

Teaching-Learning Plan

Program Level & Semester/Year: B. Pharm 7th (UG)

No. of classes per week: 4 hrs

Lecture No.	Unit / Chapter	Syllabus Topics to be Covered
02	I	Introduction of chromophores and auxochromes, spectral shifts
03	I	Solvent effect on absorption spectra, Derivation of Beer & Lambert's law, deviations
04	I	Instrumentation of UV-vis spectroscopy, Discussion on applications
05	I	Theory of Fluorimetry, Concepts of Singlet, doublet, and triplet electronic states
06	I	Factor affecting fluorescence and quenching
07	I	Instrumentation and applications of fluorimeter
08	II	Introduction to IR spectroscopy and fundamental modes of vibrations in polyatomic molecules.
09	II	Types of vibrations associated with IR detectors and applications.
10	II	Instrumentation of IR, Discussion on IR detectors and applications.
11	II	Principle of flame photometry and interferences.
12	II	Instrumentation and applications of flame photometry
13	II	Principle of AAS and interferences associated with it
14	II	Instrumentation and application of AAS
15	III	Introduction to chromatography and types of it
16	III	Elaborate absorption and partition column chromatography
17	III	Methodology and techniques of packing of column
18	III	Advantage and disadvantage of Column chromatography.

Teaching-Learning Plan

Course & Code: B. Pharm BF701T

Academic Session: 2020-21

Section: A

Mode of Delivery Used	Text / Ref. Book / Resource	Date of Lecture		Remarks	Signature of Principal
		Scheduled	Actual		
Blackboard	Instrumental methods of Chromatography by R.K. Sharma	07/09/2020	07/09/2020		
"	"	09/09/2020	09/09/2020		
"	"	10/09/2020	10/09/2020		
Powerpoint	e-Resource (Middlestone)	21/09/2020	21/09/2020		
Blackboard	Instrumental methods of analysis Chitra	22/09/2020	22/09/2020		
"	"	23/09/2020	23/09/2020		
"	"	24/09/2020	24/09/2020		
"	"	26/09/2020	26/09/2020		
"	"	28/09/2020	28/09/2020		
"	"	29/09/2020	29/09/2020		
"	"	30/09/2020	30/09/2020		
"	"	5/10/2020	5/10/2020		
"	"	9/10/2020	9/10/2020		
"	"	10/10/2020	10/10/2020		
"	"	12/10/2020	12/10/2020		
"	"	17/10/2020	17/10/2020		
"	"	19/10/2020	19/10/2020		
"	"	24/10/2020	24/10/2020		

Teaching-Learning Plan

Program Level & Semester/Year: B.Pharm 7th sem (UG)

No. of classes per week: 4 hours

Lecture No.	Unit / Chapter	Syllabus Topics to be Covered
20	III	Discussion on common detecting reagents for selective compound. R _f values
21	III	Advantages, disadvantages and application of TLC.
22	II	Difference between nephelometer and Turbidimeter principle associated with it.
23	I	Instrumentation & application of nepheloturbidometry.
24	III	Introduction to paper chromatography and methods logs adopted
25	II	Development techniques of paper chromatography
26	I	Advantages, disadvantages and application of paper chromatography
27	II	Introduction to electrophoresis and factor affecting electrophoretic mobility.
28	III	Techniques of paper, gel and capillary electrophoresis
29	II	Introduction and theory involved in gas chromatography
30	III	Instrumentation and derivatization of GC.
31	IV	Temperature programming, advantages, disadvantages and application of GC chromatography
32	IV	Introduction, and theory involved in HPLC.
33	IV	Instrumentation, and advantages of HPLC.
34	V	Introduction, classification of Ion Exchange resin.
35	V	Mechanism of Ion exchange process, factor affecting ion exchange
36	V	Methodology of IEC and its application.

Teaching-Learning Plan

Course & Code: B.Pharm; BP7017

Academic Session: 2020-21

Section: A

Mode of Delivery Used	Text / Ref. Book / e-Resource	Date of Lecture		Remarks	Signature of Principal
		Scheduled	Actual		
Blackboard	Instrumental method of analysis of drug & natural product by BP 7017	23/10/2020	23/10/2020		
"	"	31/10/2020	31/10/2020		
"	"	2/11/2020	2/11/2020		
"	"	7/11/2020	7/11/2020		
"	"	9/11/2020	9/11/2020		
"	"	14/11/2020	14/11/2020		
"	"	16/11/2020	16/11/2020		
Powerpoint	e-resource (Microchem)	21/11/2020	21/11/2020		
Blackboard	Instrumental method of analysis of drug & natural product	23/11/2020	23/11/2020		
"	"	24/11/2020	24/11/2020		
"	"	26/11/2020	26/11/2020		
"	"	28/11/2020	28/11/2020		
"	"	30/11/2020	30/11/2020		
"	"	2/12/2020	2/12/2020		
"	"	4/12/2020	4/12/2020		
"	"	7/12/2020	7/12/2020		
"	"	9/12/2020	9/12/2020		
"	"	11/12/2020	11/12/2020		

Record of Internal Assessment and Continuous Evaluation

Semester/Year:		Course:				Course Code:			Vno/GD
Roll No.	Name of Students	Assignments				Sessional Exam			
		A-1	A-2	A-3	A-4	Test-1	Test-2	Test-3	
						24	26	0	
		19	17	16		24	26		
13 & 5100 (100)		14	14	20		19	19	20	
02		15	14	17		18	19	17	
03		17	18	17		21	19	18	
04		17	16	14		26	28	26	
05		18	18	17		23	25	19	
06		14	18	20		27	27	23	
08		19	17	18		21	22	23	
09		18	18	17		24	26	21	
10		16	16	17		20	24	26	
11		14	20	19		25	23	22	
12		20	20	19		21	24	20	
13		16	17	17		21	25	13	
14		19	18	19		23	23	19	
15		17	19	18		25	26	20	
16		19	19	19		24	25	0	
17		19	19	19		22	23	0	
18		18	14	16		24	25	15	
19		19	17	16		25	25	16	
20		17	18	17		22	21	9	
21		15	16	15		27	27	9	
22		20	14	14		24	21	11	
23		17	18	14		21	25	9	
25		14	18	18		22	20	0	
26		18	18	18		21	20	0	
27		15	18	18		19	18	0	
28		20	20	18		23	23	0	
29		20	19	18		24	23	13	
30		19	14	17		26	27	0	
31		14	17	20		21	20	0	
32		20	20	19		17	15	0	
33		16	17	16		19	18	0	
34		18	18	19		20	21	0	
35		17	19	18		20	21	12	
36		14	14	14		0	0		
37									

Record of Internal Assessment and Continuous Evaluation

Semester/Year:		Course:				Course Code:			Vno/GD
Roll No.	Name of Students	Assignments				Sessional Exam			
		A-1	A-2	A-3	A-4	Test-1	Test-2	Test-3	
38		18	19	18		25	17	17	
39		19	17	16		42	28	16	
40		18	17	18		19	20	12	
41		19	19	19		24	25	13	
42		18	19	17		0	0	0	
43		17	18	17		27	17	0	
44		17	16	17		21	25	0	
45		18	18	17		26	27	19	
46		19	20	20		27	26	0	
47		18	13	18		14	23	12	
48		15	15	14		23	23	0	
49		18	17	17		22	23	21	
50		14	14	14		14	17	0	
51		18	18	17		26	27	29	
52		12	19	17		23	24	0	
53		17	16	17		26	25	19	
54		18	17	18		23	12	25	
55		14	14	14		28	26	18	
56		18	19	17		26	24	26	
58		17	18	17		26	26	29	
59		17	16	17		25	23	26	
60		18	18	17		19	17	30	
61		14	20	20		14	11	21	
62		18	17	18		25	23	26	
64		18	18	17		15	16	0	
65		16	16	17		16	15	21	
66		19	21	20		25	16	24	
68		20	20	19		26	09	26	
69		16	17	16		27	20	29	
70		12	11	19		24	4	28	
71		18	18	17		24	18	29	
72		16	16	17		14	05	06	
73		14	20	20		21	19	19	
74		20	20	19		23	27	26	
76		16	13	16		27	20	29	

Record of Internal Assessment and Continuous Evaluation

Semester/Year:

Course:

Course Code:

Roll No.	Name of Students	Assignments				Sessional Exam			Viva / GD
		A-1	A-2	A-3	A-4	Test-1	Test-2	Test-3	
77		18	18	19		23	06	21	
78		18	18	17		20	01	15	
79		16	16	17		18	03	18	
80		19	20	20		24	11	22	
81		20	20	19		21	17	18	
82		16	17	16		26	15	20	
83		18	18	19		24	10	23	
84		17	17	16		19	03	13	
85		20	20	20		22	03	16	
86		19	18	20		25	29	29	
87		16	17	17		26	21	21	
88		19	17	18		26	21	22	
89		20	20	20		28	26	20	
90		20	20	19		27	27	20	
91		16	17	16		23	25	19	
93		18	18	19		22	24	16	
94		18	18	17		28	20	0	
95		19	20	20		21	20	14	
96		18	17	18		23	22	16	
97		18	18	17		25	24	16	
98		16	16	17		15	11	0	
99		19	20	20		25	25	17	
100		19	16	18		24	26	16	
101		16	20	16		21	18	12	
1051001026		19	20	19		26	27	18	
L-01		17	17	20		21	18	12	
L-02		20	18	16		21	20	5	
L-03		18	18	18		19	20	0	
L-04		17	20	18		19	23	17	
L-05		17	19	19		22	19	13	
L-06		20	18	18		23	24	18	
L-07		15	19	16		19	19	13	
L-10		19	20	19		20	19	0	

Teaching Learning Plan

Name of the Course: Instrumental Method of Analysis (Theory)

Course Code: BP701T

Semester: B. Pharm 7th Sem.

Year: 2020-21

Name of the Teacher: Mr. Susankar Kushari & Mr. Suman Kumar

Designed Course Outcome:

1. Understand the interaction of matter with electromagnetic radiations and its applications in drug analysis.
2. Understand the chromatographic separation and analysis of drugs.
3. Perform quantitative & qualitative analysis of drugs using various analytical instruments.

Tentative Dates	Syllabus Topics to be Covered	Learning outcome	Contribution to which CO	Mode of Delivery	Assessment Tools (With tentative date of Assessment) for measuring Outcome
Unit I Lecture Aug: 04	Introduction to syllabus			<ul style="list-style-type: none"> Lecture with board work. 	
Unit I Lectures Aug: 10,11,12,17	Electronic transitions, chromophores, auxochromes, spectral shifts	<ul style="list-style-type: none"> Highlight the different electronic transitions. Discuss the examples of chromophores along with auxochromes. Explain the spectral shifts. 	CO. 1 CO. 2	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion Lecture handouts 	
Unit I Lectures Aug: 17, 18,19	Solvent effect on absorption spectra. Beer and Lambert's law, Derivation and deviations	<ul style="list-style-type: none"> Describe the effect of solvents on absorption spectra. Derivation of Beer Lambert's law 	CO. 2	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion Lecture handouts 	
Unit I Lectures Aug: 24,25,26	Instrumentation - Sources of radiation, wavelength selectors, sample cells Detectors:	<ul style="list-style-type: none"> Discuss instrumentation and applications of UV-Vis spectroscopy 	CO. 3 CO. 4	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion 	

	Photo tube, Photomultiplier tube, Photo voltaic cell. Silicon Photodiode along with applications.			<ul style="list-style-type: none"> Lecture handouts 	
Unit I Lectures Sep: 1,2,3	Theory, Concepts of singlet, doublet and triplet electronic states, internal and external conversions	<ul style="list-style-type: none"> Discuss the principle associated with fluorimetry. Elaborate concepts of singlet, doublet and triplet electronic states. 	CO. 1 CO. 3 CO. 4	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion 	
Unit I Lectures Sep: 8,9,10,15	Factors affecting fluorescence, quenching, instrumentation and applications	<ul style="list-style-type: none"> Explain factors affecting fluorescence intensity. Discuss the instrumentation and applications of spectrofluorimeter. 	CO. 1 CO. 2	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion 	1st Class Test Assessment date: 4 th -11 th September 2020 Assignment 1 Assessment date: 19 Aug 2020
Unit II Lectures Sep:21,22, 23	Introduction to fundamental modes of vibrations in poly atomic molecules, sample handling, factors affecting vibrations	<ul style="list-style-type: none"> Discuss the principle associated with IR spectroscopy. Elaborate different vibrations. 	CO. 1 CO. 3 CO. 4	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion LCD projector Lecture handouts 	
Unit II Lectures Sep: 28, 29,30	Instrumentation - Sources of radiation, wavelength selectors, detectors - Golay cell, Bolometer, Thermocouple, Thermister, Pyroelectric detector	<ul style="list-style-type: none"> Explain instrumentation and applications of IR spectroscopy 	CO. 3 CO. 4	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion Lecture handouts 	

	and applications			
Unit II Lectures Oct:5,6,7	Principle, interferences, instrumentation and applications of flame photometer.	<ul style="list-style-type: none"> • Discuss the principle of flame photometer. • Explain the different types of interferences. • Discuss the instrumentation and applications. 	CO,4	<ul style="list-style-type: none"> • Lecture with board work. • Interactive discussion • Lecture handouts
Unit II Lectures Oct:12, 13,14	Principle, instrumentation and applications of nepheloturbidimetry.	<ul style="list-style-type: none"> • Elaborate the principle of nepheloturbidimetry. • Distinguish between nephelometer and turbidimeter. • Discuss the instrumentation and applications. 	CO,3 CO,4	<ul style="list-style-type: none"> • Lecture with board work. • Interactive discussion •
Unit III Lectures Oct:19,20, 21	Introduction to different types of chromatography. Methodology, advantages, disadvantages and applications of column chromatography.	<ul style="list-style-type: none"> • Explain the different types of chromatography. • Elaborate the principle associated with adsorption and partition chromatography. • Advantages and disadvantages of column chromatography. • Discuss its applications. 	CO,1 CO,3 CO,4	<ul style="list-style-type: none"> • Lecture with board work. • Interactive discussion • Lecture handouts
Unit III Lectures Oct:26, 27, 28	Introduction, Principle, Methodology, R _f values, advantages, disadvantages and applications of TLC.	<ul style="list-style-type: none"> • Discuss the principle of TLC. • Elaborate the advantages and disadvantages of TLC. • Discuss the applications of TLC. 	CO,1 CO,3 CO,4	<ul style="list-style-type: none"> • Lecture with board work. • Interactive discussion • Lecture handouts
Unit III Lectures Nov:2,3,4	Introduction, methodology, development techniques, advantages,	<ul style="list-style-type: none"> • Discuss the principle of TLC. • Elaborate the advantages and disadvantages of 	CO,1 CO,3 CO,4	<ul style="list-style-type: none"> • Lecture with board work. • Interactive discussion

	disadvantages and applications of paper chromatography.	<p>paper chromatography.</p> <ul style="list-style-type: none"> Discuss the applications of paper chromatography. 		<ul style="list-style-type: none"> Lecture handouts 	
Unit III Lectures Nov:9,10, 11	Introduction, factors affecting electrophoretic mobility, Techniques of paper, gel, capillary electrophoresis, applications.	<ul style="list-style-type: none"> Explain the concept of electrophoresis. Discuss the factors affecting electrophoresis. Elaborate different techniques of paper, gel and capillary electrophoresis. Discuss the applications of electrophoresis. 	CO.1 CO.4	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion Lecture handouts 	<p>2nd Class Test Assessment date: 9-16th Nov 2020</p> <p>Assignment 2 Assessment date: 12 Nov.</p>
Unit IV Lectures Nov:9,10, 11	Introduction, theory, instrumentation, derivatization, temperature programming, advantages, disadvantages and applications of gas chromatography.	<ul style="list-style-type: none"> Explain the principle of gas chromatography. Discuss the instrumentation and temperature programming. Elaborate the advantages, disadvantages and applications of gas chromatography. 	CO.1 CO.3 CO.4	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion Lecture handouts 	
Unit IV Lectures Nov:16, 17,18	Introduction, theory, instrumentation, advantages and applications of HPLC.	<ul style="list-style-type: none"> Discuss the principle of HPLC. Illustrate the instrumentation of HPLC. Explain the advantages and applications of HPLC. 	CO.3 CO.4	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion Lecture handouts 	
Unit IV Lectures Nov:23,24 ,25,27	Introduction, classification, ion exchange resins, properties, mechanism of ion exchange process, factors affecting ion exchange,	<ul style="list-style-type: none"> Discuss the classification of ion exchange resin. Illustrate the mechanism of ion exchange process. 	CO.1 CO.3 CO.4	<ul style="list-style-type: none"> Lecture with board work. Interactive discussion 	

	methodology and applications	<ul style="list-style-type: none"> • Elaborate the factors affecting ion exchange process. • Explain the application of IEC. 			
Unit IV Lectures Dec:7,8,9	Introduction, theory, instrumentation and applications of gel chromatography.	<ul style="list-style-type: none"> • Discuss the theory of gel chromatography. • Illustrate the instrumentation of gel chromatography. • Discuss the applications. 	CO. 1 CO. 4	<ul style="list-style-type: none"> • Lecture with board work. • Interactive discussion 	3rd Class Test Assessment date: 14-21 Dec 2020 Assignment 2 Assessment date: 7 Dec.
Unit IV Lectures Dec:14,15, 16	Introduction, theory, instrumentation and applications of affinity chromatography.	<ul style="list-style-type: none"> • Discuss the theory of affinity chromatography. • Illustrate the instrumentation of affinity chromatography. • Discuss the applications. 	CO. 1 CO. 4	<ul style="list-style-type: none"> • Lecture with board work. • Interactive discussion • Lecture handouts 	

Submitted by:

Sig:
Subject Teacher

Verified by:

Sig:
Module Co-ordinator

Teaching Learning Plan

Name of the Course: Instrumental Method of Analysis (Practical)

Course Code: BP705P

Semester: B.Pharm 7th

Year: 2020-2021

Name of the Teacher: Mr. Susankar Kushari & Mr. Suman Kumar

Tentative Dates	Syllabus Topics to be Covered	Learning outcome	Mode of Delivery	Assessment Tools (With tentative date of Assessment) for measuring Outcome
Batch D: Aug 11 B: Aug 13	To perform the assay of Paracetamol tablet by standard curve method	<ul style="list-style-type: none"> • Explain the principle and procedure behind the assay of Paracetamol by standard curve method 	<ul style="list-style-type: none"> • Demonstration of experiments • Lab manual 	<ul style="list-style-type: none"> • Laboratory Performance. • Viva-voce • Report analysis • Evaluation of practical record Assessment date: On the day of experiment
Batch D: Aug 18 B: Aug 20	To perform the assay of Paracetamol tablet by absorptivity method	<ul style="list-style-type: none"> • Explain the procedure for performing assay of Paracetamol tablet by absorptivity method 	<ul style="list-style-type: none"> • Demonstration of experiments • Lab manual 	<ul style="list-style-type: none"> • Laboratory Performance. • Viva-voce • Report analysis • Evaluation of practical record Assessment date: On the day of experiment
Batch D: Aug 25 B: Aug 27	To determine the absorption maxima and the effect of solvents on absorption maxima of given organic compound.	<ul style="list-style-type: none"> • Explain the principle and procedure of solvent effects on organic compound 	<ul style="list-style-type: none"> • Demonstration of experiments • Lab manual 	<ul style="list-style-type: none"> • Laboratory Performance. • Viva-voce • Report analysis • Evaluation of practical record Assessment date: On the day of experiment
Batch D: Oct 11 B: Oct 13	To determine the standard curve of	<ul style="list-style-type: none"> • Explain the procedure for performing the 	<ul style="list-style-type: none"> • Demonstration of experiments 	<ul style="list-style-type: none"> • Laboratory Performance. • Viva-voce

	riboflavin by using spectrofluorimeter .	standard curve of riboflavin by using spectrofluorimeter .	<ul style="list-style-type: none"> • Lab manual 	<ul style="list-style-type: none"> • Report analysis • Evaluation of practical record Assessment date: On the day of experiment
Batch D: Oct 18 B: Oct 20	Estimation of quinine sulphate by using spectrofluorimeter	<ul style="list-style-type: none"> • Explain the procedure and principle for the Estimation of quinine sulphate by using spectrofluorimeter 	<ul style="list-style-type: none"> • Demonstration of experiments • Lab manual 	<ul style="list-style-type: none"> • Laboratory Performance. • Viva-voce • Report analysis • Evaluation of practical record Assessment date: On the day of experiment
Batch D: Oct 25 B: Oct 27	To determine the effect of quenching on quinine sulphate	<ul style="list-style-type: none"> • Explain the principle of quenching and procedure involved. 	<ul style="list-style-type: none"> • Demonstration of experiments • Lab manual 	<ul style="list-style-type: none"> • Laboratory Performance. • Viva-voce • Report analysis • Evaluation of practical record Assessment date: On the day of experiment
Batch D: Nov 3 B: Nov 5	To perform the assay of Allopurinol tablets and to find the lambda maxima of the given solution using UV spectrophotometer.	<ul style="list-style-type: none"> • Explain the procedure for assay of allopurinol tablets. 	<ul style="list-style-type: none"> • Demonstration of experiments • Lab manual 	<ul style="list-style-type: none"> • Laboratory Performance. • Viva-voce • Report analysis • Evaluation of practical record Assessment date: On the day of experiment
Batch D: Nov 10 B: Nov 12	To interpret the IR spectra of given sample	<ul style="list-style-type: none"> • Explain the principle of IR spectroscopy and distribution of frequency regions of various functional groups. 	<ul style="list-style-type: none"> • Demonstration of experiments • Lab manual 	<ul style="list-style-type: none"> • Laboratory Performance. • Viva-voce • Report analysis • Evaluation of practical record Assessment date:

Batch D: Nov 17 B: Nov 19	To compare the IR spectra of marketed furosemide and pure furosemide	<ul style="list-style-type: none"> Explain the comparison of IR spectra of marketed furosemide and pure furosemide. 	<ul style="list-style-type: none"> Demonstration of experiments Lab manual 	On the day of experiment <ul style="list-style-type: none"> Laboratory Performance. Viva-voce Report analysis Evaluation of practical record Assessment date: On the day of experiment
Batch D: Nov 25 B: Nov 27	To perform the TLC of supplied sample and detect the spot in UV fluorescence cabinet chamber,	<ul style="list-style-type: none"> Explain the procedure for TLC 	<ul style="list-style-type: none"> Demonstration of experiments Lab manual 	Laboratory Performance. <ul style="list-style-type: none"> Viva-voce Report analysis Evaluation of practical record Assessment date: On the day of experiment
Batch D: Dec 3 B: Dec 5	Isolation of plant pigments by column chromatography	<ul style="list-style-type: none"> Explain the procedure for packing of column (wet packing technique) 	<ul style="list-style-type: none"> Demonstration of experiments Lab manual 	Laboratory Performance. <ul style="list-style-type: none"> Viva-voce Report analysis Evaluation of practical record Assessment date: On the day of experiment
Batch D: Dec 10 B: Dec 12	To demonstrate HPLC instrument	<ul style="list-style-type: none"> Explain the instrumentation, principle of HPLC 	<ul style="list-style-type: none"> Demonstration of experiments Lab manual 	Laboratory Performance. <ul style="list-style-type: none"> Viva-voce Report analysis Evaluation of practical record

Submitted by:

Sig:
Subject Teacher

Verified by:

Sig:
Module Co-ordinator

GIRIJANANDA CHOWDHURY INSTITUTE OF PHARMACEUTICAL SCIENCE

1st Sessional Examination

B.Pharm 7th Semester (October, 2020)

Subject: Instrumental Method of Analysis

Subject Code: BP701T

Full Marks:30

Time:1 hr

Q.I. Answer all the questions:

1×10=10

1. The source of light for visible radiation is.....

- a) Deuterium lamp
- b) Xenon arc lamp
- c) Tungsten halogen lamp
- d) All of the above

2. Which of the following technique is associated with molecular emission?

- a) UV-vis spectroscopy
- b) Fluorescence Spectroscopy
- c) IR spectroscopy
- d) X-ray diffraction

3. The fingerprint region is.....

- a) 8-50 μm
- b) 1-20 μm
- c) 50-100 μm
- d) 100-150 μm

4. The type of transition seen in Microwaves is.....

- a) Molecular vibration
- b) Inner shell electron
- c) Molecular rotation
- d) Outer shell electron

5. Bathochromic shift means.....

- a) Increase in wavelength
- b) Increase in intensity
- c) Decrease in wavelength
- d) Decrease in intensity

6. Hooke's law is associated with.....

- a) IR
- b) UV-Vis
- c) Fluorimetry
- d) NMR

7. The C=O stretch value in case of aromatic aldehyde ranges from..... cm^{-1}

- a) 2950-2850
- b) 3200-3400
- c) 1690-1710
- d) 3000-3050

8. According to Beer's law the intensity of incident light is directly proportional to.....of the sample.

- a) Concentration
- b) Turbidity
- c) Path length
- d) Dilution

9. The mid IR ranges from..... cm^{-1} .

- a) 12,500-4000
- b) 667-50
- c) 4000-667
- d) None of the above.

10. $n \rightarrow \sigma^*$ is shown by one of the following functional groups.

- a) -OH
- b) Benzene
- c) Alkanes
- d) -CHO

Q.II. Answer any two out of three:

2×5=10

1. Derive Beer-Lambert's law.
2. Proof that $A = 2 - \log \%T$.
3. Write a short note on Quenching.

Q.III. Answer any one of two:

1×10=10

1. Write a note on principle and detectors used in UV-Vis spectroscopy.
2. Discuss Jablonski's diagram.

B. Pharm 7th Semester End Term Examination 2020
Subject: Instrumental Methods of Analysis (BP701T)

Full Marks: 75

Time: 3 hrs

A. Answer all

1×20=20

1. The cathode used in Atomic Absorption Spectroscopy is.....
 - a) Sodium
 - b) Based on the element
 - c) Tungsten wire
 - d) Silver wire
2. Silica gel is stable at pH:
 - a) 1-5
 - b) 6-9
 - c) 2-7
 - d) None of these
3. The following principle applicable in UV is.....
 - a) Molecular vibration
 - b) Molecular rotation
 - c) Electronic transition
 - d) Inner shell transition
4. Thermocouple and Thermopile detectors are composed of.....
 - a) Bismuth & antimony
 - b) Wheatstone bridge
 - c) Oxides of Mn, Co & Ni
 - d) Twin dielectric flake
5. Which of the following is an example of strong basic anion exchange resin?
 - a) Dolomite
 - b) Phenol formaldehyde
 - c) Quarternary ammonium compounds
 - d) Sulphonated polystyrene
6. In Flame Photometry Barium emits.....colour under visualisation.
 - a) Brick red
 - b) Orange
 - c) Lime green
 - d) Violet
7. In TLC, fluorescent indicator is.....
 - a) Zinc stearate
 - b) Zinc sulphate
 - c) Zinc silicate
 - d) None of these

8. Electron capture detector used in GC is composed of.....
- a) Selenium
 - b) Oxides of Mn, Co & Ni
 - c) Germanium
 - d) All of the above
9. The diameter of analytical column used in HPLC is.....
- a) 2-4.5 mm
 - b) 50-60 mm
 - c) 0.25-0.35 mm
 - d) 5-6 mm
10. The common reagent for detecting barbiturates in TLC is.....
- a) Aniline phthalate
 - b) Mercuric nitrate
 - c) Diphenyl carbazone
 - d) Antimony trichloride
11. The following reagent is used in Gel electrophoresis for protein separation?
- a) Comassie blue
 - b) Methylene blue
 - c) Alizarin
 - d) Catechol-violet
12. The most popular thickness of layer in TLC is:
- a) 0.5 mm
 - b) 0.75 mm
 - c) 0.25 mm
 - d) 2 mm
13. In reverse phase chromatography mobile phase is.....
- a) Polar
 - b) Either a) or c)
 - c) Non polar
 - d) None of the above
14. Weakly acidic cation exchange resins are useful in pH range of:
- a) 1-14
 - b) 1-12
 - c) 5-14
 - d) 1-9
15.converts sample in mist or aerosol.
- a) Nebulizer
 - c) Detector

C. Answer any two

10×2=20

1. Define Chromophore and Auxochrome with suitable examples. Explain the different electronic transitions of UV Visible spectroscopy. With diagram explain the working of a double beam UV-Visible spectrophotometer. 2+4+4=10
2. With Jablonski diagram explain the principle of Fluorescence and Phosphorescence. Mention the factors affecting fluorescence. 4+6=10
3. Write the principle of HPLC. Write diagram explain different parts and working of an HPLC instrument. 3+7=10

B. Pharm 7th Semester End Term Examination 2020
Subject: Instrumental Methods of Analysis (BP701T)

Answers to the MCQ questions

- A. 1. b
2. c
3. c
4. a
5. c
6. c
7. c
8. b
9. a
10. c
11. a
12. c
13. a
14. c
15. a
16. c
17. b
18. a
19. a
20. c

GIRIJANANDA CHOWDHURY INSTITUTE OF PHARMACEUTICAL SCIENCE2nd Sessional ExaminationB.Pharm 7th Semester(December, 2020)

Subject: Instrumental Method of Analysis

Subject Code:BP701T

Full Marks:30

Time:1 hr

Q.L. Answer all the questions:

1×10=10

1. Laminar flow burner is also called as.....

- a) Nozzle mix burner c) Premixed burner
b) Turbulent burner d) Total Consumption burner

2. Which of the following is not an application of flame photometry?

- a) Assay of metformin c) To estimate Na, Mg & Ca
b) To detect metallic ions in sample d) To determine Ca & Mg in cement.

3. In flame emission photometer.....is used for qualitative analysis.

- a) Colour c) Both a) & b)
b) Intensity d) None of the above

4. Which of the following is an advantage of grating monochromator?

- a) Dispersion is non-overlapping c) Dispersion is overlapping
b) Dispersion is linear d) Dispersion is non-linear

5. At what pressure should be the gases sealed in a tube to be maintained in Hollow cathode lamp?

- a) 1-5 torr c) 5-20 torr
b) 20-50 torr d) 50-100 torr

6. Boltzman equation is associated with.....

- a) Flame photometry c) AAS
b) IEC d) NMR

7. In AASis used as an anode.

- a) Tungsten c) Selenium
b) Element to be investigated d) Germanium

ASSIGNMENT

TOPIC - GIVE THE METHODOLOGY AND DEVELOPING TECHNIQUES
OF PAPER CHROMATOGRAPHY

SUBJECT - INSTRUMENTAL METHODS OF ANALYSIS

SUBMITTED BY

ABHIRUP MUKERJEE

B. PHARM 7TH SEMESTER

SEC 'A'

ROLL NO : 04

Paper Chromatography

- Paper chromatography is processed as the separation of sample component carried out on paper.
- Cellulose filter paper is generally used as stationary phase.
- Here, separation of components occur due to partition mechanism. The solute molecules are distributed between two liquids.

Methodology of Paper Chromatography

- Chromatographic paper of appropriate size and shape is selected. A baseline of 2cm from lower edge is drawn.
- A drop of test solution is applied as a small spot on a paper and the spot is dried.
- Mobile phase is kept in the chamber.
- The paper is placed in the chamber such that the lower edge of the paper is dipped into a solvent (mobile phase).
- Care must be taken so that the base line should not be dipped in mobile phase.
- A lid is kept on top of the chamber.
- Solvent will raise through the paper by capillary action and it reaches the spot of the test solution.
- From this action, the various sample component are moved by solvent system at various speed.
- When solvent has raised to a suitable height, the paper is taken out of the chamber and dried.
- Various spots are visualized by suitable reagents.

- R_f values are calculated for each component of sample i.e., migration parameter by using the formula,

$$R_f = \frac{\text{Distance travelled by solute from base line}}{\text{Distance travelled by solvent from base line}}$$

Development Techniques

- Paper is flexible when compared to glass plate used in TLC.
- The different types of development techniques are as follows:-

1) Ascending Development

- When paper development is done by allowing the solvent to travel up the paper i.e. solvent flow against the gravity, it is known as ascending technique.
- Here, the mobile phase is kept in a container at the bottom of the chamber. The sample are applied a few centimeters from the bottom edge of the paper.

2) Descending type development

- When development of paper is done by allowing the solvent to travel down the paper, it is known as descending technique.
- This is carried out in a special chamber where the solvent ~~to travel~~ holder is at the top. The spot is kept at the top.
- The spot is kept at the top and solvent flows down the paper.
- The advantage of this type is that the development is faster.

3) Ascending - Descending Development

- It is a hybrid of both ascending and descending technique.
- only length of separation is increased, first ascending takes place followed by descending.

4) Circular / Radial Development

- In this type, a circular filter is used. Then the various material to be analysed are placed at the centre.
- After drying the spot, the paper is fixed horizontally on a petri-dish containing the solvent so that the tongue of the paper dips into the solvent.
- The paper is covered by means of petri-dish cover. The solvent rises through the tongue.
- when the solvent has moved through a sufficient distance, the components get separated.

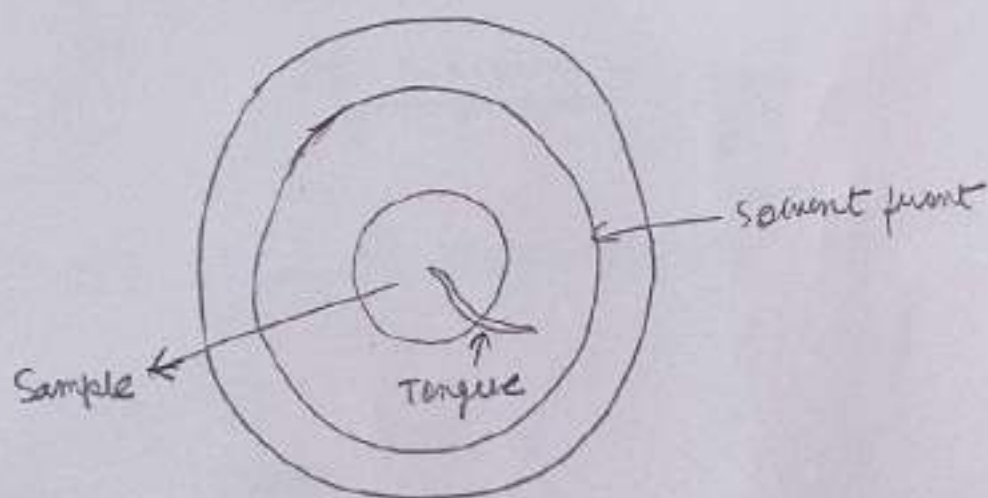


Fig: Radial paper chromatography

ASSIGNMENT

TOPIC:- Give the methodology and developing techniques of paper chromatography.

Subject:- Instrumental Methods of Analysis

Submitted by :

Abhishek Kumar Jha

B.Pharm 7th Semester

Sec - 'A'

Roll No. - 06.

Paper Chromatography

In paper chromatography, separation of sample component is carried out on paper. The cellulose filter paper is often used as stationary phase.

In this technique separation of sample components occurs due to partition mechanism. The solute molecules are distributed between two liquids.

Methodology of paper chromatography

- 1) Chromatographic paper of appropriate size and shape is selected. A base line of 2cm from lower edge is drawn.
- 2) A drop of test solution is applied as a small spot on a paper and the spot is dried.
- 3) Mobile phase is kept in the chamber.
- 4) The paper is placed in the chamber in such a way that lower edge of paper is dipped into a solvent.
- 5) Care must be taken so that the base line should not be dipped in mobile phase.

- 6) A lid is kept on top of the chamber.
- 7) Solvent will rise through the paper by capillary action and it reaches the spot by of the best solution.
- 8) From this action the various sample component are moved by the solvent system at various speed.
- 9) When solvent has raised to a suitable height, the paper is taken out of the chamber and dried.
- 10) Various spot are visualized by suitable reagents.
- 11) R_F values are calculated for each component of sample, i.e migration parameter by using the formula.

$$R_F = \frac{\text{Dist. travelled by solute from baseline}}{\text{Dist. travelled by solvent from base line}}$$

Development technique.

- Paper is flexible when compared to glass plate used in TLC.

The different types of development technique are:-

1) Ascending Development (go up)

When the development of the paper is done by allowing dips into the solvent. The paper is covered by means of petri

the solvent to travel up the paper. i.e. solvent flow against the gravity, it is known as ascending technique.

Here, mobile phase is kept in a container at the bottom of the chamber. The sample are applied on few centimeters from the edge of the paper.

2) Descending type (a downward slope)

When the development of the paper is done by allowing the solvent to travel down the paper, it is known as descending technique.

This is carried out in a special chamber where the solvent holder is at the top. Spot is kept at the top & the solvent flows down the paper.

3) Ascending - descending development

It is a hybrid of the above two technique. Only length of separation increased, first ascending takes place followed by descending.

4) Circular / Radial development

In this type, a circular filter is used, then the various material to be analysed are placed at the center. After drying the spot, the paper is fixed horizontally on a petri-dish containing the solvent so that the tongue of the paper dips into the solvent. The paper is covered by the means of a petri-dish.

dish cover. The solvent rises through the tongue. When solvent front has moved through a sufficient distance the component get separated.



ASSIGNMENT

TOPIC - Give the methodology & developing techniques of paper chromatography

SUBJECT - INSTRUMENTAL METHOD OF ANALYSIS

SUBMITTED BY - Abhishek Das

Roll - 5

B.Pharm 7th Sem

Sec A

Paper Chromatography

Paper chromatography is a technique used to separate phytochemical components. The cellulose filter is used as stationary phase. Separation occurs due to partition mechanism.

Methodology

- Chromatographic paper of appropriate size and shape is selected. A baseline of 2 cm from lower edge is drawn.
- A drop of test solution is applied as a small spot on a paper and the spot is dried.
- Mobile phase is kept in the chamber.
- The paper is placed in the chamber in such a way that lower edge of the paper is dipped into a solvent (mobile phase).
- Care must be taken so that the base line should not be dipped in mobile phase.
- A lid is kept on top of the chamber.
- Solvent will rise through the paper by capillary action and it reaches the spot of the test soln.
- From the action the various sample components are moved by solvent system at various speeds.
- When solvent has risen to a suitable height, the paper is taken out of the chamber and dried.
- Various spots are visualized by suitable reagents.
- R_f values are calculated for each component of sample by -
$$R_f = \frac{\text{Dist. travelled by solute from base line}}{\text{Dist. travelled by solvent from base line}}$$

Development Techniques

- 1) Ascending Development :- When the development of the paper is done by allowing the solvent to travel up the paper. In here the mobile phase is kept in a container at the bottom of the chamber. The sample is applied few centimeters from bottom edge.
- 2) Descending type :- When the development of the paper is done by allowing the solvent to travel down the paper. Carried out in a spherical chamber where the solvent holder is at top. The spot is kept at the top & solvent flows down the paper.
- 3) Ascending - Descending Development :- It is a hybrid of the 2 above techniques. Only length of separation increased, first ascending followed by descending.
- 4) Radial Development :- Here, a circular filter is used. The materials to be analysed are placed at centre. After drying the spot, paper is fixed horizontally on a petri dish containing the solvent so that the tongue of the paper dips into the solvent. The paper is covered by means of petri dish cover. The solvent rises through the tongue. When solvent front has moved through a sufficient distance, the component gets separated.

ASSIGNMENT

ON

INSTRUMENTAL METHOD
OF ANALYSIS

Topic:- Give the methodology and developing techniques
of paper chromatography

Name- Anurag Krishna Murthy Baruah

Roll no - 11

B. Pharm, 7th Sem

Paper Chromatography:-

In paper chromatography, separation of sample component is carried out on paper. The cellulose filter paper is often used as stationary phase.

In this technique separation of sample components occurs due to partition mechanism. The solute molecules are distributed between two liquids.

Methodology of Paper Chromatography:

- 1) Chromatographic paper of appropriate size and shape is selected. A baseline of 2 cm from lower edge is drawn.
- 2) A drop of test-solution is applied as a small spot on a paper and the spot on a paper and the spot is dried.
- 3) Mobile phase is kept in the chamber.
- 4) The paper is placed in the chamber in such a way that lower edge of the paper is dipped into a solvent (mobile phase).
- 5) Care must be taken so that the base line should not dip in mobile phase.
- 6) A lid is kept on top of the chamber.
- 7) Solvent will rise through the paper by capillary action and it reaches the spot of the test solution.

7) From this action the various sample components are moved by solvent system at various speed.

8) When solvent has raised to a suitable height, the paper is taken out of the chamber and dried.

9) Various spot are visualized by suitable height, the paper is taken out of the chamber and dried.

11) R_f values are calculated for each component of sample i.e., migration parameter by using the formula;

$$R_f = \frac{\text{Dist. travelled by solute from base line}}{\text{Dist. travelled by solvent from base line}}$$

Development technique

• Paper is flexible when compared to glass plate used in TLC.

The different types of development technique are -

1) Ascending Development (going)

When the development of the paper is done by allowing the solvent to travel up the paper i.e., solvent flow against the gravity.

Here, the mobile phase is kept in a container at the bottom of the chamber. The

sample are applied a few centimeters from the bottom edge of the paper.

2) Descending type (downward slope) :-

When the development of the paper is done by allowing the solvent to travel down the paper, it is known as descending type.

Special chamber is used where solvent holder is at the top. The spot is kept at the top and the solvent flows downwards.

3) Ascending - descending development :-

It is hybrid of the above two techniques. Only length of separation increases, first ascending takes place followed by descending.

4) Circular / Radial development :-

In this type, a circular filter is used. When one or more materials to be analysed are placed in the center. After drying the spot, the paper is fixed horizontally on a petri dish containing the solvent so that the tongue of the paper dips into the solvent. The solvent rises through the tongue. When solvent front has moved through the tongue. When solvent front has moved through a sufficient distance the component is separated.

**GIRIJANANDA CHOWDHURY
INSTITUTE OF
PHARMACEUTICAL SCIENCE**



Assignment on Instrumental Method Of Analysis

Topic: Give the methodology and developing techniques of Paper Chromatography

Submitted to-

Mr. Susankar Kushari

Faculty, GIPS

Submitted by-

Name- Bhabana Kalita

Class- B Pharm 7th, Sem (A)

Roll no- 190510011017

Paper Chromatography -

It is a technique in which separation of an unknown substance is mainly carried out by the flow of solvents on the specially designed chromatographic filter paper.

• Methodology -

In methodology of paper chromatography practical requirements as follows -

- 1) Paper as stationary phase.
- 2) Preparation of paper.
- 3) Preparation of sample.
- 4) Application of the sample.
- 5) Mobile phase as developing solvent.
- 6) Chromatographic chamber.
- 7) Development of chromatogram.
- 8) Drying of chromatogram.
- 9) Location of spot.
- 10) Quantitative analysis.
- 11) Elution.

1) Paper as stationary phase -

- For paper chromatography Whatman filter paper is used.
- This paper contains α -cellulose 98-99%, β -cellulose 0.3-1%, ether soluble matter 0.015-0.02%, mineral content 0.01-0.07%, besides this ammonia and lipophilic substance are also present.
- Choice of filter paper depends upon thickness, flow rate, purity and technique.

2) Preparation of papers -

- As grade and type of paper decided for separation, it is cut in desired size and shape.
- After deciding the direction of run on paper, origin line marked on paper.

3) Preparation of sample -

- It is important to choose proper solvent for making solution.
- Generally in mobile phase sample solution is prepared or in volatile solvents.

4) Application of the sample -

- Prepared sample applied through capillary or micropipette on origin line marked on paper.
- Very low concentration is used to avoid large zone.
- Generally size of spot should be as small as possible.

5) Mobile phase as developing solvent -

- The selection of mobile phase depends mainly on nature of sample to be separated.
- Factors should be considered at the time of selection of mobile phase like viscosity, polarity.

6) Chromatographic chambers -

- Glass chromatographic chamber is preferred for paper chromatography.
- Length of chamber depending upon length and breadth of paper and type of development.

7) Development of chromatogram -

- At the time of development of paper, paper dipped in solvent in such a way that the spot will not dip completely into the solvent.

- The solvent will run over paper by capillary action.
- After complete development paper is carefully taken out of chamber.

8) Drying of chromatogram -

- Simple hair dryer is a convenient to dry chromatograms.
- They also dried by cold or hot air depending on volatility of solvents.

9) Location of spot -

- After drying of chromatogram next step is to detect developed spots of separated samples.
- If substance are coloured easily detected visually, but for detecting colourless spots various methods are use -

(a) Physical method -

E.g - Iodine chamber, UV chamber for fluorescent compound.

(b) Chemical method -

E.g - Amino acids - Ninhydrin in acetone.
Alkaloids - Dragendorff's reagent.

10) Methods of quantitative analysis -

- Measurement of spot length.
- Planimetric measurement
- Method of Counting squares.
- Weighing the Excised Spot.
- Method of Visual Comparison.

11) Elution -

- The section of the developed filter paper is cut out and eluted with an appropriate solvent such as ethanol or chloroform.
- The elute is then made up to volume and measured in a spectrophotometer or colorimeter and compared with a standard of known concentration.

• Development Techniques -

In paper chromatography various development techniques were employed to increase ease and efficiency of operation. They are -

- 1) Ascending development.
- 2) Descending development.
- 3) Ascending - Descending development.
- 4) Circular / Radial / Disk development
- 5) Two dimensional development.

1) Ascending development -

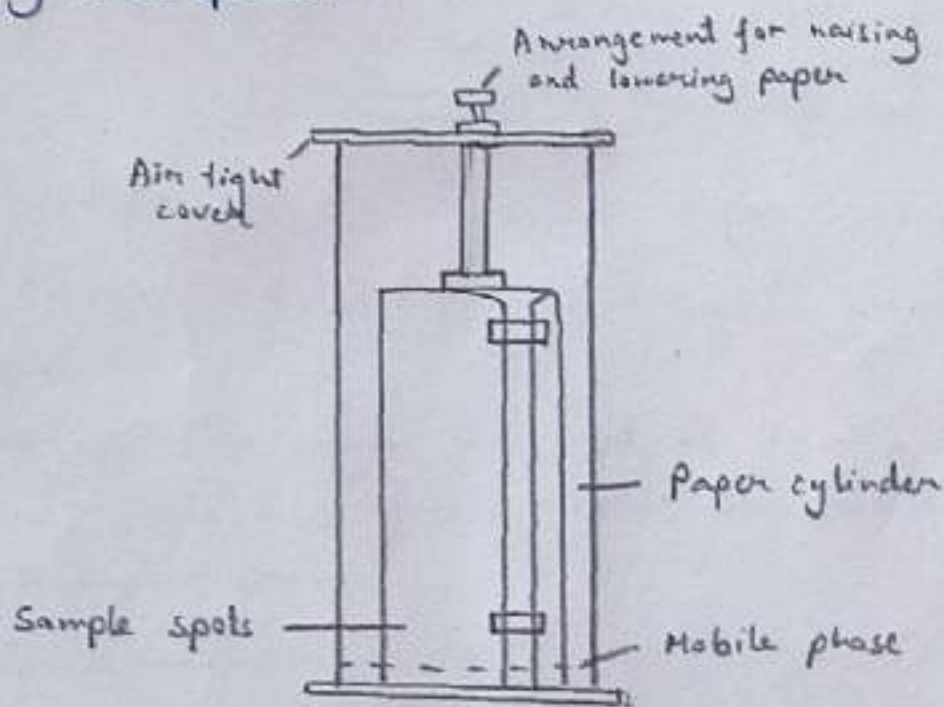


Fig: Apparatus for ascending paper chromatography

- In this technique mobile phase flows against gravity means upward direction of chromatographic paper.
- The origin line made at the bottom of the paper.
- Samples drop put on origin line and allow to air dry.
- Chromatographic chamber filled with mobile phase and saturates it with vapours of mobile phase.
- Prepared paper hangs in chromatographic chamber with thread on

held together by staples on plastic clips.

2) Descending development -

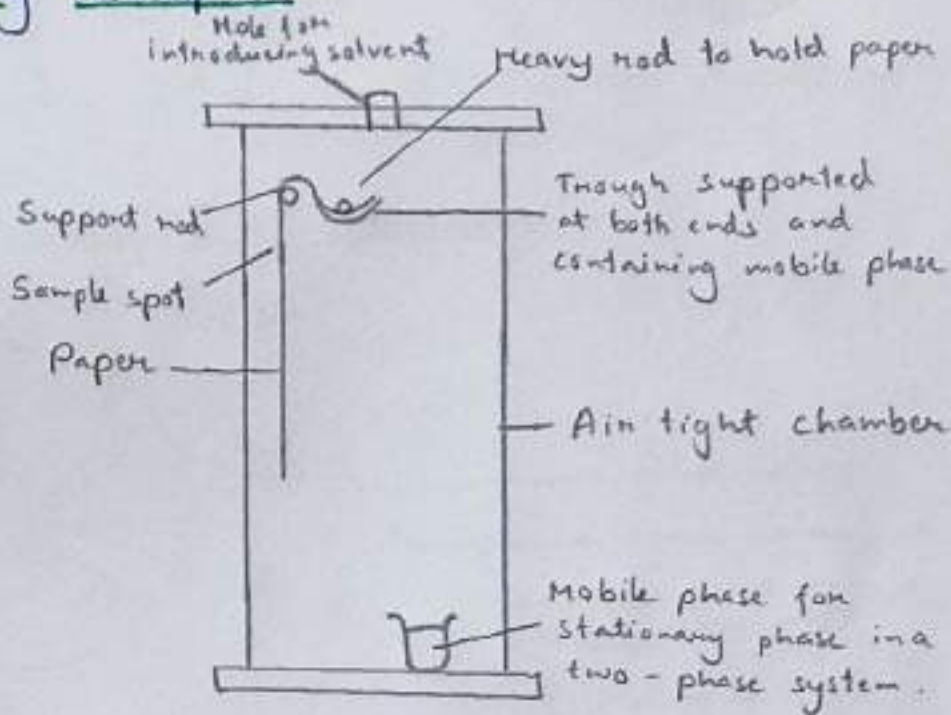


Fig: Apparatus for descending Paper Chromatography.

- This is carried out in special chamber where solvent held at the top.
- The sample spot on origin line kept at the top of paper and mobile phase flows with gravity means downward direction of chromatographic paper.
- Rate of development is faster.

3) Ascending - Descending development -

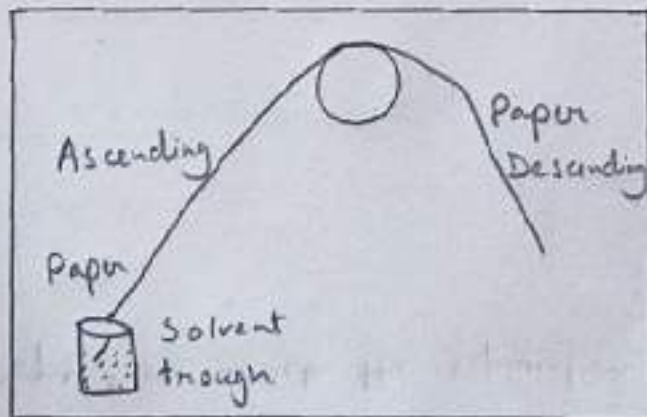


Fig: Ascending - Descending development

- This is a combination of ascending and descending type.
- The length of paper is increased and length of separation also increased by using combination of techniques.
- First ascending takes place followed by descending development.

4) Circular / Radial / Disk development -

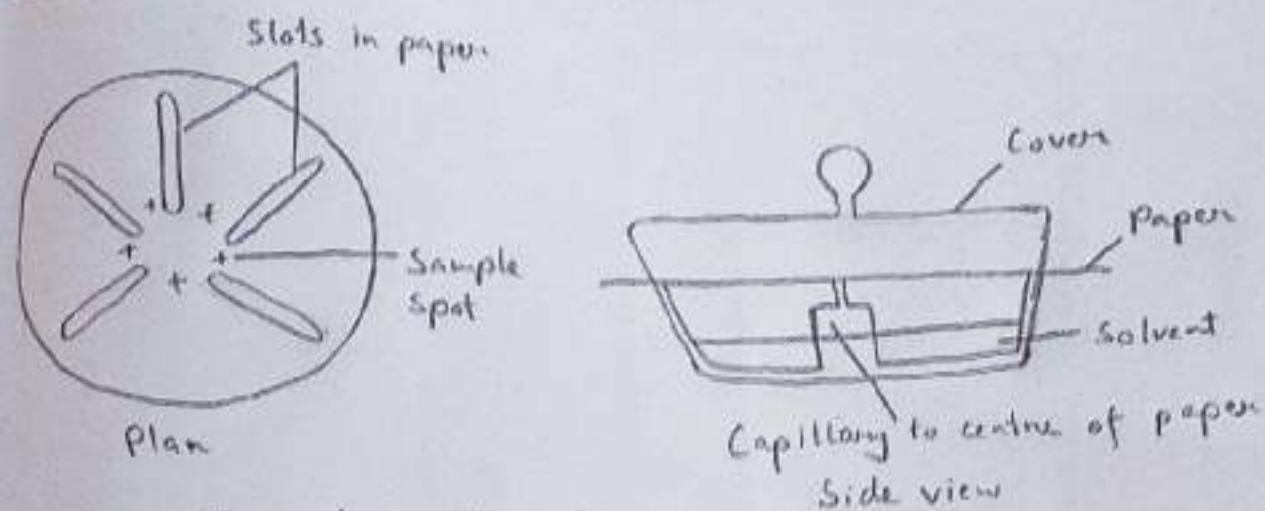


Fig: Apparatus for circular Paper Chromatography

- In this technique a circular chromatographic paper is used.
- This paper has a wick cut from centre of the paper, sample spot is applied on upper end of wick.
- The paper is then placed on edge of circular disk with wick dipping into the solvent at the bottom of the disk.

5) Two-dimensional developments -

- It is specially suitable when R_f values are very close or nearly the same.
- In this technique square or rectangular chromatographic paper is used.
- The sample is applied to one of the corners.

GIRIJANANDA CHOWDHURY UNIVERSITY

ASSIGNMENT ON INSTRUMENTAL METHOD OF ANALYSIS

Topic: - Methodology and Development
Techniques of paper chromatography.

Submitted to:

- Sir. Sushankar Kulkarni
- Assistant Professor
GCU
- Department of
Pharmaceutical
Chemistry

Submitted by:

- ~ Evaline Myllimngap
- ~ Roll-no : 190510011029
- ~ B.Pharm 7th Semester
Section - A
- ~ Date: 5/12/22

Q. Give the methodology and developing techniques of paper chromatography.

INTRODUCTION:- Paper chromatography was first introduced by the German scientist, Christian Friedrich Schonbein in 1865.

- It is a type of a planar chromatography. It is the simplest and widely used type of chromatography procedure which runs on a specialized paper.

Principle - This technique is a type of partition chromatography in which the substances are distributed between two liquids, i.e. one is the stationary liquid (usually water) which is held in the fibres of the paper and called the stationary phase; the other is the moving liquid or developing solvent and called the moving phase. The components of the mixtures to be separated migrate at different rates and appear as spots at different points on the paper.

- Originally paper chromatography was used to separate mixture of organic substances such as dyes and amino acids only. But now this method has been perfected to separate cations and anions of inorganic substances as well.

- In this technique, a drop of the test solution is applied as a small spot on a filter paper and the spot is dried. The paper is kept in a closed chamber and the edge of the filter paper is dipped into a solvent called developing solvent. As soon as the filter paper gets the liquid through its capillary axis and when it reaches the spot of the test solution, the various substances are moved by solvent system at various speeds. When the solvent has moved

these cations to a suitable height (15-18cm) the is dried and various spots are visualised by suitable reagents called visualising reagents. The movements of substances relative to the solvent is expressed in terms of R_f values i.e. migration parameters.

Types of Paper Chromatography

(a) Descending chromatography: When the development of the paper is done by allowing the solvent to travel down the paper, it is known as descending technique. The apparatus consists of a well-glass tank of suitable size and shape which is provided with a trough for the mobile phase in the upper portion. The paper with the sample is inserted with the upper end in the trough containing the mobile phase, the jar itself having equilibrated with the mobile phase prior to development.

(b) Ascending chromatography: - When the development of the paper is done by allowing the solvent to travel up the paper, it is known as ascending technique. In ascending chromatography, the mobile phase is in a suitable container at the bottom of the chamber the chamber itself. The samples are applied a few centimeters from the bottom edge of the paper suspended a book.

(c) Ascending - Descending chromatography: - It is the part of the above two techniques. In this technique a glass vessel allowing the ascending development to occur into the descending after crossing the glass.

(d) Radial Paper Chromatography: - This is also known as circular paper chromatography. In this technique a circular filter paper is used and the various material to be analysed are placed at the centre.

(e) Two-dimensional chromatography: In this, a square or rectangular paper is used. The sample is applied to one of the corners. The second development is performed at right angle to the direction of the first run. This type of chromatography can be carried out with identical solvent systems in both the directions or by two solvent systems.

METHODOLOGY

(a) Choice of the Proper Chromatographic Technique: -
- The first job is to select the mode of paper chromatographic technique i.e. ascending, descending, ascending-descending, radial or two-dimensional technique.
- The choice of technique depends upon the nature of the substances to be separated.

(b) Choice of the filter paper: -
- The filter paper plays an important role in the success of paper chromatography. The choice of paper depends on the type of problem under investigation. Whatmann filter papers are commonly used.

(c) Proper Developing Solvent: The best possible developing solvent is generally selected for the separation of substances under examination. The choice of this depends upon the simple fact that R_f values should be different of different constituents present in a mixture.

(d) Preparation of samples:- It is not possible to have any standard procedure for preparation of samples because this problem revolves around several factors of the given samples, especially the presence of components like fats, salts, proteins etc.

(e) Spotting: For ascending technique, a strip of Whatman filter paper of suitable size (25cm x 7cm) is used. A horizontal line is drawn on the paper by a lead pencil. This line is known as origin line. On the origin line, cross marks are made with a pencil in such a way that each cross is at least 2cm away from each other.

By the help of a graduated micro pipette test solutions are applied on cross marks and spots are dried cautiously by a stream of hot cold air.

(f) Drying the chromatography:- The wet chromatograms after development are dried in special cabinets which are being heated electrically with temperature controls.

(g) Visualisation:-

Visualisation of the spots can be done in

ways:

- (i) Either by chemical means or
- (ii) By using physical method.

(i) Chemical Detection: Chemical treatment can develop the colour of colourless spots on the paper. The reagents used for visualising the spots are known as chromogenic reagents and visualising reagents.

(ii) Physical Methods: Some colourless spots, when held under a UV lamp, fluoresce and reveal their existence. When the substance is coloured, the spots can be observed either by reflected or transmitted light.

(iii) Calculation of R_f values: The distance of chromatographed species is noted from its centre to the origin line. The distance of solvent front from the origin line gives the R_f values.

ASSIGNMENT ON INSTRUMENTAL METHOD OF ANALYSIS

Topic - Methodology of AND Development techniques of paper chromatography

Submitted to:-

- Mr. Sudeekar Kushari (Sir).
- Assistant professor of GCU.
- Department of pharmaceutical chemistry.

Submitted By:-

- Name: Koyal Sarkar
- Roll NO: 190510011041
- Semester: B. Pharm, 7th semester
- Branch: B. pharmacy.
- Date: 5-12-22
- University: Girijan and a Chowdhury University (GCU).

Give the Methodology and developing techniques of paper chromatography.

Introduction: of paper chromatography:-

→ paper chromatography was first introduced by German scientist Christian Friedrich Schönbein (1865). It is a partition-chromatography and liquid-liquid chromatography.

→ paper chromatography is considered to be the simplest and most widely used of the chromatographic techniques because of its applicability to isolation, identification and quantitative determination of organic and inorganic compounds.

→ paper chromatography is an analytical method used to separate colored chemicals or substances.

Types of paper chromatography - These are 2 types of paper chromatography, they are:-

- (1) Paper adsorption chromatography:- paper impregnated with silica or alumina acts as adsorbent (stationary phase) and solvent as mobile phase.
- (2) Paper partition chromatography:- Moisture/coater present in the pores of cellulose fibres present in filter paper acts as stationary phase and another mobile phase is used as solvent.

Methodology of paper chromatography:-

→ If the substance are colored they are visually detected easily.

→ But for colorless substance, physical and chemical methods are used to detect the spot.

(a) Non-specific methods (physical methods)

→ Eg:- Iodine-chamber method.

→ UV chamber for fluorescent compounds - at 254 or at 365nm.

(b) Specific Methods (Chemical methods) or specific method - examples:-

- Ferric chloride
- Nitroprusside in acetone
- Dragendorff's reagent
- 3,5-dinitrobenzoic acid

- phenolic comp. and tannins
- Amino acids
- Alkaloids
- cardiac glycosides

(c) Direct Measurement Method:-

(i) Comparison of visible spots:-

- A rough quantitative measurement
- Component in a mixture can be carried out by comparing the intensity and size of the spot with a standard substance.

(ii) Photo densitometry:-

- This method is used with the chromogram of colored compound, instrument measures quantitatively the density of spots.

(iii) Fluorimetry:-

- The compound to be fluorescent or convertible into fluorescent derivatives.

(iv) Radiotracer method:-

- The compound containing radioactive element is labeled and treated with local reagent, using Geiger Muller counter.

(v) Polarographic and Conductometric method:-

- used to measure the amount of material in the spot.

Indirect Measurement Method:-

- In this technique, the spots are cut into portions and eluted with solvent, the solutions can be analyzed by any technique of analysis like spectrophotometry, electrochemical methods, etc.

The various steps involved in paper chromatography are:-

(1) Selection of stationary phase (filter paper):-

- The stationary phase used in paper chromatography is water which get absorbed on a special type of paper made up of 98-99% cellulose fibres (α-cellulose). Other constituents (1%) include β-cellulose, pentosan, ash, emulsion, nitrogen etc.

- cellulose fibres is surrounded by a film of moisture, moisture or water molecules constitute about 22% by weight of paper.

- various types of whatmann chromatography papers are available, choice of whatmann paper depends on type of separation.

- coarse and faster paper (whatmann 21RT). It is used when substance to be separated have sufficiently wide R_F.

- Slow paper (whatmann 20). It is used for better resolution of substance with close R_F values.

- Heavy paper (whatmann 3mm).

- It is generally used for preparative purposes.

* The choice of filter paper depends upon:-

- (a) Type of analyte → qualitative or quantitative.
- (b) Reason for analysis → analytical or preparative.
- (c) Nature of the sample → hydrophilic, hydrophobic, neutral.

* After selecting or appropriate filter paper, cut into required shape (square, rectangular or circular) base line is drawn with pen about 2-2.5cm above the edge of paper and sample is applied over this line.

(2) Selection of solvent system (Mobile phase).

→ The mobile phase is generally a mixture of one or more polar organic solvents or salts and water (single phase solvent preferred).

→ Solvent system is taken in a clean & dry closed vessel preferably, a glass tall cylindrical or rectangular shape closed & saturate.

→ The choice of solvent system depends on the following criteria:-

- Solvent system should be inert. It should not react with paper and sample.
- The ratio of solvent system should change with time.
- It should be of suitable viscosity, surface tension and density.

Sample to be analyzed

Amino acids

Sugars

Hallide
(F, Cl, Br, I)

Mobile phase

n-butanol: Ac
water

(4:1:5)

ethyl acetate:
water

(2:1:2)

pyridine: water

(9:1)

(3) Preparation and Application of sample -

→ There is no standard procedure for preparation of sample if sample is solid then dissolved in suitable volatile solvent.

→ About 2-10 μ l of sample is applied at spots.

→ By help of graduated micropipette sample are applied spots are dried by a stream of hot or cold air.

(4) Development of chromatogram -

Suitable solvent system is placed in development tank.

covered it with a lid to saturate environment of tank.

The spotted paper is placed into the tank with its spotted end dipping into solvent (spot is kept slightly above the level of solvent) (paper should not touch the sides of tank).

The solvent now travel upon paper due to capillary action.

Solvent dissolves the constituents of sample which travel upwards according to their individual partition coefficients.

When solvent travels about 3/4th of the paper development is stopped by removing paper. The developed paper is known as chromatogram.

The chromatogram is immediately marked with pencil at distance travelled by solvent (known as solvent front). The paper is allowed to dry in open air or heated chamber.



(5) Visualization of chromatogram:

(a) Chemical Detection: - Chemical treatment develop colour of coloured spots on paper. The reagents used for visualizing are known as chromatogenic reagents or visualising reagents or locating agents.

- Reagent
- Ninhydrin solution
 - Aniline phthalate
 - Iodine vapour
 - Dragendorff's reagent
 - Bromo-cresol green

- Mixture
- Amino acids.
 - Sugars
 - organic bases
 - Alkaloids.
 - Acids.

(b) Physical Methods - Some coloured spots when placed under UV lamp, fluoresce, reveal their existence.

(6) Calculation of Migration parameters:

$$R_f = \frac{\text{Distance travelled by solute from origin}}{\text{Distance travelled by solvent from origin}}$$

$$R_x = \frac{\text{Distance travelled by the substance from origin}}{\text{Distance travelled by standard substance from origin}}$$

$$R_m = \log \left(\frac{1}{R_f} - 1 \right)$$

Development Technique -

- paper is flexible when compared to glass plate used in TLC. Reversal types of development are possible which increases the ease of operation.
- The paper is dipped in solvent in such a manner that the spots will not dip completely into the solvent.
- The solvent will rise up and it is allowed to run 2/3rd of paper height for better and efficient result.
- It takes from several minutes to several hours.

Different type of Development techniques -

- (1) Ascending Development
- (2) Descending Development
- (3) Ascending-Descending Development
- (4) Circular/Radial Development
- (5) Two dimensional Development

And, here, we have described the different type of development techniques.

ASSIGNMENT ON INSTRUMENTAL METHOD OF ANALYSIS

Topic :- Methodology And Development techniques of paper chromatography

Submitted to :-

- Mr. Susankar Kushari (sir)
- Assistant professor of MCO
- Department of pharmaceutical chemistry

submitted by :-

- Name :- Madhusmita Paw
- Roll no :- 190510011049
- Semester :- B. Pharm, 7th Sem
- Branch :- B. Pharmacy
- Date :- 5-12-22

Give the methodology & developing techniques of Paper Chromatography.

The various steps involved in paper chromatography are:

(1) Selection of stationary phase (filter paper):

* The stationary phase used in paper chromatography is water which get absorbed on a special type of paper made up of 99.99% cellulose fibers (α -cellulose) other constituents (about 1%) includes β -cellulose, pentosans, ash, ammonia, nitrogen etc.

* Cellulose fibers is surrounded by a film of moisture. moisture or water molecules constitute about 22% by weight of paper.

* Various types of Whatmann chromatography papers are available. Choice of Whatmann paper depends on type of separation.

* Coarser and faster paper (Whatmann 51ET)
It is used when substances to be separated have sufficiently wide apart R_f

* Slow papers (Whatmann 20)
It is used for better resolution of substances with close R_f values.

* Heavy paper (Whatmann 3mm) \Rightarrow
It is generally used for preparative purposes.

* The choice of filter paper depends upon-

- (a) Type of analysis \rightarrow qualitative or quantitative
- (b) Reason for analysis \rightarrow analytical or preparative.
- (c) Nature of the sample \rightarrow hydrophilic, hydrophobic, neutral or

Charged to suspend: enigma with vein
 * After selecting an appropriate filter paper it is cut into required shape (square, rectangular or circular) Base line is drawn with pencil about 2-3cm above the edge of paper and sample is applied over this line.

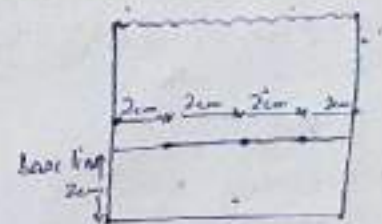
(2) Selection of Solvent System (Mobile phase):-
 * The mobile phase is generally a mixture of one or more polar organic solvents, buffer or salt and water (Single phase solvent are preferred)
 * Solvent system is taken in a clean and dry closed vessel preferably a glass tank of cylindrical or rectangular shape closed to saturate.

* The choice of Solvent system depends upon the following criteria:

- (a) Solvent system should be inert. It should not react with paper and sample.
- (b) The ratio of solvent system should not change with time.
- (c) It should be of suitable viscosity, surface tension and density.

Sample to be analyzed	Mobile phase used
Amino Acids	n-butanol : Acetic acid : water (4 : 1 : 5)
Sugars	Ethyl acetate : pyridine : water (2 : 7 : 2)
Halides (F ⁻ , Cl ⁻ , Br ⁻ , I ⁻)	Pyridine : water (9 : 1)

3) Preparation and Application of sample:-
 * There is no standard procedure for preparation of sample if sample is solid then dissolved in suitable volatile solvent.
 * About 2-10 ul of sample is applied as spots.
 * By help of graduated micro pipette sample are applied. Spots are dried by a stream of hot or cold air.



4) Development of Chromatogram:-
 Suitable Solvent system is placed in development tank. Covered it with a lid to saturate environment of tank.

The spotted paper is placed into the tank with its spotted end dipping into solvent (Spots kept slightly above the level of solvent) (Paper should not touch the sides of tank).
 The solvent now travel up on paper due to capillary action. Solvent dissolves the constituents of sample which travels upwards according to their individual partition coefficients.
 (The system should not be disturbed once development starts)
 When solvent travels about 3/4th of the paper development is stopped by removing paper. The developed paper is known as Chromatogram.

The chromatogram is immediately marked with zones at distance travelled by solvent (known as solvent front). The paper is allowed to dry in open air or heated chamber.

2) Visualizations of chromatogram:-

a) Chemical Detection: chemical treatment can develop colour of colourless spots on paper. The reagents used for visualising spots are known as Chromatogenic reagents or visualising reagents or locating agents.

Reagent	Mixture
Ninhydrin solution	Amino acids
Aniline phthalate	Sugars
Iodine vapour	Organic bases
Dragendorff's reagent	Alkaloids
Bromocresol green	Acids

b) Physical Methods:- Some colourless spots when placed under UV lamp, fluorescence reveal their existence.

(6) Calculation of Migration Parameter:-

$$R_f = \frac{\text{Distance travelled by solute from origin line}}{\text{Distance travelled by solvent from origin line}}$$

$$R_r = \frac{\text{Distance travelled by the substance from origin line}}{\text{Distance travelled by standard substance from origin line}}$$

$$R_m = \log \left(\frac{1}{R_f} - 1 \right)$$

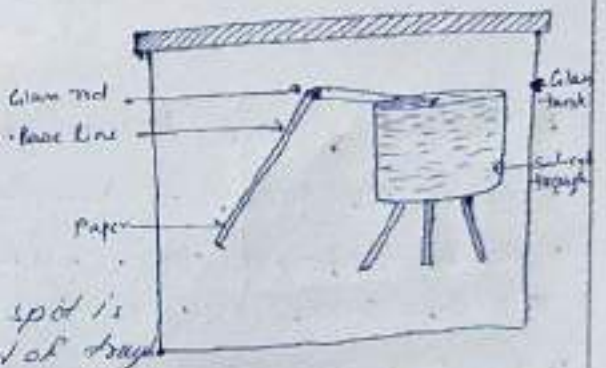
Development Techniques:-

* Depending upon the nature of components to be separated any one of the following may be opted:-

(1) Descending Chromatography:-

When development of the paper is done by allowing the solvent to travel down the paper, it is known as descending technique.

* The container consist of a well sealed glass tank of suitable size. It provided with trough for the mobile phase.



* The paper with sample spot is inserted with the upper end of trough containing mobile phase.

* The advantage of this is that the development can be continued indefinitely even though solvent run off at the other end of paper.

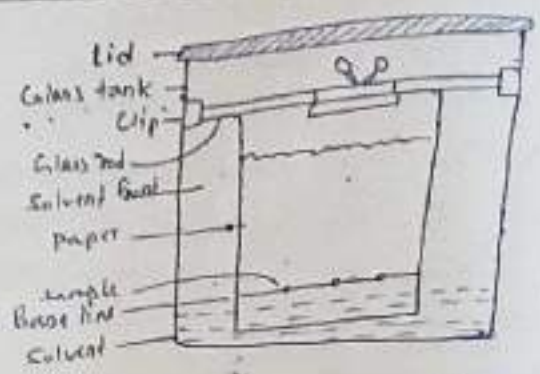
(2) Ascending Chromatography:-

When development of the paper is done by allowing the solvent to travel up the paper it is known as ascending technique.

* In ascending chromatography, the mobile phase is placed in a suitable container at the bottom of the chamber.

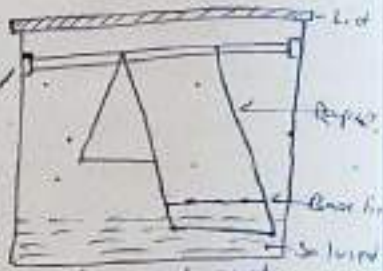
The samples are applied a few cm. from bottom edge of the suspended paper.

* The ascending technique is preferred if R_f values of various constituents are almost same.



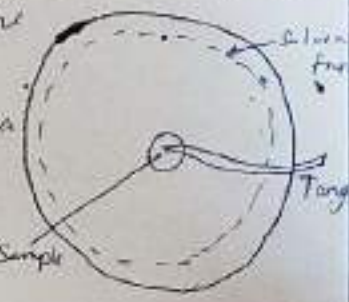
3) Ascending and Descending Chromatography:-
 * It is the hybrid of ascending and descending techniques.
 * In this technique upper part of ascending chromatography can be folded over a glass rod.

* allowing ascending development to change into the descending development after crossing glass rod.



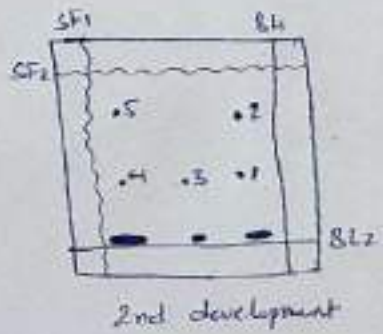
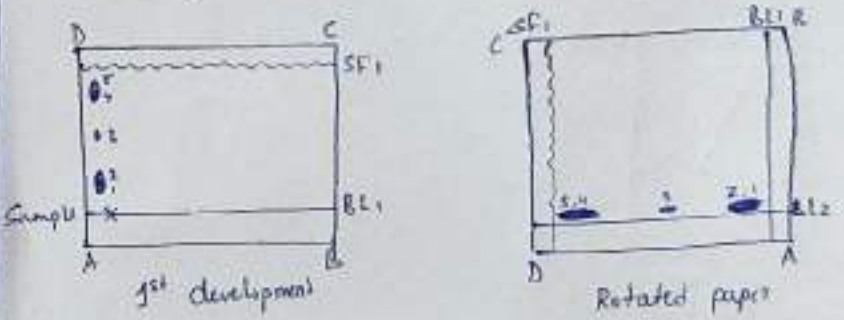
4) Radial Paper Chromatography:-

* It is also known as circular paper chromatography.
 * It is used for radial development.
 * In this technique a circular filter paper is used.
 * Various materials to be analysed are placed at the center and stirred.
 * The paper is fixed horizontally in a petri dish.
 * The paper is placed such that the lower end of the wick at the tongue touches the solvent.
 * The petri dish is then covered and solvent ascends the wick



by capillary action.
 - The solvent may also be applied continuously over the dried spot with the capillary tube.
 - When solvent front has moved a sufficient large distance components get separated in the form of concentric circular zones.

5) Two dimensional Chromatography:-
 * In this - a square or rectangular paper is used.
 * Sample is applied to one of the corners.
 * Second development is performed at right angle to the direction of the first run.
 * This type of chromatography can be carried out with identical solvent system in both directions or by two solvent systems.



GIRIJANANDA CHOWDHURY INSTITUTE OF PHARMACEUTICAL SCIENCE

PROGRAM :	BACHELOR OF PHARMACY (B. PHARM) (New Curriculum - 2017)		
UNIVERSITY :	Assam Science & Technology University		
NAME OF THE SUBJECT :	Theory:	Instrumental Method of Analysis	
SUBJECT CODE :	Theory:	BP701T	
SESSION :	Aug - December 2020		
NAME OF THE TEACHER(S) :	Mr. Susankar Kishari & Mr. Suman Kumar		

COURSE OUTCOME ASSESSMENT SHEET

ROLL NO	Theory				Theory								GRADE
	NO. OF CLASSES	NO. OF CLASSES ATTENDED	ATT. PERCENTAGE		SESSIONAL (20)				CUMULATIVE EVALUATION (20)				
					1ST SES	2ND SES	3RD SES	AVG	CE1	CE2	CE3	AVG	
170510011001	40	35	88		24	26	0	25	18	17	18	17.7	B
170510011002	40	37	92		26	27	23	27	19	19	20	19.3	A
170510011003	40	35	88		21	18	0	20	18	18	17	18.0	A
170510011004	40	37	92		22	19	18	21	17	18	17	17.3	C
170510011005	40	39	98		26	28	26	27	17	16	17	16.7	A
170510011006	40	38	95		23	25	19	24	18	18	17	17.7	D
170510011008	40	36	90		27	27	23	27	19	18	20	19.0	A
170510011009	40	34	85		21	22	23	23	18	17	18	17.7	B
170510011010	40	37	92		24	26	21	25	18	18	17	17.7	A
170510011011	40	36	90		20	14	16	22	16	16	17	16.3	B
170510011012	40	35	88		25	27	22	26	19	20	19	19.3	B
170510011013	40	36	90		26	14	20	25	20	20	19	19.7	B
170510011014	40	38	95		21	25	13	23	16	17	17	16.7	D
170510011015	40	35	88		23	23	14	23	18	18	19	18.3	B
170510011016	40	37	92		25	26	20	26	17	19	18	18.0	B
170510011017	40	36	90		24	25	0	25	19	19	19	19.0	A
170510011018	40	36	90		22	23	0	23	19	18	19	19.0	B
170510011019	40	38	95		24	25	15	25	18	19	18	18.3	C
170510011020	40	37	92		25	25	16	25	18	17	16	17.0	B
170510011021	40	38	95		22	21	9	22	17	18	17	17.3	B
170510011022	40	37	92		22	21	9	22	15	16	15	15.3	C
170510011023	40	35	88		24	21	11	23	20	19	19	19.3	C

Susankar Kishari
SIGNATURE OF THE TEACHER

ROLL NO	Theory			Theory								O/E
	NO. OF CLASSES HELD	NO. OF CLASSES ATTENDED	ATTENDANCE	SESSIONAL (20)				CONTINUOUS EVALUATION (20)				
				1ST SEM	2ND SEM	3RD SEM	AVG	CE 1	CE 2	CE 3	AVG	
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170510010027	40	34	85	21	20	0	21	18	18	18	18.0	D
170510010028	40	32	80	18	18	0	18	20	20	19	19.7	B
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170510010030	40	37	92	24	23	13	24	19	19	19	19.0	A
170510010031	40	35	88	26	27	0	27	19	17	20	18.7	A
170510010032	40	31	78	21	20	0	21	20	20	19	19.7	A
170510010033	40	34	85	17	15	0	16	16	17	16	16.3	D
170510010034	40	36	90	19	18	0	19	18	18	19	18.3	B
170510010035	40	35	88	20	21	0	21	17	19	18	18.0	A
170510010036	40	37	92	20	25	12	21	19	19	19	19.4	C
170510010037	40	36	90	0	0	0	0	19	19	19	19.0	D
170510010038	60	17	28	25	27	17	26	18	19	18	18.3	C
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170510010049	40	37	92	27	28	21	28	18	17	18		D
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Susanna Kashem
SIGNATURE OF THE TEACHER

ROLL NO	Theory			Theory										GSE
	NO. OF CLASSES HELD	NO. OF CLASSES ATTENDED	ATTENDANCE	SESSIONAL (30)				CONTINUOUS EVALUATION (20)				GSE		
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170510011086	40	38	95	25	24	29	27	20	20	20	20.0	D		

Sulata Chakravarty
SIGNATURE OF THE TEACHER

GIRIJANANDA CHOWDHURY INSTITUTE OF PHARMACEUTICAL SCIENCE

PROGRAM :	BACHELOR OF PHARMACY (B. PHARM) (New Curriculum - 2017)		
UNIVERSITY :	Assam Science & Technology University		
NAME OF THE SUBJECT :		Practical	Instrumental Method of Analysis
SUBJECT CODE :		Practical	BP705P
SESSION :	Aug - December 2020		
NAME OF THE TEACHER(S) :	Mr Jaganakar Kushari & Mr Sumen Kumar		

COURSE OUTCOME ASSESSMENT SHEET

ROLL NO	Practical				ESE	Practical																SEMESTRAL			ESE CP						
	NO. OF CLASSES	NO. OF PRESENTERS	ATTENDANCE	GRADE		CONTINUOUS ASSESSMENT (20)																1ST SEM	2ND SEM	AVG							
						EXPT 1	EXPT 2	EXPT 3	EXPT 4	EXPT 5	EXPT 6	EXPT 7	EXPT 8	EXPT 9	EXPT 10	EXPT 11	EXPT 12	EXPT 13	EXPT 14	EXPT 15	EXPT 16					EXPT 17	EXPT 18	EXPT 19	EXPT 20		
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170510011003	10	80	100	0		17	19	17	18	17	19	19	17	17	18													35	36	34	B
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170510011008	10	80	100	0		17	17	19	18	19	17	18	19	19	19													31	28	30	O
170510011009	10	80	100	0		17	17	18	17	19	18	17	16	17	19													34	38	36	A
170510011010	10	80	100	0		18	18	17	18	19	17	19	17	17	18													32	33	33	O
170510011011	10	80	100	0		10	19	18	19	18	19	19	18	17	17													37	30	29	A
170510011012	10	80	100	0		19	18	17	19	19	19	19	18	17	19													34	28	36	O
170510011013	10	80	100	0		19	19	18	19	17	19	19	19	18	19													31	28	30	A
170510011014	10	80	100	0		19	19	17	18	17	19	18	19	17	18													31	30	31	B
170510011015	10	80	100	0		19	19	18	17	17	18	17	19	18	17													26	28	27	A
170510011016	10	80	100	0		18	19	17	18	18	17	18	15	17	18													29	33	31	A
170510011017	10	80	100	0		17	17	19	17	18	18	17	17	19	17													35	37	36	A
170510011018	10	80	100	0		18	17	19	18	19	17	18	17	15	18													37	38	38	B
170510011019	10	80	100	0		17	17	18	17	19	18	18	17	18	17													38	39	39	A
170510011020	10	80	100	0		18	18	17	18	19	17	18	18	17	18													23	25	24	A
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170510011022	10	80	100	0		18	18	17	18	19	17	18	17	18	18													23	25	24	A
170510011023	10	80	100	0		17	19	18	19	18	19	18	17	17	18													26	23	23	B

Suman Kumar
SIGNATURE OF THE TEACHER

