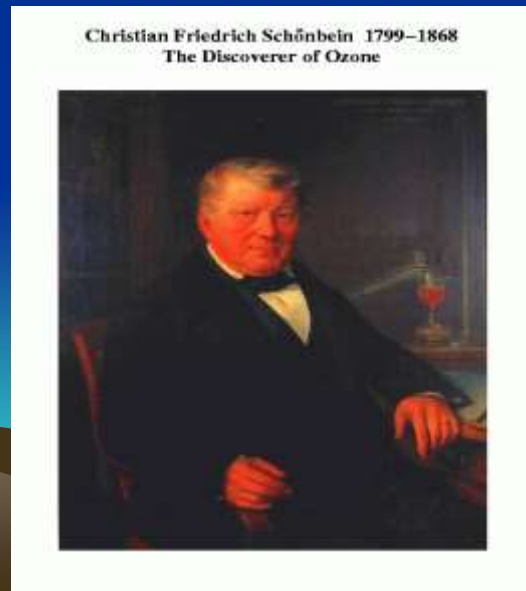


Paper chromatography



PAPER CHROMATOGRAPHY

- Paper Chromatography (PC) was first introduced by German scientist **Christian Friedrich Schonbein** (1865).
- PC 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.



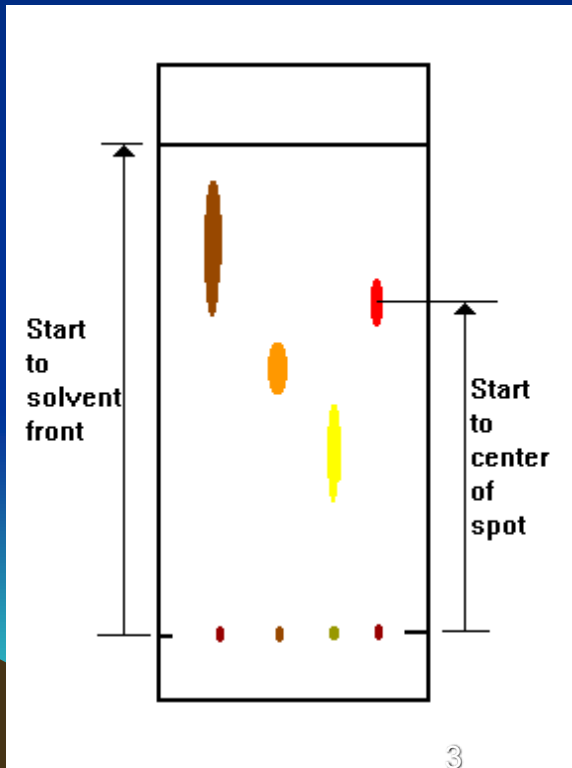
PAPER CHROMATOGRAPHY

- ANALYSIS OF UNKNOWN SUBSTANCES

It is carried out mainly by the flow of solvents on specially designed filter paper.

There are two types of paper chromatography, they are:

Paper adsorption and Paper Partition



PAPER CHROMATOGRAPHY

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 / Water present in the pores of cellulose fibers present in filter paper acts as stationary phase & another mobile phase is used as solvent

In general P.C – Paper Partition Chromatography

PAPER CHROMATOGRAPHY

PRINCIPLE OF SEPERATION

The principle of separation is mainly **partition rather than adsorption**.

Cellulose layers in filter paper contains moisture which acts as stationary phase & organic solvents/buffers are used as mobile phase

PAPER CHROMATOGRAPHY

- PRACTICAL REQUIREMENTS
- 1) Stationary phase & papers used
- 2) Application of sample
- 3) Mobile phase
- 4) Development technique
- 5) Detecting or Visualizing agents



PAPER CHROMATOGRAPHY

- STATIONARY PHASE AND PAPERS USED

Whatman filter papers of different grades like No.1, No.2, No.3, No.4, No.20, No.40, No.42 etc are used. In general this paper contains 98-99% of α -cellulose, 0.3 – 1% β -cellulose

Factors that governs the choice of paper:

- » Nature of Sample and solvents used.
- » Based on Quantitative or Qualitative analysis.
- » Based on thickness of the paper.

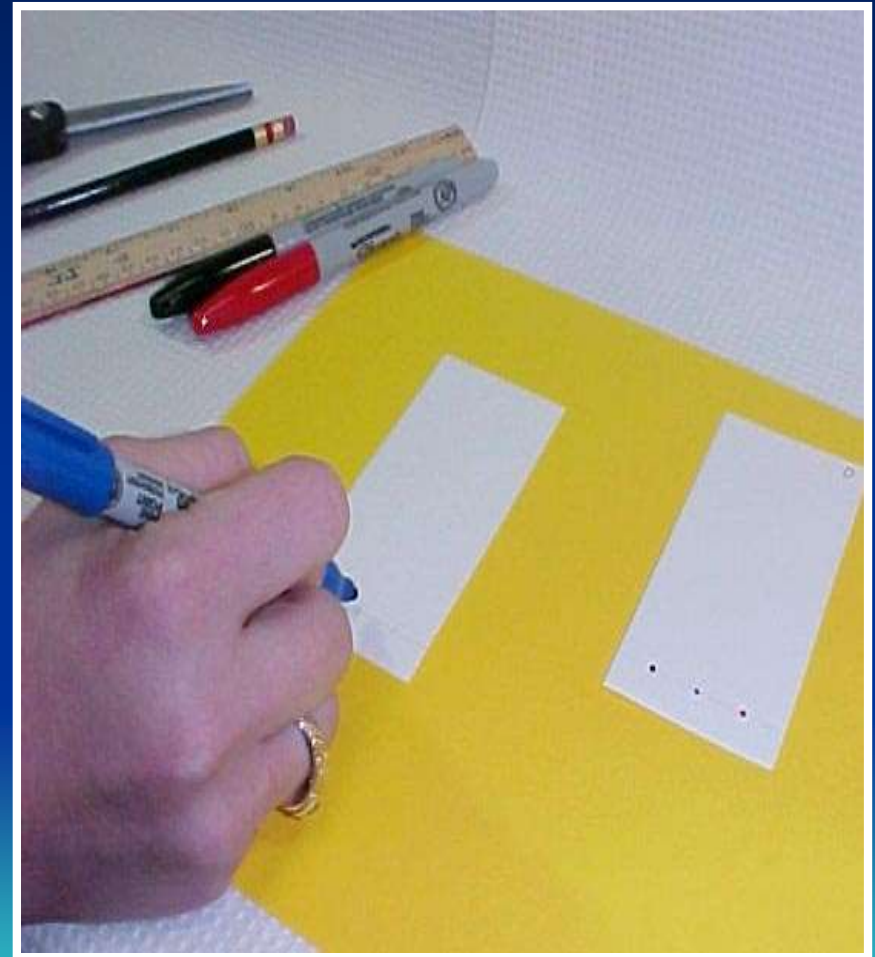


PAPER CHROMATOGRAPHY

- **Modified Papers** – acid or base washed filter paper, glass fiber type paper.
- **Hydrophilic Papers** – Papers modified with methanol, formamide, glycol, glycerol etc.
- **Hydrophobic papers** – acetylation of OH groups leads to hydrophobic nature, hence can be used for reverse phase chromatography.
- Impregnation of silica, alumina, or ion exchange resins can also be made.

PREPARATION OF PAPER

- Cut the paper into desired shape and size depending upon work to be carried out.
- The starting line is marked on the paper with an ordinary pencil **5cm** from the bottom edge.
- On the starting line marks are made **2cm** apart from each other.



PAPER CHROMATOGRAPHY

- Preparation of the solution
- Choice of suitable solvent for making solution is very important. Pure solutions can be applied direct on the paper but solids are always dissolved in small quantity of a suitable solvent.
- Biological tissues are treated with suitable solvents and their extracts obtained. Proteins can be precipitated with alcohol and salts can be removed by treatment with ion exchange resin.



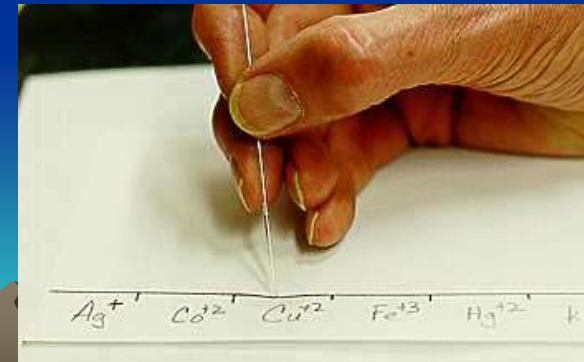
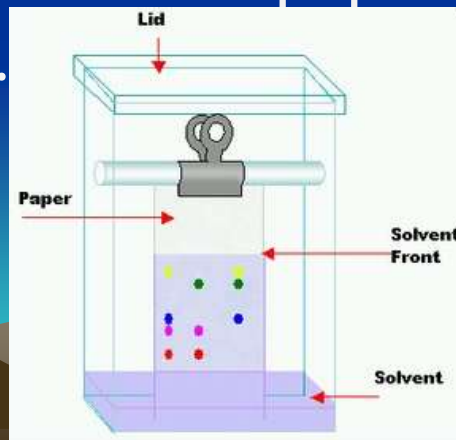
PAPER CHROMATOGRAPHY

APPLICATION OF SAMPLE

The sample to be applied is dissolved in the mobile phase and applied as a small spot on the origin line, using capillary tube or micropipette.

very low concentration is used to avoid larger zone

- The spot is dried on the filter paper and is placed in developing chamber.



Choice of the Solvent

- The commonly employed solvents are the polar solvents, but the choice depends on the nature of the substance to be separated.
- If pure solvents do not give satisfactory separation, a mixture of solvents of suitable polarity may be applied.

PAPER CHROMATOGRAPHY

- MOBILE PHASE

- Pure solvents, buffer solutions or mixture of solvents

- Examples- **Hydrophilic mobile phase**

- Isopropanol: ammonia:water 9:1:2

- Methanol : water 4:1

- N-butanol : glacial acetic acid : water 4:1:5

Hydrophobic mobile phases

dimethyl ether: cyclohexane

kerosene : 70% isopropanol

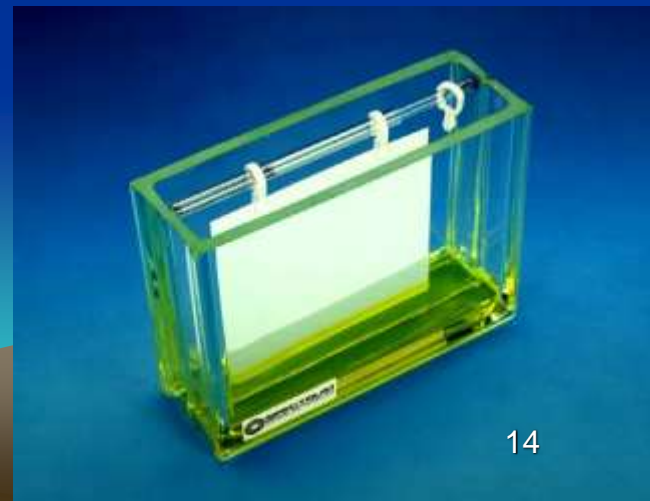


CHROMATOGRAPHIC CHAMBER

The chromatographic chamber are made up of many materials like glass, plastic or stainless steel.

Glass tanks are preferred most. They are available in various dimensional size depending upon paper length and development type.

The chamber atmosphere should be saturated with solvent vapor.



PAPER CHROMATOGRAPHY

DEVELOPMENT TECHNIQUE

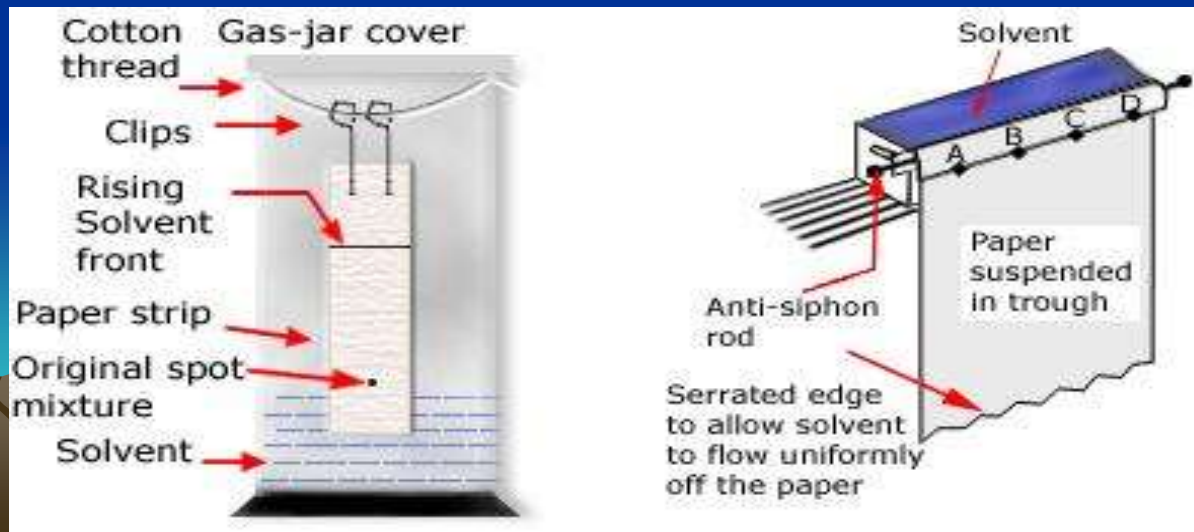
- Paper is flexible when compared to glass plate used in TLC, several types of development are possible which increases the ease of operation.
- The paper is dipped in solvent in such a manner that the spots will not dip completely into the solvent.
- The solvent will rise up and it is allowed to run $2/3^{\text{rd}}$ of paper height for better and efficient result.

PAPER CHROMATOGRAPHY

- Different types of development tech. are

1) ASCENDING DEVELOPMENT (go upward direction)

- Like conventional type, the solvent flows against gravity. The spots are kept at the bottom portion of paper and kept in a chamber with mobile phase solvent at the bottom.



PAPER CHROMATOGRAPHY

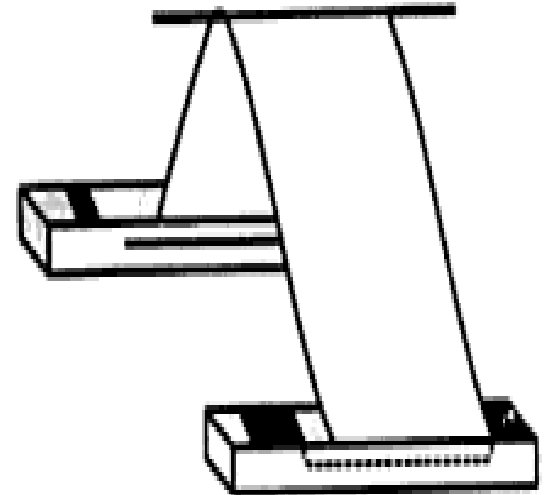
- 2) DESCENDING TYPE (a downward slope)
- This is carried out in a special chamber where the solvent holder is at the top. The spot is kept at the top and the solvent flows down the paper.
- In this method solvent moves from top to bottom so it is called descending chromatography.
- **ADVANTAGE IS THAT, DEVELOPMENT IS FASTER**

PAPER CHROMATOGRAPHY

3) ASCENDING - DESCENDING DEVELOPMENT

A **hybrid** of above two technique is called ascending-descending chromatography.

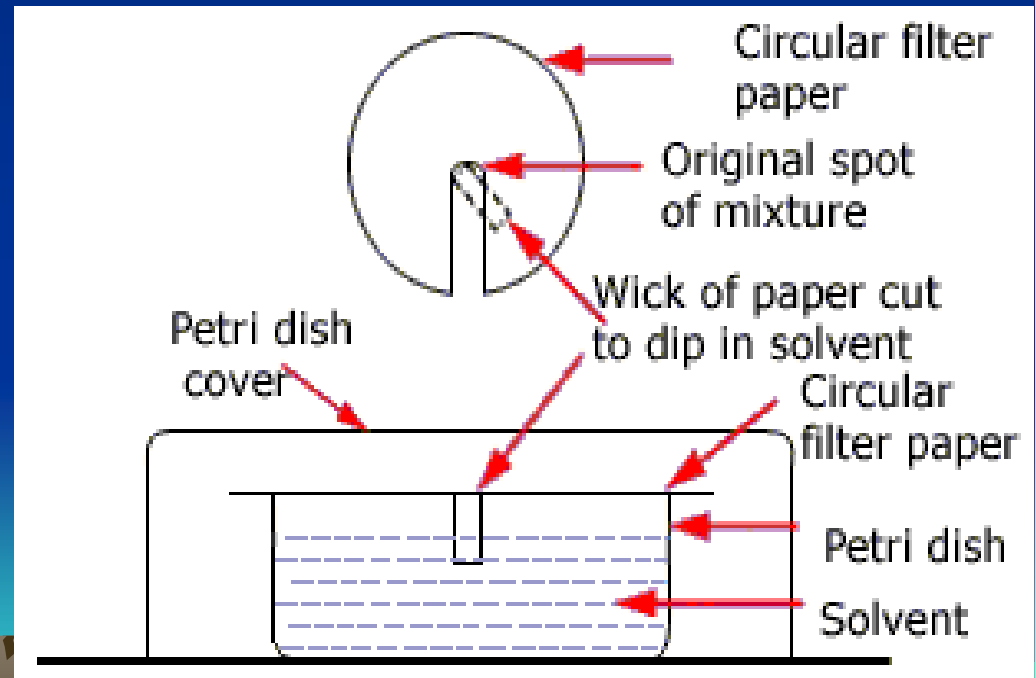
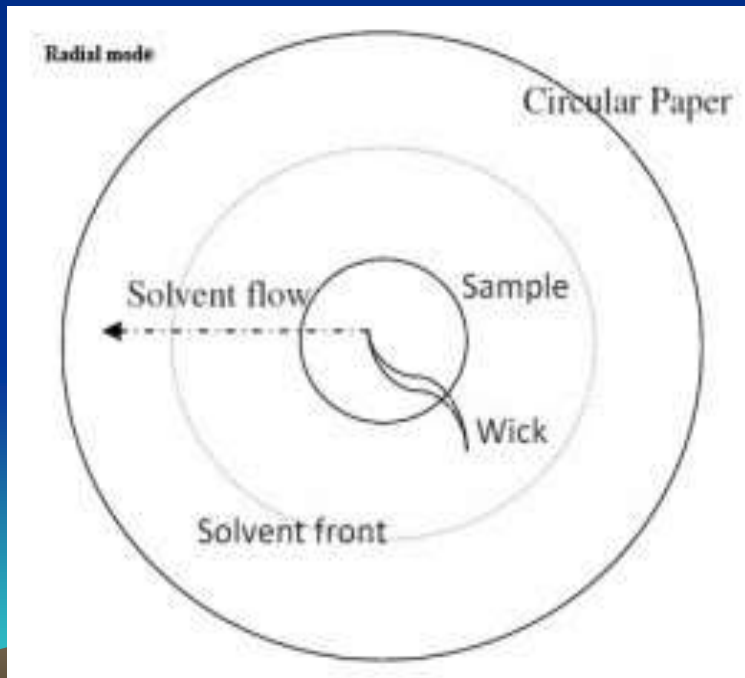
Only length of separation increased, first ascending takes place followed by descending



PAPER CHROMATOGRAPHY

4) CIRCULAR / RADIAL DEVELOPMENT

Spot is kept at the centre of a circular paper. The solvent flows through a wick at the centre & spreads in all directions uniformly.



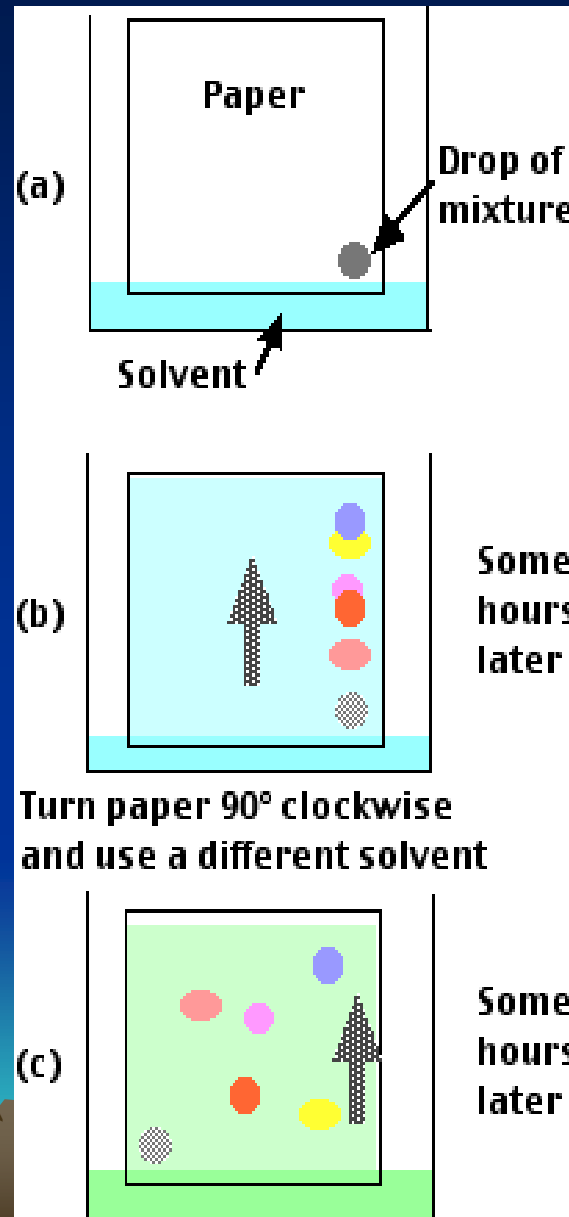
PAPER CHROMATOGRAPHY

5) TWO DIMENSIONAL DEVELOPMENT

In this method the paper is developed in one **direction** and after development, the paper is developed in the **second direction** allowing more compounds to be separated into individual spots.

in the second direction, either **same solvent/different solvent system** can be used for development.

TWO DIMENSIONAL DEVELOPMENT



PAPER CHROMATOGRAPHY

DRYING OF CHROMATOGRAM

- After the solvent has moved a certain distance for certain time the chromatogram is taken out from the tank & position of the solvent front is marked with a pencil.
- They are dried by cold or hot air depending on volatility of solvents. A simple hair dryer is a convenient device to dry chromatograms.



PAPER CHROMATOGRAPHY

DETECTING / VISUALISING AGENTS

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)

E.g. iodine chamber method,

UV chamber for fluorescent compounds – at 254 or at 365nm.

(b) Specific methods (Chemical methods) or Spraying method - examples,

- Ferric chloride
- Ninhydrin in acetone
- Dragendroff's reagents
- 3,5 dinitro benzoic acid
- Phenolic comp. & tannins
- Amino acids
- Alkaloids
- Cardiac glycosides



Following detecting tech. can also be categorized as

- **1) Destructive techniques**
- Specific spray reagents, samples destroyed before detection e.g. – ninhydrin reagent
- **2) Non-destructive techniques**
- For radio active materials - Geiger Muller counter
- uv chamber, iodine chamber

QUANTITATIVE ESTIMATIONS

The method can be divided into two main groups

1. Direct techniques-
2. Indirect techniques-

PAPER CHROMATOGRAPHY

- Direct Measurement Method

- (i) Comparison of visible spots
- A rough quantitative measurements
- Component in a mixture can be carried out by comparing the intensity and size of the spot with a standard substance.
- (ii) Photo densitometry
- The method is used with the chromatograms of colored compound, instrument which measures quantitatively the density of the spots.



PAPER CHROMATOGRAPHY

- (iii) Fluorimetry
- The compound to be determined by fluorimetry must be fluorescent or convertible into fluorescent derivatives.
- (iv) Radiotracer Method
- The compound containing radioactive element is labeled and treated with locating reagent. Using Geiger Muller counter.
- (v) Polarographic & Conductometric methods
- Used to measure the amount of material in the spot

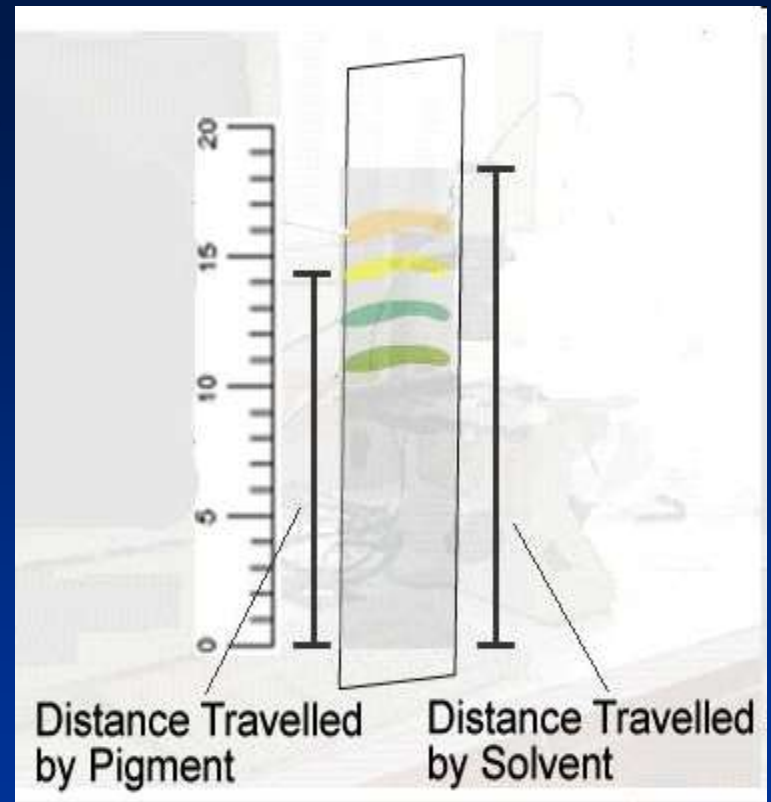
PAPER CHROMATOGRAPHY

- Indirect Measurement Method

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

Rf VALUE (Retardation Factor)

In paper chromatography the results are represented by Rf value which represent the movement or migration of solute relative to the solvent front.



The Rf value is calculated as :
$$\frac{\text{Distance travelled by the solute}}{\text{Distance travelled by the solvent front}}$$

Factors affecting Rf VALUE

- i. The temperature
- ii. The purity of the solvents used
- iii. The quality of the paper, adsorbents & impurities present in the adsorbents
- iv. Chamber saturation techniques, method of drying & development
- v. The distance travelled by the solute & solvent
- vi. Chemical reaction between the substances being partitioned.
- vii. pH of the solution

R_x VALUE

- In many cases it has been observed that the solvent front is run off the end of the paper. R_x value is thus used,
- It is the ratio of distance travelled by the sample and the distance travelled by the standard. R_x value is always closer to 1.

Sources of Error

- 1. Error during application of the spots
- Apply minimum volume of the concentrated solution in order to avoid diffusion through the paper which leads to poor separation
- Spots should be approximately of the same diameter.
- 2. Development
- Improper adjustment of the paper in the tank leads to this error so the paper should be held vertically.
- Do chamber saturation
- 3. Detection
- The spraying methods affect the final result

APPLICATIONS

- Separation of mixtures of drugs
- Separation of carbohydrates, vitamins, antibiotics, proteins, etc.
- Identification of drugs
- Identification of impurities
- Analysis of metabolites of drugs in blood , urine

ADVANTAGES OF P.C

Simple ,rapid ,inexpensive ,excellent resolving power

PRECAUTIONS IN P.C

**Establishing the vapor solvent equilibrium
Stability of solvent mixture is first ensured**

A vibrant and cheerful graphic with a blue sky and white clouds background. The words "Thank You" are written in a large, blue, rounded font with a white outline. The text is surrounded by several red hearts of varying sizes. In the foreground, there are red tulips and two butterflies: one orange and black, and another yellow and black. Green leaves and branches are visible in the upper corners.

Thank You