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Dr. K.V. Subba Reddy Institute of Pharmacy

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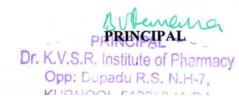
E-mail: principalkvsrip@gmail.comwww.drkvsrip.ac.in

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2.3 Student centric	methods such as exp	eriential learning, participative learning and problem solving methodologies are used
for enhancing learn	ing experiences and	teachers used ICT Enabled tools including online resources for effective teaching and
learning process	(AY: 2021-22)	

INDEX

S. No	Description	Page No
1	Sample Copy of LCD Projector	2
2	Sample Copy of chalk and board	3
3	Sample Copy of LCD projector used classes	4
4	Sample Copy of online teaching	5-6
5	Sample Copy of blended teaching	7
6	Sample Copy of Edu blogs	8
7	Sample Copy of industrial training& internship	9-15
8	Sample Copy of hands on learning	16-17
9	Sample Copy of assignment	18-20
10	Sample Copy of innovative model making	21
11	Sample Copy of project based learning	22-85
12	Sample copy of case study learning	86-101
13	Sample Copy of E-learning	102-109
14	Sample Copy of workshop & guest lectures	110
15	Sample Copy of pharmacy seminars	111
16	Sample Copy of group discussion& debates	112-117
17	Sample Copy of bridge courses	118-123
18	Sample Copy of remedial classes	124-149
19	Sample Copy of lab sessions	150-152
20	Sample copy of quiz	153
21	Sample copy of think pair share	154
22	Sample copy of club activities	155
23	Sample copy of problem solving methods	156-159
24	Sample copy of virtual labs	160-163





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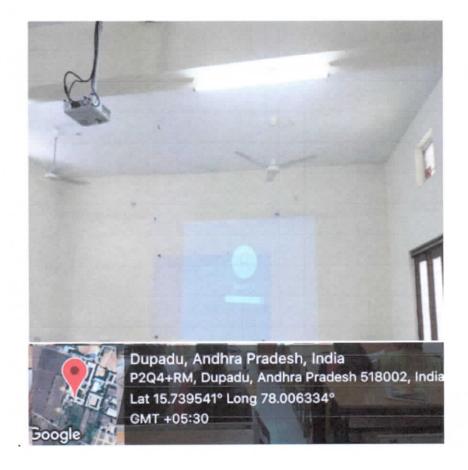
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LCD PROJECTOR:

Spacious, well -lit and well -ventilated classrooms and seminar hall [with ICT] with comfortable seating arrangement are available for smooth conduct of UG and PG theory sessions. Besides the conventional teaching aids, classrooms are also equipped with state-of-the-art audiovisual technology viz. digital smart board, LCD Projectors, thus stepping up the teaching-learning experience to next higher level. Well -thought positioning of aisle in the classroom ensure proper interaction between teacher and students



The list of various ICT enabled pedagolocial initiatives used by the faculty members for improving teaching and learing experience are as follows:

Chalk and board: To convey basics, critical information, backgrounds, theories



Fig1.1:Mrs.Anuradha Assistant professor of Department of Mathematics explaining concepts of Matrix for Pharm D I year students of pharmacy using chalk and board



Fig: 1.2Mrs.B.Jhansi, Assistant professor of department of Pharmaceutical Chemistry explains the concepts of Acid-Base Reaction for B.Pharmacy I year students of pharmacy using chalk and board

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Fig 1.2 Mr.Harish, Assistant professor of department of Pharmacy Practice explains the topic of ICF using LCD projector.



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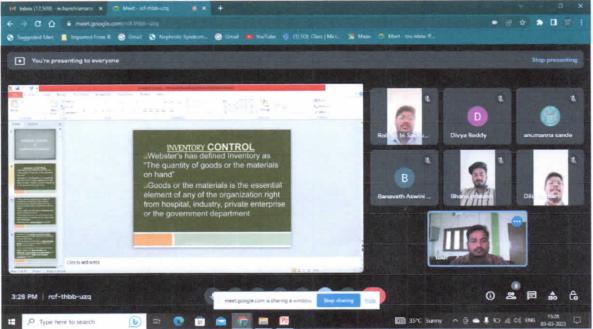
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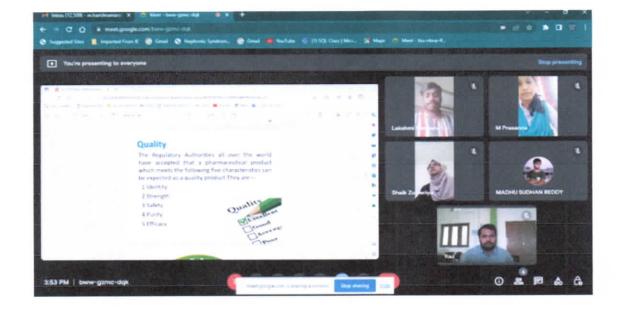
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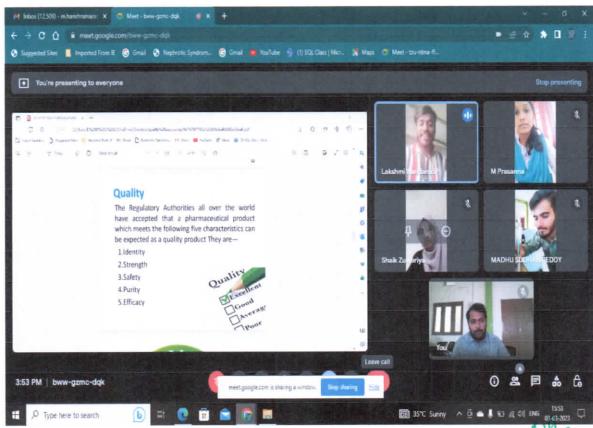
ONLINE TEACHING:

Especially during Covid-19 pandemic situation made a swift transition from classroom to online teaching by using platforms like Google meet, Google classroom, ZOOM, CISCO, WEB-EX, and You tube.











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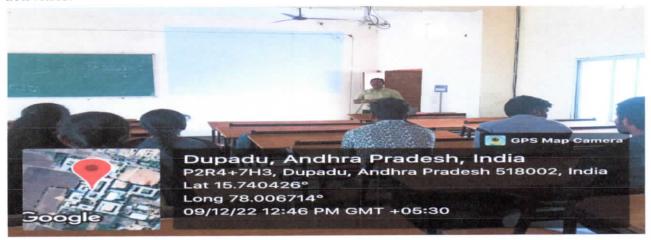
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BLENDED TEACHING:

Teaching aid techniques such as video lectures and PowerPoint presentations are being used for the implementation of active learning strategies such as collaborative and individual learning activities.



Mr.P.T.Nagaraju, M.pharm department of pharmaceutical analysis explaining basic concepts in analysis to B.Pharm I Year students.





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Edu-Blogs: Faculty blogs and channels are used as instructional potential of online resources to students

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INDUSTRIAL TRAINING 14/06/2022- 30/06/2022

Industrial training, where a student undertakes a period of training at an organization usually during a semester break, acts as an important part in preparing the student for a professional career. With the actively involved preparation, the student learns about the industrial demands, skill set and work ethics.

Every year students experience industrial training as per the syllabus prescribed in JNTUA and gain knowledge of various departments like Formulation and research department (FR&D) i.e., Manufacturing of Tablets, capsules and injection preparations etc., Quality Assurance, Quality control, Clinical research, Drug regulatory affairs, Hospital Pharmacy etc. as such III year B.Pharmacy students have visited KRISH CARE FORMULATIONS& BRILLIANT BIO PHARMA(P)LTD, Hyderabad, Telangana-5000 72, India and received hands on experience about industry affairs.







Date:14/06/2022

TO WHOM SO EVER IT MAY CONCERN

This is to certify that Mr. G. TARUN KUMAR (18ER1R0016) S/O G.RAMU is bonafied student of DR.K.V.SUBBAREDDY INSTITUTE OF PHARMACY, KURNOOL. He has undergone industrial training in our organization from 14.06.2022 to 30.06.2022 as part of partial fulfillment of his B. Pharmacy course.

During the training, he had interacted with Quality control, Quality Assurance & Production departments and acquired basic knowledge in these areas.

During this aforesaid period, we found his hard working, sincere and learning attitude

For KRISHCARE FORMULATIONS

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11-6-56/2/25, Opp. IDPL Factory, Balanagar, Hyderabad 500 037

Cell: +91-9959666898 Telefax 040-23060195 Fimail krishcare Richardson com





Date:14/6/2022

TO WHOM SO EVER IT MAY CONCERN

This is to certify that Ms.P.SALIMA (18ER1R0038) D/O P.BASHA is bonafied student of DR.K.V.SUBBAREDDY INSTITUTE OF PHARMACY, KURNOOL. She has undergone industrial training in our organization from 14.06.2022 to 30.06.2022 as part of partial fulfillment of his B. Pharmacy course.

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Date:14/6/2022

TO WHOM SO EVER IT MAY CONCERN

This is to certify that Mr.S. MALIK BASHA(18ER1R0044) S/O S.GOUSE MOHIDDIN is bonafied student of DR.K.V.SUBBAREDDY INSTITUTE OF PHARMACY, KURNOOL. He has undergone industrial training in our organization from 14.06.2022 to 30.06.2022 as part of partial fulfillment of his B. Pharmacy course.

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Date:14/6/2022

TO WHOM SO EVER IT MAY CONCERN

This is to certify that Ms.H.ZAKIYA AFRIN (18ER1R0063) D/O H.KALIMULLA is bonafied student of DR.K.V.SUBBAREDDY INSTITUTE OF PHARMACY, KURNOOL. She has undergone industrial training in our organization from 14.06.2022 to 30.06.2022 as part of partial fulfillment of his B. Pharmacy course.

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INTERNSHIP:

DEPARTMENT OF PHARM.D STUDENTS INTERNSHIP REPORT

Academic Year: 2016-22 Batch: 2016

Internship Title:

The Pharm.D institutions are aligned with adjacent hospitals in order to provide real-time training including ward rounds. Students are acknowledged with practical in clinical pharmacy and medical pharmacy. After completion of the course, students are gained with the skills in diagnosis, therapeutic usage of drugs, treatment and selection of diseases

Internship Period: 15-06-2020 to 14-06-2021.

Company/ Organization: Kurnool General Hospital, Kurnool

Students Feedback:

	FACTORS ON FEEDBACK	Very Poor	Poor	Satisfactory	Good	Excellent
FACTOR	R: WARD ROUNDS					
1.	The tasks given are related to the subjects that I have learned in the institution.				5	2
2.	The tasks given are Challenging and testing your mettle.			1	1	5
3.	The tasks given were able to complete within the given duration.				3	4
ACTO	R: CLINICAL AND MEDICAL PHARMACY S	KILLS				
4.	The work environment is suitable for training of industrial trainee.			1	6	
5.	The colleagues provide good support and respective of industrial trainees.			1	4	2
ACTO	R: IN DIAGNOSIS SKILLS					
6.	The training attended has prepared me well to work in the industry in terms of knowledge.			1	3	3
7.	The overall learning in the institution had helped me to go through industrial training with confidence.				4	3
FACTO	R: THERAPEUTIC USAGE OF THE DRUGS					
8	The training attended has prepared me well to work in the industry in terms of knowledge.			1	3	3
9	The overall learning in the institution had helped me to go through industrial training with confidence.					43

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	■ Good	5	1	3	6	4	3	4
	■ Excellent	2	5	4		2	3	3

Impact Analysis:

The Internship help the students in the following ways.

- ✓ All the students got Valuable Work Experience
- ✓ Students can present papers in journals/conferences.
- ✓ Students can get placements in Companies.
- ✓ Students can easily choose right career.
- ✓ They will also get multiple Job opportunities.

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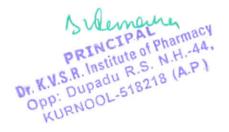
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HANDS - ON LEARNING

B. Pharmacy, Pharm.D & Pharm.D (PB) and M. Pharmacy students gain knowledge intheory by taking part in the pragmatic learning in various laboratories located within the institute. Faculty has designed various experiments according to the syllabus assigned by the JNTUA. Students gain practical awareness through live activities and handling the instruments such as Pharmaceutical Analysis instruments operation UV – Visible Spectrophotometry, HPLC, Pharmacology laboratory experiments with animals, and Pharmaceutical Chemistry etc.,



III/II Pharm.D students learning about Hot Air Oven in the Microbiology Lab.





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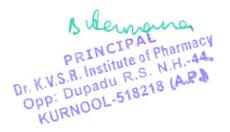
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I/II B.Pharmacy student learning about instrumentation at Pharmaceutical Analysis Laboratory.



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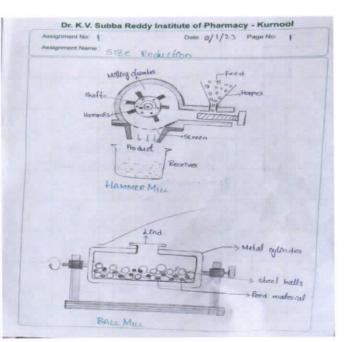
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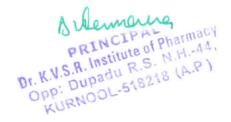
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ASSIGNMENTS

Assignments are the part of the internal examination evaluation process, in which would be immense value as additional learning instruments. Many types of assignments can be given to students of all such as essays, literature reviews, critical reviews, reflective journals, annotated bibliographies and case studies, depends upon the need and learning situations. It implies a task for students to accomplish the aim of learning particular contents, concepts or relationships etc., in this text, learning assignments involve students' independent information seeking and use of a wide range of information resources which are available for them. So every semester/year students are assigned with 2 or 3 topics per each subject regarding to their syllabus and asked to gather more relevant information. Allotment of marks to students is given according to their task completion. Through this student are enriched with knowledge regarding of topic, proof reading of and presentation techniques.

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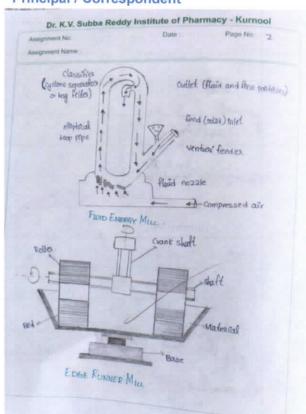
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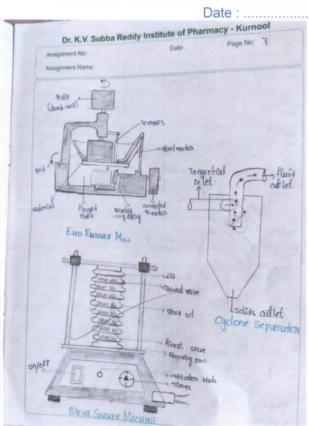
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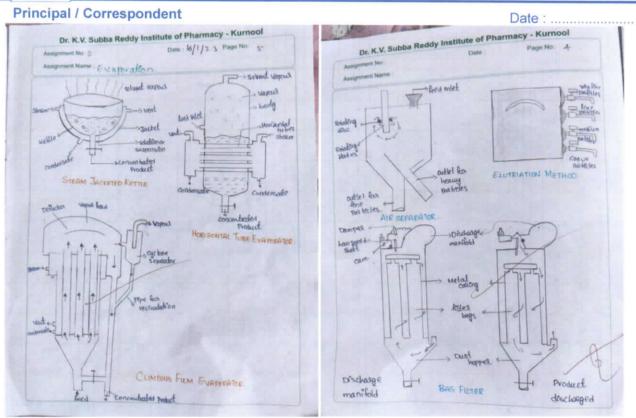


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INNOVATIVE MODEL MAKING:

Students are guided to prepare scientific models through Pharma Expo. The main aim of conducting Pharma Expo is to integrate knowledge within a course, a major, or across an entire Program of study to associate the concepts. This enables students to understand and remember the relationship among various concepts, structures or species.



Model making competition conducted by R.Naga Anjaneyulu IN 2021



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Dr. K.V. Subba Region Tentione of Pharences

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Project based learning is a teaching method were students gain knowledge and skill by
working for an extended period of time to investigate and respond to an authentic, engaging
and complex questions, problems or challenges. It is not only provides opportunities for
students to collaborate with or drive their own learning, but also teaches those skills such as
problem solving and helps to develop additional skills integral to their future such as critical
thinking and time management. So every year IV B. Pharmacy, V year Pharm.D and II year
M. Pharmacy students are allotted with a project under the supervision/guidance of faculty to
be completed within an academic year. Marks are allotted to projects according to their
performance, project results, presentation and viva-voice. Research and review articles of
their projects are published by students in various national and international journals.

PHARIMD V YEAR PROJECT SATCHES:

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	numbers 17ER1T0001		Comparing the outcome in patients of Acute	KVSP/IRB/2021-	100 mine
1	17ER1T0002 17ER1T0003	Dr. G. Nagarajan	Pancreatitis with and Without Prophylactic	2022/Piarm.D/PROJ/021	8-1
	17ER1T0004		Antibiotics in a Tectiary Care Hospital		, /
4	17ER1T0005		The burden of polypharmacy and pattern of co-	KVSP/IRB/2021-	1.19
	17ERIT0007	Dr. B.V. Ramana	morbidities among Chronic Kidney Disease	2022/Phurm. D/PROJ/022	
	17ER1T0008 17ER1T0009		patients in Tertiary Care Hospital		0
3	17ER/T0010	De C. Managian	Analysis of Feto-malernal outcome in	KVSP/IRB/2021-	0-6
	17ER1T0011	Dr. G. Nagamjan	Eclampsia in a Tertiary Care Hospital	2022/Pharm.D/PROJ/023	1
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4.	17ER1T0015 17ER1T0024	Dr. K. Chandrasekhar	manifestations and management of complicated	2022/Pharm.D/PRO1/024	OF S
	17ER1T0016		A Prospective observational study on single vs.	KVSP/IRB/2021-	
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	17ER/T0014		department of a tertiary care hospital	2022 Pharm.D/PRO1/026	101
	17ER1T0025		A descriptive comparison study of	KVSP/IRB/2021-	
7	17ER1T0027 17ER1T0023	Dr. Renuka Tejeswini	effectiveness of cophalosporins and	2022/Pharm.D/PROJ/027	A
		rejeswiii	pencillin antibiotics in community	20.22(1.1111111.D/4.1503/02.7)	ape .
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B.PHARMACY PROJECT LIST (AY:21-22)

Date :				
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		NAME OF THE				SIGNATURE
Batch	Reg.no	STUDENT	GUIDE	TITLE	DEPARTMENT	
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XIII	18ER1R0061	EDIGA SITA SIVA PRASAD GOUD	E. HONEY	INVESTIGATION OF INVITRO ANTI	PHARMACOLOGY	
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XV	18ER1R0087	SHAIK MOHHAMED JAKEER	DR.M.SPURTH Y MITHRA	PHYTOCHEMICAL SCREENING &ANTI MICROBIAL	PHARMACY PRACTICE	
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Principal / Correspondent

Date:

ANALYTICAL METHOD DEVELOPMENT AND VALIDATION OF DAPAGLIFLOZIN IN PURE FORM BY USING UV SPECTROPHOTOMETRY"

A thesis submitted to

JAWAHARLAL NEHRU TECHNOLOGICAL UNIVERSITY

ANANTAPUR



In partial fulfilment of the requirements for the award of degree in

BACHELOR OF PHARMACY

Submitted By

P.SANOVAR KHATUN (18ER1R0036)

P.MAHESH (18ER1R0037)

P.SALIMA (18ER1R0038)

R.VASAVI (

(18ER1R0039)

R.TEJA (18ER1R0040)

S.NUSRATH SHAISTHA (18ER1R0094)

Under the guidance of

Mr. P.T. Nagaraju M. Pharm.

Department of Pharmaceutical Analysis



DR.K.V.SUBBAREDDYINSTITUTEOFPHARMACY



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Principal / Correspondent

Dat	e:		

CERTIFICATE

This is to certify that the dissertation entitled "ANALYTICAL METHOD DEVELOPMENT AND VALIDATION OF DAPAGLIFLOZIN IN PURE FORM BY USING UV SPECTROPHOTOMETRY" was done for the partial fulfillment for the award of degree of bachelor of pharmacy in Pharmaceutical Analysis has been carried out by P.SANOVARKHATUN (18ER1R0036), P.MAHESH (18ER1R0037), P.SALIMA (18ER1R0038), R.VASAVI (18ER1R0039), R.TEJA (18ER1R0040), S.NUSRATH SHAISTHA (18ER1R0094). under the guidance and supervision of Mr. P.T. NAGARAJU, M. Pharm, at DR.K. V. SUBBA REDDY INSTITUTE OF PHARMACY, Dupadu, Kurnool, during the period 2021-2022. It is further certified that this work or any part of this has not been submitted in part or full for the award of any degree or fellowship.

DATE:

PLACE: Dupadu

Principal

Dr. G. NAGARAJAN, M.Pharm, Ph.D.

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Dupadu, Kurnool, A.P – 518002.





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Principal / Correspondent

Date :	

CERTIFICATE BY SUPERVISOR

This is to certify that the dissertation entitled ANALYTICAL METHOD DEVELOPMENT AND VALIDATION OF DAPAGLIFLOZIN IN PURE FORM BY USING UV SPECTROPHOTOMETRY" was done for the partial fulfillment for the award of degree of bachelor of pharmacy in Pharmaceutical Analysis has been carried out by P.SANOVARKHATUN (18ER1R0036), P.MAHESH (18ER1R0037), P.SALIMA (18ER1R0038), R.VASAVI (18ER1R0039), R.TEJA (18ER1R0040), S.NUSRATH SHAISTHA (18ER1R0094) under the guidance and supervision of Mr. P.T. NAGARAJU, M... Pharm, at DR.K.V.SUBBAREDDY INSTITUTE OF PHARMACY, Dupadu, Kurnool, during the period 2021-2022. It is further certified that this work or any part of this has not been submitted in part or full for the award of any degree or fellowship.

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Mr. P.T. NAGARAJU, M..Pharm,

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SENDORSEMENT BY PRINCIPAL

This is to certify that the research project entitled "ANALYTICAL METHOD DEVELOPMENT AND VALIDATION OF DAPAGLIFLOZIN IN PURE FORM BY USING UV SPECTROPHOTOMETRY" was carried out by P.SANOVARKHATUN (18ER1R0036), P.MAHESH (18ER1R0037), P.SALIMA (18ER1R0038), R.VASAVI (18ER1R0039), R.TEJA (18ER1R0040), S.NUSRATH SHAISTHA (18ER1R0094) students of B .pharm 4 th year during the academic year 2021 - 2022, we hereby, certify the adequacy of work to the board examiners, in partial fulfillment for the award of degree of BACHELOR OF PHARMACYas prescribed by Jawaharlal Nehru Technological University Anantapur, Anantapuramu, Andhra Pradesh, has been carried under the supervision of Mr.P.T.NAGARAJU.

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DECLARATION BY CANDIDATES

We P.SANOVARKHATUN (18ER1R0036), P.MAHESH (18ER1R0037), P.SALIMA (18ER1R0038), R.VASAVI (18ER1R0039), R.TEJA (18ER1R0040), S.NUSRATH SHAISTHA (18ER1R0094) pursued admission in B. Pharm 4th year program for the academic year 2019-2020, at Dr. K. V.Subba Reddy Institute of Pharmacy, Dupadu, Kurnool (Affiliated to JNT University Anantapur, Anantapuramu) and in connection with partial fulfilment for award of degree of Bachelor of Pharmacy. We hereby declare that the entitled research work "ANALYTICAL METHOD DEVELOPMENT AND VALIDATION OF DAPAGLIFLOZIN IN PURE FORM BY USING UV SPECTROPHOTOMETRY" was carried out by us undertheguidance of Mr. P.T. NAGARAJU, M., Pharm., Asst. Professor, Department of Pharmaceutical Analysis. We express no conflict of interest for due submission of this work to board of examiners for evaluation process.

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AN EXPRESION OF GRATITUDE

With God All Things Are Possible"

Milestones in life are achieved, not by individual efforts but by blessings and guidance of elders, near and dear ones. This project is the product of collective wisdom and experience of all those who have shared their views far beyond those found within the covers of book. We therefore take this opportunity to express our acknowledgements to all of them.

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We express our deep sense of gratitude to Dr.K.V.Subba Reddy, Chairman of Dr.K.V.Subba Reddy Group of institutions and Srmt. K.Vijayalakshmamma Correspondent of Dr.K.V.Subba Reddy Group of institutions supporting us in providing all equipments and re quired things. And our sincere thanks to Dr.G.Nagarajan M.Pharm, Ph.D. Prinicpal, Dr.K.V.Subba Reddy Institute of Pharmacy who has been a constant guide to us throughout this work and taught us the nuances of experimentation and the interpretation of results which has shaped up our dissertation in to its presentform.

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CONTENTS

S.NO	TITLE	PAGE NO.
1	INTRODUCTION	1
2	OBJECTIVE	20
3	REVIEW OF LITERATURE	22
4	DRUG PROFILE	28
5	MATERIALS AND INSTRUMENTS	30
6	METHODOLOGY	32
7	RESULTS AND DISCUSSION	36





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LIST OF TABLES

Date :

S.NO	TITLE	PAGE NO.
1.1	SELECTION OF SOLVENT	6
5.1	MATERIALS	30
5.2	INSTRUMENTS	31
7.1	UV SPECTROSCOPY METHOD	36
7.2	CALIBRATION DATA OF DAPGLIFLOZIN	37
7.3	INTRADAY PRECISION	38
7.4	INTER DAY PRECISION	39
7.5	OBSERVATION FOR ACCURACY STANDARD(50µg/ml)	40
7.6	OBSERVATION FOR ACCURACY STANDARD 80%(45µg/ml)	40
7.7	OBSERVATION FOR ACCURACY STANDARD100% (55µg/ml)	41
7.8	OBSERVATION FOR ACCURACY STANDARD100% (65µg/ml)	41
7.9	FOR ACCURACY SUMMARY	42
7.10	OBSERVATION OF LIMIT OF DETECTION	42
7.11	OBSERVATION OF LIMIT OF QUANTIFICATION	42

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rincipal /	Correspondent	Date :
7.12	FOR RUGGEDNESS	43
7.13	FOR230 AND 234 WAVELENGHTS	43
7.14	ROBUSTNESS SUMMERY	44





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LIST OF FIGURES

Date	

S.NO	TITLE	PAGE NO.
1.1	INSTRUMENTION	4
7.1	SELECTION OF WAVE LENGTH	33
7.2	CALIBRATION CURVE OF DAPAGLIFLOZIN	37





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LIST OF ABBREVIATION

Abbreviation	Full Name
AR	Analytical Reagent
BP	Boiling Point
MW	Molecular Weight
FDA	Food and Drug Administration
ICH	International Conference of Harmonization
ml	Millilitre
μg	Microgram
M	Molar
L	Litre
Min	Minute
LR	Laboratory reagent
RSD	Relative Standard Deviation
UV	Ultraviolet
nm	Nanometer
ng	Nanogram Nanogram
mg	Milligram
NMT	Not More Than
S	Slope



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S/N	Signal to noise ratio	
r^2	Correlation Coefficient	
LOD	Limit of Detection	
LOQ	Limit of Quantitation	
Q.C	Quality Control	
V/v	Volume by volume	
W/v	Weight by volume	



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ABSTRACT

simple. specific, accurate and precise Spectroscopy method A was developed and validated for the estimation of dapagliflozin in Pure form. The Standard solution was prepared by weighing 100 mg of dapagliflozin in 100 ml volumetric flask with 0.1N Nitric Acid. The final Standard solution was made to produce 1000 µg / ml with 0.1N Nitric Acid. Further dilutions were prepared as per procedure and were scanned at 232 nm. The linearity was found in the concentration range of 10-60 µg / ml. The Correlation coefficient was 0.996. The regression equation was found to be Y = 0.043 X - 0.336. The method was validated for linearity, accuracy, precision, limit of detection, limit of quantitation and ruggedness robustness. The limit of detection and limit of quantitation for estimation of dapaliflozin was found to be 5.36 (µg / ml) and 17.08 (µg / ml), respectively. The percentage recovery of dapaliflozin was found to be in the range of $98.49 \pm 0.0001 - 101.3 \pm 0.003$. Proposed method can be successfully applied for the quantitative determination of dapagliflozin in pharmaceutical Pure form





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1. INTRODUCTION

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Analytical chemistry ⁽¹⁾ is often described as the area of chemistry responsible for characterizing the composition of matter, both qualitatively (what is present) and quantitatively (how much is present). Analytical chemistry is not a separate branch of chemistry, but simply the application of chemical knowledge.

Pharmaceutical Analysis ⁽²⁾ is the branch of chemistry involved in separating, identifying and determining the relative amounts of the components making up a sample of matter. It is mainly involved in the qualitative identification or detection of compounds and quantitative measurements of the substances present in bulk and pharmaceutical preparation.

The technique ⁽³⁾ employed in quantitative analysis is based upon the quantitative performance of suitable chemical reactions and either measuring the amount of reagent needed to complete the reaction, or ascertaining the amount of reaction product obtained.

Quality ⁽⁴⁾ is important in every product or service but it is vital in medicine as it involves life. Unlike ordinary consumer goods there can be no "second quality" in drugs. Quality control is a concept, which strives to produce a perfect product by series of measures designed to prevent and eliminate errors at different stages of production.

Physico-chemical methods ^(5, 6) are used to study the physical phenomenon that occurs as a result of chemical reactions. Among the Physico-chemical methods, the most important are optical (Refractometry, Polarimetry, Emission, Fluorescence methods of analysis, Photometry including Photocolorimetry and Spectrophotometry covering UV-Visible and IR regions and Nephelometry or Turbidimetry) and chromatographic (Column, Paper, TLC, GLC, HPLC) methods. Methods such as Nuclear Magnetic Resonance and Para Magnetic Resonance are becoming more and more popular. The combination of Mass Spectroscopy with Gas Chromatography and Liquid Chromatography are the most powerful tools available. The chemical methods include the gravimetric and volumetric procedures which are based on



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complex formation; acid - base, precipitation and redox reactions. Titrations in non-aqueous media and complexometry have also been used in pharmaceutical analysis.

The number of new drugs is constantly growing. This requires new methods for controlling their quality. Modern pharmaceutical analysis must need the following requirements.

- 1. The analysis should take a minimal time.
- 2. The accuracy of the analysis should meet the demands of Pharmacopoeia.
- 3. The analysis should be economical.
- 4. The selected method should be precise and selective.

These requirements are met by the Physico-chemical methods of analysis, a merit of which is their universal nature that can be employed for analyzing organic compounds with a diverse structure. Of them, Visible Spectrophotometry is generally preferred especially by small scale industries as the cost of the equipment is less and the maintenance problems are minimal.

Instrumental methods of Chemical analysis:

Instrumental method is an exciting and fascinating part of chemical analysis that interacts with all areas of chemistry and with many other areas of pure and applied sciences. Analytical instrumentation plays an important role in the production and evaluation of new products and in the protection of consumers and environment. This instrumentation provides lower detection limits required to assure safe foods, drugs, and water air. Instrumental methods are widely used by Analytical chemists to save time, to avoid chemical separation and to obtain increased accuracy.

SPECTROSCOPY (7)

Spectroscopy is the measurement and interpretation of Electro Magnetic Radiation (EMR) absorbed



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Date : or emitted when the molecule or atoms or ions of a sample move from one energy state to another energy state. This change may be from ground state to excited state or excited state to ground state. At ground state, the energy of a molecule is the sum of rotational, vibrational and electronic energy. In other words, Spectroscopy measures the changes in rotational, vibrational and / or electronic energies.

Ultraviolet Spectroscopy:

Ultraviolet Spectroscopy is concerned with the study of absorption of UV radiation which ranges from 200 nm to 400 nm. Any molecule has n, π or σ combination of these electrons. These bonding (σ and π) and non bonding (n) electrons absorb the characteristic radiation and undergoes transition from ground state to excited state.

Visible Spectroscopy (Colorimetry):

Colorimetry is concerned with the study of absorption of visible radiation whose wavelength ranges from 400 nm to 800 nm. Any coloured substance will absorb radiation in this wavelength region. Coloured substances absorb light of different wavelength in different manner and hence we get an absorption curve (absorbance Vs wavelength). In this absorption curve, the wavelength at which maximum absorption of radiation takes place is called as λ_{max} . This λ_{max} is characteristic or unique for every coloured substance and this is a qualitative aspect, useful in identifying the substance.





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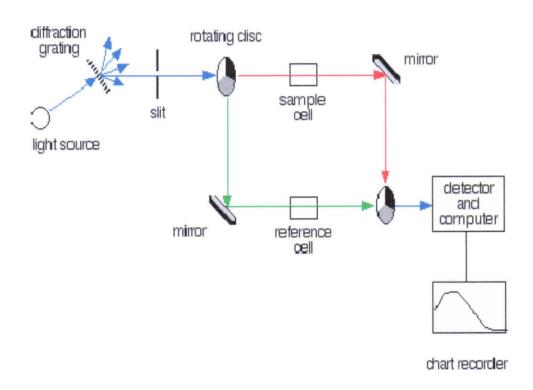


Fig No: 1.1. UV Visible double beam Spectrophotometer Laws governing absorption of radiation:

Beer's law (related to concentration of absorbing species)

'The intensity of a beam of monochromatic light decreases exponentially with increase in the concentration of absorbing species arithmetically'.

Lambert's law (related to thickness / path length of absorbing species)

'The rate of decrease of intensity (monochromatic light) with the thickness of the medium is directly proportional to the intensity of incident light'.

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Limitations of Beer-Lambert's Law:

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The Beer-Lambert law is rigorously obsessed provided a single species gives rise to the observed absorption. However the law may not obsessed when,

- 1. Different forms of the absorbing molecules are in equilibrium.
- 2. Solute and solvent from association complexes.
- 3. There is a thermal equilibrium between ground electronic state and a low lying excited state.
- 4. The compounds are charged by irradiation (fluorescent compounds).

Deviations from Beer's Law:

According to Beer's law, a straight line passing through the origin should be obtained, when a graph is plotted between absorbance (A) and concentration (C). Deviation from the law may be positive or negative, according to whether the resulting curve is concave upwards or concave downwards. The deviations from the Beer's law may be due to interaction of the solute molecules with each other or with the solvent or may be due to instrumental factors. The most important reasons that cause deviations are

- 1. Negative deviation can always be expected when the illumination is not monochromatic.
- 2. The presence of impurities that fluoresce or absorb at the required absorption wavelength.
- 3. Environmental errors such as solvent, temperature and pressure.





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- Date :
- Chemical factors such as change in pH and chemical equilibrium, presence of complexing agent, competitive metal ion reactions and concentration dependence.
- 5. Refractive index of sample.
- Instrumental errors such as radiation, stability of radiation source, stability of slit control and electronics and reliability of the optical parts.

Choice of solvent (8):

Several solvents used in Ultraviolet Spectroscopy with their cut off wavelengths are listed in Table: 1.1.

A suitable solvent for UV Spectroscopy should meet the following requirements.

a. It should not absorb radiations in the region under investigation.





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b. It should be less polar, so that it has minimum interaction with the solute molecules.

c. The solvent used should be of high purity.

Table No: 1.1. List of common solvents used in UV Spectroscopy

Solvents	Cut off wavelength(nm)
Acetonitrile	190
Water	191
Cyclohexane	195
Hexane	201
Methanol	203
Ethanol	204
Ether	215
Methylene dichloride	220
Chloroform	237
Carbon tetrachloride	257

Detectors:

A detector is a transducer that convert EMR into an electron flow and subsequently, into a current flow or voltage in the readout circuit. Photoelectric or Photo multiplier tubes are



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Date:.... generally used as detectors. The detector must have the following important requirements. It must respond to radiant energy over a broad wavelength range.

- a. It should be sensitive to low levels of radiant power.
- b. It should rapidly respond to the radiation and produce an electrical signal that can be readily amplified.
- It should have relatively low noise level (for stability). c.
- d. The signal produced is directly proportional to the power of beam striking it.





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For Accuracy:

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Statistical procedures and representative calculations:

	The consistence	y and	suitability	of th	e developed	method	are	substantiated	through	the
statisti	cal analysis like	stand	ard deviation	on, rel	ative standar	rd deviati	ion.			

The second secon
☐ Standard Deviation = ☐ =
Where, $\mathbf{x} = \text{Sample}$,
x_i = Mean value of
samples, $n = Number of$
samples
\square Relative Standard Deviation = $\square / x_i \times 100$
\square Molar extinction coefficient (mol ⁻¹ cm ⁻¹) = A / C ×
LWhere, $A = Absorbance$ of drug
C = Concentration of drug, L = Path length
- G
□ Sandell's sensitivity (mcg / cm ² / 0.001 absorbance units) = $C / A \times C$
0.001 Where, C = Concentration of drug





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Principal / Correspondent A = Absorbance of drug	Date :
A - Absorbance of drug	
Unit- $(mcg / cm^2 = 0.001 absorbance)$	
Coefficient of variance (σ) = $\sum (x - x^{-})^{2} / n-1$	
\square Regression equation $y = a + bx$	
\square Slope = y / x	
Where, $x = Concentration$	
y = Absorbance, a = Intercept	
\square Limit of detection: (DL) = 3.3 × \square / S	
Units- (µg / ml)	
Where, \Box = Standard deviation of the response.	
S = Slope of the calibration curve.	
The slope S may be estimated from the calibration curve of the analyte.	
The estimate of \square may be carried out in a variety of ways.	
□ Limit of quantitation (QL) = $10 \times \square$ /	
INST. OS	

Date :



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SUnit- (µg / ml)

Where, \Box = Standard deviation of the response

S = Slope of the calibration curve

The slope S may be estimated from the calibration curve of the analyte. The estimation of \Box may be carried out in a variety of ways.

Basic criteria for new method development of drug analysis (10)

- The drug or drug combination may not be official in any pharmacopoeias,
- A proper analytical procedure for the drug may not be available in the literature due to patent regulations,
- Analytical methods may not be available for for the drug in the form of aformulation due to the interference caused by the formulation Excipients,
- Analytical methods for the quantification of the drug in biological fluids may notbe available,
- Analytical methods for a drug in combination with other drugs may not be available,



· Limit of Detection

All of the above mentioned methods were developed and validated statistically to ensure their Accuracy, Precision, Linearity, Ruggedness and other analytical method validation parameters as mentioned in the various guidelines.

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formulation. The recovery study was carried out at \(\subseteq \text{ve different levels and results were found to be satisfactory. Conclusion: The results of estimation and validation parameters like accuracy, precision, ruggedness, linearity and range were studied for all the developed methods and were found to be within limits. The proposed method can be adopted for routine quality control for estimation of drug in formulation.

Key words: Dapagli \(\subseteq \text{czin}, \text{UV-Visible spectroscopy}, \text{Validation}, \text{Derivative}, \(\text{Area under curve} \)

Area under curve Spectrophotometry ABSTRACT

Aim: Simple, precise and accurate UV-spectrophotometric methods for estimation of Dapagli ozin were developed and validated as per ICH guidelines. Experimental and Results: These methods includes Calibration curve, Area under curve (AUC), First and Second order derivative method based on measurement of absorbance at a selected wavelengths using UV-visible on measurement of absorbance at a selected wavelengths using UV-visible spectrophotometer with 1 cm matched quartz cell and methanol with water as a solvent. All developed methods obeyed Beer's-lambert's law in the concentration range of 5-40 μg/mL, with correlation coef cient value less than 1. The percent amount of drug estimated by these methods was nearly 100%, found to be in good agreement with label claim of marketed tablet formulation. The recovery study was carried out at we different levels and results were found to be satisfactory. Conclusion: The results of estimation and validation parameters like accuracy, precision, ruggedness, linearity and range were studied for all the developed methods and were found to be within limits. The proposed method can be adopted for routine quality control.

range were studied for all the developed methods and were found to be within limits. The proposed method can be adopted for routine quality control for estimation of drug in formulation.

Key words: Dapagli □ ozin, UV-Visible spectroscopy, Validation, Derivative, Area un Aim: Simple, precise and accurate UV-spectrophotometric methods for estimation of Dapagli □ ozin were developed and validated as per ICH guidelines. Experimental and Results: These methods includes Calibration curve, Area under curve (AUC), First and Second order derivative method based on measurement of absorbance at a selected wavelengths using UV-visible spectrophotometer with 1 cm matched quartz cell and methanol with water as a solvent. All developed methods obeyed Beer's-lambert's law in the concentration range of 5-40 μg/mL, with correlation coef □ cient value less than 1. The percent amount of drug estimated by these methods was nearly 100%, found to be in good agreement with label claim of marketed tablet formulation. The recovery study was carried out at □ ve different levels and results were found to be satisfactory. Conclusion: The results of estimation and validation parameters like accuracy, precision, ruggedness, linearity and

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Lambert's Law In The Concentration Range Of 5-40 Mg/Ml, With Correlation Coefocient Value Less Than
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Agreement With Label Claim Of Marketed Tablet Formulation. The Recovery Study Was Carried Out At
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And Validation Parameters Like Accuracy, Precision, Ruggedness, Linearity And Range Were Studied For
All The Developed Methods And Were Found To Be Within Limits. The Proposed Method Can Be
Adopted For Routine Quality Control For Estimation Of Drug In Formulation.

K. Bhavyasri.et.al^[14]

A New, Simple, Precise, Accurate, Reproducible And Economic Stability Indicating UV Spectroscopic Method Was Developed And Validated For Simultaneous Estimation Of Dapagliflozin And Metformin In Pure And Combined Pharmaceutical Dosage Form. The UV Spectrophotometric Estimation Of Dapagliflozin And Metformin Was Determined Using The Q Absorption Ratio Method At 222 Nm And 232 Nm Respectively Using Water As Diluent. The Linearity Ranges For Dapagliflozin And Metformin Was 2 – 32 Mg/Ml And 1 – 20μg/Ml Respectively With Their Correlation Coefficient Values (R2) 0.999. The Percentage Recovery At Various Concentration Levels Varied From 96.82 - 99.8 % For Dapagliflozin And 98.15 To 99.35 % For Metformin Confirming That The Method Is Accurate. LOD And LOQ For Dapagliflozin Was Found To Be 0.0241μg/Ml And 0.0293 Mg/Ml And For Metformin 0.0732 Mg/Ml And 0.0890 Mg/Ml. In The Precision Study, The% RSD Value Was Found To Be 0.1845 % And 0.2052 % For Dapagliflozin And Metformin Respectively. Degradation Studies Were Performed, Both The Drugs Were Found To Be Degraded In Acid By Using Hydrochloric Acid, In Base Using Sodium Hydroxide Solution, In Peroxide Using Hydrogen Peroxide Solution, Temperature And In Light By Exposing It To UV Light In UV Chamber. The Results For Estimation Of Dapagliflozin



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And Metformin Hydrochloride And Validation Parameters Like Accuracy, Precision, Ruggedness,
Linearity Were Studied For The Method And Were Found To Be Within The Limits. The Developed
Method Was Free From The Interferences Due To Excipients Present In The Formulation And It Can
Be Used For Routine Quality Control Analysis.

4.Gajanan Vithoba Mante.et.al^[15]

Aim: Simple, Precise And Accurate UV-Spectrophotometric Methods For Estimation Of Dapagliflozin Were Developed And Validated As Per ICH Guidelines. Experimental And Results: These Methods Includes Calibration Curve, Area Under Curve (AUC), First And Second Order Derivative Method Based On Measurement Of Absorbance At A Selected Wavelengths Using UV-Visible Spectrophotometer With 1 Cm Matched Quartz Cell And Methanol With Water As A Solvent. All Developed Methods Obeyed Beer's-Lambert's Law In The Concentration Range Of 5-40 Mg/Ml, With Correlation Coefficient Value Less Than 1. The Percent Amount Of Drug Estimated By These Methods Was Nearly 100%, Found To Be In Good Agreement With Label Claim Of Marketed Tablet Formulation. The Recovery Study Was Carried Out At Five Different Levels And Results Were Found To Be Satisfactory. Conclusion: The Results Of Estimation And Validation Parameters Like Accuracy, Precision, Ruggedness, Linearity And Range Were Studied For All The Developed Methods And Were Found To Be Within Limits. The Proposed Method Can Be Adopted For Routine Quality Control For Estimation Of Drug In Formulation.

5. Mohd.Zameeruddin. et.al^[16]

Aim Simple, Precise And Accurate UV-Spectrophotometric Simultaneous Equation Method For Estimation Of Dapagliflozin And Saxagliptin Were Developed And Validated As Per ICH Guidelines.



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Experimental And Results The Objective Of The Work Is To Develop UV Spectroscopic Method For Simultaneous Estimation Of Dapagliflozin (DAPA) And Saxagliptin (SAXA). This Method Involve Solving Of Simultaneous Equations Based On Measurement Of Absorbance At Two Wavelengths 223 Nm And 212 Nm. Both The Drugs Obey The Beer's Law In The Concentration Ranges 4-24 μg/Ml And 5-50 μg/Ml Respectively. Results Of The Methods Were Validated Statistically. Novel, Simple, Sensitive, Rapid, Accurate And Economical Spectrophotometric Methods Have Been Developed For Simultaneous Estimation Of Dapagliflozin And Saxagliptin .The Method Can Be Used To Estimate The Amount Of Dapagliflozin And Saxagliptin In Mixture Containing Dapagliflozin And Saxagliptin.

6. Dr. K. Bhavyasri. et.al^[17] A New, Simple, Precise, Accurate, Reproducible And Economic Stability Indicating Spectroscopic Method Was Developed And Validated For Simultaneous Estimation Of Dapagliflozin And Metformin In Pure And Combined Pharmaceutical Dosage Form. The UV Spectrophotometric Estimation Of Dapagliflozin And Metformin Was Determined Using The Simultaneous Equation Method At 222 Nm And 232 Nm Respectively. The Linearity Ranges For Dapagliflozin And Metformin Were 2 – 32 Mg/Ml And 1 – 20μg/Ml Respectively With Their Correlation Coefficient Values (R2) 0.999. The Percentage Recovery At Various Concentration Levels Varied From 96.82 - 99.8 % For Dapagliflozin And 98.15 To 99.35 % For Metformin Confirming That The Method Is Accurate. LOD And LOQ For Dapagliflozin Was Found To Be 0.0241 Mg/Ml And 0.0293 Mg/Ml And For Metformin 0.0732 Mg/Ml And 0.0890 Mg/Ml. In The Precision Study, The% RSD Value Was Found To Be 0.1845 % And 0.2052 % For Dapagliflozin And Metformin Respectively. Degradation Studies Were Performed, Both The Drugs Were Found To Be Degraded In Acid, Base, Peroxide, Temperature And Light. The Proposed Method Can Be Applied Successfully For The Simultaneous Estimation Of Both Drugs In Quality Control Laboratories.





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7. M. R. Usman.et.al^[18] In The Present Work, A Simple, Rapid, Sensitive, Precise, And Reproducible, Specific UV Spectrophotometric Method For The Determination Of Saxagliptin (SAXA) And Dapagliflozin (DAPI) In Bulk Drug And Pharmaceutical Dosage Form Were Developed And Validated. Methods: A Simple Double Beam UV Spectrophotometric Method Has Been Developed And Validated With Different Parameters Such As Linearity, Precision, Repeatability, The Limit Of Detection (LOD), Limit Of Quantification (LOQ), Accuracy As Per ICH Guidelines. Results: UV-Visible Spectrophotometric Method, Measurement Of Absorption At A Maximum Wavelength In 10 Ml Methanol And Volume Make With Water Solvent System As Reference SAXA And DAPI Were Found To Be At 224 Nm And 274 Nm Respectively. The Drug Obeyed The Beer's Law And Showed A Good Correlation. Beer's Law Was Obeyed In The Concentration Range 2-10 μg/Ml For Saxagliptin And 4-20 μg/Ml For Dapagliflozin, Respectively With A Correlation Coefficient Was 0.999. The LOD And LOQ Of Saxagliptin Were Found To Be 0.040 Mg/Ml And 0.01230 Mg/Ml, Dapagliflozin Was Found To Be 0.1230 Mg/Ml And 0.5460 Mg/Ml, Respectively. Percentage Assay Of SAXA And DAPI In Tablets. Conclusion: The Proposed Method Is Simple, Precise, Accurate, And Reproducible Can Be Used For Routine Analysis Of Saxagliptin And Dapagliflozin In Bulk And Tablet Dosage Form.

8.Karuna Priya Chitra.et.al^[19] Dapagliflozin (DAP) Is Indicated For The Management Of Diabetes Mellitus Type 2, And Functions To Improve Glycemic Control In Adults When Combined With Diet And Exercise1. DAP Is An Inhibitor Of Sodium-Glucose Cotransporter 2 (SGLT2) Responsible For The Majority Of The Reabsorption Of Filtered Glucose From The Tubular Lumen. By Inhibiting SGLT2, DAP Reduces Reabsorption Of Filtered Glucose And Lowers The Renal Threshold For Glucose, And Thereby Increases Urinary Glucose Excretion2. In Present Work, A Selective, Specific, Sensitive And Economical UV Spectroscopic Method Has Been Developed For The Estimation Of Dapagliflozin In Bulk And Its Pharmaceutical Dosage Forms. An Absorption Maximum Was Found To Be At 233.65 Nm. Dapagliflozin Obeyed Beer's Law In The Concentration Range From 10-35 μg / Ml. Proposed



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Method Was Validated According To ICH Guidelines And Values Of Accuracy, Precision And Other
Statistical Analysis Were Found To Be In Good Accordance With The Prescribed Values With
Correlation Coefficient Of 0.9998. The Percentage Recovery Of Dapagliflozin Ranged From 99.7 In
Pharmaceutical Dosage Form. Results Of The Analysis For Accuracy, Precision, LOD, LOQ And Were
Found To Be Satisfactory. The Proposed Method Is Simple, Rapid And Suitable For The Routine
Quality Control Analysis.

9. R. Aswini.et.al[²⁰] Dapagliflozin And Saxagliptin Are Very Effectively Used Treatment For Type II Diabetes. They Are Very Potent Inhibitors Of Renal Glucose Reabsorption And Dipeptidyl Peptidase Protein 4(DPP-4) And Sodium Glucose Transport Protein 2 And Also They Are Called As DPP4 & SGLT2 Inhibitors. They Are Generally Administered As Tablets. Determination Of Dapagliflozin And Saxagliptin In Pharmaceutical Dosage Form And Bulk Form, Several Analytical Methods Including UV, HPLC, LC-MS And HPTLC Techniques Has Been Developed. Methods Indicating Human Plasma Stability And Impurity Profiling Are Also Described For Both Drugs. For Qualitative And Quantitative Estimation Of Dapagliflozin And Saxagliptin, These Analytical Methods Can Be Used And It Can Also Be Used For Its Related Degradants In Bulk Formulations And Biological Fluids. The Following Study Depicts The Review On Analytical Methods Which Includes Estimating The Antidiabetic Drugs.

10.G. V. Mante.et.al^[21] Rapid, Precise And Accurate RP-HPLC Method For Estimation Of Dapagliflozin From Its Tablet Dosage Form Was Developed And Validated As Per ICH Guidelines. The Chromatographic Separation Was Achieved By Isocratic Mode With A Mixture Of Acetonitrile: 0.1% Triethylamine (Ph-5.0) In The Ratio Of 50:50v/V As Mobile Phase Using Princeton C18column At Flow Rate Of 1ml/Min And Detection Wavelength Of 224nm. Using Optimized Chromatographic Conditions, Retention Time Of Drug Was Found To Be 5.163min. The Proposed Method Obeyed Beer's-Lambert's Law In The Concentration Range Of 10-70μg/Ml, With Correlation Coefficient Value 0.999.The Mean Percent Amount Of Drug Estimated Was 100.57%, Found To Be Good In Agreement With Label Claim Of Marketed Tablet Formulation. The Validation Parameters Like Accuracy,



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Precision,	Ruggedness,	Robustness,	Linearity	And	Range	Were	Studied	For	Proposed	Method	And
Were Fou	nd To Be Wit	hin Limits. S	stress Test	ing U	nder V	arious	Conditio	ns S	uch As Ph	(Acid/B	ase),
Oxidation	, Temperature	, Light, Humi	dity, Etc.	Was A	Also Ca	rried C	Out				

11. Shilpi Pathak. et.al^[22] Dapagliflozin Is An Antidiabetic Drug That Works On The Kidneys Of Reabsorption Of Glucose In Kidneys By The Sodium-Glucose Co-Transporter Offer. It Is Used In Patients With Type 2 Diabetes. It Is Administered As Tablets. It Has Several Analytical Papers For Estimation Of Active Pharmaceutical Ingredient (API) Or Drug Formulation By Reverse Phase-High Performance Liquid Chromatography (RP-HPLC) And Ultraviolet Spectroscopy (UV). It Is Very Challenging To Use Of Chemicals, Drugs, And Solvent Of Separation Methods Used In The Pharmaceutical Product To Green Chemistry. This Review Mostly Used Dihydrogen Phosphate Buffer And Other Toxic Reagents For Estimation And These Agents Harm Instruments, As Well As, Environment And A Lot Of Waste So That Novel Analytical Techniques For Quantifying And Defining Dapagliflozin Should Be Built As Easy As Possible And Secure For The Individual And The Community. This Review Pays Attention To The Critical Condition Of Physicochemical, Properties, Action, And Aims To Focus On Different Analytical Methods For The Estimation Of Dapagliflozin In Pharmaceutical Formulations.

4. DRUG PROFILE [23]

MOLECULAR FORMULA: C21H25CLO6







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CHEMICAL NAME

STRUCTURE

:BMS -512148;[1S]-1,5-anhydro-1-C-{4-chloro -3-[4-ethoxy

Phenyl] methyl]phenyl}-D-glucitol

CATEGORY : Antidiabetic

MOLECULAR WEIGHT: 408.873mg/ml

MELTING POINT : 74-78 °C

DESCRIPTION :

SOLUBILITY :ethanol, DMSO,Dimethyl formamide





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PHARMACOKINETIC DATA

BIOAVAILADILITY : ORALLY

PROTIN BINDING : 91%

HALF LIFE : 12.9hrs

METABOLISM : HEPATIC

EXCRETION : RENAL EXCRETION

MECHANISM OF ACTION

upadu

- Sodium-glucose cotransporter 2, expressed in the proximal renal tubules, is responsible for the majority
 of the reabsorption of filtered glucose from the tubular lumen
- Dapagliflozin is an inhibitor of SGLT2. By inhibiting SGLT2, Dapagliflozin reduces reabsorption of filtered glucose and thereby promotes urinary glucose excretion
- Dapagliflozin also reduces sodium reabsorption and increases the delivery of sodium to the distal tubule.
 This may influence several physiological functions including, but not restricted to, lowering both preand afterload of the heart and downregulation of sympathetic activity, and decreased intraglomerular pressure, which is believed to be mediated by increased tubuloglomerular



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Uses: Used to treat type-2 diabetes mellitus.

5. MATERIALS AND INSTRUMENTS

The following materials used were either AR/LR grade or the Possible Pharma grade available as supplied by the manufacturer or supplier without further purification or investigation.

Table No: 5.1 Materials

S. No	Materials	Source
1	Dapagliflozin	Aurobindo pharma Pvt.Ltd, Jadacherla.
2	Methanol	Merck Mumbai Ltd, Mumbai.
3	Glacial acetic acid	Merck Mumbai Ltd, Mumbai.
4	Sulphuric acid	Merck Specialities Pvt Ltd, Mumbai.
5	Sodium hydro oxide	Merck Specialities Pvt Ltd, Mumbai.
6	Nitric acid	Merck Specialities Pvt Ltd, Mumbai.
7	Water	Nishanth, Kurnool
8	Ethanol	Merck Specialities Pvt Ltd, Mumbai.





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9	Choloroform	Merck Specialities Pvt Ltd, Mumbai.	

Table No: 5.2 Instruments

S. No	Equipments	Source
1	UV Spectrophotometer	Analytical Technologies Limited 2080N
2	Sonicator	Wensar

6. METHODOLOGY

6.1 UV SPECTROSCOPY

Analytical Technologies Limited UV-VIS 2080N Spectrophotometer was used with 1c m matched quartz cells. The data processing was performed using UV- probe software.

UV method development:

The parameters for the development were as follows

- 1] Linearity
- 2] Accuracy
- 3] Precision
- 4] Robustness





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- 5] Ruggedness
- 6] Limit of detection
- 7] Limit of quantification

Selection of solvents:

In order to select suitable solvent for determination of Dapagliflozin various solvent methanol, glacial acetic acid, sodium hydroxide, sulphuric acid, Nitric acid, ethanol tried for the solubility studies and it was found that Dapagliflozin was freely soluble in 0.1N HNO₃ the present investigation distilled water was selected as a solvent.

Selection of wavelength:

10mg/ml of Dapagliflozin was scanned in the range of 200-400nm.





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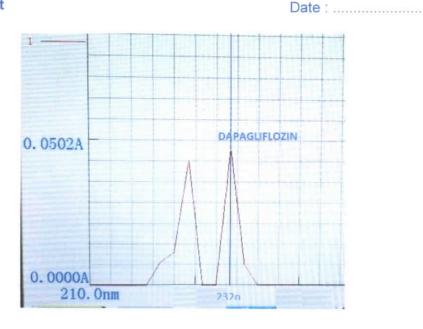


Fig.No. 6.1 UV spec

VALIDATION OF THE METHOD:

The method was validated in terms of parameters like linearity, accuracy, precision, LOD, LOQ, ruggedness, and robustness.

Preparation of 0.1n nitric acid:

Take 63ml of Concentrated Nitric Acid and make upto to 1000 ml with water.

Preparation of stock solution:

100mg of Dapagliflozin was dissolved in 0.1n nitric acid in a 100 ml volumetric flask and solution was made upto volume with nitric acid.

Preparation of standard working solution:



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10ml of standard solution	was dissolved in	100ml of volumetric	flask and the solution	was made up to
volume with 0.1N HNO ₃ .				

1.LINEARITY

To evaluate the linearity, serial dilution of analyte were prepared from the standard working solution was diluted with solvent to get a series of concentration ranging from 10,20,30,40,50 and 60 micro gram\ml. the prepared solution were filtered through whatman filter paper [NO.41]. Calibration curve was constructed by plotting the absorbance on y-axis against the concentration on x-axis [table no;7.2].

2.PRECISION

The precision of analysed method was studied by analysis of multiple sampling of homogeneous sample.

The precision is expressed as standard devition [or] relative standard deviation. The presicion of the method was demonstrated by intra-day and inter- day variation studies.

2.1 Intraday-Precision:

In The Intraday Studies, the Standard Solutions (40mg/ml) Was Analysed for 6 Times in different time Interval with in day . %RSD was Calculated presented in table 7.3

2.2 Inter day Precision:

In the Inter-day variation studies , the standard solution (40mg/ml) was Analysed for 6 times n different days . %RSD was Calculated Presented In 7.4

3. Accuracy:



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Recovery Studies By The Standard Addition Method Performed with a View to Justify the accuracy of
proposed method . previously analysed sample of Dapagliflozin (45,55and 65microg/ml) were spiked
with 80,100,120% extra Dapagliflozin standard and the mixture were analysed by the proposed method.
The experiment was performed in triplicate and recovery of the pure. %RSD was calculated and
reported in table 7.9.

4. Sensitivity:

The sensitivity of measuring of Dapagliflozin by use of the proposed method was estimated in terms of limit od detection [LOD] and the limit of quantitation {LOQ}. The LOD and LOQ were calucated by the use of equation LOD=3.3 x σ /s /s and LOQ=10 x σ /s where σ is the standard deviation of response and S is the slope of the calibration curve LOD and LOQ values are reported in table7.10 and 7.11

5.RUGGEDNESS:

Ruggedness is the measure of the reproducibility of a test result under normal expected operating condition from instrument to instrument and analyst to analyst. The ruggedness of the method was determined by carrying out the experiment by different operations. The result of ruggedness testing is reported in the table 7.12

6.ROBUSTNESS:

Robustness is a measure of capacity of a method to remain unaffected by small but deliberate variation in the method condition, and is indication of the reliability of the method. A method is robustness, if it is unexpected by small changes in operating condition. To determine the robustness of this method, the experimental condition where deliberately altered at 3 different levels and responses were evaluated. Variation of wave length[230nm and 234nm] had no significant effect and the absorbance of 40 μ g/ml Solution, indicating that the method was robustness. The result are shown in table 7.13 & 7.1





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7. RESULTS & DISCUSSION

UV SPECTROSCOPY METHOD

Table No: 7.1 Characteristic parameters of Dapagliflozin for the proposed UV spectroscopy method:

Parameters	UV
Calibration range (µg/ml)	10-60(μg/ml)
Wavelength	232nm
Regression equation (y*)	0.0435x
Slope	0.0435
Correlation co efficient(r ²)	0.994
LOD (µg/ml)	5.36
LOQ (μg/ml)	17.08

 $Y^*=bx+a$ where x is the concentration of Dapagliflozin in $\mu g/ml$ and Y is the absorbance at the respective λmax .

7.1VALIDATIONOFANALYTICALMETHOD:



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Validation of an analytical method is the process to establish by laboratory studies that the performance characteristic of the method meets the requirement for the intended analytical application. Performance characteristic were expressed in terms of analytical parameters.

1. Linearity:

Calibration graph were plotted using absorbance of standard drug versus concentration of standard drug solution. Linear regression data showed a good linear relationship over a

Concentration range 10-60µg/ml.

Table No: 7.2. Calibration data of Dapagliflozin

S. No.	Concentration (µg/ml)	Absorbance
1	0	0
2	10	0.0522
3	20	0.0884
4	30	0.1276



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5	40	0.1703
6	50	0.2122
7	60	0.2673

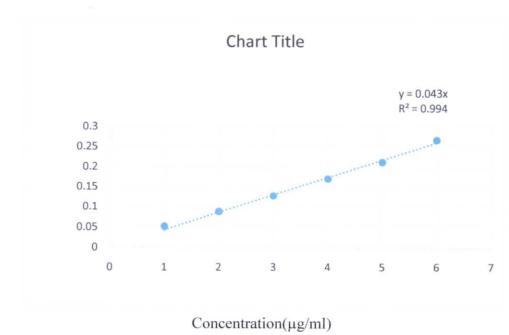


Fig: 7.1 Calibration curve of Dapagliflozin

Observation:



absorbance

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- 1. The correlation coefficient for Dapagliflozin was found to be 0.994 respectively.
- 2. The linearity range for Dapagliflozin was found to be10-60μg/ml.

2. Precision:

Table No: 7.3. Intraday precision

0.03	Conc		Absorbance							0/ DC
0.S.N 0	μg/m l	1	2	3	4	5	6	AVG	SD	%RS D
1	40	0.162 6	0.163 5	0.164 6	0.167 7	0.170	0.169 7	0.16605	0.003523	1.1216
2	40	0.176	0.173	0.178 5	0.179 6	0.174 9	0.175	0.17631 7	0.002353	1.3345
3	40	0.172	0.176	0.175 5	0.172 7	0.173	0.177 8	0.17463	0.002221	1.5180
4	40	0.171	0.176	0.178	0.175	0.173	0.172 5	0.17538	0.002796	0.5942
5	40	0.170 6	0.171	0.172 5	0.173 6	0.174 5	0.175	0.17301 7	0.001911	1.1045



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							6			
6.	40	0.172	0.173	0.174	0.179	0.178	0.175	0.17563	0.00257	1.466

Acceptance criteria:

%RSD of the six replicate injections should not more than 2.0%

Table No: 7.4: Inter day precision result

	Conc			Absor	bance						
S.No	μg/ml	1	2	3	4	5	6	AVG	SD	%RSD	
1	40	0.1789	0.1777	0.1752	0.1766	0.1745	0.1796	0.176417	0.001653	0.9369	
2	40	0.1723	0.1745	0.1756	0.1789	0.1740	0.1755	0.175133	0.002202	1.2573	
3	40	0.1718	0.1711	0.1758	0.1798	0.1765	0.1746	0.175083	0.003212	1.8345	
4	40	0.1743	0.1721	0.1756	0.1725	0.1759	0.1774	0.174167	0.001569	0.9008	
5	40	0.1746	0.1759	0.1785	0.1796	0.1764	0.1752	0.177067	0.001815	1.0250	
6	40	0.1733	0.1725	0.1713	0.1791	0.1752	0.1769	0.174433	0.002749	1.575	





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Acceptance criteria:

%RSD of the six replicate injections should not more than 2.0%

2. Accuracy:

Table No: 7.5.Observation for accuracy standard(50µg/ml)

S. No	Concentration (μg/ml)	Absorbance
1	Set-1	0.2127
2	Set-2	0.2073
3	Set-3	0.2098
4	AVG	0.2099
5	SD	0.002702
6	%RSD	1.28

Table No: 7.6Observation for accuracy for 80% (45μg/ml)



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S. No	Concentration (μg/ml)	Absorbance
1	Set-1	0.1886
2	Set-2	0.1876
3	Set-3	0.1893
4	AVG	0.1885
5	Result	44.90
6	%Rec	101.3
7	SD	0.00085
8	%RSD	0.453

Table No: 7.7. Observation for accuracy standard 100% (55μg/ml)

S. No	Concentration (μg/ml)	Absorbance
1	Set-1	0.2345
2	Set-2	0.234



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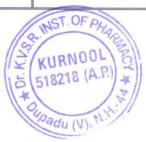
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3	Set-3	0.2335
4	AVG	0.234
5	Result	55.74
6	%Rec	98.49
7	SD	0.0005
8	%RSD	0.0117

Table No: 7.8. Observation for accuracy standard 120% (65µg/ml)

S .No	Concentration (µg/ml)	Absorbance
1	Set-1	0.2563
2	Set-2	0.2691
3	Set-3	0.2881
4	AVG	0.271167
5	Result	64.57
6	%Rec	99.8



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7	SD	0.016
8	%RSD	0.405

Table No: 7.9. For accuracy summary

Sample (%)	Initial amount(µg/ml)	Amount added(μg/ml)	Amount recovered(µg/ml)	%Recovery ±SD*	%RSD
80	40	5	44.90	101.3±0.003	0.453
100	50	5	55.74	98.49±0.000	0.011
120	60	5	64.57	99.8±0.016	0.405

^{*}Average of three determinations

Acceptance criteria:

- 1. %Recovery should be within the range of 98-102%
- 2. %RSD should be less than 2.

4. Sensitivity:

Limit of detection of (LOD) and limit of quantitation (LOQ) were determined from standard and slope method as per ICH guidelines, for Dapagliflozin LOD was found to be $5.63\mu g/ml$ and LOQ was found to be $17.07\mu g/ml$.

Table No: 7.10. Observation of Limit of Detection:



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S.No	Slope	SD of precision	LOD				
1	0.043	0.0025	5.63				

Table No: 7.11. Observation of Limit of quantitation:

S.No	Slope	SD of precision	LOQ
1	0.043	0.0025	17.07

5. Ruggedness:

Table No: 7.12. For Ruggedness(Analyst to Analyst)

	Analys	t-1	Analyst-2						
S. No	Concentration(µg/ml)	Absorbance	Concentration (μg/ml)	Absorbance					
1	40	0.1743	40	0.1711					
2	40	0.1708	40	0.1738					
3	40	0.1722	40	0.1705					
	AVG	0.1724	AVG	0.1718					
	SD	0.001762	SD	0.001758					
	%RSD	1.022	%RSD	1.023					



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Acceptance criteria:

%RSDofthesixreplicateinjectionsshouldnotmorethan2.0%

6. Robustness:

Table No: 7.13. For 230and 234 wavelengths

S. No	Concentration	Absorbance (at 230 nm)	Absorbance (at 234 nm)
1	Set-1	0.1425	0.1609
2	Set-2	0.1469	0.1589
3	Set-3	0.1438	0.1601
	AVG	0.1444	0.1599
	SD	0.002260	0.001007
	%RSD	1.56	0.6297

Table No: 7.14. Robustness Summary

S. No	Condition	Modification	Mean absorbance±SD*	%RSD for absorbance



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1	Wavelength(nm)	230	0.144±0.0022	1.56
•	Wavelength(mn)	234	0.159±0.0010	0.62

^{*}Average of the three determination.

Acceptance criteria:

%RSD should not be more than 2.





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DISCUSSION

In the present work, an attempt was made to provide a newer, sensitive, simple, accurate and low cost UV-Visible Spectroscopic method. It is successfully applied for the determination of Dapagliflozinin pharmaceutical preparations without the interferences of other constituent in the formulations.

The optimum wavelength for detection was 232nm at which better detector response for the drug were obtained. The calibration was linear in concentration range of $10\text{-}60\mu\text{g/ml}$ in the Table 7.2 for Dapagliflozin respectively. The sensitivity for the drug has been calculated and the LOD and LOQ of the Dapagliflozin was found to be $0.56\,\mu\text{g/ml}$ and $1.7\,\mu\text{g/ml}$ in the Table 7.10 & Table 7.11.

The low values of % R.S.D. indicate the method is precise and accurate. The mean recoveries were found in the range of 98-102% in the Table 7.9 for Dapagliflozin respectively.

Ruggedness of the proposed methods was determined by analysis of aliquots from homogeneous slot by different analysts, using similar operational and environmental conditions; the % R.S.D. reported was found to be less than 2 % in the Table 7.12.

The proposed method was validated in accordance with ICH parameters and the results of all methods were very close to each other as well as to the label value of commercial pharmaceutical formulation. Therefore, there is no significant difference in the results achieved by the proposed method.



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Hence it is suggested that the proposed is UV / VIS Spectrophotometric method can be effectively applied for the routine analysis of Dapagliflozinin bulk.

8. CONCLUSION

For routine analytical purpose it is always necessary to establish method capable of analysing huge number of samples in a short time period with due accuracy and precision.

Dapagliflozin is not official in Pharmacopoeia. There is few analytical methods appeared in the literature for the determination of the Dapagliflozin. In literature review we have method only for the estimation of the above drugs of concern in individually or in combination of others. In view of the above, a simple and specific analytical method was planned to develop with sensitivity, accuracy, precision and economical.

In the present investigation of UV spectrophotometric method for the quantitative estimation of Dapagliflozin in pure drug has been developed and validated.

The proposed UV method is more sensitive, accurate and precise and is suggested for routine analysis.





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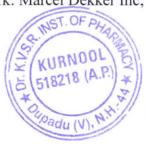
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CASE STUDY LEARNING

Case studies are a written description of a real-life problem or situation. Only the facts are provided, usually in chronologic sequence similar to what would be encountered in a patient care setting. The use of cases actively involves the students in the analysis of facts and details of the case in the traditional format called SOAP analysis, by selection of a solution to the problem and defense of his or her solution through discussion of the case details. In the case based learning students use their recall of previously learned information to solve clinical case. The case method is used primarily to develop the skills of self-learning, critical thinking, problem identification, and decision making. Working on subsequent cases with similar problems reinforces information recall. Case studies in the health sciences provides the personal history of an individual patient and information about 1 or more health problems that must be solved. The students work through the facts of the case, analyze the available data, gather more information, develop hypotheses, consider possible solutions, arrive at the optimal solution and consider the consequences of the learner's decisions. The use of the case studies and other active learning strategies will enhance the development of essential skills necessary to practice pharmacy in any setting, including community, ambulatory care, primary care, health systems. Long term care. Home health care, managed care and the pharmaceutical industry.



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PHARM.D ROASTERS YEAR WISE(21-22)

PHARM.D-II YEAR ROASTERS

ROLL NO NAME OF STUDENT	JAN-FEB	MARCH-APRIL		JULY	AUGUST	SEPTEMBER	OCTOBER
SERITUSS2 DADIPINENI LEELAVATHI	(2022)	(2022)	(2022)	(2022)	(2022)	(2022)	(2022)
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DERITORA DEPATI MAHA AISHWARYA	PAEDIATRICS	MM-7	MM-2	PATIENT	MM-I	PAEDIATRICS	MM-3
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DERITOROT HEBBARE SRAVANI BAL							
DERITOOOS JINKA LAVANYA			MM-7				
DERITHORS KALLUR SALSIREESHA	MM-3	PAEDIATRICS		MM-2	PATIENT	MM-1	PAEDIATRICS
DERITORIO KAMALAPURAM SUPRAJA BAI		(CHMW-1,2)		3131-2	COUNSELLING		(CHMW-3,4)
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0ERIT0017 SUVARNA SRAVANI							
0ERIT0018 VAVILALA ARCHANA				PAEDIATRICS			PATIENT
GERITOGIO VISANAKARRA TRIVENI	MM-I	(CHMW-3,4)	MM-3	(CHMW-1,2)	MM-7	MM-2	COUNSELLING
0ER1T0020 BOGYAM MANASA				(-11,111-1,12)			
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0ER1T0025 SHAIK IRSHAD NUFSHRUN	COCHSECULIO						
0ER1T0026 VANGALA SAMYUKTHA							
DERITO027 MOLLA KIFAYATHULLA							
DERITO028 K.S.PRADHIYUMNA						PAEDIATRICS	
DERITO029 SYED ABRAR AHMED	MM-2	PATIENT	MM-1	PAEDIATRICS	MM-0	(CHMW-1,2)	3171-1
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20ER1T0003	D.GAMANA SREE	
20ER1T0004	D.MAHA AISHWARYA	
20ER1T0005	G.SUCHARITHA	



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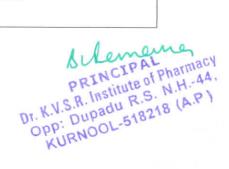
20ER1T0006	G.KAVYA	
BATCH-II(JAN -FEE	3)	

ROLL NUMBER	STUDENT NAME	WARD
20ER1T0007	H.SRAVANIBAI	MM3
20ER1T0008	J.LAVANYA	
20ER1T0009	K.SAI SIRISHA	
20ER1T0010	K.SUPRAJA BAI	
20ER1T0011	K.HIMA BIMDU	
ROLL NUMBER	STUDENT NAME	WARD
20ER1T0012	M.BHAVYA	PEDIATRIC(CHMW-3,4)
20ER1T0013	M.HEMALATHA	
20ER1T0014	M.SUMALATHA	
20ER1T0015	M.CHITRA	
20ER1T0016	M.AMULYA	

ROLL NUMBER	STUDENT NAME	WARD	
20ER1T0017	SUVARNA SRAVANI	MM-I	
20ER1T0018	V.ARCHANA		
20ER1T0019	V.TRIVENI		
20ER1T0020	B.MANASA		
20ER1T0021	E.SUMANJALI		

ROLL NUMBER STUDENT NAME		WARD
20ER1T0022	E.CHARITHA	PATIENT COUNSELLING
20ER1T0023	G.NAGA SUREKHA	
20ER1T0024	G.KIRTHI	
20ER1T0025	S.IRSHAD NUFSHRUN	
20ER1T0026	V.SAMYUKTHA	

ROLL NUMBER	STUDENT NAME	WARD	
20ER1T0027	M.KIFAYATHULLA	MM-2	
20ER1T0028	K.S.PRADHYUMNA		



Date :



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20ER1T0029	S.ABRAR AHMED	
20ER1T0030	S.NASIR AHMED	
20ER1T0031	V.VIJAYA TEJA	
20ER1T0032	S.KALID	

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KURNOOL-518278 (A.P.)



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Principal / Correspondent

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PHARM.D III YEAR ROASTER(2021-2022)

	NAMES OF STUDENTS	Aug-Sept (2021)	Oct-Nov	Dec-Jan(2022)	Feb-March	April-May	June	
SER170002	Alefiya kotawala							
9ER1T0003	A.Keerthi							
19ER3T0004	B.Sowjanya	8AM-7	DVL	PATIENT COUNSELING	MM-3	MM-1	MM-2	
19ER1T0005	B.Aswini Bai							
19ER1T0006	C.Sushma							
19ER1T0009	G Mounika							
19ER1T0010	G .Naga vinugna	MM-2	MM-7	The same of the sa	PATIENT COUNSELING	MM-3	MM-1	
19ER1T0012	K. Meghana	MW-2	MM-7	DVL		NAMO-3	MM9-3	
19ER1T0013	P. Naga Sruthi							
19ER1T0014	Shaik Rizwana				DVL			
19ER3T0016	B. Mounika	MM-2	MM-2	MM-7		PATIENT COUNSELING	MM-3	
19ER1T0024	P. Sudha	Dene-2	MM-Z	MINE-7	DVL	PATERI COMISELINO	min. s	
19ER1T0025	S.Navya Stree							
19ER1T0026	T. Khasid							
195R1TD028	Valmiki Venkateshwari						The second second	
19ER1T0029	Vangala Naga divya	MM-3	MM-1	MM-2	MM-7	DVL	PATIENT COUNSELING	
19ER1T0007	C. Mehaboob Desai							
19ER1T0001	A. Dilip Kumar							
19ER1T0011	K.Sai Chaitanya							
19ER1T0015	V. Vamsi Krishna							
19ER1T0015	A. Sai Kumar Reddy	PATIENT COUNSELING	MM-3	MM-1	MM-2	MM-7:	DAT	
19ER1T0017	8. Venkatesh							
19ER1T0019	K. Veera Reddy							
19ER1T0020	K. Satyanarayana charan						MM-7	
19ER1T0021	M. Hansmontha	DVL	PATIENT COUNSELING	G MM-3	MM-1	MM-2		
19ER1T0023	P. Siva Gangadhar		The second second					
19ER1T0027	T. Shanu Prakash							
KURNOÖL.	PRINCIPAL PRINCIPAL 5.R. Institute of Pharmacy 5.D. Dupadu R.S. N.H-7. JERNOOL-518216 (A.P.)	IACV				SUPERINTENDENT GOVERNAMENT GENEROOL	ENERAL HOSPITAL	





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HOSPITAL DUTIES (AUG-SEP) (AY: 21-22)

ROLL NUMBER	STUDENT NAME	WARD	
19ER1T0002	K.ALEFIYA	MM-7	
19ER1T0003	A.KEERTHI		
19ER1T0004	B.SOWJANYA		
19ER1T0005	B.ASHWINI BAI		
19ER1T0006	C.SUSHMA		

ROLL NUMBER	STUDENT NAME	WARD
LOED LTOOOD	G.MOUNIKA	MM-2
19ER1T0009	G.NAGA VINUGNA	
19ER1T0010	G.NAGA VINOGNA	
19ER1T0012	K.MEGHANA	
19ER1T0013	P.NAGA SRUTHI	

ROLL NUMBER	STUDENT NAME	WARD	
19ER1T0014	SHAIK RIZWANA	MM-I	
19ER1T0018	B.MOUNIKA		
19ER1T0024	P.SUDHA KALYANI		
19ER1T0025	S.NAVYA SREE		

Schemener PRINCIPAL Dr. K.V.S.R. Institute of Pharmacy Opp: Dupadu R.S. N.H.-44 KURNOOL-518218 (A.P.)

Date :



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	STUDENT NAME	WARD
ROLL NUMBER		
	T.KHASID	MM-3
19ER1T0026		
	V.VENKATESWARI	
19ER1T0026		
	V.NAGA DIVYA	
19ER1T0026		
19EK110026	CAMPILL BOOK BEGAL	
	C.MEHABOOB DESAI	
19ER1T0026		
	A.DILIP	
19ER1T0001		

ROLL NUMBER	STUDENT NAME	WARD	
19ER1T0011	K.SAI CHAITANYA	PATIENT COUNSELLING	
19ER1T0015	V.VAMSI KRISHNA		
19ER1T0016	A.SAI KUMAR REDDY		
19ER1T0017	B.VENKATESH		
19ER1T0019	K.VEERA REDDY		

ROLL NUMBER	STUDENT NAME	WARD	
19ER1T0020	K.V.S.CHARAN	DVL	
19ER1T0021	M.HANUMANTH		
19ER1T0023	P.SIVA GANGHADHAR		
19ER1T0027	T.BHANU PRAKASH		

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Principal / Correspondent

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PHARM.D IV YEAR(2021-2022)

	NAMES OF STUDENTS	PHARM D IV YEA Aug-Sept (2021)	Oct-Nov	Dec-Jan(2022)	Feb-March	April-May	June	
ROLL NUMBERS	BERAMADEVI	wng-zehr (sner)	Oct 1101					
ISER1T0001						The same of the sa		
1SER1T0002	G.SAI VILASIIINI	MM-1	MM-7	MM-3 MM-4	PAEDIATRICS	PSYCHIATRIC	MM-2	
18ER1T0003	ELIKHITA							
18ERIT0004	LDIVYASREE							
18ER1T0005	JHEMALATHA							
18ER1T0007	M.MANIPRASANNA							
18ER1T0008	M.RADHIKA	MM-2	MM-1	MM-7	MM-3 MM-4	PAEDIATRICS	PSYCHIATRI	
18ER1T0010	P.ASMA	- Maria	34.31-1					
18ER1T0011	S.TAHURA BATUL							
18ER1T0013	S.ZUWERIYA SULTANA D.DURGA PAVITRA	-						
18ER1T0014	S.VANI					100		
18ER1T0015	TJAHNAVI	PSYCHIATRIC	MM-2	MM-1	MM-7	MM-3 MM-4	PAEDIATRIC	
18ERITU016	UJOSHITHA	131011011111	10000					
18ER1T0017	V.TEJASWINI							
1SERITO01S	V.MALLESHWARI							
18ER1T0019	J.ZUHA KHANAM							
18ER1T0021	KKEERTHANA	PAEDIATRICS	PSYCHIATRIC	PSYCHIATRIC	MM-1	MM-7	MM-3 MM-	
18ER1T0023	K.DIVYA SIRISHA	-			MM-2	MM-I	MM-7	
18ER1T0025	S.ASMA PARVEEN							
18ER1T0026	E.ROHINI VADANA							
18ER1T0029	S.SANA AFREEN							
ISERITO030	S SANA AFREEN	MM-3 MM-4	PAEDIATRICS					
18ER1T0011	S.BABA ARSHID							
18ER1T0006	K SHANMUKA GANESH							
18ER1T0009	M.SUJAN DORA							
18ER1T0020	LSAIKIRAN							
18ER1T0024	M PRANEETH PREMPAUL	MM-7	MM-3 MM-4	PAEDIATRICS	PSYCHIATRIC	MM-2	- MM-1	
18ER119927	P.ANWAR BASHA	3131-7	and the second s		The second second			
18ER1T0028	RAVIJAY VIKAS							
17ER1T0026	S.NEHAL				-			
KURNOOL	PRINCIPAL PRINCIPAL OR KYSR Institute of Phatia Opp Dupadu RS NH-7 Opp Dupadu RS NH-7	MACY				INTENDENT RIMENT GENERAL	HOSPITAL.	

HOSPITAL DUTIES (AUG-SEP) (AY 21-22)

REG.NO	NAME OF THE STUDENT	WARD	
18ER1T0001	B.RAMA DEVI	MM-1	
18ER1T0002	G.SAI VILASINI		
18ER1T0003	E.LIKITHA		
18ER1T0004	J.DIVYA SREE		
18ER1T0005	J.HEMA LATHA		

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KURNOOL-518218 (A.P.)

Date :



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Princi	pal	Corres	pond	lent
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REG.NO	NAME OF THE STUDENT	WARD	
18ER1T0007	M.RADHIKA	MM-2	
	P.ASMA		
18ER1T0008			
	S.TAHURA BATUL		
18ER1T0010			
18FR1T0013	S.ZUWERIYA SULTHANA		

REG.NO	NAME OF THE STUDENT	WARD	
18ER1T0014	D.DURGA PAVITHA	PSYCHIATRY	
18ER1T0015	S.VANI		
18ER1T0016	T.JAHNAVI		
18ER1T0017	U.JOSHITHA		
18ER1T0018	V.TEJASWINI		

REG.NO	NAME OF THE STUDENT	WARD	
18ER1T0019	V.MALLESWARI	PEDIATRICS	
	J.ZUHA KHANUM		
18ER1T0020	K.KEERTHANA		
18ER1T0022	K. DIVYA SIRISHA		
18ER1T0023	S.ASMA PARVEEN		
18ER1T0025			

REG.NO	NAME OF THE STUDENT	WARD	
18ER1T0026	E.ROHINI VANDANA	MM-3,MM-4	
18ER1T0029	S.SANA AFREEN		
18ER1T0030	S.SANAAFREEN		
18ER1T0011	S.BASHA ARSHID		





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Principal / Correspondent		Date :	
18ER1T0006	K.SHANMUKH GANESH		
REG.NO	NAME OF THE STUDENT	WARD	
18ER1T0009	M.SUJAN DORA	MM-7	
18ER1T0020	I.SAI KIRAN		
18ER1T0024	M.PRANEETH PREM PAUL		
18ER1T0027	P.ANWAR BASHA		
18ER1T0028	R.VIJAY KUMAR		
18ER1T0026	S.NEHAL		

PRINCIPAL

Dr. K.V.S.R. Institute of Pharmacy

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KURNOOL-518218 (A.P.)



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Principal / Correspondent

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PHARM.D V YEAR(2021-2022)

TERTITION D. CHETHANA	n April Psychiatry	MM-7
TERITO002 D-SANDYA	Psychiatry	MAA-7
TERTITION D. CHETHANA	Psychiatry	MM-7
TERTITION		
17ERIT10007 K.BHAVANI 17ERIT10008 M.PRIYANKA MMA-7 General surgery Pediatrics MM-1 MM		
TERITORS M. PRIYANKA MAA-7 General surgery Pediatrics MAA-1 MAA		
17ERIT0009 M. MAMATHA	2 001	Psychiatry
		3674
ITERITIONS N. VIJAYA		
ITERITORI S. NAVYASREE	_	
17ERIT9012 SRUKSAR SAMREEN		
17ERITORIS S. SURASINI Psychiatry MM.7 General surgery Pediatrics	MM-1 MM-7	DVI
17ERIT0015 S. SHAFIYA TABASSUM		
17ERIT0016 SHAIK WASIMA NASKEEN		
ITERITORI7 AMBATI PAVITRA		
L'ERITSOIS G. LEORON REBEKHA		
17ERIT0019 J PAVITRA DVI. Psychiatry MM-7 General turget	ry Pediatrics	MM-1 MM-7
17ERIT0021 M. VAISHNAVI		
17ERIT0022 NSHINV SUSAN		
ITERIT0023 P. DIVYA		
17ERTT0025 RUMSHA RUKSAR		
17ERIT0827 SHAIKTHASLEEM MM-1 MM-2 DVL Psychiatry MM-7	General surgery	Pediatrics
17ERIT0028 T. YAMMINI		
ITERIT0029 V.SAMYUKTIIA		
17ERIT0014 S. VISWANATH REDDY		
17ERIT0020 K.LAL REDOY		
17ERT19024 R. NEERAJ Fedicitics MM-1 MM-2 DVL Psychology	y MW-3	General surgery
17ERIT0030 Y KARTHIK KUMAR		
16Y01T0023 SHOAIB ABDUR RAQUEER		

HOSPITAL DUTIES (AUG-SEP) (AY 21-22)

REG.NO	NAME OF THE STUDENT	WARD
17ER1T0001	B.SOWJANYA	GENERAL SURGERY
17ER1T0002	D.SANDYA	
17ER1T0003	D.CHETHANA	
17ER1T0004	D.SAILAJA	
17ER1T0005	K.SUPRAJA	

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Date :



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Princip	oal / Corr	espondent
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REG.NO	NAME OF THE STUDENT	WARD	
17ER1T0006	K.LAKSHMI CHARITHA	MM-7	
17ER1T0007	K.BHAVANI		
17ER1T0008	M.PRIYANKA		
17ER1T0009	M.MAMATHA		
17ER1T0010	N.VIJAYA		

REG.NO	NAME OF THE STUDENT	WARD	
17ER1T0011	N.NAVYA SREE	PSYCHIATRY	
17ER1T0012	S.RUKSAR SAMREEN		
17ER1T0013	S.SUHASINI		
17ER1T0015	S.SHAFIYA THABASSUM		
17ER1T0016	A.WASIMA NASREEN		

REG.NO	NAME OF THE STUDENT	WARD	
17ER1T0017	A.PAVITHARA	DVL	
17ER1T0018	G.LEOHON REBEKHA		
17ER1T0019	J.PAVITHRA		
17ER1T0021	M.VAISHNAVI		
17ER1T0022	N.SHINY SUSAN		





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PHARM.D VI YEAR(2021-2022)

T0007 T0008 T0010	G NAVEEN BABU ASAD AU KHAN M RAMANJANEYULU	PSYCHIATRIC					
T0010	MRAMANIANEYULU	DEVENDATION					
10011		PENCHUATRIC					
			PEADIATRICS	GENERAL SURGERY	MM-7	5555-6	FM5-3
	M.RAM SUNIL	Parconarios					
T0017	D.YAKUB BASHA						
T0003	P.IAVEED						
T0006	5 MANIKANTA						
10020	3 LOKESH ROYAL	FM-2 FM-8	PSYCHIATRIC	PEADIATRICS	GENERAL SURGERY	N5N1-2	MM-2
110021	5 SUBBA REDDY						
170004	C.KEERTH!						
170013	ALSAI DEEPIKA						
110029	LSWETHA MADHURIMA	MAN 8	FM-7 FM-8	PSYCHIATRIC	PEADIATRICS	GENERAL SURGERY	MM-7
170015	D. SHARON BOSE						
170016							
10026			MM II	150-7 550-8	PSYCHIATRIC	PEADIATRICS	
1T0027		MM-7					GENERAL SURGERY
	S SHABANA TASAFEN						
170014	5 SUMERA IRHAM						
170030	D. JAYAROOPA VANI						
170018	B.BHAVANA			MM-E	FM-7 FM-8	PSYCHIATRIC	
170019	SRI KRISHNA JYOTIKA						
170021	M.PRIYANIALI	GENERAL SURGERY	5454-7				PEADIATHICS
170024	LVAISHNAVI						
110025	V.HARI SUSHUMA						
170001	B AKHILA						
170002	M.AKSHITHA					FM-7 FM-8	
170005	LASYA	PEADIATRICS	GENERAL SURGERY	MNI-7	MM-8		PSYCHIATRIC
		PERMITTING					
170009	P.PREMA LATHA						
	170006 170020 170020 170024 170013 170015 170016 170027 170027 170028 170018 170018 170018 170021 170021 170021 170021 170021	170006 S. MANKANTA 170020 S. LORES H. ROYAL 170023 S. SUBBA REDDY 170024 S. SUBBA REDDY 170013 S. SUBBA REDDY 170013 AS AS I OCERNICA 170013 C. SWATTHA MADHURIMA 170016 M. V. NAGA VASAVI LATHA 170026 L. (*Frie" N. MANASA 170027 R. MANASA 170027 R. MANASA 170028 S. SHARRAN ATASAFEN 170018 B. BHAVANA 170018 B. BHAVANA 170019 S. SUBBRA SHAN YOTHA 170021 M. PRIVANSAIS 170021 M. PRIVANSAIS 170025 V. MARI SUSHUMA 170025 V. MARI SUSHUMA 170005 B. ARHINAN V. TIPOOTS V. MARI SUSHUMA 170005 V. MARI SUSHUMA 1700010 B. ARHINAN V. TIPOOTS V. MARI SUSHUMA T. V. MARI SUSHUMA T. V. MARI SUSHUMA V. M	170006 S. MANIKANTA TODO	170006 S. MANIKANTA FM. 2 FM. 8 PSYCHIATRIC T00213 S. SUBBA REDOY T10024 S. SUBBA REDOY T10024 S. SUBBA REDOY T10025 S. SUBBA REDOY T10026 S. SUBBA REDOY T10026 S. SUBBA REDOY T10026 S. SUBBA REDOY S. SUBBA	170006 S. MANIKANTA FM. 2 FM. 8 PSYCHIATRIC PEADIATRICS 170024 S. SURBA REDOY TOO23 S. SURBA REDOY TOO24 S. SURBA REDOY TOO25 S. SURBA REDOY TOO25 S. SURBA REDOY TOO25 S. SURBA REDOY TOO26 S. SURBA REDOY TOO26 S. SURBA REDOY TOO26 S. SURBANA TASAETN MM. 2 FM. 7 FM. 8 FM. 7 FM. 8 TOO26 S. SURBANA TASAETN MM. 7 MM. 8 FM. 7 FM. 8 TOO27 S. SURBANA TASAETN MM. 7 MM. 8 FM. 7 FM. 8 TOO28 S. SURBANA TASAETN MM. 7 MM. 8 FM. 7 FM. 8 TOO29 S. SURBANANA TOO21 S. SURBANANA TOO21 S. SURBANANA TOO21 S. SURBANANA TOO22 M. PRIVANIANI TOO25 S. SURBANANA TOO25 M. PRIVANIANI TOO25 V. MAR SUSHUMA TOO25 V. MAR SUSHUMA TOO25 V. MAR SUSHUMA S. SURBANANA S. SUSHUMA S. SURBANANA S. SURBAN	170006 S. MANIKANTA FM. 2 FM. 8 PSYCHIATRIC PEADIATRICS GENERAL SURGERY 170024 S. SUBBA REDDY 170024 S. SUBBA REDDY 170024 S. SUBBA REDDY 170025 S. SUBBA REDDY 170025 S. SUBBA REDDY 170025 S. SUBBA REDDY 170026 S. SUBBA REDDY 170026 S. SUBBA REDDY 170027 R. ANANASA 170028 S. SUBBANASA 170028 S. SUBBANASA 170029 S. SUBBANASA SUSHIMAN 170029 S. SUBBANASA SUSHIMA 170029 S. SUBBANASA SUSHIMAN S. SUBBANASA SUSHIMAN S. SUBBANASA SUSHIMA S. SUBBANASA SUSHIMA S. SUBBANASA S. SUBBANASA	100006 S. MANIKANTA FM-2 FM-8

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Opp: Dupadu R.S. N.H.-44,
Opp: Dupadu R.S. (A.P.)
KURNOOL-518218 (A.P.)

Date :



16ER1T0016

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Principal / Correspondent

REG NUM	NAME OF THE STUDENT	WARD	
16ER1T0004	C.KEERTHI	MM-8	
16ER1T0013	M.SAI DEEPIKA		
16ER1T0029	L.SWETHA MADHURIMA		
16ER1T0015	D.SHARONE ROSE		
	M.V.NAGA VASAVI LATHA		

REG NUM	NAME OF THE STUDENT	WARD
16ER1T0026	L.GRACE	MM-2
16ER1T0027	K.MANASA	
16ER1T0028	S.SHBHANA	
16ER1T0014	S.SUMERA	
16ER1T0030	D.JAYA RUPA VANI	

REG NUM	NAME OF THE STUDENT	WARD
16ER1T0018	B.BHAVANA	GENERAL SURGERY
16ER1T0019	SRI KRISHNA JYOTHIKA	
16ER1T0021	M.PRIYANJALI	
16ER1T0024	L.VAISHNAVI	
16ER1T0025	V.HARI SUSHMA	

	NAME OF THE STUDENT	WARD
REG NUM		



Date :



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1.CED 1.TO001	B.AKHILA	PEDIATRICS	
16ER1T0001			
	M.AKSHITHA		
16ER1T0002			
	LASYA		
16ER1T0005			
	P.PREMA LATHA		
16ER1T0009			
	SADIYA SAMREEN		
16ER1T0012			

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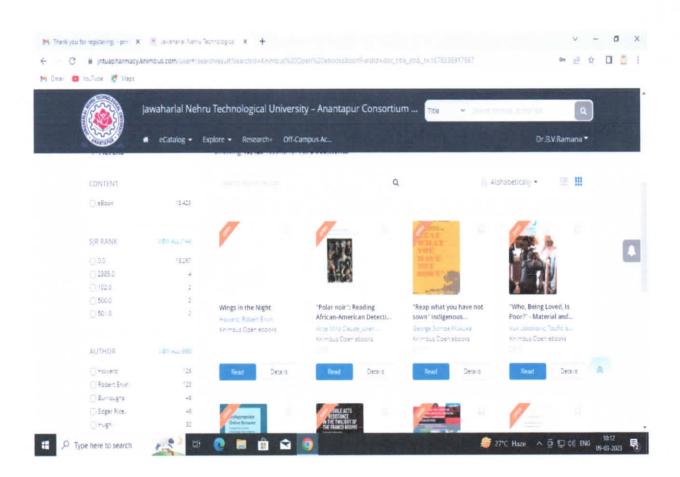
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OPP: Dupadu R.S. N.H. A.A.

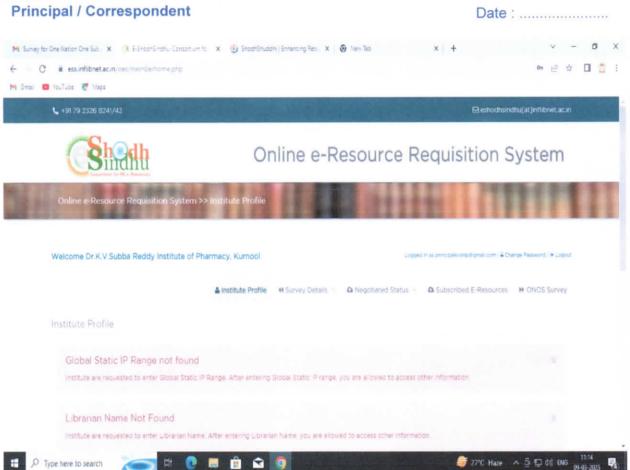
OPP: Dupadu S18218 (A.P.)

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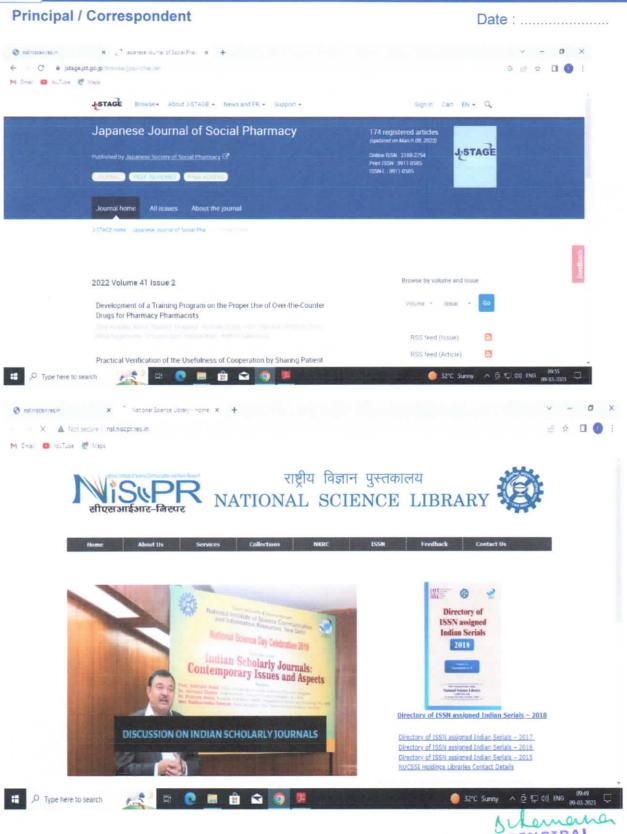
Dr. N. P. Dungdu S. S. 18 (A.P.)

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Date :



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■ GPAT 2021 - Previous Year Ques X + ← → C # youtube.com/watch?v=WQD_HRCDECc 단 년 ☆ 뭐 🛛 🧯 : = NouTube gpat 2020 Q You Top chat replay v Siddharth Shankar Nag start fast sir Screat Superstar 1 we will earn after we win? Himanshu Gupta Good evening sir **PREVIOUS YEAR** SRIRAM YADAV M 19P0P026 good evening sir Ram h good evening sir QUESTION Drx sing vlog hi vikrant chaudhary ram ram sir ji 25 MCQs V Bhumika Patili good evening sir Explanation Dramma vlog gd evening 3 Nikhilesh Maruthi 74 Egarly waiting sir Ishan Upadhyay 🕠 🗘 🐧 🥻 🥻 000000022 Abhisek Triparthi Krishna Patil good evening Sir... WouHub good evening GPAT 2021- Previous Year Question Paper Discussion with Explanation | Part-I (A Kartik Ramteks | Odlevening siz GPAT DISCUSSION CENTER 6 (\$) Thanks ... 15 1.9K (7) A Share ● 34°C Sunny ^ 40 40 ENG 11:52 30-63-2023 P Type here to search

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Number of teachers and students using the library per day over last year

Last page of accession register	Method of computing Per day usage of library	No. of users using library through e- access per day	No. of teachers accessing library per day	No. of students accessing library per day	Certified E- copy of the ledger of the data for 5days	Certified screenshots of the data for 5 days for online access
-	Gate register	195	41	355	_	-

Percentage per day usage of library by teachers and students:

Number of students=650

Number of teachers=51

Formula= (Number of teachers and students using the library perday/Total number of teachers and students) X 100

 $= (591/650) \times 100$

=90.92





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Data												
Date												

WORKSHOP & GUEST LECTURES:

	AY @ 2021-22	
1.	One day Webinar on Quality Standards of AYUSH recommended drugs for Covid-19	Dr. M. Kalyan Kumar Raju
2.	One day conference on Research in Ayurvedic Product and Quality assurance and quality control in Phytopharmaceuticals.	Dr. Md. Jamal Basha
3.	One day conference on the Role of Artificial Intelligence During Covid-19	Dr. K. Srikanth (NIPER-SAS)
4.	Conference on "Role of stem cells in Pancreatic regeneration and Reversal of Diabetes"	Dr. K. P. Shameem (USA)
5.	Webinar on "Academic ethics and Plagiarism"	Dr. M. Gulshan Kumar
6.	One day conference on Recent Development and Future challenges of Nuclear Science in Health Sector	Dr. S. Banesh (IIT)
7.	One day webinar on Regulatory Affairs and international marketing	Mr. Rahul Singh Gurzar
8.	Workshop on Advances in Biotechnology at BioEra	Mr. K. Kamabagiri Swamy (BIOCON)
9.	A webinar on immunology and immune technology	Mr. S. Siva Kumar (NOVARTIS)
10.	One day conference on biodiversity, climate change &suitable agriculture towards food security	Mr. K. Narender Goel
11.	One day conference on strategies for global competitions &economic growth.	Mr. T. Prashant Kumar
12.	One day Webinar on synthetic and pharmaceutical chemistry	Dr. Md Shameer (Georgia State University)
13.	One day conference on Recent Advances in Health Sciences	Dr. K. Pavan Kumar Raju
14.	One day Conference on Amalgamation of recent Pharmaceutical Developments in ayurveda	Dr. I. Rupesh (Ayurveda- Tirupati)
15.	Conference on the role of pharmacists in the prevention and management of sleep disorders	Dr. Y. Ganesh
16.	One day conference on Harmonization and Global Efforts for Safe and Effective Generic Drugs	Mr. E. Kiran Goud
17.	Workshop on pharmaceutical quality assurance and quality control and quality management	Dr. C. Mallikarjun Rao
18.	One day conference on stress and stress management	Dr. K. Amarnath





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PHARMACY SEMINARS:

		DR.K.V.SUBBREDDY INSTI	TUTE OF PHARMACY, DUPADU, KURNOOL		
S.No.	Holl No.	Name of the Student	en by B.Pharmary Students in Academic Year 2019-20		
1	1912(19.000)	II SUPRIVA BAL	Title	Presentation Date	
200	THEN INDOLL	D SREYA REDDY	CHEAC DISEASE CHED AND ADDLESS ENT PSYCHIATRY	9/2/2019	
3 1 1	19ER180036	GHARBHA	CHRONIC KIDNEY DISEASE	9/2/2019	
4	191 K180025	K NAVANEL IIIA	CHRONIC LETKEMIA	9/2/2019	
10.3000	191R1R0076	PARJUNIKA	COPD	9/2/2019	
6	THERTROOMS	C KISHORE BABU	COLORECTAL CANCER	92/2019	
10-11	-19ER1R0093	KUSCHA MANASA	CORNEA AND EXTERNAL DISEASE	9-2/2019	
×	TRERIBOOMS	GOLLA TARUN KUMAR	CORONAVIRUS (COVID-19)	1/30/2020	
-	18ER1R0017	GOTTA YELLASWAMY	DENTAL AND ORAL HEALTH	1/30/2020	
18	INFRIENCIS.	GUDIMARALLA KAVIN KUMAR	DEPRESSION	1/3W202U	
	TREBURDOCA	KURUVA SURESH	DIABITIC MICROVASCIT AR COMPLICATIONS	1/30/2020	
152	INFRIRONS.	NARRALIKHITA	DIALYSIS	1/30/2020	
(3)	181 R 1R 00 15	PALLAPU SANDYA RANI	ELECTRONIC HEALTH RECORDS	1/30/2020	
14	188'S I BOO37	PICHIGUNILA MAHUSH	EMERGING AND RESMERGING INVECTIOUS DISEASES	1/30/2020	
15	17ER1R0002	A SUNTINA	END-STAGE RENAL DISEASE	6/13/2018	
Te	(TERTROGOV	BOYA VIDYAVANI	GALLBLADDER AND BILLIARY DISEASE	6/15/2018	
17	17ER (ROUG)	DANDU MADHUSUBHAN	GENOMIC MEDICINE	n 13/2018	
	LTERIROGIA	DASHAVATHPRAVALLIKA	GERD	6/11/2018	
18			ERECTILE DYSFUNCTION	6/13/2018	
19	17ER1RI022	GOLLA MADRO SHEKAR	ETHICS	6.13/2018	
20	178.9(1900)62	BOMMARA BHARGAVA REDDY	HEAD AND NECK CANCER	6/13/2018	
21	17FR1R0067	GITTAIAH HEMALATHA	HEADACHI	8/23/2017	
22	telRikouts -	MALLEPOGU MAHESH	HEALTH DIVERSITY	8/23/2017	
23	16ERTROOTS	PAGIDIMARY	LUNG CANCER	8/23/2017	
34	16ER1R0025	BUKKEM PRATHYUSHA	LYME DISPASE	8/23/2017	
25	IGERTRO026	BREEMAGUNTLA PRAVEEN RAJ	OBIGYN AND WOMEN'S HEALTH NURSING	W/23/2017	
26	161R1R0034	DUDEKULA SHAIKAHAMED	OBESITY AND WEIGHT MANAGEMENT	8/23/2017	
25	16ER:R0036	SHAIK SHAKEER	PATIENT SAFETY	8/23/2017	
28	165R1R0979	YELLALA SRUTEJA			
			given by Phaem.D Students in Academic Year 2019-20	and the second second	
		Name of the Student	Title	Presentation Date	
5.76	Roll No.	B ASWINI BAL	INSULIN THERAPY	2/7/2020	
1	TVERTTOROS	C SUSHMA	INTEGRATIVE MEDICINE	2/7/2020	
2	19ER110000		INTERVENTIONAL CARDIOLOGY & SURGERY	2/7/2020	
- 3	191.R110009	G MOUNIKA	RENAL CELL CARCINOAIA	2/7/2020	
4	PPERITION1	K SALCHAITHANYA	REPRODUCTIVE ENDOCRINGLOGY	2/7/2020	0
5	19ER1 T0015	V. VAMSI KRISHNA	RESIDENTS	3/7/2026	1
6	1963(1700)7	BIVENEATISH	RESTLESS LEGS SYNDROME	7/7/2020	CONTREIP'S
7	19ER110018	GANTA EDIGA LIKHTIDIA	RHEUMATORD ARTHRITIS	4.11-2019	Daganu B.S

The second second	DATE OF TAXABLE	JAKKA IRVYA SRIJ	SCHIZOPHULNIA I	4/1/2019
10	SHEEL STANFT	MI NAGALA MANI PRANANNA	PEDIATRIC DERMATOLOGY	4/11/2019
100	286 82 1000-9	MICRIAGRANI SUJANA DORA	HODGKIN'S LYMPHONG	4/17/2019
THE REAL PROPERTY.	25EW1T(803	SRAIK FAHLSKA BATES	PEDIATRIC OPPUTIAL MOLOGY	4/18/2019
- 1/13kmm	\$ 84.00 1 TOO 2 7	UNUSERE ROSMITTA	URINARY INCONTINUNCE & OAR	40 1/2019
114	F. RELECTIONS	KANTATHAMKEERIHANA	MECHANICAL VENTILATION	4/11/2010
S. Barrier	1753,119661	DARA SHAILAIA	MEDICAL LIBESTYLE	6/24/2019
100	3.75302 T00006	V V AMST KRISHNA	MEDICAL MALFRACTICE AND LEGAL ISSUES	6-24-2019
3.40	ETERTIMOT	KATIKE BHAVANI	MEDICAL PRACTICE MANAGEMENT	6/24/2019
100000000000000000000000000000000000000	TERRETON T	AMBATIPAVITRA	SULP DISORDERS	6-34-2019
11114	124361744010	JPANITHRA	SOCIAL DETERMINANTS OF BEALTH	0/24/2019
211	PERTIONS.	PULLDINYA	SPINAL DISORDERS	6/24/2019
100000000000000000000000000000000000000	FFER I TONOS	RUMBRARUKSAR	PEPTIC LL CER DISEASE	621/2019
33	INCRETIONS:	PATHAN JAVEED	PERIOPERATIVE NURSENG	11/2/2017
25 17	16EE/10004	CHENCHI REDIDVGARI KEERTHI	PSEUSIONIA	11/2/2017
A CONTRACTOR	160 RATIONS.	ASAD ALIKHAN	MODANE	162/2017
THE REAL PROPERTY.	PREMITTRODA	PERUSIALATIKA	MINIMALLY INVASIVE GASTROINTES IN AUSTRICIBLY (MIGS)	11/2/2017
26	TOURS TOUT 40	SITAIK SUMERA BUHAM	MODERATE TO SEVERE ASTRMA	11/2/2017
25	16EB) T0016	INTA VENKALA NAGA VASAVILAT	JOINT DISORDERS	11/2/2017
79. 1	160RITIOIS	BOOYAM BRAYANA	KEINEY A PANCREAS TRANSPLANT	11/2/2017
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DEPARTMENT OF PHARMACY (2021-2022)

CIRCULAR

DATE: 24/03/2021

All the faculty and students here by informed that the Group Discussion Program will be conducted for B pharmacy & Pharm.D students on the 26th March 2021 as per the following schedule. All the faculty & students are instructed to note the same and attend the program as per the schedule.

Copy to faculty members

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Principal / Correspondent

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Group Discussion Session

AY: 2021-2022

Group Discussions help to check students 'interactive skills and how effective they are in communicating with people. The GD is to check how one behave, participate and contribute in a group, how much importance do you give to the group objective as well as your own, how well do you listen to viewpoints of others and how open-minded are you in accepting views contrary to your own. The aspects which make up a GD are verbal communication, non-verbal behavior, and conformation to norms, decision-making ability and cooperation. The following skills are required in GD

- ♦ Communication Skills
- Interpersonal Skills
- ♦ Leadership Skills
- ♦ Motivational Skills
- ◆ Team Building Skills
- ◆ Tolerance
- ♦ Clarity over Ambiguity
- ♦ Divergent Thinking
- ♦ Listening skills
- ♦ Presentation Skills
- ♦ Analytical / Logical skills

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Topic:

Animals should not be used in medical research:

Imprecise results from animal experiments may result in clinical trials of biologically faulty or even harmful substances, thereby exposing patients to unnecessary risk and wasting scarce research resources. Animal toxicity studies are poor predictors of toxic effects of drugs in humans.

Animals should not be used for biomedical research because the experiments are cruel and inhumane. Although animal testing brings more medical advancement and less human-based experiments, it is an expensive way of researching that produce imprecise outputs and at the same time is a practice of animal cruelty.



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Date :

Using Animals for Testing: Pros Versus Cons

- Improves human health: ...
- · Helps ensure safety of drugs: ...
- Alternative methods of testing do not simulate humans in the same way. ...
- · Some substances tested, may never be used for anything useful: ...
- It is very expensive: ...
- Animals and humans are never exactly the same.



Disadvantages:

- Animal experiments are time-consuming and expensive.
- Animal experiments don't accurately mimic how the human body and human diseases respond to drugs, chemicals or treatments.

Animals are very different from humans and, therefore, react differently Batch of 21 B. Pharm students attended the session conducted on 26/03/21

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DEPARTMENT OF PHARMACY (2021-2022)

CIRCULAR

DATE: 22/04/2021

All the faculty and students here by informed that the Debate Program will be conducted for B pharmacy & Pharm.D students on the 24/04/2021 as per the following schedule. All the faculty & students are instructed to note the same and attend the program as per the schedule.

Copy to faculty members

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Debate

AY: 2021-2022

Debate is a process that involves formal discourse on a particular topic, often including a moderator and audience. In a debate, arguments are put forward for common opposing viewpoints. Debates have historically occurred in public meetings, academic institutions, debate halls, coffee houses, competitions, and legislative assemblies. Debates has also been conducted for educational and recreational purposes, usually associated with educational establishments and debating societies.

- 1.Develop excellent oral and written communication skills.
- 2.Develop excellent critical thinking skills.
- 3. Develop effective tools for research, organization and presentation.
- 4. Develop strategies to overcome fears of public speaking.
- 5.Discover the confidence and desire to participate in all academic classes.

Benefits:

- Confidence Belief in themselves and their abilities, and the desire to participate in all classes.
- Curiosity The passion of discovery through effective tools for research, organization and presentation.
- Critical Thinking How to explore the world through the lens of an inquisitive mind.
- Communication Oral & written skills and strategies for lively yet respectful discussions & disagreements.
- Control Eliminate the fears of public speaking.
- Creativity The desire to explore, create and invent.
- Camaraderie Meet like-minded peers at tournaments and build healthy bonds of competition.
- Leadership Self-motivation and the ability to delegate assignments and manage peers.

Debates teaches individuals the importance of being prepared to listen. First, it trains people in the mental preparation of listening – having a listening plan. During a debate you listen for specific things, points you want to answer, weakness in logic, supporting material and key points.

Summary

Using debates in the classroom provides students the opportunity to explore real-world topics and issues. Debates also engage students through self reflection and encourage them to learn from their peers

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Topic:

Are books better than television?

All the research says reading a book is good for you. Better even than listening to an audiobook or reading one on an e-reader. It reduces stress, promotes comprehension and imagination, alleviates depression, helps you sleep and may contribute to preventing Alzheimer's. Reading is active; watching TV is passive.



Reading has a positive effect on our mental health, while watching TV has the exact opposite effect. Reading can reduce stress, lower our blood pressure, our heart rate and muscle tension. On top of the knowledge boost reading provides us with, it also has a healing effect on our mental state.





Dr. K.V. Subba Reddy Institute of Pharmacy

(Approved by AICTE, P.C.I. New Delhi & Permanently Affiliated to JNTUA Anantapuramu, MOU with Government General Hospital & KMC, Kurnool) Opp: Dupadu R.S., N.H - 44, KURNOOL - 518 218, A.P. INDIA. E-mail: principalkvsrip@gmail.com www.drkvsrip.ac.in

Principal / Correspondent

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Date:11/10/2021

DEPARTMENT OF PHARMACY (2021-2022)

CIRCULAR

All the I year B.Pharmacy faculty and students here by informed that the bridge course will be conducted for I year B.Pharmacy & Pharm.D students on 13/10/2021 to 17/10/2021 as per the following schedule. All the faculty & students are instructed to note the same and attend the classes as per the schedule.

S.NO	DATES	TIME	SUBJECT	FACULTY
1	13/10/2021 TO 17/10/2021	9-10AM	PHARMACEUTICAL CHEMISTRY	R. MOHANA PRIYA
2	13/10/2021 TO 17/10/2021	10-11AM	PHARMACEUTICAL ANALYSIS	K.SARA SIRISHA
3	13/10/2021 TO 17/10/2021	11-12AM	MATHEMATICS/BIOLOGY	V.ANU RADHA
4	13/10/2021 TO 17/10/2021	2-3PM	COMMUNICATION SKILLS	S.SREE LEKHA

COPY TO FACULTY MEMBERS

Dr. K. PRINCIPAL

Dr. K. PRINCIPAL R.S. N.H.-44.

Opp: Dupadu R.S. N.H.-44.

KURNOOL-518218 (A.P.)

PRINCIPAL

Dr. K.V.S.R. Institute of Pharmacy
Opp: Dupadu R.S. N.H.-44.

KURNGOL-518218 (A P.)



Dr. K.V. Subba Reddy Institute of Pharmacy

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Principal	/ Correspondent
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PHARMACEUTICAL CHEMISTRY:

S.NO	TOPIC NAME	NO.OF HOURS
1	Introduction to pharmaceutical chemistry	2
2	Basics of chemistry	2
3	General preparations and methods of compounds	2
4	methods of radio pharmaceuticals preparations	2
5	Basic tests for purity of compounds	2
	TOTAL	10

PHARMACEUTICAL ANALYSIS:

S.NO	TOPIC NAME	NO.OF HOURS
1	Basic concepts of analysis	2
2	Basic theories of titration methods	2
3	Different preparations and methods of standardization	
4	Concepts of oxidation & reduction	2
5	Methods of electro chemical methods of analysis	2
	TOTAL	10

& Werthelina Institute of Pharmacy PRINCIPAL Dupadu R.S. N.H. 44. Dr. K.V.S.R. Institute of Pharmacy 518218 (A.P.)

Opp: Dupadu R.S. N.H.-44, KURNOOL-518218 (A.P.)



Dr. K.V. Subba Reddy Institute of Pharmacy

(Approved by AICTE, P.C.I. New Delhi & Permanently Affiliated to JNTUA Anantapuramu, MOU with Government General Hospital & KMC, Kurnool) Opp: Dupadu R.S., N.H - 44, KURNOOL - 518 218, A.P. INDIA. E-mail: principalkvsrip@gmail.com www.drkvsrip.ac.in

Principal /	Corres	pondent

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MATHEMATICS:

S.NO	TOPIC NAME	NO.OF HOURS
1	Theory and their application in pharmacy	2
2	Solving different types of problems by applying theory	2
3	Basic introduction of Logarithms	
4	Basic introduction of Matrices and Determinant	2
5	Applications in solving chemical kinetics and pharmacokinetic equations	2
	TOTAL	10

COMMUNICATION SKILLS:

TOPIC NAME	NO.OF HOURS
Learning Basic communication skills	2
basic learning of listening skills	2
Basic under standing of writing skills	
Learning of presentation skills	2
Learning of interview skills	2
TOTAL	10
	Learning Basic communication skills basic learning of listening skills Basic under standing of writing skills Learning of presentation skills Learning of interview skills

PRINCIPAL

Dr. K.V.S.R. Institute of Pharmacy

PRINCIPAL OL 518213 (A.P.)

Dr. K.V.S.R. Institute of Pharmacy

Opp: Dupadu R.S. N.H.-44.

KURNOOL-518218 (A.P.)

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Di. K.V.S.R. Institute of Pharmacy Opp: Dupadu R.S. N.H.-44, KURNOOL-518218 (A.F.)

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Dr. K.V. Subba Reddy Institute of Pharmacy

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Principal / Correspondent

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1210	. 4			
Date		 	20.00	

Date:20/10/2021

DEPARTMENT OF PHARMACY (2021-2022)

CIRCULAR

All the II year B.Pharmacy faculty and students here by informed that the bridge course will be conducted for II year B.Pharmacy later entry students on 13/10/2021 to 17/10/2021 as per the following schedule. All the faculty & students are instructed to note the same and attend the classes as per the schedule

S.NO	DATES	TIME	SUBJECT	FACULTY
1	13/10/2021 TO 17/10/2021	9-10AM	PHARMACEUTICAL CHEMISTRY	Dr. K. CHANDRA SEKHAR
2	13/10/2021 TO 17/10/2021	10-11	COMPUTER APPLICATIONS	S PAVAN KUMAR

COPY TO FACULTY MEMBERS

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Dr. K.V. PRINCIPAL Opp: Dupadu K.S. N.H.-44, KURNOOL-518218 (A.P.)

PRINCIPAL

Dr. K.V.S.R. Institute of Pharmacy
Opp: Dupadu R.S. N.H.-44.

KURNGOL-518218 (A P)



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Principal / Correspondent

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PHARMACEUTICAL CHEMISTRY:

S.NO	TOPIC NAME	NO.OF HOURS
1	Introduction to pharmaceutical chemistry	2
2	Basics of chemistry	2
3	General preparations and methods of compounds	2
4	methods of radio pharmaceuticals preparations	2
5	Basic tests for purity of compounds	2
	TOTAL	10

COMPUTER APPLICATIONS:

S.NO	TOPIC NAME	NO.OF HOURS
1	Types of applications of computers in pharmacy	2
2	Concepts of information system and softwares used in pharmacy	2
3	Web technologies (HTML,XML,CSS,MS ACCESS)	2
4	Objectives of Bio informatics	2
5	Data analysis in pre clinical development	2
	TOTAL	10

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Dr. K.V.S.R. Institute of Rharmacy (that a of Pharmacy

Opp: Dupadu R.S. N.H.-44,

KURNOOL-518218 (A.P.)

KURNOOL-518218 (A.P.)

DR.K.V.SUBBAREDDY INSTITUTE OF PHARMACY, DUPADU, KURNOOL(AY:21-22)

PHARMACEUTICAL CHEMISTRY:

S.NO	NAME OF THE STUDENT	13/10/21	14/10/21	15/10/21	16/10/21
21ER5R0001	J.DEEPTHI	V	V		
21ER5R0002	V.SAI TARUN KUMAR	V	V	V	V
21ER5R0003	S.SUMANTH	V	1		
21ER5R0004	B.MOHAMMED THOWSIF	V	V		
21ER5R0005	S.MUBEENA BHANU	V	V	V	V
21ER5R0006	B.NIKIL	$\sqrt{}$	J	$\sqrt{}$	V
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COMPUTER APPLICATIONS

S.NO	NAME OF THE STUDENT	13/10/21	18/10/21	15/10/21	16/10/2
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21ER5R0002	V.SAI TARUN KUMAR	V.	V	V	V
21ER5R0003	S.SUMANTH	V	V		V
21ER5R0004	B.MOHAMMED THOWSIF	V .	V	V	V
21ER5R0005	S.MUBEENA BHANU	V	V	V	V
21ER5R0006	B.NIKIL			V	V
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Opp: Dupadu R.S. N.H.-44,
KURNOOL-518218 (A.P.)



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Principal / Correspondent

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DEPARTMENT OF B.PHARMACY

CIRCULAR

Date: 18/09/2021

All the students of II, III & IV B.Pharmacy I semester & Pharm.D who have Secured Less than 50% in I Mid Examinations are hereby informed to attend the Remedial Classes from 5:00PM to 6:00PM during 20.09.2021 to 25.10.2021 I request all the students to utilize this opportunity to improve your percentage.

COPY TO FACULTY MEMBERS

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DRINCIPAL PRINCIPAL Pharmacy Opp: Dupadu R.S. N.H.-44, KURNOOL-518218 (A.P.)

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Dr. K.V.S.R. Institute of Pharmacy
Opp: Dupadu R.S. N.H.:44.

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Department of B. Pharmacy

Following are the students from B.Pharmacy II, III & IV years I semester who need to attend remedial classes without fail.

		II Year I Sem		
20ER1R0009	20ER1R0029	20ER1R0036	20ER1R0039	20ER1R0056
20ER1R0057	20ER1R0089	20ER1R0093	20ER1R0097	20ER1R00A6

		III Year I Sem		
19ER1R0008	19ER1R0014	19ER1R0020	19ER1R0021	19ER1R0022
19ER1R0036	19ER1R0039	19ER1R0049	19ER1R0060	19ER1R0070

	IV Year I Sem		
_18ER1R0008	18ER1R0013	18ER1R0015	-18ER1R0017
18ER1R0034			
	•	18ER1R0008 18ER1R0013	- TODATIONIS

Dr. K.V.S.N. Institute of Pharmacy
Opp: **PRINCIPAL** N.H.-44,
KURNOOL-518218 (A.P.)

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Dr. K.V.S.R. Institute of Pharmacy
Opp: Dupadu R.S. N.H.-44.
KURNOOL-518218 (A P.)



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Data	
Date	

Department of B. Pharmacy

Timetable for Remedial Classes

Year & Sem: II - I

Academic Year: 2021-2022

Day	Timings	Faculty	Subject	Venue	Cian
Monday	5:00PM - 6:00PM	R.MOHANA PRIYA	ORGANIC CHEMISTRY	Room No.106	Muiyer
Tuesday	5:00PM - 6:00PM	B.V.RAMANA	PHYSICAL PHARMACY	Room No.107	av-I

Year & Sem: III - I

Academic Year: 2021-2022

Day	Timings	Faculty	Subject	Venue	Sign
Monday	5:00PM - 6:00PM	K.CHANDRASE	MEDICINAL	Room No.110	~
		KHAR	CHEMISTRY		D.
Tuesday	5:00PM - 6:00PM	S.RAJESH RAJA	PHARMACO LOGY	Room No.111	2

Year & Sem: IV - I

Academic Year: 2021-2022

Day	Timings	Faculty	Subject	Venue	Sign
Monday	5:00PM - 6:00PM	J.GOPALA KRISHNA	INSTRUMENTAL METHODS OF ANALYSIS	Room No.208	Jey Krelm
Tuesday	5:00PM - 6:00PM	S.K.RUBINA	NDDS	Room No.210	Sipro

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Dr. K.V.S.R. Institute of Pharmacy
Opp: Dupaduo S. Dupadu R.S. N.H.-44.

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Department of B. Pharmacy

Attendance for Remedial Classes

Course Name: ORGANIC CHEMISTRY

Year: II - I

Course Faculty: R.MOHANA PRIYA

A.Y: 2021-2022

S. No.	Roll No.	20-09-2021	27-09-2021	04-10-2021	11-10-2021	18-10-2021	25-10-2021
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Department of B. Pharmacy

Attendance for Remedial Classes

Course Name: PHYSICAL PHARMACY

Year: II - I

Course Faculty: DR.B.V.RAMANA

A.Y: 2021-2022

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Department of B. Pharmacy

Attendance for Remedial Classes

Course Name:

MEDICINAL CHEMISTRY

Year: III-I

Course Faculty: DR.K.CHANDRA SEKHAR

A.Y: 2021-2022

S. No.	Roll No.	20-09-2021	27-09-2021	04-10-2021	11-10-2021	18-10-2021	25-10-2021
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Date	 	 	

Department of B.Pharmacy Attendance for Remedial Classes

Course Name: PHARMACOLOGY -I

Year: III-I

Course Faculty: S.RAJESH RAJA

A.Y: 2021-2022

S. No.	Roll No.	21-09-2021	28-09-2021	05-10-2021	12-10-2021	19-10-2021	26-10-2021
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5	19ER1R0022	V	4	×	/	~	V
6	19ER1R0036	1	X	1	V		5
7	19ER1R0039	~	/	V	X	V	
8	19ER1R0049		V	1	~	~	
9	19ER1R0060	V	Y	1	V	~	
10	19ER1R0070	/	V	V	×	V	V
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PRINCIPAL Dr. K.V.S.R. Institute of Pharmacy Opp: Dupadu R.S. N.H.-44, KURNOOL-518218 (A P)



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Department of B.Pharmacy

Attendance for Remedial Classes

Course Name: INSTRUMENTATION METHODS OF ANALYSIS

Year: IV-1

Course Faculty: J.GOPALA KRISHNA

A.Y: 2021-2022

S. No.	Roll No.	20-09-2021	27-09-2021	04-10-2021	11-10-2021	18-10-2021	25-10-2021
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Dr. K.V.S.R. Institute of Pharmacy
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Department of B. Pharmacy

Attendance for Remedial Classes

Course Name: NDDS

Year: IV-1

Course Faculty: S.K.RUBINA

A.Y: 2021-2022

S. No.	Roll No.	21-09-2021	28-09-2021	05-10-2021	12-10-2021	19-10-2021	26-10-2021
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Dr. K.V.S.R. Institute of Pharmacy

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Dr. K.V.S.R. Institute of Pharmacy
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Department of Pharm.D

Following are the students from PHARM.D II, III & IV years who need to attend remedial classes without fail.

		II Y	ear		
20ER1T0002	20ER1T0005	20ER1T0016	20ER1T0017	20ER1T0020	20ER1T0028

		III Y			
19ER1T0005	19ER1T00015	19ER1T00018	19ER1T0021	19ER1T0024	19ER1T0025
19ER1T0027	19ER1T0029				

		IV Year		
18ER 1T0003	18ER 1T0014	18ER 1T0019	18ER 1T0027	

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Dr. K.V.S.R. Institute of Pharmacy

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Department of Pharm.D

Timetable for Remedial Classes

Year & Sem: II

Academic Year: 2021-2022

Day	Timings	Faculty	Subject	Venue	Sign
Monday	5:00PM - 6:00PM	DR. M. SPURTHI MYTHRA	PT-I	Room No.106	\$
Tuesday	5:00PM - 6:00PM	DR.S.RAJESH RAJA	P.COL-I	Room No.107	Rajed

Year & Sem: III

Academic Year: 2020-21

Day	Timings	Faculty	Subject	Venue	Sign
Monday	5:00PM - 6:00PM	DR.S. KUSUMA KUMARI	PT-II	Room No.110	40
Tuesday	5:00PM - 6:00PM	DR.K. CHANDRASEKH AR	MC	Room No.111	W

Year & Sem: IV

Academic Year: 2020-21

Day	Timings	Faculty	Subject	Venue	Sign
Monday	5:00PM - 6:00PM	SURESH BABU	Biostatistics & Research Methodology	Room No.208	SE
Tuesday	5:00PM - 6:00PM	B.ARUNA	CT	Room No.210	(8 1:

Dr. K.V.S.R. Institute of Pharmacy Opp: Dupadu R.S. N.H.-44, Opp: Dupadu R.S. N.H.-44, WINOOL-518218 (A.P.)

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Department of Pharm.D

Attendance for Remedial Classes

Course Name: PT-I

Year: II

Course Faculty: DR. M. SPURTHI MYTHRA

A.Y: 2021-2022

S. No.	Roll No.	20-09-2021	27-09-2021	04-10-2021	11-10-2021	18-10-2021	25-10-2021
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Dr. K.V.S.R. Institute of Pharmacy Opp: Dupadu R.S. N.H.-44, KURNOOL-518218 (A P)



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Department of Pharm.D

Attendance for Remedial Classes

Course Name: PHARMACOLOGY-I

Year: II

Course Faculty: DR.S.RAJESH RAJA

A.Y: 2021-2022

S. No.	Roll No.	21-09-2021	28-09-2021	05-10-2021	12-10-2021	19-10-2021	26-10-2021
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5	20ER1T0020	V					
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Department of Pharm.D

Attendance for Remedial Classes

Course Name: PT -II

Year: III

Course Faculty: DR.S. KUSUMA KUMARI

A.Y: 2021-2022

S. No.	Roll No.	20-09-2021	27-09-2021	04-10-2021	11-10-2021	18-10-2021	25-10-2021
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5	19ER1T0018	V		V		V	
6	19ER1T0021	1	V	1	V		
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Department of Pharm.D

Attendance for Remedial Classes

Course Name: MEDICINAL CHEMISTRY

Year: III

Course Faculty: DR.K.CHANDRASEKHAR

A.Y: 2021-2022

S. No.	Roll No.	21-09-2021	28-09-2021	05-10-2021	12-10-2021	19-10-2021	26-10-2021
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Principal / Correspondent

Date:.....

Department of Pharm.D

Attendance for Remedial Classes

Course Name: BIOSTATISTICS & RESEARCH METHODOLOGY Year: IV

Course Faculty: SURESH BABU

A.Y: 2021-2022

S. No.	Roll No.	20-09-2021	27-09-2021	04-10-2021	11-10-2021	18-10-2021	25-10-2021
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Opp: Dupadu R.S. N.H.-44,
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Department of Pharm.D

Attendance for Remedial Classes

Course Name: BIOSTATISTICS & RESEARCH METHODOLOGY Year: IV

A.Y: 2021-2022

Course Faculty: SURESH BABU

S. No.	Roll No.	20-09-2021	27-09-2021	04-10-2021	11-10-2021	18-10-2021	25-10-2021
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Dr. K.V.S.H. Institute of Pharmacy Opp: Dupadu R.S. N.H.-44, KURNOGL-518218 (A.P.)

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Dr. K.V.S.R. Institute of Pharmacy

Opp: Dupadu R.S. N.H.-44.

KURNOOL-518218 (A.P.)

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Principal	/ Correspondent

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Department of Pharm.D

Attendance for Remedial Classes

Course Name: CLINICAL TOXICOLOGY

Year: IV

Course Faculty: B.ARUNA

A.Y: 2021-2022

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S. No.	Roll No.	21-09-2021	28-09-2021	05-10-2021	12-10-2021	19-10-2021	26-10-2021
1	18ER 1T0003	X					
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Department of B.Pharmacy

Circular

Date: 09/04/2022

All the students of II, III & IV B.Pharmacy II Semester who have Secured Less than 50% in I Mid Examinations are hereby informed to attend the Remedial Classes from 5:00PM to 6:00PM during 11/04/2022to 17/05/2022. I request all the students to utilize this opportunity to improve your percentage.

Copy to faculty members:

Dr. K.V.PRINCIPALOT Pharmacy Opp: Dupadu R.S. N.H.-44, KURNOOL-518218 (A.P.)

Dr. K.V.S.R. Institute of Pharmacy Opp: Dupadu R.S. N.H.-44. KURNOOL-518218 (A.P.)



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Department of B.Pharmacy

Following are the students from B. Pharmacy II, III & IV years II semester who need to attend remedial classes without fail.

		II Year II Sem		
20ER1R0009	20ER1R0029	20ER1R0036	20ER1R0039	20ER1R0056
20ER1R0057	20ER1R0089	20ER1R0093	20ER1R0097	20ER1R00A6

		III Year II Sem	1	
19ER1R0008	19ER1R0014	19ER1R0020	19ER1R0021	19ER1R0022
19ER1R0036	19ER1R0039	19ER1R0049	19ER1R0060	19ER1R0070

		IV Year II Sem		
18ER1R0001	18ER1R0008	18ER1R0013	18ER1R0015	18ER1R0017
18ER1R0018	18ER1R0034		+	

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Dr. K.V.S.R. Institute of Pharmacy

Dani Dupadu R.S.N.H.-44.

RURNOOL-5-PRINCIPAL

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Date :

Department of B.Pharmacy

Timetable for Remedial Classes

Year & Sem: II -II

Academic Year: 2021-2022

Day	Timings	Faculty	Subject	Venue	Sign
Monday	5:00PM - 6:00PM	DR.J.GOPALA KRISHNA	MC-I	Room No.102	4
Tuesday	5:00PM - 6:00PM	E.HONEY	PHARMACO LOGY-I	Room No.102	211

Year & Sem: III - II

Academic Year: 2021-22

Sign
A
- 101

Year & Sem: IV - II

Academic Year: 2021-22

Day Timings Faculty Subject Venue Sign Monday 5:00PM - 6:00PM P.T.NAGARAJU MOOCS Room No.104 -II Tuesday 5:00PM - 6:00PM K.SARA SIRISHA MOOCS Room No.104 -III

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F 1			
1210			
Date	 	 	

Department of B.Pharmacy

Attendance for Remedial Classes

Course Name: MC-I

Year: II - II

Course Faculty: J.GOPALA KRISHNA

A.Y: 2021-2022

S. No.	Roll No.	11-04-2022	18-04-2022	25-04-2022	2-05-2022	9-05-2022	16-05-2022
1	20ER1R0009	*		1	1		10 05 2022
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4	20ER1R0039				~		1
5	20ER1R0056	V					
6	20ER1R0057	×	1	X			V
7	20ER1R0089						
8	20ER1R0093	1/		N.C.	. /		X
9	20ER1R0097		X				
10	20ER1R00A6	V					V
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Department of B.Pharmacy

Attendance for Remedial Classes

Course Name: PHARMACOLOGY-I

Year: II - II

Course Faculty: E.HONEY

A.Y: 2021-2022

S. No.	Roll No.	12-04-2022	19-04-2022	26-04-2022	03-05-2022	10-05-2022	17-05-2022
1	20ER1R0009	V		1			
2	20ER1R0029		1/				N/
3	20ER1R0036	/	V	/			7
4	20ER1R0039	/			X		
5	20ER1R0056		1	X			
6	20ER1R0057	*X;		1			V
7	20ER1R0089	-/		~		2/	
8	20ER1R0093	X	V				
9	20ER1R0097	~	- /			3/	
10	20ER1R00A6			~	/		1
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Dr. K.V.S.R/Institute of Pharmacy
Opp: PRINCIPAL
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Data	6.5			
Date		 	 	

Department of B.Pharmacy

Attendance for Remedial Classes

Course Name: MC-III

Year: III-II

Course Faculty: R.MOHANA PRIYA

A.Y: 2021-2022

S. No.	Roll No.	11-04-2022	18-04-2022	25-04-2022	2-05-2022	9-05-2022	16-05-2022
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4	19ER1R0021	×	1/	1	~		1
5	19ER1R0022		1	~	- /	1	
6	19ER1R0036				×		
7	19ER1R0039					×	1
8	19ER1R0049	~		1			1
9	19ER1R0060						×
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Department of B.Pharmacy

Attendance for Remedial Classes

Course Name: BIO TECHNOLOGY

Year: III-II

Course Faculty: DR.S.KUSUMA KUMARI

A.Y: 2021-2022

S. No.	Roll No.	12-04-2022	19-04-2022	26-04-2022	03-05-2022	10-05-2022	17-05-2022
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5	19ER1R0022	X	V	V	1/	1	-
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7	19ER1R0039		X	~		V	V
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9	19ER1R0060			×		V	~
10	19ER1R0070			V		×	~
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Department of B.Pharmacy

Attendance for Remedial Classes

Course Name: MOOCS-II

Year: IV-II

Course Faculty: P.T.NAGARAJU

A.Y: 2021-2022

S. No.	Roll No.	11-04-2022	18-04-2022	25-04-2022	2-05-2022	9-05-2022	16-05-2022
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3	18ER1R0015		4	1/		1/	1
4	18ER1R0017	×	1	V	1/	1/	1/
5	18ER1R0018		X	1/			
6	18ER1R0034	/	V	X	~	~	V
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Opp: Dupadu R.S. N.H.-44, - KURNOOL-\$18218 (A.P.)

Dr. K.V.S.R. Institute of Pharmacy Opp: Dupadu R.S. N.H.-44 KURN@OL-518218 (A.P.)



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Department of B.Pharmacy

Attendance for Remedial Classes

Course Name: MOOCS-III

Year: IV-II

Course Faculty: K.SARA SIRISHA

A.Y: 2021-2022

S. No.	Roll No.	12-04-2022	19-04-2022	26-04-2022	03-05-2022	10-05-2022	17-05-2022
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4	18ER1R0017			×	. /		
5	18ER1R0018	V	/			X	
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Dr. K.V. PRINCIPAT harmacy Opp: Dupadu R.S. N.H.-44 KURNOOL-518218 (A.P.)

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Department of Pharmacy

Laboratory Sessions

Branch: B.Pharmacy Regulation: R19



Fig 1.1 Experimental Lab on Limit test in Pharmaceutical Inorganic Laboratory



Fig 1.2 Drying of powder in hot air oven in Physical Pharmaceutical Laborator

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Date :

Department of Pharmacy

Laboratory Sessions

Branch: Pharm.D Regulation: R19



Fig 1.3 Experimental Lab on Titration in Pharmaceutical Analysis Laboratory

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OPP. Dir OL-513218 (A.P.)

KURWOL-513218 (A.P.)

Date:.....



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Fig 1.4 Experimental Lab in Medicinal Chemistry Laboratory

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Dr. E.S. N.H.-44,
Option 5182188 (A.P.1)



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Date :

Pharma Quiz

The Department of Pharmacy has conducted a **Pharma Quiz Competition** in the college auditorium on 18th March 2021 under Pharmacy students Chapter in which four different teams with five participants in each group gave tough battle to each other.

Mrs.S.Lakshmi Garu presented mementos to the winners.





Fig 1: Presenting Mementos to the winners by Principal

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Department of Pharmacy

Think Pair Share Session

AY: 2021-22

Think-Pair-Share (TPS) is a cooperative learning activity that can work in varied size classrooms and in any subject. Instructors pose a question, students first THINK to themselves prior to being instructed to discuss their response with a person sitting near them (PAIR).

In other words, Think-pair-share (TPS) is a collaborative learning strategy where students work together to solve a problem or answer a question about an assigned reading. This strategy requires students to (1) think individually about a topic or answer to a question; and (2) share ideas with classmates. Discussing with a partner maximizes participation, focuses attention and engages students in comprehending the reading material.

- It helps students to think individually about a topic or answer to a question.
- It teaches students to share ideas with classmates and builds oral communication skills.
- It helps focus attention and engage students in comprehending the reading material.



Fig.1: Students during a Think Pair Share Session

Batch of 52 Pharmacy students attended the session conducted on 7/4/2021.

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CLUB ACTIVITIES:

This club activities is organized by students as well as faculty members is to promote & enhance activity in educational institution .this kind of activities empower the students analytical interpretation skills. It provide familiarize members with emerging trends in the field of pharmacy. It provide environment for learning inter disciplinary academic research.





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Opp: Dupadu R.S. (A.P.)
KURNOOL-518218 (A.P.)

Dr. K. V. SUBBA REDDY INSTITUTE OF PHARMACY

(Approved by AICTE & PCI New Delhi, Permanent Affiliated to JNTU Anantapur, Anantapuramu, MoU with Government General Hospital, KMC, Kurnool)

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E-mail:principalkvsrip@gmail.com

Cell: +919440282181, +919704333789

Fax: 08518-287618

DATE:07/06/2021

DEPARTMENT OF PHARMACY (2021-2022)

CIRCULAR

All the faculty and students here by informed that Work Shop on Training and Handling of Dissolution Apparatus will be conducted for III- & IV-year B. pharmacy students on 09/06/2021 to 10/06/2021 as per the following schedule. All the faculty & students are instructed to note the same and attend the classes as per the schedule.

Copy to faculty members

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KURNOOL-518218 (A P)

Date :



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PROBLEM SOLVING METHODS Dr.K.V.Subba Reddy institute of pharmacy conducted training on handling of dissolution apparatus for B.Pharmacy III&IV Year students on 21/08/2021 to 24/08/2021. This workshop is very helpful to acquire knowledge, conceptual under standing, and skills to solve problems and make informed decision in scientific context. Eminent speaker S.K.RUBINA Madam, M. Pharmacy, Assistant professor Department of pharmaceutical analysis.





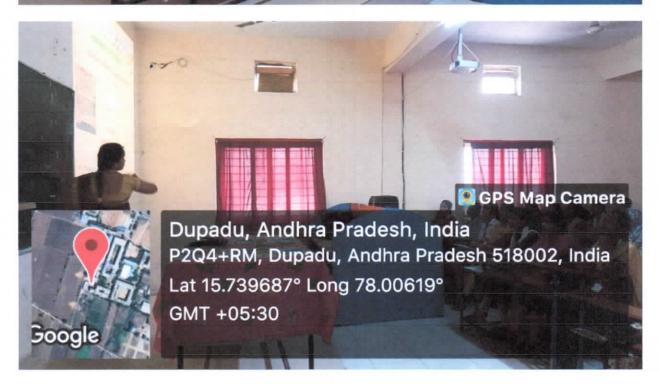


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Principal / Correspondent

Date : GPS Map Camera Dupadu, Andhra Pradesh, India P2Q4+RM, Dupadu, Andhra Pradesh 518002, India Lat 15.739766° Long 78.006073° GMT +05:30 Google



S.K. Rubina garu discussing about working of dissolution apparatus

Dr. K.V.S.R. Institute of Pharmacy Opp: Dupadu R.S. N.H.-44, KURNOOL-518218 (A.P.)



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Date :



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Principal / Correspondent

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P2Q4+RM, Dupadu, Andhra Pradesh 518002, India
Lat 15.73955° Long 78.006326°

GMT +05:30

Dr. N.S. P. Dupadu 518218 (A.R.)



Dr. K.V. Subba Reddy Institute of Pharmacy

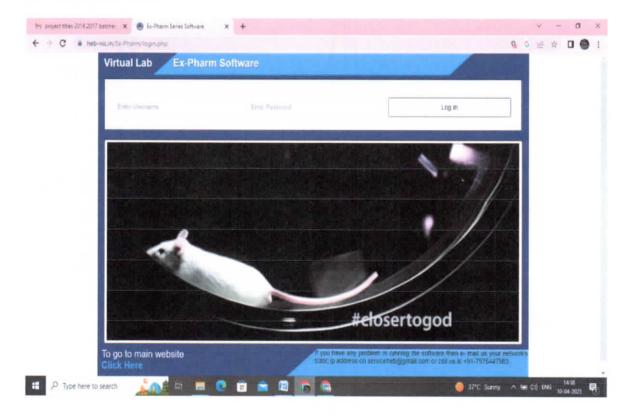
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VIRTUAL LABS

Virtual labs are used to provide remote asses to stimulation based labs in various disciplines of sciences and pharmacy departments. By using these virtual labs to enthuse students to conduct experiments by arousing their curiosity. This would help them in learning basic & advanced concepts through remote experimentation. This would help them to provide complete learning management system around the virtual labs where the students/ teachers can avail the various tools for learning, including additional web resources, video lectures, animated demonstrations and self examination.



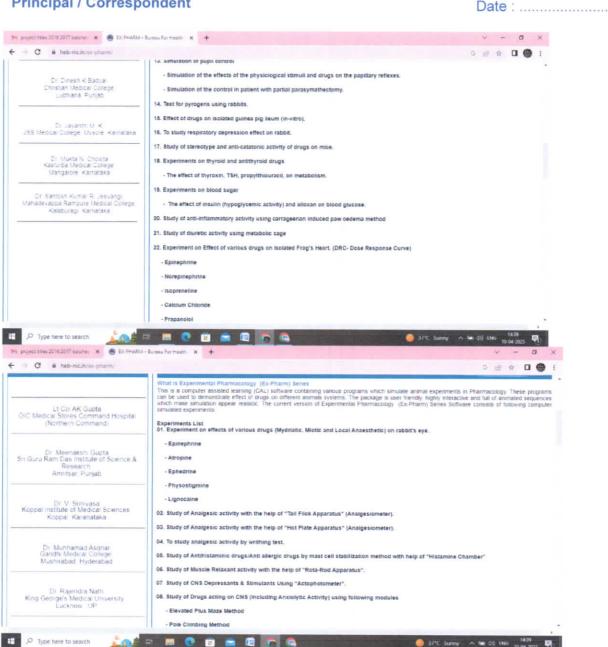




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B.Pharmacy II year students participative learning in pharmacology laboratory





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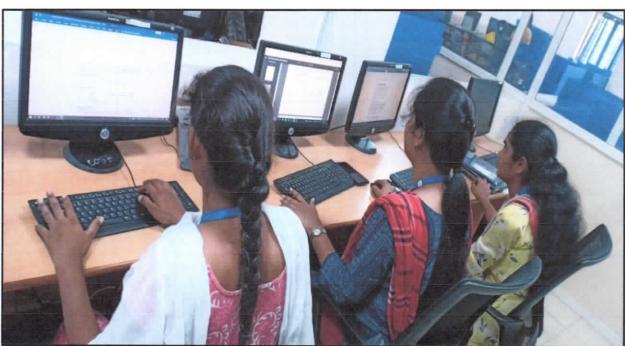


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