2016

THE MASTER OF PHARMACY (M. PHARM.) COURSE REGULATION 2014

(Based on notification in the Gazette of India No. 362, Dated December 11, 2014)

SCHEME AND SYLLABUS



PHARMACY COUNCIL OF INDIA

Combined Council's Building, Kotla Road, Aiwan-E-Ghalib Marg, New Delhi-110 002. Website: www.pci.nic.

COURSE STRUCTURE AND SYLLABUS For M. PHARM

MPH R 18 Regulations

(Applicable for batches admitted from 2018-2019)



JAWAHARLAL NEHRU TECHNOLOGICAL UNIVERSITY: KAKINADA KAKINADA - 533 003, Andhra Pradesh, India

Table of Contents

S.No.	Content	Page.No.
S.NO.	Regulations	05
1.	Short Title and Commencement	05
2.	Minimum qualification for admission	05
3.	Duration of the program	05
4.	Medium of instruction and examinations	05
5.	Working days in each semester	05
6.	Attendance and progress	05
7.	Program/Course credit structure	05
8.	Academic work	06
9.	Course of study	06
10.	Program Committee	18
11.	Examinations/Assessments	18
12.	Promotion and award of grades	30
13.	Carry forward of marks	30
14.	Improvement of internal assessment	30
15.	Reexamination of end semester examinations	30
16.	Allowed to keep terms (ATKT)	31
17.	Grading of performances	31
18.	The Semester grade point average (SGPA)	31
19.	Cumulative Grade Point Average (CGPA)	32
20.	Declaration of class	32
21.	Project work	32
22.	Award of Ranks	33
23.	Award of degree	33
24.	Duration for completion of the program of study	33
25.	Revaluation I Retotaling of answer papers	33
26.	Re-admission after break of study	33
27.	Pharmaceutics (MPH)	34
28.	Industrial Pharmacy (MIP)	51
29.	Pharmaceutical Chemistry (MPC)	66
30.	Pharmaceutical Analysis (MPA)	84
31.	Pharmaceutical Quality Assurance (MQA)	102
32.	Pharmaceutical Regulatory Affairs (MRA)	120
33.	Pharmaceutical Biotechnology (MPB)	140
34.	Pharmacy Practice (MPP)	158
35.	Pharmacology (MPL)	176
36.	Pharmacognosy (MPG)	195
37.	Research Methodology & Biostatistics (MRM)	213



असाधारण

EXTRAORDINARY

भाग III-खण्ड 4

PART III - Section 4

प्रधिकार से प्रकाशित PUBLISHED BY ALTHORITY

Tl. 3621 No. 3621 नई दिल्ली, बहरमतिवार, दिसम्बर 11, 2014/अग्रहायण 20, 1936

NEW DELHI, THURSDAY, DECEMBER 11, 2014/AGRAHAYANA 20, 1936

PHARMACY COUNCIL OF INDIA

NOTIFICATION

New Delhi, the 10th December, 2014

The Master of Pharmacy (M.Pharm) Course Regulations, 2014

No. 14-136/ 2014-PCL—In exercise of the powers conferred by Sections 10 and 18 of the Pharmacy Act, 1948 (8 of 1948), the Pharmacy Council of India, with the approval of the Central Government hereby makes the following regulations; namely—

<u>CHAPTER –I: REGULATIONS</u>

1. Short Title and Commencement

These regulations shall be called as "The Revised Regulations for the Master of Pharmacy (M. Pharm.) Degree Program - Credit Based Semester System (CBSS) of the Pharmacy Council of India, New Delhi". They shall come into effect from the Academic Year 2016-17. The regulations framed are subject to modifications from time to time by the authorities of the university.

2. Minimum qualification for admission

A Pass in the following examinations

- a) B. Pharm Degree examination of an Indian university established by law in India from an institution approved by Pharmacy Council of India and has scored not less than 55 % of the maximummarks (aggregate of 4 years of B. Pharm.)
- b) Every student, selected for admission to post graduate pharmacy program in any PCI approved institution should have obtained registration with the State Pharmacy Council or should obtain the same within one month from the date of his/her admission, failing which the admission of the candidate shall be cancelled.

Note: It is mandatory to submit a migration certificate obtained from the respective university where the candidate had passed his/her qualifying degree (B.Pharm.)

3. Duration of the program

The program of study for M.Pharm. shall extend over a period of four semesters (two academic years). The curricula and syllabi for the program shall be prescribed from time to time by Phamacy Council of India, New Delhi.

4. Medium of instruction and examinations

Medium of instruction and examination shall be in English.

5. Working days in each semester

Each semester shall consist of not lessthan 100 working days. The odd semesters shall be conducted from the month of June/July to November/December and the even semesters shall be conducted from the month of December/January to May/June in every calendar year.

6. Attendance and progress

A candidate is required to put in at least 80% attendance in individual courses considering theory and practical separately. The candidate shall complete the prescribed course satisfactorily to be eligible to appear for the respective examinations.

7. Program/Course credit structure

As per the philosophy of Credit Based Semester System, certain quantum of academic work viz. theory classes, practical classes, seminars, assignments, etc. are measured in terms of credits. On satisfactory completion of the courses, a candidate earns credits. The amount of credit associated with a course is dependent upon the number of hours of instruction per week in that course. Similarly the credit associated with any of the other academic, co/extra- curricular activities is dependent upon the quantum of work expected to be put in for each of these activities per week/per activity.

7.1. Credit assignment

7.1.1. Theory and Laboratory courses

Courses are broadly classified as Theory and Practical. Theory courses consist of lecture (L) and Practical (P) courses consist of hours spent in the laboratory. Credits (C) for a course is dependent on the number of hours of instruction per week in that course, and is obtained by using a multiplier of one (1) for lecture and a multiplier of half (1/2) for practical (laboratory) hours. Thus, for example, a theory course having four lectures per week throughout the semester carries a credit of 4. Similarly, a practical having four laboratory hours per week throughout semester carries acredit of 2.

The contact hours of seminars, assignments and research work shall be treated as that of practical courses for the purpose of calculating credits. i.e., the contact hours shall be multiplied by 1/2. Similarly, the contact hours of journal club, research work presentations and discussions with the supervisor shall be considered as theory course and multiplied by 1.

7.2. Minimum credit requirements

The minimum credit points required for the award of M. Pharm. degree is 95. However based on the credit points earned by the students under the head of co-curricular activities, a student shall earn a maximum of 100 credit points. These credits are divided into Theory courses, Practical, Seminars, Assignments, Research work, Discussions with the supervisor, Journal club and Co-Curricular activities over the duration of four semesters. The credits are distributed semester-wise as shown in Table 14. Courses generally progress in sequence, building competencies and their positioning indicates certain academic maturity on the part of the learners. Learners are expected to follow the semester-wise schedule of courses given in the syllabus.

8. Academic work

A regular record of attendance both in Theory, Practical, Seminar, Assignment, Journal club, Discussion with the supervisor, Research work presentation and Dissertation shall be maintained by the department/teaching staff of respective courses.

9. Course of study

The specializations in M.Pharm program is given in Table 1.

Table – 1: List of M.Pharm. Specializations and their Code

S. No.	Specialization	Code
	Specialization	Code
1.	Pharmaceutics	MPH
2.	Industrial Pharmacy	MIP
3.	Pharmaceutical Chemistry	MPC
4.	Pharmaceutical Analysis	MPA
5.	Pharmaceutical Quality Assurance	MQA
6.	Pharmaceutical Regulatory Affairs	MRA
7.	Pharmaceutical Biotechnology	MPB
8.	Pharmacy Practice	MPP
9.	Pharmacology	MPL
10.	Pharmacognosy	MPG

The course of study for M.Pharm specializations shall include Semester wise Theory & Practical as given in Table -2 to 11. The number of hours to be devoted to each theory and practical course in any semester shall not be less than that shown in Table -2 to 11.

Table – 2: Course of study for M. Pharm. (Pharmaceutics)

Course Code	Course	Credit Hours	Credit Points	Hrs./ wk	Marks
	Seme	ster I			
MPH101T	Modern Pharmaceutical Analytical Techniques	4	4	4	100
MPH102T	Drug Delivery System	4	4	4	100
MPH103T	Modern Pharmaceutics	4	4	4	100
MPH104T	Regulatory Affair	4	4	4	100
MPH105PA	Pharmaceutics Practical I	6	3	6	75
MPH105PB	Pharmaceutical Practical II	6	3	6	75
-	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650
	Seme	ster II			
MPH201T	Molecular Pharmaceutics (Nano Tech and Targeted DDS)	4	4	4	100
MPH202T	Advanced Biopharmaceutics & Pharmacokinetics	4	4	4	100
MPH203T	Computer Aided Drug Delivery System	4	4	4	100
MPH204T	Formulation Development of Pharmaceutical and Cosmetic Products	4	4	4	100
МРН205РА	Pharmaceutics Practical	6	3	6	75
MPH205PB	Pharmaceutics Practical IV	6	3	6	75
-	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650

Table – 3: Course of study for M. Pharm. (Industrial Pharmacy)

Course Code	ole – 3: Course of study for M. Pha Course	Credit Hours	Credit Points	Hrs./ wk	Marks		
	Semester I						
MIP101T	Modern Pharmaceutical Analytical Techniques	4	4	4	100		
MIP102T	Pharmaceutical Formulation Development	4	4	4	100		
MIP103T	Novel drug delivery systems	4	4	4	100		
MIP104T	Intellectual Property Rights	4	4	4	100		
MIP105PA	Industrial Pharmacy Practical I	6	3	6	75		
MIP105PB	Industrial Pharmacy Practical II	6	3	6	75		
-	Seminar/Assignment	7	4	7	100		
	Total	35	26	35	650		
	Semesto	er II					
MIP201T	Advanced Biopharmaceutics and Pharmacokinetics	4	4	4	100		
MIP202T	Scale up and Technology Transfer	4	4	4	100		
MIP203T	Pharmaceutical Production Technology	4	4	4	100		
MIP204T	Entrepreneurship Management	4	4	4	100		
MIP205PA	Industrial Pharmacy Practical III	6	3	6	75		
MIP205PB	Industrial Pharmacy Practical IV	6	3	6	75		
-	Seminar/Assignment	7	4	7	100		
	Total	35	26	35	650		

Table – 4: Course of study for M. Pharm. (Pharmaceutical Chemistry)

Table – 4: Course of study for M. Pharm. (Pharmaceutical Chemistry)								
Course	Ca.,,,,,,	Credit	Credit	Hrs./	Maulia			
Code	Course	Hours	Points	wk	Marks			
	Semester I							
1 1/10/11/11	Modern Pharmaceutical Analytical Techniques	4	4	4	100			
MPC1012T	Advanced Organic Chemistry -I	4	4	4	100			
MPC1031	Advanced Medicinal chemistry	4	4	4	100			
MPC104T	Chemistry of Natural Products	4	4	4	100			
MPC105PA	Pharmaceutical Chemistry Practical I	6	3	6	75			
MPC105PB	Pharmaceutical Chemistry Practical II	6	3	6	75			
-	Seminar/Assignment	7	4	7	100			
	Total	35	26	35	650			
	Seme	ster II						
MPC201T	Advanced Spectral Analysis	4	4	4	100			
	Advanced Organic Chemistry -II	4	4	4	100			
MPC203T	Computer Aided Drug Design	4	4	4	100			
MPCOMAT	Pharmaceutical Process Chemistry	4	4	4	100			
MPC205PA	Pharmaceutical Chemistry Practical III	6	3	6	75			
МРС105РВ	Pharmaceutical Chemistry Practical IV	6	3	6	75			
-	Seminar/Assignment	7	4	7	100			
	Total	35	26	35	650			

Table – 5: Course of study for M. Pharm. (Pharmaceutical Analysis)

Course Code	– 5: Course of study for M. Phai Course	Credit Hours	Credit Points	Hrs./wk	Marks		
	Semester I						
MPA101T	Modern Pharmaceutical Analytical Techniques	4	4	4	100		
MPATOTI	Advanced Pharmaceutical Analysis	4	4	4	100		
MPA103T	Pharmaceutical Validation	4	4	4	100		
MPA104T	Food Analysis	4	4	4	100		
MPA105PA	Pharmaceutical Analysis Practical I	6	3	6	75		
MPA105PB	Pharmaceutical Analysis Practical II	6	3	6	75		
-	Seminar/Assignment	7	4	7	100		
	Total	35	26	35	650		
	Semes	ter II					
MPA201T	Advanced Instrumental Analysis	4	4	4	100		
MPA202T	Modern Bio-Analytical Techniques	4	4	4	100		
MPA203T	Quality Control and Quality Assurance	4	4	4	100		
MPA204T	Herbal and Cosmetic Analysis	4	4	4	100		
MPA205PA	Pharmaceutical Analysis Practical III	6	3	6	75		
MPA205PB	Pharmaceutical Analysis Practical IV	6	3	6	75		
-	Seminar/Assignment	7	4	7	100		
	Total	35	26	35	650		

Table – 6: Course of study for M. Pharm. (Pharmaceutical Quality Assurance)

Course Code	Course of study for M. Pharm. (F	Credit Hours	Credit Points	Hrs./wk	
	Semes	ster I			
MQA101T	Modern Pharmaceutical Analytical Techniques	4	4	4	100
MQA102T	Quality Management System	4	4	4	100
MQA103T	Quality Control and Quality Assurance	4	4	4	100
MQA104T	Product Development and Technology Transfer	4	4	4	100
MQA105PA	Pharmaceutical Quality Assurance Practical I	6	3	6	75
MQA105PB	Pharmaceutical Quality Assurance Practical II	6	3	6	75
-	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650
	Semes	ter II			
MQA201T	Hazards and Safety Management	4	4	4	100
MQA202T	Pharmaceutical Validation	4	4	4	100
MQA203T	Audits and Regulatory Compliance	4	4	4	100
MQA204T	Pharmaceutical Manufacturing Technology	4	4	4	100
MQA205PA	Pharmaceutical Quality Assurance Practical III	6	3	6	75
MQA205PB	Pharmaceutical Quality Assurance Practical IV	6	3	6	75
-	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650

Table – 7: Course of study for M. Pharm. (Regulatory Affairs)

Course Code	Course	Credit Hours	Credit Points	Hrs./ wk	Marks
	Seme	ster I			
MRA101T	Good Regulatory Practices	4	4	4	100
MRA102T	Documentation and Regulatory Writing	4	4	4	100
MRA103T	Clinical Research Regulations	4	4	4	100
MRA104T	Regulations and Legislation for Drugs & Cosmetics, Medical Devices, Biologicals & Herbals, and Food & Nutraceuticals In India and Intellectual Property Rights	4	4	4	100
MRA105PA	Regulatory Affairs Practical I	6	3	6	75
MRA105PB	Regulatory Affairs Practical II	6	3	6	75
	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650
	Seme	ster II			
MRA201T	Regulatory Aspects of Drugs & Cosmetics	4	4	4	100
MRA202T	Regulatory Aspects of Herbal & Biologicals	4	4	4	100
MRA203T	Regulatory Aspects of Medical Devices	4	4	4	100
MRA204T	Regulatory Aspects of Food & Nutraceuticals	4	4	4	100
MRA205PA	RegulatoryAffairsPracticalIII	6	3	6	75
MRA205PB	Regulatory Affairs Practical IV	6	3	6	75
	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650

Table – 8: Course of study for M. Pharm. (Pharmaceutical Biotechnology)

Course	S: Course of study for M. Pharm. Course	Credit	Credit	Hrs./	Marks
Code	Seme	Hours	Points	wk	
		ster i			
MPB101T	Modern Pharmaceutical Analytical Techniques	4	4	4	100
MPB102T	Microbial And Cellular Biology	4	4	4	100
MPB103T	Bioprocess Engineering and Technology	4	4	4	100
MPB104T	Advanced Pharmaceutical Biotechnology	4	4	4	100
MPB105PA	Pharmaceutical Biotechnology Practical I	6	3	6	75
МРВ105РВ	Pharmaceutical Biotechnology Practical II	6	3	6	75
-	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650
	Semes	ter II			
MPB201T	Proteins and protein Formulation	4	4	4	100
MPB202T	Immunotechnology	4	4	4	100
MPB203T	Bioinformatics and Computer Technology	4	4	4	100
MPB204T	Biological Evaluation of Drug Therapy	4	4	4	100
МРВ205РА	Pharmaceutical Biotechnology Practical III	6	3	6	75
МРВ205РВ	Pharmaceutical Biotechnology Practical IV	6	3	6	75
-	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650

Table – 9: Course of study for M. Pharm. (Pharmacy Practice)

Course Code	ble – 9: Course of study for M. Ph Course	Credit Hours	Credit Points	Hrs./wk	Marks
	Semest	er I			
MPP101T	Clinical Pharmacy Practice	4	4	4	100
MPP102T	Pharmacotherapeutics-I	4	4	4	100
MPP103T	Hospital & Community Pharmacy	4	4	4	100
MPP104T	Clinical Research	4	4	4	100
MPP105PA	Pharmacy Practice Practical I	6	3	6	75
MPP105PB	Pharmacy Practice Practical II	6	3	6	75
-	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650
	Semeste	er II			
MPP201T	Principles of Quality Use of Medicines	4	4	4	100
MPP102T	Pharmacotherapeutics II	4	4	4	100
MPP203T	Clinical Pharmacokinetics and Therapeutic Drug Monitoring	4	4	4	100
MPP204T	Pharmacoepidemiology & Pharmacoeconomics	4	4	4	100
MPP205PA	Pharmacy Practice Practical III	6	3	6	75
MPP205PB	Pharmacy Practice Practical IV	6	3	6	75
-	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650

Table – 10: Course of study for (Pharmacology)

Course Code	Course	Credit Hours	Credit Points	Hrs./wk	Marks
	Seme	ster I			
MPL101T	Modern Pharmaceutical Analytical Techniques	4	4	4	100
MPL102T	Advanced Pharmacology-I	4	4	4	100
MPL103T	Pharmacological and Toxicological Screening Methods-I	4	4	4	100
MPL104T	Cellular and Molecular Pharmacology	4	4	4	100
MPL105PA	Pharmacology Practical I	6	3	6	75
MPL105PB	Pharmacology Practical II	6	3	6	75
-	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650
	Semes	ster II			
MPL201T	Advanced Pharmacology II	4	4	4	100
MPL202T	Pharmacological and Toxicological Screening Methods-II	4	4	4	100
MPL203T	Principles of Drug Discovery	4	4	4	100
MPL204T	Experimental Pharmacology practical- II	4	4	4	100
MPL205PA	Pharmacology Practical III	6	3	6	75
MPL205PB	Pharmacology Practical IV	6	3	6	75
-	Seminar/Assignment	7	4	7	100
	Total	35	26	35	650

Table – 11: Course of study for M. Pharm. (Pharmacognosy)

Course Code	Course	Credit Hours	Credit Points	Hrs./wk	Marks		
	Semester I						
MPG101T	Modern Pharmaceutical Analytical Techniques	4	4	4	100		
MPG102T	Advanced Pharmacognosy-1	4	4	4	100		
MPG103T	Phytochemistry	4	4	4	100		
MPG104T	Industrial Pharmacognostical Technology	4	4	4	100		
MPG105PA	Pharmacognosy Practical I	6	3	6	75		
MPG105PB	Pharmacognosy Practical II	6	3	6	75		
-	Seminar/Assignment	7	4	7	100		
	Total	35	26	35	650		
	Semes	ter II					
MPG201T	Medicinal Plant biotechnology	4	4	4	100		
MPG102T	Advanced Pharmacognosy-II	4	4	4	100		
MPG203T	Indian system of medicine	4	4	4	100		
MPG204T	Herbal cosmetics	4	4	4	100		
MPG205PA	Pharmacognosy Practical III	6	3	6	75		
MPG205PB	Pharmacognosy Practical IV	6	3	6	75		
-	Seminar/Assignment	7	4	7	100		
	Total	35	26	35	650		

Table – 12: Course of study for M. Pharm. III Semester (Common for All Specializations)

Course Code	Course	Credit Hours	Credit Points
MRM301T	Research Methodology and Biostatistics*	4	4
-	J ournal club	1	1
-	Discussion / Presentation (Proposal Presentation)	2	2
-	Research Work	28	14
	Total	35	21

^{*} Non University Exam

Table – 13: Course of study for M. Pharm. IV Semester (Common for All Specializations)

Course Code	Course	Credit Hours	Credit Points
-	Journal Club	1	1
-	Research Work	31	16
-	Discussion/Final Presentation	3	3
	Total	35	20

Table – 14: Semester wise credits distribution

Semester	Credit Points						
	26						
II	26						
III	21						
IV	20						
Co-curricular Activities (Attending Conference, Scientific Presentationsand Other Scholarly Activities)	Minimum=02 Maximum=07*						
Total Credit Points	Minimum=95 Maximum=100*						

^{*}Credit Points for Co-curricular Activities

Table – 15: Guidelines for Awarding Credit Points for Co-curricular Activities

Table – 13. Guidelines for Awarding Credit Points for Co-curricular Activities							
Name of the Activity	Maximum Credit Points Eligible / Activity						
Participation in National Level Seminar/Conference/Workshop/Symposium/ Training Programs (related to the specialization of the student)	01						
Participation in international Level Seminar/Conference/Workshop/Symposium/ Training Programs (related to the specialization of the student)	02						
Academic Award/Research Award from State Level/National Agencies	01						
Academic Award/Research Award from International Agencies	02						
Research / Review Publication in National Journals (Indexed in Scopus / Web of Science)	01						
Research / Review Publication in International Journals (Indexed in Scopus / Web of Science)	02						

Note: International Conference: Held outside India; International Journal: The Editorial Board Outside India

10. Program Committee

The M. Pharm. programme shall have a Programme Committee constituted by the Head of the Institution in consultation with all the Heads of the departments.

The composition of the Programme Committee shall be as follows:

A teacher at the cadre of Professor shall be the Chairperson; One Teacher from each M.Pharm specialization and four student representatives (two from each academic year), nominated by the Head of the institution.

Duties of the Programme Committee:

Periodically reviewing the progress of the classes.

Discussing the problems concerning curriculum, syllabus and the conduct of classes.

Discussing with the course teachers on the nature and scope of assessment for the course and the same shall be announced to the students at the beginning of respective semesters.

- l. Communicating its recommendation to the Head of the Institution on academic matters.
- 2 The Programme Committee shall meet at least twice in a semester preferably at the end of each sessional exam and before the end semester exam.

11. Examinations/Assessments

The schemes for internal assessment and end semester examinations are given from Table-16.

11.1. End semester examinations

The End Semester Examinations for each theory and practical course through semesters I to IV shall be conducted by the respective university except for the subject with asterix symbol (*) for which examinations shall be conducted by the subject experts at college level and the marks/grades shall be submitted to the university.

^{*}The credit points assigned for extracurricular and or co-curricular activities shall be given by the Principals of the colleges and the same shall be submitted to the University. The criteria to acquire this credit point shall be defined by the colleges from time to time.

Tables – 16: Schemes for internal assessments and end semester (Pharmaceutics- MPH)

]	nternal A	ssessment		End So Ex	T-4-1		
Course Code	Course	Continues	Session	nal Exams	Total	Marks	Durati	Total Marks	
		Mode	Marks	Duration	Totai	Warks	on		
		SEMI	ESTER I						
MPH101T	Modern Pharmaceutical Analytical Techniques	10	15	1Hr	25	75	3Hr	100	
MPH102T	Drug Delivery Systems	10	15	1Hr	25	75	3Hr	100	
MPH103T	Modern Pharmaceutics	10	15	1Hr	25	75	3Hr	100	
MPH104T	Regulatory Affairs	10	15	1Hr	25	75	3Hr	100	
MPH105PA	Pharmaceutics Practical I	10	15	3Hr	25	50	3Hr	75	
МРН105РВ	Pharmaceutics Practical II	10	15	3Hr	25	50	3Hr	75	
-	Seminar/Assignment	-	-	-	-	-	-	100	
		Total						650	
		SEME	STER II						
MPH201T	Molecular Pharmaceutics (Nano Tech and Targeted DDS)	10	15	1 Hr	25	75	3Hr	100	
MPH202T	Advanced Biopharmaceutics & Pharmacokinetics	10	15	1Hr	25	75	3Hr	100	
MPH203T	Computer Aided Drug Delivery System	10	15	1Hr	25	75	3Hr	100	
MPH204T	Formulation Development of Pharmaceutical and Cosmetic Products	10	15	1Hr	25	75	3Hr	100	
MPH205PA	Pharmaceutics Practical I	10	15	3Hr	25	50	3Hr	75	
МРН205РВ	Pharmaceutics Practical I	10	15	3Hr	25	50	3Hr	75	
-	Seminar/Assignment	-	-	-	-	-	-	100	
	Total								

Tables – 17: Schemes for internal assessments and end semester (Industrial Pharmacy- MIP)

	- 17: Schemes for internal			sessment	,	End S				
Course Code	Course	Continues	Session	nal Exams	Total	Marks	Duration	Total Marks		
		Mode	Marks	Duration	Total	WILKS	Burucion			
SEMESTER I										
MIP101T	Modern Pharmaceutical Analytical Techniques	10	15	1Hr	25	75	3Hr	100		
MIP102T	Pharmaceutical Formulation Development	10	15	1Hr	25	75	3Hr	100		
MIP103T	Novel Drug Delivery Systems	10	15	1Hr	25	75	3Hr	100		
MIP104T	Intellectual Property rights	10	15	1Hr	25	75	3Hr	100		
MIP105PA	Industrial Pharmacy Practical I	10	15	3Hr	25	50	3Hr	75		
MIP105PB	Industrial Pharmacy Practical II	10	15	3Hr	25	50	3Hr	75		
-	Seminar/Assignment	-	-	-	-	-	-	100		
		Total						650		
		SEME	STER II							
MIP201T	Advanced Biopharmaceutics and Pharmacokinetics	10	15	1 Hr	25	75	3Hr	100		
MIP202T	Scale up and Technology Transfer	10	15	1Hr	25	75	3Hr	100		
MIP203T	Pharmaceutical Production Technology	10	15	1 Hr	25	75	3Hr	100		
MIP204T	Entrepreneurship Management	10	15	1Hr	25	75	3Hr	100		
MIP205PA	Industrial Pharmacy Practical III	10	15	3Hr	25	50	3Hr	75		
MIP205PB	Industrial Pharmacy Practical IV	10	15	3Hr	25	50	3Hr	75		
-	Seminar/Assignment	-	-	-	-	-	-	100		
		Total						650		

 $Tables-18: Schemes \ for \ internal \ assessments \ and \ end \ semester \ (Pharmaceutical \ Chemistry-MPC)$

			ernal Ass	sessment		End S	Total Marks			
Course Code	Course	Continues	Session	Sessional Exams		Marks		Duration		
		Mode	Marks	Duration	Total	Warks	Duration			
SEMESTER I										
MPC101T	Modern Pharmaceutical Analytical Techniques	10	15	1Hr	25	75	3Hr	100		
MPC102T	Advanced Organic Chemistry – I	10	15	1Hr	25	75	3Hr	100		
MPC103T	Advanced Medicinal Chemistry	10	15	1Hr	25	75	3Hr	100		
MPC104T	Chemistry of Natural Products	10	15	1Hr	25	75	3Hr	100		
MPC105PA	Pharmaceutical chemistry Practical I	10	15	3Hr	25	50	3Hr	75		
MPC105PB	Pharmaceutical chemistry Practical II	10	15	3Hr	25	50	3Hr	75		
	Seminar/Assignment	-	-	-	-	-	-	100		
		Total						650		
		SEME	STER II							
MPC201T	Advanced Spectral Analysis	10	15	1Hr	25	75	3Hr	100		
MPC202T	Advanced Organic Chemistry II	10	15	1Hr	25	75	3Hr	100		
MPC203T	Computer Aided Drug Design	10	15	1Hr	25	75	3Hr	100		
MPC204T	Pharmaceutical Process Chemistry	10	15	1 Hr	25	75	3Hr	100		
MPC205PA	Pharmaceutical chemistry Practical III	10	15	3Hr	25	50	3Hr	75		
MPC205PB	Pharmaceutical chemistry Practical IV	10	15	3Hr	25	50	3Hr	75		
	Seminar/Assignment	-	-	-	-	-	-	100		
		Total						650		

 $Tables-19: Schemes \ for \ internal \ assessments \ and \ end \ semester \ (Pharmaceutical \ Analysis-MPA)$

	1	1V1	PA)						
		Int	ernal Ass	sessment		End S E			
Course Code	Course	Continues	Session	nal Exams	m . 1			Total Marks	
		Mode	Marks	Duration	Total	Marks	Duration		
SEMESTER I									
MPA101T	Modern Pharmaceutical Analytical Techniques	10	15	1Hr	25	75	3Hr	100	
MPA102T	Advanced Pharmaceutical Analysis	10	15	1Hr	25	75	3Hr	100	
MPA103T	Pharmaceutical Validation	10	15	1Hr	25	75	3Hr	100	
MPA104T	Food Analysis	10	15	1Hr	25	75	3Hr	100	
MPA105PA	Pharmaceutical Analysis Practical I	10	15	3Hr	25	50	3Hr	75	
MPA105PB	Pharmaceutical Analysis Practical II	10	15	3Hr	25	50	3Hr	75	
	Seminar/Assignment	-	-	-	-	-	-	100	
		Total						650	
		SEME	STER II						
MPA201T	Advanced Instrumental Analysis	10	15	1 Hr	25	75	3Hr	100	
MPA202T	Modern Bio-Analytical Techniques	10	15	1Hr	25	75	3Hr	100	
MPA203T	Quality Control and Quality Assurance	10	15	1Hr	25	75	3Hr	100	
MPA204T	Herbal and Cosmetic Analysis	10	15	1Hr	25	75	3Hr	100	
MPA205PA	Pharmaceutical Analysis Practical III	10	15	3Hr	25	50	3Hr	75	
MPA205PB	Pharmaceutical Analysis Practical IV	10	15	3Hr	25	50	3Hr	75	
	Seminar/Assignment	-	-	-	-	-	-	100	
		Total						650	

 $Tables-20: Schemes \ for \ internal \ assessments \ and \ end \ semester \ (Pharmaceutical \ Quality \\ Assurance- \ MQA)$

		Assuran	ernal Ass	,		End Semester Exams		
Course Code	Course	Continues	Session	nal Exams				Total Marks
		Mode	Marks	Duration	Total	Marks	Duration	
		SEMI	ESTER I					
MQA101T	Modern Pharmaceutical Analytical Techniques	10	15	1Hr	25	75	3Hr	100
MQA102T	Quality Management System	10	15	1Hr	25	75	3Hr	100
MQA103T	Quality Control and Quality Assurance	10	15	1Hr	25	75	3Hr	100
MQA104T	Product Development and Technology Transfer	10	15	1Hr	25	75	3Hr	100
MQA105PA	Pharmaceutical Quality Assurance Practical I	10	15	3Hr	25	50	3Hr	75
MQA105PB	Pharmaceutical Quality Assurance Practical II	10	15	3Hr	25	50	3Hr	75
	Seminar/Assignment	-	-	-	-	-	-	100
		Total						650
		SEME	STER II					
MQA201T	Hazards and Safety Management	10	15	1Hr	25	75	3Hr	100
MQA202T	Pharmaceutical Validation	10	15	1Hr	25	75	3Hr	100
MQA203T	Audits and Regulatory Compliance	10	15	1Hr	25	75	3Hr	100
MQA204T	Pharmaceutical Manufacturing Technology	10	15	1Hr	25	75	3Hr	100
MQA205PA	Pharmaceutical Quality Assurance Practical III	10	15	3Hr	25	50	3Hr	75
MQA205PB	Pharmaceutical Quality Assurance Practical IV	10	15	3Hr	25	50	3Hr	75
	Seminar/Assignment	-	-	-	-	-	-	100
Total								650

 $Tables-21: Schemes \ for \ internal \ assessments \ and \ end \ semester \ (Pharmaceutical \ Regulatory \ Affairs-MRA)$

		In	ternal As	sessment			Semester xams	
Course Code	Course	Continues Mode	Session Marks	nal Exams Duration	Total	Marks	Duration	Total Marks
		SEMES	TER I					
MRA101T	Good Regulatory Practices	10	15	1Hr	25	75	3Hr	100
MRA102T	Documentation and Regulatory Writing	10	15	1Hr	25	75	3Hr	100
MRA103T	Clinical Research Regulations	10	15	1Hr	25	75	3Hr	100
MRA104T	Regulations and Legislations for Drugs & Cosmetics, Medical Devices, Biologicals & Herbals, and Food & Nutraceuticals in India and Intellectual Property Rights	10	15	1Hr	25	75	ЗНг	100
MRA105PA	Regulatory Affairs Practicals I	10	15	3Hr	25	50	3Hr	75
MRA105PB	Regulatory Affairs Practicals II	10	15	3Hr	25	50	3Hr	75
	Seminar/Assignment	-	-	-	-	-	-	100
		Total						650
		SEMEST	ER II					
MRA201T	Regulatory Aspects of Drugs and Cosmetics	10	15	1Hr	25	75	3Hr	100
MRA202T	Regulatory Aspects of Herbal & Biologicals	10	15	1Hr	25	75	3Hr	100
MRA203T	Regulatory Aspects of Medical Devices	10	15	1Hr	25	75	3Hr	100
MRA204T	Regulatory Aspects of Food Neutraceuticals	10	15	1Hr	25	75	3Hr	100
MRA205PA	Regulatory Affairs Practicals III	10	15	3Hr	25	50	3Hr	75
MRA205PB	Regulatory Affairs Practicals IV	10	15	3Hr	25	50	3Hr	75
	Seminar/Assignment	-	-	-	-	-	-	100
		Total						650

 $Tables-22: Schemes \ for \ internal \ assessments \ and \ end \ semester \ (Pharmaceutical \ Biotechnology-MPB)$

G		Int	ernal Ass	essment			Semester xams	Total
Course Code	Course	Continues	Session	nal Exams	Total	Marks	Duration	Total Marks
		Mode	Marks	Duration	Total	WILLIAM	Duration	
		SEM	ESTER I					
MPB101T	Modern Pharmaceutical Analytical Techniques	10	15	1Hr	25	75	3Hr	100
MPB102T	Microbial and Cellular Biology	10	15	1Hr	25	75	3Hr	100
MPB103T	Bioprocess Engineering and Technology	10	15	1Hr	25	75	3Hr	100
MPB104T	Advanced Pharmaceutical Biotechnology	10	15	1Hr	25	75	3Hr	100
MPB105PA	Pharmaceutical Biotechnology Practical I	10	15	3Hr	25	50	3Hr	75
MPB105PB	Pharmaceutical Biotechnology Practical II	10	15	3Hr	25	50	3Hr	75
	Seminar/Assignment	-	-	-	-	-	-	100
		Total						650
		SEMI	ESTER II	I				
MPB201T	Proteins and Protein Formulation	10	15	1Hr	25	75	3Hr	100
MPB202T	Immunotechnology	10	15	1Hr	25	75	3Hr	100
MPB203T	Bioinformatics and Computer Technology	10	15	1Hr	25	75	3Hr	100
MPB204T	Biological Evaluation of Drug Therapy	10	15	1Hr	25	75	3Hr	100
MPB205PA	Pharmaceutical Biotechnology Practical III	10	15	3Hr	25	50	3Hr	75
MPB205PB	Pharmaceutical Biotechnology Practical IV	10	15	3Hr	25	50	3Hr	75
	Seminar/Assignment	-	-	-	-	-	-	100
		Total						650

Tables – 23: Schemes for internal assessments and end semester (Pharmacy Practice- MPP)

G		Int	ternal Ass	sessment		End S Ex	Total		
Course Code	Course	Continues	Session	nal Exams	Total	Mada	Duration	Total Marks	
		Mode	Marks	Duration	Totai	Marks	Duration		
		SEME	STER I						
MPP101T	Clinical Pharmacy Practice	10	15	1Hr	25	75	3Hr	100	
MPP102T	Pharmacotherapeutics - I	10	15	1Hr	25	75	3Hr	100	
MPP103T	Hospital & Community Pharmacy	10	15	1Hr	25	75	3Hr	100	
MPP104T	Clinical Research	10	15	1Hr	25	75	3Hr	100	
MPP105PA	Pharmacy Practice Practical I	10	15	3Hr	25	50	3Hr	75	
MPP105PB	Pharmacy Practice Practical II	10	15	3Hr	25	50	3Hr	75	
	Seminar/Assignment	-	-	-	-	-	-	100	
		Total						650	
		SEMES	STER II						
MPP201T	Principles of Quality Use of Medicines	10	15	1Hr	25	75	3Hr	100	
MPP202T	Pharmacotherapeutics - II	10	15	1Hr	25	75	3Hr	100	
MPP203T	Clinical Pharmacokinetics and Therapeutic Drug Monitoring	10	15	1Hr	25	75	3Hr	100	
MPP204T	Pharmacoepidemiology & Pharmacoeconomics	10	15	1Hr	25	75	3Hr	100	
MPP205PA	Pharmacy Practice Practical III	10	15	3Hr	25	50	3Hr	75	
MPP205PB	Pharmacy Practice Practical IV	10	15	3Hr	25	50	3Hr	75	
	Seminar/Assignment	-	-	-	-	-	-	100	
	Total								

Tables – 24: Schemes for internal assessments and end semester (Pharmacology- MPL)								
		Internal Assessment				End Semester Exams		
Course Code	Course	Continues	Session	nal Exams	Total	Marks	Duration	Total Marks
		Mode	Marks	Duration	Total	Marks	Duration	
SEMESTER I								
MPL101T	Modern Pharmaceutical Analytical Techniques	10	15	1Hr	25	75	3Hr	100
MPL102T	Advanced Pharmacology - I	10	15	1Hr	25	75	3Hr	100
MPL103T	Pharmacology and Toxicology Screening methods- I	10	15	1Hr	25	75	3Hr	100
MPL104T	Cellular and Molecular Pharmacology	10	15	1Hr	25	75	3Hr	100
MPL105PA	Pharmacology Practical I	10	15	3Hr	25	50	3Hr	75
MPL105PB	Pharmacology Practical II	10	15	3Hr	25	50	3Hr	75
	Seminar/Assignment	-	-	-	-	-	-	100
		Total						650
		SEME	STER II					
MPL201T	Advanced Pharmacology - II	10	15	1Hr	25	75	3Hr	100
MPL202T	Pharmacology and Toxicology Screening methods- II	10	15	1Hr	25	75	3Hr	100
MPL203T	Principles of Drug Discovery	10	15	1Hr	25	75	3Hr	100
MPL204T	Experimental Pharmacology Practical II	10	15	1Hr	25	75	3Hr	100
MPL205PA	Pharmacology Practical III	10	15	3Hr	25	50	3Hr	75
MPL205PB	Pharmacology Practical IV	10	15	3Hr	25	50	3Hr	75
	Seminar/Assignment	-	-	-	-	-	-	100

Total

Table	es – 25: Schemes for interr	nal assessme	nts and	end seme	ster (P	harmaco	ognosy- M	PG)
		Int	ternal Ass	sessment			Semester xams	
Course Code	Course	Continues	Session	Sessional Exams		Marks		Total Marks
		Mode	Marks	Duration	Total	lviaiks	Duration	
		SEMI	ESTER I					
MPG101T	Modern Pharmaceutical Analytical Techniques	10	15	1Hr	25	75	3Hr	100
MPG102T	Advanced Pharmacognosy - I	10	15	1Hr	25	75	3Hr	100
MPG103T	Phytochemistry	10	15	1Hr	25	75	3Hr	100
MPG104T	Industrial Pharmacognostical Technology	10	15	1Hr	25	75	3Hr	100
MPG105PA	Pharmacognosy Practical I	10	15	3Hr	25	50	3Hr	75
MPG105PB	Pharmacognosy Practical II	10	15	3Hr	25	50	3Hr	75
	Seminar/Assignment	-	-	-	-	-	-	100
		Total						650
		SEME	ESTER II					
MPG201T	Medicinal Plant Biotechnology	10	15	1Hr	25	75	3Hr	100
MPG202T	Advanced Pharmacognosy - II	10	15	1Hr	25	75	3Hr	100
MPG203T	Indian system of Medicine	10	15	1Hr	25	75	3Hr	100
MPG204T	Herbal Cosmetics	10	15	1Hr	25	75	3Hr	100
MPG205PA	Pharmacognosy Practical III	10	15	3Hr	25	50	3Hr	75
MPG205PB	Pharmacognosy Practical IV	10	15	3Hr	25	50	3Hr	75
	Seminar/Assignment	-	-	-	-	-	-	100
		Total						650

Tables – 26: Schemes for interna	l assessments anden o	dsemesterexaminations	(Semester III& IV)

140105 20	s. Schemestor internal	Internal Assessment			End Semester Exams			
Course Code	Course	Conti		ssional Exams	Tot	Mark	Durati	Total Marks
		nuous Mode	Mark s	Durati on	al	S	on	
		SEI	MESTE	R III				
MRM30 1T	Research Methodology and Biostatistics*	10	15	1 Hr	25	75	3 Hrs	100
-	Journal club				25	·	-	25
_	Discussion / Presentation (Proposal Presentation)			-	50		·	50
-	Research work*					350	1 Hr	350
		To	tal					525
		SEN	MESTE	R IV				
-	J ournal club	·			25	-	-	25
-	Discussion / Presentation (Proposal Presentation)	-			75	·	·	75
	Research work and Colloquium		-		-	400	1 Hr	400
Total							500	

^{*}Non University Examination

11.2. Internal assessment: Continuous mode

The marks allocated for Continuous mode of Internal Assessment shall be awarded as per the scheme given below.

Table – 27: Scheme for awarding internal assessment: Continuous mode

Theory				
Criteria	Maximum Marks			
Attendance (Refer Table – 28)	8			
Student – Teacher interaction	2			
Total	10			
Practical				
Attendance (Refer Table – 28)	10			
Based on Practical Records, Regular viva voce, etc.	10			
Total	20			

Table – 28: Guidelines for the allotment of marks for attendance

Percentage of Attendance	Theory	Practical
95 – 100	8	10
90 – 94	6	7.5
85 – 89	4	5
80 - 84	2	2.5
Less than 80	0	0

11.2.1. Sessional Exams

Two sessional exams shall be conducted for each theory / practical course as per the schedule fixed by the college(s). The scheme of question paper for theory and practical sessional examinations is given in the table. The average marks of two sessional exams shall be computed for internal assessment as per the requirements given in tables.

12. Promotion and award of grades

A student shall be declared PASS and eligible for getting grade in a course of M.Pharm.programme if he/she secures at least 50% marks in that particular courseincluding internal assessment.

13. Carry forward of marks

In case a student fails to secure the minimum 50% in any Theory or Practical course as specified in 12, then he/she shall reappear for the end semester examination of that course. However his/her marks of the Internal Assessment shall be carried over and he/she shall be entitled for grade obtained by him/her on passing.

14. Improvement of internal assessment

A student shall have the opportunity to improve his/her performance only once in the sessional exam component of the internal assessment. The re-conduct of the sessional exam shall be completed before the commencement of next end semester theory examinations.

15. Reexamination of end semester examinations

Reexamination of end semester examination shall be conducted as per the schedule given in table 29. The exact dates of examinations shall be notified from time to time.

Table – 29: Tentative schedule of end semester examinations

Semester	For Regular Candidates	For Failed Candidates
I and III	November / December	May / June
II and IV	May / June	November / December

16. Allowed to keep terms (ATKT):

No student shall be admitted to any examination unless he/she fulfills the norms given in 6. ATKT rules are applicable as follows:

A student shall be eligible to carry forward all the courses of I and IIsemesters till the III semester examinations. However, he/she shall not be eligible to attend the courses of IV semester until all the courses of I, II and III semesters are successfully completed.

A student shall be eligible to get his/her CGPA upon successful completion of the courses of I to IV semesters within the stipulated time period as per the norms.

Note: Grade AB should be considered as failed and treated as one head for deciding ATKT. Such rules are also applicable for those students who fail to register for examination(s) of any course in any semester.

17. Grading of performances

17.1. Letter grades and grade points allocations:

Based on the performances, each student shall be awarded a final letter grade at the end of the semester for each course. The letter grades and their corresponding grade points are given in Table -30.

Table–30: Letter grades and grade points equivalent to Percentage of marks and performances.

Percentage of Marks Obtained	Letter Grade	Grade Point	Performance
90.00 – 100	0	10	Outstanding
80.00 – 89.99	A	9	Excellent
70.00 – 79.99	В	8	Good
60.00 – 69.99	С	7	Fair
50.00 - 59.99	D	6	Average
Less than 50	F	0	Fail
Absent	AB	0	Fail

A learner who remains absent for any end semester examination shall be assigned a letter grade of AB and a corresponding grade point of zero. He/she should reappear for the said evaluation/examination in due course.

18. The Semester grade point average (SGPA)

The performance of a student in a semester is indicated by a number called 'Semester Grade Point Average' (SGPA). The SGPA is the weighted average of the grade points obtained in all the courses by the student during the semester. For example, if a student takes five courses (Theory/Practical) in a semester with credits C1, C2, C3 and C4 and the student's grade points in these courses are G1, G2, G3 and G4, respectively, and then students' SGPA is equal to:

$$SGPA = \begin{array}{c} C_1G_1 + C_2G_2 + C_3G_3 + C_4G_4 \\ \\ C_1 + C_2 + C_3 + C_4 \end{array}$$

The SGPA is calculated to two decimal points. It should be noted that, the SGPA for any semester shall take into consideration the F and ABS grade awarded in that semester. For example if a learner has a F or ABS grade in course 4, the SGPA shall then be computed as:

$$SGPA = \begin{array}{c} C_1G_1 + C_2G_2 + C_3G_3 + C_4* \ ZERO \\ \\ C_1 + C_2 + C_3 + C_4 \end{array}$$

19. Cumulative Grade Point Average (CGPA)

The CGPA is calculated with the SGPA of all the IV semesters to two decimal points and is indicated in final grade report card/final transcript showing the grades of all IV semesters and their courses. The CGPA shall reflect the failed statusin case of F grade(s), till the course(s) is/are passed. When the course(s) is/are passedby obtaining a pass grade on subsequent examination(s) the CGPA shall only reflect the new grade and not the fail grades earned earlier. The CGPA is calculated as:

$$CGPA = \begin{array}{c} C_1S_1 + C_2S_2 + C_3S_3 + C_4S_4 \\ \\ C_1 + C_2 + C_3 + C_4 \end{array}$$

where C_1 , C_2 , C_3 ,... is the total number of credits for semester I,II,III,... and S_1 , S_2 , S_3 ,... is the SGPA of semester I,II,III,....

20. Declaration of class

The class shall be awarded on the basis of CGPA as follows:

First Class with Distinction = CGPA of 7.50 and above First Class = CGPA of 6.00 to 7.49 Second Class = CGPA of 5.00 to 5.99

21. Project work

All the students shall undertake a project under the supervision of a teacher in Semester III to IV and submit a report. 4 copies of the project report shall be submitted (typed & bound copy not less than 75 pages).

The internal and external examiner appointed by the University shall evaluate the project at the time of the Practical examinations of other semester(s). The projects shall be evaluated as per the criteria given below.

Evaluation of Dissertation Book:

Objective(s) of the work done		50Marks
Methodologyadopted		150 Marks
Results and Discussions		250 Marks
Conclusions and Outcomes		50 Marks
	Total	500 Marks

Evaluation of Presentation:

Presentation of work		100 Marks
Communicationskills		50 Marks
Question and answer skills		100 Marks
	Total	250 Marks

22. Award of Ranks

Ranks and Medals shall be awarded on the basis of final CGPA. However, candidates whofail in one or more courses during the M.Pharm program shall not be eligible for award of ranks. Moreover, the candidates should have completed the M. Pharm program in minimum prescribed number of years, (two years) for the award of Ranks.

23. Award of degree

Candidates who fulfill the requirements mentioned above shall be eligible for award of degree during the ensuing convocation.

24. Duration for completion of the program of study

The duration for the completion of the program shall be fixed as double the actual duration of the program and the students have to pass within the said period, otherwise they have to get fresh Registration.

25. Revaluation I Retotaling of answer papers

There is no provision for revaluation of the answer papers in any examination. However, the candidates can apply for retotaling by paying prescribed fee.

26. Re-admission after break of study

Candidate who seeks re-admission to the program after break of study has to get the approval from the university by paying a condonation fee.

Website: www.jntuk.edu.in Email: dap@jntuk.edu.in



Phone: 7032894555

Directorate of Academic Planning

JAWAHARLAL NEHRU TECHNOLOGICAL UNIVERSITY KAKINADA KAKINADA-533003, Andhra Pradesh, INDIA

(Established by AP Government Act No. 30 of 2008)

Lr. No. JNTUK/DAP/RAC/I Year/M.Pharmacy/2022-23

Date: 08-12-2022

Dr. KVSG Murali Krishna.

M.E. Ph D.

Director, Academic Planning JNTUK, Kakinada

All the Principals of Affiliated Colleges.

JNTUK, Kakinada.

Revised Academic Calendar of I Year M. Pharmacy Academic year 2022-23

LSEME	STER		
Description	From	To	Weeks
Commencement of Class Work	12.12.2022	,	
Induction Classes	12.12.2022	17.11.2022	1 W
l Unit of Instruction	19:12.2022	11.02,2023	8 W
1 Mid Examinations	06.02.2023	11.02.2023	
Il Unit of Instructions	13.02.2023	08.04.2023	874
II Mid Examinations	03.04.2023	08.04.2023	
Preparation & Practicals	10.04.2023	15.04.2023	1W
End Examinations	17.04.2023	29.04.2023	5 May
Commencement of Il Semester Class Work	01.05.2023		
H SEME	STER		
Commencement of Class Work	01.05.2023		
I Unit of Instructions	01.05.2023	24.06.2023	SW
I Mid Examinations	26.06.2023	24.06.2023	
If Unit of Instructions	26.06.2023	19.08.2023	877
II Mid Examinations	14.08.2022	19.08.2023	
Preparation & Practicals	21.08.2023	26,08,2023	1.W
End Examinations	28.08.2023	10.09.2023	2 W
Commencement of Class Work	12,09,2023		

Director Academics & Planning JNTUK Kakinac

femic Planning MYUK Kakinada

to the Hon'ble Vice Chancellor, JNTUK Copy to the

Copy to P Copy to

JNTUK

Copy to

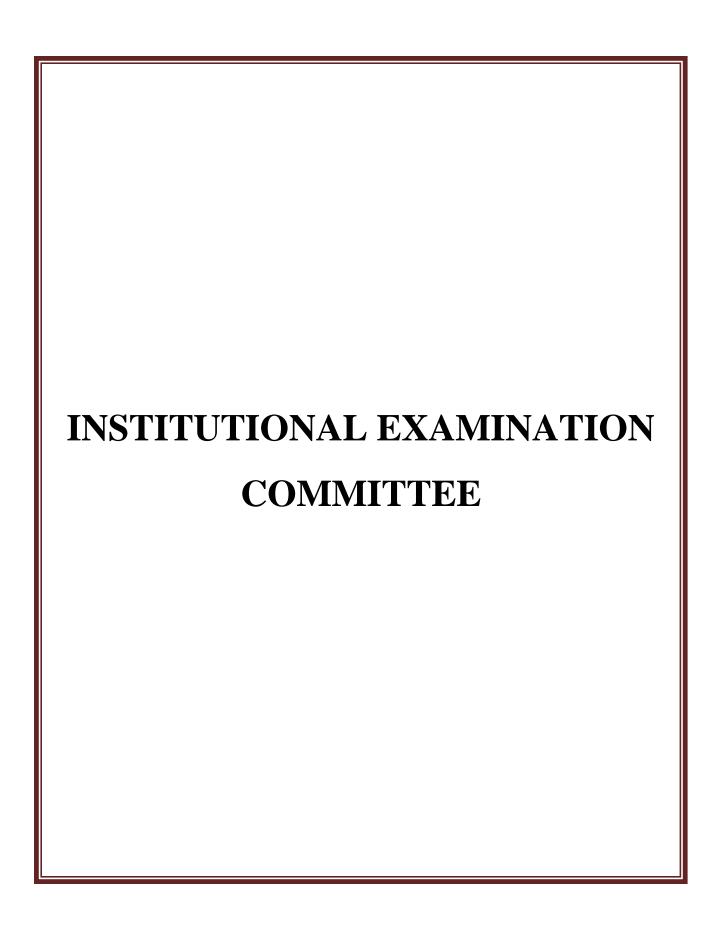
Audit, JNTUK

Copy to

on, JNTUK

VIJAYA INSTITUTE OF PHARMACEUTICAL SCIENCES FOR WOMEN

NIKEPADU, VIJAYAWADA 528 198



VIJAYA INSTITUTE OFPHARMACEUTICAL SCIENCES FOR WOMEN Enikepadu, Vijayawada – 521108

Date: 26-07-2021

OFFICE ORDER

INSTITUTIONAL EXAMINATION COMMITTEE

The Institutional Examination Committee for the academic year 2021 - 2022 is constituted as follows and it is effective for a period of 06-09-2021 to 06-08-2022. Following staff members are appointed as Institutional Examination Committee.

S.NO	NAME	DESIGNATION	POSITION	SIGNATURE
1	Dr. K. Padmalatha	Principal	Chairman	Matts
2	Mr. S. Venkateswara Rao	Assoc. Professor	College Examination	C. Vertiserla
e solo ch			Officer	
3	Mr. A. Jayarami Reddy	Assoc. Professor	Member	Anady
4	Mrs. A.V.S. Hima bindu	Asst. Professor	Member	148
5	Dr. N. Prathibha	Asst. Professor	Member	Postiti
6	Dr. S. Sundar	Professor	Member	124

Functions and Responsibilities:

- 1. Ensure proper dissemination of information with regard to examination among all the stakeholders' viz. students / faculty / non teaching staff / university authorities etc.
- 2. Receive and submission of exam notification / schedule from JNTUK web portal.
- 3. To ensure proper organization of in semester assessments / sessional / end semester examinations in the college.
- 4. Ensure proper communication with JNTUK with regards to examination and fulfillment of university circulars.
- 5. Appoint alternative external senior supervisor / chairman / internal examiners / external examiners for conduct of end semester theory / practical examination with permission of university authorities.
- 6. Record and issue the answer books and other exam related stationary to the invigilators / internal examiners 30 minutes before start the exam
- 7. Download and print the appropriate number of question papers at least 20 minutes before the commencement of the exam and maintaining absolute confidentiality
- 8. Resolve students / faculty / university grievances with regards to examinations.
- 9. Uploading internal theory / practical examination marks on JNTUK web portal.

ENIKEPADU

10. Maintain records with regards to conduct of examination and results.

Copy to: 1. Establishment File

2. Concerned Faculty member

DIA Padmalatha
PARINCIPAL

VIJAYA INSTITUTE OF

PHARMACEUTICAL SCIENCES FOR WOMEN
ENIKEPADU, VIJAYAWADA - 521 108



JAWAHARLAL NEHRU TECHNOLOGICAL UNIVERSITY KAKINADA

UNIVERSITY EXAMINATION CENTER, KAKINADA

M. PHARMACY I SEMESTER (PCI REGULATION) I MID EXAMINATIONS, FEBRUARY - 2023

TIME TABLE

TIME: 10:00 AM TO 12:00 NOON

ENIKEPADU, VIJAYAWADA - 521 108

BRANCH & SPECIALIZATION	06-02-2023 (Monday)	07-02-2023 (Tuesday)	08-02-2023 (Wednesday)	09-02-2023 (Thursday)
PHARMACEUTICAL CHEMISTRY (02)	Modern Pharmaceutical Analytical Techniques (MPC101T)	Advanced Organic Chemistry –I (MPC102T)	Advanced Medicinal Chemistry (MPC103T)	Chemistry of Natural Products (MPC104T)
PHARMACEUTICS (03)	Modern Pharmaceutical Analytical Techniques (MPH101T)	Drug Delivery Systems (MPH102T)	Modern Pharmaceutics (MPH103T)	Regulatory Affairs (MPH104T)
PHARMACOLOGY (06)	Modern Pharmaceutical Analytical Techniques (MPL101T)	Advanced Pharmacology-I (MPL102T)	Pharmacological and Toxicological Screening Methods-I (MPL103T)	Cellular and Molecular Pharmacolog (MPL104T)
PHARMACOGNOSY (07)	Modern Pharmaceutical Analytical Techniques (MPG101T)	Advanced Pharmacognosy-1 (MPG102T)	Phytochemistry (MPG103T)	Industrial Pharmacognostical Technology (MPG104T)
PHARMACY PRACTICE (08)	Clinical Pharmacy Practice (MPP101T)	Pharmacotherapeutics-I (MPP102T)	Hospital & Community Pharmacy (MPP103T)	Clinical Research (MPP104T)
INDUSTRIAL PHARMACY (09)	Techniques (MIP1011)	Development (MIP102T) KEPADU KEPADU	Novel drug delivery systems (MIP103T)	Intellectual Property Rights (MIP104T) PAL
	Termina de la companya della companya della companya de la companya de la companya della company	A * Wallet	VIJAYA INST	LULE OF
		A * *	PHARMACEUTICAL SCIE	NCES FOR WOMEN

	97, 60		4 174 4 4 4 4	
BRANCH & SPECIALIZATION		07-02-2023 (Tuesday)	08-02-2023 4 WC (Wednesday)	
PHARMACEUTICAL REGULATORY AFFAIRS (13)	Good Regulatory Practices (MRA101T)	Documentation and Regulatory Writing (MRA 102T)	Clinical Research Regulations (MRA103T)	Regulations and Legislation for Drugs & Cosmetics, Medical Devices, Biologicals & Herbals, and Food & Nutraceuticals In India and Intellectual Property Rights (MRA104T)
PHARMACY QUALITY ASSURANCE (15)	Modern Pharmaceutical Analytical Techniques (MQA101T)	Quality Management System (MQA102T)	Quality Control and Quality Assurance (MQA103T)	Product Development and Technology Transfer (MQA104T)
PHARMA CEUTICAL ANALYSIS (16)	Modern Pharmaceutical Analytical Techniques (MPA101T)	Advanced Pharmaceutical Analysis (MPA102T)	Pharmaceutical Validation (MPA103T)	Food Analysis (MPA1041)

- NOTE: (i) If Government declares holiday on any of the above dates, the examinations will be conducted as usual
 - (ii) Any omissions or clashes in this Time Table may please be informed to the Controller of Examinations immediately.
 - (iii) The Principals are requested to inform the University, if any other substitute subjects that are not included in the above time table immediately

Date: 27-01-2023

ENIKEPADU

VIJAYA INSTITUTE OF PHARMACEUTICAL SCIENCES FOR WOMEN ENIKEPADU. VIJAYAWADA-521 108.

Controller of Examinations (PG)

VIJAYA INSTITUTE OF PHARMCEUTICAL SCIENCES FOR WOMEN ENIKEPADU, VIJAYAWADA – 521108.

I M. Pharm I Sem I Mid Exams Invigilation Duties, Feb-2023

Timings : 10:00 AM TO 12:00 PM

Exam Dates	Staff Name	Staff Signature
06.02.2023 (Monday)	Mrs. K. V. R. Rajeswari	Tring
07.02.2023 (Tuesday)	Ms. B. Lekhya	Libbur
08.02.2023 (Wednesday)	Dr. B. Dhanush	Ph
09.02.2023 (Thursday)	Mrs. K. V. R. Rajeswari	/ Luigh

Exams Incharge

(Dr. S. Venkateswara Rao) EXAMS-INCHARGE

VIJAYA INSTITUTE

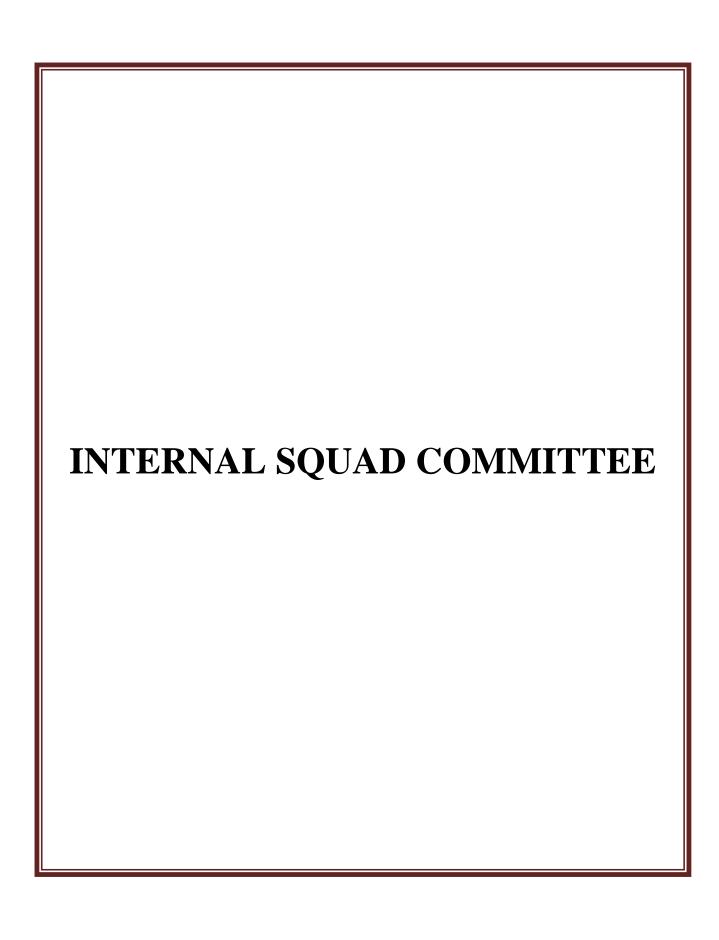
HARMACEUTICAL SCIENCES FOR WOMEN MIKEPADU VIJAYAWADA 821 108

Principal

(Dr. K. Padmalatha)

VIJAYA INSTITUTE OF

PHARMACEUTICAL SCIENCES FOR WOMEN ENIKEPADU, VIJAYAWADA - 521 168



VIJAYA INSTITUTE OFPHARMACEUTICAL SCIENCES FOR WOMEN Enikepadu, Vijayawada – 521108

Date: 26-07-2021

OFFICE ORDER

INTERNAL SQUAD COMMITTEE

The Internal Squad Committee has been constructed for smooth conduct of sessional / end semester examinations for the academic year 2021 – 2022 for the period of 06-09-2021 to 06-08-2022. Following staff members are appointed as Internal Squad Committee.

S.NO	NAME	DESIGNATION	POSITION	SIGNATURE
1	Dr. K. Padmalatha	Principal	President	(alts
2	Mr. S. Venkateswara Rao	Assoc. Professor	Chairman	S. Vanduk
3	Mr. A. Jayarami Reddy	Asst. Professor	Member	Meason
4	Mrs. A.V.S. Hima bindu	Asst. Professor	Member	HE
5	Mrs. Ch. Anupama Swathi	Asst. Professor	Member	A

Responsibilities:

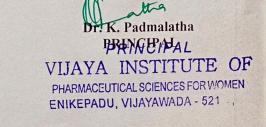
- 1. Strict checking of unfair means is sole responsibility of members of committee.
- 2. Before the start of examination, the committee members should check every student.
- 3. Care should be taken by committee members, that the students should not carry mobile phones, calculator or any sort of electronic material inside the examination hall.
- 4. Check weather students are carrying hall tickets by committee members to maintain environment of examination. Any issue related to the unfair means should immediately report to the principal or college examination officer.

aceutica

ENIKEPADU

Copy to: 1. Establishment File

2. Concerned Faculty member



I MID

ATTENDANCE SHEET FOR I MID EXAMINATIONS

COURSE: M. Pharm

Date of Examination: 08.02.23

Time: 10.00 AM TO 12.00 PM

Room No: 01

Subject Name: Pharmacological & Toxicological Screening Methods-I

Subject Code: MPL103T

No. of Students Present: 02

No. of Students Absent: 0

S.No.	Hall Ticket No.	Name of the Student	Answer Booklet Serial No.	Signature of the Student
1	227N1S0601	BOYALAPALLI PRASANNA	7N220001	B. Pasanna
2	227N1S0602	CHALAMALA RAMYANJALI		CH Ramyarjali
3	227N1S0603	SHAIK HAFSA	7N220003	- ABSENT -

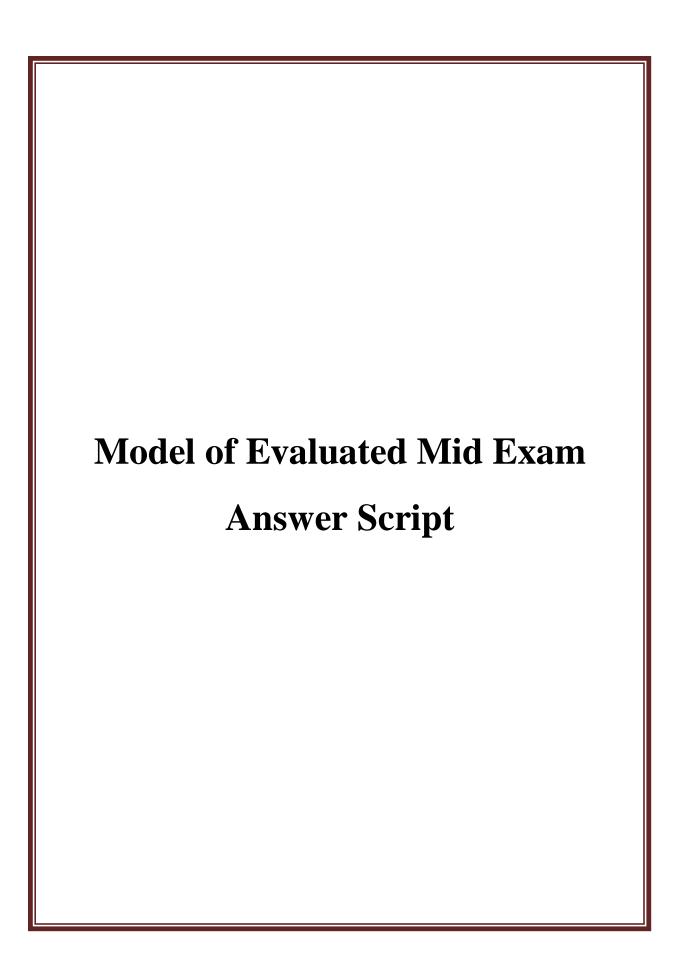
Signature of the Invigilator: B- Dhaery

Name of the Invigilator: B. DHANUSH.

Designation: ASCT. PROFESSOR

Signature of the Principal PRINCIPAL VIJAYA INSTITUTE O.

PHARMACEUTICAL SCIENCES FOR WOMEN UKEPADU, VIJAYAWADA - 521 108



SRK FOUNDATION'S

VIJAYA INSTITUTE OF PHARMACEUTICAL SCIENCES FOR WOMEN

ENIKEPADU, VIJAYAWADA



2022 - 2023

SESSIONAL BOOK

Name

: B. Pasanna

Class

: 1st M. Pharmacy [pharmacology dept].

Roll No.

: 227NIS0601

Subject

: Pharmacological & Toxicological Screening models-I

Internal	Objective	Subjective	Assignment	Total	Staff Sign	Student Sign
1		21		21	KIKSU	B. Pagganna
II		2342	_	23/2	(MED)	B. Almana

Final Average: 22

NELNCERAL Staff Sign

HOD Sign

I M. Pharmacy | I sem I mid Examination Su: Pharmacol cal & Toxicolo cal Screenin Predinical Evaluation of drugs 131 Alzheimer's disease. Invitro 1)-Adenylcyclase activity In vivo 1) Step down a) Two Compartment Test 3) Radial arm mage Study 4) Water maze Study Step down Mice 81 but either Sex Test Std | Test compound orally Animal is placed on a platform containing tectangular box with floor grids, these grids are attached to Shocking device to deliver the foot shock. Measure Step down assay latency period.

After finished the experiment the animal Step down 31 remain is recorded.

Iwo Compartment Test Mice 81 rat either Sex Et should contain soxsocm rectangular box with temoval ceiting 35° cm this is connected with Small box 15x15 cm with black walk. These two Compartments has transparent Slight doll is present between of the compastments Illuminating with 100 w bulb at the centre of large Compartment

latency is measured.

Radial arm mage Study

chice of rat efther sex

apparature is wooden, consists lelevated as 8 dadial arms mage with 56 long length, som wide, 2 cm height

-Animal is placed on the apparatur & made is attached to make.

During test animal food should be provide per day at once, west should be maintained at 85.

Trained the animals to run in mage to catch the Pellets.

Training is terminated after & choices, the animal obtained maximum of pellets with minimum of extens

Errors is measure.

Water make Study water tank filled with 20 depth a water tank filled with 20 depth a Apparatur is circular & it is divided I distributed into 4 parts Served as Starting point, with

Thes apparatus is divided into 4 quardants. Small avandant is fixed in the any one of the quardant

The quardant (small) is placed on entire training secons.

The animal is allow in water upto 60-90 sec. to findout the quardent.

Trained rate are identify the greatdant in less than

After experiment, quardant is removed & allow the animal in water to swim for 30 seconds.

	Screening Procedures for parasympathomimetic
	an vitro
	D'Guinea pig Bleum.
	2) Isolated eye of rodent.
	3) To detect anticholine sterase activity.
	2n vivo
	I) To detect anticholinesterase activity in Kat.
	Guinea pig aleum
-	Gainea pig (290-500 gm)
	Killed by Stanning
	Abdomen cut & mid ihasion
	T.
	Remove pancreas Bntestine
	Entertine is cut half I pass through glass is inserted
	înto î⊢ ·
	o o o o the free of
-	Tyrode Solution is passed through & efficient the free of
	Substances.
-	Cut the Pieces into (2-3) cm.

one side piece is attached to tissue clamp & insented Into 10-20 ml of tisue bath & attached to wanter writer lever mas produces confractions 10- do times. J Ach (0.01mg) is added. Et produces 70-901. of maximal responses To defect anticholinesterase activity Thiolester Ach is used as Substrate To detect the Inhibita of engyme activity 2.89 ml of phosphate buffer. 10 ml Sample. O.IMI DINB. All are maded & incubate for 10 minutes Add Substrate & measure the spectrometer by using Spectrometry. lo defect anticholinesterage in cat Cat (1-2 kg) anesthesized with pentobarbitone.

Test- compound DV administer.

Caratoid artery is canulated to detect BP

L

occur different doses at chokinesterase activity
as follows.

at dose X: No detectable effects

to 10-20 mm ttg.

at dose 4x: Fall in BP to 150-100 mm Hg

- 1) Pritial 1se in respiration & Use in respiration.
- 2) Salivary & Branchial Secretions Seen.
- 3) Urnation -

Screening procedures for parkinson's disease

In vitro

DExperiments using rate Straital Slices.

- 2) Dopamine Stimulated adenyla anticholinesterate activity En vivo
- D Tremonine & oxetremonine in rate mice.
- 2) MPTP model in monteys.
- 3) Reservine antagonism in mice
- 4) 6- Hydroxy dopamine induced neostraital lises in rate

(18-22g) Male CMRI mice.
- 10 O - 10
Test compound I stranded is given orally.
1
After 24 hrs Oxitremonine (0.15 mgkg) Subcutaneously
administer.
1
Rectal Temperature is measured before administration
& 1, 2, 4, 6 hrs after administration.
<i>T</i>
Record the Signs like tremors, Stupus etc
PTP model in monkeys
8 Adult thesus monteys
(5-8 kg)
. _
Over a period of 8-10 days &u doses in mptp is administered.
is administered.
↓
Developed parkinson's Symptoms.
[small
This is reversible, when test compound L-Dopa is administered
This is reverable, when text comp
L - Dopa is administered
\checkmark
Check the actions [signs.

	Resempine antagonism in mice.
1	Take NMR1 mice (20-259)
	1
	Administer drug 2P (10 mg/kg)
	Resexpine (o.imglkg)
	3 Chservation for ag his.
	Les expine inject.
	Produce Horizontal movements for every 10 minutes
	Rearing & Grooming is recorded by expert observer.
4	Preclinical Screening pulnciples of Sympatholytic drug
	2n vitro
	D' Micrating membrane prolapse in cale.
	> cop acrosses gre antagonism of mouse egg.
	Bavivo
	1) Vas deference of ral
	3) Straitel Ship of cat
	3) To access the B, & B2 advenorecuptors of agonism &
	antagonism.

Vas deference of val Rat (1- 4.8 gm) killed by Stunning Cut the abdomen & make the midline incision to dissect the Vas deference. Treame is suspend in tream both (tyrode, actided, No is added Phentolamine is used as Standard (1. reduction of activity of the advenoreceptors. Straited Strip of cat Cat (a-3 kg) Anesthesized. Suspended in organ bath (Krebs, aeraled, 38°c). Tension is added 0.5 mg, magnitude 5-6 times, then NA/A is added then after add test

Phentolamine is used as come
Phentolamine is used as Standard drug (.1. reductions of agonism & antagonism.
NI.
Miciating remembrane prolapse in cake.
Cats (Group 6- a gormale)
(2-3 kg)
V. "
Anesthesized with pentobarbitone.
Dest compound advenaline is administered
at produces the relaxant activity
Compare the animals with Std.
Continue the process with other animals
awith different concentrations.
Anesthesia
It is the drug to produce reversible loss of Sensation
87 Consiousness.
-> Anesthesia is done by experts in lab
-> So many conesthitics are available mainly used
D Enhalata anesthetics.
2) Intravenous anesthetics.
Daralysed for some-time with artificial respiration is prod
Paralysed to some-time with artificial respiration is prod

-> This is done at Experimental lab. Tension is avoid by patient, animal also feels fearless. - Adequate results are not come due to tension. -> care should be taken by expertian. -> Helshe has experienced in cutting Stills insertion Stills. -> Hardle the animal with appropriate care. -> cruelty is avoid deal carefully. -> once animal is anestherized incision is made & cut, take the part it should be placed at tissue Suspended bath (81gan bath). -) Done the Stitches almost care. -) Provide the nutritional food of saline liquids to recovery of animal.) Tablets will be dissolve & given through Oral route -) Rf any cases it should not recover & unhealthy it should be immediately done by euthanasia. Euthanasia The animal is gentle killing on death is called euthanasia. -) This is done for experimental work or research or

	fermentation of organs. in labs.
	-> This should be done by painters.
	-> Euthanesia is done by ethical process.
	not in Cruelty tolm.
1	Experiments are done almost without harming leotheresia. For euthenesia it has some rules bugulations. Reason is compulsory for euthenesia is done.
1	For euthanesia it has some rules Buequiations.
	Reason is Compulsory for euthaneria is done.
	maintain records
6.	Différent Strains and species of laboratory animals. Rat:
	This the Small animal in laboratory it should be
	educated , very sensitive to drugs.
	inbred rate used.
	a) wistar rats
	6) Albino rate .
	Thead is wide, tail is long which is longer than body
	Albino rate
	Head is long, narrow body tail is equal to body.
	Some Characteristics of val
	-> Rate has doesn't Vomiting center that's why it can't
*	Vomit
	> No gall bladder.

	-> Mainly used for tetratogenicity, mutagenicity & Carcinogenicity.
	Carrinogenicity.
	(V lous e
	-) This is the smallest animal in laboratory, available as
	Cheap -
	-> Sensitive to drugs.
	Determine by using teralogenicity.
	Fillnea pig
	This is the I I O I I CM
	This is the docte & deal conefully.
	> But is sensitive to hickomines & produce Severe brancho constrict a asphyxis.
	-> used to 1-a sphyxis.
	-) used in biolesay of digitalis & local anesthetics.
	thannister inflammatory dough identification.
	Two types hammsters are present
	1) Golden hammster.
	2) Crusene hammster.
	This is churty body, Short legs, to toes in back, 4 tous
	in front.
	Golden hammster is used for vivology, cancer.
	Cocurene hammeter is used for antidoabetic, antipyretics.
	mainly for immunology.
	Rabbit
	Reupealand rabbit is mostly used. Bet has long ears.
1	of has long ears.
	- It is used for corresponding & telephogenicity.

i

.1

-> Mainly used for bioassay of digitalis.
Montey
Structurally resemble to human being physically.
-) This brain is close to human brain.
-> renainly on I have to human bearn.
Cat Neuro disorders.
cat is used for BP trees - + (131
morphine is Produce unconsiousness in cat
-> Caratoid autery to Carl
Frog Frog artery is fixed to measure the bp.
- 1 Rt is aquatic amphibian animal.
The mainly used for Neuro muscular junction publishes
HE I Wello muscular junctions published
Frankgenic animals.
Mouse
-> The first animal
This is done by altering the gene of mother &
insert into baby rat- (egg), it should produce.
large size baby (size of moderse) than mother.
Goat
Take goal milk & make them milk for Ophan
babies to feed -
street: (Dolly is the first clone baby).
Sheep: (Dolly is the first clone baby). - wool is used for making textiles, pharmaceutical uses. Chicken
-) an ovum an new molecule is inserted large size her is

Produce & it should be produce more eggs.

Fich

Debra fishes are newly produced.

Cath Buffalo

These milk are used for feed for in many

products

I M. Pharmacy | I Sen II mid Exams Sub Pharmacological and Toxicological Screening methods - I 2. Screening methods of Antidiabetic ") Pancreatumy of dogs 2) Alloxan induced diabetics (Rabbit, Rat, Dog) 3) Streptozotocin induced diabetics (Rat). 4) - Hormone induced Diabetic mellitus Pancreatarny of Dogs > Open abdomine Male Beagil Dogs -> - Anaestherized with with care and Pento Sodium barbitone (12-16 Kg) with proper IV (30 mg/kg) & placed Surgical Skilk on back Pancreas is brought Botto ends of Small vessels into opeasting field of pancreas + Pancreas are & Separate the mesentic are ligated ligated attachments After 19-20 has The pylonic & splenic The mysentry body Pancreas is of pancreas & tail parts of pancreas dissected out is Separated & are delievered into by Sacrificing the animal wounds. The pancreas is The pancreatic The pancreatic tiesue of the only one part branch of Splenic Splenic vessel is is to be dissected vessels are doubly lighted ligated

1	50	
Finally pancreas is . to be dissected out	The abdominal wall & Superficial layer of Skin is Statured.	treated with post operative
	3rd day give milk Q teem to normal feed.	1 litre of glucore Q 10 Eu of retard Ansulin . Q 3P of
<	Ensulin is replaced with retard insulin.	mckami≥ol.
Allowan induced dia	7	
Rabbit New Zealan Albino to (2.5-4kg	(5 tog 100 ml) is	- mic hypoglyce.
#V	100-150 mg/kg IV	Doug is used for further Screening.
(150-2009)	Alloxan monohydrate -	-) Hyperglycemic
og → Male Beagil sale (12-16 kg)	-> Iv 60 mg/kg -> Olloxan mohonydrate	Jireated with glucose & Canned food
25		Treated with single Sc retard Ansulin

	Streptozotown induced diabe	tics
1	Albino Test drug admi rate through IV (150-2009) (60 mg/kg)	inister -> Initially increase glucose level in 150-200 mm H bcz increase blood Volume in 31
34 34		1
	After 12-14 Beverity & Symptoms days of Sheeptosotocin Steady State concentration the animals are undergo for Screening tests.	doe After 15-4 hrs. glucoco levels increase because of increase in serum of glucose.
	Homone induced diabetics	. * :
	Deoxymethasone -> Du 2.5 m (Long acting Carticosteroid)	nglkg -> 200m of diabetics.
3	Anti cancer drugs	
	=0	
	an vitro	
	i) Triptan blue viability test	
	2) Alamol blue assay. In vivo In vivo Liver cancer induced by diether Carcinogen induced models.	•
1	Conserver induced models	0 1.
7) caranoger mance	*
	Viral infect illiaces	
	-) Transplantath models.	
	I Hollow fibre	

by diethyl amino bengene -> tel Childel Migh > divided into a Rak vehicle of the his (150-800g) groups (distilled water) -> Each group has -) the link down with 5 animals (Test & control) dough DANGE & TIM After 6 hrs Dismo 1 twiled Dissect In 15th After 4- 5 Observe week < मान ०२ । नि See the live months Livertums 12 weeks Colour. into gray Colour 7-Any patches ___ Identify is Seen is ___ Live ANOVA . Values ? Checked. malignacy Douameters Carcinogen induced models Mice only for single dose of 2.5 mg of DHMB in acctone & 6-10 pg of TPA in 0-2 ml in acetone Percent of Caranogen incidence & multiplicity of treatment - Compared with DNMB controls. Trest compound is given through interperitornally & The percent of Carcinogen is usually 100%. of DNMB

DNMB alone give it induce Carcinogens.

Reduce carcinogen is identify by reduction of symptoms

Vival infects models

- The mouse mammary Tumous Virus (mmTV) is isolated in Jackson laboratory identified as "Non-Chromosomal factor". I produce tumous in CH3 Strain in mice.
- -) Some Viruses Cause Cancer via integrate in Certain cells.

Some virusus cause tumour by oncogenes.

Jabelson mikaratne leutemia Virus.

Monoley mikuraine Sarcoma Virus.

- Engineering virus are used now routienly tol.

Transplantata Virus (models)

- -) Turnous cells on tiesues are implanted in a hoset
- Ectopic Implanted organ than in different organ.

 Orthotopic Implanted organ into analogus organ

 into original turnous.

an vivo hollow fibre assay 1.
-> In vivo Screening tool introduced in 1955 by NCI
-) le human turnour cell lines (Breast, cancer, colon,
malemia le ovacy)
-) Ofter in vivo treatment fibres are removed & analyse
in viteo.
-> In vivo assessed availability.
Drack 1
Preclinical screening principles of antiasthamatic daugs
In vitro
Resolated Guinea não Luna Strips.
Resolated Guinea pig Lung Strips.
2) Psolated Guinea pig Trachea.
1) Bronchospasmolytic activity in Guinea pig-
Describe Sparmolytic outstity in Guinea pig- 2) Broncho overadivity of Guinea pig.
desolated Guinea pig Trachéa.
Albino Guinea pig (300-350g)
(300-3509)
20 M 10 M
Sacrificed with con necrosis.
Sectorice a with Con Decrosis
V -
Entire Trachea is removed le cut înto individual
ornge.
. O

All rings are held together by silk ! thread.
Mounted to Sigan both containing kreb's Solution & buffer Solution at 37°c & tension is added
& buffer Solution at 37°c & tension is address
Bath is bubbled by adding Carbogen.
J.
Isomesentic contractile is measured by using
polygeaph.
Spasmogen is added.
1
A Fest drug is added (Beopranil 1 mg/kg)
Obs a construction and Construction
Obtain Constant contractile add Spasmagen
Add test drug
4
Measure Constant Confractile Obtain at
maximum level.
Bronchospasmolytic activity in Guinea pig.
Male Guinea pig (200-300 g)
(200-300 g)
Anestherized with pentobarbitone.

Anestherized is not much deep avoid spontaneous respirato Jugular vein is connulated by for test dough Spasmogen. Caratoid outery is measured for bp Trachea is cannelated by two way Trachea. I) one trachea for transducer bp 2) one trachea for respirato Artifical aersol is pumped at 190-200 mm in 1 man Stroke Measure & record the lung which is not taken. Add Spasmogen (Histamine, HCl) Contractile is produced & by adding sparmagen B test drug in 10-15 mints interval Broncho overactivity of Guinea pig : Doine Gennea pig (300-350g)

a consiste of Enhalato acrosol boxes for B & c.
↓
Rate is placed in tox -1 & treated with across
& De ultrasound nebuliques.
J
Box B is passed way to Box C.
1 hietamine
an Box c produce 0.1.1. Solven of Hel with altroses
nebuliger.
1.
produce Convulsions.
J
Emmediately remove animals from box.
Pharmacological Screening methods of antiulcer dougs
D) pyrotic tigator of sale.
2) Stress ulcer models
a) Restrain induced ulcers.
b) cold water immersion ulcers.
c) Stress & NSAIDS induced relicers
d) Swimming , Strew ulcers.
3) Histamine induced gastric ulcers.
4) Acetic acid induced gastric ulcers.
5) Reserpine induced Chronic gastoic ulcers.

5.

I) pylonic light of Rate > Anestherized by wistar rate .__ , fasting-fo) ether 46 hrs but (150-200g) given water 1 inch abdominal Pylorous is lifted incision, below Carefully without sciphoid process disturbing the blood Supply. contente of Stomach are drained out by Stomach is open graduated centrifuge along greater Curvature Acetic acid 0.01N Nacl Measure The is used to centifuge Seventy of lessions . UI = UH+ US+Up X10-1 2) Stress ulcer models a) Resembine induced ulcers lest compound administer , Fasted for 36 Orally (150-200 g) Measure Geep Straining Stomach is open & gastoic relicers In ag his t dissected out by Stockning animal.

Cold-water immersion induced ulcers.
rate Pasted for Text compound administer
(150-2009) le hours animale are dipped into
Evan blue is water by restraining
(30 mg/kg) Vertically at
is given by 1 hour
tail vein
Next day diesect
Provide formal - out by open
Saline overnight Stornach along
guater curvature
• 1:
Lesions are wash with warm
measured by water
helping of evan blue
colour.
Stores & NSAIDS induced ulcers
wister - Fasted for - Test drug administer.
Du-36 hours
1100-20091
Stomach is Ocal Ni in 30 ml of
Stomach is 0.01 N in 30 ml of
open along inject (20)
greater
, Cuntature
· 1
Measure lesions of
gastric ulcers.

-1\ 0
d) Swimming induced ulcers
Albino rate -) Fasted for - administer test doug
111-6 % 1 100 1 100 100 100 100 100 100 100 1
Dulan woller -to with
arec for 5 hrs.
rales s
Stomach is Ofter 5-4 his animals
Open along - are Strained.
gleater curvature.
. L
Lesions is measured as
0 - Lexions is absent
1'- Lesions c 1 mm.
2 - Lerians c 1-2 mm
3 - Lesions C 2-4 mm
5 - Lesions (more than 4 mm
Hickory and Danie Warre
Historine induced gastric ulcers
wiset-
Guinea _ Facting for _ Hickamine - Hickamine toxicity
Guinea Facting for - Historine This trainine toxisty Pig (200-250 g) (200-250 g) Facting for - Historine toxisty Chaptine is Given before le
(200-250 g) given before &
after drug
treatment 0
1
A something of the
measured along gleater
Culvature

Acetic acid induced gastric ulcers.
-1 / / / / / / / / / / / / / / / / / / /
(150-200 gmg) - fasted for - Autic and (0.01) in (150-200 gmg) 24 hrs 30 ml of injection is
(150-200 gmg) - Jasted for - Autic and (0.01) 11) (150-200 gmg) 24 hrs 30 ml of injection is given at mucosomal layer of Stomach
J. Jagler of
These are typically produce gastric ulcers
These are typically produce gastric ulcers Chronic ulcers by penetrating.
regenerated with
healing.
Reserpine induced chronic pulcers -
Wistar rats provide Std Before expt (150-2009) diet Before expt
(150-2009) diet withdrawn
liquid taken
1 mg/kg - Criven test doug
is given Texapine au.
Stomach is open along
gréates convature &
Messus County of
Measure Seventy of
legions.
ii

& Pharmacological Screening methods of antihypertensive agents and hepatoprotective drugs an vivo I) Two kidney 1 clip method in sp rate 2) one kidney I clip method in SD rock 3) Salt sensitive dant rate 1) Fouctose induced hypertension in wistor rate 5) DOCA Salt rate 6) Tail cuff method. Two Kidney 1 clip method in Sprague Drawley rate -> The arrivery is consmicted on only one side with the other astery left untouched. -) This result in Sustained increase in BP. Initially, Salt Q water retent is more because the contact of other kidney being infact -> In this Situate the Kidney resultant in andiotensin dependent. -) The increased angiotensin (1) is released from aldosterone from advenal medulla. Salt & water retn is male -> This results in decrease renin products
-> This is a volume dependent process (Hy extension)

one cloney I cho method in sprak	W5
Here one briefy is constricted by only one side	other
1	
Britially within a few hours by increase.	
. 1	
There is no constricts of conora lateral tidney & r	70
Pressure d'arens l'numuresis.	
, 4	
Salf Q Hao terenth is more	
1	60
plasma venin usually normal.	
Hypertansion is volume dependent.	
Salt Sensitive dahl rate	
	70-
No. (711)	ntice.
(150-2009) Salt Contain the upto 1 month of provide this Meas	bP
-food 1.	cuse bp
before &	after
Measure Sacrifice & doug to	
Organ size increase the text	14.
geoup animals	
damage the organ.	

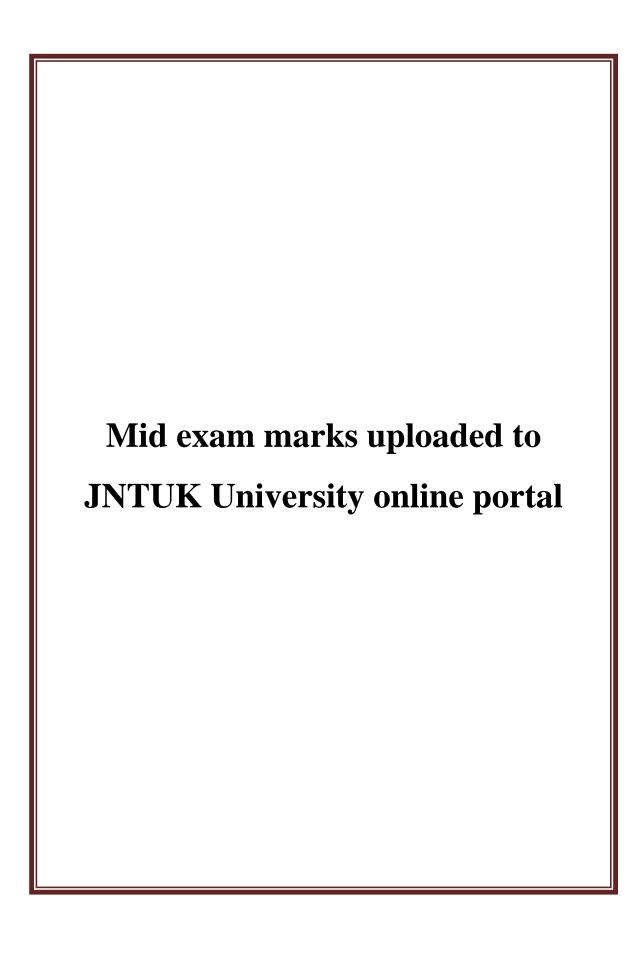
Fructore induced hypertension in wistor rate Wistar rate - feed with Observe, Produce (150-200g) ansulin Canned food fluid intake registance Dointing behaviou Body weight Test doug adminis Measure bp, Fest by Brisulin 8 Triglycerides before ANOUA . & after doug' treatment. DOGA Salt rate wistar rate _s one kidney Don king the Puovide replaced with (180-200 g) Salt content is removed water upto Saline 1 month ancrease Bp Measure before & after doug treatment -Tail cuff method Charles River Produce artifical Annestherized by male rats hypertension by 2P 0.8 ml of 41. (800-350q) expose both kidneys of Chloral hydrate & place the clips of both rend arteries.

cust-
-> After 15-6 weeks - A inflatable wise is
hypertension is attached at tail
attained base.
1
inflatable cuff is approximately
reach 300 mm offg.
J
pressure of inflatable is slowly
removed & press bp is detected
& record in polygeaph.
-40
Administer Less dans 20 Bi
Administer test drug IP -101
& times at in alternative days
decreased bp
↓ '
Evaluato
Day 1: predrug & his post drug
Day 3: predrug, & has post drug.
De in a bis post dough Q & boss
Day is: Preday 2 his post daugs Q 2 has
postding.
U

Mid exam marks scored by students are entered in the Mother register

Pharmacology I MiPh / I Sem (2022-23)

		Sub: Pharmacological & Toni	cological	Screenf	ng Methods	-I IMPLI037362			
SINO	Register No	Name of the Student						Practical Marke	
<i>C</i> .	V	V	Imid	Dimid	0 0				
	287NISO601	BOYANAPAUT PRASANNA	21	2)	22	M			
2	COBOLINESS	CHALAMALA RAMYANJALT	19	20	20	0			
3	227NISD603	SHATK HAFSA	0	0	0	P			
					t.	R			
						A			
				7					
		S-Venychish		(1)	1	,			
		EXAMS-INCHARGE VIJAYA INSTITUTE		PRO	VCIPAL				
		PHARMACEUTICAL SCIENCES FOR WO		RMACEUTICAL	SCIENCES FOR WOME	C			
			N	IKEPADU.VIJ	AYAWADA 521 10	A			
		• ,				L			
			1	1	1	10.70 (I.		





JAWAHARLAL NEHRU TECHNOLOGICAL UNIVERSITY KAKINADA

FINAL PDF for M.Pharm I Semester Internal Marks
College: VIJAYA INSTITUTE OF PHARMACEUICAL SCIENCES FOR WOMEN:7N

Date:19-05-2023

	•	1	1			
HTNO	SUBJECT	MID_1	MID_2	SEMINAR	FINAL	SUB_TYPE
227N1S1601	MPA101T	25	25	0	25	Τ
227N1S1602	MPA101T	24	24	0	24	T
227N1S1603	MPA101T	24	25	0	25	T
227N1S1604	MPA101T	0	0	0	0	T
227N1S1605	MPA101T	25	25	0	25	T
227N1S1606	MPA101T	25	25	0	25	T
227N1S1607	MPA101T	24	25	0	25	T
227N1S1608	MPA101T	24	24	0	24	T
227N1S1601	MPA102T	25	25	0	25	T
227N1S1602	MPA102T	21	24	0	23	T
227N1S1603	MPA102T	23	25	0	24	T
227N1S1604	MPA102T	0	0	0	0	T
227N1S1605	MPA102T	22	24	0	23	T
227N1S1606	MPA102T	24	25	0	25	T
227N1S1607	MPA102T	24	24	0	24	Τ
227N1S1608	MPA102T	20	21	0	21	T
227N1S1601	MPA103T	23	25	0	24	Τ
227N1S1602	MPA103T	22	25	0	24	T
227N1S1603	MPA103T	23	25	0	24	Τ
227N1S1604	MPA103T	0	0	0	0	Τ
227N1S1605	MPA103T	22	25	0	24	T
227N1S1606	MPA103T	24	25	0	25	T
227N1S1607	MPA103T	25	25	0	25	T
227N1S1608	MPA103T	22	25	0	24	T
227N1S1601	MPA104T	25	25	0	25	T
227N1S1602	MPA104T	24	25	0	25	T
227N1S1603	MPA104T	25	25	0	25	T
227N1S1604	MPA104T	0	0	0	0	T
227N1S1605	MPA104T	25	25	0	25	Τ
227N1S1606	MPA104T	25	25	0	25	T
227N1S1607	MPA104T	25	25	0	25	Τ
227N1S1608	MPA104T	20	22	0	21	T
227N1S1601	MPA105PA	25	25	0	25	L
227N1S1602	MPA105PA	24	24	0	24	L
227N1S1603	MPA105PA	25	25	0	25	L
227N1S1604	MPA105PA	0	0	0	0	L
227N1S1605	MPA105PA	25	25	0	25	L
227N1S1606	MPA105PA	25	25	0	25	L
227N1S1607	MPA105PA	25	25	0	25	L
227N1S1608	MPA105PA	25	25	0	25	L
227N1S1601	MPA105PB	25	25	0	25	L
227N1S1602	MPA105PB	25	24	0	25	L

LITNO	CUDIFOT	MID 4	MID 2	CEMINAD	FINIAL	CUD TYPE
HTNO	SUBJECT	MID_1	MID_2	SEMINAR	FINAL	SUB_TYPE
227N1S1603	MPA105PB	25	25	0	25	L
227N1S1604	MPA105PB	0	0	0	0	L
227N1S1605	MPA105PB	25	25	0	25	L
227N1S1606	MPA105PB	25	25	0	25	L
227N1S1607	MPA105PB	25	25	0	25	L
227N1S1608	MPA105PB	25	25	0	25	L
227N1S1601	MPA106S	0	0	98	98	S
227N1S1602	MPA106S	0	0	85	85	S
227N1S1603	MPA106S	0	0	98	98	S
227N1S1604	MPA106S	0	0	0	0	S
227N1S1605	MPA106S	0	0	97	97	S
227N1S1606	MPA106S	0	0	98	98	S
227N1S1607	MPA106S	0	0	85	85	S
227N1S1608	MPA106S	0	0	85	85	S
227N1S0301	MPH101T	25	25	0	25	<i>T</i>
227N1S0302	MPH101T	24	25	0	25	<i>T</i>
227N1S0303	MPH101T	24	24	0	24	T
227N1S0304	MPH101T	25	23	0	24	Τ
227N1S0305	MPH101T	24	24	0	24	T
227N1S0306	MPH101T	25	25	0	25	T
227N1S0307	MPH101T	24	25	0	25	T
227N1S0308	MPH101T	25	25	0	25	T
227N1S0309	MPH101T	24	25	0	25	Τ
227N1S0310	MPH101T	25	25	0	25	T
227N1S0311	MPH101T	23	22	0	23	T
227N1S0312	MPH101T	25	25	0	25	T
227N1S0313	MPH101T	0	0	0	0	T
227N1S0314	MPH101T	21	25	0	23	Τ
227N1S0301	MPH102T	25	24	0	25	T
227N1S0302	MPH102T	25	22	0	24	Τ
227N1S0303	MPH102T	24	23	0	24	T
227N1S0304	MPH102T	25	23	0	24	Τ
227N1S0305	MPH102T	24	24	0	24	T
227N1S0306	MPH102T	25	23	0	24	Τ
227N1S0307	MPH102T	21	22	0	22	Τ
227N1S0308	MPH102T	25	23	0	24	T
227N1S0309	MPH102T	22	23	0	23	<i>T</i>
227N1S0310	MPH102T	24	25	0	25	<i>T</i>
227N1S0311	MPH102T	18	23	0	21	<i>T</i>
227N1S0312	MPH102T	24	25	0	25	<i>T</i>
227N1S0313	MPH102T	0	0	0	0	<i>T</i>
227N1S0314	MPH102T	20	23	0	22	<i>T</i>
227N1S0301	MPH103T	24	25	0	25	<i>T</i>
227N1S0302	MPH103T	25	25	0	25	Τ
227N1S0303	MPH103T	25	25	0	25	T
227N1S0304	MPH103T	25	24	0	25	Τ
227N1S0305	MPH103T	25	25	0	25	T
227N1S0306	MPH103T	25	25	0	25	T
227N1S0307	MPH103T	25	25	0	25	Τ
227N1S0308	MPH103T	24	25	0	25	Τ

UTNO	SUB IECT	MID 4	MID 2	CEMINAD	FINAL	SUB TYPE
HTNO	SUBJECT	MID_1	MID_2	SEMINAR		SUB_TYPE
227N1S0309	MPH103T	24	25	0	25	<i>T</i>
227N1S0310	MPH103T	25	25	0	25	<i>T</i>
227N1S0311	MPH103T	24	22	0	23	<i>T</i>
227N1S0312	MPH103T	25	25	0	25	<i>T</i>
227N1S0313	MPH103T	0	0	0	0	<i>T</i>
227N1S0314	MPH103T	24	25	0	25	<i>T</i>
227N1S0301	MPH104T	22	25	0	24	<i>T</i>
227N1S0302	MPH104T	22	25	0	24	<i>T</i>
227N1S0303	MPH104T	22	25	0	24	T
227N1S0304	MPH104T	23	25	0	24	T
227N1S0305	MPH104T	24	25	0	25	Τ
227N1S0306	MPH104T	23	25	0	24	Τ
227N1S0307	MPH104T	19	25	0	22	T
227N1S0308	MPH104T	20	25	0	23	T
227N1S0309	MPH104T	21	25	0	23	T
227N1S0310	MPH104T	20	25	0	23	T
227N1S0311	MPH104T	15	23	0	19	Τ
227N1S0312	MPH104T	19	25	0	22	Τ
227N1S0313	MPH104T	0	0	0	0	Τ
227N1S0314	MPH104T	20	25	0	23	T
227N1S0301	MPH105PA	22	24	0	23	L
227N1S0302	MPH105PA	24	24	0	24	L
227N1S0303	MPH105PA	23	23	0	23	L
227N1S0304	MPH105PA	24	23	0	24	L
227N1S0305	MPH105PA	25	25	0	25	L
227N1S0306	MPH105PA	23	23	0	23	L
227N1S0307	MPH105PA	22	23	0	23	L
227N1S0308	MPH105PA	23	23	0	23	L
227N1S0309	MPH105PA	22	23	0	23	L
227N1S0310	MPH105PA	22	23	0	23	L
227N1S0311	MPH105PA	22	21	0	22	L
227N1S0312	MPH105PA	23	24	0	24	L
227N1S0313	MPH105PA	0	0	0	0	L
227N1S0314	MPH105PA	22	23	0	23	L
227N1S0301	MPH105PB	21	23	0	22	L
227N1S0302	MPH105PB	24	23	0	24	L
227N1S0303	MPH105PB	21	22	0	22	L
227N1S0304	MPH105PB	22	22	0	22	L
227N1S0305	MPH105PB	24	23	0	24	L
227N1S0306	MPH105PB	23	23	0	23	L
227N1S0307	MPH105PB	22	22	0	22	L
227N1S0308	MPH105PB	23	22	0	23	L
227N1S0309	MPH105PB	22	22	0	22	L
227N1S0310	MPH105PB	22	23	0	23	L
227N1S0311	MPH105PB	21	22	0	22	L
227N1S0312	MPH105PB	22	23	0	23	L
227N1S0313	MPH105PB	0	0	0	0	L
227N1S0314	MPH105PB	22	23	0	23	L
227N1S0301	MPH106S	0	0	85	85	S
227N1S0302	MPH106S	0	0	95	95	S

LITNO	CUDIFOT	MID 4	MID 2	CEMINAD	FINIAL	CUD TYPE
HTNO	SUBJECT	MID_1	MID_2	SEMINAR	FINAL	SUB_TYPE
227N1S0303	MPH106S	0	0	80	80	S
227N1S0304	MPH106S	0	0	85	85	S
227N1S0305	MPH106S	0	0	98	98	S
227N1S0306	MPH106S	0	0	80	80	S
227N1S0307	MPH106S	0	0	85	85	S
227N1S0308	MPH106S	0	0	83	83	S
227N1S0309	MPH106S	0	0	95	95	S
227N1S0310	MPH106S	0	0	95	95	S
227N1S0311	MPH106S	0	0	80	80	S
227N1S0312	MPH106S	0	0	85	85	S
227N1S0313	MPH106S	0	0	0	0	S
227N1S0314	MPH106S	0	0	85	85	S
227N1S0601	MPL101T	25	25	0	25	<i>T</i>
227N1S0602	MPL101T	22	25	0	24	<i>T</i>
227N1S0603	MPL101T	0	0	0	0	<i>T</i>
227N1S0601	MPL102T	24	24	0	24	<i>T</i>
227N1S0602	MPL102T	22	24	0	23	T
227N1S0603	MPL102T	0	0	0	0	Τ
227N1S0601	MPL103T	21	22	0	22	T
227N1S0602	MPL103T	19	20	0	20	T
227N1S0603	MPL103T	0	0	0	0	T
227N1S0601	MPL104T	25	25	0	25	T
227N1S0602	MPL104T	23	24	0	24	Τ
227N1S0603	MPL104T	0	0	0	0	T
227N1S0601	MPL105PA	24	24	0	24	L
227N1S0602	MPL105PA	24	24	0	24	L
227N1S0603	MPL105PA	0	0	0	0	L
227N1S0601	MPL105PB	25	24	0	25	L
227N1S0602	MPL105PB	25	24	0	25	L
227N1S0603	MPL105PB	0	0	0	0	L
227N1S0601	MPL106S	0	0	98	98	S
227N1S0602	MPL106S	0	0	97	97	S
227N1S0603	MPL106S	0	0	0	0	S
227N1S1701	MRA101T	21	25	0	23	Τ
227N1S1702	MRA101T	22	25	0	24	T
227N1S1703	MRA101T	24	25	0	25	T
227N1S1704	MRA101T	24	25	0	25	<i>T</i>
227N1S1705	MRA101T	23	25	0	24	<i>T</i>
227N1S1706	MRA101T	25	25	0	25	<i>T</i>
227N1S1701	MRA102T	25	25	0	25	T
227N1S1702	MRA102T	25	25	0	25	<i>T</i>
227N1S1703	MRA102T	25	25	0	25	T
227N1S1704	MRA102T	25	25	0	25	<i>T</i>
227N1S1705	MRA102T	25	25	0	25	Τ
227N1S1706	MRA102T	25	25	0	25	T
227N1S1701	MRA103T	21	22	0	22	Τ
227N1S1702	MRA103T	19	22	0	21	T
227N1S1703	MRA103T	22	21	0	22	T
227N1S1704	MRA103T	24	23	0	24	Τ
227N1S1705	MRA103T	24	24	0	24	Τ

HTNO	SUBJECT	MID_1	MID_2	SEMINAR	FINAL	SUB_TYPE
227N1S1706	MRA103T	23	24	0	24	Τ
227N1S1701	MRA104T	22	23	0	23	Τ
227N1S1702	MRA104T	23	22	0	23	Τ
227N1S1703	MRA104T	21	22	0	22	Τ
227N1S1704	MRA104T	25	25	0	25	Τ
227N1S1705	MRA104T	25	25	0	25	Τ
227N1S1706	MRA104T	25	25	0	25	T
227N1S1701	MRA105PA	21	21	0	21	L
227N1S1702	MRA105PA	22	21	0	22	L
227N1S1703	MRA105PA	22	21	0	22	L
227N1S1704	MRA105PA	24	24	0	24	L
227N1S1705	MRA105PA	24	24	0	24	L
227N1S1706	MRA105PA	23	24	0	24	L
227N1S1701	MRA105PB	24	24	0	24	L
227N1S1702	MRA105PB	24	24	0	24	L
227N1S1703	MRA105PB	25	24	0	25	L
227N1S1704	MRA105PB	24	24	0	24	L
227N1S1705	MRA105PB	24	24	0	24	L
227N1S1706	MRA105PB	25	24	0	25	L
227N1S1701	MRA106S	0	0	85	85	S
227N1S1702	MRA106S	0	0	80	80	S
227N1S1703	MRA106S	0	0	80	80	S
227N1S1704	MRA106S	0	0	98	98	S
227N1S1705	MRA106S	0	0	98	98	S
227N1S1706	MRA106S	0	0	95	95	S

Controller of Examinations

Date:19-05-2023

Verified by: PRINCIPAL