Draw a well labelled diagram of cell and	Draw a well labelled diagram of cell and	
	discuss its anatomy and physiology	discuss its anatomy and physiology	
3	Classify different types of tissues with	Classify different types of tissues with	
	examples	examples	
4	Explain the function of skin	Explain the function of skin	
5	Classify human bones	Classify human bones	
6	What do you mean by joints	What do you mean by joints	
7	Explain the composition and function of	Explain the composition and function of	
	blood	blood	
8	Discuss the human blood grouping system	Discuss the human blood grouping system	
9	Sketch the well labelled diagram of lymph	Sketch the well labelled diagram of lymph	
	node and explain its functions	node and explain its functions	
10	Differentiate between sympathetic and	nd Differentiate between sympathetic and	
	parasympathetic system	parasympathetic system	
11	Discuss anatomy and physiology of	of Discuss anatomy and physiology of	
	tongue	tongue	
12	Explain the cranial nerve with examples	Explain the cranial nerve with examples	
13	Explain in detail about anatomy of heart	Explain in detail about anatomy of heart	
14	Provide brief information of conduction		
	system of heart	system of heart	
15	List out the disorders of heart	List out the disorders of heart	





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List of Tutorials (F. Y. B. Pharmacy)

Academic Year 2023-24

BP102T Pharmaceutical Analysis – Theory

- 1. Scope and application pharmaceutical analysis
- 2. Methods of expressing concentration
- 3. Primary and secondary standards
- 4. Errors, accuracy and precision
- 5. Pharmacopoeia, impurities and limit test
- 6. Theories of acid-base titration
- 7. Indicators and theories of incicators with neutralisaion curve
- 8. Non-aqueous titration
- 9. Precipitation titration
- 10. Complexometric titration
- 11. Gravimetric titration
- 12. Diazotization titration
- 13. Types of redox titration with principle and application
- 14. Conductivity of cell and conductometry with application note on reference and indicator electrode of potentiometer.
- 15. Ilkonic equation with construction and working of dropping mercury electrode and rotating platinum electrode



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Tutorial NO : 01 05 of 2023
(P. 1) Write down Scope + Application of P.A.
At a The purpose of pharmocutical Analysis is to identify Substances, purify them. described the molecular structure of Chemical Compounds that makes up pharmaceutical to determine how these Compounds are Combined to make up a pharmaceutical product
o Scoped 3-
To become a pharmaceutical Analyst, I fudent must posses knowledge in the following areas biology, Chemistry, physics & mathematics.
The ideal Cardidate for the position of pharmaceutical analyst coill posses a degree in pharmaceutical anglysis, pharmacelogy, Chemistry or any related subject.



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	A 2.18 18 18 18
٥	Application :-
	P C Compating according
0	Classification of a Compound according
	to its Chemical Properties.
F _ X I-	Fra Cameourd
•	Analysis of Mixture for Compounds.
,0	# separating of a Compound from
	mixture
0	purification, identification 4
	Characterization of Compounds.
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	Compound.
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	S sol
X	C>
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List of Tutorials (F. Y. B. Pharmacy)

Academic Year 2023-24

BP103T Pharmaceutics- I Theory

Sr. No.	Tutorials
1.	Flow chart of solid dosage forms classification.
2.	Flow chart of liquid dosage forms classification.
3.	Give a model prescription.
4.	Give the formulae in posology.
5.	Parts of prescription and handling of prescription
6.	Disadvantages of powders
7.	Advantages of powders.
8.	Differentiate between lotions and liniments.
9.	Disadvantages of suspensions.
10.	Advantages of suspensions.
11.	Disadvantages of suppositories
12.	Advantages of suppositories.
13.	Write in short about ORS powder.
14.	What do you mean by displacement value?
15.	Give the formula for the preparation of Paracetamol pediatric elixir

are

Mr. S. N. Jain

Principal

Dr. S. B. Bari Principal

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Tutorial No-5	
Parts of Prescriptions and Handling of prescriptions.	-I
· ·	@Ir
Prescriptions - Prescriptions is the	① N. ② Fa
order written by the physician	3 U
Prescriptions is the order written by the physician or any other medical practitioner to the pharmacist to compount or dispense specific medication for individual patients or owner	@ Su
for individual patients or owner	Su
of the animal.	
Parts of Prescriptions:	al distribution
C Heading 8.	bJi
Theoding &. Theoding &. Thout date invalid	©T:
b) Information about prescriber	- In
Name of patient. Address Age	e]
prescriptions. b) Information about prescriber c) Information about patient- Name of patient, Address, Age and Gender / Sex of patient	Ы.
	c]
2 Superscription 3. R, symbol which means you take.	(00)
9000	15/10/17/6/21

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ndling_	-It also represent [God of Healing].
oner 1	(3) Tracriptions: (1) Name of drug (Generic or Trade Name) (2) Formulations of drugs (3) Unit dosage of drugs
nt on	Subscription :- Subscription means information given by physician to pharmacist
Jer	al Quantity to be dispensed / Amt to be use
	6 Transcription / Signa 3.
ĭ	- In formation given by physician to
ge _	Transcription / Signa : In formation given by physician to patient regarding: a) Instruction about the amount of drug to be taken.
10	of Frequency of the close to be taken
ich	6 Signature:

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List of Tutorials (F. Y. B. Pharmacy)

Academic Year 2023-24

BP104T Pharmaceutical Inorganic Chemistry – Theory

- 1. Explain & enlist various sources & types of impurities.
- 2. Describe in details-limit test of arsenic
- 3. i) Write a note on IP ii) Limit test of iron
- 4. i) Explain in details theory of acid and base ii) Define buffer & explain mechanism of action of buffers
- 5. Explain in details buffer action of acid and base
- 6. Explain in details methods of adjusting tonicity and pH
- 7. i) Write in detail the buffer system of the body ii) Define anti-carries agents. Explain its activity
- 8. i) Define GI agents, classify acidifiers. Write a note on NH4Cl. ii) Define antacids and classify them.
 - i) Define classify cathartics with examples. Describe MgSO4
- 9. ii) Define antimicrobial agents with example.
- 10. Write in details i) Boric acid ii) Iodine
- 11. Define & Classify expectorants
- 12. i) Discuss in detail haematinics ii) Define & classify poison & antidote with example.
- 13. i) What is cyanide poisoning? Explain its treatment. ii) Discuss in detail astringent.
- 14. Discuss in detail radioactive materials
- 15. Write in details radioactive decay



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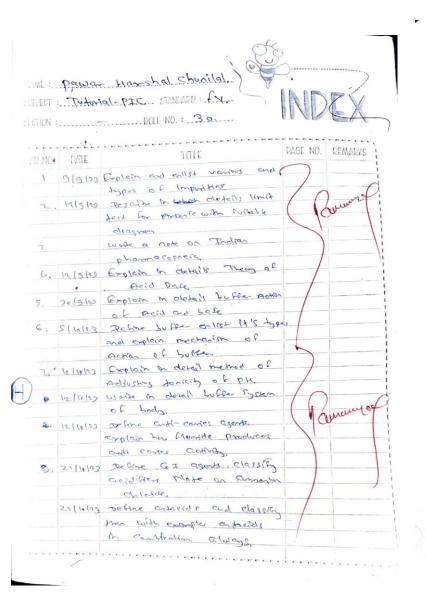


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List of Tutorials (F. Y. B. Pharmacy)

Academic Year 2023-24

BP104T Pharmaceutical Inorganic Chemistry – Theory



Sr. No.	Date	Title (1)
		Till dates martinal council.
		of Horman backen
5.	28 (10/19	Define and clossify Catherthy with
		Suitable example Describe mg
	20(10(25	action action (cabial agents with ex.
		white on detail mechanism of coster of
		antimicrobial agents.
OF	7/10/23	1) Done acid
		i) locking with 11's preparation
11.	25/11/23	- Classify expectionents
		the shired and charical proportion
		Can Amarila Chloride
	50 (11)53	Deline emeter with mechanism of
		action describe monograph or cati
		C. Inhale.
12.	26/11/23	Discuss a detail harmaticies
	26 (11/23	Define and classify poison and arridary for
		explain with Sultable example 15
13	2112123	what is cynicle poisoning 2 explain the
400-7		treatment.
((3115153	what do you mean by ractio active substance directs in details
	•	about neasurement of radioactive
		write pharmaceutical application
15.		of radio active Substance
		0
		Caronder
		h23
		13.12.2020



Dr. S. B. Bari

Principal A L

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	Tutorial -1.
0.1)	Explain an various source and the type
7	Source of Topurities , Acarding to JCH
	guidlines an impurity
•	Reagent
•	Method Consist in new days
	5-leads
•	
9)	packages - Lat
•	Storense
0	Raw moderial ? Impurities from raw material may be carried through transfacturing process and contaminate the final product. Ex. & Rock Salt (Casa, + mgcl) - Nacl. Rock Salt Contains Small amount of Calcius Sulphate and magnessium chloride, New Nacl. prepared from the Source may Contain
(3)	Reagent in the reagent, used in manufacturing are not completely remove by warding, then the may find fine product.
	To above reaction ammoniated chloride prepared contains, ammoniated hydroxides

Proprito, Corta
Now if is and
with water the
Gral product. It may contaminate the
Method :
There are various trethod used for
There are various treathed used for
Contain Drugs a multiple step synthesis
Process is a used, which producer inter
Mediated Compounds
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this intermediate Compound community
will contaminate the final product.
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water, but sametimes for reducing Cost
water that Contains
we use Softened water that contain we use Softened water Imputly that co
contaminate the final product.
Contemina
Top water : Coatain Cast, right, Nat et
as Impurity. alst as as incl
softened water in Contain Mat, cr as infl
impurity.
Distilled water ! Best but costly.

Some explace contamination! To industrial arrea admosphere is a contaminated with dust particle are home. General with admosphere contaminated. See Dustry them and got contaminated. Soi! Mach reach with admosphere contaminated that a why it show not leapt open for long three. There's why most of the industries build in guter arreas where pollution is very late. The user solvents and aggest may under the user solvents and contain at a sense of the users of the user		[*************************************
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Contaminated with dust particles are some some some some some some some som		- 1 chief correct common spread 1/1
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Exi. Mach ready with atmospheric Co. Front Contaminated that's why it show foot copt open for long time. That's why most of the industries build in outer aneas where pollution is very low. © Readien with vessel: During manufacturing process source the uses Solvents and reagent may under reaction with we vessel and Contaminate final producted and contaminate final producted some vessel and contaminate final producted some vessel and contaminate final producted some vessel that are manufacturing to the contaminate of the contami		
First contaminated that's why it show First why most of the Industries That's why most of the Industries build in outer areas where pollution is try low. Co reaction with vessel: During manufacturing process source the uses solvents and reagent may unless reaction with we vessel and contaminate final products Contain Areais as impurity The interpolation of impurity and vessel that are manufacturing processes and and a series as impurity and a vessel that are manufacturing processes and a series as impurity. The last vessel that are manufacturing processes and a series as impurity and a series as impurity. The packaging are error in the processes such as a series in the series and a series as a series in the series are series.		
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That's why most of the industries build in guter areas where pollution is very low. Co Reaction with vessel: During manufacturing process source the user Solvati and reagent may under reaction with we vessel and containate final products Ex. Iron vessel Condain Areanic as impurity in Iron vessel may contain Tron and areans as impurities.		e land ofted
Duild in outer areas where pollution is build in outer areas where pollution is try low. C Reaction with vessel: During manufacturing process source the uses Solvats and reagent may under reaction with we vessel and contaminate final productor Cx. Iron vessel Contain Argonic as impurition in Iron vessel may contain Tren and consens as impurities. Deckaging are error 1.		not kept open for long time
Ex Iron vessel may center Tren and Then vessel may contain Argenic as impurition for the manufacturing process source The user Solvents and reagent may under the user Solvents and reagent may under the user Solvents and reagent may under final producted Condain Argenic as impurition final production companies that are manufactured in them vessel may center Tren and answers as impurition.		i of the lodustries
During manufacturing process source During manufacturing process source the user Solvents and reagent may under reaction with we vessel and contain nate final productor Companies that are manufactured in manufacturing the area of the manufacturing that are manufacturing the source of impurities. The wassel may contain the and and answers as impurities.		That's why hard where pollution is
© Reaction with vessel! During manufacturing process source The user Solvents and reagent may unless reaction with neversel and contaminate final products Ex. Iron ressel Condain Arconic as impurity Then vessel may contain Tron and ansate as impurities.		build in guter areas
The user Solvents and reagent may under the user Solvents and reagent may under reaction with the vessel and containants final products Ex. Iron ressel Contain Argenic as impuri for inorganic comparate that are manif in Iron vessel may contain Fron and consents as impurities. To packaging an error 1.		very low.
The user Solvents and reagent may under the user Solvents and reagent may under reaction with the vessel and containants final products Ex. Iron ressel Contain Argenic as impuri for inorganic comparate that are manif in Iron vessel may contain Fron and consents as impurities. To packaging an error 1.	0	Reaction with vessel:
the user Solvate and reagant the reaction with we versel and contain nate final production of Contain Argenic as impurition. The local versel that contain Tran and contain as Impurition. The local versel that contain Tran and contain are manufactured as impurition.		
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Final productor Condain Angenic as impusion for Iron vessel Condain Angenic as impusion in Iron vessel may certain Iron and angenic as impusition. To packaging an error !		reaction with vervessel and contaminate in
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oncere as impurities. Deckessing an error!		· · · · · · · · · · · · · · · · · · ·
D packaging an error !.		in Iron vessel may center Tron and
1 packaging an error!		ancente as impunities.
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E-lines packed in Similar Containers		tollet of the come sample Continues le
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labelles trans also cause major package		do potential source of classic major packaging
in series that		10 Selling 1 min
X 7 8 9 2.		X 7765.

	Paga (ks.) Oda
(2)	Storage Condition 1 "
	Alle man other of final a
	After preparation of final products H Should be Stored in appropriate
	Containe depoding upon
-)	Madure of material.
	Dotch size
	Quentu
	Congrally Maderials like place
	Iron Stainker Steel and aluminium
	used for storage Improper storage
	to reaction with those materials and
	Contant nation to final product &
	//
6)	Types of Impurity:
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	horganic Impusity.
3	Residual colvents
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0	anganic impurities!
	O organic impurities basically arise dust
	Synthesis, purification and storage of
	drug Subsidance



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List of Tutorials (F. Y. B. Pharmacy) Academic Year 2023-24

BP201T Human Anatomy and Physiology-II

- 1. Describe in detail of nephron with neat labelled diagram.
- 2. Describe in detail of nephron with neat labelled diagram.
- 3. Write in detail structure and function of brain.
- 4. Draw a diagram of digestive system and describe phases of digestion.
- 5. Describe in detail structure and function of liver and stomach.
- 6. Describe in detail structure and function of lungs.
- 7. Write in detail note on salivary gland.
- 8. Describe in detail mechanism of respiration.
- 9. Draw diagram of urinary system and describe physiology of urine formation.
- 10. Describe in detail structure and function of kidney.
- 11. Describe in detail structure and function of nephron.
- 12. Describe in detail anatomy and physiology of Adrenal gland.
- 13. Describe in detail Protein synthesis.
- 14. Describe in detail male and female reproductive system.
- 15. Detail note on Menstrual cycle and spermatogenesis.

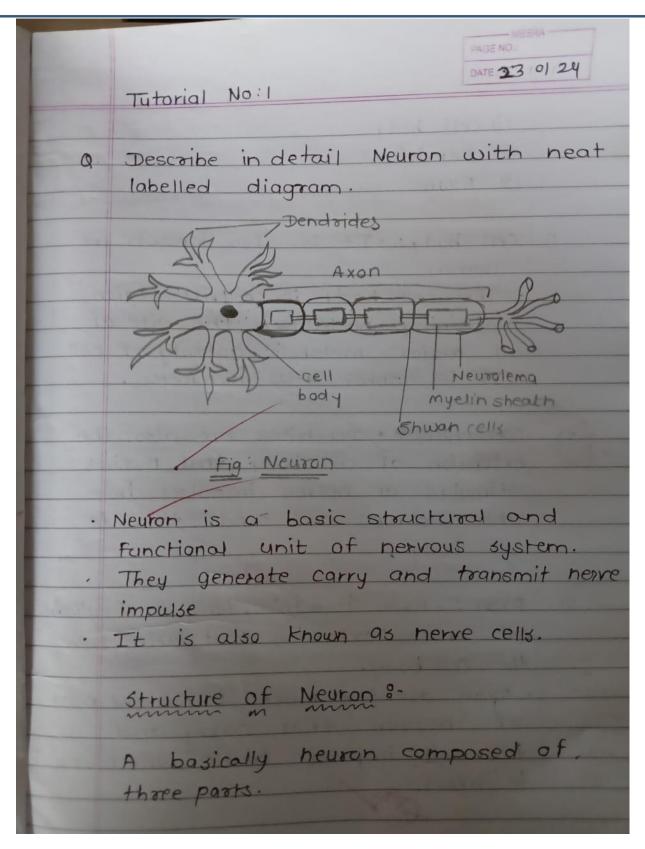




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12.5	EC NO PORTO DE LA CONTRACTOR DE LA CONTR
	① Cell body ② Dendrites ③ Axon.
	(ell body: Tt is also known as soma. The cell body is central region
	containing nucleus is the site of all major metabolic activity of cell. Tt is approx 4-100 micrometer.
(2)	Dendrides: Dendrites are also the extension of cell body that receives stimulus or nerves impulses from other neurons and sends them to cell body.
3)	Axon: Axon is a thin, long and cylindrical process / extension that axise from the cell body. Axon are the most important part of neuron that carry and transmit impulse frame one neuron to other.





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BP202T Pharmaceutical Organic Chemistry I – Theory

Tutorial	Tutoria	al Questions
number	A Division (NPP)	B Division (NPP)
1	Explain hybridization. Give a brief idea about sp3, sp2, and sp hybridization.	Explain hybridization. Give a brief idea about sp3, sp2, and sp hybridization.
2	Explain E1 and E2 reactions. Factors affecting E1 and E2.	Explain E1 and E2 reactions. Factors affecting E1 and E2.
3	Explain Markovniffs and antimorkoniffs rule.	Explain Markovniffs and antimorkoniffs rule.
4	Write a note on electrophilic addition reaction	Write a note on electrophilic addition reaction
5	Give SN1 and SN2 mechanisms.	Give SN1 and SN2 mechanisms.
6	Give a short note on Aldol condensation and benzoin condensation reaction	Give a short note on Aldol condensation and benzoin condensation reaction
7	Give a short note on the Cannizaro reaction and Perkin condensation reaction	Give a short note on the Cannizaro reaction and Perkin condensation reaction
8	Give structure and uses of formaldehyde	Give structure and uses of formaldehyde
9	Explain the acidity of carboxylic acid, and the effect of substituent on it.	Explain the acidity of carboxylic acid, and the effect of substituent on it.
10	Give structure and uses of acetic acid and tartaric acid	Give structure and uses of acetic acid and tartaric acid
11	Give factors affecting SN1 and SN2 mechanism	Give factors affecting SN1 and SN2 mechanism
12	Difference between E1 and E2 mechanism	Difference between E1 and E2 mechanism
13	Difference between SN1 and SN2 reaction	Difference between SN1 and SN2 reaction
14	Give structure and uses of Amphetamine	Give structure and uses of Amphetamine
15	Give brief introduction to aliphatic amine	Give brief introduction to aliphatic amine





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BP204T Pathophysiology – Theory

Tutorial	Tutorial (Questions
number	A Division (PSB)	B Division (PSB)
1	Differentiate between reversible and	Differentiate between reversible and
	irreversible cell injury	irreversible cell injury
2	Discuss about apoptosis related to its	Discuss about apoptosis related to its
	molecular mechanisms	molecular mechanisms
3	Explain cellular adaptations	Explain cellular adaptations
4	Discuss the mechanism of inflammation	Discuss the mechanism of inflammation
	in detail	in detail
5	Explain the chronic renal failure	Explain the chronic renal failure
6	Discuss the pathophysiology of CCF	Discuss the pathophysiology of CCF
7	Explain the pathophysiology of diabetes	Explain the pathophysiology of diabetes
8	Explain the pathophysiology of epilepsy	Explain the pathophysiology of epilepsy
9	Explain the pathophysiology of	Explain the pathophysiology of
	hepatitis	hepatitis
10	Explain the pathophysiology of cancer	Explain the pathophysiology of cancer
11	Explain the pathophysiology of	Explain the pathophysiology of
	tuberculosis	tuberculosis
12	Explain the pathophysiology of AIDS	Explain the pathophysiology of AIDS
13	Explain the pathophysiology of	Explain the pathophysiology of
	rheumatoid arthritis	rheumatoid arthritis
14	Explain the pathophysiology of peptic	Explain the pathophysiology of peptic
	ulcer	ulcer
15	Explain the wound healing	Explain the wound healing



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List of Tutorials (F. Y. B. Pharmacy)

Academic Year 2023-24

BP203T Biochemistry – Theory

- 1. Biomolecules
- 2. Bioenergetics
- 3. Glycolysis and citric acid cycle
- 4. HMP shunt and glycogen metabolic pathways
- 5. Oxidation phosphorylation and electron transport chain
- 6. Oxidation of fatty acids
- 7. Formation and utilization of ketone bodies
- 8. Cholesterol biosynthesis and disorders of lipid metabolism
- 9. Transamination and decarboxylation
- 10. Urea cycle and its disorders
- 11. Catabolism of phenylanaline and triosine and their disorders
- 12. Catabolism of heme and protein metabolism disorders
- 13. Purine and pyrimidine biosynthesis and organisation of mammalian genome
- 14. Structure and replication of RNA and DNA and RNA transcription
- 15. Enzymes and Coenzymes



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	PAGENO: 1
	DATE: 02/02/2024
	Tutorial - 1
*	Biomolecules ccarbohydrate, lipid, protein).
\longrightarrow	· Biomolecules are molecules that
	occurs naturally inside living
	organism
	Generally most of the biomolecules
	contains carbon as major element.
•	Other than carbon, biomolecules
	generally contain H, N, O, p, s.
	(Tupeso F
	biomolerules
) carbohy drates.
,	2) Lipids
	27
	3) proteins
	4) Mudeic acid.



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	PAGE NO.: 2
ز	carbohydrates:
	They are most abudant naturally occurring organic compounds in the nature.
	They are simply defined as the biomolecules containing carbon hydrogen and exugen in matio 1:2:1 they are also known as hydrogen of carbons).
	since most of carbohydrates are sweet in taste hence they are also known as sugar i.e sucrose
•	orbohydrates are also called Saccharides.
•	Genral Formula: Cn (H20n).
	Sugar like ribose deoxybose Forms genetic material dHA and RNA.



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	TAGING 3
2)	Lipid5:
	The world lipid is derived from
	greek word 'lipos' which means
	fat.
•	tipids can be defined as organic
	substances that are reactively
	insoluble in water and soluble
	in organic Solvents couch as
	ether, Chlorofrom of benzene.
•	They are hydrophoblic in nature
	unlike proteins, nucleic acid
	carpphydrates. lipid are not
	polymer.
	example: Fats and oils.
	cells membrance or plasma membr
	ance is made up of lipids.
	They are precuoss of hormones.
	ceg restosteorne, progesteron,.
	estrogen).



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	PAGE NO: 4
3)	proteins: proteins are most abudant organic molecules in living system.
,	They constitude about 50% of day cellular weight.
	They are essential for the structure function regulation of body's tissue and organs.
	proteins are made up of smaller unit called Amino acid.
P	2.02.024



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