Standard Solutions

# Introduction

When carrying out any quantitative work in chemistry, it is important to know the concentration of any solutions you use. Too great a concentration and some reactions will become dangerous , too low a concentration and some reactions will not work.

For most practical work, it is possible to get by making reasonably accurate solutions in the normal way. For instance weighing out the appropriate number of moles of solid and diluting to the required volume – or diluting a solution of concentrated acid. For analytical work, however (and this does crop up at Advanced Higher level), you sometimes need to know concentrations more accurately. You do this by a process called standardisation.

There is a certain number of chemicals that can be used as what are called primary standards. That means that they can be used as references or yardsticks to enable you to benchmark other reagents that will react with them.

Features of a primary standard include:

* **High purity** – if your compound is not pure, it is no use as a standard because you can never know its exact concentration.
* **Stability** (low reactivity) – if your compound is not stable, then you will only be able to use it as a reference if it is freshly obtained and even then you may not be entirely sure.
* **Low hygroscopicity and efflorescence** – similar to above. If your compound is absorbing or losing water, you cannot know its molecular mass accurately.
* **High solubility** – you need to be able to make a reasonably concentrated solution.
* **High molecular mass** – the higher the molecular mass, the more you weigh out to make a solution and so any error is proportionally less.
* Ideally, but not essentially, it should be of **low toxicity**, be **readily available** (and not too expensive) and not be too hazardous for the environment.

# Some Common Primary Standards

Here are a few common chemicals that are suitable for use as primary standards

|  |  |  |
| --- | --- | --- |
| Sodium carbonate | Potassium bromate | Sodium chloride |
| Potassium iodate | Benzoic acid | Potassium dichromate |
| Sulphanilic acid | Sulphamic acid |  |

# Preparing a standard solution

1. Work out the amount of the primary standard you need for the molarity and volume of your solution.
2. Roughly weigh out slightly more than this,.
3. Dry your solid in an oven - usually at about 110 – 120°C.
4. Allow to cool in a dessicator
5. Weigh out the exact amount of solid needed to make your solution and put it into a volumetric flask.

*Make sure you find the mass of solid by subtraction rather than taring the balance – this will help to eliminate any systematic errors (see Weighing).*

1. Dissolve in about ¾ of the final amount of distilled/deionised water\* (ideally boiled out and cooled to remove dissolved gases as well)

*\* Or other solvent – though it is not common to make up standard solutions in solvents other than water.*

1. Use more of the distilled water to wash out the weigh boat and add the washings to the flask.
2. Top up to the mark on the volumetric flask to get your standard solution.

*It is a good idea to make up quite a concentrated solution that you can then dilute down – that way you will minimise any weighing errors.*

# Specific Examples

|  |  |
| --- | --- |
| Solution to be standardised | Standards used |
| Aqueous strong acids | Standard sodium carbonate solution (using methyl orange as an indicator) |
| Ethanoic and other weak acids | Pre-standardised sodium hydroxide (using phenolphthalein as an indicator). |
| Alkalis | Pre-standardised solution of hydrochloric acid (standardised as above)  OR Benzoic acid (using phenolphthalein as an indicator)  OR Sulphamic acid (using phenolphthalein as an indicator) |
| Sodium Thiosulphate | Potassium iodate (or bromate) to release iodine and titrate against it (using starch as an indicator near the end-point). Each iodine molecule reacts with 3 moles Na2S2O3 |
| Silver Nitrate | Standard sodium chloride (Using 5% potassium chromate solution as an indicator). Titrate until the first colour change. |
| potassium permanganate | Standard ethanedioic (oxalic acid) No indicator needed. 2 moles of potassium permanganate react with 5 moles of ethanedioic acid. |

This is just an overview. You will find details on how to standardise the above solutions on the SSERC website under the name of the solution to be standardised. Details of others can be found in books such as Vogel’s ‘Handbook of Quantitative Inorganic Analysis’. If you do not have access to a suitable book and can’t find the information on our website, get in touch and we’ll find out for you.