Chapter one: QUALITATIVE ANALYSIS.

Introduction

Qualitative analysis is the identifications of cations and anions in given samples of substances. Compounds can only be identified by carrying out sample tests which identify the ions present. The identification follows a scientific procedure in which conclusions (or deductions) are made on observations from tests or experiments.

Qualitative analysis reactions are usually carried out in tests tubes using small quantities of reagents, even if reagents are in excess. Large quantities of reagents are wasteful and small quantities usually give observable and reliable results. Large quantities can also result in dangerous reaction which cannot be controlled and give unreliable results or wrong observations and conclusions.

This section forms question number two of chemistry paper 545/3 or paper 545/4. It is mainly based on the chemistry of salts which is learnt in senior two. Students are therefore advised to revise the chemistry of salts in order to present quality work in this section. Safety is the most important aspect for students of chemistry in the laboratory; therefore students should take caution and responsibility. Students should revise all rules and regulations of the laboratory before performing practical.

Qualitative analysis requires a student to carry out a number of sample tests on one or more substances mixed, the observation is recorded from which deduction about the substance is made.

Unless the quantity is specified, volumes of about 1cm³ for liquids and almost 0.2g for solids are sufficient to use in qualitative test tube reactions. The reactions occur mainly in aqueous solution. The results can positive or negative. The positive indicates the expected result is present and negative test indicates that the expected result is absent.

The reactions with both dissolved and undissolved ions produce various precipitates and gases. Each ion has its own particular properties which can be used to identify it.

The tests are qualitative and require careful observations deductive and thinking. The qualitative analysis in this case will be limited to the following ions.

The cations considered at this level include:

White substance

- 1. Zinc ions Zn^{2+}
- 2. Lead (ii) ions Pb²⁺
- 3. Aluminium ions Al³⁺
- 4. Calcium ions Ca²⁺
- 5. Ammonium ions NH₄⁺
- 6. Magnesium ions Mg²⁺

Coloured substances

- 7. Copper (ii) ions Cu²⁺
- 8. Iron (ii) ions Fe^{2+}
- 9. Iron (iii) ions Fe³⁺

At this level, the anions considered include:

- 1. Sulphate ions SO₄²-
- 2. Chloride ions Cl⁻
- 3. Carbonate ions CO₃²-
- 4. Nitrate ions NO₃
- 5. Iodide ions I
- 6. Bromide Br
- 7. Sulphite ions SO_3^{2-}

During qualitative analysis there are three major phase namely.

- > Preliminary tests which give a clue about the composition the compound given.
- Precipitation reactions of cations (confirmatory tests for cations)
- Precipitation reactions of anions (confirmatory tests for anions)

PRELIMINARY TESTS.

These are tests carried out to give you a clue of the possible cations and anions present in a given substance. The word probably is used. These include, colour of the substance, smell and solubility of the substance in water, action of heat, formation of precipitates and gases. In some qualitative analysis exercises, a short list of ions can be given as present and in this case the ions identified from the preliminary tests alone.

However, in others, preliminary tests alone are not sufficient to completely identify the ions. Further tests would have to be performed before the ion could be clearly identified; an additional or extra test which identifies an anion or cation conclusively is known as **confirmatory test**. A confirmatory test clearly identifies an ion. There can be more than one confirmatory test for a particular ion. On such test is sufficient to confirm the ion.

The following notes are useful in the identification of ions.

- a) **Colour**: cations in their solid or dissolved aqueous solution have characteristic colours.
 - (i) Mg^{2+} , Ca^{2+} , Ba^{2+} , Al^{3+} , Zn^{2+} , NH_4^+ , and Pb^{2+} are colourless or white solid.
 - (ii) Cu^{2+} is blue.
 - (iii) Fe²⁺ is green
 - (iv) Fe^{3+} is yellow or brown.
 - (v) NO_3^- , SO_4^{2-} , Cl^- , or hydrated compound is a crystalline substance.

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b) Action of heat on solids

When solids are heated they decompose and gases are usually given off. These gases usually give useful information about the anions present in the unknown substance. A wide range of different gases can be given off and all must be identified with physical and chemical tests. Most gases can be identified by one or a combination of the following properties.

The colour, smell, effect on litmus paper, action on splint and the usual confirmatory tests can carried out on some gases e.g. for SO₂, is a colourless gas with an irritating smell, it turns moist blue litmus paper red and then bleached, and turns acidified dichromate solution from orange to green.

Procedure

- ➤ Put a small portion of the unknown substance in a clean dry test tube.
- ➤ Heat gently at first the more strongly.
- ➤ Observe the colour changes on the solid and observe the gases and vapours given off and identify them by their colours, smell, effect on litmus and their confirmatory test.
- Record the colour of the residues left after the heating which is usually an oxide of the metal and some of the metal oxides have certain characteristic colours. These can also be used to give useful information about the cation present in the unknown substance.

The table below shows the possible observations and deductions that can made when a substance is heated.

Observations	Deduction
A colourless liquid forms on the upper parts of the	Water of crystallization
test tube which turns white anhydrous copper (II)	Substance is hydrated, HCO ₃ -, OH-,
sulphate blue and blue cobalt (II) chloride paper	or HSO ₄ probably present.
pink.	
A reddish brown gas with an irritating smell and	NO ₂ is evolved probably NO ₃
turns moist blue litmus paper red is given off	present
A colourless gas which turns moist blue litmus	SO ₂ is evolved probably SO ₃ ²⁻ or
paper red and acidified potassium dichromate from	SO ₄ ²⁻ present
orange to green	
A colourless gas with a chocking/pungent smell	NH ₃ is given off and NH ₄ ⁺ is present
which turns moist red litmus paper blue and forms	
dense white fumes with Conc. HCI	
A colourless gas which turns damp blue litmus	CO ₂ is given off and CO ₃ ²⁻ or HCO ₃ ⁻
paper red and lime water milky	present
Colourless gas with an irritating smell, turns blue	HCl gas probably Cl ⁻ present
litmus paper red and forms dense white fumes with	
Conc. NH ₃	

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Greenish yellow gas with a very sharp irritating	Cl ₂ gas evolved probably Cl ⁻ present
smell bleaches damp litmus paper.	
white (smoky) fumes with chocking smell turns	SO ₃ gas evolved probably SO ₄ ²⁻ or
blue litmus paper red and forms a white precipitate	HSO ₄ present
with BaCI ₂ or Ba(NO ₃) solution	
Colour of the residue (Observation)	Possible metal oxide formed
	(Deduction)
The residue is black	Cu, FeO, Fe ₃ O ₄ is formed probably
	Cu ²⁺ , Fe ²⁺ , Fe ³⁺ , probably present.
Yellow residue when hot and turns white on cooling	ZnO is formed probably Zn ²⁺ present
Red-brown residue when hot and turns yellow when	PbO is formed probably Pb ²⁺ present
cold	
Pale green solid leaves a reddish brown residue	Fe ₂ O ₃ formed, Fe ²⁺ salt oxidized to
after heating	Fe^{3+}
Green solid leaves black solid after heating	CuO or FeO formed hence Cu ²⁺ or
	Fe ²⁺ present
A yellow substance forms a reddish brown residue	Fe ₂ O ₃ formed probably Fe ³⁺ present
White sublimate forms on the walls of the test tube	NH ₄ ⁺ probably present
Cracking sound heard (decrepitation occurs)	Possibly Pb(NO ₃) ₂ present.

Notes

- ✓ Litmus paper must be damp
- ✓ Which testing for gases with litmus paper, do not touch the mouth of the inner walls of the test tube. Hold the test paper in the mouth of test tube without touching the tube.
- ✓ Have the litmus paper, delivery tube, and other necessary apparatus ready before a reagent is poured onto a test substance. Any gases produced may diffuse away quickly.\

General rules on solubility of common salts.

This information can be used to predict formation of precipitates.

Salts	Soluble	Insoluble
Nitrates	All nitrates are soluble in	None
	water	
Chlorides	All chlorides are soluble	-lead (II) chloride (only
	except lead (II) chloride and	soluble in hot water)
	silver chloride	-silver chloride
Sulphates	All sulphates are soluble	-lead (II) sulphate
	except lead (II) sulphate,	-barium sulphate
	barium sulphate and calcium	-calcium sulphate is only
	sulphate	sparingly soluble
Carbonates	Carbonates of group 1 and	All carbonates are insoluble
	ammonium carbonate	except those group 1 and
		ammonium carbonate
Hydrogen carbonates	All are soluble	None
Sulphites	Sulphites of group 1 and	All sulphites are insoluble
	ammonium sulphite	except those of group 1 and
		ammonium sulphite

NB:

- > Sparingly soluble salts in most cases are a mixture of a soluble and an insoluble salt.

 During filtration, a soluble salt forms the filtrate and an insoluble salt forms the residue.
- ➤ Dilute acids are used to dissolve solids that do not dissolve in water. In case the reaction is slow warm the mixture.

Observation	Deduction
Solid dissolves in acid with bubbles	An insoluble salt of CO ₃ ²⁻ present
of a colourless gas that forms a white	
precipitate with lime water (turns	
lime water milky)	
Substance dissolves in an acid on	An insoluble compound
warming	Probably O ²⁻ or Cl ⁻ present

Group Activity

- 1. You are provided with the following substances labeled
 - A $Pb(NO_3)_2$
 - B ZnCO₃
 - C FeSO₄.7H₂O
 - D CuSO₄.5H₂O
 - E NH₄Cl
- a) Heat each solid using a small portion strongly until no further change occurs on the solid
- b) Observe the colour changes on each solid and identify the gases by colour, smell, effect on litmus paper and a confirmatory test if any.

PRECIPITATION REACTIONS AND CONFIRMATORY TESTS OF THE CATIONS

Precipitation is due to formation of insoluble metal hydroxides, chlorides, sulphates.

Sodium hydroxide and ammonium hydroxide (ammonia solution) are commonly used to detect cations in solutions by precipitating them out as insoluble metal hydroxide.

Some insoluble hydroxides will dissolve when excess reagent is added. **Therefore, it is** important to record the observation with few drops and when excess reagent is added.

In case the precipitate dissolves then record the colour of the solution formed.

Examples of these reactions include;

a) Copper (II) ions Cu²⁺

(i) Using sodium hydroxide solution, a blue precipitate insoluble in excess is formed.

Equation: $Cu^{2+}_{(aq)} + 2OH^{-}_{(aq)} \longrightarrow Cu(OH)_{2(s)}$

(ii) (Confirmatory test): using aqueous ammonia solution, a blue precipitate soluble in excess forming a deep blue solution.

Equation: $Cu^{2+}_{(aq)} + 2OH^{-}_{(aq)} \longrightarrow Cu(OH)_{2(s)}$

In excess ammonia solution, the blue precipitate dissolves forming a deep blue solution. The deep blue solution is a complex ion, tetra amine copper (ii) ions.

Equation: $Cu(OH)_{2(s)} + 4NH_{3(aq)} \longrightarrow Cu(NH_3)_4^{2+}_{(aq)} + 2OH_{(aq)}^{-}$

b) Iron (II) ions Fe²⁺

(i) Using sodium hydroxide solution, a dirty green precipitate insoluble in excess is formed, turns brown on standing. This is because oxygen in air oxidizes iron (II) ions to iron (III) ions.

Equation: $Fe^{2+}_{(aq)} + 2OH^{-}_{(aq)} \longrightarrow Fe(OH)_{2(s)}$

(ii) Using ammonia solution forms a dirty green precipitate insoluble is excess, turns brown on standing due to aerial oxidation of iron (II) ions to iron (III) ions.

Equation: $Fe^{2+}_{(aq)} + 2OH^{-}_{(aq)} \longrightarrow Fe(OH)_{2(s)}$

(iii) (Confirmatory test): using potassium hexacyanoferate (III) solution, a dark blue precipitate is formed.

c) Iron (III) ions Fe³⁺

(i) Using sodium hydroxide solution, a brown precipitate insoluble in excess is formed.

Equation: $Fe^{3+}_{(aq)} + 3OH^{-}_{(aq)} \longrightarrow Fe(OH)_{3(s)}$

(ii) Using ammonia solution, a brown precipitate insoluble in excess is formed.

Equation: $Fe^{3+}_{(aq)} + 3OH^{-}_{(aq)} \longrightarrow Fe(OH)_{3(s)}$

(iii) (**Confirmatory test**): Using potassium hexacyanoferate (II) solution, a dark blue precipitate is formed.

(iv) (**Confirmatory test**): Using potassium thiocyanate solution; a deep red solution is formed.

d) Zinc ions Zn²⁺

(i) Using sodium hydroxide solution, a white precipitate soluble in excess forming a colourless solution.

Equation:
$$Zn^{2+}_{(aq)} + 2OH^{-}_{(aq)} \longrightarrow Zn(OH)_{2(s)}$$

The white precipitate zinc hydroxide dissolves because it is amphoteric therefore reacts with sodium hydroxide forming a soluble complex ion, Called zincate ion.

Equation:
$$Zn(OH)_{2(s)} + 2OH^{-}_{(aq)} \longrightarrow Zn(OH)_4^{2-}_{(aq)}$$

(ii) (Confirmatory test): using ammonia solution, a white precipitate soluble in excess forming a colourless solution.

Equation:
$$Zn^{2+}_{(aq)} + 2OH^{-}_{(aq)} \longrightarrow Zn(OH)_{2(s)}$$

The precipitate dissolves because zinc hydroxide reacts with ammonia forming a soluble complex ion, called tetra amine zinc ions.

Equation:
$$Zn(OH)_{2(s)} + 4NH_{3(aq)} \longrightarrow Zn(NH_3)_4^{2+}{}_{(aq)} + 2OH_{(aq)}^{-}$$

e) Lead (II) ions Pb²⁺

(i) Using sodium hydroxide solution, a white precipitate which dissolves in excess forming a colourless solution.

Equation:
$$Pb^{2+}_{(aq)} + 2OH^{-}_{(aq)} \longrightarrow Pb(OH)_{2(s)}$$

The precipitate dissolves because lead (II) hydroxide is amphoteric and therefore

reacts with sodium hydroxide solution forming a soluble complex ion called plumbate ion.

Equation:
$$Pb(OH)_{2(s)} + 2OH^{-}_{(aq)} \longrightarrow Pb(OH)_4^{2-}_{(aq)}$$

(ii) Using ammonia solution, a white precipitate insoluble in excess ammonia is formed.

Equation:
$$Pb^{2+}_{(aq)} + 2OH^{-}_{(aq)} \longrightarrow Pb(OH)_{2(s)}$$

(iii) (**Confirmatory test**): using potassium iodide solution, a bright yellow precipitate is formed.

Equation:
$$Pb^{2+}_{(aq)} + 2I_{(aq)} \longrightarrow PbI_{2(s)}$$

(iv) (Confirmatory test): using dilute hydrochloric acid and warm, a white precipitate soluble on warming and re-appears on cooling. Equation: $Pb^{2+}_{(aq)} + 2CI_{(aq)} \longrightarrow PCI_{2(s)}$

(v) (**Confirmatory test**): using potassium dichromate solution, a yellow precipitate is formed.

Equation:
$$Pb^{2+}_{(aq)} + CrO_4^{2-}_{(aq)} \longrightarrow PbCrO_{4(s)}$$

(vi) Using dilute sulphuric acid, a white precipitate is formed. Equation: $Pb^{2+}_{(aq)} + SO_4^{2-}_{(aq)} \longrightarrow PbSO_{4(s)}$

(vii) Using sodium carbonate solution, a white precipitate is formed. Equation: $Pb^{2+}_{(aq)} + CO_3^{2-}_{(aq)} \longrightarrow PbCO_{3(s)}$

f) Aluminium ions Al³⁺

(i) Using sodium hydroxide solution, a white precipitate which dissolves in excess forming a colourless solution.

Equation: $Al^{3+}_{(aq)} + 3OH^{-}_{(aq)} \longrightarrow Al(OH)_{(s)}$

The precipitate dissolves because the Aluminium hydroxide is amphoteric and reacts with sodium hydroxide forming a soluble complex ion called Aluminate ion.

Equation: $Al(OH)_{3(S)} + OH^{-}_{(aq)} \longrightarrow Al(OH)_{4^{-}(aq)}$

- (ii) Using ammonia solution, a white precipitate insoluble in excess is formed. Equation: $Al^{3+}_{(aq)} + 3OH^{-}_{(aq)} \longrightarrow Al(OH)_{3(s)}$
- (iii) (**Confirmatory test**), 2 drops of litmus solution, test solution if not acidic add dilute hydrochloric acid and then add dilute ammonia solution until the solution is just acidic.

Observation: Blue lake forms by absorbing Al(OH)₃ precipitate in aqueous NH₃ leaving the solution almost colourless.

g) Ammonium ions NH₄⁺

Add sodium hydroxide and heat the mixture. Smell the mixture, Ammonium ion form no precipitate. On warming, acolourless gas with a chocking smell and turns moist red litmus paper blue is given off (**This is the Confirmatory test for NH**₄⁺)

Equation: $NH_4^+_{(aq)} + OH^-_{(aq)} \longrightarrow NH_{3(g)} + H_2O_{(l)}$

h) Calcium ions Ca²⁺, Barium ions Ba²⁺ and Magnesium ion Mg²⁺

Using sodium hydroxide solution, these ions form a white precipitate insoluble in excess.

$$Ca^{2+}_{(aq)} + 2OH_{(aq)} \longrightarrow Ca(OH)_{2(s)}$$

$$Ba^{2+}_{(aq)} + 2OH^{-}_{(aq)} \longrightarrow Ba(OH)_{2(s)}$$

$$Mg^{2+}_{(aq)} + 2OH^{-}_{(aq)} \longrightarrow Mg(OH)_{2(s)}$$

NB:

- ❖ In carrying out tests, it is important to use known reagents and conditions that give characteristics results such as formation of gases and precipitates.
- ❖ The test must be practical and the student should be familiar with the expected result in order to make correct conclusions.
- ❖ Knowledge of the reagent and expected observations are important in order to make conclusions. A student who does not know what the reagent is testing for cannot make correct conclusions from the observations. Therefore knowledge of the test reagents, what they test for and the expected result is important in identifying ions e.g. When dilute sodium hydroxide is added to a solid substance and the mixture warmed, if a pungent smelling gas is produced, it indicates the presence of NH₄⁺ ions, if both the test and expected observation is not known, the correct conclusion of NH₄⁺ cannot be drawn easily.

Forms of unknowns and tests

The substance that appears in practical examinations may be single compounds or mixture of compounds, and may contain more than two ions. Tests can be performed on unknown solutions, solids or their mixture. Insoluble solids can be dissolved with dilute or concentrated acids. The acids react with the insoluble substance to form their soluble ions on which tests can be carried out in solution.

Example: group activity

Solution A contains Pb²⁺or AI³⁺ or Zn²⁺.

Carryout the following tests and identify the ions present.

- (a) Add dil. NaOH in drops and the in excess.
- (b) Add dil. NH₃ in drops and then in excess
- (c) Add dil. HCl, heat the mixture well and allow it to cool.
- (d) Carry out a test a test of your own choice to confirm the cation present in A.

Record the results obtained in the table below.

	Tests	Observations	Deductions
(a)	Dil. NaOH was added to A drop-wise until in excess.		
(b)	Dil. NH ₃ was added to A drop-wise until in excess.		

(c)	Dil. HCl was added to A, the mixture was heated and then cooled.	
(d)	Carry out the test to confirm the cation in A	

Note:

- ✓ When an unknown is given and it is required to find the ions in it, it requires a systematic elimination of the ions through preliminary and further tests. A confirmatory test may be necessary to determine the ion conclusively. Specific tests are carried out and the ions present deducted from the results.
- ✓ This requires a student to have good knowledge of what the reagents test for and the conditions of reactions and what observations to expect.
- ✓ The tests are always based on small amount of reactants, small quantities give observations and reliable results than the large quantities.

v

Marking logical observations and deductions.

- > Care should be taken to add initially one or two drops of the reagent and then excess if that is what is required.
- ➤ Observations of the effect of adding the reagents drop-wise and then in excess can lead to the drawing of correct observations. For example, in a test, addition of drops of dilute sodium hydroxide solution to a given solution may produce a precipitate, the correct deduction is that Mg²+, Ca²+, Zn²+, Pb²+ and Al³+ ions are present. However on addition of excess alkali, the precipitate dissolves. The correct deduction now is Zn²+,Pb²+ and Al³+ ions are present. Ca²+ and Mg²+ ions are eliminated because Mg(OH)₂ and Ca(OH)₂ precipitates do not dissolve in excess sodium hydroxide solution. They are not amphoteric.
- Further tests can be carried out to discover which of the ions, Zn²⁺,Pb²⁺ and Al³⁺ ions are present.
- A deduction can therefore cover one or more ions and you must record all the ions which are pointed out by a certain observation.

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General solubility rules.

- (a) Water soluble compounds
 - NO₃-, all nitrates of common metals are soluble in water.
 - ❖ All salts of Na⁺, K⁺ and NH₄⁺ are soluble in water.
 - ❖ SO₄²-, all sulphates are soluble except CaSO₄, PbSO₄, and BaSO₄.
 - ♣ All chloride (Cl⁻), bromide (Br⁻) and iodide (I⁻) are soluble except those of Pb²+ and Ag⁺.
 - ❖ HCO₃-, all hydrogen carbonates are soluble.
- (b) Mainly water insoluble compounds.
 - ❖ CO₃²⁻, all carbonates are insoluble except those of group 1 metals and (NH₄)₂CO₃.
 - ❖ All oxides (O²-) and hydroxides are insoluble except those of group 1 metals, Na⁺, K⁺ and NH₄⁺.
 - SO₃²-, all sulphites are insoluble except those group 1 elements and (NH₄)₂SO₃.

NB: These rules are useful in prediction formation of precipitate in qualitative analysis.

Formation of precipitates.

Precipitates are formed when two solutions containing ions that form an insoluble compound are mixed

The ions come together to form the precipitates or crystals e.g. lead (II) sulphate precipitate from a mixture of lead (II) nitrate solution and aqueous solution of a sulphate, e.g. aqueous sodium sulphate.

$$Pb(NO_3)_2 \ {}_{(aq)} + Na_2SO_4 \ {}_{(aq)} \longrightarrow PbSO_4 \ {}_{(s)} + 2NaNO_4 \ {}_{(aq)}$$

If the cation is known, e.g. Pb^{2+} in this case the anion present can be predicted. On the other hand if the anion is known the cation can be predicted.

The prediction is based general knowledge of solubility of substance in water. It is important to state the colour and nature of the precipitate formed.

Insoluble salts however precipitate from solution. Note the colour and nature of the precipitate.

Example

Aqueous solutions of the following salts are mixed together in the test tube.

- (a) $Pb(NO_3)_{2 \text{ (aq)}}$ and $Na_2SO_4 \text{ (aq)}$
- (b) $ZnCl_{2 (aq)}$ and $Zn(NO_4)_{2 (aq)}$
- (c) Na_2CO_3 (aq) and $Zn(NO_3)_2$ (aq)
- (d) KI $_{(aq)}$ and Pb(NO₃)_{2 $_{(aq)}$}

Solution:

(a)
$$Pb(NO_3)_{2 (aq)}$$
 $\longrightarrow Pb^{2+}_{(aq)} + 2NO_3^{-}_{(aq)}$
 $Na_2SO_4_{(aq)}$ $\longrightarrow 2Na^+_{(aq)} + SO_4^{2-}_{(aq)}$

All nitrates are soluble, therefore NaNO₃, cannot be precipitated but all sulphates are soluble excess PbSO₄,BaSO₄ and CaSO₄. Therefore PbSO₄ will be precipitated.

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- (b) No precipitate will be formed, because all the ions present, Zn²⁺, Cl⁻, and NO₃⁻, no cation and anion pair to form an insoluble salt. Possible combinations of Zn(NO₃)₂ and ZnCl₂ are all soluble salts.
- (c) The ions present are Na⁺, CO₃²⁻, Zn²⁺ and NO₃⁻. CO₃²⁻ all insoluble except only those of group 1 and (NH₄)₂CO₃. Therefore ZnCO₃ will be precipitated because it is insoluble in water.

These generalizations are useful in predicting formation of precipitates in qualitative analysis.

For example, if the addition of an aqueous solution of lead (II) nitrate to an unknown solution produces a white precipitate, it indicates the unknown solution cations one or more types of anions which form an insoluble compound with Pb^{2+} ions. These are Cl^- , SO_4^{2-} , SO_3^{2-} or CO_3^{2-} .

The following generalizations can be derived from the results of the above experiment.

- ✓ Any soluble salt and any of the three soluble carbonates, Na₂CO₃, K₂CO₃ and (NH₄)₂CO₃ forms a white precipitate of an insoluble metal carbonate. Because all carbonates are insoluble except those of group 1 and ammonium carbonate.
- ✓ Any soluble salt + lead (II) nitrate forms a precipitate of an insoluble lead (II) salt because most of lead (II) salts are insoluble except lead (II) nitrate. The precipitates are identified by their colours.
- ✓ Any barium salt and any soluble sulphate forms a white precipitate of barium sulphate.
- ✓ Any soluble sulphite and any soluble barium salt forms a white precipitate of barium sulphite.
- ✓ Any soluble chloride and silver nitrate solution produces a white precipitate if silver chloride.

These generalizations will help you answer many related questions on theory paper.

Sample question

Aqueous solution of the following salts were mixed together in the test tubes. State what was observed and write the ionic equations for the reaction that took place in each case.

- (a) CaCl_{2(aq)} and NaCO_{3(aq)}
- (b) NaI $_{(aq)}$ and Pb(NO3)2 $_{(aq)}$
- (c) $Ba(NO_3)_2$ and $H_2SO_4\ \mbox{\tiny (aq)}$

Solution:

(a) Observation: white precipitate is formed.

Equation: $Ca^{2+}_{(aq)} + CO_3^{2-}_{(aq)} \longrightarrow CaCO_3_{(s)}$

(b) Observation: yellow precipitate is formed

Equation: $Pb^{2+}_{(aq)} + 2I^{-}_{(aq)} \longrightarrow PbI_{2(s)}$

(c) Observation: white precipitate is formed

Equation: $Ba^{2+}_{(aq)} + SO_4^{2-}_{(aq)} \longrightarrow BaSO_{4(s)}$

Summary of the reactions of the cations

Reagent	Observation	Deduction
Sodium hydroxide	A white precipitate insoluble in excess	Ca ²⁺ , Ba ²⁺ , or Mg ²⁺
Solution	sodium hydroxide solution	Present
	A white precipitate soluble in excess	Zn ²⁺ , AI ³⁺ or Pb ²⁺
	forming a colourless solution	Present
	A blue precipitate insoluble in excess	Cu ²⁺ present
	sodium hydroxide solution	
	A dirty green precipitate insoluble in	Fe ²⁺ present
	excess sodium hydroxide solution	
	A brown precipitate insoluble in	Fe ³⁺ present
	excess sodium hydroxide solution	
	No observable change, on warming, a	NH ₃ gas evolved
	colourless gas with a pungent chocking	NH ₄ ⁺ present
	smell, turns damp red litmus paper	
	blue and forms dense white fumes with	
	Con. HCI	
Ammonia	A white precipitate insoluble in excess	Pb^{2+} , Mg^{2+} or AI^{3+}
Solution	ammonia solution	Present
	A white precipitate soluble in excess	Zn ²⁺ present
	ammonia solution forming a colourless	
	solution	
	A dirty green precipitate insoluble in	Fe ²⁺ present
	excess ammonia solution	
	A blue precipitate soluble in excess in	Cu ²⁺ present
	excess ammonia solution forming a	
	deep blue solution	
	A brown precipitate insoluble in	Fe ³⁺ present
	excess ammonia solution	
Potassium	A bright yellow precipitate is formed	Pb ²⁺ present
Iodide solution	No observable change	Al ³⁺ present
	White precipitate in a brown solution	Cu ²⁺ present
Potassium	A dark blue precipitate formed	Fe ³⁺ present
Hexacyanoferrate(II)	Brown precipitate is formed	Cu ²⁺ presents
solution		-
Potassium	Deep red solution formed	Fe ³⁺ present
thiocyanate solution		

Potassium	A dark blue precipitate formed	Fe ²⁺ present
hexacyanoferrate (III)		
solution		
Dilute sulphuric acid	A white precipitate formed	Pb ²⁺ , Ba ²⁺ or Ca ²⁺
		Present
	No observable change	Mg ²⁺ present
Aluminium thiocynanate	Deep red solution	Fe ³⁺ present
Hydrogen peroxide and	Effervescence occurs and a colourless	O ₂ , evolved
warm	gas which relights a glowing splint.	Fe ²⁺ present.
	Blue solution turns brown	
Copper (II) sulphate and	Solution changes from blue to	Cu ²⁺ present
zinc powder	colourless.	
	A brown solid is formed	

ACIDIFYING THE FILTRATE

Back ground information

This test involves a reaction between two cations in solution with a reagent mostly sodium hydroxide or ammonia solution.

One of the cations forms a precipitate which does not dissolve and the other forms a precipitate which dissolves in excess reagents forming a soluble complex.

On filtering, the insoluble precipitate forms the residue on the filter paper while the soluble complex drops down and is collected as the filtrate.

Acidifying this filtrate involves freeing the cation in the complex. The acid is added drop-wise until the solution is just acidic. This is indicated by formation of a precipitate until it just dissolves, immediately stop adding the acid because the solution is now just acidic. Adding excess acid may affect your results.

In most cases a **precipitate appears until it just dissolves** as the acid is being added drop-wise to the filtrate.

Note

- ❖ Always add excess reagent (NaOH or NH4OH) in order to dissolve the soluble precipitate.
- ❖ Shake if necessary for the soluble precipitate to dissolve.

If the above is not done, the filtrate collected will not contain the complex and acidifying it will show no change and the solution obtained will give negative tests only.

Worked out examples.

(a) Using sodium hydroxide

This reagent is used to separate two cations, one of which forms a precipitate that does not dissolve in excess i.e. Fe^{2+} , Cu^{2+} , Ca^{2+} , Ba^{2+} , Fe^{3+} and one which dissolves in excess sodium hydroxide i.e. Zn^{2+} , Pb^{2+} or Al^{3+}

You are provided with a substance B which contains two cations and one anion. Carryout the following tests on B to identify the ions present. Record your observations and deductions in the table below.

Tests	Observations	Deductions
(a)Heat a spatula end full of	Colourless gas that turns	CO ₂ gas evolved
B in a dry hard test tube	moist blue litmus paper red	CO ₃ ²⁻ probably present
strongly until no further	and lime water milky.	
change.	A black residue is formed	CuO formed, Cu ²⁺ present.
(b)To one spatula end full of B in the test tube, add dilute	B dissolves in the acid with bubbles of a colourless gas	CO ₂ is given off, CO ₂ ² - present
nitric acid to dissolve. To the solution obtained, add sodium hydroxide drop-wise	that turns moist blue litmus paper red and lime water	
until in excess and shake. Filter and keep both filtrate	milky. A blue precipitate insoluble in excess.	Cu ²⁺ present.
and the residue.	A blue residue and a	
Experiment for separating	colourles filtrate are formed	Zn ²⁺ , Pb ²⁺ , Al ³⁺ present in the
the two cations	on filtering.	filtrate.
i.e Zn ²⁺ (soluble) from Cu ²⁺		
(insoluble)		
(c)To 2cm ³ of the filtrate, add	A white precipitate forms and	Zn ²⁺ present
dilute nitric acid until the	dissolves in the acid	
solution is just acidic.		
Divide the resultant solution		
into 2 portions.		
Experiment for freeing the		
cation from the complex		
i.e Zn ²⁺ from [Zn(OH)4] ²⁻		
(i)To the first portion, add	A white precipitate soluble in	Zn ²⁺ , Pb ²⁺ , Al ³⁺ , present
sodium hydroxide drop-wise	excess forming a colourless	
until in excess.	solution.	
(ii)To the second portion, add	A white precipitate soluble in	Zn ²⁺ present
ammonia solution drop-wise	excess forming a colourless	
until in excess.	solution.	

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(d)Wash the residue with distilled water and add dilute nitric acid. Divide the solution into 2 portions.	Residue dissolves forming a blue solution.	Cu ²⁺ present.
(i)To the first portion, add sodium hydroxide drop-wise until in excess.	A blue precipitate insoluble in excess.	Cu ²⁺ present
(ii)To the second portion, add ammonia solution drop-wise until in excess.	A blue precipitate soluble in excess forming a deep blue solution.	Cu ²⁺ present

Identify the; anion present: CO_3^{2-} Cations present: Cu^{2+} and Zn^{2+}

(b) Using ammonia solution

This reagent is used to separate two cations of which one dissolves in excess ammonia solution e.g. Zn^{2+} or Cu^{2+} and the other does not dissolve in excess ammonia e.g. Pb^{2+} , Al^{3+} , Fe^{2+} , Fe^{3+} etc.

You are provided with substance B which contains two cations and one anion. Carryout the following tests on B to identify the ions present. Record your observations and deductions in the table below.

Tests	Observations	Deductions
(a)Heat a spatula end full of	Colourless gas that turns	CO ₃ ² - gas evolved
B in a dry hard test tube	moist blue litmus paper red	CO ₃ ²⁻ probably present
strongly until no further	and turns lime water milky.	
change	Residue is reddish brown	Fe ₂ O ₃ formed
		Fe ²⁺ probably present
(b)To one spatula end full of	B dissolves in the acid with	CO ₂ is given off.
B in the test tube, add dilute	bubbles of a colourless gas	CO ₃ ²⁻ present
nitric acid to dissolve. To the	that turns lime water milky.	
resultant solution, add		
ammonium hydroxide	A dirty green precipitate	An insoluble salt of Fe ²⁺
solution drop-wise until in	insoluble in excess.	present in the residue and
excess and shake. Filter and	A green residue and blue	Cu^{2+} , present in the filtrate
keep both the filtrate and	filtrate is formed on filtering.	-
residue.		
Experiment for separating		
the two cations.		
i.e Cu ²⁺ (soluble in		
ammonia) from Fe ²⁺		
(insoluble in ammonia)		

(c)To 2cm ³ of the filtrate, add	A blue precipitate forms and	Cu ²⁺ present
dilute nitric acid until the	dissolves in the acid	
solution is just acidic.		
Divide the resultant solution		
into 2 portions.		
Experiment for freeing the		
cation from the complex		
i.e Cu ²⁺ from [Cu(NH ₄)] ²⁺		
(i)To the first portion, add	A blue precipitate insoluble in	Cu ²⁺ present
sodium hydroxide drop-wise	excess.	
until in excess.		
(ii)To the second portion, add	A blue precipitate soluble in	Cu ²⁺ present
ammonia solution drop-wise	excess forming a deep blue	
until in excess.	solution.	
(d)Wash the residue with	Residue dissolves to form a	Fe ²⁺ present
distilled water and add dilute	green solution	
nitric acid.		
Divide the resultant solution		
into 2 portions.		
(i)To the first portion, add	A dirty green precipitate	Fe ²⁺ present
sodium hydroxide drop-wise	insoluble in excess. Turns	Fe ²⁺ oxidized to Fe ³⁺
until in excess.	brown on standing	
(ii)To the second portion, add	A dirty green precipitate	Fe ²⁺ present
ammonia solution drop-wise	insoluble in excess. Turns	Fe ²⁺ oxidized to Fe ³⁺
until in excess.	brown on standing.	

The cation present Cu^{2+} and Fe^{2+} .

The anion present CO_3^{2-}

PRECIPITATION REACTIONS OF ANIONS

Detection of anions is based on;

Precipitation of anions out of solution as insoluble salts of SO₄²⁻, Cl⁻, CO₃²⁻. Example;

- (a) Insoluble sulphates include;
 - lead (II) sulphate, barium sulphate and calcium sulphate.
- (b) Insoluble chlorides include;
 - lead (II) chloride and silver chloride.
- (c) Insoluble carbonates include;

All metal carbonates will form precipitates except sodium carbonate, potassium carbonate and

Ammonium carbonate which are soluble.

Test reagent for anions in solution, observations and deductions

Reagent	Observa	tion	Deduction
Lead (II) nitrate solution	White pro	ecipitate is formed	Cl ⁻ , SO ₄ ²⁻ , CO ₃ ²⁻ or SO ₃ ²⁻
		_	present
lead (II) nitrate solution +	White pro	ecipitate insoluble in	Cl ⁻ , and SO ₄ ²⁻ present
dil. HNO ₃	dilute nit	ric acid	_
	White pro	ecipitate soluble in	SO ₄ ²⁻ present
	dilute nit	ric acid	
Lead (II) nitrate solution +	white pre	cipitate insoluble in	
dilute nitric acid + heat	dilute nit	ric acid	
	(i)	White precipitate	SO ₄ ²⁻ absent
		dissolves on	PbCl ₂ salt formed
		heating and	Cl ⁻ confirmed present
		reappears on	
		cooling.	
	(ii)	White precipitate	Cl ⁻ absent
		does not dissolve	PbSO ₄ salt formed
		on heating	SO ₄ ²⁻ confirmed present
Barium nitrate or barium	White pro	ecipitate is formed	SO_4^{2-} , CO_3^{2-} , SO_3^{2-} present
chloride solution			
Barium nitrate solution +	White pre	ecipitate insoluble in	SO ₄ ²⁻ confirmed
dilute nitric acid	dilute nit	ric acid	
silver nitrate solution +	(i)	White precipitate	Cl ⁻ confirmed
dilute nitric acid		formed	
	(ii)	Pale yellow	Br ⁻ confirmed
		precipitate formed	
	(iii)	Yellow precipitate	I ⁻ confirmed

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Preliminary tests for anions

(a) Test: Action of dilute HCl or H₂SO₄ acid.

Add dilute acid to the solid substance in the test tube. If there is no reaction warm gently but do not boil.

Observat	ion	Deduction
(i)	Effervescences of a colourless, odourless gas	CO ₂ gas evolved, CO ₃ ²⁻ or
	that that turns lime water milky and damp blue	HCO ₃ present
	litmus paper red	
(ii)	In the cold or on heating, a colourless gas with	SO ₂ gas evolved, SO ₃ ²⁻ present
	a pungent smell turns blue litmus paper red	
	and bleaches it. It turns acidified potassium	
	dichromate solution from orange to green	
(iii)	Evolution of bad smelling gas (rotten egg	H ₂ S gas evolved, S ² -present
	smell), turns blue litmus paper red and turns	
	paper soaked in lead (II) Ethanoate solution	
	black	
(iv)	Solid dissolves in acid but no gas evolved	CO_3^{2-} , HCO_3^{-} or S^{2-} absent
		Cl ⁻ , SO ₄ ²⁻ , NO ₃ ⁻ , Br ⁻ or I ⁻
		probably present

(b) Test: Action of concentrated sulphuric acid.

Add cold concentrated sulphuric acid to the solid substance in the test tube. If there is no reaction warm gently but do not boil.

Obser	vation	Deduction
1.	Vigorous effervescence of a colourless,	CO ₂ gas produced
	oudourless gas that turns lime water milky	CO ₃ ²⁻ confirmed
2.	I the cold or on heating, colourless gas with a	SO ₂ gas
	pungent smell turns blue litmus paper red and	SO ₃ ²⁻ present
	bleaches it. It turns acidified potassium	
	dichromate solution from orange to green	
3.	Evolution of a bad smelling gas (rotten egg	H ₂ S gas evolved
	smell), turns blue litmus paper red and turns	S ²⁻ present
	paper soaked in lead (II) Ethanoate solution	
4.	Effervescence of fuming, pungent smelling	HCl gas
	chocking gas. Gas turns moist blue litmus paper	Cl ⁻ present
	red and forms dense white fumes with Conc.	
	NH ₃	
5.	Frothy effervescence of red brown vapour,	Br ₂ gas produced
	pungent smelling fuming gas. Gas turns blue	Br- present
	litmus paper red and reddish brown liquid is	
	formed	
6.	Colourless, pungent fuming gas. Gas turns blue	HI gas, I ⁻ present
	litmus paper red. Black solid is formed.	

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7. Colourless, pungent smelling vapour forms.	HNO ₃ vapour and NO ₂ gas
Vapour turns blue litmus paper red and fumes in	produced
air. Red brown vapour forms on heating.	NO ₃ ⁻ present.
8. Solid dissolves, no gas produced, no precipitate	Cl ⁻ , Br ⁻ , I ⁻ and NO ₃ ⁻ probably
forms.	absent
	SO ₄ ²⁻ probably present

Note: concentrated sulphuric acid oxidizes some of the hydrogen bromide and hydrogen iodide to red drown liquid bromine and violet iodine respectively.

$$2HBr_{(g)} + H_2SO_{4(l)} \longrightarrow Br_{2(g)} + SO_{2(g)} + 2H_2O_{(l)}$$

Confirmatory tests for the anions.

a) Sulphate ions SO₄²-

(i) Add a few drops of lead (II) nitrate solution or lead (II) Ethanoate a white precipitate is formed.

Equation: $Pb^{2+}_{(aq)} + SO_4^{2-}_{(aq)} \longrightarrow PbSO_{4(s)}$

(ii) Add a few drops of barium nitrate solution followed by dilute nitric acid, a white precipitate insoluble in the acid is formed.

Equation: $Ba^{2+}_{(aq)} + SO_4^{2-}_{(aq)} \longrightarrow BaSO_{4(s)}$

(iii) Add a few drops of barium chloride followed by dilute hydrochloric acid, a white precipitate insoluble in the acid is formed.

Equation: $Ba^{2+}_{(aq)} + SO_4^{2-}_{(aq)} \longrightarrow BaSO_{4(s)}$

b) Carbonate ions CO₃²-

(i) Add a few drops of lead (II) nitrate solution followed by dilute nitric acid, a white precipitate soluble in the acid with bubbles of a colourless gas that turns lime water milky.

Equation: $Pb^{2+}_{(aq)} + CO_3^{2-}_{(aq)} \longrightarrow PbCO_{3(s)}$

(ii) Add a few drops of dilute mineral acid preferably dilute nitric acid, bubbles or effervescences of a colourless gas that turns lime water milky.

Equation: $2H^+_{(aq)} + CO_3^{2-}_{(aq)}$ \longrightarrow $CO_{2(g)} + H_2O_{(l)}$

(iii) Add lead (II) nitrate or lead ethanoate solution, white precipitate of lead (II) carbonate is formed which dissolves in dilute acid forming a colourless solution.

Equation: $Pb^{2+}_{(aq)} + CO_3^{2-}_{(aq)} \longrightarrow PbCO_3_{(s)}$

(iv) Add silver nitrate solution, a white precipitate of silver nitrate is formed. The precipitate dissolves in aqueous ammonia or dilute nitric acid forming a clourless solution.

Equation: $Ag^{+}_{(aq)} + CO_3^{2-}_{(aq)} \longrightarrow Ag_2CO_3_{(s)}$

c) Nitrate ions NO₃ Brown ring test

(i) Add freshly prepared iron (II) sulphate solution followed by concentrated sulphuric acid carefully down the walls of the test tube in tilted position so that the acid sinks to the bottom a brown ring test forms at the junction.

d) Chloride ions Cl

(i) Add a few drops of lead (II) nitrate solution and warm, a white precipitate dissolves on warming and reappears on cooling.

Equation: $Pb^{2+}_{(aq)} + Cl^{-}_{(aq)} \longrightarrow PbCl_{(s)}$

(ii) Add a few drops dilute nitric acid followed by silver nitrate solution, a white precipitate is formed. The precipitate darkens on standing in light readily dissolves in ammonia solution forming a complex.

Equation: $Ag^{+}_{(aq)} + Cl^{-}_{(aq)} \longrightarrow AgCl_{(s)}$

e) <u>Iodide ions I</u>

(i) Add lead (II) nitrate solutions, a yellow precipitate is formed. Equation: $Pb^{2+}_{(aq)} + 2I^{-}_{(aq)} \longrightarrow PbI_{2(s)}$

f) Hydrogen carbonate ions, HCO3

(i) Boil the solution of the unknown and add a few drops of phenolphthalein indicator.

Observation:

Bubbles of colourless gas are formed which turn lime water milky. The resultant solution turns phenolphthalein indicator pink.

(ii) Add magnesium sulphate solution and boil.

Observation:

A white precipitate is formed on boiling.

g) Sulphite ions SO₃²-

(i) Add lead (II) nitrate solution followed by dil. HNO₃. White precipitate is formed. The precipitate is soluble in dil. HNO₃.

Equation: $Pb^{2+}_{(aq)} + SO_3^{2-}_{(aq)} \longrightarrow PbSO_{3(s)}$ $PbSO_{3(s)} + 2H^{+}_{(aq)} \longrightarrow SO_{2(g)} + H_2O_{(l)} + Pb^{2+}_{(aq)}$

(ii) Add aqueous barium chloride or barium nitrate followed by dil. HNO₃. A white precipitate soluble in dil. HNO₃ confirms sulphite ions.

Equation: $Ba^{2+}_{(aq)} + SO_3^{2-}_{(aq)} \longrightarrow BaSO_{3(s)}$ $BaSO_{3(s)} + 2H^+_{(aq)} \longrightarrow SO_{2(g)} + H_2O_{(l)} + Ba^{2+}_{(aq)}$

h) Bromide ions Br

- (i) Add lead (II) nitrate solution. A yellow precipitate is formed Equation: $Pb^{2+}_{(aq)} + 2Br_{(aq)} \longrightarrow PbBr_{2(s)}$
- (ii) Add silver nitrate solution followed by dilute nitric acid. Cream or light yellow precipitate insoluble in dil. HNO₃ and darkens on standing in light. Equation: $Ag^+_{(aq)} + Br^-_{(aq)} \longrightarrow AgBr_{(s)}$

Sample question

- 1. Name the reagent that can be used to distinguish between the following pairs of ions in aqueous solution and state what would be observed when each of the following ion is treated with the reagent named
 - (a) SO_3^{2-} (aq) and SO_4^{2-} (aq)

Reagent: Barium nitrate solution or Barium chloride solution followed by dil. Nitric acid or hydrochloric acid.

Observation: SO_3^{2-} (aq) - A white precipitate soluble in dilute acid.

 SO_4^{2-} (aq) - A white precipitate insoluble in dilute acid.

Note:

Similarly lead (II) nitrate solution followed by nitric acid can be used and the observations are as above

Alternatively, we can use dil. HCl or H₂SO₄.

Observation: SO₃²- Effervescence is produced

SO₄²- No effervescence is produced

(b) HCO_3^{2-} (aq) and CO_3^{2-} (aq)

Reagent: Magnesium chloride solution or magnesium sulphate solution

Observation: CO₃²-(aq) White precipitate of MgCO₃ is formed

HCO_{3 (aq)} No precipitate is formed in cold but on heating a white

precipitate is formed.

Equations: $Mg^{2+}_{(aq)} + CO_3^{2-}_{(aq)} \longrightarrow MgCO_{3(s)}$ $Mg^{2+}_{(aq)} + HCO_3^{-}_{(aq)} \longrightarrow Mg(HCO_3)_{2 (aq)}$

Explanations

When the solution is heated, magnesium hydrogen carbonate decomposes to form a white precipitate of magnesium carbonate.

$$Mg(HCO_3)_{2 (aq)} \longrightarrow MgCO_{3(s)} + CO_{2(g)} + H_2O_{(l)}$$

(c) **Reagent**: silver nitrate solution.

Observation: Cl⁻ White precipitate of AgCl is formed

I Yellow precipitate of AgI is formed

Or we can use lead (II) nitrate solution

Observation: Cl⁻ White precipitate of PbCl₂ is formed

I yellow precipitate of PbI₂ is formed

WORKED EXAMPLES

Example 1

You are provided with substance **Z** which contains **one cation** and **two anions**. Carry out the following tests to identify the cations and anions in **Z**. identify any gases evolved.

Tests	Observations	Deductions
Heat a spatula end-full of Z in a dry test tube until there is no further change	A colourless gas evolved, gas turns moist blue litmus paper red, gas turns lime water milky Residue is yellow when hot, white when cold	Gas is CO Hence CO ₃ ²⁻ , HCO ₃ - present Residue is ZnO Hence Zn ²⁺ present
Dissolve two spatula end-full of Z in distilled water, filter and keep both the filtrate and residue	Partialy dissolves forming a colourless solution and a white residue	Zn ²⁺ , Pb ²⁺ , Al ³⁺ , Ba ²⁺ , Mg ²⁺ , Ca ²⁺ , probably present
Divide the filtrate into four equal portions, To the first portion, and sodium hydroxide solution drop wise until excess.	White precipitate, dissolves in excess forming a colourless solution	Zn ²⁺ , Pb ²⁺ , Al ³⁺ , probably present
To the second portion, add ammonia solution drop wise until excess.	White precipitate, soluble in excess forming a colourless solution.	Zn ²⁺ , confirmed present
To the third portion, add 3 drops of lead (II) nitrate.	White precipitate.	SO ₄ ² -, Cl ⁻ , SO ₃ ² -, CO ₃ ² -, probably present
Carry out a test of your own to confirm the anion in Z To the fourth portion, add barium nitrate solution followed by dilute nitric acid.	White precipitate insoluble in dilute nitric acid.	SO ₄ ²⁻ , confirmed present
Dissolve the residue in dilute hydrochloric acid and divide the resultant solution into two equal portions.	Dissolves with effervescence evolving a colourless gas, gas turns moist blue litmus paper red, gas turns lime water milky. A colourless solution is formed	Gas is CO ₂ Hence CO ₃ ²⁻ , confirmed present
To the first portion, add sodium hydroxide solution drop wise until excess	White precipitate soluble in excess forming a colourless solution.	Zn ²⁺ , Pb ²⁺ , Al ³⁺ , probably present
To the second portion, add ammonia solution drop wise until excess	White precipitate, soluble in excess ammonia solution forming a colourless solution	Zn ²⁺ , confirmed present

Identify the;

- (i) Cation in \mathbb{Z} . $\mathbb{Z}n^{2+}$
- (ii) Anions in **Z**. CO_3^{2-} , SO_4^{2-}

Example 2

You are provided with substance **X** which contains **one cation** and **two anions**. Carryout the following tests to identify the cations and anions present in **X**.

Test	Observation	Deduction
(a)To 2 spatula end-full of X	X is white powdery	Al ³⁺ , Zn ²⁺ , Pb ²⁺ , probably
add about 5cm ³ of water and	crystalline solid, partiary	present
shake well. Filter and keep	soluble forming a colourless	CO ₃ ²⁻ , NO ₃ ⁻ , SO ₄ ²⁻ , CI ⁻ ,
both the residue and the	filtrate and a white residue.	probably present
filtrate.		
(b)Divide the filtrate into four	A white precipitate, soluble in	Al ³⁺ , Zn ²⁺ , Pb ²⁺ , probably
portions.	excess forming a coloueless	present
(i)To the first portion, add	solution.	
dilute sodium hydroxide		
solution drop wise until in		
excess		
(ii)To the second portion, add	A white precipitate insoluble	Pb^{2+} , Al^{3+} , Mg^{2+} , probably
dilute aqueous ammonia drop	in excess.	present
wise until in excess.		
(iii)To the third portion, add	A yellow precipitate is	Pb ²⁺ confirmed present
drops of potassium iodide	formed	
solution		
(iv)To the fourth portion, add	A white precipitate was	Gas is NO ₂ ,
copper fillings followed by	formed on addition of the	NO ₃ confirmed present
conc. sulphuric acid and boil	acid. On boiling, a brown gas	
	which turned moist blue	
	litmus paper red was given	
	off.	
	A green blue solution was	
	formed.	
(c)Wash the residue in (a)	Effervescence with evolution	Gas is CO_2 ,
above and place it in a test	of a colourless gas, gas turns	CO ₃ ²⁻ , confirmed present
tube. Add dilute HCI	moist blue litmus paper red	
	and lime water milky.	

- (d) (i) Cation in \mathbf{X} . Pb²⁺
 - (ii) Anions in \mathbf{X} . NO_3^- and CO_3^{2-}

Example 3

You are provided with substance **Y** that contains **one cation** and **one anion**. Carry out the following tests to identify the cations and anions in **Y**. identify any gases that may be evolved. Record your observations and deductions in the table below.

Tests	Observations	Deductions
a)Appearance of Y	Y is a green crystalline substance	Fe ²⁺ , Cu ²⁺ probably present
b)Place a spatula end-full of Y in a dry test tube then heat strongly until there is no further change	A colourless vapour that condenses to a colourless liquid is given off. The liquid turns white anhydrous copper	Water of crystallization from hydrated salt or HCO ₃ -, OH
	(II) sulphate blue. A colourless gas with an irritating smell is evolved, gas turns blue litmus paper red,	Gas is HCl, Cl ⁻ probably present
	gas forms misty fumes in air and dense white fumes with conc. Ammonia solution. Black residue is formed	CuO, FeO, Fe ₃ O ₄ , Cu ²⁺ , Fe ²⁺ , Fe ³⁺ , probably present
c) To a spatula end-full of Y in a test tube add 5cm ³ of distilled water and shake. Divide the solution into 4	Dissolves forming a pale blue solution	Cu ²⁺ probably present
portions. (i)To the first portion, add sodium hydroxide solution drop wise until in excess	Pale blue precipitate insoluble in excess sodium hydroxide.	Cu ²⁺ present
(ii)To the second portion, add dilute ammonia solution drop wise until in excess	Blue precipitate soluble in excess forming a deep blue solution	Cu ²⁺ confirmed present
(iii)To the third portion, add lead (II) nitrate solution.	White precipitate is formed	Cl ⁻ , SO ₄ ²⁻ , CO ₃ ²⁻ , SO ₃ ²⁻ , present
(iv)Use the fourth portion to carry out a test of your own choice to confirm the anion in Y		
Test To the fourth portion, add 2-3 drops of silver nitrate solution followed by dilute nitric acid	White precipitate insoluble in nitric acid	Cl ⁻ confirmed present

Identify the;

Cations in **Y**: Cu²⁺ Anions in **Y**: Cl⁻

Example 4

You are provided with substance **A** that contains **two cations** and **anions**. Carry out the following tests to identify the cations and anions in **A**. identify any gases the may be evolved. Record your observations and deductions in the table below.

Tests	Observations	Deductions
a) Place two spatula end-full	Partiary dissolves forming a	Pb ²⁺ , Zn ²⁺ , NH ₄ +, Al ³⁺
of A in a test tube, add 5cm^3	colourless filtrate and white	probably present.
of water, shake vigorously	residue.	
and filter. Keep both the	No observable change, but a	NH ₃ gas evolved
filtrate and the residue.	colourless gas with an	NH ₄ ⁺ confirmed present
Divide the filtrate into 3	irritating pungent smell is	
portions	evolved. The gas turns red	
(i)To the first portion, add	litmus paper blue and forms	
excess sodium hydroxide	dense white fumes with conc.	
solution then heat and smell.	HCI.	
(ii)To the second portion, add	White precipitate does not	Cl ⁻ absent
lead (II) nitrate solution and	dissolve on heating	SO ₄ ²⁻ present
heat		
(iii)Use the third portion to		
carry out a test of your choice		
to confirm the anion in A		
Test		
To the third portion, add 2-3	White precipitate insoluble in	SO ₄ ²⁻ confirmed present
drops of Barium chloride	the acid	304 commined present
solution followed by dilute	the deld	
hydrochloric acid.		
b) Wash the residue with little	White residue turns yellow	CO ₂ gas evolved
distilled water then divide it	when hot and then white on	CO ₃ ²⁻ present
into two parts.	cooling.	r
(i)Strongly heat the first part	A colourless gas that turns	
until no further change occurs	blue litmus paper pink and	
	lime water milky is evolved.	
c) To the second portion of	The residue dissolves with	CO ₂ gas evolved,
the residue, add 3cm ³ of	rapid effervescence of a	CO ₃ ²⁻ confirmed present
dilute hydrochloric acid.	colourless gas that turns	
Divide the mixture into 2	damp blue litmus paper pink	
portions.	and lime water milky.	
(i)To the first part of the	A colourless solution is	Zn^{2+} , Pb^{2+} , Al^{2+} , present
mixture, add sodium	formed.	
hydroxide solution drop wise	A white precipitate soluble in	
until in excess.	excess NaOH _(aq) forming a	
	colourless solution	

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(ii)To the second portion, add	White precipitate soluble in	Zn ²⁺ confirmed present.
ammonia solution drop wise	excess forming a colourless	
until in excess.	solution.	

Identify the:

Cations in **A**: NH₄⁺ and Zn²⁺ Anions in **A**: SO₄²⁻ and CO₄²⁻

Example 5

You are provided with a substance **T** which contains **one cation** and **one anion**. Carry out the following tests on **T** to identify the ions present. Identify any gases evolved. Record your observations and deductions in the table below.

Tests	Observations	Deductions
Note the appearance of T	T is green crystalline solid	Fe ²⁺ , Cu ²⁺ present
		NO ₃ -, SO ₄ ² -, Cl ⁻ probably
		present
Heat a spatula end-full of T in	Colourless vapour condenses	Water of crystallization from
a dry test tube strongly until	to form a colourless liquid	hydrated salt or HCO ₃ or
no further change.	which turns white anhydrous	OH-
	copper (II) sulphate blue.	
	Colourless gas that turns	
	moist blue litmus paper red,	SO ₂ gas evolved,
	gas turns acidified potassium	SO ₄ ²⁻ present
	dichromate from orange to	
	green.	
	Green solid turns to reddish	Fe ₂ O ₃ formed
TD 1 1 . C . 11 . C . TD	brown.	Fe ³⁺ probably presents
To one spatula end-full of T	Dissolves forming a pale	Fe ²⁺ , Cu ²⁺ probably present
in the test tube, add 5cm ³ of distilled water and shake.	green solution	
Divide the solution into 5		
portions.		
To the first portion, add	Dirty green precipitate	Fe ²⁺ present
sodium hydroxide solution	insoluble in excess and turns	re present
until in excess	brown on standing.	
To the second portion, add	Dirty green precipitate	Fe ²⁺ present
ammonia solution drop wise	insoluble in excess and turns	Te present
until in excess.	brown on standing.	
To the third portion, add	A dark blue precipitate is	Fe ²⁺ confirmed present
potassium hexacyanoferrate	formed	1
(III).		
To the fourth portion, add	A white precipitate is formed	Cl ⁻ , SO ₄ ²⁻ , CO ₄ ²⁻ , SO ₃ ²⁻
lead (II) nitrate solution.		present.

To the fifth portion, add	White precipitate insoluble in	SO ₄ ²⁻ confirmed present.
Barium nitrate solution	nitric acid.	
followed by nitric acid.		

Identify the;

Cation in **T**: Fe²⁺ Anion in **T**: SO₄²⁻

Example 6

You are provided with a substance **W** which contains **one cation** and **two anions**. Carry out the following tests on **W** to identify the ions present. Identify any gases evolved. Record your observations deductions in the table below.

Tests	Observations	Deductions
Note the appearance of W	W is a green crystalline solid	Fe ²⁺ , Cu ²⁺ present
		SO ₄ ²⁻ , NO ₃ -, Cl-, CO ₃ ²⁻
		probably present.
Heat a spatula end-full of W	Colourless vapour condenses	Water of crystallization from
in a dry clean test tube	to form a colourless liquid	hydrated salt, HCO ₃ -, OH- or
strongly until no further	which turns white anhydrous	HSO ₄ ⁻ present.
change	copper (II) sulphate blue.	CO ₂ gas evolved,
	A colourless gas that turns	CO ₃ ²⁻ probably present.
	moist blue litmus paper red	
	and lime water milky is given	CuO, FeO, F ₃ O ₄ formed
	off.	Cu ²⁺ , Fe ²⁺ , Fe ³⁺ probably
	Green solid turns to black.	present
T one spatula end-full of W	Partiary dissolves forming a	Cu ²⁺ , Fe ²⁺ probably present.
in the test tube and 5cm ³ of	blue filtrate and a green	
distilled water and shake.	residue.	
Filter and keep the residue.		
Divide the filtrate into 4		
portions.		2.
To the first portion, add	A blue precipitate insoluble	Cu ²⁺ present.
sodium hydroxide solution	in excess sodium hydroxide	
drop wise until in excess.	solution	
To the second portion, add	A blue precipitate soluble in	Cu ²⁺ confirmed present
ammonia solution drop wise	excess forming forming a	
until in excess.	deep blue solution.	
To the third portion, add lead	A white precipitate is formed.	SO ₄ ²⁻ , Cl ⁻ , SO ₄ ²⁻ , CO ₃ ²⁻ , SO ₃ ²⁻
(II) nitrate solution.		probably present.
To the fourth portion, add	A white precipitate insoluble	CI ⁻ confirmed present.
silver nitrate solution	in nitric acid.	
followed by nitric acid.		
Wash the residue with cold	Residue dissolves with	CO ₂ ² - gas evolved.
water and add nitric acid.	bubbles of a colourless gas	CO_3^{2-} confirmed present.

Divide the solution into 2	that turns damp blue litmus	
portion	paper pink and lime water	
	milky.	
To the first portion, add	Blue precipitate insoluble in	Cu ²⁺ present.
sodium hydroxide solution	excess.	
drop-wise until in excess.		
To the second portion, add	Blue precipitate soluble in	Cu ²⁺ present.
ammonia solution drop-wise	excess forming a deep blue	
until in excess.	solution.	

Identify the;

Cation in W: Zn²⁺

Anions in W: CO₃²⁻ and Cl⁻

Example 7

You are provided with substance **M** which contains **one cation** and **one anion**. Carry out the following tests to identify the cation and anion present in **M**. Identify any gases evolved and record the observations and deductions in the table below.

Tests	Observations	Deductions
(a)Heat a spatula end-full of	A colourless gas that turns	CO ₂ gas evolved
M strongly in a test tube until	moist blue litmus paper red	CO_3^{2-} present.
no further change.	and lime water milky.	
	Brown residue when hot and	Residue is PbO.
	yellow when cold.	Pb ²⁺ present.
(b)Dissolve a spatula end-full	Colourless gas that turns	CO ₂ gas evolved
of M in dilute nitric acid.	moist blue litmus paper red	CO ₃ ²⁻ confirmed present.
Divide the solution obtained	and lime water milky.	
into 3 parts.	Colourless solution foemed.	
(i)To the first part, add	A white precipitate soluble in	Pb ²⁺ , Zn ²⁺ , Al ³⁺ probably
sodium hydroxide solution	excess.	present.
drop-wise until in excess.		
(ii)To the second part, add	A white precipitate insoluble	Al^{3+} , Pb^{2+} probably present.
dilute aqueous ammonia	in excess.	
solution drop-wise until in		
excess.		
(iii)Use the third part to carry		
out a test of your choice to		
confirm the cation in M .		
Test		
To the third part add	A yellow precipitate is	Pb ²⁺ confirmed present.
potassium iodide solution.	formed.	

Identify:

- (i) The cation in \mathbf{M} . Pb²⁺
- (ii) The anion in \mathbf{M} . CO_3^{2-}

PRACTICAL SCHEDULES

Practical 1

You are provided with substance **H** which contains **one cation** and **one anion**. Carry out the following tests to identify the cation and anion in **H**. Identify any gases evolved. Record your observation and deductions in the table below.

Tests	Observations	Deductions
(a)Heat a spatula end-full of H in		
a dry test tube until there is no		
further change.		
(b)Dissolve two spatula end-full		
of H in 5cm ³ of distilled water.		
Divide the solution into 4 equal		
portions.		
(i)To the first portion, add		
sodium hydroxide drop-wise		
until in excess.		
(ii)To the second portion, add		
ammonia solution drop-wise		
until in excess.		
(iii)To the third portion, add 3		
drops of lead (II) nitrate solution.		

(iv)Use the fourth portion to		
carry out a test of your own to		
confirm the anion in H .		
•••••		
•••••		
T1 (C 4		
Identify the;		
Cation in H		
	•••••	
Anion in H		

Practical 2

You are provided with substance **P** which contains **two cations** and **one anion**. Carry out the following tests to identify the ions in **P**. Identify any gases evolved and record your observations and deductions in the table.

Tests	Observations	Deductions
(a)Heat a spatula end-full of P in		
a dry test tube until there is no		
further change.		
(b)Dissolve two spatula end-full		
of P in 5cm ³ of distilled water.		
Filter and keep both the filtrate		
and the residue. Divide the		
solution in four equal portions.		
(NT 41 6" 4 4" 11		
(i)To the first portion, add		
sodium hydroxide solution drop- wise until in excess.		
wise until ill cacess.		
(ii)To the second portion, add		
ammonia solution drop-wise		
until in excess.		

(iii)To the third portion, add 3	
drops lead (II) nitrate solution.	
(iv)Use the fourth portion to	
carry out the test of your own to	
confirm the anion in P .	
(c)Dissolve the residue in dilute	
sulphuric acid and warm. Divide the resultant solution into two	
equal portions.	
o qualifornitation	
(i)To the first portion, add	
sodium hydroxide solution drop-	
wise until in excess.	
(ii)To the second portion, add ammonia solution drop-wise	
until in excess.	
Girai in encessi	
Identify the:	
Cations in P	
Anion in P	

Practical 3

You are provided with substance **X** which contains **one cation** and **two anions**. Carry out the following tests to identify the cation and anions in **X**. Identify any gases that may be evolved. Record your observations and deductions in the table below.

Tests	Observations	Deductions
(a)Heat a spatula end-full of X		
in a dry test tube until there is		
no further change.		

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(b)Dissolve the two spatula end-full of X in 5cm ³ of distilled water. Filter and keep both the filtrate and the residue. Divide the solution in four equal portions.	
(i)To the first portion, add sodium hydroxide solution drop-wise until in excess.	
(i)To the second portion, add ammonia solution drop-wise until in excess.	
(ii)To the third portion, add lead ethanoate solution.	
(iii)To the fourth portion, Barium chloride solution followed by dilute hydrochloric acid.	
(c)Dissolve the residue in dilute nitric acid and warm. Divide the resultant solution in two equal portions.	
(i)To the first portion, sodium hydroxide solution drop-wise until in excess.	
(ii)To the second portion, add ammonia solution drop-wise until in excess.	
Identify the: Cation in X	

Practical 4

You are provided with substance \mathbf{Q} which contains **two cations** and **one anion**. Carry out the following tests to identify the cations and anion in \mathbf{Q} . Identify any gases evolved. Record your observations and deductions in the table below.

Tests	Observations	Deductions
(a)Heat a spatula end-full of Q in dry test tube until there is no further change.		
(b)Dissolve two spatula endfull of Q in 5cm ³ of distilled water.		
(c)To the resultant solution, add sodium hydroxide solution drop-wise until in excess. Filter and keep both the filtrate and the residue.		
(d)Add dilute nitric acid to the filtrate until the solution becomes acid. Divide the resultant solution into three equal portions.		
(i)To the first portion, add sodium hydroxide drop-wise until in excess.		
(ii)To the second portion, add 3 drops of lead (II) nitrate solution.		

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(iii)To the third portion, add barium chloride solution followed by dilute hydrochloric acid.	
(e)Dissolve the residue in dilute nitric acid and warm. Divide the resultant solution into three equal portions.	
(i)To the first portion, add sodium hydroxide solution drop-wise until in excess.	
(ii)To the second portion, add ammonia solution drop-wise until in excess.	
(iii)Use the third portion to confirm the cation in Q	

L	en	4:	fx,	+1	٠.
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Cations in Q	 	 	 				
Anion in Q	 	 	 			 	

Practical 5

You are provided with substance **Z** which contains **one cation** and **two anions**. Carry out the following tests to identify the cation and anions in **Z**. Identify any gases evolved and record your observations and deductions in the table below.

(a)Heat a spatula end-full of Z in a dry test tube gently and later strongly until there is no further change.	
(b)Dissolve two spatula end-full of Z in 5cm ³ of distilled water. Divide the solution into five portions.	
(i)To the first portion, add sodium hydroxide solution dropwise until in excess.	
(ii)To the second portion, add ammonia solution drop-wise until in excess.	
(iii)To the third portion, add 3 drops of potassium iodide solution.	
(iv)To the fourth portion, add 3 drops of lead (II) nitrate solution.	
(v)Use the fifth portion to carry out a test of your own choice to confirm the anion in Z .	

Identify the:	
Cations in X	

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Practical 6

Anion in X.....

You are provided with substance **T** which contains **two cations** and **two anions**. Carry out the following tests to identify the cations and anions in **T**. Identify any gases evolved. Record your observations and deductions in the table below.

Tests	Observations	Deductions
(a)Heat a spatula end-full of T in a dry test tube until there is no		
further change.		
(L)Discolars true souts 1, and full		
(b)Dissolve two spatula end-full of T in 5cm ³ of distilled water.		
Then add sodium carbonate solution.		
Filter and keep both the filtrate		
and the residue. Divide the filtrate into four equal portions.		
(i)To the first portion, add		
sodium hydroxide solution drop- wise until in excess.		
(ii)To the second portion, add 3		
drops of lead (II) nitrate solution		

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(iii)Use the third portion to carry	
out the test of your own choice	
to confirm the anion in T .	
to commin the amon in 1.	
(iv)To the fourth portion, add	
barium chloride solution	
followed by dilute hydrochloric	
acid.	
(c)Wash the residue with	
distilled water and dissolve in	
dilute nitric acid until it	
dissolves. Divide the resultant	
solution into three equal	
portions.	
portions.	
(i)To the first portion, add	
sodium hydroxide solution drop-	
wise until in excess.	
wise until ill excess.	
(ii)To the second portion, add	
ammonia solution drop-wise	
until in excess.	
until ill excess.	
(iii)To the third or other and d	
(iii)To the third portion, add	
potassium iodide solution.	
	<u> </u>
T	
Identify the:	
Cations in T	
Anions in T	

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You are provided with substance **Y** which contains **two cations** and **one anion**. Carry out the following tests to identify the cations and anion in **Y**. Identify any gases evolved. Record your observations and deductions in the table below.

Tests	Observations	Deductions
(a)Heat a spatula end-full of Y		
in dry test tube until there is no		
further change.		
(h)Dissolve two spetule and full		
(b)Dissolve two spatula end-full of Y in 5cm ³ of distilled water.		
of 1 in Join of distinct water.		
(i)To about 1cm ³ of the solution,		
add 3 drops lead (II) nitrate		
solution.		
(ii)Use 1cm ³ of the solution to		
carry out a test of your own to		
identify the cation in Y .		
(c)To the rest of the solution, in		
(b) add sodium hydroxide		
solution drop-wise until in		
excess.		
Filter and keep both the filtrate		
and the residue.		
(d) Ad diluta HCI asid drap wise		
(d)Ad dilute HCI acid drop-wise to the filtrate until it is just		
acidic. Divide the resultant		
solution into three portions.		
portions.		
	I .	

(i)To the first portion, add sodium hydroxide solution dropwise until in excess.		
(ii)To the second portion, add ammonium hydroxide solution drop-wise until in excess.		
(iii)To the third portion, add potassium iodide solution.		
(e)Dissolve the residue from (c) in dilute HCI acid and divide the resultant solution into two portions.		
(i)To the first portion, add sodium hydroxide solution dropwise until in excess.		
(ii)To the second portion, add ammonia solution drop-wise until in excess.		
Identify the:	1	
Cations in Y		

Cations in \mathbf{Y}	 	 						
Anion in V								

You are provided with substance **M** that contains **one cation** and **one anion**. Carry out the following tests and identify the cation and anion in **M**. identify any gases evolved. Record your observations and deductions in the table below.

Tests	Observations	Deductions
(a)To a spatula end-full of M ,		
add 5cm ³ of distilled water and		
shake. Divide the resultant		
solution into five parts.		
(i)To the first part, add sodium		
hydroxide solution drop-wise		
until in excess.		
(ii)To the second part, add		
ammonia solution drop-wise		
until in excess.		
(:::)To the third next odd load		
(iii)To the third part, add lead (II) nitrate solution.		
(11) mitate solution.		
(iv)To the fourth part of the		
solution, carry out the test of		
your own choice to confirm the		
anion in M .		

Identify the:
Cation in M
Anion in M

You are provided with substance **V** which contains **one cation** and **one anion.** Carry out the following tests to identify the cation and anion in **V**. Identify any gas(es) that may be evolved. Record your observations and deductions in the table below.

Tests	Observations	Deductions
(a)Heat a spatula end-full of V in a	Observations	2 caacaaa
dry test tube.		
(b)Add 5cm ³ of distilled water to		
one spatula end-full of V and		
shake. Divide the resultant		
solution into five parts.		
solution into five parts.		
(i)To the first part, add sodium		
hydroxide solution and warm.		
njaromae soration and warm.		
(ii)To the second part, add dilute		
ammonia solution drop-wise until		
in excess.		
in cheess.		
(iii)To the third part, add lead (II)		
nitrate solution and warm then		
allow it to cool.		

(iv)To the fourth part, add silver		
nitrate solution followed by dilute	1	
nitric acid.		
(v)Use the fifth part to carry out a		
test of your own choice to confirm	1	
the anion in V .	1	
	1	
	1	
	1	
	1	
	1	
	L	
Identify the:		
Cation in V		
Anion in V		

You are provided with substance **W** which contains **one cation** and **one anion**. Carry out the following tests on W to identify the cation and anion in **W**. Identify any gases given off. Record your observations and deductions in the table below.

Tests	Observations	Deductions
(a)Heat a spatula end-full of W in a dry test tube until there is no further change.		
(b)To a spatula end-full of W, add 5cm ³ of distilled water and shake to dissolve. Divide the resulting solution into four parts.		

(i)To the first part, add sodium solution hydroxide drop-wise until in excess.	
(ii)To the second part, add ammonia solution drop-wise	
until in excess.	
(iii)To the third part, add lead (II) nitrate solution.	
(iv)To the fourth part, add barium nitrate solution followed by dilute nitric acid.	

Identify the:
Cation in W
Anion in W

You are provided with substance \mathbf{Q} which is a mixture of ammonium chloride and copper (II) carbonate. Complete the table below by filling in the expected observation and deductions when the tests are carried out on \mathbf{Q} .

Tests	Observations	Deductions
(a)Heat a spatula end-full of Q in a dry test tube strongly until there is no further change.		
(b)To one spatula end-full of Q in a test tube, add 5cm ³ of distilled water and shake. Filter, keep both the filtrate and the residue. Divide the filtrate in 3 portions.		
(i)To the first portion, add sodium hydroxide solution dropwise until in excess and warm.		
(ii)To the second portion, add lead (II) nitrate solution.		

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(iii)To the third portion, carry	
out a test of your own choice to	
identify the anion in \mathbf{Q} .	
(c)Wash the residue with	
distilled water, transfer it in a	
test tube and add dilute nitric	
acid. Divide the resultant	
solution into two portions.	
solution into two portions.	
(i)To the first portion, add	
sodium hydroxide solution drop-	
wise until in excess.	
(ii)To the second portion, add	
ammonia solution drop-wise	
until in excess.	

Cations in **Q** NH4⁺ and Cu²⁺ Anions in **Q** Cl⁻ and CO₃²⁻

Practical 12 (UNEB PAPER 3 2019)

You are provided with substance \mathbf{Q} , which contains **two cations** and **one anion**. Carry out the following tests to identify the cations and the anion present in \mathbf{Q} . Identify any gas(es) that may be evolved. Record your observations and deductions in the table below.

Tests	Observations	Deductions
(a)Dissolve one spatula end-full		
of \mathbf{Q} in about 5cm^3 of water. Add		
excess sodium hydroxide		
solution; shake well and filter.		
Keep both the filtrate and the		
residue.		
(b)To the filtrate, add dilute nitric		
acid until the solution is just acid.		
Divide the acidic solution into		
five portions.		
(i)To the first portion of the		
acidified solution, add dilute		
sodium hydroxide drop-wise		
until in excess.		
(ii)To the second portion of the		
acidified solution, add aqueous		
ammonia drop-wise until in excess.		
excess.		
(iii)To the third portion of the		
acidified solution, add potassium		
iodide solution.		
(iv)To the fourth portion of the		
acidified solution, add lead (II)		
nitrate solution and warm.		
(v)Use the fifth portion of the		
acidified solution to carry out a		
test of your own to confirm the anion in Q .		
amon in Q.		
	ı	

/\D' 1 .1 .1 .1	
(c)Dissolve the residue in a minimum amount of dilute	
sulphuric acid and divide the	
resultant solution into two parts.	
resultant solution into two parts.	
(i)To the first part of the solution	
add sodium hydroxide solution	
drop-wise until in excess.	
(ii)To the second part of the	
solution, add 1 small piece of	
zinc granules and leave the solution to stand for 5 minutes.	
Divide the solution into two	
portions and use them for part (d)	
portions and use them for part (u)	
(d) (i) To the first portion of the	
solution, add sodium hydroxide	
drop-wise until in excess.	
(ii)To the second portion of the	
solution, add aqueous ammonia	
drop-wise until in excess.	
T	
The cations in Q are	
_	

The cations in	Q are
The anion in O	is.

Chapter two: VOLUMETRIC ANALYSIS

Volumetric analysis is the technique of finding the concentration of the solution. In this technique the solution of the known concentration is added repeatedly to the one whose concentration is to determine until there is just enough of it to neutralize the other of the unknown concentration. This method is called titration. During titration, the volume of standard solution that completely reacts with the measured volume of the solution under the analysis is recorded. The titration is repeated to obtain consistent tires.

From consistent tires values, an average volume of the standard solution is calculated. By use of appropriate stoichiometry (reaction ratio) obtained from the balanced equation between the reacting substance in the experiment, the concentration of the solution under analysis can be calculated.

Volumetric analysis is performed using the following important apparatus. These include burette, pipette, conical flask, beakers, retort stand and clamp, volumetric flasks, weighing balance. Quantitative (volumetric) analysis requires the knowledge and application of knowledge on mole concept. Therefore the following terms are useful in volumetric analysis.

1. Relative formula mass (RFM).

This is the mass of one molecule of an element or compound on a scale on which one atom of carbon-12 is exactly 12 units. It is determined by adding the atomic mass of the element in the compound.

Relative formula mass has no units.

E.g Calculate the relative formula mass of Na_2CO_3 . (Na = 23, C = 12, O = 16)

RFM of
$$Na_2CO_3 = (23x2) + (12x1) + (16x3)$$

= $46 + 12 + 48$
= 106

2. Molarity of the solution (concentration in moles per litre) or molar concentration.

This refers to the number of moles of the compound contained in a litre (1000cm³) of a solution.

Therefore, 0.2M of HCI means 1000cm³ of HCI contains 0.2 moles.

3. Concentration in grams per litre or grams per cubic decimeters.

This refers to the mass of the compound dissolved to make 1 litre of the solution. When 4g of sodium carbonate are dissolved to make 1 litre of solution, we say that the solution has a concentration of 4gl⁻¹ or 4gdm⁻³ of sodium carbonate.

Therefore, Concentration in grams per litre = Molarity x Relative formula mass.

Example

53.5g of sodium chloride were dissolved to make $2000cm^3$ of the solution, calculate the concentration of sodium chloride in grams per litre. (Na = 23, Cl = 35.5)

Solution

RFM of NaCl =
$$23 + 35.5$$

Therefore molar mass of NacCl = 58.5g

58.5g of NaCl contain 1 mole

1g of NaCl will contain $\frac{1}{58.5}$ moles

53.5g of NaCl will contain
$$\frac{1 \times 53.5}{58.5}$$
 moles

2000cm³ of solution contains 0.9145 moles of NaCl

$$1 \text{cm}^3$$
 of solution will contain $\frac{0.9145}{2000}$ moles of NaCl

$$1000 \text{cm}^3$$
 of solution will contain $\frac{0.91445 \times 1000}{2000}$ moles of NaCl 0.4573M

1 moles of NaCl weighs 58.5g

0.4573 moles will weigh
$$\frac{0.4573 \times 58.5}{1}$$
 g 26.75 gl⁻¹

4. A standard solution.

This is a solution whose concentration is known. The concentration can be:

- (i) In moles of the compound in a litre of solution (Molarity).
- (ii) In grams of the compound in a litre of solution.
- (iii) In grams of the compound in a given volume usually 250cm³ or 500cm³.

A standard solution is used to standard solutions whose concentration is not known (unstandard solution) during a process called volumetric analysis.

ACID-BASE TITRATION

Acid-base titration is the titration between an acid and an alkali. The reaction taking place in this titration is called **neutralization reaction**. The reaction is represented by the equation below.

$$H^{+}_{(aq)} + OH^{-}_{(aq)} \longrightarrow H_2O_{(l)}$$

The progress of the reaction is determined by adding an indicator to the solution in the conical flask.

At the end point, the indicator shows a sharp colour change. Students should take care when the end point is about to be reached. Add the solution from the burette drop by drop to avoid over shooting.

At this level, the indicator solutions used for titration are phenolphthalein and methyl orange indicators.

They show the following colours in acid and base media.

Indicator	Colour in acid medium	Colour in alkali medium
Phenolphthalein	Colourless	Purple
Methyl orange	Red/Pink	Yellow

Worked Examples

Example 1

You are provided with:

BA1, which is potassium hydroxide solution with unknown concentration.

BA2, which is 0.2M sulphuric acid.

You are required to determine the concentration of BA1

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA1** into a clean conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA2** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Results:

Volume of pipette used 25.0 cm³

Experiment	1	2	3
Final burette reading (cm ³)	15.20	17.60	21.60
Initial burette reading (cm ³)	0.00	2.00	6.00
Volume of BA2 used (cm ³)	15.20	15.60	15.60

Titre values of **BA2** used for average

15.60cm³ and 15.60cm³

Average volume of **BA2** used.

$$\frac{15.60 + 15.60}{2}$$

$$15.60 \text{cm}^3$$

(a) Write the equation for the reaction that took place between **BA1** and **BA2**.

$$2KOH_{(aq)} + H_2SO_{4(aq)} \longrightarrow K_2SO_{4(aq)} + 2H_2O_{(l)}$$

(b) Calculate the molarity of potassium hydroxide.

1000cm³ of solution contains 0.2 moles H₂SO₄

 1cm^3 of solution will contain $\frac{0.2}{1000}$ moles of H_2SO_4

15.60cm³ of solution will contain $\frac{0.2x15.60}{1000}$ moles of H₂SO₄

From the equation,

1 mole of sulphuric acid reacts 2 moles of potassium hydroxide.

0.00312 moles of sulphuric acid will react with $\frac{2 \times 0.00312}{1}$ moles of potassium hydroxide

25.0cm³ of solution contains 0.00624 moles of potassium hydroxide

 1cm^3 of solution will contain $\frac{0.00624}{25.0}$ moles of potassium hydroxide

 1000cm^3 of solution will contain $\frac{0.00624 \times 1000}{25.0}$ moles of potassium hydroxide 0.25 M

NB: M is the same as moles per litre, moles per 1000cm³ and moles per dm³

(c) Determine the concentration of KOH in grams per litre. (K = 39, O = 16, H = 1)

RFM of KOH =
$$(39 \times 1) + (16 \times 1) + (1 \times 1)$$

39 + 16 + 1
56

1 mole KOH weighs 56g

0.25 moles of KOH will weigh
$$\frac{0.25 \times 56}{1}$$
 g 14 gl⁻¹

Example 2

You are provided with:

FA1, which is a solution containing 3.7 grams per litre of a metal hydroxide M(OH)₂

FA2, which is 0.1M dilute hydrochloric acid

You are required to determine the atomic mass of metal M in M(OH)₂.

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **FA1** into a conical flask. Add 2-3 drops of methyl orange indicator and titrate with **FA2** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Results

Volume of pipette used 25.0 cm³

Experiment	1	2	3
Final burette reading(cm ³)	25.00	27.10	28.20
Initial burette reading (cm ³)	0.00	2.00	3.00
Volume of FA2 used (cm ³)	25.00	25.10	25.20

Tire values of **FA1** used for average

25.10cm³ and 25.20cm³

Average volume of FA1 used

$$\frac{25.10 \times 25.20}{2}$$

$$25.15 \text{cm}^{3}$$

(a) Write the equation for the reaction that took place between FA1 and FA2.

$$M(OH)_{2 (aq)} + HCl_{(aq)} \longrightarrow MCl_{2 (aq)} + 2H_2O_{(l)}$$

(b) Calculate the number of moles of FA1 that reacted with FA2.

1000cm³ of solution contains 0.1 moles of HCl

$$1 \text{cm}^3$$
 of solution will contain $\frac{0.1}{1000}$ moles of HCl

$$25.15$$
cm³ of solution will contain $\frac{0.1 \times 25.15}{1000}$ moles of HCl

0.002515moles of HCl

From the equation,

2 moles of HCl reacted with 1 mole of M(OH)₂

0.002515 moles of HCl will react with
$$\frac{1 \times 0.002515}{2}$$
 moles of M(OH)₂

0.0012575 moles of M(OH)₂

- (c) Determine the molarity of **FA1**.
 - 25.0cm³ of solution contains 0.0012575 moles of M(OH)₂

 $1 cm^3$ of solution will contain $\frac{0.0012575}{25.0}$ moles of M(OH)₂

 $1000 cm^3 \ of \ solution \ will \ contain \ \frac{0.0012575 \ x \ 1000}{25.0} \ moles \ of \ M(OH)_2$

0.05M

(d) Calculate the formula mass of FA1 and hence the relative atomic mass of metal M in $M(OH)_2$.

$$(H = 1, O = 16)$$

0.05 moles contain 3.7g M(OH)₂

1 mole will contain
$$\frac{3.7 \times 1}{0.05}$$

74g

Or

Use of formula

$$RFM = \frac{\text{Conc. in } gl^{-1}}{\text{Molarity}}$$
$$= \frac{3.7}{0.05}$$
$$= 74$$

$$M(OH)_2 = RFM$$

$$M + (16 \times 2) + (1 \times 2) = 74$$

 $M + 32 + 2 = 74$

$$M + 34 = 74$$

$$M = 74 - 34$$

$$M = 74$$

$$M = 40$$

Example 3

You are provided with:

BA1, which is a solution containing 4.0g solution hydroxide dissolved to make 500cm³ of the solution.

BA2, which is a solution containing 0.1M acid H_nX .

You are required to determine the basicity of an acid H_nX.

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA1** into a conical flask. Add 2-3 drops phenolphthalein indicator and titrate with **BA2** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Results

Volume of pipette used 25.0 cm³

Experiment	1	2	3
Final burette reading (cm ³)	25.40	25.00	27.00
Initial burette reading (cm ³)	0.00	0.00	2.00
Volume of BA2 used (cm ³)	25.40	25.00	25.00

Titre values of **BA2** used for average.

25.00cm³ and 25.00cm³

Average volume of BA2 used

$$\frac{25.00+25.00}{2}$$
25.00cm³

(a) Calculate the molarity sodium hydroxide (Na = 23, O = 16, H = 1)

500cm³ of solution contains 4.0g of NaOH

 1cm^3 of solution will contain $\frac{4.0}{500}$ g of NaOH

1000cm³ of solution will contain $\frac{4.0 \times 1000}{500}$ g NaOH

RFM of NaOH =
$$(23 \times 1) + (16 \times 1) + (1 \times 1)$$

= $23 + 16 + 1$
= 40

40g of NaOH contains 1 mole

1g of NaOH will contain
$$\frac{1}{40}$$
 moles

8g of NaOH will contain
$$\frac{1 \times 8}{40}$$
 moles

(b) Calculate the number of moles of BA1 that reacted

1000cm³ of solution contains 0.2 moles of NaOH

$$1 \text{cm}^3$$
 of solution will contain $\frac{0.2}{1000}$ moles of NaOH

$$25.0$$
cm³ of solution will $\frac{0.2 \times 25.0}{1000}$ moles of NaOH

0.005 moles of NaOH

(c) Calculate the number of moles of the acid that reacted.

1000cm³ of solution contains 0.1 moles of the acid

$$1 \text{cm}^3$$
 of solution will contain $\frac{0.1}{1000}$ moles of the acid

25.00cm³ of solution will contain
$$\frac{0.1 \times 25.00}{1000}$$
 moles of the acid

0.0025 moles of the acid

(d) Determine the basicity of the acid H_nX and hence write the equation of reaction between the acid and the base.

$$NaOH : H_nX$$

Mole ratio
$$\frac{0.005}{0.0025}$$
 : $\frac{0.0025}{0.0025}$

2 . 1

$$2NaOH_{(aq)} + H_2X_{(aq)} \longrightarrow Na_2X_{(aq)} + 2H_2O_{(l)}$$

Example 4

You are provided with the following;

BA1, which is a solution made by dissolving 4.8g of metal hydroxide MOH in one litre.

BA2, which is a 0.0625 moll⁻¹ sulphuric acid.

You are required to determine the relative atomic mass of M in MOH

Procedure:

Pipette 25cm³ (or 20cm³) of **BA1** into a clean conical flask and add 2-3drops of phenolphthalein indicator. Titrate this mixture with **BA2** from the burette. Repeat the titration until you obtain consistent results.

Record your results in the table below.

Results:

Volume of pipette used 25.0 cm³

Final burette reading (cm ³)	24.10	48.10	24.00
Initial burette reading (cm ³)	0.00	24.10	0.00
Volume of BA2 used (cm ³)	24.10	24.00	24.00

Volumes of BA2 used for calculating the average

24.00cm³ and 24.00cm³

Average volume of BA2 used

$$\frac{24.00+24.00}{2}$$

$$\frac{48.00}{2}$$

$$24.00 \text{cm}^3$$

Questions:

- (a) Calculate the;
 - (i) Number of moles of BA2 that reacted,

1000cm³ of solution contains 0.0625 moles of sulphuric acid.

 1cm^3 of solution will contain $\frac{0.0625}{1000}$ moles of sulphuri aid

 24.00cm^3 of solution will contain $\frac{0.0625 \times 24.00}{1000}$ moles of sulphuric aid 0.0015 moles of sulphuri acid

(ii) Number of moles of MOH in BA1 that reacted

$$2MOH_{(aq)} + H_2SO_{4\;(aq)} \xrightarrow{\hspace*{1cm}} M_2SO_{4\;(aq)} + 2H_2O_{\;(l)}$$

From the equation;

1 mole of sulphuric acid reacted with 2 moles of MOH

0.0015 moles of sulphuric acid will react with $\frac{2 \times 0.0015}{1}$ moles of MOH

0.003 moles of MOH

(iii) Molarity of BA1.

25.0cm³ of solution contains 0.003 moles of MOH.

 1cm^3 of solution will contain $\frac{0.003}{25.0}$ moles of MOH

 1000cm^3 of solution will contain $\frac{0.003 \times 1000}{25.0}$ moles of MOH

0.12M

(iv) Formula mass of MOH.

0.12 moles of MOH contains 4.8g

1 mole of MOH will contain
$$\frac{4.8}{0.12}$$
 g $40g$

(v) Relative atomic mass of M. (O = 16, H = 1)

MOH =
$$40$$

 $(M \times 1) + (16 \times 1) + (1 \times 1) = 40$
 $M + 16 + 1 = 40$
 $M + 17 = 40$
 $M = 40 - 17$
 $M = 23$

Example 5

You are provided with the following:

GA1, which is a solution made by dissolving 12.6g of a monobasic acid HNO_X to make 1 litre of solution.

GA2, which is a 0.2M sodium hydroxide solution.

You are required to determine the value of x in HNO_X

Procedure:

Pipette 25cm³ (or 20cm³) of **GA2** into a conical flask and add 2-3 drops of phenolphthalein indicator. Titrate this mixture with **GA1** from the burette. Repeat the titration until you obtain consistent results.

Record your results in the table below.

Results:

Volume of pipette used 25.0 cm³

Experiment	1	2	3
Final burette reading (cm ³)	25.20	25.00	25.00
Initial burette reading (cm ³)	0.00	0.00	0.00
Volume of GA1 used (cm ³)	25.20	25.00	25.00

Volumes of GA1 used to calculate the average

25.00cm³ and 25.00cm³

Average volume of **GA1** used

25.00+25.00

50.00

 25.00cm^3

Questions:

(a) Write the ionic equation for the reaction between the acid in **GA1** and sodium hydroxide in **GA2**.

$$H^+_{(aq)} + OH^-_{(aq)} \longrightarrow H_2O_{(l)}$$

- (b) Calculate the;
 - (i) Moles of sodium hydroxide in GA2 that reacted with acid.

1000cm³ of solution contains 0.2 moles of sodium hydroxide

 1cm^3 of solution will contain $\frac{0.2}{1000}$ moles of sodium hydroxide

 25.0cm^3 of solution will contain $\frac{0.2 \times 25.0}{1000}$ moles of sodium hydroxide

0.005 moles of sodium hydroxide

(ii) Number of moles of HNO_X in **GA1** that reacted.

From the equation;

1 mole sodium hydroxide reacted with 1 mole of HNO_X

0.005 moles of sodium hydroxide will react with $\frac{1 \times 0.005}{1}$ moles of HNO_X 0.005 moles of HNO_X

(iii) Molarity of GA1.

25.00cm³ of solution contain 0.005 moles of HNO_X

 1cm^3 of solution will contain $\frac{0.005}{25.00}$ moles of HNO_X

 1000cm^3 of solution will contain $\frac{0.005 \times 1000}{25.00}$ moles of HNO_X 0.2M

- (iv) Formula mass of HNO_X
- 0.2 moles of HNO_X contain 12.6g

1 mole of HNO_X will contain $\frac{12.6}{0.2}$ g 63g

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(v) Value of x.
$$(O = 16, H = 1, N = 14)$$

$$HNO_X = 63$$

$$(1 x1) + (14 x1) + (X x 16) = 63$$

$$15 + 16X = 63$$

$$16X = 63 - 15$$

$$16X = 48$$

$$X = 3$$

Example 6

You are provided with the following

BA1, which is a solution containing 10g of solid Y.10H₂O in 500cm³ of solution.

BA2, which is 0.1M hydrochloric acid solution.

(Ratio of $Y.10H_2O$: HCl is 1:2 and Y = 202)

You are required to determine the percentage purity of solid Y.10H₂O

Procedure:

Pipette 25cm³ (or 20cm³) of **BA1** into a clean conical flask. Then add 2-3 drops of methyl orange indicator and titrate with solution **BA2** from the burette. Repeat the titration until you obtain consistent results. Enter your results in the table below.

Results:

Volume of pipette used 25.0 cm³

Final burette reading (cm ³)	15.40	30.90	46.30
Initial burette reading (cm ³)	0.00	15.40	30.90
Volume of BA2 used (cm ³)	15.40	15.50	15.40

Titre values used for calculating average volume of BA2.

15.50cm³ and 15.40cm³

Average volume of BA2

15.50+15.40

2

15.45cm³

Questions:

- (a) Calculate the;
 - (i) Moles of hydrochloric acid in **BA2** that reacted 1000cm³ of solution contains 0.1 moles of hydrochloric acid

 1cm^3 of solution will contain $\frac{0.1}{1000}$ moles of hydrochloric acid

15.45cm³ of solution will contain $\frac{0.1 \times 15.45}{1000}$ moles of hydrochloric acid

0.001545 moles of hydrochloric acid

(ii) Moles of $Y.10H_2O$ in **BA1** that reacted with hydrochloric acid From the mole ratio;

2 moles of the hydrochloric acid reacted with 1 mole of Y.10H₂O

0.001545 moles of hydrochloric acid will react with $\frac{1 \times 0.001545}{2}$ moles of Y.10H₂O

0.000772 moles of Y.10H2O

(iii) Molarity of BA1 and hence the percentage purity of the solid, $Y.10H_2O$

25.0cm³ of solutions contain 0.000772 moles of Y.10H₂O

 1cm^3 of solution will contain $\frac{0.000772}{25.0}$ moles of Y.10H₂O

 $1000 cm^3 \ of \ solution \ will \ contain \ \frac{0.000772 \ x \ 1000}{25.0} \ moles \ of \ Y.10H_2O$ 0.03088M

RFM of $Y.10H_2O = 202 + 180$ = 382gs

1 mole of Y.10H₂O weighs 382g

0.03088 mole of Y.10H₂O weighs 382 x 0.03088

11.7gl⁻¹

Also

 $500cm^3$ of **BA1** contain 10g of $Y.10H_2O$

 $1 \text{cm}^3 \text{ of } \mathbf{BA1} \text{ will contain } \frac{10}{500} \text{ g of } \text{Y.}10\text{H}_2\text{O}$

 $1000 \text{cm}^3 \text{ of } \textbf{BA1} \text{ will contain } \frac{10 \times 1000}{500} \text{ g of } \text{Y}.10 \text{H}_2 \text{O}$

Therefore the percentage purity of $Y.10H_2O = \frac{11.7 \times 100}{20} \%$

58.5%

NB: Percentage impurity = 100 – (percentage purity)

PRACTICAL SCHEDULES

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You are provided with:

BA1, which is 0.1M hydrochloric acid

BA2, which is sodium hydroxide

You are required to determine the concentration of solution of BA2 in

- (i) Moldm⁻³
- (ii) gl⁻¹

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA2** in a conical flask. Add 2-3 drops of phenolphthalein indicator and titrate with **BA1** from the burette.

Repeat the procedure to obtain consistent results.

Record your results in the table below.

Results

Experiment	1	2	3
Final burette reading			
(cm^3)			
Initial burette reading			
(cm^3)			
Volume of BA1 used			
(cm^3)			

Titre values of BA1 used for average					
	lume of BA1 used			 	•••••
Questions					
Calculat	te the;				
(i)	Number of moles of	of BA1 that rea	cted.		
	• • • • • • • • • • • • • • • • • • • •			 	
	• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •		 	

(ii) Number of moles of BA2 that reacted.		
(iii) Concentration of BA2 in Moldm ⁻³		
(iv) Concentration BA2 in gl ⁻¹		

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You are provided with:

BA3, which is 0.2M solution of hydrochloric acid

BA4, which is a made by dissolving 10.6g of sodium carbonate to make 1 litre if solution.

You are required to determine the concentration of BA3 in

- (i) $Mold m^{-3}$
- (ii) gl⁻¹

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA4** into a conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA3** from the burette.

Repeat procedure to obtain consistent results.

Record your results in the table below.

Titre values of **BA3** used for average

Volume of pipette used		cm	1^3	
Experiment	1	2	3	
Final burette reading (cm ³)				
Initial burette reading (cm ³)				
Volume of BA3 used (cm ³)				
	•	•	•	

Average	e volume of BA3 us	sed				
						• • • • • • • • • • • • • • • • • • • •
Questic (a) (i)	ons: Calculate the; Number of mole	es of BA4 that re	eacted (Na = 2	3, C = 12, O =	16)	

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(ii)	Number of moles of BA3 that reacted.
•••••	
` ′	Concentration of BA3 in moldm ⁻³
	Concentration of BA3 in gl^{-1} (Cl = 35.5, H = 1)

You are provided with:

BA1, which is 0.2M hydrochloric acid solution.

BA2, which is a salt solution made by dissolving 10.6g of M₂CO₃ in 1 litre of solution.

You are required to determine the relative atomic mass of metal M in M₂CO₃.

Procedure:

Results:

Pipette 25.0cm³ (or 20.0cm³) of **BA2** into a conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA1** from the burette.

Repeat procedure to until you obtain consistent results.

Record your observations in the table below.

Volume of pipette used			cm ³	
Experiment	1	2	3	
Final burette reading (cm ³)			
Initial burette reading (cm ³)				
Volume of BA1 used (cm ³)				
Titre values of BA1 used Average volume of BA1				
Questions:				
(a) Calculate the;				
	moles of BA1	that reacted		
		•••••		• • • • • • • • • • • • • • • • • • • •

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	(ii) 	Write the equation reaction between BA1 and BA2
	(iii)	Write the mole ratio of the reaction
(b)	Calcu	late the number of moles BA2 that reacted
(c)	Calcu	late the molarity of BA2
(4)		late the relative formula mass of M-CO-
(u)		late the relative formula mass of M ₂ CO ₃

(e) Determine the relative at	omic mass c	of M ($C = 12$,	O = 16)	
		• • • • • • • • • • • • • • • • • • • •		
Practical 4				
You are provided with:				
BA3, which is a solution contain	ing 1.7g of	OH⁻ ions per	litre	
BA4 , which is a dibasic acid H_2	X containing	9.8gl ⁻¹		
You are required to determine th	e atomic ma	ass of X in the	e acid (H = 1, $O = 16$)
Procedure:				
Pipette 25.0cm ³ (or 20.0cm ³) of 3	RA3 into a a	conical flask	Add 3 drops phenolr	hthalein indicator
and titrate with BA4 from the bu		comear mask.	rad 5 drops phenoip	minarem mareator
Repeat the procedure until you o		tent results.		
Record your results in the table b				
·				
Results:			2	
Volume of pipette used	•			
Experiment	1	2	3	
Final burette reading (cm ³)				
Initial burette reading (cm ³)				
Volume of BA4 used (cm ³)				
			<u> </u>	
Titre values of BA4 used for av	rerage			
	• • • • • • • • • • • • • • • • • • • •			
Average volume of BA4 used				
Write the ionic equation for the r	reaction bety	ween BA3 and	d BA4	
	• • • • • • • • • • • • • • • • • • • •			

Qu	estion	ns:
		alculate the;
	(i)	Molarity of BA3
	(::)	Number of males of DA2 that reacted
	(ii)	Number of moles of BA3 that reacted
	(iii)	Number of moles BA4 that reacted
	(111)	rumber of moles b /14 that reacted
	(iv)	Molarity of BA4

(v)	Relative atomic mas	ss of X in t	the acid		
•••••			• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	
	_				
Practical					
-	rovided with:		af dibasia sai	d V all O asa	litus of colution
	ch is a solution contai ch is made dissolving		=	-	
	equired to determine t	_	-		of the solutions
	f Y.nH ₂ O reacts 2 mo				= 90)
(1 111010 01	1 1		- 11) WI 0111WO, 111	111111111111111111111111111111111111111	<i>></i> • <i>/</i>
Procedur	e:				
Pipette 25	$.0 \text{cm}^3 \text{ (or } 20.0 \text{cm}^3 \text{) or}$	f BA6 in a	conical flask.	Add 3 drops of	f phenolphthalein indicator
and titrate	with BA5 from the b	ourette.			
Repeat pro	ocedure until you obta	ain consist	ent results.		
Record yo	our results in the table	below.			
.					
Results:	C : 44 -			3	
	f pipette		2		
Experi		1	2	3	
Final b	ourette reading (cm ³)				
_	burette reading				
(cm ³)	e of BA5 used				
(cm ³)	ie of Brie used				
		•		•	
Titre value	es of BA5 used for av	erage			
Average v	volume of BA5 used				

iestior	ns:
(a) C	Calculate,
(i)	The molarity of sodium hydroxide
• • • • • • • • • • • • • • • • • • • •	
• • • • • • •	
• • • • • • •	
(ii)	The number of moles of sodium hydroxide that reacted
•••••	
(iii)	The number of moles of Y.nH ₂ O that reacted
(111)	The humber of moles of 1.htt20 that reacted
• • • • • • • • • • • • • • • • • • • •	
• • • • • • • • • • • • • • • • • • • •	
• • • • • • • • • • • • • • • • • • • •	
(iv)	The molarity of Y.nH ₂ O
(' /	

(v)	The value of value	e of n in Y.nl	H_2O		
Practical 6					
You are pro	vided with:				
-	is 0.3M sodium	hydroxide so	lution		
BA2, which	is 0.2M solution	of acid H _n A			
You are req	uired to determin	e the stoichio	metry reaction be	etween the acid H	nA and sodium
hydroxide.					
Procedure:					
Pipette 25.0	$0 \text{cm}^3 \text{ (or } 20.0 \text{cm}^3\text{)}$	of $BA1$ into	conical flask. Ad	d 2-3 drops of me	thyl orange
indicator an	d titrate with BA ?	2 from the bu	rette.		
Repeat the p	procedure until yo	ou obtain con	sistent results		
Record you	r results in the tab	ole below.			
Results:					
Volume of 1	pipette used			cm ³	
Experim	ent	1	2	3	
Final but	rette reading				
(cm ³)					
	rette reading				
(cm ³)	of BA2 used				
(cm ³)	of BAZ used				
(CIII)					
Titre values	of BA2 used for	average			
Average vo	lume of BA2 used	 1		• • • • • • • • • • • • • • • • • • • •	
		-			
		• • • • • • • • • • • • • • • • • • • •			• • • • • • • • • • • • • • • • • • • •

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()	HECTIONC	•
v	uestions	•

(i)	Number of moles of sodium hydroxide that reacted
(ii)	Number of moles of H _n A that reacted
(L) T	Determine the mede action of the solid II. A to coding budgeride
	Determine the mole ratio of the acid H _n A to sodium hydroxide
	Determine the value of n in H _n A
	Setermine the value of h in Fig. 1
(4) 1	Write the equation for reaction
(u) \	Write the equation for reaction

You are provided with:

BA1, which is 0.2 moles per litre of the acid H_nX

BA2, which is 0.5M solution of sodium hydroxide

You are required to determine the basicity, n of the acid H_nX

Procedure:

Pipette 25.0cm^3 (or 20.0cm^3) of **BA2** into a conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA1** from the burette

3

Repeat the procedure until you obtain consistent results

1

Record your results in the table below.

Result	S	:
--------	---	---

Experiment

	burette reading			
(cm ³))			
Initia	l burette reading			
(cm ³				
Volu	me of BA1 used			
(cm ³))			
Titre val	lues of BA1 used for a	verage		
Average	volume of BA1 used			
Questio				
-	Calculate the;			
-		of sodium hydro	oxide that reacted	
(a) (Calculate the;	of sodium hydro	oxide that reacted	
(a) (Calculate the;	of sodium hydro	oxide that reacted	
(a) (Calculate the;	of sodium hydro		
(a) (Calculate the;	of sodium hydro		
(a) (Calculate the;	of sodium hydro		
(a) (Calculate the;	of sodium hydro		

(ii)	The number of moles of H _n X that reacted
• • • • • • • • • • • • • • • • • • • •	
• • • • • • • •	
• • • • • • • • • • • • • • • • • • • •	
(b) De	etermine,
(i)	The mole ratio of the acid H_nX to sodium hydroxide
	······································
(ii)	The basicity, n of the acid H _n X
• • • • • • • • • • • • • • • • • • • •	
• • • • • • • •	
(c) W	rite an ionic equation of the reaction between the acid and the base (sodium hydroxide)

You are provided with the following:

BA1, which is a solution containing 12.6gl⁻¹ of a dibasic acid of formula H₂C₂O₄ .xH₂O.

BA2, which is 0.1M sodium hydroxide solution.

You are required to determine the value of x

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA2** into a clean conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA1** from the burette.

Repeat the procedure until you obtain consistent results.

1

Record your results in the table below.

Experiment

Final burette reading

(cm ³)			
Initia	al burette reading			
(cm ³))			
	me of BA1 used			
(cm^3))			
Titre val	lues of BA1 used for a	average		
1100 100	ides of Bill dsed for t	reruge		
••••••			 	
Δverage	volume of BA1 used		 	
Average	volume of DAT used			
	• • • • • • • • • • • • • • • • • • • •		 	•••••
		•••••	 	
Questio				
Calculat				
(i)	Number of moles	of BA2 reacted		
••••			 	
••••			 • • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •
••••			 	

(ii)	Number of moles of BA1 that reacted with BA2
(iii)	Concentration moll ⁻¹ of BA1
(iv)	The R.M.M of the acid and hence the value of x.
•••••	

3

Practical 9

You are provided with:

BA1, which is a solution of acid H_nX of concentration 0.625M.

BA2, which is a 0.5M sodium hydroxide solution.

You are required to determine the reaction ratio for BA1 and BA2.

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA2** into a clean conical flask. Add 2-3 drops of phenolphthalein indicator and titrate with **BA1** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Resul	lts:
-------	------

Experiment

	i mai saistie isaamg (em)			
	Initial burette reading (cm ³)			
	Volume of BA1 used (cm ³)			
Ti 	tre values of BA1 used for ave	erage	 	
 A'	verage volume of BA1 used		 	
Q	uestions:			
	alculate the;			
	(i) Number of moles of	BA1 that reacted		

(ii)	Number of moles					
(iii)	The value of n (b	easicity of H	$(X_1$			
	The mole ratio.	• • • • • • • • • • • • • • • • • • • •			•••••	
Practica	1 10					
	provided with:					
-	ich is a solution of	an impure ac	eid H ₂ X of cond	centration 2gd	m^{-3} .	
BA2, wh	ich is a solution of	sodium hydr	ogen carbonate	e made by diss	solving 4.2g	in 1dm ⁻³ .
You are	required to determin	ne the percer	ntage purity of	BA1.		
Procedu						
_	5.0cm ³ (or 20.0cm ³			al flask. Add 2	2-3 drops me	thyl orange
	and titrate with BA ne procedure until y					
-	our results in the ta		nsistem results	•		
record y	our results in the tu	ore below.				
Results:						
Volume	of pipette used				cm ³	
Expe	riment	1	2	3		
Final (cm ³)	burette reading					
Initial (cm ³)	burette reading					
Volum (cm ³)	ne of BA1 used					

Ti	Titre values of BA1 used for average			
		volume of DA1 wood		
Α	verage	volume of BA1 used		
• •				
• •				
O	uestion	is:		
•		alculate the;		
	(i)	Number of moles of BA2 that reacted		
	• • • • • •			
	• • • • • • •			
	• • • • • • •			
	•••••			
	••••			
	••••			
	• • • • • • •			
	(ii)	Number of moles of BA1 that reacted		
	• • • • • •			
	• • • • • •			
	• • • • • • •			
	• • • • • •			
	• • • • • •			
	• • • • • •			
	(iii)	Molarity of BA1 $(X = 88g)$		

You are provided with the following:

BA3, which is 0.04M hydrochloric acid solution

BA4, which is a solution made by dissolving 3.6g of impure Na₂CO₃.10H₂O to make 500cm³ of a solution.

You are required to determine the percentage impurity of **BA4**.

Procedure:

Results:

Pipette 25.0cm³ (or 20.0cm³) of **BA4** into a conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA3** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Volume of pipette used	cm ³
Experiment 1 2 3	

Final burette reading (cm³)

Initial burette reading (cm³)

Volume of **BA3** used (cm³)

Titre values of BA3 used for average Average volume of BA3 used					
Question (a) V	ons: Write an equation for the read				
	Calculate the;				
(i)	Number of moles of BA3				

(ii)	Number of moles of Na ₂ CO ₃ .10H ₂ O in 250cm ³ of a solution.
•••••	
(iii)	Percentage purity of Na ₂ CO ₃ .10H ₂ O.
•••••	
•••••	(Na = 23, C = 12, O = 16, H = 1)

You are provided with the following:

BA3, which is a 0.2M monobasic acid.

BA4, which is a solution made by dissolving 11.6g of Na₂CO₃.nH₂O to make a litre of a solution.

You are required to determine the value n

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA4** into a conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA3** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Resul	ts:
-------	-----

icsuits.				
Volume of pipette used			cm ³	
Experiment	1	2	3	
Final burette reading				
(cm^3)				
Initial burette reading				
(cm ³)				
Volume of BA3 used (cm ³)				
Fitre values of BA3 used for the value of BA3 used for the values of	ed			
Questions:				
(a) Calculate the;(i) Moles of BA3 th	not manatad			
(i) Moles of BA3 th	iai reacted			
			• • • • • • • • • • • • • • • • • • • •	
			• • • • • • • • • • • • • • • • • • • •	
			• • • • • • • • • • • • • • • • • • • •	
			• • • • • • • • • • • • • • • • • • • •	

(ii)	Moles of Na ₂ CO ₃ .10H ₂ O in 1000cm ³ of a solution.
(iii)	Formula mass of BA4 and hence the value of n.
	(Na = 23, C = 12, O = 16, H = 1)

You are provided with the following:

BA1, which is a solution containing 6.2g of Y.nH2O in one litre

BA2, which is 0.1M hydrochloric acid

(Y =106, 1 mole of Y.nH₂O reacts with 2 moles of hydrochloric acid)

Resul	lte•
Kesu	us.

You are required to determine the	ne number	of moles of wat	ter of crystallizati	on in BA1
Procedure:				
Pipette 25.0cm ³ (or 20.0cm ³) of	BA1 into	a clean conical	flask. Add 2-3 dr	ops of
phenolphthalein indicator and ti	trate with	BA2 from the b	urette.	
Repeat the procedure until you o	obtain con	sistent results.		
Record your results in the table	below.			
Results:				
Volume of pipette used			cn	1^3
Experiment	1	2	3	
Final burette reading (cm ³)				
Initial burette reading (cm ³)				
Volume of BA2 used (cm ³)				
Average volume of BA2 used				
Questions:		•••••		
(a) Calculate the:				
(i) Number of moles of	BA2 that 1	reacted.		

(ii)	Number of moles of BA1 that reacted with the acid
• • • • • • • • • • • • • • • • • • • •	
• • • • • • • • • • • • • • • • • • • •	
• • • • • • • • • • • • • • • • • • • •	
• • • • • • • • • • • • • • • • • • • •	
(iii)	Concentration of BA1 in moles per litre
	<u>.</u>
• • • • • • • • • • • • • • • • • • • •	
(1V)	Value of \mathbf{n} in Y.nH ₂ O.
• • • • • • • • • • • • • • • • • • • •	
•••••	

You are provided with the following:

BA1, which is a solution containing 12.5gl⁻¹ of a mixture of anhydrous sodium carbonate and sodium chloride.

BA2, which is a 0.1M hydrochloric acid solution.

You are required to determine the composition of sodium chloride in **BA1**.

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA1** into a clean conical flask. Add 2-3 drops methyl orange indicator and titrate with **BA2** from the burette.

Repeat the procedure until you obtain consistent results

Record your results in the table below.

R	esults:			
V	olume of pipette used			cm ³
	Experiment	1	2	3
	Final burette reading (cm ³)			
	Initial burette reading (cm ³)			
	Volume of BA2 used (cm ³)			

Titre val	es of BA2 used for average
	olume of BA2 used
Question	s: lculate the;
(i)	Number of moles of BA2 that reacted.

(ii)	Number of moles of BA1 that reacted with BA2 .
	Concentration in moles per litre of RA1
	Concentration in moles per litre of BA1
	Percentage of sodium chloride in BA1 .

You are provided with:

BA1, is an aqueous solution of XCO₃ prepared by dissolving 2.65g of it in 250cm³ of distilled water.

BA2, is an aqueous solution of hydrochloric acid prepared by dissolving 7.3g of it in a litre of distilled water.

You are required to determine the relative atomic mass of X in XCO₃.

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA1** into a clean conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA2** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Results:

Experiment

(C1	nal burette reading m ³)			
Ini	itial burette reading m ³)			
	olume of BA2 used m ³)			
Titre	values of BA2 used for a	verage		
Avera	age volume of BA2 used			
_	tions:			
(a) Calculate the;	~! - 25 5 H -		
_) Calculate the;	Cl = 35.5, H =	1)	
(a) Calculate the;	Cl = 35.5, H =	1)	
(a) Calculate the;	Cl = 35.5, H =	1)	
(a) Calculate the;	Cl = 35.5, H =	1)	
(a) Calculate the;	Cl = 35.5, H =	1)	

(ii)	Number of moles of BA2 that reacted
• • • • • • • • • • • • • • • • • • • •	
(iii)	Number of BA1 that reacted (BA1 : BA2 = $1 : 2$)
• • • • • • • • • • • • • • • • • • • •	
• • • • • • • • • • • • • • • • • • • •	
(iv)	Concentration of BA1 in moles per litre
(b) De	etermine the;
(i)	Relative formula mass of XCO ₃
• • • • • • • • • • • • • • • • • • • •	
• • • • • • • • • • • • • • • • • • • •	
• • • • • • • •	

(ii) Value of X in XCO ₃	(C - 12, C - 1)	16)			
(II) Value of X III ACO3	(C = 12, O = 1)	10)			
			•••••	,	
			•••••	,	
Procedure 16					
You are provided with:					
BA1 , which is a sodium hydrox	ide solution of	molarity 0.21	M		
BA2 , which is made by dissolvi		•			
You are required to determine the		=	c of solution.		
Procedure:	ie busierty, x o	or ucid II _X I.			
Pipette 25.0cm ³ (or 20.0cm ³) of	BA1 into a co	nical flask A	dd 2-3 drops of pher	nolphthalein	
indicator and titrate with BA2 fr			add 2 3 drops of photo	тогринатен	
Repeat the procedure until you of					
Record your results in the table		in results.			
Results:	ociow.				
Volume of pipette used			cm^3		
Experiment Experiment	1	2	3		
	1	2	3		
Final burette reading (cm ³)					
Initial burette reading					
(cm ³)					
Volume of BA2 used (cm ³)					
Titre values of BA2 used for avo	orogo				
Title values of BA2 used for avo	erage				
Avarage volume of PA2 used				• • • • • • • • • • • • • • • • • • • •	
Average volume of BA2 used					

estion (a) W	Write an equation for the reaction between the acid H_XY and so	dium hydroxide.
	Calculate the;	
(i)	Number of moles of BA1 that reacted	
(ii)	Number of moles of acid H _X Y that reacted with sodium hyd	roxide.
(c) D	Deduce the molarity BA2 .	
(d) D	Determine the value x in the acid H_XY (H = 1, Y = 96)	

You are provided with:

BA1, which is a solution made by dissolving 3.35g of solid Na₂Y in 250cm³ of distilled water. **BA2**, which is 0.1M hydrochloric acid.

You are required to determine the ratio of reaction between Na_2Y and hydrochloric acid (Y = 96, Na = 23)

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA1** into a conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA2** from the burette.

2

3

1

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Experiment

F	inal burette reading				
	cm^3)				
I	nitial burette reading				
(cm^3)				
	olume of BA2 used				
	cm^3)				
		ı	I.	1	_
Titre	e values of BA2 used for a	verage			
1111	values of Briz asea for a	verage			
• • • • •		• • • • • • • • • • • • • • • • • • • •		 	
• • • • •				 	
Ave	rage volume of BA2 used				
	etions•			 	
 Que	stions:			 	
 Que (a) Calculate the;			 	
 Que (
 Que (a) Calculate the;				
 Que (a) Calculate the;			 	
 Que (a) Calculate the;				
 Que (a) Calculate the;				
 Que (a) Calculate the;				
 Que (a) Calculate the;				
 Que (a) Calculate the;				
 Que	a) Calculate the;				

(ii)	Number of moles of Na ₂ Y in BA1
(iii)	Number of moles of hydrochloric acid in BA2 that reacted with Na ₂ Y
(b) D	Determine the ratio of reaction between Na ₂ Y and hydrochloric acid.
• • • • • •	

You are provided with the following.

BA1, is a solution of sodium carbonate of molarity 0.1M

BA2, is a solution of hydrochloric acid of molarity 0.2M

You are required to determine the reaction ratio between sodium carbonate and hydrochloric acid.

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA1** into a clean conical flask. Add 2-3 drops methyl orange indicator and titrate with **BA2** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Results:					
Volume of pipette used			c	m^3	
Experiment	1	2	3		
Final burette reading (cm ³))				
Initial burette reading (cm ³)					
Volume of BA2 used (cm ³)				
Questions: (a) Determine the;					
(i) Number of moles (of sodium c	arbonate that rea	acted with the a	cid	
(1) I variable of moles (or sourant co				

(ii)		·			
• • • •					
••••	•••••				
••••					
••••					
••••					
(iii					
••••					
••••					
Praction					
	which is a solution cont	aining 16.0g of a	n impure sample	e of sodium carbon	ate -10-water,
	$0_3.10\mathrm{H}_2\mathrm{O}$	£ 1 1. 1	: 1		
	which is a 0.1M solution	•		a a ula a ura e a	
Y ou ar	e required to determine	tne percentage p	urity of socium	carbonate.	
_	25.0cm ³ (or 20.0cm ³)			2-3 drops of methy	vl orange
	or and titrate with BA4				
	the procedure until you		it results.		
Record	l your results in the tabl	e below.			
Result	s:				
Volum	e of pipette used			cm ³	
Fina (cm	al burette reading				
Init (cm	ial burette reading				
	ume of BA4 used			_	
(cm					

Titre values of BA4 used for average	
Average volume of BA4 used	
Questions:	
carbonate.	
rerage volume of BA4 used lestions: (a) Write the ionic equation for the reaction between hydrochloric acid and sodium	
(ii) The mass of solution carbonate in one inte. (iva = 25, $C = 12$, $O = 10$, $\Pi = 1$)	

(c) Determine the percentag	e purity of sodiun	n carbonate in the	sample.	
•••••				,
Practical 20				
You are with the following:				
BA1 , which is made by dissolvi	ng 5.3g of a carbo	onate of the Y ₂ CO	3 in 1 litre of th	ne solution.
BA2 , which is 0.1M hydrochlor				
You are required to determine the		mass of Y		
Procedure: Pipette 25.0cm³ (or 20.0cm³) of indicator and titrate with BA2 fr Repeat the procedure until you of Record your results in the table	rom the burette. obtain consistent r		lrops of methyl	orange
Results:				
Volume of pipette used			cm ³	
Final burette reading (cm ³)				
Initial burette reading (cm ³)				
Volume of BA2 used (cm ³)				
Titre values of BA2 used for ave	erage			
Average volume of BA2 used.			•••••	•••••
	• • • • • • • • • • • • • • • • • • • •			

estion (a) W	Nrite equation for the reaction that took place.
• • • • • •	
	alculate the;
(i)	Number of moles of BA2 that reacted
(1)	Trumber of moles of 2122 that reacted
(ii)	Number of moles of Y ₂ CO ₃ that reacted
(iii)	Molar concentration of Y_2CO_3 in BA1 .
(c) D	Determine the relative atomic mass of Y

You are provided with the following

BA1, which is 0.02M hydrochloric acid

BA2, which is a solution made by dissolving 1.05g of a metal carbonate MCO₃ to make 0.5 litres of aqueous solution.

You are required to determine the relative atomic mass of metal, M in the metal carbonate.

P	r	Λ	c	Δ	h	11	r	Δ	•
L	ı	v	v	U	u	u	L	c	٠

Pipette 25.0cm³ (or 20.0cm³) of **BA2** into a conical flask. Add 3drops of phenolphthalein indicator and titrate with **BA1** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Result:

Fina					
	burette reading (cm ³)				
Initia (cm ³	al burette reading				
Volu	me of BA1 used (cm ³)				
Titre va	lues of BA1 used for ave	•			
	e volume of BA1 used				
Questio			can DA1 and D		
_	Write the equation for the	e reaction betw	een DAT and D	0 A 2.	
-	Write the equation for the	e reaction betw			
(a) \	Write the equation for the determine: The number of mole				

(ii)	The number of moles metal carbonate MCO ₃ that reacted.
	The concentration in moles per litre of the metal carbonate MCO ₃ .
	alculate the;
(i)	Relative molecular mass of MCO ₃
(ii)	The relative atomic mass of M in the metal carbonate MCO ₃ .
•••••	
•••••	

You are provided with the following:

CA1, which is a solution of carbonate ions, CO₃²-

CA2, which is a solution containing 16.1gl⁻¹ of a metal sulphate, MSO₄.

CA3, which is a 0.2M hydrochloric acid solution.

You are required to determine the atomic mass of M in MSO₄.

The carbonate ions in solution react with an aqueous solution of the metal sulphate according to the equation.

$$CO_3^{2-}(aq) + MSO_4(aq) \longrightarrow MCO_3(s) + SO_4^{2-}(aq)$$

The metal carbonate produced reacts with dilute hydrochloric acid according to the equation.

$$MCO_{3(s)} + 2HCl_{(aq)} \longrightarrow MCl_{2(aq)} + H_2O_{(l)} + CO_{2(g)}$$

Procedure:

- (a) Pipette 25.0cm³ (or 20.0cm³) of **CA1** into a conical flask.
- (b) Using a 50cm³ measuring cylinder, transfer 25cm³ of **CA2** into the conical flask in (a) .shake the contents in the conical flask and allow to stand, label the resultant mixture **CA4**.
- (c) Add 3-4 drops of methyl orange indicator to the resultant mixture in **CA4** in (b) and then titrate using solution **CA3** from the burette until you reach the end point.
- (d) Repeat the procedure (a) to (c) to obtain consistent results.

Record your results in the table below.

Results:

Volume of pipette used	 	cn
Final burette reading (cm ³)		
Initial burette reading (cm ³)		
Volume of CA3 used (cm ³)		

Titre	e val	lues	of	CA3	use	d fo	r av	erag	ge									
Ave										 	 	••••	 • • • • •	 	 	 	 	 ••••
• • • • •																		
• • • • •																		
					. .					 	 		 	 	 	 	 	

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Questions:

(a) C	Calculate the;
(i)	Number of moles of hydrochloric acid in CA3 used.
(ii)	Number of moles of MCO ₃ that reacted with HCl.
	Name have a formation of model and all the data are and d
(iii)	Number of moles of metal sulphate that reacted
(iv)	
(b) D	Determine the atomic mass of M in MSO ₄ . $(S = 32, O = 16)$

You are provided with:

BA1, is an aqueous solution of XCO₃ prepared by dissolving 5.3g of it in 500cm³ of distilled water.

BA2, is an aqueous solution of 0.2M hydrochloric acid.

You are required to determine the value of X in XCO₃.

Procedure:

Results:

Pipette 25.0cm³ (or 20.0cm³) of **BA1** into a conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA2** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Volume	e of pipette used		 	cm ³
Fina	l burette reading (cm ³)			
Initi	al burette reading (cm ³)			
Volu	ime of BA2 used (cm ³)			
Titre va	llues of BA2 used for avera	nge	 	
Average	e volume of BA2 used		 	
Questic	ons:		 	
(a)	Calculate the,			
(i)	Number of moles of B .	A2 that reacted.		
••••				
• • • • •			 	

(ii)	Number of moles of BA1 that reacted (BA1 : BA2 = 1: 2)
	Concentration of BA1 in moles per litre.
	Relative formula mass of XCO ₃
(v)	Value of X in XCO ₃

You are provided with the following:

BA1, which is a solution made by dissolving 3.45g of a hydrated salt X.nH₂O in 250cm³ of water.

BA2, which is a 0.1M hydrochloric acid.

You are required to determine the value of n in the salt.

Procedure:

Pipette 25cm³ (or 20cm³) of **BA1** into a conical flask. Add 2-3 drops of methyl orange indicator

and titrate w	with BA2 from the bu	ırette.			
Repeat the p	procedure until you o	btain consistent	results		
Record your	r results in the table l	below.			
Results:					
Volume of p	pipette used			cm ³	
Final but	rette reading (cm ³)				
Initial bu	rrette reading (cm ³)				
Volume	of BA2 used (cm ³)				
Titre values	of BA2 used for ave	erage			
	lume of BA2 used				
Questions:					
	culate the;			_	
(i)	Number of moles	of hydrochloric	acid that reacte	d.	
••••		•••••		• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •
••••					
••••		•••••			
••••		•••••			

	(ii)	Number of moles of X.nH ₂ O that reacted. (1 mole of X.nH ₂ O reacts with 2 moles of hydrochloric acid).
	(iii)	Number of moles of X.nH ₂ O in 250cm ³ of BA1 .
(b)		mine the value of n in X.nH ₂ O. (H = 1, O = 16, X = 106)
• • • •		

You are provided with:

BA1, which is a solution containing 0.1 moles per dm³ of a monobasic acid, Q.

BA2, which is a solution made by dissolving 5.0gl⁻¹ of substance Y, which is an impure sodium hydroxide.

You are required to determine the percentage purity of Y.

Procedure:

Pipette 25.0cm³ (20.0cm³) of **BA2** into a conical flask. Add 2-3 drops of methyl orange indicator and titrate with **BA1** from the burette.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Final burette reading (cm³)

Resu	lts:

rette reading (cm ³)				
of BA1 used (cm ³)				
of BA1 used for avera	ge			
ume of BA1 used	•••••	•••••		
	f sodium hydro	xide that reacted	1	
	of BA1 used (cm ³) of BA1 used for avera ume of BA1 used	of BA1 used (cm³) of BA1 used for average ume of BA1 used ulate the; Number of moles of sodium hydro	of BA1 used (cm³) of BA1 used for average ume of BA1 used ulate the; Number of moles of sodium hydroxide that reacted	of BA1 used (cm ³) of BA1 used for average ume of BA1 used

	(ii)	Concentration of sodium hydroxide per litre of BA2 .
	(iii)	Mass of sodium hydroxide in one litre of BA2 . (H = 1; O = 16; Na = 23)
(b)	Detern	nine the percentage purity of Y.

You are provided with FA1 and FA2.

FA1, is a solution made by dissolving 2.0g of sodium hydroxide to make 500cm³ of solution.

FA2, is a solution made by dissolving 13.2g of an impure dibasic acid H₂Y per litre of solution.

You are required to find the percentage purity of the acid.

Procedure:

Results:

Pipette 20cm³ or 25cm³ of **FA1**, transfer it into a clean conical flask. Titrate it against **FA2** from the burette using phenolphthalein indicator.

Repeat the procedure until you obtain consistent results.

Record your results in the table below.

Volume of pipette used	cm ³
Final burette reading (cm ³)	
Initial burette reading (cm ³)	
Volume of FA2 used (cm ³)	
Values of FA2 used to calculate the average	ge volume
Average volume of FA2 used	
Questions: (a) Write equation for the reaction between	
(b) Calculate the number of moles of F	FA1 that reacted with FA2.

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(c) Calculate the number of moles of FA2 present in a litre of the solu	ution.
(d) Calculate the mass of the pure acid in the sample. Hence the perce $(Y = 96, H = 1)$	entage purity

You are provided with the following solutions.

BA1, which is a solution made by dissolving 8.05g of potassium hydroxide in a little of distilled water and solution made up to 500cm³.

BA2, which is a 0.25M solution of an acid H_nY , prepared by dissolving 15.75g of acid in 500cm^3 of solution.

You are required to determine the basicity n, of acid H_nY , and hence the value of Y (K = 39, O = 16, H = 1)

Procedure:

Pipette 25.0cm³ (or 20.0cm³) of **BA1** in a conical flask. Add 2-3 drops of phenolphthalein indicator.

Titrate with **BA2** from the burette until when you obtain consistent results.

Record your results in the table below.

V	olume of pipette used			cm ³
	Experiment	1	2	3
	Final burette reading (cm ³)			
	Initial burette reading (cm ³)			
	Volume of BA2 used (cm ³)			

Titı	re valı	ues used to calculate	average volum		
		volume of BA2		 	
Qu	estior			 	
Cal	culate	e the;			
	(i)	Molarity of BA1 .			

(ii)	Moles of BA1 that reacted.
(iii)	Moles of BA2 that reacted.
•••••	
(iv)	Value of n in H _n Y.
(v)	Value of Y in H _n Y
•••••	

You are provided with the following.

BA1, which is a solution containing 0.1 moles of an acid H_nX per litre of solution.

BA2, which is a solution prepared by dissolving 1.95g of sodium hydroxide in 500cm³ of distilled water.

You are required to determine the basicity of the acid (value of n in H_nX).

-					•				
ν	r	n	n	Δ	n	11	r	Δ	•
		.,	٠.						•

Results:

Pipette 25cm³ (or 20cm³) of **BA2** into a clean conical flask.

Add 2-3 drops of phenolphthalein indicator and titrate with **BA1** from the burette.

Record your results in the table below.

Repeat the titration until you obtain consistent results.

Volume of pipette used			cm ³	
Final burette reading (cm ³)				
Initial burette reading (cm ³)				
Volume of BA1 used (cm ³)				
			<u> </u>	
Values of used to calculate avera	ge volume of B A	A1 used.		
Average volume of BA1 used.				
Questions:				
(a) Calculate the;				
(i) Molarity if BA2 .	(Na = 23, O = 16)	5, H = 1)		

(ii)	Number of moles of BA2 that reacted.
• • • • • •	
(iii)	Moles of BA1 that reacted.
(iv)	Mole ratio of BA1 : BA2 ; hence determine the value of n in H_nX .
Write	the equation of reaction between BA1 and BA2.

You are provided with the following solutions.

 $\boldsymbol{BA1}$ is a solution made by dissolving 2.65g of a carbonate X_nCO_3 to make $250cm^3$ of solution.

BA2 is 0.2M hydrochloric acid solution.

You are required to determine the value of \mathbf{n} in the carbonate by titration.

•				-	1			
	'n	n	CE	\mathbf{n}	ш	r	Δ	•
		₹,	u		w			•

Pipette 25cm³ or 20cm³ of **BA1** into a clean conical flask. Add 2 drops of methyl orange indicator and titrate the mixture with **BA2** from a burette. Repeat the titration and record your results in the table below.

Volume of pipette used			c	m ³
Burette readings	1	2	3	
Final reading (cm ³)				
Initial reading (cm ³)				
Volume of BA2 used (cr	m ³)			
Average volume of BA2 use	ed			
Questions:				
(a) Calculate,	C 1 CD	40.1		
(i) The number	of moles of B	A2 that reacted.		

(ii)	The number of moles of X_nCO_3 in $1000cm^3$ of solution. $[X_nCO_3:HCl=1:2]$
 (iii)	Determine the;
(i)	Molar mass of the carbonate
•••••	
(ii)	Value of n in the carbonate X_nCO_3 (X = 27, C = 12, O = 16)

You are provided with the following;

BA1 which is a solution containing 2g of sodium hydroxide in 500cm³ of solution.

BA2 is a 0.06M solution of an unknown acid **Q**.

You are required to determine the mole ratio between the acid and sodium hydroxide.

-	_					•				
Н	ν	r	n	c	Δ	М	u	r	Δ	•
н	L		ι,	·		u	u		L	

Pipette 25cm³ (or 20cm³) of **BA1** into a conical flask. Add 2-3 drops of phenolphthalein indicator. Titrate it with **BA2** from the burette.

Repeat the titration until you obtain consistent results.

Record your results in the table below.

T					
Results: Volume of pipette used			,	2m ³	
volume of pipette used				J111	
Final burette reading (cn	n ³)				
Initial burette reading (cm ³)					
Volume of BA2 used (cr	m ³)				
Volumes used to determine	average volume	of BA2			
Average volume of BA2 use	ed.				
Questions:					
(a) Calculate the number	er of moles of:				
(i) Sodium hydr	roxide used. (Na	= 23, O = 16, H	= 1)		
()					
					•••

(ii) Acid used.				
(b) Determine the mole ra	tio between acid			
•••••				
BA1, is a solution made by dissolution. BA2, is 0.05M hydrochloric at You are required to determine Procedure: Pipette 25cm³ or 20cm³ of BA indicator and titrate with BA2 table below.	cid solution. the number of m 1 into a clean tith from the burette	noles of water of ration flask, add a	crystallization, r 2 drops of methy tion and record y	n in the carbonate. yl orange your results in the
Volume of pipette used			cn	13
Burette readings	1	2	3	
Final reading (cm ³)				
Initial reading (cm ³)				
Volume of BA2 used (cm ³)				
Titre values of BA2 used for c	calculating averag	ge volume		

Avera	ge vol	ume of BA2 used.
Quest	ions:	
_	Calcı	ılate
(4)	(i)	The number of moles of hydrochloric acid that reacted.
	(1)	The named of moles of hydroemoric dela that redeted.
	•••••	
	•••••	
	•••••	
	•••••	
	(ii)	The number of moles of the carbonate that reacted.
	(11)	
	•••••	
	• • • • • •	
	• • • • • •	
	• • • • • •	
(b)) Detei	rmine the value of $\bf n$ in the carbonate $X_2CO_3.nH_2O$. [X = 23, C = 12, H = 1]
	• • • • • •	

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Practical 32

You are provided with the following solutions.

BA1, which is a solution made by dissolving 2.65g of a salt M₂X in 250cm³ of solution.

BA2, is 0.25M hydrochloric acid.

You are required to determine the formula mass of X in the salt by titration.

n.		
Pr	ጉራልብ	ma.
111	vecu	lure:

Pipette 25cm³ or 20cm³ of **BA1** into a conical flask, add 2 drops of methyl orange indicator and titrate with **BA2** from the burette. Repeat the titration and record your results in the table below.

Volume of pipette used			cm ³	
Burette readings	1	2	3	
Final burette reading (cm ³)	1			
Initial burette reading (cm ³				
Volume of BA2 used (cm ³))			
Titre values of BA2 used for continuous and the continuous section and th				
Questions:				
(a) Calculate the number of	of moles of B .	A2 that reacted.		

(b)	Determine the molarity of the salt $(M_2X : HCl = 1 : 2)$
(c)	Determine the value of X ($M = 23$).

You are provided with the following;

BA1, which is a 0.067M sodium hydroxide solution

BA2, which is a solution containing 4.2g of an organic acid **Q** in 500cm³ of solution.

You are required to determine the mole ratio of reaction between sodium hydroxide and Q.

Procedure:

Pipette 20/25cm³ of **BA1** into a clean conical flask then add 2-3 drops of phenolphthalein indicator.

Titrate **BA1** with **BA2** from the burette until the solution just turns colourless.

Repeat the above procdure until you obtain consistent results. Record your results in the table blelow.

V	olume of pipette used			cm ³
			Γ	Γ
	Experiment	1	2	3
	Final burette reading (cm ³)			
	Initial burette reading (cm ³)			
	Volume of BA2 used (cm ³)			

Γitre values used to calculate the average volume of BA2 used							
Average volume of BA2 used							
Questions:							
(a) Calcu	late the:						
(i)	Concentration of BA2 in mol/dm ³						

(ii)	Number of moles of acid in the average volume of BA2 that reacted
(iii)	Number of moles of BA1 that reacted with BA2 .
) Deter	rmine the mole ratio of reaction between sodium hydroxide and acid Q

Chapter three: THERMO-CHEMISTRY

Practical experiments on thermochemistry are mainly on heat of neutralization. When an acid is added to a base, heat energy is produced. This detected by a rise in temperature which is measured by a thermometer.

When reactions take place, bonds are either formed or broken. As a result there is a change in temperature, chemical reactions result into energy changes. The heat change that occur s during a chemical reaction referred to as a enthalpy, ΔH

There are two types of reactions classified depending on heat change accompanying the reaction. These are endothermic reactions and exothermic reactions

Endothermic reactions

These are reactions which occur with absorption of heat from the surrounding. In these reactions, energy must be supplied for the reactants to be converted to products. The energy (temperature) of the products falls below that of initial energy (temperature). The enthalpy change is positive implying that energy (heat) is absorbed from the surrounding e.g. when sodium thiosulphate dissolves in water.

$$Na_2S_2O_{3(s)} + (aq)$$
 $\longrightarrow Na_2S_2O_{3(aq)}$, $\Delta H = +ve$

Exothermic reactions

These are reactions which occur with evolution/liberation of heat to the surrounding. The temperature (heat) rises above the initial temperature (heat). The enthalpy change accompanying the reaction is negative. This implies that heat is given out to the surrounding e.g.

$$C_{(s)} + O_{2(g)} \longrightarrow CO_{2(g)} \Delta H = -ve, \Delta H = -393 \text{Kjmol-}$$

-393Kjmol- implies that 393Kj of energy is liberated when 1 mole of carbon is completely burnt in oxygen to form carbon dioxide

Examples of heat changes

***** Enthalpy of neutralization

This is the quantity of heat evolved when one mole of hydrogen ions completely reacts with one mole of hydroxyl ions to form one mole of water

$$H^{+}_{(aq)} + OH^{-}_{(aq)} \longrightarrow H_{2}O_{(1)} \Delta H = -57.3 \text{Kjmol}^{-1}$$

Neutralization reaction is exothermic meaning that energy is given out. When a strong acid is neutralized with a strong alkali, the heat change is approximately 57.3Kjmol⁻¹. However when the alkali or acid is weak, the enthalpy change is less than 57.3Kjmol because heat is absorbed to dissociate the weak alkali or acid.

***** Enthalpy of solution

Heat of solution is the quantity of heat liberated or absorbed when one mole of a substance is completely dissolved in water to form an infinitely dilute solution. The quantity of heat change depends on the amount of solvent used.

$$NaOH_{(s)} + (aq) \longrightarrow Na^+ (aq) + OH^- (aq)$$

\(\) Heat of precipitation/displacement.

This is the quantity of heat liberated when one mole of a given substance is completely displaced by another substance e.g. when iron filings are added to a solution of copper (II) sulphate, a blue solution of CuSO4 turns to pale green and a brown solid is deposited at the bottom. The pale green solution is due to formation of iron (II) sulphate whereas the brown solid is due to formation of copper metal.

$$\begin{array}{ccc} CuSO4_{(aq)} + Fe_{(s)} & \longrightarrow & FeSO4_{(aq)} + Cu_{(s)} \\ Ionically, & & & \\ Cu^{2^+}{}_{(aq)} + Fe_{(s)} & \longrightarrow & Fe^{2^+}{}_{(aq)} + Cu_{(s)} \end{array}$$

Note. Iron is able to displace copper from its ions because iron is higher than copper in the electrochemical series.

A .	4 •	• 4	1
Ac	u	vity	7 J

Activi	ty 1
a)	When 25cm ³ of 0.2M sulphuric acid was added to 100cm ³ of 1M sodium hydroxide solution, the temperature of the resultant solution increased by 12.6 °C. Write the ionic equation for the reaction that took place.
b)	Calculate the enthalpy of neutralization sodium hydroxide (S.H.C = 4.2 Kjmol ^{-1 0} C ⁻¹ , density of solution = 1 gcm ⁻³)
Activi	ty 2
	50cm ³ of 2M hydrochloric acid and 50cm ³ of 2M sodium hydroxide both at 22 °C were mixed in a plastic beaker. The mixture was stirred and maximum temperature obtained as 35 °C. (S.H.C = 4.2Kjmol ⁻¹ °C ⁻¹ , density of solution = 1gcm ⁻³)
	Explain why a plastic beaker was used for mixing

(b) Calculate the heat of the reaction
(c) Explain precisely what would happen to the value of heat of reaction calculated above it 50ml of 2M ammonia solution was used instead of sodium hydroxide.
PRACTICAL SCHEDULES Practical 1
You are provided with the following:
 BA5, which is 2M sodium hydroxide solution.
 BA6, which is hydrochloric acid solution.
Plastic cup.
• Thermometer.
Plastic beaker.
You are required to determine the enthalpy of neutralization of hydrochloric acid by sodium
hydroxide
Procedure:
(a) Measure 50cm ³ of BA5 into the plastic cup
(b) Record the temperature, t ₁ of the solution
(c) Wash the thermometer with distilled water
(d) Measure 50cm ³ of BA6 into the plastic beaker
(e) Measure and record the temperature, t ₂
(f) Now add all the 50cm ³ of BA6 into the alkali in the plastic cup
(g) Stir the mixture carefully with the thermometer and record the highest temperature t ₃
Results:
Volume of BA5 used
Volume of BA6 used
Total volume of the mixture.
Temperature of BA5 , t ₁
Temperature of BA6 , t ₂
Average initial temperature.

	emperature, t ₃ nine the temperature change
Quest	ions:
_	Write an equation for the reaction between BA5 and BA6
(b)	Calculate the number of moles of sodium hydroxide that reacted.
(0)	
	10.1
(c)	Calculate the heat evolved (S.H.C = 4.2 Kjmol ⁻¹ 0 c ⁻¹ , density of solution = 1 gcm ⁻³)
(4)	Coloulate the moler heat of neutralization of hydrochleric said by sodium hydrovide
(u)	Calculate the molar heat of neutralization of hydrochloric acid by sodium hydroxide.
(e)	If 2M ethanoic acid was used instead of 2M HCl, what would be the heat change as
(0)	compared to the value calculated above? Explain your answer.

You are provided with the following,

- **BA1**, which is 2M NaOH solution
- **BA2**, which is 2M HCl solution
- Plastic cup
- Burette
- Thermometer

You are required to determine the enthalpy of neutralization of NaOH by HCl

D.,	oce	J.,		_
Pr	м.н	411	re	•

- a) Pipette 25cm³ **BA1** into a clean plastic cup. Note and record the temperature, t₁
- b) Note and record the temperature t_2 of solution **BA2** and put in the burette
- c) Add 5cm³ of **BA2** from the burette to **BA1**, stir and record the maximum temperature of the mixture, t₃
- d) Repeat procedure (c) at 5cm³ interval until the total of 50m³ of **BA2** has been added
- e) Record the results in the table below

esults: itial temperature of BA1, t ₁ itial temperature of BA2, t ₂										
verage temperature of the mixtu	re									•••••
Volume of BA2 used (cm ³)	5	10	15	20	25	30	35	40	45	50
Maximum temperature,t ₃ (⁰ C)										
Temperature change $t_3 - \underline{t_1 + t_2} (^0c)$										
(a) Plot a graph of temperature (b) Use your graph to determine graph								ze BA	1 fron	n the

	(c)	Calculate the molarity of BA2								
										• • • • • • • • • • • • • • • • • • • •
		Calculate the enthalpy of neutralization of solution = 1 gcm ⁻³)	ation o	f BA1	by B	A2 (S	.H.C :	= 4.2H	Kjmol	^{-1 0} c ⁻¹ , density
										• • • • • • • • • • • • • • • • • • • •
Pr	actic	al 3								
		e provided with the following:								
		which is a solution containing 54.5g	of a n	nixture	of po	tassiu	ım hy	droxid	de and	l potassium
	-	e per litre of solution.								
		which is a 0.5M sulphuric acid. The required to determine the percental	age of	notace	ium cı	ılnhat	a in T	1		
	oced	-	age or	potass	iuiii Si	прпа	.c III L	<i>)</i> A1.		
		Measure and record the temperatur	re of D	A1 .						
		Pipette 25.0cm^3 (or 20.0cm^3) of D								
		Measure 10cm ³ of DA2 using a me								-
		beaker containing DA1 . Gently sting temperature attained by the mixture.		a thern	nomet	er and	l reco	rd the	high	est
		Repeat procedures (b) and (c) using		0 25	30 an	d 35c	m ³ of	DA2.		
		Enter your results in the table below	-	.0, 25,	oo un	u 33 c .	01	D.12.		
	` ′	•								
	sults								2	
VC	lume	e of pipette used						cı	n ³	
	Volu	ume of DA2 used (cm ³)	0	10	15	20	25	30	35	
-	Tem	perature of solution mixture (⁰ C)								

Questions:

DA	at a graph of temperature of solution mixture (along the vertical axis) against volume of 2 used (along the horizontal axis)
	termine from your graph, the maximum volume of DA2 required to react with DA1 .
••••	
 Ca	culate the;
	i) Maximum number of moles of sulphuric acid that reacted.
	······································
	ii) Number of moles of potassium hydroxide in 20.0cm ³ (or 25.0cm ³) of DA1
	Concentration of DA1 in grams of potassium hydroxide per litre of solution. (H = $1, O = 16, K = 39$)

	iv) Percentage of potassium sulphate in DA1 .
Practi	ical 4
You a	re provided with the following:
BA1 ,	which is a solution containing 1M sodium hydroxide solution
BA2,	which is a solution containing 1M sulphuric acid
You a	re required to determine the enthalpy of neutralization of sulphuric acid.
Proce	dure:
	Using a measuring cylinder, measure 50cm ³ of BA1 into a clean plastic beaker. Note and
	record its initial temperature $T_1{}^0$ C. Wash the thermometer and dry it.
2.	Using another measuring cylinder, measure 25cm^3 of BA2 . Note and record its initial temperature $T_1{}^0$ C.
3.	•
Recul	ts of the experiment
	temperature of BA1
	temperature of BA2
	temperature of mixture
Quest	
	Write the equation for the reaction between BA1 and BA2
(4)	, write the equation for the reaction setween Bill that Bill
(b)	Calculate the quantity of heat produced during the reaction (S.H.C = $4.2 \text{Kjmol}^{-10} \text{c}^{-1}$, density of solution = 1gcm^{-3})

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•		
(c) C	Calculate the molar heat of neutralization of the acid by sodium hy	droxide
•		
(d) E	Explain why a plastic beaker is used instead of glass or metallic cu	p.
•		
(e) E	Explain why sodium hydroxide is used instead of ammonium hydr	oxide soluti
•		
(f) It	f ammonium hydroxide is used, would you expect the heat of neu	ralization to
	greater, equal or less. Give a reason for your answer.	irunzunon t
	·	
•		• • • • • • • • • • • • • • • • • • • •

You are provided with the following:

0.5M copper (II) sulphate

Zinc powder

 $(S.H.C = 4.2 \text{Kjmol}^{-10} \text{c}^{-1}, \text{ density of solution} = 1 \text{gcm}^3)$

You are required to determine the enthalpy of displacement of copper (II) sulphate by zinc

Procedure:

a) Using a measure cylinder, transfer 50cm^3 of 0.5 M solution of copper (II) sulphate into a plastic cup. Note and record the temperature, t_1 of the solution

b)	Add 2g of zinc powder and stir well but carefully with the thermometer. Note and record
	the highest temperature,t ₂ attained by the mixture
Result	is:
Initial	temperature of solution, t ₁
Final t	emperature of the mixture,t ₂
Tempe	erature change, $t_2 - t_1$
Questi	ions:
a)	State what was observed
b)	Write the equation for the reaction that took place
a)	Coloulate the number of moles of conner (II) sulphote solution
()	Calculate the number of moles of copper (II) sulphate solution
d)	Calculate the number of moles of zinc in $2g$ ($Zn = 65$)

e)	Calculate the number of moles of zinc that reacted
f)	Calculate the heat change that occurred in the reaction. S.H.C = 4.2 Kjmol ⁻¹⁰ c ⁻¹ , density of solution = 1 gcm ⁻³)
g)	Calculate the enthalpy change in Kjmol ⁻¹
h)	If magnesium instead of zinc was used, would the enthalpy change for the reaction be greater, less or equal to that you have calculated in (g) above. Explain your answer.

You are provided with the following:

- Plastic beaker
- Measuring cylinder
- Distilled water
- Thermometer
- 8g of **X**

You are required to determine the enthalpy of solution salt **X**

Procedure:

- a) Using the measuring cylinder, measure 90cm³ of distilled water into a plastic beaker
- b) Record the temperature of the water in the beaker for every minute for 4 minutes
- c) At the fourth minute, add salt **X** and stir continuously to dissolve. Record the temperature every half a minute for 3 minutes.

Record your results in the table below.

Results:

Time	1.0	2.0	3.0	4.0	4.5	5.0	5.5	6.0	6.5	7.0	7.5
(min)											
Temp											
(^{0}C)											

Questions:

١.	D1 .	1	•		• .	. •
0	\ Plota	aranh	Δ t	tamparatura	againet	tima
а		$\gamma_1 a_1 n_1$	()1	temperature	against	\mathbf{u}

D)	Extrapolate	tne graph to	find the change in temperature.	
----	-------------	--------------	---------------------------------	--

Calculate the heat of solution from the quantity of water used. (S.H.C = 4.2 KJmol 10 C 1 , density of solution = 1 gcm $^{-1}$)

d)	State whether the reaction is exothermic or endothermic, give a reason for your answer

You are provided with the following Solution **J** which is a basic solution Solution **D** which is an acidic solution You are required to determine:

- i) the molarity of the acid
 - ii) The molar heat of neutralization of the acid and the base

Procedure:

- (a) Using a measuring cylinder, measure 150cm³ of **J** and transfer into a 250cm³ beaker. Add 50cm³ of water, mix and label this **BA1**.
- (b) Transfer 100cm³ of **D** into another 250cm³ beaker using a measuring cylinder. Add 100cm³ of water. Mix and label this **BA2**.
- (c) Measure and record the initial temperature of **BA1**
- (d) Run 25cm³ of **BA1** from the burette into a dry plastic beaker.
- (e) Using a measuring cylinder transfer at once 10cm³ of **BA2** into a plastic beaker containing **BA1**. Stir with the thermometer and record the highest temperature attained by the mixture.
- (f) Repeat the procedures (d) to (e) using 20, 30, 40 and 50cm³ of **BA2**
- (g) Record your results in the table below.

Initial temperature of **BA1**.....

Volume of BA2 used (cm ³)	10	20	30	40	50
Highest temperature of the mixture (⁰ c)					

Questions:

	Plot a graph of highest temperature attained against the volume of BA2 . From the graph determine; (i) The volume of BA2 required to neutralize 25.0cm ³ of BA1
	(i)The maximum temperature change for the reaction
	Calculate the molarity of $\bf BA2$ (1 mole of the base reacts with 1 mole of the acid, Molarity of $\bf BA1 = 1.5M$
(d) L	Determine the;
	(i) Maximum heat evolved during the reaction (S.H.C of solution = $4.2jg^{-10}c^{-1}$,
	density of mixture = 1gcm^{-3})

(ii) Molar heat of reaction be	atryaan	tha aa	id and t	ha haga			
(ii) Word heat of feaction of							
Practical 8							
You are provided with the following:							
M which is an acidic solution							
N which is an alkaline solution							
You are required to determine the mola	rity of	an acid	lic solut	ion M			
 (a) Using a measuring cylinder, me add 90cm³ of distilled water and (b) Transfer 190cm³ of N into anoth 10cm³ of distilled water and mix (c) Transfer BA1 into a burette. Ru (d) Add at once 10cm³ of BA2 from 25cm³ of BA1. Gently stir with attained by the mixture. (e) Repeat the procedures (c) to (d) (f) Record your results in the table 	d mix. I her 250 x. Labe n 25cm n the m the the	Label to cm ³ be a l this shaded the casuring the casuri	his solu eaker us olution A1 into ng cylin eter and	tion B A ing a m BA2 a plasti der into	easuring c beake a plast the high	g cylinde r. ic beaker	er. Add containin
Volume of BA2 used (cm ³)	10	15	20	25	30	35	40
Temperature of the mixture (°C)							
Questions: (a) Plot a graph of temperature the second (b) From the graph, determine the next temperature in the second (c) and the second (c) are second (c) and the second (c) are secon		_				o raget w	ith 25am³

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(c) **BA1**

(d)	Calculate the number of moles of alkali that reacted (molarity of $\mathbf{BA2} = 1.9\mathbf{M}$)
(e)	Calculate the number of moles of the acid that reacted (ratio of acid to alkali = $1:2$)
(f)	Calculate the molarity of the acid

You are provided with the following

BA1 which is a solution containing 50g/l of a mixture of sodium hydroxide and sodium sulphate. Solution **D** which is 1M sulphuric acid

You are required to determine the percentage of sodium sulphate in the mixture

Procedure:

- (a) Using a measuring cylinder, measure 70cm³ of **D** and transfer into a 250cm³ beaker. Add 70cm³ of water, mix and label this **BA2**.
- (b) Fill the burette with **BA2**.

Volume of **BA1** used (cm³)

- (c) Pipette 25.0cm³ (or 20.0cm³) of **BA1** into a plastic beaker. Note and record its temperature, t₁.
- (d) Run 5cm³ of **BA2** from the burette into a plastic beaker containing **BA1**. Stir the mixture with a thermometer and record the highest temperature, t₂ attained.
- (e) Determine the change in temperature, T for the reaction.
- (f) Repeat the procedure (c) to (e) using volume 10, 15, 20, 25, 30 and 35cm³ of **BA2**.
- (g) Record your results in the table below.

Volume of BA2 used (cm ³)				
Temperature of BA1 , t ₁ (⁰ C)				
Temperature of the mixture, t ₂ (⁰ C)				
Change in temperature, T (⁰ C)				

Ouestions:

CSU	ions.
(a)	Plot a graph of change in temperature, T against volume of BA2
(b)	From the graph state the highest change in temperature
(c)	Write the equation for the reaction that took place

(d) Determ	mine the volume of BA1 and BA2 that gives the highest temperature
	ate the number of moles of:
(i)	BA2 that reacted.
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···	\ D 4.1 d 4 3
(11)	BA1 that reacted.
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•••	
(f) Calcul	Molarity of BA1
•••	
•••	
•••	
•••	
•••	

(:	ii) Concentration of BA1 in g/l
•	
•	
(iii)Percentage of sodium sulphate in the mixture. (Na = 23, O = 16, S = 32, H = 1)
•	
	rmine the enthalpy of neutralization during the reaction. (S.H.C of solution = $4.2j$ density = $1gcm^{-3}$)
	density = 1gcm ⁻³)
10c-1,	density = 1gcm ⁻³)
10c-1,	density = 1gcm ⁻³)
10c-1,	density = 1gcm ⁻³)
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10c-1,	density = 1gcm ⁻³)
10c-1,	density = 1gcm ⁻³)
10c-1,	density = 1gcm ⁻³)

Practical 10.

You are provided with the following:

BA1, which is 1.8M H_nX solution.

BA2, which is 2M sodium hydroxide solution.

You are required to determine the value of n in H_nX

Procedure:

- (a) Using a measuring cylinder, measure exactly 40cm³ of **BA2** and place it into a plastic cup or beaker
- (b) Using a burette, transfer 10cm³ of **BA1** into another plastic cup or beaker.
- (c) Place a thermometer in the cup or beaker containing **BA1** and transfer all the **BA2** in the beaker containing **BA1** while stirring carefully using the thermometer and record the maximum temperature reached.
- (d) Repeat procedure (a) to (c) for volume of **BA2** 35, 30, 25, 20, 15, 10, and 5cm³ with the corresponding volumes of **BA1** given in the table. Enter your results in the table below.

Results:

volume of sodium hydroxide /cm ³	40	35	30	25	20	15	10	5
Volume of H _n X /cm ³	10	15	20	25	30	35	40	45
% of H _n X	20	30	40	50	60	70	80	90s
Temperature /°C								

(a) P	lot a graph of temperature against percentage of H _n X.
(b) C	Calculate the volume of H _n X and sodium hydroxide that reacted to give the maximum
	emperature rise.

(c)	Deduce the value of $\bf n$ and $\bf X$ (BA1 contains 70g of $\bf H_n \bf X$ per litre).

Practical 11 (WAKISSHA PAPER 3 2013)

You are provided with the following;

BA1, which is a solution containing 3.15g of a hydrated dibasic acid H₂X.nH₂O, in 250cm³ of the solution.

BA2, which is a 0.2M sodium hydroxide solution.

You are required to determine the percentage of water of crystallization, **n**, in the hydrated acid. (1 mole of acid reacts with 2 moles of sodium hydroxide)

Procedure:

- (i) Pipette 25.0cm³ (or 20.0cm³) of **BA2** into a plastic beaker. Measure and record the initial temperature.
- (ii) Fill the burette with **BA1**, then run 10cm³ of **BA1** from the burette into the beaker containing **BA2**. Gently stir the solution using the thermometer, and record the maximum temperature attained by the mixture.
- (iii)Repeat procedure (ii) above until 30cm³ of **BA1** has been added.
- (iv)Record your results in the table below.

Results:

Volume of BA1 added (cm ³)	0	10	15	20	25	30
Maximum temperature attained (⁰ C)						
Temperature rise (⁰ C)						

Questions:

 (a) i) Plot a graph of temperature rise (along vertical axis) against volume of BA1 added (along horizontal axis). ii) From your graph, determine the volume of BA1 required for complete neutralization of BA2. 	ion
(b) Calculate the, (i) Number of moles of BA2 that reacted.	

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	(ii) Concentration of the hydrated acid in BA1 in moles per dm ³
	(iii)Molar mass of the hydrated acid.
(c) i) Determine the value of n in the hydrated acid (H = 1, O = 16, X = 88)
(0) 1,	
(ii)Calculate the percentage of water of crystallization in the acid H ₂ X.nH ₂ O.
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Practical 12 (WAKISSHA PAPER 3 2015)

You are provided with the following,

Metal **Q** measuring about 7cm

Z which is a 0.2M solution of dilute acid, HX

You are required to determine the mass of metal ${\bf Q}$

Procedure

- (i) Measure and cut exactly 6cm of metal **Q** provided.
- (ii) Using a measuring cylinder, measure 40cm³ of solution **Z** and transfer it into a clean plastic beaker, note and record the initial temperature of this solution in the table provided below.
- (iii)Place metal \mathbf{Q} into the solution \mathbf{Z} in the plastic beaker and simultaneously start the clock.
- (iv)Note and record the temperature of the contents in the beaker after every 30 seconds in the table below.

Results table

Time (seconds)	0	30	60	90	120	150	180	210
Temperature (⁰ C)								

Questions:	
(a) Plot a	graph of temperature against time.
(b) Determ	mine the highest temperature rise.
•••••	
* *	ate the amount of heat produced during the reaction. (Assume: Specific heat ty of solution = $4.2J/g/^{0}$ C, density of solution = $1g/\text{cm}^{3}$)
• • • • • • • • • • • • • • • • • • • •	
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(d) Given that th Calculate the	e molar heat of reaction between metal ${\bf Q}$ and acid ${\bf Z}$ is 1600KJ/mol,
	per of moles of Q that reacted
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•••••	
(ii) Mass	of \mathbf{Q} used in this experiment ($\mathbf{Q} = 24$)
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Chapter four: PRACTICAL EXPRIMENTS ON RATES OF REACTIONSRate of reaction

some reactions take place almost instantaneously like explosion of hydrogen in air mixture when lighted splint is inserted in it, a concept described as burning with pop sound. Other reactions are slow e.g. rusting of iron in which a noticeable change is observed after appreciable time. Some reactions show changes in colours of reactants or of products with time. For example nitrogen oxide reacts with oxygen to form nitrogen dioxide which is brown. The gradual change in colour can be measured with time, and this can indicate rate of reaction.

The change in concentration with time can be followed by titration, you will also deal with reactions in which colourless gases are evolved.

During a chemical reaction, the concentration of products increases whereas that of reactants decreases. The change in concentration with time is what is termed as rate of reaction. Therefore rate of reaction is the measure of the change in quantity of reactants or products with time.

Rate of reaction is precisely defined as speed/rate at which reactants are turned into products

In determining the rate of a chemical reaction, the amount of product formed in a given period of time or the amount of reactants consumed in a given period of time is measured.

Then the rate is expressed as amount of products formed per unit time or the amount of reactants consumed per unit time.

Rate = amount of products formed

Time

Rate = amount of reactants consumed

Time

Activities for students

Activity 1

10cm length of magnesium ribbon was cleaned with sand paper. 50cm3 of 0.1M dilute hydrochloric acid was measured into a clean conical flask and corked. A rubber cork was then removed and 10cm magnesium ribbon added, the stop watch was started and flask corked immediately. The set up was organized as shown below and gas evolved measured in the syringe in intervals of 5 seconds. The date was obtained and tabulated, use the values to answer questions.

Time (s)	Volume of gas evolved
	(cm ³)
0	0
5	20
10	40
15	60
20	75
25	85
30	85
35	85
40	85
45	85

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	Plot a graph of volume of gas evolved against time Use your graph to find the rate reaction at 13 seconds
(a)	What volume of the is evolved at 16 seconds
(0)	What volume of the is evolved at 16 seconds
(d)	Describe the graph you have drawn in (a) above
(e)	Explain precisely the shape of the graph
(f)	Write a balanced equation for the reaction

Activity 2

50cm3 of hydrochloric acid was measured into the conical flask. The flask and its contents were placed on a direct weighing balance. Drop of magnesium ribbon was added to acid in flask and the timer started. Mass of the flask and its contents was taken in intervals of 5 seconds until all the magnesium ribbon was reacted.

The values were tabulated; use the table below to answer questions.

Time (s)	Mass of flask + its contents
0	65.50
5	65.45
10	65.35
15	65.30
20	65.27
25	65.25
30	65.25
35	65.25
40	65.25

(a)	Plot a graph of volume of gas evolved against time
(b)	Use your graph to find the rate of reaction at 17 seconds
(c)	What is the mass of flask and contents at 13 seconds
(-)	
<i>(</i> 1)	D 7 4 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
(a)	Describe the graph you have drawn in (a) above

(e)	Explain the shape of the graph
(f)	Write a balanced equation for the reaction

FACTORS THAT AFFECT THE RATE OF REACTION

The major factors that affect the rate of reaction include temperature, concentration, surface area, pressure, light and catalyst. We shall pick temperature, surface area and concentration for practical purposes, however others can be taught in theory.

Effect of temperature on the rate of reaction.

The rate of reaction increases with increase in temperature. Increase in temperature increases the molecular velocities of which subsequently increase the kinetic energy of reacting particles. This increases frequency of interaction/collision of the reacting molecules to form products.

Note.

For reaction to take place, particles must collide with sufficient kinetic energy. When the temperature is increased, particles gain energy and interact with sufficient average minimum energies. This implies that activation energy of the reaction increases and so is the rate of reaction. Activation energy is the energy barrier which must be overcome by the reacting species before the reaction can proceed.

Practical A

- You are provide with the following,
- **BA1**, which is 0.16M sodium thiosulphate solution
- **BA2**, which is 2M hydrochloric acid solution
- 2 conical flasks
- 2 measuring cylinders
- Stop clock
- A sheet of white paper

You are required to determine the rate of reaction when a thiosulphate reacts with dilute hydrochloric acid.

Procedure

- (i) Using a clean measuring cylinder, transfer 50cm³ of **BA1** into a clean conical flask. Measure and record its temperature (room temperature)
- (ii) Using a black or blue pen, mark a cross on the sheet of white paper.
- (iii)Add 5cm³ of **BA2** into a flask and at the same time start the stop clock, swirl the mixture and place on the sheet of white paper with the cross in the middle
- (iv)Look down through the mixture at the cross and stop the clock when the cross just disappears. Record the time taken for the cross to just disappear in the middle.
- (v) Measure another 50cm³ of **BA1** into a clean conical flask, this time heat the **BA1** to a temperature of 30 °C. Add 5cm³ of **BA2** and immediately start the stop clock, swirl the mixture and place it on a sheet of white paper with the cross in the middle. Note and record the time t, it takes for the cross to disappear.
- (vi)Repeat the procedure (iii) to (v) but this time heat **BA1** to 40 0 C, 50 0 C, 60 0 C respectively before adding **BA2**. Record your results in the table below.

Temperature (⁰ C)	Time t, (s)	$^{1}/_{t}$ (s ⁻¹)
Room temperature		
30		
40		
50		
60		

Questions:

(a) Plot a graph of
(i) Time against temperature
(ii) $\frac{1}{t}$ against temperature
(b) From your graph in (a) (i) determine the rate of reaction at 20 seconds

(c)	Explain the shapes of the graphs
(d)	What is your conclusion from the two graphs
(e)	State and explain how the rate of reaction varies with temperature in (a) (ii) above.
(f)	Write a balanced ionic equation for the reaction that occurred between BA1 and BA2.

Effect of concentration on the rate of reaction

The rate of reaction increases with the increase in concentration of the reactant species. When concentration of reactants increase in a given volume of the reaction vessel, the chances of collision of reacting particles consequently increase resulting into increased reaction rate. The greater the concentration of reactants, the greater the number of interactions between reacting particles, therefore more effective collisions occur with minimum activation energy hence faster rate of reaction.

However the reverse is true for concentration of products.

Activity 3

10cm length of magnesium ribbon was cleaned with sand paper. 50cm^3 of 0.1 M dilute hydrochloric acid was measured into a clean conical flask and corked. A rubber cork was then removed and 10cm magnesium ribbon added, the stop watch was started and flask corked immediately. The set up was organised as shown below and gas evolved measured in the syringe in intervals of 5 seconds. The above procedure was repeated using 0.2 M hydrochloric acid. The date was obtained and tabulated, use the values to answer questions.

Time (s)	Volume of gas evolved	Volume of gas
	using 0.1M HCl	evolved
		using 0.3M HCl
0	0.0	0.0
120	6.5	10.0
240	13.0	20.0
360	20.0	25.5
480	24.5	29.5
600	27.0	32.0
720	29.5	32.0
840	32.0	32.0

Questions:

(a)	Plot a graph of the same	axes the volume	of gas liberated	d using 0.1M and	l 0.3M HCl acid
	against time				

(b)	Explain the shapes of the graphs	
		•
		•
		•

(c)	Find the rate of reaction for both curves at 410 seconds.	
		•

Reaction of sodium thiosulphate and dilute hydrochloric acid

Sodium thiosulphate reacts with dilute hydrochloric acid to form a white/yellow precipitate of sulphur according to the following equation.

$$Na_2S_2O_3$$
 $(aq) + 2HCl(aq) \longrightarrow NaCl(aq) + S(s) + SO_2(g) + H_2O(l)$

The ionic equation is,

$$S_2O_3^{2-}$$
 (aq) + 2H⁺ (aq) \longrightarrow S(s) + SO₂ (g) + H₂O(l)

The effect of changing concentration of thiosulphate on the rate of reaction by carrying out separate experiments in which the concentration of hydrogen ions (acid) is maintained constant and concentration of thiosulphate is varied. In separate experiments, the volume of the solution is kept constant. The beaker of the same size is used to mix the solutions and the beaker is placed on top of a sheet of paper marked with a black cross. Looking from above, the black cross disappears after precipitation of a specific amount of sulphur. In separate experiments the extent of reaction when the cross disappears will be the same but the time for the cross to disappear will inversely depend on the concentration of thiosulphate ions. Consequently the reciprocal of the time for the cross to disappears in each experiment measures the average rate of formation of the precipitate quantity per second.

A graph of concentration of sodium thiosulphate against time begins with a steep slope which gradually becomes less steep before finally flattening out towards the horizontal axis. The gradient of the curve at any point measures the rate of reaction at the time represented by the point on the graph. The shape of the graph shows that the rate of reaction decreases as the concentration of sodium thiosulphate decreases.

A graph of sodium thiosulphate against $^{1}/_{t}$ is a straight line within the limits of experimental errors. The reciprocal of the time is the measure of rate of reaction for the appropriate concentration. At a fixed temperature, the rate of reaction is directly proportional to the concentration of sodium thiosulphate solution provided that enough acid is used to react with the thiosulphate.

The effect of temperature on the rate of reaction between sodium thiosulphate and dilute hydrochloric acid can be performed by heating the solution to certain temperature before adding the acid. A graph of $^1\!/_t$ against temperature can be plotted. The results from such an experiment will show that the higher the temperature, the greater the rate of reaction. A fairly small rise in temperature has a large influence on the rate of reaction.

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Practical B

You are provided with following:

A piece of magnesium ribbon labeled T

FA1, which is dilute sulphuric acid

A stop clock

You are required to determine the rate of reaction between magnesium and sulphuric acid at different concentrations.

Procedure:

- (i) Cut T into 5 equal pieces of 2cm each
- (ii) Using a measuring cylinder, measure 50cm³ of **FA1** into a clean glass beaker
- (iii)Drop one piece of **T** in the beaker and simultaneously start the stop clock. Swirl the solution continuously ensuring that **T** is always inside the solution.
- (iv) Note and record the time t, in seconds taken for **T** to completely dissolve.
- (v) Wash the beaker and repeat the procedures (ii) to (iv) using 40cm³, 30cm³, 20cm³ and 10cm³ of **FA1** but in each case make the total volume of the solution to 5cm³ by adding water from the burette.

(vi)Record your results in the table below including the of $\frac{1}{t}$ for each reaction

Volume of FA1 (cm ³)	50	40	30	20	10
Volume of water added (cm ³)	0	10	20	30	40
Time, t (s)					
1/ _t (s ⁻¹)					

Questions:

(a)	Plot a graph of volume of FA1 against $\frac{1}{t}$
(b)	Write and ionic equation for the reaction
(c)	What does $\frac{1}{t}$ represent
, ,	

d) Fron	the graph;
(i) Determine the rate of reaction $\frac{1}{t}$ when the volume of FA1 is 25.5cm ³
(:	ii) Determine the volume of FA1 , the rate of reaction is 4.25×10^{-2} s ⁻¹
(1	iii)What is the effect of concentration on the rate of the reaction?

Practical C

You are provided with the following,

- **BA1** which is a solution containing 0.2M thiosulphate ions
- **BA2** which is a solution containing 2M hydrochloric acid
- Distilled water
- 2 conical flasks
- 2 measuring cylinders
- Stop clock
- Sheet of white paper

You are required to determine the rate of reaction between thiosulphate and hydrochloric acid

Procedure

- a) Make a cross with a blue or black pen on the sheet of white paper provided
- b) Using a measuring cylinder, measure 60ml of **BA1** into a conical flask
- c) Add 10cm³ of **BA2** into the **BA1** in the conical flask and immediately start the stop clock
- d) Swirl the mixture to mix the two solutions and place it on the cross
- e) Look through the solution mixture from above, record the time taken for the cross to just disappear
- f) Into another clean conical flask, measure 50cm³ of **BA1**
- g) Add 10cm³ of distilled water to it and swirl
- h) Then add 10cm³ of **BA2** to the solution, swirl and place it on the cross
- i) Look from above, record the time t, taken for the cross to just disappear
- j) Repeat procedures (d) (i) but this time measure volumes of **BA1** as shown in the table below. Add a volume of distilled water to ensure that the total volume of **BA1** added to the flask remains 60cm^3
- k) Record your results in the table below.

Exp no.	Vol of BA1 (cm ³)	Vol of water added (cm ³)	Vol of BA2 (cm ³)	Time (s)	1/ _t (s ⁻¹)
1	60	0	10		
2	50	10	10		
3	40	20	10		
4	30	30	10		
5	20	40	10		

BA1 reacts with BA2 according to the equation.

Questi	ons:
(a)	Why does the cross disappear in the experiment?
(b)	Write the balanced equation for the reaction that took place between BA1 and BA2 .
(c)	Plot a graph of
	(i) Volume of BA1 against time, t
	(ii) Volume of BA1 against $\frac{1}{t}$
(d)	How does the concentration of BA1 affect the time for the cross to disappear.
(e)	Determine the rate of reaction between BA1 and BA2 and state the units.
	Fne

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Practice, Continuous reading and hard work makes chemistry easier.