

### Separation

<b>Pure substances</b>	Contain only ONE thing - have fixed melting and boiling points
<b>Mixtures</b>	Contain MORE THAN ONE thing, do not have exact properties, formed by a physical change so they can be easily separated

### Purification as part of drug synthesis

Synthesis	Purification	Characterisation
Weigh up starting materials	Extraction	NMR
Set up a reaction	Recrystallisation	IR
Monitor reaction	Distillation	MS
Work up reaction to start purification	Chromatography	RF
	Solid-phase extraction	Melting point

### Separation techniques

1. Distillation	Used to separate the components of a LIQUID mixture by vapourising, condensing vapours and then collecting the liquid condensate. <b>Separation is a result of different boiling points</b>
2. Filtration	Physically separated solids from liquids Especially used for insoluble solids in liquids
3. Centrifugation	Uses centrifugal force that spins the samples fast Separated even fine solid matter from liquids

### Separation techniques (cont)

- |                      |  |
|----------------------|--|
| 4. Recrystallisation | 1. Dissolve impure crude material in minimum volume of solvent |
|                      | 2. Filter any insoluble material                               |
|                      | 3. Allow solution to cool slowly                               |
|                      | 4. Crystals will form  |

### Polarity

#### Definition of solubility

Ability of a solvent to dissolve in a solute, depending on the nature of the forces acting between the solute and solvent  
- **LIKE DISSOLVES LIKE**

e.g. **Ionic** compounds are more soluble in **polar** solvents  
**Covalent** compounds are more soluble in **non-polar** solvents

### Solvent mixtures

Immiscible: if the mixtures are too different, they will NOT mix, and will form separate layers or 'phases'

Partially miscible: mixtures are not too different but not too similar, they may mix at determinate proportions depending on the compositions

Miscible: if they are similar, they mix in EVERY proportion, forming one layer or 'phase'

### Liquid-liquid Separation

#### What is liquid-liquid separation?

A method in which 2 molecules initially present in one sample can be separated by giving them the choice of different immiscible solvents

- Very common when isolating or purifying a product
- Can be used to extract natural products
- Normally done in workup stage after a reaction

#### Criteria of organic solvent for liquid-liquid extraction

- SHOULD readily dissolve the substance to be extracted
- SHOULD NOT react with the substance to be extracted
- SHOULD NOT react with or be miscible with water (usual second solvent)
- SHOULD have a low boiling point so it can be easily removed from the products

Common extraction solvents are dichloromethane or ethyl acetate

### Chromatography

What is chromatography?

Physical method of separation where the components to be separated are distributed between 2 phases: the stationary and mobile phase

Common types:

- TLC (thin layer)
- Column
- HPLC (high performance liquid)
- GC (gas)
- SEC (size exclusion)
- Ion exchange
- Chiral

### Chromatography - definitions

Analyte	Substance being separated (or analysed)
Mobile phase	Phase that moves in a definite direction - consists of sample being separated and the solvent that moves the sample through the column
Stationary phase	The substance fixed in place for the chromatography procedure
Eluent	Solvent <b>entering</b> column
Elate	Solvent <b>leaving</b> column
Elution	Process of passing liquid through chromatography

### DIFFERENT METHODS & RETENTION OF ANALYTES

1. **Adsorption chromatography**  
Separation is based on differences between the **adsorption affinities** of the sample analytes for the surface of a solid-stationary phase  
Based on **non-covalent** interactions  
**Used for organic molecules**

### DIFFERENT METHODS & RETENTION OF ANALYTES (cont)

2. **Partition chromatography**  
Separation based on differences between the **solubility** of the sample analytes in the mobile and stationary phases → stationary phase = immobilised liquid  
Based on **non-covalent** interactions  
**Used for organic polar molecules**
3. **Exclusion chromatography**  
Separation based on **exclusion effects** e.g. differences in size and shape  
**used for proteins and nucleic acids**
4. **Gel electrophoresis**  
Specialised example of size-exclusion chromatography which uses agarose/polyacrylamide and passes electricity through  
Separates based on **size and shape**  
**Used for proteins and nucleic acids**  
Visualised by using UV and stains



## DIFFERENT METHODS & RETENTION OF ANALYTES (cont)

**5. Ion exchange chromatography**  
Separation based on **differences in ion exchange affinities** → e.g. differences in charge size and type  
Based on **ionic forces**  
**Used for cations, anions, proteins, peptides, amino acids and nucleic acids**  
- Cation exchangers = negative stationary phase  
Anion exchangers = positive stationary phase

**6. Thin layer chromatography**  
Separation based on **K<sub>x</sub> values**  
Polar stationary phase = thin layer of silica spread over a glass plate  
Stationary phase is placed in the mobile phase  
Analyte moves up the plate and components are spread based on their K<sub>x</sub> values  
**Lower R<sub>f</sub> = compound is MORE polar**  
**Higher R<sub>f</sub> = compound is LESS polar**  
**Increased eluent polarity can increase the R<sub>f</sub>**  
More polar = stronger interactions in the **normal** phase  
Less polar = stronger interactions in the **reverse** phase

**7. Flash column chromatography**  
Used for large amounts of material (5-25mg) when TLC is not effective

## INSTRUMENTAL CHROMATOGRAPHY

**1. High performance liquid chromatography**  
Analytical use: identify multiple analytes in complex mixtures  
Preparative use: purification  
**2. Gas chromatography**  
Separated by their **volatility**  
Forces the analyte through a column of the stationary phase by the gas mobile phase at a high pressure and temperature

## Quantitative analysis

Single point calibration	Multiple point calibration	Internal standard
1. A solution containing a known concentration of the compound to be measured in the sample is injected → relation concentration obtained = AUC	1. Several solutions containing different concentrations of compound to be measured are injected → AUC obtained	1. One solution containing a known concentration of the compound to be measured AND a known concentration of internal standards are injected to find a Response factor
2. The sample with unknown concentration is injected → new AUC is compared with previous to find concentration in the sample	2. The sample with unknown concentration is injected → new AUC is compared with previous to find concentration in the sample	2. The sample and the same known concentration of internal standard are injected to find the AUC and Response factor

## Separation in pharmacy

### Drugs in the clinic must be pure:

1. Impurities can be harmful and cause side effects
2. Impurities can alter the ability to formulate a drug correctly
3. Impurities can also affect the stability and shelf-life of the drug

### Drugs in development MUST be pure:

This is to ensure that it is the drug that brings about a therapeutic effect and not any impurities

- Before any new compound is tested, it is purified and characterised

Separation is also part of the quality control and monitoring process

## Importance of ionisation

Some functional groups can be charged, and this depends on:

- whether the molecule is an acid or a base
- the pH of the molecule

### Physiological pH = 7.4

we need to know what groups are charged at this pH as it will help us determine:

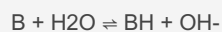
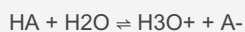
- types of drug-target binding interactions
- solubility (uptake and distribution)
- potential salt forms

#### e.g. 1 - Carboxylic acids

#### e.g. 2 - Aliphatic amines

Acids react with water:

Bases react with water:



pKa of carboxylic acids ~ <5

pKa of aliphatic amines ~ >8

These functional groups are almost ALWAYS ionised at physiological pH

pKa & pH = measures of dissociation

If the pKa = pH, then the functional group is 50% ionised

## Partitioning

Partition Distribution of a solute between 2 solvents

Compound is present in BOTH phases according to its relative solubility in both

Solute distributes itself between the 2 liquids in accordance with its **partition coefficient**

## Partitioning (cont)

**Partition coefficient** Ratio of the concentration of the solute in one liquid OVER the concentration in the other

Dynamic equilibrium exists between the 2 liquids, temperature dependent

## Partitioning in pharmacy

Relative hydrophobic/hydrophilic properties of a drug are crucial as it influences:

- solubility
- adsorption
- distribution
- metabolism
- excretion

**Drug too polar?** It will not cross the cell membranes across the gut wall

**Drug too lipophilic?** Drug will be poorly absorbed, so will likely be taken up into the fat tissue and not circulated

## LogP

$$P = \frac{\text{Concentration of drug in octanol}}{\text{Concentration of drug in water}}$$

LogP: used as a measure of hydrophobicity of a drug

Hydrophobic/hydrophilic character is measured using partition

- Hydrophobic molecules prefer the **octanol layer** → high P
- Hydrophilic molecules prefer the **water layer** → low P

LogP measures only unionised forms of the drug between octanol and water

LogD is used to represent relative distributions of all species, charged or uncharged

## Distribution Coefficient (K<sub>x</sub>)

$$K_x = \frac{C(\text{stat})}{C(\text{mob})}$$

C(stat) = conc of X in **stationary** phase

C(mob) = conc of X in **mobile** phase

Each compound will have a different K<sub>x</sub>

Chromatographic separations can be altered by changing the nature of the stationary and/or mobile phase(s)

### Solid-phase extraction

Used in sample preparation to remove matrix interferences such as proteins

Concentrates sample of interest

Solid phase in SPE is similar to the stationary phase used in chromatography

Active substances can be retained or unretained

### SPE vs. Chromatography

- Column is much smaller for SPE
- Specialised columns are more common
- Analytes are typically strongly retained on the SPE column, then impurities are washed away
- ↳ In chromatography, both the analyte and impurities are passed through at different rates
- SPE is used for sample preparation, then chromatographic methods follow

### Stationary phases

#### Normal phase

Stationary phase = polar  
Mobile phase = non-polar

Components elute in order of **increasing** polarity

Used in TLC and flash column

**More** polar compounds are retained stronger

More polar solvents increase elution

Solvents = the complete range but rarely greater than 20% for very polar solvents

A strong solvent for the normal phase is a weak solvent for the stationary phase

#### Reverse phase

Stationary phase = non-polar  
Mobile phase = polar

Components elute in order of **decreasing** polarity

Used in HPLC

**Less** polar compounds are retained stronger

Less polar compounds increase elution (e.g. methanol, ethanol)

Solvents = very polar solvents (e.g. water, methanol)

### Component retention

For TLC, retention is expressed in terms of  $R_f$

For column, retention is expressed in terms of column volumes (CV)

**The relationship between  $R_f$  and CV are reciprocal**

Lower  $R_f$  = greater CV → Low  $R_f$  if preferred as it increases analyte contact time and improves chances of component separation or resolution

To separate adjacent compounds, a large  $\Delta CV$  is desired

### Order of elution

TLC and HPLC depend on **polarity**

- Normal phase: polar retained more, non-polar elutes first
- Reverse phase: non-polar retained more, polar elutes first

Gas chromatography depends on **volatility or molecular weight**

- Highly volatile or low MW elutes first

Size exclusion depends on **molecular size**

- Low size retained more, largest size elutes first

Ion exchange depends on **charge**

- Complementary charges are retained more



By **eyeeyuu**

[cheatography.com/eyeeyuu/](https://cheatography.com/eyeeyuu/)

Not published yet.

Last updated 15th August, 2022.

Page 5 of 5.

Sponsored by **ApolloPad.com**

Everyone has a novel in them. Finish Yours!

<https://apollopad.com>