

Analytical Chemistry

Is a branch of chemistry which is concerned with identifying the constituents of the chemical and the possibility of quantification or qualitative.

Analytical chemistry is divided into two branches:

1. Qualitative analytical chemistry:

It is concerned with the knowledge of constituent elements types of the compound, It also looks at how the elements or materials are separated and identified and is concerned with the external appearance of the compound such as color and smell.

2- Quantitative Analytical Chemistry:

It is concerned with quantifying the elements or compounds in a sample.

Which in turn are classified into two parts:

A.Volumetric analysis:

It is one of the ways by which the amount of elements or compounds in a sample solution is estimated (it is better to say how it is estimated).

B. Quantitative weight analysis:

It is based on the estimate of the material by weighing it or by any other method. The material is estimated as a weight in grams or its parts or percentage in a particular model and is divided into:

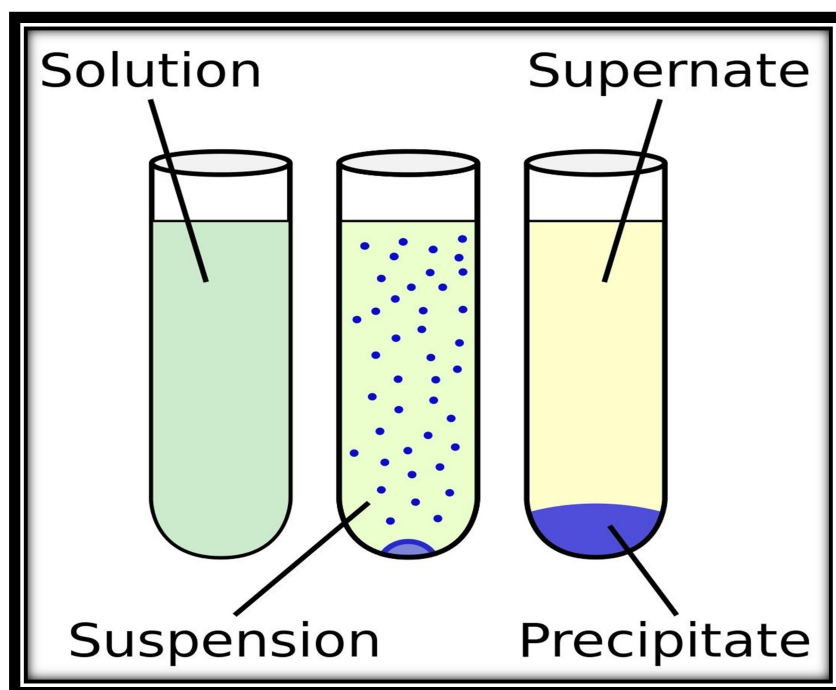
- 1- Precipitation method.
- 2- Evolution volatilization.
- 3- Specific gravimetric methods.
- 4 - Methods of electrostatic deposition Electro gravimetric methods.

NOTE :

we prefer the methods of volumetric analysis more than the methods of weight analysis, knowing that the latter is more accurate than the previous one this is due to the slow of the weight analysis.

The precipitation method is the best method used in weight analysis.

The sample is transferred to the solution to be analysed in a suitable manner. The element to be evaluated is precipitated as a non-dissoluble compound, after which the precipitate is filtered and washed well and then burned and dried.



Properties of precipitants in the methods of weight analysis:

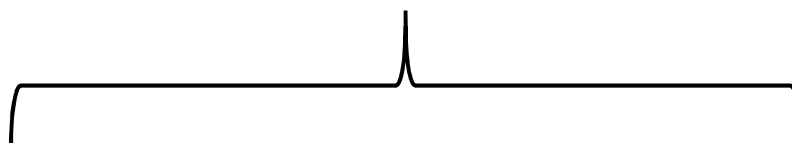
- 1- Must be known chemical formula.
- 2- The solubility of precipitant is very low.
- 3- The crystals of the precipitant of an appropriate size (large).
- 4- The precipitate shall be free of contaminants that cannot be washed or removed during drying of the precipitate.
- 5- The precipitate shall remain stable at the drying temperature (any thermal constant).
- 6- The Molecular weight of the precipitant is large compared with the material to be estimated to avoid the experimental error.

The precipitate factor

Is the chemical factor that causes the precipitation of the soluble matter to be estimated, forming precipitant or low soluble salts in the water .



The precipitate factor is consists of two types:



1- Selectivity

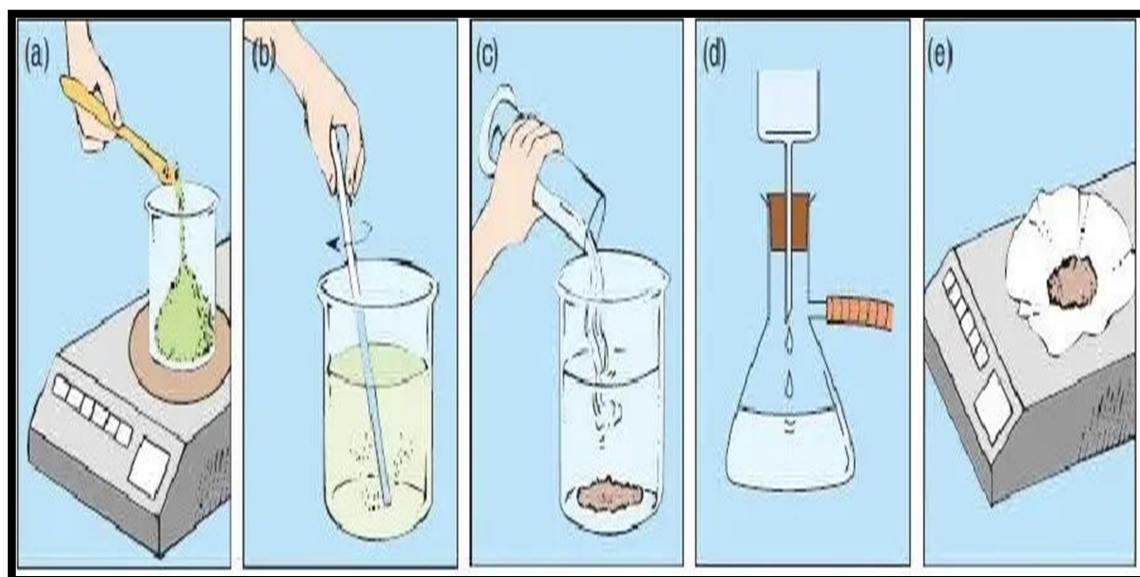
Works on the deposition of a group of ions

2- Specificity

Works on depositing only one ion without other substances.

Quantitative weighting steps:

- 1- Preparation of sample and dissolution.
- 2- Precipitation of analyte.
- 3- Digestion or aging of the solution.
- 4- Filtration of precipitate .
- 5- Washing of precipitate.
- 6- Drying and ignition.
- 7- Weighing and calculation.



1- Preparation of solution

This is done by selecting a suitable weight from the sample to be estimated and then dissolving the sample by selecting the appropriate solvent, As well as to create a suitable conditions for the experiment, such as pH of solution and solution temperature.

2- Precipitation of analyte :

The precipitation process of the selected components from the rest of the components in the solution is done by adding the precipitant factor.

The precipitate process consists of two steps :

First :

The formation of small particles (1-100) nm, called nuclei.

Second:

The nuclei growth into large particles. The nuclei do not appear directly after the addition of the precipitated agent but often is noticed after a period called the induction period. The duration of the sediment varies with the sediment difference and ranges from the second parts of silver chloride to a few minutes in the case of barium sulphate.

3. Digestion of precipitation:

It is the process of heating the precipitant with the standard solution (usually 90 or 95 C° and leaving it a period of time.

Interest from digestion of precipitation :

- 1- Increased the solubility of small particles (Help the crystals growth).
- 2- Leave the precipitate with the standard solution for a period of time allowing these particles to be precipitate on the large particles.
- 3- helps the grouping of colloidal precipitant.

4 – Filtration of precipitant :

The purpose of the filtration process is to separate the precipitation from the standard solution and the filtration process is easier and faster if the precipitation crystals are large while this process becomes more difficult as the crystals are small.

It is used for this purpose:

- 1- Ashless few ash leaves.
- 2- the base of asbestos furnished on the inner bottom of the crucible.
- 3- crucible have porosity for filtration.

The fast filtration is best to prevent post filtration, which considers one of the types of contaminants.

5- Washing of precipitant:

Precipitation processes are usually carried out by solutions containing undesirable ions other than the ions that are to be precipitate.

Therefore, the precipitants that are mostly polluted so the precipitant washing process are necessary to remove the undesirable ions from the surface of the precipitation.

6- Drying and burning:

Drying processes are often carried out at temperatures below 250 C° almost between (110-150 C°) with a drying oven.

The burning process takes place at a temperature higher than 250 C° and up to 1200 C°. Here, a gas lamp is used, and then a burner with a temperature of 1200 C°.

7- Weight:

After finishing the process of burning or drying the precipitant, transfer the precipitate to the desiccator and leave it in the desiccator until it is cooled and have taken the laboratory temperature of.

General rules for obtaining a precipitate with large crystals:

- 1- precipitation from diluted solutions.
- 2- Make the acidic function of the solution suitable for the precipitation process because some materials prefer to be deposited in the acidic middle, while other materials prefer precipitation in the middle of the base.
- 3- Slow addition with continuous stirring for the purpose of avoiding the occurrence of a condition above the local saturation.
- 4- In the precipitation of hydroxides, it is preferable to use buffer solutions for the purpose of avoiding the development of high local PH centers and this can be avoided by precipitation from homogeneous solutions.
- 5- It is preferable to digest the precipitant if there is no risk of post-precipitation.
- 6- Wash the precipitant with a suitable solution do not react with the precipitate and do not increase its solubility and prefer to wash the precipitant on several small batches.

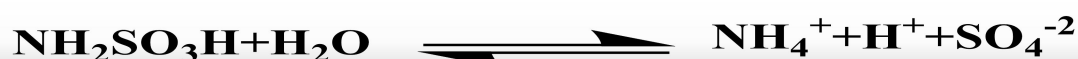
Precipitation From Homogenous Solution :

A- Is one of the precipitation methods in which the precipitation agent is composed slowly, uniformly and homogeneously in the solution either through the hydrolysis of a substance or the interaction of two substances with each other such as:

B- Oxalates are generated from the hydrolysis of dimethyl or diethyl oxalate, which is used as a precipitating agent for calcium.



C- Sulfates where sulfate ion can be generated from hydrolysis of sulfuric acid.



Advantages of precipitation of a homogeneous solution :

- 1- The crystals are slowly deposited so that they are large and regular so that they can be easily filtered.
- 2- The formed Precipitate is thermally stable so it is easy to dry.
- 3- The deposit shall be Empty of impurities.

Disadvantages of precipitation of a homogeneous solution :

- 1- Precipitation requires a homogeneous solution for a long time compared to normal Precipitation methods.
- 2- A high-cost method because Precipitators and solvents are used compared to what method normal Precipitation.

Contamination of precipitation

When there are substances with a foreign solution having a solubility product equal to or similar to the material to be precipitate if these strange materials precipitate with the material to be estimated, leading to contamination of the precipitation.

It gets in two cases:

1- Co precipitation

Includes all Contaminants obtained during the composition of the precipitation required to be analyzed and includes:

- A- True precipitation.
- B- Occlusion.
- C- Adsorption.

2- post precipitation

Which is the Contamination of the precipitation after its formation. The precipitate is composed of pure and solid crystals during precipitation and then it is contaminated by precipitator of another substance. This type of contamination can be eliminated by filtration immediately after precipitation.

Calculation

The calculations for weighted analysis Depends on two things:

- 1- The sample.
- 2- Material to be estimated.

The two are then linked to find the percentage and concentration of the material required.

$$\text{Percentage of the material } X(\%) = \frac{\text{weight of material}}{\text{weight of the sample}} \times 100$$

$$\text{weight of material (X)} = \frac{\text{gravimetric factor (G.F)}}{\text{weight precipitate (Wt. PPt)}}$$

Gravimetric Factor

Is the weight of the formula or the atomic weight of the material to be analyzed to the weight of the formula of the known weight precipitate and does not depend on the weight of the original form which is a fixed quantity.

$$\text{Gravimetric Factor (G.F)} = \frac{\text{A.wt or F.wt of (x)}}{\text{F.wt of ppt}} \times \frac{a}{b}$$

Note:

If there is a common atom in the ridge and the place (except oxygen), we must multiply the numerator or the place or both with a number or two different numbers so that the number of this common atom is equal in the numerator and denominator.

Q-1- What is the weight for the indium sample if you know that the percentage for the indium that precipitate in the form of In_2O_3 is (34.5%) , the gravimetric factor for the indium is(0.82) and the weight of the precipitate is (1g)? A.wt for In =114.8 , O=16.

$$\text{Wt of In} = \text{G.F} \times \text{Wt of In}_2\text{O}_3$$

$$= \frac{\text{A.Wt of In}}{\text{M.wt of In}_2\text{O}_3} \times \frac{2}{1} \times 1 \text{ g} = \frac{114.8 \times 2}{114.8 \times 2 + 16 \times 3} = 0.82 \text{ g}$$

$$\% \text{ In} = \frac{\text{Wt of In}}{\text{Wt of Sample}} \times 100 \quad \longrightarrow \quad 34.5 / 100 = \frac{0.82}{\text{Wt of sample}}$$

$$\text{Wt of Sample} = 2.3 \text{ g}$$

Q-2- If the weight of the lead sample is (3 g) was precipitated in the form of the PbSO_4 and the weight of the precipitate is (2 g) , what is the percentage for the precipitated substance in form of PbSO_4 ? If you know that the atomic weights of $\text{Pb} = 207.2$, $\text{S} = 32$ And $\text{O} = 16$

$$\text{Wt of Pb} = \text{G.F} \times \text{Wt of Pb SO}_4$$

$$= \frac{\text{A.Wt of Pb}}{\text{M.Wt of PbSO}_4} \times 2 \text{ g} = 1.36 \text{ g} , \text{ \%Pb} = \frac{\text{Wt of Pb}}{\text{Wt of Sample}} \times 100$$

$$\text{\% Pb} = \frac{1.36}{3} \times 100 = 45.5 \%$$

Q-3- What is the gravimetric factor for the substance that have a weight (0.37 g) if you know that the weight of the precipitate is (0.5) grams?

$$\text{Wt of Substance} = \text{G.F} \times \text{Wt of precipitate}$$

$$0.37 = \text{G.F} \times 0.5 \quad \longrightarrow \quad \text{G.F} = 0.74$$

Q-4- Calculate the weight of the chloride ion that was precipitated on the form of the silver chloride if you know that the weight of the precipitate is (0.53 g) ,atomic weights of the silver (108) and the chlorine (35.5)?

$$\text{Wt of Cl} = \text{G.F} \times \text{Wt of AgCl}$$

$$= \frac{\text{A.Wt of Cl}}{\text{M.Wt of AgCl}} \times 0.53 \text{g} = 0.13 \text{ g}$$

Q-5- A sample of non-pure sulfate salt 4 g was precipitated in the form of barium sulphate after it was dissolved in (100 ml) of distilled water and added the precipitated agent. Calculate the molarity of the sulfate ion in the non -pure salt ,If you know that the weight of the precipitate is (2.5 g), Atomic weights for S= 32, O= 16 and Ba= 137.3?

$$\text{Wt of SO}_4 = \text{G.F} \times \text{Wt of BaSO}_4$$

$$= \frac{\text{M.Wt of SO}_4}{\text{M.Wt of BaSO}_4} \times 2.5 \text{ g} \quad \longrightarrow \quad 1.02 \text{ g}$$

$$\text{M for SO}_4 = \frac{\text{Wt SO}_4}{\text{M.Wt SO}_4} \times \frac{1000}{\text{V(Ml)}} = 0.10$$

Example:

An impure sample of barium weighs (1 gm) that was precipitated as barium chloride after adding the precipitate, so the weight of the precipitate after burning processes (0.3 gm) Calculate the percentage of barium in the sample?

$$\% \text{Ba} = \text{Wt of Ba} / \text{Wt of Sample} \times 100$$

$$\text{Wt of Ba} = \text{G.F} \times \text{Wt of BaCl}_2$$

$$= 137.3 / 137.3 + 2 \times 35.5 \times 0.3 = 0.19 \text{ gm}$$

$$\% \text{Ba} = 0.19 / 1 \times 100 = 19.8\%$$

Experiment No.1

Determination the ratio of water of hydration in crystalised $\text{BaCl}_2 \cdot \text{XH}_2\text{O}$

Introduction

The water crystallization can be determined using sample weight change as a result of its water content ejection by heating it.

This method can be only used in the following cases:

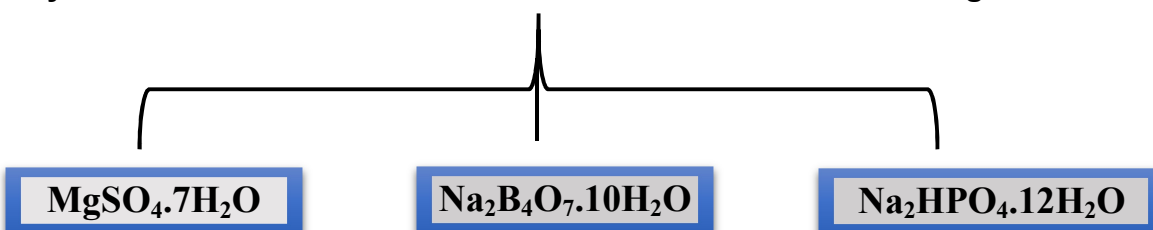
- 1- When water is the primary part to be volatilized only.
- 2- When the precipitation weight does not change as a result of Oxidation or hydrolysis.

The Barium chloride salt loses all crystallized water at a temperature range from (100-120) centigrade, high temperatures can be used for this purpose considering that the salt does not break at high temperatures up to (800-900) centigrade.



Barium dichloride crystals shows a little tendency to absorb water in the spaces of the molecules of the substance, which means that any change in weight represents a great precision water of crystallization.

Crystallization water can be determine for each of the following salts:



Materials and tools

1- Crucible.

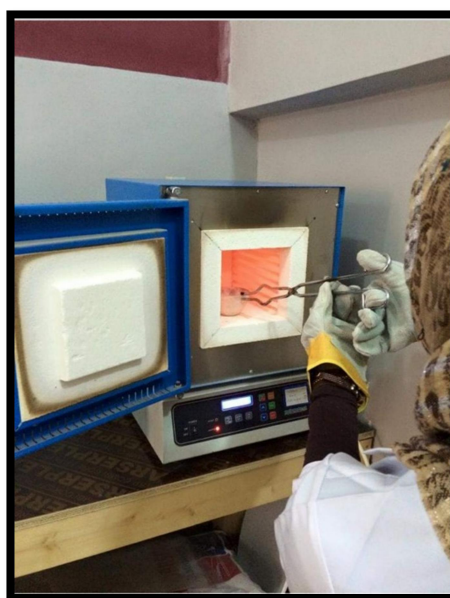
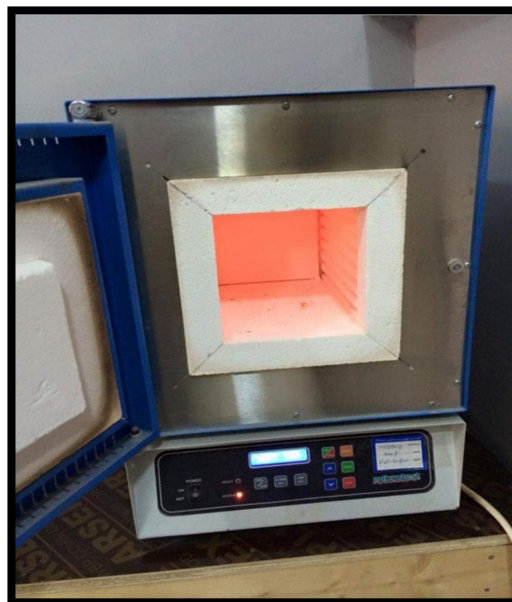
2- Desiccator.

3- Oven & Furnace.

4- Barium Chloride crystals $\text{BaCl}_2 \cdot \text{XH}_2\text{O}$.

Procedure

- 1- Heat the crucible inside the oven and then leave for a (30) minutes , cool in a desiccator for (15) minutes and then weigh after cooling.
- 2- Put in the crucible (0.5) gm of the sample and weigh it again (aqueous Barium Chloride $\text{BaCl}_2 \cdot \text{XH}_2\text{O}$).
- 3- Put the crucible inside the burning furnace for half an hour.
- 4- Let the crucible and its contents to be cooled in a desiccator and then weight it.
- 5- From weight loss after burning process, calculate the prcentage of water in aqueous Barium Chloride and Calculate the water molecule number in aqueous Barium Chloride ($\text{BaCl}_2 \cdot \text{XH}_2\text{O}$) .



Calculation:

$$\% \text{ of H}_2\text{O} = \text{Wt of H}_2\text{O} / \text{Wt of Sample} \times 100 \quad (\text{الملح المائي})$$

$$= \frac{\text{wt of analyte with crucible before heating} - \text{wt of analyte with crucible after heating}}{\text{wt of analyte with crucible before heating} - \text{wt of empty crucible}} \times 100$$

$$1 \text{ gm} - 0.8 \text{ gm} / 1 \text{ gm} \times 100 = 20 \%$$

to calculate the number of water molecules we used the following law :

$$\frac{\text{wt of salt (BaCl}_2 \cdot \text{XH}_2\text{O})}{\text{M.wt}} = \frac{\text{wt of salt (BaCl}_2\text{)}}{\text{M.wt}}$$

$$1 \text{ gm} / 208 + 18 \text{ X} = 0.8 \text{ gm} / 208$$

$$\text{X} = 2.3$$

Important notice

The wt of BaCl₂.XH₂O its coming from :

Wt salt with crucible before heating - wt of empty crucible

And the wt of BaCl₂ its coming from :

Wt salt with crucible after heating - wt of empty crucible

Discussion Questions

- 1- What are the conditions required in a materials that water crystallization can be determined? Give an example.
- 2- Why empty Crucibles need to be heated?
- 3- Define crystallized materials? Can some of them be used in chemical analyzes?
- 4- What is the purpose of the crucible cooling after heating in a desiccator?

Experiment No.2

Determination of Chloride as Silver Chloride

Introduction

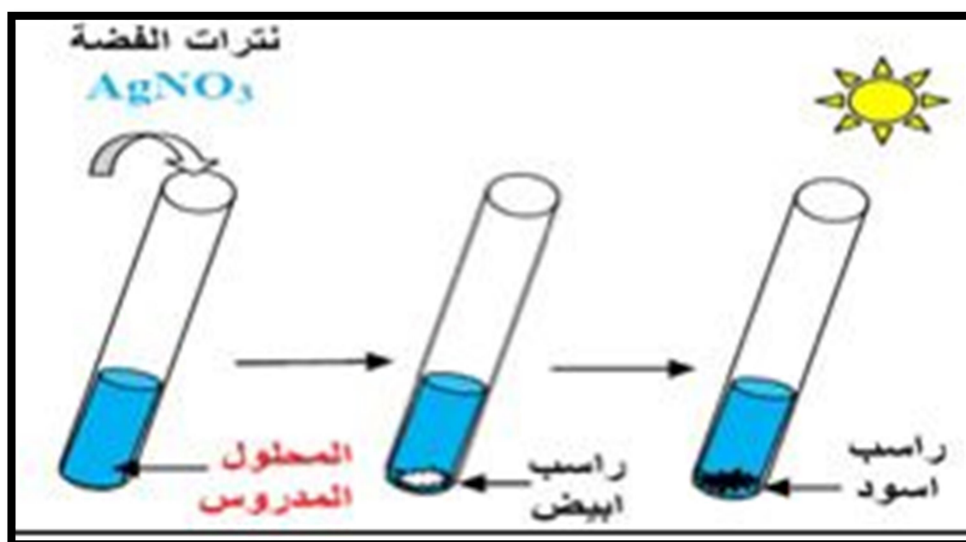
The chloride can be determined in a soluble form in water by adding an increase of silver nitrate (AgNO_3) solution to the chloride solution diluted with diluted nitric acid (HNO_3) and then the precipitated (AgCl) weight after filtration, washing, and drying :



The purpose of nitric acid addition to the solution is to prevent the deposition of some silver salts such as Silver carbonate (Ag_2CO_3) and silver phosphate (Ag_3PO_4), which are deposited from the neutral or base solution. If the CO_3^{2-} and PO_4^{3-} ions are found in the solution, the acid solution Obtainable filtrate (AgCl) easily. The heating and stirring of the

solution containing AgCl; precipitates the colloidal particles formed in principle and the solution becomes clear after it has been reversed.

The properties of precipitate are white salt that is low in solubility and water solubility (about 0.0014 g / L) at room temperature but increases with temperature increase. This salt is sensitive to light and to reducing factors.

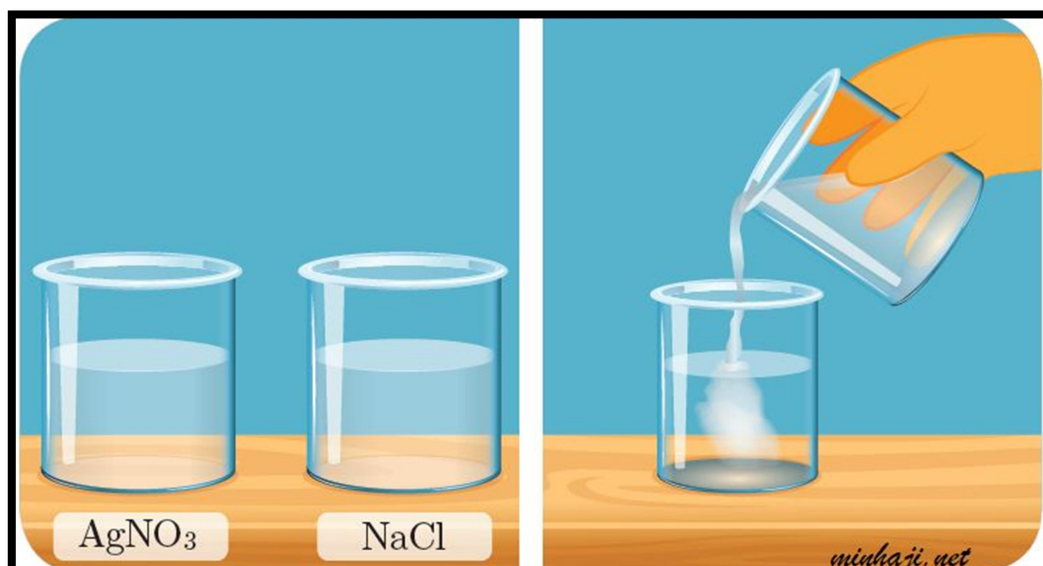
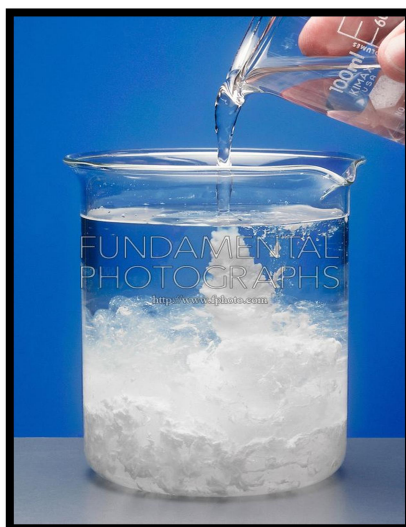
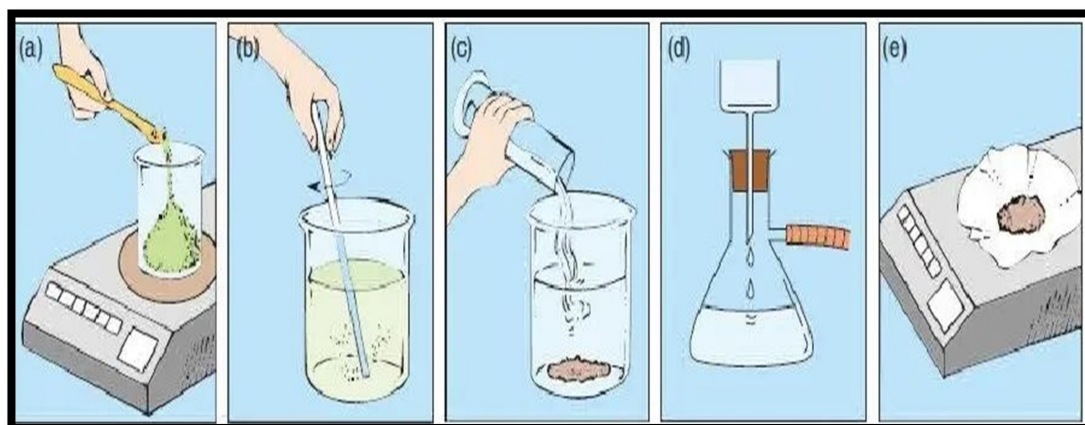


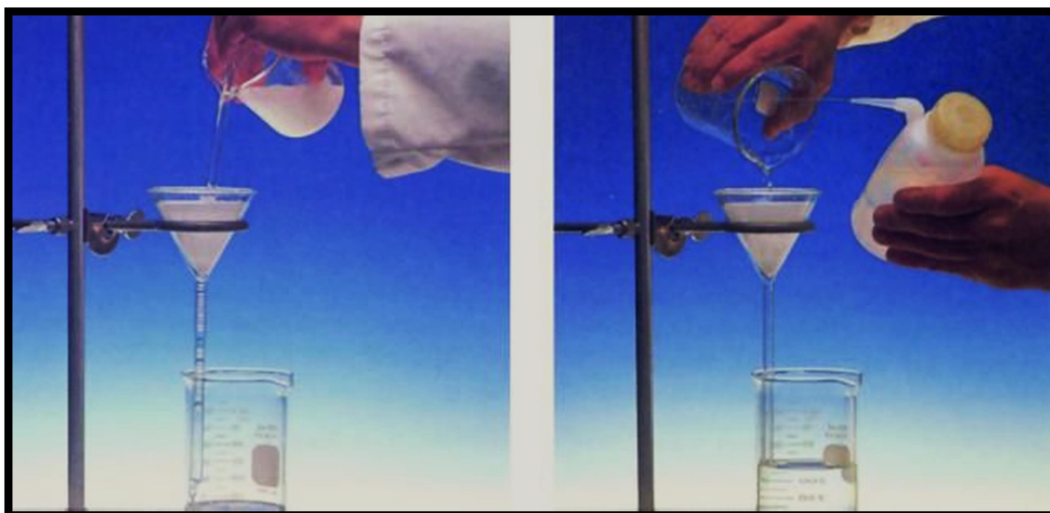
Materials

- 1 - Dissolved chloride salt in water.
- 2- Diluted nitric acid (1: 1).
- 3 - Silver nitrate solution (0.1N)
- 4- Diluted Nitric acid (0.015 N).

Procedure

- 1- Accurately weigh (1.0 g) of chloride sample. Transfer weight to a Baker (250 ml) with a glass rod and glass watch as a cover, add about 75 ml of distilled water and dissolved salt by stirring.
- 2- Add about (1.0 mL) of diluted nitric acid (1: 1). The experiment should be conducted in a dark place (dim light).
- 3- Deposition of chloride in the form of silver chloride is achieved by adding about (20 mL) of silver nitrates (0.1 M), the addition should be slow with continuous stirring of the cold solution.
- 4- Heat the mixture quickly until it reaches near boiling point with continuous stirring.
- 5- Remove the baker from the heat and leave the precipitate stabilized (2 - 3 minutes), make sure that the precipitation process is complete by adding a few drops of silver nitrate solution to the pure liquid. If no additional deposit is shown, the baker is placed in a dark place and the solution is left to settle for one hour before filter.
- 6- Prepare the filtration apparatus, filter the solution, wash the precipitate (on the filter paper) with cold diluted nitric acid (0.015N, 5ml) and add small amount until the silver nitrate is removed and then wash the precipitate by a little quantity of distilled water .
- 7- Dry the precipitate at 110-130 oc and then weigh it. Calculate the weight of chloride and then the percentage.





Calculation

Wt of Sample = 1.0 gm

Wt of precipitate AgCl = 0.6 gm

If you know A.wt of Ag= 108 , Cl = 35.5

Calculate the percentage of Cl in the sample ?

% Cl = Wt of Cl / Wt of Sample \times 100

Wt of Cl = G.F \times Wt of AgCl

$$= 35.5/108+35.5 \times 0.6 = 0.247 \times 0.6 = 0.148\text{gm}$$

$$\% \text{cl} = 0.148 / 1.0 \times 100 = 14.82 \%$$

Discussion

- 1- Why the experiments are done in a dark place?
- 2- The diluted nitric acid is used as a washing agent for the residue and does not use diluted hydrochloric acid?

- 3- What are the specifications of the formed silver chloride precipitate?
- 4- The precipitate filtration is done at room temperature or less?
- 5- Calculate the weight of silver chloride residue that can be obtained from a solution (100 mL) dissolved in (0.2 g) of magnesium chloride?
- 6- Why wash the precipitate with distilled water before the drying ?