

Dilutions and Titrations

- Sometimes you know the concentration of a solution but it's not what you require
 - Therefore, you must dilute the stock solution
 - The amount of substance (mol) is the **same in the diluted sample as the undiluted sample**
 - The volume is the only variable that changes
- To dilute a sample, **take the volume of the original solution and add a known volume of solvent** to produce a more dilute solution
 - The undiluted and diluted sample has the following equations respectively:
 - **Moles = $C_1 \times V_1$**
 - **Moles = $C_2 \times V_2$**
 - Therefore: **$C_1V_1 = C_2V_2$**
- For example, if you have 100ml of a 2M HCL solution and you add 900ml of H₂O, what is the concentration of the diluted sample?
 - It will be $0.1 \times 2 = 1(c)$
 - It will then be $0.2/1 = c$
 - $C = 0.2 \text{ mol L}^{-1}$
- **You can also work out the volume needed to give a certain concentration in a diluted solution**, for example:
 - If the concentration of the undiluted sample is 2M, the volume is 0.2L and the concentration of the diluted solution is 0.2M then the volume of the new solution will be:
 - $(0.2 \times 2)/0.2$ which is 2L
- Sometimes you need to dilute a solution by a significant amount, and so this is done using **serial dilutions**
 - This is done by **adding small dilutions to the stock solution** and then taking that as the **new stock solution**. This is then **repeated and yields many different sample dilutions**
- **Titration** is used to directly measure the concentration of a sample
 - This is done using a **series of chemical reactions to measure amounts in a sample**
 - The **production of a product** is key in this process
 - **A burette is used to deliver the second reactant to a flask** and an **indicator** is used to **detect the endpoint of the reaction**
- A **coupled reaction** is when the product from one reaction is then used straight way as the ingredient for another reaction, for instance $A+B=C$ and $C+D=E$, therefore you cannot get E without A or B
- **UV spectroscopy** can be used to detect the concentration of a sample:
 - **The amount of light that passes through the sample is indirectly proportional to the concentration**
 - In a **low concentration**, **much light** passes through the sample
 - In a **high concentration**, **little light** passes through
 - The **Beer-Lambert** law states that the **absorbance of light is equal to the molar absorptivity ($L \text{ mol}^{-1} \text{ cm}^{-1}$) (ϵ) x the concentration (mol L^{-1}) (c) x the path length (cm) (l):**

$$A = \epsilon cl$$