$H_{3}CO + 2e = H_{2}^{+} + 20H^{-})$ ¹ م (² م 2 ·OH, Cu; H20 $= H_{20} + 20H^{-} - Sn(0H)_{1}$ Ģ H_2O H_3CO OH Fe - 2e -+ Fe 2+ K 2H20 80 0.00 V(SOy) = 2H; Unp = Dosp In/In*//p 000 $k: \mathcal{P} \mathcal{B}^{2+} \mathcal{L} \mathcal{P} = \mathcal{P} \mathcal{B};$ 2e⁻ 2e = H2+20H NH2 NН V Br2=VBr3; N NH, CH CН 60-4e = 40n; H 2H20+02+2H20+4e= 40H; CH 50 $(V0_2)N0_3 + 5N0_2 + 3H_20_3$ Ν OM3 0H2 a, Na $V + Br_2 = V Br_3; Jn / Zn^2 // P$ 02+2H20+4e= 40H; NOH3 H3CO. H-0,4 0,2 0,6 $V_{\rm P} H N O_3 = V O_2 N O_3 + N O_2 + H_2 O_3$ 1 H2+20H 02+2H,0 СH $\mathcal{V}_{np} = \mathcal{V}_{oS}$ CH3 NH2 2H20+ Q $2H_{2}O$ 0H202 0/H ٥ 2H20+2e=rc21 +204 $\sin^{+}20H = in(0H)_{2} NH_{3}$ ΗN CH 3 $\Delta G = \Delta H \quad \exists H_2 SO_4 = V(SO_4)_2 + 2H; CH_3$ Cu $Sn^{2+}+2e^{-}=Sn$ N'OH3 Sn+02 + 2HO -TZS $Sn^{2+}+2e^{-}=Sn$ K (_) NH2 CH3COONa+H2O CI/NatNa3 0 + Pp^{2r} = Zn²⁺ + PB; $V_{r}: H_{2} + J_{2} = 2 H J$ $CH_3 PO_4^3 - 1,16 HPO_3^2 - 1,50 O_2 + 2H_2 0 + 4e$ Zn $H_3(VF_6)$ H+Sn⁺ $V + J_2 = V J_2$; $H_2 P O_2 - 205 P_4 - 0.19$ $Sn + O_2 + 2H_2 O_2 + 4e = 40 H + Sn^+$ 40 $V_{6}: 2HJ = H_{2}+J_{2}$ $Na_2 SO_4 \rightleftharpoons 2 Na^+ + SO_4^2$ CH 3 $T_{4}S = 412 - (298, 41, 10^{-3})$



CHAPTER ANALYSIS



TIME



Usually tested in MCQs
 Tested as add-on to other chapters

• Relatively straight forward chapter

2 **key** concepts

• 3 advanced concepts

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ightarrow Salts, Fuels & Crude Oil



- Light overall weightage
- Constitute to **0.5%** of marks for past 5 year papers

FRACTIONAL

DISTILLATION

7 DIFFERENT SEPARATION TECHNIQUES



DISTILLATION

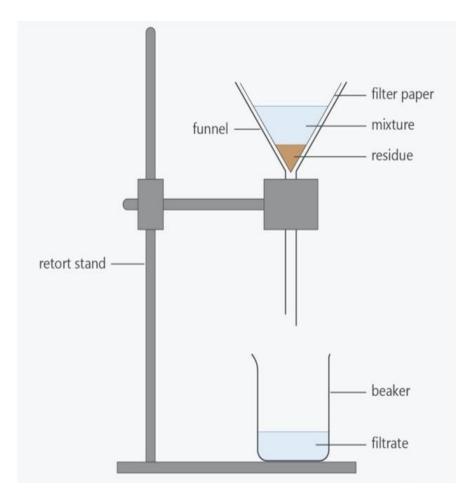
PURE SUBSTANCE VS IMPURE MIXTURE

	Pure Substance	Impure Mixture
Definition	Only one type of substance	Two or more substances
Physical properties	Fixed proportion	Any ratio
	Fixed M.P. & B.P.	Have a range of M.P. & B.P.
	Single spot on chromatogram	Multiple spots on a chromatogram

COMPOUND VS MIXTURE

	Compound	Mixture
Formation	Chemically combined	Physically combined
Separation technique	Can be separated using chemical methods (Decomposition, electrolysis, reduction with carbon)	Can be separated using physical methods (separation techniques)
Composition	Fixed ratio	Any ratio
m.p/b.p	Fixed mp & bp	Have a range of M.P. & B.P.

FILTRATION



1) Put a filter paper onto the filter funnel.

2) Pour the mixture into the filter funnel.

3) The **residue** will remain in the filter funnel while the **filtrate** will be collected in the beaker.

Magnetic Roller

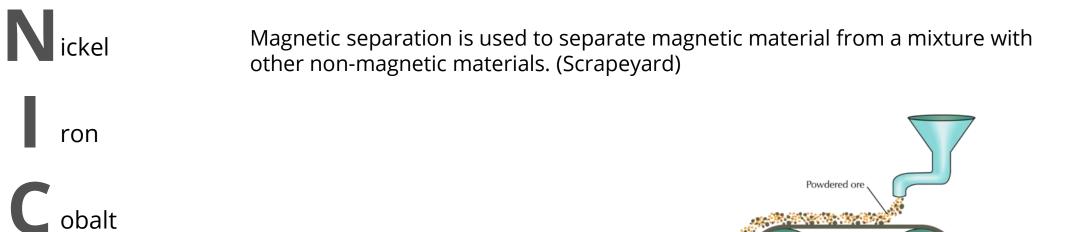
Moving belt

MUST KNOW

MAGNETIC SEPARATION

Magnetic Materials

teel



St' Nics girls are pretty right? Know a Nicholas that is handsome? Are you attracted? That's right, magnetic material.

7

Many students are confused when it comes to these 2 separation techniques:

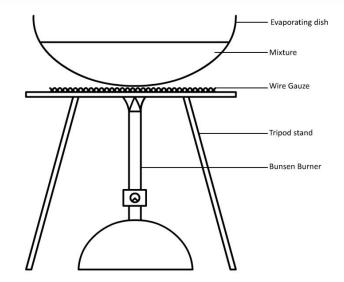
KEY CONCEPT

How exactly are they different and in which scenarios do we use them?

TWO METHODS EVAPORATION TO DRYNESS CRYSTALLISATION



EVAPORATION TO DRYNESS



For this method, the solution is heated until all the **water is** evaporated completely, leaving behind only the solid.

However, a **limitation** to this method is that it **cannot be used for solids that decompose on heating**.

For example, sugar decomposes upon heating, salt however does not.

*A saturated solution contains the maximum amount of solute that can be dissolved in the solvent at a particular temperature.

MUST KNOW

CRYSTALLISATION

Crystallisation is different from evaporation to dryness.

The aim of crystallisation aim is not to evaporate ALL of the water, but rather it focuses on heating it till saturation. After which, it is left to cool in order to obtain the crystals.

This is used for crystals that decompose upon heating to be collected. (ie: sugar)

Steps:

1) Heat the solution till saturation.

2) Allow the saturated solution to cool and pure solid crystals will form slowly.

3) Filter to collect the crystals.

4) Wash the crystals with cold distilled water and dry between sheets of filter paper.



Ultimately, what it boils down to the nature of the salt.

The critical question to ask is:

Will the solute decompose under heating?

If yes, use crystallisation.

If not, use evaporation to dryness.

Example:

Sugar decomposes under heat, crystallisation is the correct choice.

Salt has high melting & boiling point, evaporation to dryness will get the job done.

EVAPORATION TO DRYNESS VS CRYSTALLISATION

When do we do simple distillation and when do we do fractional distillation?

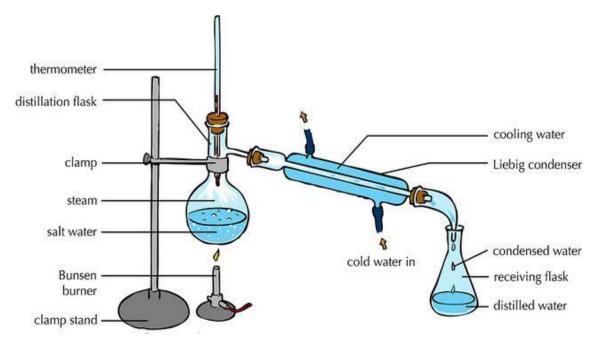
Is one method better than the other?

FATHER & SON SIMPLE DISTILLATION & FRACTIONAL DISTILLATION

KEY CONCEPT



SIMPLE DISTILLATION

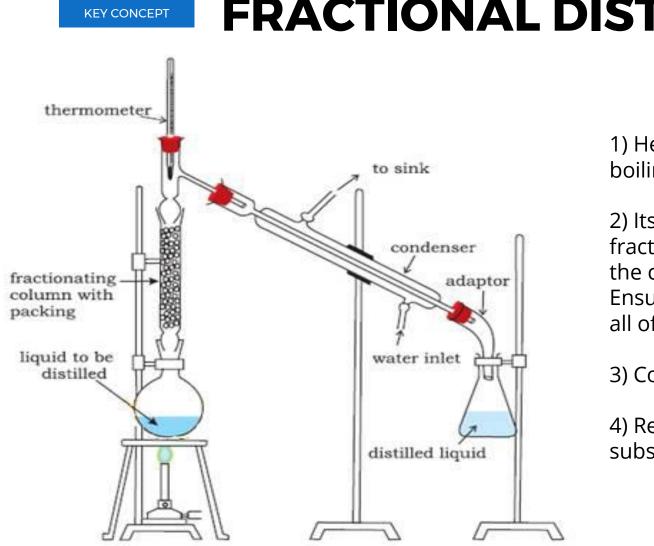


KEY CONCEPT

1) Heat the solution in a round-bottomed flask for even heat distribution. Boiling chips are added to prevent vigorous movement of liquid which ensure smooth boiling.

2) The water boils and water vapour rises and enters the condenser. The water vapour cools down in the condenser and is collected as a distillate.

3) The **distillate** is collected in a conical flask to prevent spillage.



FRACTIONAL DISTILLATION

1) Heat the mixture. The liquid with the lowest boiling point will be the first to be vaporised.

2) Its vapour will rise up to the top of the fractionating column, cooled as it passes through the condenser and be collected as the distillate. Ensure the temperature remains the same until all of first vapour has condensed.

3) Collect the distillate in the conical flask.

4) Repeat the process and collect the different substances that were in the mixture.

ADVANCED

things to note

Understanding the science behind fractional distillation

Difference in boiling points

The rationale behind fractional distillation is the miscible liquids in the solution having a minimum of **at least 10°C** difference in boiling point.

By boiling the liquids at their respective boiling point, it allows us to separate them.

Purpose of fractionating column

A fractionating column contains a large number of glass beads, creating **a larger surface area for condensation of vapours** for substances that have yet to reach their boiling point. This would only allow the intended vapour to escape.

Purpose of thermometer

The thermometer is placed at the tip of the fractionating column, right before the gas enters the condenser.

By doing so, we can **monitor the temperature of the gas that is escaping** accurately, allowing us to adjust the intensity of the heat accordingly.

KEY CONCEPT

So in which situations is simple distillation used and which situations do we use fractional distillation?

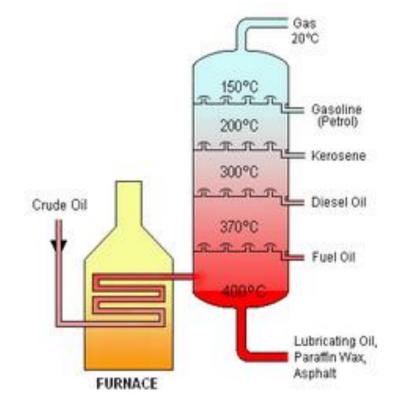
FATHER & SON SIMPLE DISTILLATION & FRACTIONAL DISTILLATION



*Key concept from Fuel and Crude Oil chapter in Organic Chemistry.



APPLICATION: OIL REFINERY



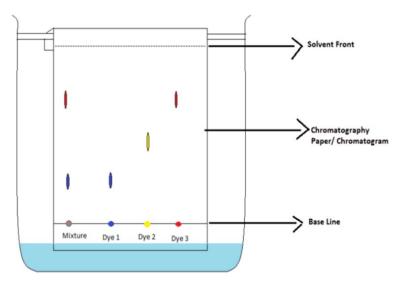
See you again soon!

CHROMATOGRAPHY

Chromatography is used to separate and identify miscible solutes that are dissolved together. (Food substances, dye etc...)

The different components can be identified by comparing their **Rf values (retention factor).**

Chromatography works under this principle: **the rate at which a particular solute moves relative to the solvent is fixed.**



ADVANCED

things to note

Chromatography is a method used to separate and identify small amounts of solutes that are dissolved in solvents.

Substance solubility in solvent

A **substance's solubility in a particular solvent** is commonly the main reason which results in **differing R_f values**.

Locating Agent

For colourless substances, a **locating agent** is required **to make the colourless solution visible.**

Knowing the names of specific locating agents is not needed. Yay!

Precaution

Starting line should be drawn with pencil instead of ink as the ink may dissolve in the solute, causing the results to be inaccurate. Commonly tested!

For more notes & learning materials, visit: <u>www.overmugged.com</u>

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