Resolution Number : AC/II (20-21).2.RUS15

S.P. Mandali's

Ramnarain Ruia Autonomous College

(Affiliated to University of Mumbai)



Syllabus for Semester V & VI Program: B.Sc. (Drugs & Dyes) Program Code :(RUSACDD)

(Credit Based Semester and Grading System with effect from the academic year 2023-24)



PROGRAM OUTCOMES

S. P. Mandali's Ramnarain Ruia Autonomous College has adopted the Outcome Based Education model to make its science graduates globally competent and capable of advancing in their careers. The Bachelors Program in Science also encourages students to reflect on the broader purpose of their education.

РО	Description			
A stud	A student completing Bachelor's Degree in Science program will be able to:			
	Recall and explain acquired scientific knowledge in a comprehensive manner and			
PO 1	apply the skills acquired in their chosen discipline. Interpret scientific ideas and			
	relate its interconnectedness to various fields in science.			
	Evaluate scientific ideas critically, analyse problems, explore options for practical			
PO 2	demonstrations, illustrate work plans and execute them, organise data and draw			
	inferences.			
	Explore and evaluate digital information and use it for knowledge upgradation.			
PO 3	Apply relevant information so gathered for analysis and communication using			
	appropriate digital tools.			
PO 4	Ask relevant questions, understand scientific relevance, hypothesize a scientific			
104	problem, construct and execute a project plan and analyse results.			
	Take complex challenges, work responsibly and independently, as well as in			
PO 5	cohesion with a team for completion of a task. Communicate effectively,			
	convincingly and in an articulate manner.			
PO 6	Apply scientific information with sensitivity to values of different cultural groups.			
100	Disseminate scientific knowledge effectively for upliftment of the society.			
	Follow ethical practices at work place and be unbiased and critical in interpretation			
PO 7	of scientific data. Understand the environmental issues and explore sustainable			
	solutions for it.			
	Keep abreast with current scientific developments in the specific discipline and			
PO 8	adapt to technological advancements for better application of scientific knowledge			
	as a lifelong learner.			



PROGRAM SPECIFIC OUTCOMES

	Description
A stude	nt completing Bachelor's Degree in Science program with applied component
as Drug	s & Dyes will be able to:
PSO 1	Appreciate the vista of applications of chemistry in the fields of drugs and dyes
PSO 2	Become aware of the ways in which the science has, and can be applied to real
	problems.
PSO 3	Become cognizant of the important contributions of chemistry in the two fields
	of drugs and dyes, and apply their knowledge of molecules and the way in
	which they prefer to behave in specific situations.
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PROGRAM OUTLINE

	SEMESTER V			
Course Code	Unit	Course Title / Unit Title	Credits	
RUSACDD501		Drugs and Dyes	\sim	
		General Introduction to Drugs	N.	
	Ι	Routes of Drug Administration and Dosage Forms		
		Pharmacodynamic agents		
		Anti-Neoplastic Drugs		
		Anti HIV Drugs		
	II	Cardiovascular Drugs		
		Antidiabetic Agents		
		Anti parkinsonism Drugs		
		Drugs for Respiratory System		
		Introduction to Dyestuff Chemistry		
	III	Classification of dyes based on constitution	2	
		Classification Based on Application		
		Intermediates		
	IV	Preparation of intermediates Dyeing method of cotton fibres		
RUSACPDD501		Practical	2	
RUSACPDD501		Practical	2	
mai				



Course Code	Unit	Course Title/Unit Title	Credits
RUSACDD601		Drugs & Dyes	
		Drug Discovery, Design and Development	
		Drug Metabolism	$\mathbf{\mathcal{C}}$
	Ι	Chemotherapeutic Agents	
		Antibiotics	
		Antimalarials	
		Anti-inflammatory Drugs	
		Antiamoebic Drugs	
	тт	Antitubercular Drugs	
	II	Antileprotic Drugs	
		Drug Intermediates	
		Nano particles in Medicinal Chemistry	
		Colour and chemical constitution of dyes	2
	тт	Non-textile Uses of Dyes	
	III	Optical brighteners	
		Organic Pigments	
		Synthesis of specific dyes and their uses	
	IV	Types of fibres and classes of dyes applicable to	
	1.	them	
		Ecology and toxicity of dyes	
RUSACPDD601		Practical	2
RUSACPDD601			2
na	×		
			Page



Semester V Course Code: RUSACDD501 <u>Course Title: Drugs & Dyes</u> Academic year 2020-21

Course Outcomes:

After o	completing the course, the learner will be able to -
CO 1	Understand various pharmacodynamic agents with respect to their chemical
	structure, chemical class, therapeutic uses, and side effects.
CO 2	Understand different routes of drug administration.
CO 3	Describe the metabolism of drugs inside the human body.
CO 4	Enlist different routes of drug administration.
CO 5	Classify dyes based on their constitution and application.
CO 6	Correlate color and chemical constitution of dyes.
CO 7	Write the reactions involved in the synthesis of some representative drugs and dye
	intermediates.

DETAILED SYLLABUS

Course Code		Drugs & Dyes	Credits-02
RUSACDD501	Unit	Unit Title	Lectures
	I	1.1 General Introduction to Drugs	(6L)
		1.1.1 Definition of a drug, Requirements	
		of an ideal drug, Classification of	
		drugs (based on therapeutic action).	
		1.1.2 Nomenclature of drugs: Generic	
		name, Brand name, Systematic name	
		1.1.3 Definition of the following	(2L)
		medicinal terms; Pharmacon,	
		Pharmacophore, Prodrug, Half-life	
		efficiency, LD50, ED50,	(7L)
		Therapeutic Index.	



1.1.4 Brief idea of the following terms:
Receptors, Drug-receptor
interaction, Drug Potency,
Bioavailability, Drug toxicity, Drug
addiction, Spurious Drugs,
Misbranded Drugs, Adulterated
Drugs, Pharmacopoeia.
1.2. Routes of Drug Administration and
Dosage Forms
1.2.1 Oral and Parenteral routes with
advantages and disadvantages.
1.2.2 Formulations, Different dosage forms
(emphasis on sustained release
formulations.)
1.3. Pharmacodynamic agents
A brief introduction of the following
pharmacodynamic agents and the study
with respect to their chemical structure,
chemical class, therapeutic uses, and side
effects
1.3.1 CNS Drugs:
Classification based on pharmacological
actions, Concept of sedation and hypnosis,
anaesthesia. Phenobarbitone (Barbiturates –
mode of action), Phenytoin (Hydantoins),
Trimethadione (Oxazolidinediones),
Piracetam (Pyranones), Midazolam,
Alprazolam (Benzodiazepines)
Methylphenidate (Piperidines)
Chlorpromazine (Phenothiazines)
Fluoxetine (Phenyl propyl amines)



		RUIA COLLEGE Explore o Experience o Excel
	Synthesis of Trimethadione,	0
	Methylphenidate, Phenytoin.	
	1.3.2 Analgesics and Antipyretics	
	Morphine (Phenanthrene alkaloids),	
	Tramadol (Cyclohexanols), Aspirin	
	(Salicylates), Paracetamol (p-	
	Aminophenols), Synthesis of Tramadol,	
	Paracetamol.	
I	I 2.1 Anti-Neoplastic Drugs	<u>y</u>
	2.1.1. Idea of malignancy; Types of Cancer,	
	Causes of cancer, Treatment of	(3L)
	cancer (surgery, radiation therapy,	
	chemotherapy).	
	2.1.2. Chemotherapeutic agents used in the	
	treatment (Structures not	
	expected):Lomustine (Nitrosoureas),	(2L)
	Mitomycin C (Antibiotics),	
	Vincristine; vinblastine; (
	mechanism of action), Cisplatin (
	mechanism of action), Fluorouracil	(3L)
	(Pyrimidines)	
	2.1.3. Synthesis of 5-Fluorouracil from	
	urea.	
	2.2 Anti-HIV Drugs	(2L)
-0.5	2.2.1. Introduction of AIDS and HIV,	
	pathogenecity, Symptoms of AIDS,	
	mode of transmission, prevention,	(2L)
	Diagnosis and treatment	
	2.2.2. Reverse transcriptase inhibitors	
γ	(AZT, Stavudine (Pyrimidines), DDI	(3L)
	(Purines)	
	2.3 Cardiovascular drugs	



	2.3.1. Introduction, Classification based on
	pharmacological action
	2.3.2. Enalapril (-amino acids), Isosorbide
	dinitrate (Nitrates), Atenoldol
	(Aryloxy propanol amines),
	Nifedipine (Pyridines), Furosemide
	(Sulfamyl benzoic acid), Synthesis of
	Furosemide, Atenolol
	2.3.3. Drug Therapy and Renin-
	Angiotensin System.
	2.4 Antidiabetic Agents
	2.4.1. Introduction and types of diabetes;
	Insulin therapy
	2.4.2. Antidiabetic agents - Glibenclamide
	(sulphonyl ureas – mode of action),
	Metformin (Biguanides)
	2.5 Antiparkinsonism Drugs
	2.5.1. Introduction
	2.5.2. Procyclidine hydrochloride
	(Pyrrolidines), Ethopropazine
	hydrochloride (Phenothiazines),
	Laevodopa (alpha-amino acids)
	Synthesis of Levodopa from Vanillin.
.	2.6 Drugs for Respiratory System
	2.6.1. General idea of Expectorants;
\$ O-	Mucolytes; Bronchodilators
	Decongestants and Antitussives
<u>~</u> 0-	2.6.2. Bromhexine hydrochloride (Phenyl
	methyl amines), Salbutamol, Pseudo-
	ephedrine (Phenyl ethyl amines)
	Oxymetazoline (Imidazolines)
	Codeine Phosphate (Opiates)
	<u> </u>



	Synthesis of Salbutamol
III	3.1 Introduction to Dyestuff Chemistry
	3.1.1 Important landmark in the history of
	dyes
	3.1.2 . Natural colouring matter and their
	limitations: e.g., Heena, Turmeric,
	kesar, Chlorolphyll, Indigo, Alizarine (5L)
	from roots of madder plants,
	Logwood. Tyrian Purple.
	3.1.3. Synthetic Dyes: Important
	milestones, i.e. Mauve,
	Diazotization, aniline Yellow, Congo
	Red, Synthesis and structure of
	Indigo, disperse Dye, fluorescent
	Brighteners, procion reactive Dyes,
	Remazole Dyes. (Emphasis on Name
	of the Scientist and dyes and the year (5L)
	of the discovery is required and
	structure is not expected.
	3.1.4.Definition of dyes, Properties i.e.
	colour, Chromophore and
	Auxochrome, Solubility, Linearity,
	Coplanarity, fastness properties,
	substantivity, Economic viability.
	3.1.5. Explanation of nomenclature of (5L)
	commercial dyes with at least one
	example. Suffixes-G, O, R, B, 6B,
	GK, 3GK, 6GK, L, S Explanation:
	naming of dyes by colour index(two
	examples)
10	3.2. Classification of dyes based on
	constitution



		RUIA COLLEGE Explore e Experience e Exel
	(Examples are mentioned below with	
	structures)	
	Nitro Dyes-Napyhol yellow S, Nitroso Dye-	
	Gambine Y, Azo Dyes-	
	(a) Monoazo Dyes- Metanil yellow	
	(b) DiazoDyes- Napthol Blue Black	
	(c)Triazodyes -Chloroamine Green B,	
	Diphenymethane Dyes-Auramine G,	
	Triphenyl methane	
	Dyes-	
	(a) Malachite Green Series- Naphthalene	
	green V (b) Magenta Series- Acid Magenta	
	(c) Rosolic acid series-Chrome Violet,	
	Heterocyclic Dyes, Xanthene-Rhodamine	
	6G, Acridines-Acriflavine, Azines-	
	Safranine B, Oxazines-Capri blue,	
	Thiazines-Methylene Green, Quiolines-	
	Quinoline Yellow, Thiazoles-Primuline,	
	Benzoquinones and naphthaquinones –	
	Napthazarin, Anthraquinone Dyes-	
	Indanthrene, Turquoise Blue 3GK,	
	Indigoids-Indigo Caramine, Pthacyanines-	
	Sirius Light green FFGL	
	3.3 Classification Based on Application	
	Definition, fastness properties &	
	applicability on substrates examples with	
	structures (a) Acid Dyes- Orange II, (b)	
	Basic Dyes-methyl violet, Victoria Blue B	
	(c) Direct cotton Dyes- Benzofast Yellow	
	5GL (d) Azoic Dyes-Diazo components;	
0	Fast yellow G,Fast orange R. Coupling	
	components. Naphthol AS, Naphthol ASG	



	(e) Mordant Dyes-Erichrome Black A,	XX
	Alizarin. (f) Vat Dyes- Indanthrene brown	
	RRD, Indanthrene Red 5GK. (g) Sulphur	
	Dyes- Sulphur Black T (no structure) (h)	
	Disperse Dyes-Celliton Fast brown 3R,	
	perlon fastblue FFR (i) Reactive Dyes-	
	cibacron Brillant Red B,procion briilant	
	Blue HB.	
IV	4.1 Intermediates	
	4.1.1. A brief idea of Unit processes	
	4.1.2. Introduction of primary	
	intermediates, unit processes	
	4.1.3. Nitration, Sulphonation,	()
	Halogenation, Diazotization: 3	(5L)
	different methods, importance,	
	Ammonolysis, Oxidation	
	N.B.: Definition, Reagents	
	Examples with reaction conditions	
	(mechanism is not expected)	(7 L)
	4.2 Preparation of the following	
V	Intermediates.	
	4.2.1 Benzene derivatives:	
	Benzenesulphonic acid; 1,3-	
	Benzenedisulphonic acid; phenol;	
	resorcinol; sulphanilic acid; o-,m-,p-	
	chloronitrobenzenes; o-,m-,p-	(3L)
	nitroanilines; o-,m-p- phenylene	
	diamines; Naphthol ASG.	
	4.2.2 Naphthalene derivatives: α,β -	
	Naphthols; α,β -Naphthylamines;	
	Schaeffer acid, Tobias acid;	

Q



	Naphthionic acid; N.W. acid; Clev-6- acid; H acid; Naphthol AS. 4.2.3 Anthracene derivatives: 1Nitroanthraquinone; 1Aminoanthraquinon e;2-Aminoanthraquinone; 2- Methylanthraquinone; anthraquinone-1- sulphonic acid; 1-Chloroanthraquinone; Chloroanthraquinone; Benzanthrone. 4.3 Dyeing Method of Cotton Fibres 4.3.1 Direct dyeing, Vat dyeing, Mordant dyeing, Disperse dyeing 4.3.2 Forces binding of dyes to the fibres: Ionic forces, Hydrogen bonds, Van- der-Wall's forces, Covalent linkages.	
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Semester V Practical

RUSACPDD501		Drugs & Dyes	Credits
		Drug preparation	
	1.	Preparation of Methyl Salicylate from Salicylic Acid	
	2.	To write the monograph of Paracetamol and Aspirin	
		from I.P.	
		Drug Estimation	
	1.	Estimation of Ibuprofen	02
	2.	Estimation of Acid neutralizing capacity of antacid	02
		Dyes Preparation	
	1.	Preparation of Orange-II	
	2.	Preparation of p-Nitroacetanilide from Acetanilide	
		Dyes Estimation	
	1.	Estimation of Primary amino group by diazotization	

References:

- 1. Medical Chemistry by V K Ahluwalia, Madhu Chopra, Ane's Books Pvt. Ltd.
- 2. Organic Chemistry of Drug Discovery and Drug Design Richard B. Silvermann
- 3. Medicinal Chemistry Shreeram and Yogeshwari (Pearson)
- 4. Chemistry of dyes and principles of dyeing, Shenai V.A., Sevak publications, 1973



MODALITY OF ASSESSMENT

Theory Examination Pattern:

A)	Internal Assessment 40%	40 Marks
Sr No	Evaluation type	Marks
1	One Assignment	10
2	One class Test (multiple choice questions / objective/ drawing	20
	structure of drugs and dyes)	
3	Active participation in class	05
4	Overall conduct, participation in curricular and co-curricular	05
	activities.	

B) External Examination – 60% (60 Marks) Semester End Theory Examination-

- i. Duration- These examinations shall be of two hours duration
- **ii. Theory question paper pattern**: There shall be **four questions** each of **15 marks**, one on each unit. All questions shall be compulsory with internal choice within the questions.

Questions	Options	Marks	Questions on	
Q.1) a)	Any 3 out of 5	12	Unit I	
Q.1) b)	Any 1 out of 2	03	Unit I	
Q.2) a)	Any 3 out of 5	12	Unit II	
Q.2) b)	Any 1 out of 2	03	Unit II	
Q.3) a)	Any 3 out of 5	12	Unit III	
Q.3) b)	Any 1 out of 2	03	Unit III	
Q.4) a)	Any 3 out of 5	12	Unit IV	
Q.4) b)	Any 1 out of 2	03	Unit Iv	

Practical Examination Pattern:

A) Internal Examination

Particulars	Marks
Journal	05
Experimental Work	30
Active Participation	05
Total	40



B) External Examination: Semester end practical examination 60 M

Sr.No.	Particulars	Marks
1)	Laboratory Work	25 + 25
2)	Viva- Voce	05 + 05
	Total	60

PRACTICAL BOOK / JOURNAL:

The students are required to present a duly certified journal for appearing at the practical examination, failing which they will not be allowed to appear for the examination.

In case of loss of Journal and/ or Report, a Lost Certificate should be obtained from Head/ Coordinator / In charge of the department; failing which the student will not be allowed to appear for the practical examination.

Overall Examination and Marks Distribution Pattern

Course	RUSACDD501		Total
	Internal	External	
Theory	40	60	100
Practical's	40	60	100



SEMESTER VI Course Code: RUSACDD601 <u>Course Title: Drugs & Dyes</u> Academic year 2020-21

Course Outcomes :

After s	After studying the course, the learner will be able to:		
CO 1	Outline the principles involved in drug designing and metabolism of drugs inside		
	the human body.		
CO 2	Classify various chemotherapeutic agents with respect to their chemical structure,		
	chemical class, therapeutic uses, and side effects		
CO 3	Compare the relation between color and chemical constitution of dyes.		
CO 4	Explore various applications of dyes.		
CO 5	Write the reactions involved in the synthesis of some representative drugs and dye		
	intermediates.		

DETAILED SYLLABUS

Course Code		Drugs & Dyes	Credits-02
RUSACDD601	Unit	Unit Title	Lectures
	Ι	1.1 Drug Discovery, Design and	(5L)
		Development	
		1.1.1 Discovery of a Lead compound:	
		Screening, drug metabolism studies and	
•	\sim	clinical observation.	
		1.1.2 Drug development from Natural Sources:	
. ()	•	Anti infective agents	
		Anti cancer agents	
		CNS agent	(4L)
		1.1.3 Development of drug:	
\mathcal{A}		The Pharmacophore identification,	
		modification of structure or functional	(6L)
U			



group, Structure activity relationship	0
(Benzodiazepines, Sulphonamides).	
1.1.4 Structure modification to increase	5
potency: Homologation, Chain branching,	
Ring-chain transformation, Extension of	
the structure.	
1.1.5 Computer assisted drug design.	
1.2 Drug Metabolism	
1.2.1. Introduction, Absorption, Distribution,	
Bio-transformation, Excretion.	
1.2.2. Different types of chemical	
transformation of drugs with specific example	
1.3 Chemotherapeutic Agents	
Study of the following chemotherapeutic agents	
with respect to their chemical structure,	
chemical class, therapeutic uses, and side	
effects.	
1.3.1 Antibiotics	
Definition, Characteristics and properties of:	
Amoxicillin; Cloxicillin (lactum antibiotics)	
Cephalexin (Cephalosporins), Doxycycline	
(Tetracyclines), Gentamycin	
(Aminoglycosides), Ciprofloxacin	
(Quinolones)	
Synthesis of Ciprofloxacin	
1.3.2 Antimalarials	
Types of malaria: Symptoms; pathological	
detection during window period (Life cycle of	
the parasites not o be discussed) Chloroquine	
(3-Amino quinolines) Paludrine (Biguanides)	
Pyrimethamine (Diamino pyrimidines)	
Artemether (Benzodioxepins)	
	 (Benzodiazepines, Sulphonamides). 1.1.4 Structure modification to increase potency: Homologation, Chain branching, Ring-chain transformation, Extension of the structure. 1.1.5 Computer assisted drug design. 1.2 Drug Metabolism 1.2.1. Introduction, Absorption, Distribution, Bio-transformation, Excretion. 1.2.2. Different types of chemical transformation of drugs with specific example 1.3 Chemotherapeutic Agents Study of the following chemotherapeutic agents with respect to their chemical structure, chemical class, therapeutic uses, and side effects. 1.3.1 Antibiotics Definition, Characteristics and properties of: Amoxicillin; Cloxicillin (lactum antibiotics) Cephalexin (Cephalosporins), Doxycycline (Tetracyclines), Gentamycin (Aminoglycosides), Ciprofloxacin (Quinolones) Synthesis of Ciprofloxacin 1.3.2 Antimalarials Types of malaria: Symptoms; pathological detection during window period (Life cycle of the parasites not o be discussed) Chloroquine (3-Amino quinolines) Paludrine (Biguanides)



	Following combination to be discussed	
	(i) Sulfadosine-Pyrimethamine	
	(ii) Atremether-Lumefantrine (no	
	structure)	
	Synthesis of Paludrine.	
II	2.1 Anti-inflammatory Drugs	(2L)
	2.1.1. Mechanism of inflammation and various	
	inflammatory conditions.	
	2.1.2. Prednisolone, Betamethasone (Steroids),	(3L)
	Aceclofenac (N-Aryl anthranilic acids),	
	Mefanic Acid (N-Aryl anthranilic	
	acids). Synthesis of Aceclofenac.	
	2.2 Antiamoebic Drugs	(2L)
	2.2.1. Types of Amoebiasis	
	2.2.2. Metronidazole; Diloxamide furoate	
	(Furans)	
	2.2.3. Following combination therapy to be	(3L)
	discussed:	
	Ciprofloxacin-Tinidazo	
	Synthesis of Metronidazole	
	2.3 Antitubercular Drugs	
	2.3.1. Types of Tuberculosis; Symptoms and	
	diagnosis of Tubeculosis.	(2L)
	2.3.2. General idea of Antibiotics used in their	
	treatment.	
	2.3.3. Streptomycin, Rifampin, PAS	
	(Aminosalicylates), Isoniazide	
	(Hydrazides),	
	Pyrazinamide (Pyrazines), (+) Ethambutol	(3L)
	(Aliphatic diamines)	
3	Synthesis of Ethambutol.	
	2.4 Antileprotic Drugs	



	2.4.1. Introduction, Types	0
	2.4.2 Classification of anti-leprotic agents	
	Ethionamide (Thioamides), Dapsone	
	(Sulfonamides), Clofazimine	
	(Phenazines)	
	Synthesis of Dapsone	(0)
	2.4.3. Following combination therapy to be	
	discussed for the treatment of	
	Tuberculosis and Leprosy:	
	(i) Rifampin + Ethambutol +	
	Pyrazinamide	
	(ii) Rifampin + Isoniazide +	
	Pyrazinamide	
	(iii) Rifampin + Clofazimine +	
	Ethionamide.	
	2.5 Drug Intermediates: Synthesis and uses	
	(i) 2,4,5-Triamino-6-hydroxypyrimidine	
	from Guanidine.	
	(ii) 3-Chloro-5-sulphonyl amino anthranilic	
	acid from 3-Chloro-2-toludine	
	(iii) p-[2'-(5-Chloro-2-methoxy	
	benzamido) ethyl]-	
	benzenesulphonamide from Methyl-5-	
2.0	chloro-2-methoxybenzoate	
	(iv) 4-(p-Chlorophenyl)-4-	
	hydroxypiperidine from 4-	
	Chloroacetophenone.	
20	(v) p-Acetyl amino benzenesulphonyl	
	chloride from Aniline	
	(vi) Epichlorohydrine from propene.	
	2.6 Nano particles in Medicinal Chemistry	



	2.6.1. Introduction, Carbon nano particles	Explore o Experience o Excel
	(structures), Carbon nano tubes:	
	Functionalisation for Pharmaceutical	
	applications. Targeted drug delivery in	
	vaccine (Foot and mouth disease) Use in	
	Bio-physical treatment.	
	2.6.2 Gold nano particles in treatment of cancer,	
	Parkinsonism, Alzheimer.	
	2.6.3. Silver nano particles: Antimicrobial	
	activity.	
III	3. 1 Colour and chemical constitution of dyes	(5L)
	3.1.1 Absorption of visible light, colour of	
	wavelength absorbed, complementary	
	colour.	
	3.1.2 Relation between colour and chemical	
	constitution. (i) Armstrong theory	(6L)
	(quinonoid theory) and its limitations (ii)	
	Valence Bond theory; Comparative	
	study and relation of colour in the	
	following classes of compounds/dyes:	(2L)
	Benzene, Nitrobenzene, Nitroanilines,	
	Nitrophenols, Benzoquinones, Azo,	
	Triphenyl methane, Anthraquinones (iii)	(2L)
	Molecular Orbital Theory.	
-0	3.2. Non-textile Uses of Dyes	
	Structural features of the substrate, fastness and	
	other property requirements and main classes of	
	dyes used to be mentioned as applicable. (Two	
	examples with structures for each of the	
	following.) 1. Leather 2. Paper 3. Foodstuff 4.	
	Cosmetics 5. Medicinal 6. Biological Stains 7.	



		Explore • Experience • Excel
	Indicator & Analytical Reagents 8. Coloured	
	Smokes & Camouflage colours 9. Laser Dyes.	
	3.3 Optical Brighteners	
	General idea and important characteristics of	
	optical brighteners, one example eachwith	
	structure of the following classes: Stilbene,	
	Coumarin, Heterocyclic vinylene derivatives,	
	Diaryl pyrazolines, Naphthalimide derivatives.	
	3.4 Organic Pigments	
	General idea, distinguish between dyes and	
	pigments, important characteristics of organic	
	pigments, Toners, Lakes, Classification of	
	organic pigments with suitable examples, i.e.	
	Ionic Pigments-Lake of acid and basic dyes.	
	Non-ionic pigments-Azo, Indigoid,	
	Anthraquinone, Quinacridone, Phthalocyanine	
	(Copper phthalocyanine).	
I	V 4.1 Synthesis of Specific Dyes and their Uses	(12L)
	i. Orange IV from sulphanilic acid	
	ii. Eriochrome Black T from β -	
	naphthol	
	iii. Eriochrome Red B by using ethyl	
•. •	aceto acetate and 1-amino-2-	
	naphthol-4-sulphonic Acid.	
s' 0-1	iv. Direct Deep Black EW by using	
	benzidine, H acid, aniline, and m-	
0	phenylen diamine.	
	v. Congo Red from nitrobenzene	
\sim	vi. Diamond Black F by using 5-amino	
	salicylic acid, N.W. acid and α -	
U.	naphthylamine.	
P		(1L)



	vii	Malachite Green by using
	V 11.	benzaldehyde and N,N- (2L)
		dimethylaniline.
	viii.	Auramine O from dimethylaniline
		Methylene Blue by using 4-amino-
		N,N-dimethylaniline and N,N-
		dimethylaniline
	х.	Safranine T by using o-toluidine and
		aniline
	xi.	Pararosaniline by using p-toluidine
		and aniline
	xii.	Alizarine Cyanine Green G by using
		phthalic anhydride and p-
		cholorophenol
	xiii.	Indanthrene from anthraquinone
	xiv.	Disperse Yellow 6G from
		benzanthrone
	XV.	Indigo from aniline
	xvi.	Eosine by using phthalic anhydride
		and resorcinol
	xvii.	Bismark Brown from m-
		phenylenediamine.
		s of Fibres and Classes of Dyes
	Applicabl	
		on to the following types of fibres
\$° O- '		tures and classes of dyes applicable to
		Wool, Silk, Polyester.
	-	gy and Toxicity of Dyes
		ence to the textile dyes, food colours,
	benzidine	etc.
AN		
Y		
		Page 23



Semester VI Practical RUSACPDD601 Drugs & Dyes Credits **Drug Preparation** Preparation of Aspirin from Salicylic Acid 1. **Drug Estimation** Estimation of Tincture of Iodine 1. **Dye Preparation** Preparation of m-dinitrobenzene 1. 02 2. Preparation of m-nitroaniline **Dye Estimation** Estimation of Methyl Orange/ Eriochrome Black 1. T/Eosin/Congo Red by colorimetry Dyeing of fabric (cotton)by Direct Dyeing or by Vat Dyeing.

References:

- 1. Medical Chemistry by V K Ahluwalia, Madhu Chopra, Ane's Books Pvt. Ltd.
- 2. Organic Chemistry of Drug Discovery and Drug Design Richard B. Silvermann
- 3. Medicinal Chemistry Shreeram and Yogeshwari (Pearson)
- 4. Chemistry of dyes and principles of dyeing, Shenai V.A., Sevak publications, 1973

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MODALITY OF ASSESSMENT

Theory Examination Pattern:

A)	Internal Assessment 40%	40 Marks
Sr No	Evaluation type	Marks
1	One Assignment	10
2	One class Test (multiple choice questions / objective/ drawing structure of drugs and dyes)	20
3	Active participation in class	05
4	Overall conduct, participation in curricular and co-curricular activities.	05

B) External Examination – 60% (60 Marks)

Semester End Theory Examination-

iii.Duration- These examinations shall be of two hours duration

iv. Theory question paper pattern: There shall be four questions each of 15 marks, one on each unit. All questions shall be compulsory with internal choice within the questions.

Questions	Options	Marks	Questions on	
Q.1) a)	Any 3 out of 5	12	Unit I	
Q.1) b)	Any 1 out of 2	03	Unit I	
Q.2) a)	Any 3 out of 5	12	Unit II	
Q.2) b)	Any 1 out of 2	03	Unit II	
Q.3) a)	Any 3 out of 5	12	Unit III	
Q.3) b)	Any 1 out of 2	03	Olint III	
Q.4) a)	Any 3 out of 5	12	Unit IV	
Q.4) b)	Any 1 out of 2	03	Unit IV	

Practical Examination Pattern:

A) Internal Examination

Particulars	Marks
Journal	05
Experimental Work	30
Active Participation	05
Total	40



B) External Examination: Semester end practical examination

(60 Marks)

Sr.No.	Particulars	Marks
1)	Laboratory Work	25 +25
2)	Viva- Voce	05 + 05
	Total	60

PRACTICAL BOOK / JOURNAL:

The students are required to present a duly certified journal for appearing at the practical examination, failing which they will not be allowed to appear for the examination.

In case of loss of Journal and/ or Report, a Lost Certificate should be obtained from Head/ Coordinator / In charge of the department; failing which the student will not be allowed to appear for the practical examination.

Overall Examination and Marks Distribution Pattern

Course	RUSACDD601		Total
	Internal	External	
Theory	40	60	100
Practical's	40	60	100