

I. Purpose - a concise statement of the qualitative or quantitative goals in an experiment,

II. Data - Direct qualitative or quantitative observations that require no manipulation,

	sample	mass (g)	%
A	.		
	.		
	.		
B	.		
	.		
	.		

C_4H_8O

C_4H_2O

trial trial

III. Calculations - Any mathematical manipulation of data,

$$\% \text{ composition} = \frac{\text{mass of compound}}{\text{mass of mixture}} \times 100\%$$

$$\% \text{ yield} = \frac{\text{mass obtained}}{\text{theoretical mass}} \times 100\%$$

$$\% \text{ error} = \frac{\text{experimental value} - \text{theoretical value}}{\text{theoretical value}} \times 100\%$$

% error for temperature must always be calculated in Kelvin.

IV. Conclusions - A summary of the results of an experiment \Rightarrow that addresses the goals as presented in the purpose statement,

V. Discussion -

Chromatography - A technique by which a mixture of compounds can be separated on the basis of their polarity.

inside coated w/ polar material

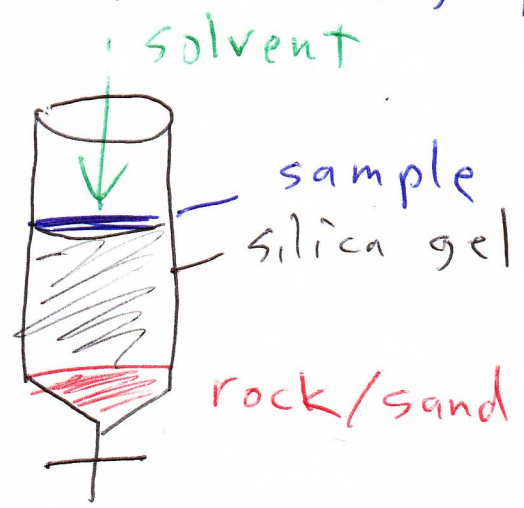


~~and~~ stationary phase - an immobile support that contains a compound that can separate the components of a mixture passed through it.

mobile phase - a gas ~~or~~ liquid that carries the mixture to be separated through the stationary phase.

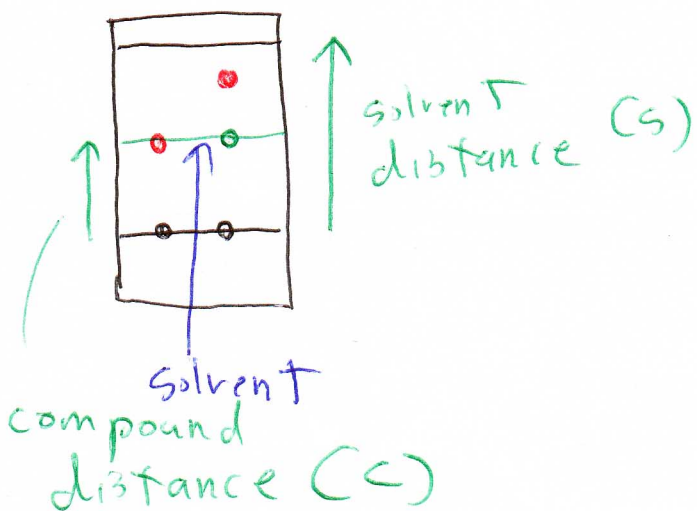
Gas chromatography (GC) - A sample is volatilized (evaporated) and passed through a column (a tube) that is coated on the inside with some form of separatory material (stationary phase). The sample is pushed through the column by an inert gas (such as N₂ or Ar) called the carrier gas (mobile phase).

Column chromatography



TLC - thin-layer chromatography

#3

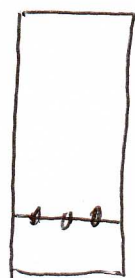


$$R_f - \text{Retention factor} \\ = \frac{\text{compound distance (c)}}{\text{solvent distance (s)}}$$

Preparing TLC plates

* TLC plates must always be marked in pencil, since the ink of a pen would travel with the development solvent,

* Do not press too hard with a pencil on a TLC plate since the silica gel can be scraped off,



- Draw a line across the bottom of the plate parallel to the bottom roughly 1 cm from the bottom,

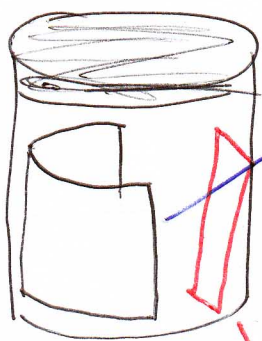
- The plate is then spotted with one or more compounds.

Development

TLC solvents - A solvent or solvent mixture is chosen so that it most effectively separates the components of the sample being separated.

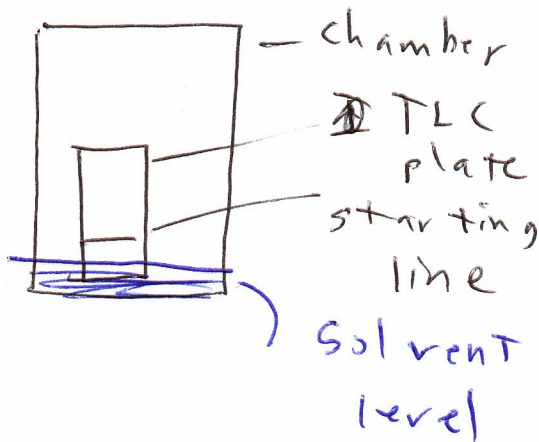
- hexanes / ethyl acetate → more polar
- chloroform / acetic acid

Development chamber



Filter paper - used to ensure that the interior of the chamber is fully saturated w/ solvent to prevent solvent evaporating off the plate.

The TLC plate must not touch the filter paper, otherwise solvent could flow sideways across the plate.



The level of solvent in the chamber must be below the start line.

Once the solvent nears the top of the plate, remove the plate from the chamber

and immediately mark the position of the solvent front.

Visualization

Some forms of silica gel contain a dye that's responsive to UV light. If a spot contains a compound that absorbs UV light, the spot will appear dark, since UV light would not reach the dye. (non-destructive)

chemical stain - the TLC plate is dipped in a compound that will react with the spots on the plate. (destructive)

~~phenol~~ p-anisaldehyde

