

PREPARATION OF LYOPHILIC AND LYOPHOBIC SOLUTION

Principle:- The lyophilic sols are directly formed by mixing and shaking the substance with a suitable liquid. Lyophobic sols cannot be prepared by direct mixing and shaking, special methods are employed to prepare lyophobic sols.

Chemicals required

- 1] Egg albumin
- 2] Sodium chloride solution (5g in 100ml water)
- 3] Ferric chloride solution (2g in 100ml water)
- 4] Aluminium chloride solution (2g in 100ml water)
- 5] Starch 1gms - 500mg
- 6] Arsenious oxide (0.2g in 100ml water)

Procedure:-

A. Preparation of lyophilic sol.

I Egg albumin sol

1] Prepare 100ml of 5% (w/v) solution of NaCl in water in a 250ml beaker.

2] Break one egg in a porcelain dish

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and pipette out the albumin and pour it in sodium chloride solution. Stir well to ensure that the sol is well prepared.

II Starch / gum Sol

- 1] Take 100ml of distilled water in a 250ml beaker and boil it
- 2] Make a paste of 500mg starch or gum in hot water and transfer this paste in 100ml of boiling water with constant stirring up to water boiling and stirring for 10 minutes. cool. ensure that sol is well prepared.

B Preparation of lyophobic sol

Ferric hydroxide / Aluminium hydroxide

- 1] Take 100ml of distilled water in a 250ml beaker and boil it
- 2] Add 2g of ferric chloride / aluminium chloride powders to boiling water and stir it well.
- 3] Take 100ml of distilled water in another 250ml beaker and boil it
- 4] Pour 10ml of ferric chloride / aluminium chloride solution prepared in step (ii)

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and transfer it into the boiling water with constant stirring keep the water boiling till brown/cubeta sol is obtained

II Arsenious Sulphide Sol.

1] Take 100ml of distilled water in 250ml beaker

2] Add 0.2g of arsenious oxide to it and boil the content of the beaker.

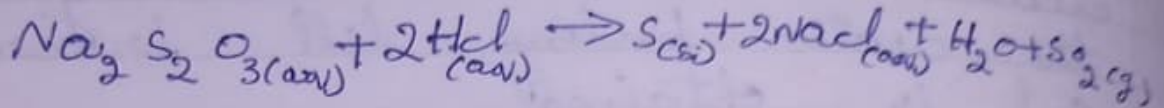
3] Cool and filter the solution.

4] Pass hydrogen sulphide (H_2S) gas through the filtered solution till it obtains the smell of H_2S (use Kipp's apparatus to pass hydrogen sulphide gas)

5] expel H_2S gas from the sol by slow heating and filter it

6] label the filtrate as arsenious sulphide sol.

Equation:



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EFFECT OF CONCENTRATION ON RATE OF REACTION

Aim: To study the effect of concentration on the rate of reaction between Sodium thiosulphate and hydrochloric acid.

Principle:- Sodium thiosulphate solution reacts with dilute hydrochloric acid and forms a sulphur precipitate of sulphur the time taken for a certain amount of sulphur to form can be used to indicate the rate of reaction.

Apparatus and chemical required.

100ml conical flask. 50ml measuring cylinder thermometer and a Stopwatch
0.1M Sodium thiosulphate solution 1M hydrochloric acid

Procedure:-

I Five 100ml clean conical flasks are taken and labeled as 1, 2, 3, 4, 5.

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2] 10ml of 0.05M Sodium thiosulphate solution is taken in the conical flask labeled 1

3] 10ml of dilute hydrochloric acid is added and immediately stop clock is started

4] The flask is swirled to mix the solution and placed on paper marked with a cross.

5] Marked cross is observed from the top. when the cross is just invisible time is noted.

6] Experiment is repeated by adding 10ml of HCl to the conical flask 2,3,4,5 containing 10ml of different concentration of Sodium thiosulphate solution as tabulated below.

[Different concentration of Sodium thiosulphate solution are made by mixing different volume of the Sodium thiosulphate solution with water as shown in the table]

A graph is plotted between the concentration of Sodium thiosulphate and the time taken for the marked cross to become just invisible.

It is clear from the graph that

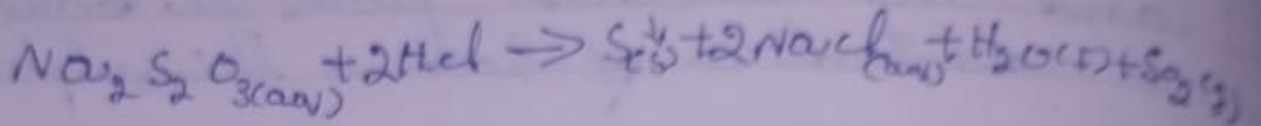
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rate of reaction between NO_2 , SO_2 , O_3 and HCl decreases with the increase in concentration of Sodium thiosulphate.

Result:- Rate of reaction between Sodium thiosulphate and hydrochloric acid decrease with the concentration of Sodium thiosulphate.

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Equation



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EFFECT OF TEMPERATURE ON RATE OF REACTION

Aim: To Study effect of temperature on rate of reaction between Sodium thiosulphate and hydrochloric acid.

Principle: The rate of reaction depends upon the temperature of the reaction generally rate of reaction increase with increase in temperature rate of reaction between Sodium thiosulphate and hydrochloric acid is studied by measuring the time taken for precipitation of Sulphur.

Apparatus and chemical required

100ml conical flask 25ml and 10ml measuring cylinder wire gauze tripod stand thermometer and a stop watch
0.1M Sodium thiosulphate Solution
and 1M Hydrochloric acid

Procedure

I 50ml of 0.1M Sodium thiosulphate Solution

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SL NO	Temperature in °C	Time taken for the printing to become invisible (s)
1	30°	166
2	40°	84
3	50°	62

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is taken in a clean 100-ml conical flask and kept in thermostat of 30°C

2] 10 ml of 1M hydrochloric acid is added and immediately stop clock is started

3] The flask is swirled to mix the solution and placed on paper marked with a cross.

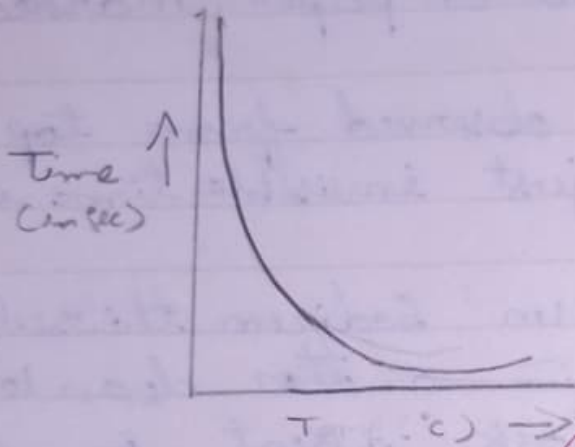
4] Marked cross is observed from top. when the cross is just invisible time is noted

5] Again 50 ml of 1M Sodium thiosulphate solution is taken in another clean 100 ml conical flask the flask is kept in a thermostat at a temperature of the solution is noted after placing it on a piece of printed paper. Immediately 10 ml of 1M HCl is added and a stop watch is started the time taken for the precipitation of Sulphur enough for to make printing on the paper just invisible is noted.

6] The same experiment is repeated at 50°C and 60°C and using fresh Sodium thiosulphate solution & HCl each time taken for paper mark to become just invisible is noted.

Observation

Volume of 0.1M Sodium thiosulphate



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Solution taken each time = 50ml

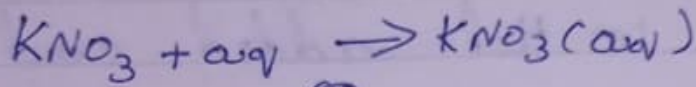
Volume of 1M HCl added each time = 10ml

A graph between the time v/s temperature is plotted and from shape of the curve

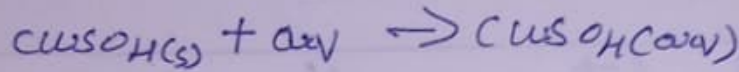
Result:-

Rate of reaction between Sodium thiosulphate and hydrochloric acid increase with the increase in temperature.

Reaction.



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DETERMINATION OF ENTHALPY OF SOLUTION

Aim :- To determine the enthalpy of solution of potassium nitrate & crystals of copper sulphate.

Principle :- A known amount of potassium nitrate dissolved in water the heat absorbed @ evolved is experimentally determined. Amount of heat absorbed @ evolved when one mole of potassium nitrate is dissolved in excess of the solvent at a given temperature is calculated. It gives the enthalpy of solution.

Apparatus and chemical required.
Polythene bottle fitted with thermometer potassium nitrate @ copper sulphate crystals.

Procedure.

100 cm³ of water is taken in a polythene bottle fitted with thermometer and stirrer about of

observation and calculation.

Initial temperature = $T_1, K = 301K$

Final temperature = $T_2, K = 300K$

change in temperature = $(T_1 - T_2), K = 1K$

Mass of KNO_3 taken = $w, g = 2.834g$

volume of the solution = $100cm^3$

Molecular mass of $KNO_3 = 101.1g\ mol^{-1}$

mass of the solution = $100g = 0.1kg$

Specific heat of solution = $4200\ J\ kg^{-1}\ K^{-1}$

Heat absorbed during the process

Mass \times Sp. heat \times change in temperature

$$= 0.1 \times 4200 \times (T_1 - T_2)$$

$$= 0.1 \times 4200 \times 1$$

$$= 420\ J$$

Amount of heat absorbed when one mol of potassium nitrate is dissolved in water

$$= \frac{420 \times 101.1}{w}$$

$$= 14983\ J$$

$$= 14.98\ kJ$$

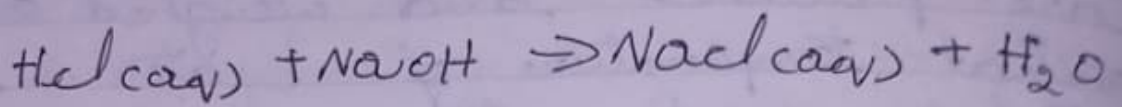
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2.3 g of KNO_3 (or) CuSO_4 crystals are powdered and weighed accurately. Initial temperature of water is noted. powdered potassium nitrate is quickly added to polystyrene bottle containing water and the bottle is immediately stoppered and solution is stirred well. The minimum temperature attained is noted. Enthalpy of solution is then calculated.

Result: Enthalpy of solution of potassium nitrate = $\Delta H = \text{kJ mol}^{-1}$

Note: To determine the enthalpy of solution of copper sulphate in the above experiment replace potassium nitrate by copper sulphate [molar mass of $\text{CuSO}_4 \cdot 5\text{H}_2\text{O} = 249$]

Reaction:-



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DETERMINATION OF ENTHALPY OF NEUTRALIZATION

Aim: To determine the enthalpy of neutralization of strong acid (HCl) with a strong base (NaOH).

Principle: A known amount of acid is neutralised by an equivalent determined amount of heat liberated per gram equivalent of an acid neutralized is calculated. It gives the enthalpy of neutralization.

Apparatus and chemical required.
Polythene bottle fitted with thermometer and stirrer. INHCl, INNaOH.

Procedure:

100 cm³ of INHCl is taken in a polythene bottle fitted with thermometer and stirrer. Both the solutions are kept in a

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Observation and calculation

$$\text{Initial temperature} = T_1 K = 301 K (28^\circ C)$$

$$\text{Final temperature} = T_2 K = 307.7 K (34.3^\circ C)$$

$$\text{Rise in temperature} = (T_2 - T_1) K = 6.83 K$$

Total Volume of the Solution - 200 cm^3

$$\text{Mass of the Solution} - 200 \text{ g} = 0.2 \text{ kg}$$

Specific heat of Solution - $4200 \text{ J/kg}^\circ\text{C}$

Heat liberated during the process
 $Q = \text{mass} \times \text{S.P Heat} \times \text{change in temperature}$

(i.e. when 0.1 gram eq. wt of HCl is neutralized)

$$Q = 0.2 \times 4200 \times (T_2 - T_1)$$

$$Q = 0.2 \times 4200 \times 6.83$$

$$Q = 5737.2 \text{ J}$$

Amount of heat liberated $Q = \frac{Q_1}{0.1}$

when one gram equivalent $Q = 57372 \text{ J}$

mass of HCl is neutralized $Q = 57.3 \text{ kJ}$

both for 15 minutes to attain the temperature of water both cubes both the solution attain steady temperature initial temperature is noted let it be T_1

Sodium hydroxide solution is quickly added to polythene bottle containing HCl solution and the bottle is shaken well the maximum temperature attained is noted let it be T_2

Result: Enthalpy of neutralization of HCl with NaOH is

$$H = -57.3 \text{ kJ mol}^{-1}$$

For Daniel cell Nernst equation is as follows

$$E_{\text{cell}} = E_{\text{cell}}^{\circ} + \frac{2.303RT}{nF} \log \frac{[Cu^{2+}]}{[Zn^{2+}]}$$

$$E_{\text{cell}} = E_{\text{cell}}^{\circ} + \frac{0.59}{2} \log \frac{[Cu^{2+}]}{[Zn^{2+}]} \text{ at } 298\text{K}$$

$$E^{\circ}_{\text{cell}} = E^{\circ}_{\text{C}} - E^{\circ}_{\text{A}} = E^{\circ}_{\text{Cu}} - E^{\circ}_{\text{Zn}}$$

$$= 0.34 - (-0.76)$$

$$= +1.10\text{V}$$

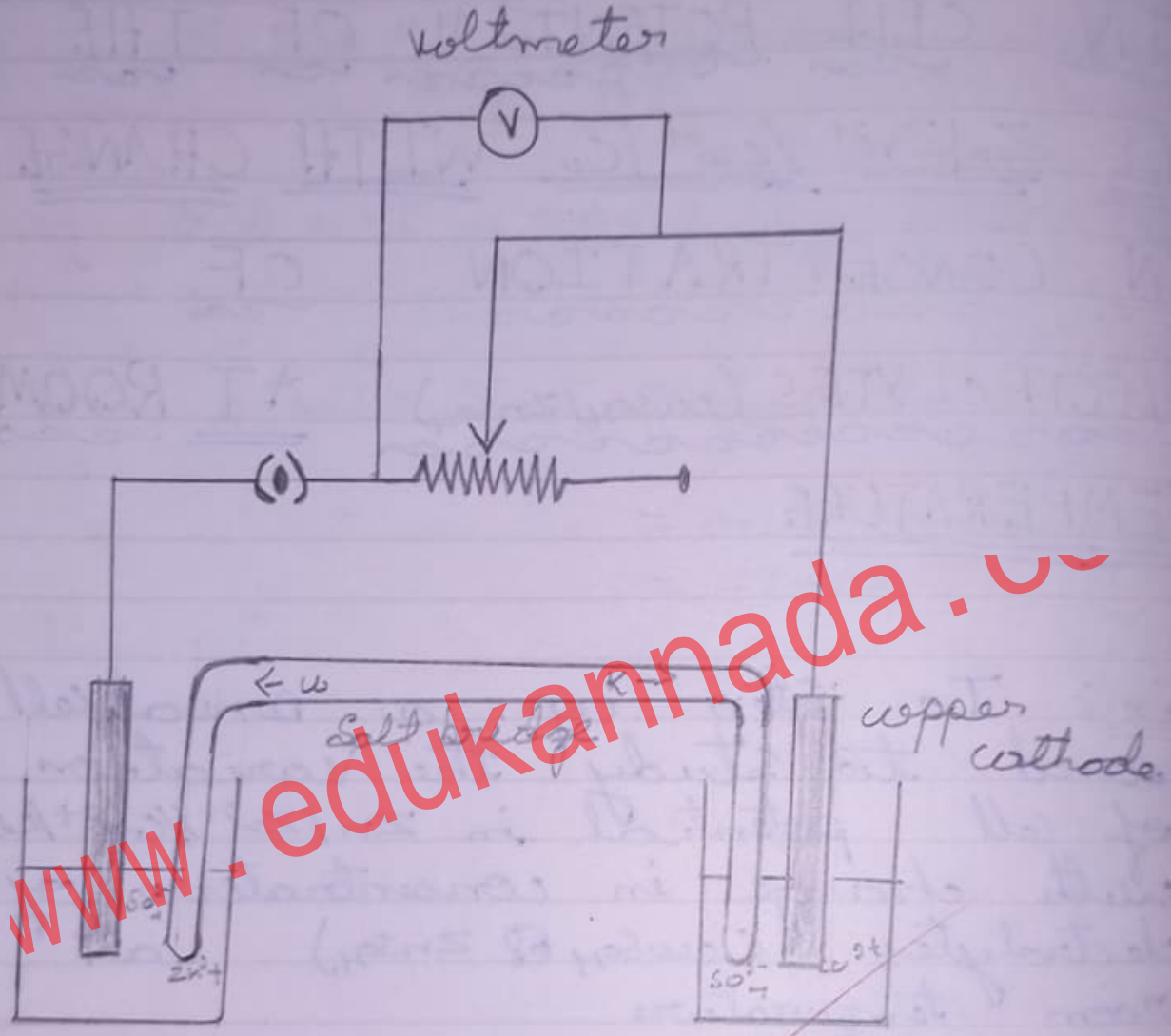
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TO STUDY THE VARIATION
IN CELL POTENTIAL OF THE
CELL $Zn/Zn^{2+} / Cu^{2+} / Cu$ WITH CHANGE
IN CONCENTRATION OF
ELECTROLYTES ($CuSO_4 / ZnSO_4$) AT ROOM
TEMPERATURE

Aim: To set up a Daniel cell and to study the variation of cell potential in $Zn/Zn^{2+} / Cu^{2+} / Cu$ with change in concentration of electrolytes ($CuSO_4$ or $ZnSO_4$) at room temperature.

Theory: Daniel cell is $Zn/ZnSO_4 / CuSO_4 / Cu$

The redox reaction of Daniel cell can be carried out in the laboratory either by using a salt bridge or porous pot. Here we shall carry out the redox reactions



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using KCl Salt bridge. which will prevent the mixing of two solutions. The complete redox reaction involves the two half cell reactions. In the Daniel cell is the concentration of $ZnSO_4$ solution is $1M$ & that of $CuSO_4$ solution is $1M$ the cell potential is 1.1 Volts. However if the concentration of $ZnSO_4$ solution and $CuSO_4$ solution are different then the cell potential can be calculated using Nernst equation.

Materials:- A strip of copper & a strip of zinc a large beaker bowl @ a porous pot a plastic tube cotton 100g of copper sulphate ($CuSO_4$) 100g of Zinc sulphate ($ZnSO_4$) Distilled water, a voltmeter (two cables alligator) clips KCl salt bridge

Procedure:-

- 1] About $100cm^3$ of $1M$ $ZnSO_4$ solution is taken in a porcelain hood. A zinc rod is dipped in it.
- 2] About $100cm^3$ of $1M$ $CuSO_4$ solution is taken in a small beaker. A copper rod is dipped in it. A salt bridge of

Trial concentration of $ZnSO_4$ solution concentration of $CuSO_4$ solution observed emf of the cell (volt) calculated E_{cell} from Nernst equation

A	1	1M	1M
	2	0.4M	1M
	3	0.2M	1M
	4	0.1M	1M

B	1M	0.4M
	1M <td>0.2M </td>	0.2M
	1M <td>0.1M </td>	0.1M

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KCl is used to connect the two solutions thus Daniell cell is formed

3] The copper plate with +ve polarity and zinc plate with the -ve polarity are connected to the voltmeter as shown

4] As soon as the circuit is completed electric current will begin to flow and then the voltmeter reading is noted which indicates the e.m.f. of the cell. This is reading No. 1

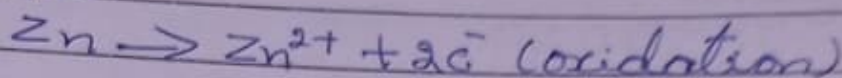
5] The above procedure is repeated while keeping the concentration of copper sulphate solution as 1M but changing the concentration of zinc sulphate solution by dilution to 0.05, 0.02 and 0.01M. Then reading 2, 3 and 4 are recorded.

6] Again the above procedure is repeated while keeping the concentration of $ZnSO_4$ solution as 1M but changing the concentration of $CuSO_4$ solution by dilution to 0.05, 0.02 and 0.01M. Thus reading 5, 6 and 7 are recorded.

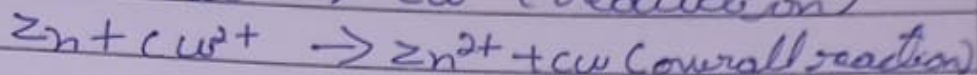
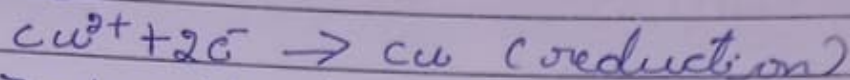
The reaction at the electrodes can be represented by the equation:

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At: anode



At: cathode

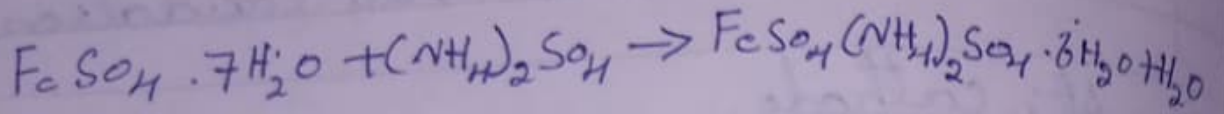
conclusions

Increase in $[\text{Cu}^{2+}]$ increases the emf of the cell

Increase in $[\text{Zn}^{2+}]$ decreases the emf of the cell.

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Reaction:-



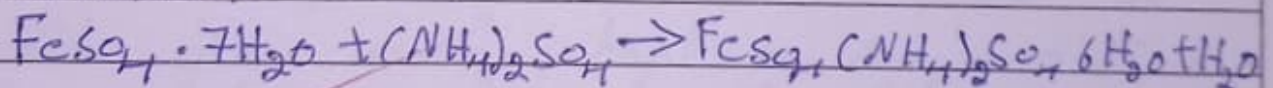
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PREPARATION OF FERROUS AMMONIUM SULPHATE CRYSTALS

Aim: To prepare crystals of ferrous ammonium sulphate or Mohr's salt.

Theory: Mohr's salt i.e. ferrous ammonium sulphate $[FeSO_4(NH_4)_2SO_4 \cdot 6H_2O]$ is a double salt. It can be prepared by mixing equal volume of equimolar solutions of $FeSO_4 \cdot 7H_2O$ and $(NH_4)_2SO_4$ of same concentration and cooling. Crystals of Mohr's salt are formed.

Reactions:-



Apparatus and chemicals required

Two 25ml beakers, a glass rod, china dish, funnel stand wire gauze tripod stand and filter paper.
Ferrous sulphate ammonium sulphate, dilute sulphuric acid and ethanol.

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Procedure:-

- 1] 14g of ferrous sulphate and 6.5g of ammonium sulphate are taken in a clean beaker
- 2] About 50ml of distilled water is boiled in another beaker for about 5 minutes to expel dissolved air.
- 3] About 2ml of dilute sulphuric acid is added to the above water and it is poured into the first beaker containing $FeSO_4$ and $(NH_4)_2SO_4$. The mixture is stirred with a glass rod to dissolve the salts.
- 4] The solution is filtered to remove any suspended impurities.
- 5] The filtrate is collected in a clean china dish and concentrated on a low flame until the crystallization point is reached.
- 6] The china dish is covered with a watch glass and kept in an undisturbed place for cooling. Well defined crystals will be obtained after some time.
- 7] The mother liquor is removed and the crystals in the china dish are washed with a small quantity of ethanol.
- 8] The crystals are dried by pressing between the folds of rough filter.

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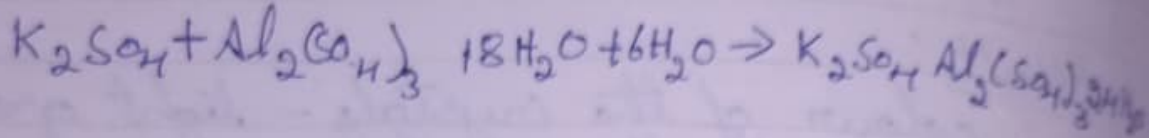
paper and weighed. the colour and shapes of the crystals are observed

Result :-

colour of the crystals - light green
shape of the crystals - monoclinic
yield (mass of the crystal obtained) - 11.865g

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Reactions



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PREPARATION OF POTASH ALUM

Aim: To prepare crystals of potash alum

Theory: Potash alum is a double salt commonly known as filterite has the formula $K_2SO_4 \cdot Al_2(SO_4)_3 \cdot 24H_2O$. It can be prepared by the crystallization of a mixture of equal volume of equimolar solutions of $Al_2(SO_4)_3 \cdot 18H_2O$ and K_2SO_4 .

Apparatus and chemical required

Two 25ml beakers, a glass rod, china dish, funnel stand, wire gauze, tripod stand and filter paper. Potassium sulphate, aluminium sulphate and concentrated sulphuric acid.

Procedure:-

1. 7g of potassium sulphate is taken in a clean beaker. It is dissolved in a minimum amount

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of distilled water with gentle heating.

2] 26.5g of aluminium sulphate is taken in another beaker about 3-4 drops of concentrated sulphuric acid are added about 20ml of distilled water is added to the beaker and heated on a wire gauze with occasional stirring. The solution is thus filtered to remove the turbidity.

3] The two clear solutions are mixed in a china dish. The dish heated gently on a wire gauze and the solution is concentrated to the crystallization point.

4] The dish is covered and cooled without any disturbance.

5] The crystals formed are separated by decanting the mother liquor. Crystals are washed with very small quantity of ice cold water.

6] The crystals are then dried between the folds of filter paper and weighed. The colour and shape of the crystals are observed and noted.

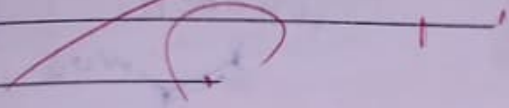
Result :-

colour of the crystals = colourless

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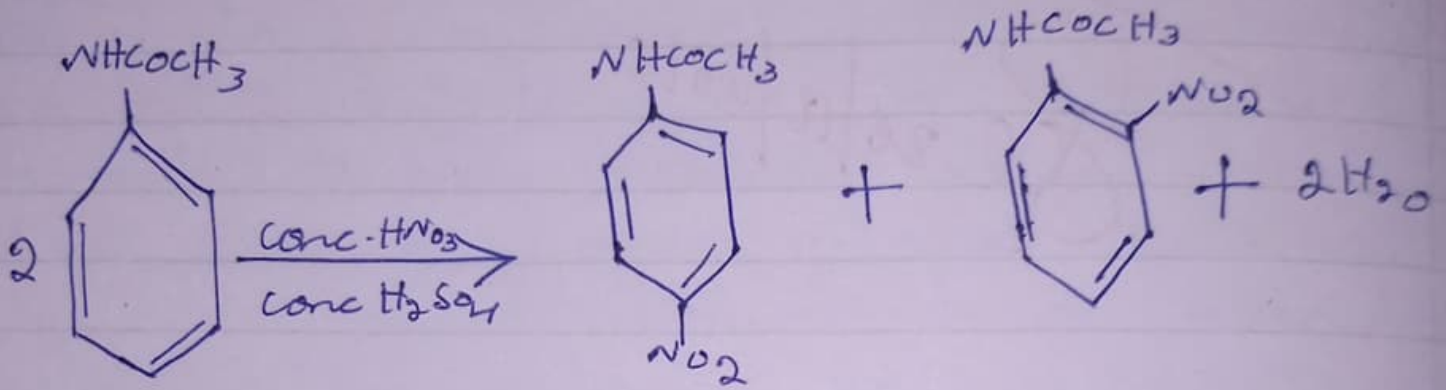
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shape of crystals = octahedral
yield = 10.484g.



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Chemical reaction.



Acetanilide

p-nitro acetanilide o-nitro acetanilide (major)

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PREPARATION OF P-NITROACETANILIDE OF From ACETANILIDE

Aim:- To prepare a pure sample of p-nitroacetanilide.

Apparatus and chemical required.

25ml round bottom flask ice bath
beaker measuring cylinder funnel
and filter paper acetanilide con
Nitric acid Sulphuric acid glacial
acetic acid and rectified spirit

Procedure:-

- 1] 10g of powdered acetanilide is placed in a 250ml of round bottomed flask and 10ml of glacial acetic acid is added.
- 2] The mixture is stirred well and 20ml of conc Sulphuric acid is added to it.
- 3] The flask is placed in ice bath and cooled.
- 4] 4ml of cold conc Nitric acid is added drop by drop with constant

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stirring.

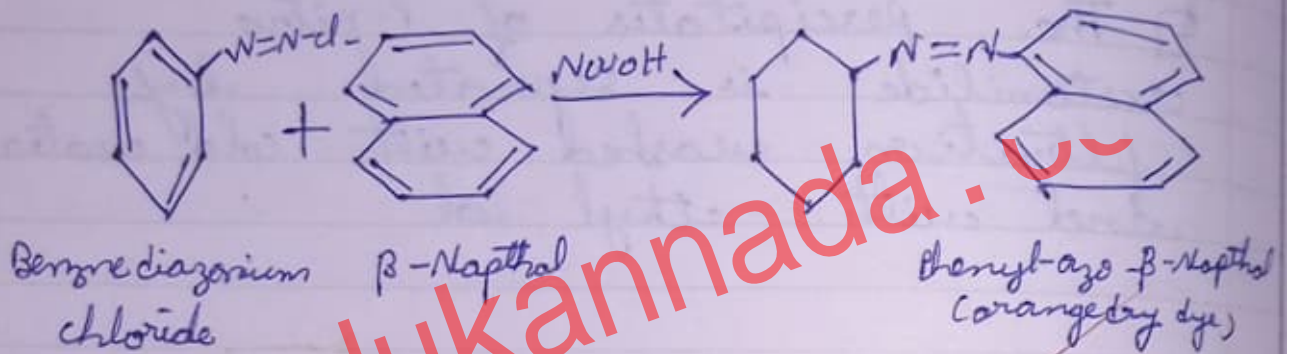
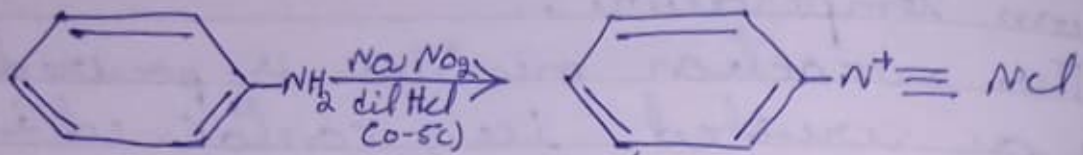
5] The flask is removed from the ice bath and allowed to stand for about 15 mins at room temperature.

6] The reaction mixture is poured on a crushed ice contained in a beaker and allowed to stand for 15 mins.

7] The precipitate of p-nitroacetanilide is separated by filtration washed with cold water and with ethyl alcohol.

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Chemical reaction



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PREPARATION OF PHENYL-AZO-β-NAPHTHOL

(an azo dye)

Aim: To prepare phenyl-azo-β-naphthol.

Apparatus and chemical required.

Conical flask beaker funnel
measuring cylinder ice bath. Aniline
β-naphthol sodium nitrite conc. HCl
Sodium hydroxide glacial acetic
acid and Spirit.

Procedure

- 1] 2.5 ml of pure aniline 8ml of conc. HCl and 8ml of water are mixed in a beaker
- 2] The solution is cooled to 0-5°C in an ice bath.
- 3] To the cold solution a solution of sodium nitrite is slowly added Benzene diazonium chloride solution is formed.
- 4] 22.5ml of 10% NaOH solution is taken in a beaker & 4.0g of pure β-naphthol is added
- 5] The cold benzene diazonium

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chloride solution is added with constant stirring to the β -naphthol
b) The mixture is allowed to stand for 10 min. an orange red dye is filtered.

Result :- Yield = 5.4 g

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SYSTEMATIC QUALITATIVE
ANALYSIS OF SIMPLE INORGANIC
SALT

Preliminary test

State :- Amorphous Solid

colour :- white

Solubility :- Soluble in cold dil HCl

Detection of acid radicals.

Detection of I group acid radical

Experiment	Observation	Inference
Salt + dil HCl	brick efferv.	I group acid radical CO_3^{2-} is present.

confirmatory test for CO_3^{2-}

Experiment	Observation	Inference
Salt + water, heat strongly pass the	lime water does not turn milky	CO_3^{2-} is confirmed

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through lime
water

Detection of basic radicals

Detection of zero group basic radical (NH_4^+)

Test for NH_4^+

Experiment	Observation	Inference
Salt + NaOH Solution heat	No pungent Smelling gas	Zero group basic radical NH_4^+ is absent.

Detection of first group basic radical

Experiment	Observation	Inference
Original Solution + dil. HCl	No precipitate	I group basic radical are absent.

Detection of Second group basic radical

Experiment	Observation	Inference
Original Solution + dil. HCl	No precipitate	II group basic radicals are absent

Detection of third group basic radical

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Experiment	Observation	Inference
original solution + NH_4Cl (Solid) + NH_4OH (excess)	no precipitate	III group basic radical are absent

Detection of fourth group basic radical

Experiment	Observation	Inference
original solution + NH_4Cl (Solid) + NH_4OH (excess) + H_2S	no precipitate	IV group basic radicals are absent.

Detection of fifth group basic radical

Experiment	Observation	Inference
original solution + NH_4Cl (Solid) + NH_4OH (excess) + $(\text{NH}_4)_2\text{CO}_3$	A white ppt is formed	V group basic radical present Ba^{2+} , Ca^{2+} or Sr^{2+} may be present

Confirmatory test for Ba^{2+} , Ca^{2+} or Sr^{2+}

Experiment	Observation	Inference
original solution + dilute acetic acid		

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K_2CO_3 Solution	Yellow ppt. is not obtained	Ba^{2+} is not confirmed.
original solution + $(NH_4)_2SO_4$ solution	no white precipitate is obtained	SO_4^{2-} is not confirmed.
original solution + NH_4OH + ammonium oxalate	white precipitate is obtained	Ca^{2+} is confirmed

Flame test

Experiment

Observation

Inference

Salt + conc. HCl + $MgSO_4$, ignite in platinum wire to the non-luminous flame

Brick red colour is imparted

Ca^{2+} is confirmed.

Report: The given inorganic salt contains

acid radical is CO_3^{2-}

base radical is Ca^{2+}

Hence, given salt is $CaCO_3$

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Salt No 2

Preliminary tests

State :- crystalline
 colour :- white

Solubility :- Soluble in cold water
 original solution :- Salt + water

Detection of acid radicals

Detection of I group acid radicals

Experiment	Observation	Inference
Salt + dil. HCl	no gas is evolved	I group acid radicals are absent.

Detection of II group acid radicals

Experiment	Observation	Inference
Salt + Conc. H_2SO_4 in dry test tube	Reaction takes place in cold with evolution of gas	II group acid radical (Cl or Br) may be present.
Add in testing 2	white fumes evolved	Cl is present

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confirmatory test for chloride

Experiment	Observation	Inference
Salt + potassium dichromate crystal + conc. H_2SO_4 heat	Reddish brown vapour of chromyl chloride are evolved	Cl is confirmed
Pass the vapour into the NaOH solution	Yellow solution is obtained	
Yellow solution + dilute acetic acid + lead acetate solution	Yellow precipitate is formed.	

Silver nitrate test for Cl⁻ radical

Experiment	Observation	Inference
Salt solution + dil HNO_3 + Silver nitrate solution	A curdy white precipitate is formed which is soluble in NH_4OH	Cl is confirmed.

Detection of basic radical.

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Detection of zero group basic radical (NH_4^+)

Test for NH_4^+

Experiment

Observation

Inference

Salt + NaOH
Solution heat

Crises a pungent
Smelling gas
which gives dense
white fumes

Zero group
basic radical
 NH_4^+ is
present

Confirmatory test for NH_4^+ (Nessler's
Seign's test)

Experiment

Observation

Inference

Original Solution
+ Nessler's Seign's
+ Excess of NaOH

A brown
precipitate
is formed

NH_4^+ is
confirmed

Report:- The given inorganic salt contains
A acid radical Cl^-
B) Basic radical NH_4^+
Hence the given salt is NH_4^+Cl^-

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Salt no: - 3

~~Preliminary test~~

State: - Amorphous Solid

colour: - white

Solubility: - Salt + dilute HCl

Original Solution: Salt + dilute HCl

~~Detection of acid radical~~

~~Detection of I group acid radical~~

Experiment	Observation	Inference
Salt + dil HCl	No black precipitate	I group acid radical are absent

~~Detection of II group acid radical~~

Experiment	Observation	Inference
Salt + conc. H_2SO_4 in dry test tube	It does not take place in cold condition Dense white fumes are evolved	II group acid radical not present It is present.

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Confirmatory test for chloride

Experiment	Observation	Inference
Salt + potassium dichromate crystal + conc. H_2SO_4 heat	Reddish brown vapour are formed	Cl^- is confirmed.
Pass the vapour into the $NaOH$ solution	Yellow solution is obtained	
Yellow solution + dilute acetic acid + lead acetate solution	Yellow precipitate is obtained	

Silver nitrate test for Cl^-

Experiment	Observation	Inference
Salt solution + dil HNO_3 + Silver nitrate solution	A curdy white precipitate is formed which is soluble in NH_4OH	Cl^- is confirmed

Detection of basic radical

Detection of zero group basic radical

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Experiment	observation	Inference
Salt + NaOH Solution heat	No pungent Smelling gas	Zero group basic radical NH_4^+ is absent

Detection of first group basic radical

Experiment	observation	Inference
Original Solution + dil HCl	No white precipitate	I group basic radical are absent

Detection of II group basic radical

Experiment	observation	Inference
Original Solution + dil HCl + H_2S	No precipitate	II group basic radicals are absent.

Detection of III group basic radical

Experiment	observation	Inference
Original Solution + NH_4Cl (Solid) + NH_4OH excess	no precipitate	III group basic radical are absent.

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Detection of fourth group basic radical

<u>Experiment</u>	<u>Observation</u>	<u>Inference</u>
original solution + NH_4Cl (Solid) + NH_4OH (excess) + H_2S	no precipitate	IV group basic radicals are absent.

Detection of fifth group basic radical

<u>Experiment</u>	<u>Observation</u>	<u>Inference</u>
original solution + NH_4Cl (Solid) + NH_4OH (excess) + $(\text{NH}_4)_2\text{CO}_3$	A white ppt is obtained	V group basic radical Ba^{2+} , Ca^{2+} or Sr^{2+} may be present

<u>Confirmatory test for Ba^{2+}, Ca^{2+} or Sr^{2+}</u>		
<u>Experiment</u>	<u>Observation</u>	<u>Inference</u>
original solution + dil acetic acid + K_2CrO_4 solution	Yellow ppt is obtained	Ba^{2+} is confirmed.

<u>Flame test</u>		
Salt (ion. Hal) make paste, ignite on platinum wire to the non-luminous flame	Apple green colour is imparted	Ba^{2+} is confirmed.

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Report - The given inorganic salt contains

A] Acid radical is Cl^-

B] Basic radical is Ba^{2+}

Hence the given salt is BaCl

(1)

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Salt no:- H

Preliminary test

State :- crystalline solid

colour :- white

Solubility :- Soluble in cold water

original solution - salt + cold water

Detection of acid radical

Experiment	Observation	Inference
Salt + dil HCl	No brisk appearance	I group acid radical are absent.

Detection of II group acid radicals.

Experiment	Observation	Inference
Salt + conc. H_2SO_4 in a dry test tube	rxn take place in cold condition	II group acid radical Cl or Br^- may be present.
	dense white fumes are evolved	Cl is present

confirmatory test for chloride

Experiment	Observation	Inference

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Salt + potassium
dichromate crystal
+ conc. H_2SO_4 , heat

reddish brown
vapour are
formed

Pass the vapour into
the $NaOH$ Solution

Yellow Solution
is obtained

Cl^- is
confirmed

Yellow Solution +
dilute acetic acid +
lead acetate Solution

Yellow precipitate
is obtained

Silver nitrate test for Cl^- or Br^- radicals.

Experiment

Observation

Inference

Salt Solution + equal
dil. HNO_3 +
Silver nitrate Solution

A curdy white
precipitate is
formed

Cl^- is
confirmed.

Detection of Basic Radical.

Detection of zero group basic radical (NH_4^+)

Experiment

Observation

Inference

Salt + $NaOH$
Solution
heat

No pungent
Smelling
gas

zero group
basic radical
 NH_4^+ is
absent.

Detection of first group basic radical

Experiment	observation	Inference
original solution + dil HCl	No white precipitate	I group basic radicals are absent.

Detection of second group basic radical

Experiment	observation	Inference
original solution + dil HCl + H_2S	No precipitate	II group basic radicals are absent.

Detection of third group basic radical

Experiment	observation	Inference
original solution + NH_4Cl (solid) + NH_4OH (excess)	no precipitate	III group basic radical are absent.

Detection of fourth group basic radical

Experiment	observation	Inference
original solution		

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NH_4Cl (Solid) + NH_4OH (aq) + $(\text{NH}_4)_2\text{CO}_3$	no precipitate	V group basic radical are absent.
---	-------------------	---

Detection of Sixth group basic radical

Exp: Since the given radical is not present in the previous groups it is present in VI group. It may be Mg^{2+}

Confirmatory test for Mg^{2+}

Experiment	Observation	Inference.
original solution of NH_4Cl (Solid) + NH_4OH + Sodium dihydrogen ortho phosphate	A white ppt is formed	VI^{th} G.B.R is Present Mg^{2+} is confirmed

Report :- The given inorganic salt.

- A) Acid radical is Cl^-
B) Basic radical is Mg^{2+}

Hence the given salt is MgCl_2

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Salt No. - 5
 ~~~~~

Preliminary test  
 ~~~~~

State: Crystalline
 colour: white

Solubility: Soluble in cold water
 original solution = salt + cold water

Detection of acid radical
 ~~~~~

Detection of I group acid radical  
 ~~~~~

Experiment	Observation	Inference
Salt + dil HCl	No gas is evolved	I group acid radicals are absent.

Detection of II group acid radical
 ~~~~~

| Experiment                                      | Observation              | Inference          |
|-------------------------------------------------|--------------------------|--------------------|
| Salt + conc. $H_2SO_4$<br>in a dry test<br>Tube | Redish vapour is evolved | $Br^-$ is present. |

Confirmatory test for  $Br^-$   
 ~~~~~

Experiment	Observation	Inference
------------	-------------	-----------

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Salt Solution + equal
Volume of Cl_2 water +
clear Shake & allow
to settle

Orange red
globule is
formed

Br^- is
confirmed.

Detection of basic radical.

Detection of zero group basic radical (NH_4^+)

Experiment	Observation	Inference
Salt + NaOH Solution heat	give a dense white fumes with a strong ammoniacal smell	Zero group Basic radical NH_4^+ is present.

Confirmatory test for NH_4^+

Experiment	Observation	Inference
original solution + Nessler's reagent + excess of NaOH	A brown precipitate is formed	NH_4^+ is confirmed.

Report :- The given inorganic salt contains

A) Acid radical is Br^-

B) Basic radical is NH_4^+

Hence the given salt is $\text{NH}_4^+ \text{Br}^-$

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Salt no 6

Preliminary test

State :- Amorphous Solid

colour :- white

Solubility :- Soluble in cold water

original solution = Salt + cold water

Detection of acid radical

Detection of I group acid radical

Experiment	Observation	Inference
Salt + dil HCl	no precipitate appearance	I group acid is absent.

Detection of II group acid radical

Experiment	Observation	Inference
Salt + conc H ₂ SO ₄ in a dry test tube	no rxn take place	II group acid radical Cl ⁻ & Br ⁻ is absent
Add copper turnings heat strongly	no rxn take place	II group acid radical is absent

Detection of III group acid radical

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Experiment	Observation	Inference
Salt Solution + BaCl_2 Solution add excess of dil. HCl	A white precipitate is formed	III group acid radical (CO_3^{2-}) is present & confirmed.

Detection of basic radical

Detection of zero group basic radical (NH_4^+)

Experiment	Observation	Inference
Salt + NaOH Solution test	no pungent smelling gas	zero group basic radical NH_4^+ is absent.

Detection of first group acid radical

Experiment	Observation	Inference
original Solution + dil. HCl	no white precipitate	I group basic radical are absent.

Detection of second group basic radical

Experiment	Observation	Inference
original Solution + dil. HCl + H_2S	No precipitate	II group basic radical are absent.

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Detection of third group basic radical

Experiment	Observation	Inference
original solution + NH_4Cl (Solid) + NH_4OH (excess)	A gelatinous white ppt is formed	III group basic radical is present

Confirmatory test for Al^{3+}

Experiment	Observation	Inference
original solution + NaOH solution	white ppt is formed	Al^{3+} is confirmed.
clear solution + Solid NH_4Cl heat	which is soluble in excess of NaOH Gelatinous white ppt reappears	

Report :- The given inorganic salt contains
 A) Acid radical is SO_4^{2-}
 B) Basic radical is Al^{3+}
 Hence the given salt is $\text{Al}_2(\text{SO}_4)_3$

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Salt no 8
~~~~~Preliminary test  
~~~~~State :- Amorphous solid  
colour :- white.

Solubility :- Soluble in cold water

Original Solution = salt + cold water

Detection of acid radicals
~~~~~Detection of F group acid radical  
~~~~~

Experiment	Observation	Inference
Salt + dil HCl	no brisk effervescence	F group acid radical is absent.

Detection of II group acid radical
~~~~~

| Experiment                               | Observation       | Inference                        |
|------------------------------------------|-------------------|----------------------------------|
| Salt + conc $H_2SO_4$ in a dry test tube | no rxn take place | II group acid radical is absent. |
| Add Cu turnings & heat strongly          | no rxn take place | II group acid radical is absent. |

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### Detection of III group acid radical

Test for  $SO_4^{2-}$

| Experiment                                                                | Observation                                                           | Inference                                                               |
|---------------------------------------------------------------------------|-----------------------------------------------------------------------|-------------------------------------------------------------------------|
| Salt Solution +<br>BaCl <sub>2</sub> Solution<br>add excess of<br>dil HCl | white ppt is<br>insoluble in<br>excess of dilute<br>hydrochloric acid | III G A R is<br>Present<br><br>$SO_4^{2-}$ is<br>present &<br>confirmed |

### Detection of basic radicals

#### Detection of zero group basic radical ( $NH_4^+$ )

| Experiment                    | Observation                | Inference                                          |
|-------------------------------|----------------------------|----------------------------------------------------|
| Salt + NaOH/<br>Solution heat | No pungent<br>Smelling gas | Zero group<br>basic radical<br>$NH_4^+$ is absent. |

#### Detection of first group basic radical

| Experiment                     | Observation             | Inference                               |
|--------------------------------|-------------------------|-----------------------------------------|
| Original Solution +<br>dil HCl | No white<br>precipitate | I group basic<br>radical are<br>absent. |

#### Detection of Second group basic radical

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Experiment  
original Solution  
+  $\text{NH}_4\text{Cl}$  (Solid)  
+  $\text{NH}_4\text{OH}$  (excess)

Observation  
no  
precipitate

Inference  
III group  
basic radicals  
are absent.

Detection of fourth group basic radical

Experiment

Observation

Inference

original Solution  
+  $\text{NH}_4\text{Cl}$  (Solid)  
+  $\text{NH}_4\text{OH}$  (excess)  
+  $\text{H}_2\text{S}$

no  
precipitate

IV group  
basic radical  
are absent

Detection of fifth group basic radical

Experiment

Observation

Inference

original Solution  
+  $\text{NH}_4\text{Cl}$  (Solid)  
+  $\text{NH}_4\text{OH}$  (excess)  
+  $(\text{NH}_4)_2\text{CO}_3$

no precipitate

V group  
basic radical

Detection of Sixth group radical

Since the given (Salt) radical is not present in the previous groups, it is present in VI group. It may be right



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Confirmatory Test for  $Mg^{2+}$ 

| Experiment                                                                          | Observation           | Inference                                                            |
|-------------------------------------------------------------------------------------|-----------------------|----------------------------------------------------------------------|
| Original Solution + $NH_4Cl$ (solid) + $NH_4OH$ + Sodium dihydrogen ortho phosphate | A white ppt is formed | $Mg^{2+}$ is confirmed<br><del>VI<sup>th</sup> B.P. is Present</del> |

Report:- The given inorganic salt contains

A) Acid Radical is  $SO_4^{2-}$

B) Base Radical is  $Mg^{2+}$

Hence, the given salt is  $MgSO_4$

## Observation

Burette :-  $KMnO_4$  Solution  
conical flask :-  $10\text{cm}^3$  FAS Solution + 17ml  
tube full  
Indicator :-  $KMnO_4$  itself (Self Indicator)  
End point :- colourless to permanent  
pale pink

## Tabular column:

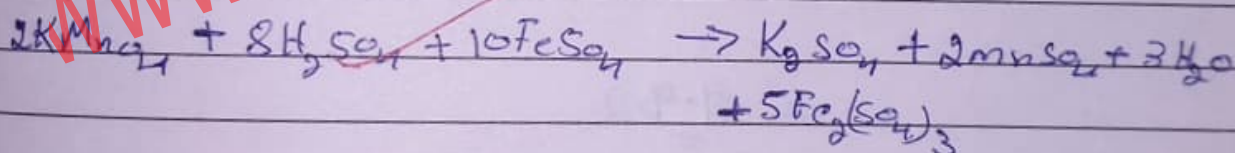
| Trial number                                       | I   | II    | III |
|----------------------------------------------------|-----|-------|-----|
| Final Burette Reading                              | 9.5 | 9.5 ✓ |     |
| Initial Burette Reading                            | 0.0 | 0.0   |     |
| Volume of $KMnO_4$ Solution added in $\text{cm}^3$ | 9.5 | 9.5   |     |



To determine the concentration of  $KMnO_4$  Solution by titrating it against 0.1M Standard Solution of ferrous Ammonium Sulphate [FAS]

Aim: To determine the concentration or molarity of  $KMnO_4$  Solution by titrating it against 0.1M Standard Solution of ferrous Ammonium Sulphate [FAS]

Principle: A known volume of Standard FAS Solution (Ferrous Salt Solution) is titrated against potassium permanganate Solution in acid medium in the reaction ferrous Sulphate gets oxidized to ferric Sulphate &  $KMnO_4$  gets reduced to colourless manganous Sulphate



By knowing the volume of  $KMnO_4$  Solution required to oxidize ferrous Sulphate from the titration the molarity of  $KMnO_4$  Solution is calculated.

Procedure: A 10 cm<sup>3</sup> pipette is washed with water, & then rinsed with given FAS Solution. A Burette is washed with water & then rinsed with  $KMnO_4$  Solution. It is

Calculation:-

$$\frac{M_1 V_1}{\alpha_1} = \frac{M_2 V_2}{\alpha_2}$$

$\text{KMnO}_4$  FAS

where  $\alpha_1$  &  $\alpha_2$  are the stoichiometric coefficients of  $\text{KMnO}_4$  &  $\text{Fe}$  in the balanced chemical equation

$$\alpha_1 = 1 \quad \& \quad \alpha_2 = 5$$

$$\therefore M_{\text{KMnO}_4} = \frac{\alpha_{\text{KMnO}_4} \times M_{\text{FAS}} \times V_{\text{FAS}}}{\alpha_{\text{FAS}} \times V_{\text{KMnO}_4}}$$

$$= \frac{1 \times 0.1 \times 10}{5 \times 9.5} = 0.02 \text{ M}$$

$$= \underline{0.02}$$



filled with  $KMnO_4$  solution & fixed to a stand. Initial Burette reading is noted. The upper meniscus is Solution.  $10\text{ml}^3$  of  $FeSO_4$  solution is pipetted out into a clean conical flask. Dilute sulphuric acid is added. The solution is titrated against the  $KMnO_4$  solution. The Burette till the solution turns permanent pale pink colour. Burette reading is noted. Titrations are repeated for concordant values. In this titration  $KMnO_4$  acts as self indicator.

Result: Molarity of  $KMnO_4 = 0.02\text{ M}$ .



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Aim: Test for the functional group present in organic compound

1] Test for unsaturation

| Experiment                                                                                                          | Observation                            | Inference                                 |
|---------------------------------------------------------------------------------------------------------------------|----------------------------------------|-------------------------------------------|
| 1] Baeyer's test<br>organic compound is dissolved in water or acetone + a few drop of 1% alkaline $KMnO_4$ solution | pink colour of $KMnO_4$                | The given organic compound is unsaturated |
| 2] Bromine water test<br>organic compound is dissolved in $CCl_4$ + 2% solution of bromine in $CCl_4$ dropwise      | orange red colour of bromine discharge | The given organic compound is unsaturated |

2] Test for Alcoholic (R-OH) group:-

|                                                                                                               |                                  |                      |
|---------------------------------------------------------------------------------------------------------------|----------------------------------|----------------------|
| 1] Ceric ammonium nitrate test<br>organic compound + few drops of ceric ammonium nitrate solution shaken well | A pink or red colour of $KMnO_4$ | Alcoholic (OH) group |
|---------------------------------------------------------------------------------------------------------------|----------------------------------|----------------------|

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## 2) Esterification

organic compound + acetic acid + 2 drops of conc.  $H_2SO_4$  keep it in boiling water bath for 5 minutes pour the mixture in  $Na_2CO_3$  solution

Δ fruity  
Smell

Alcoholic  
(-OH)  
group  
present

3) Test for Phenolic ( $Ar-OH$ ) groupa) Neutral  $FeCl_3$  test

organic compound in a test tube + a few drops of neutral  $FeCl_3$  solution

Δ violet  
colour

The given  
organic  
compound  
contains a  
phenolic group

## b) Phthalic test

organic compound + phthalic anhydride + conc.  $H_2SO_4$  heated and added to  $NaOH$

A pink  
colour

The given  
organic compound  
contains a  
phenolic group

4) Test for Aldehydic ( $-CHO$ ) group

## a) 2,4-DNP test

organic compound in a test tube + 2,4-Dinitrophenyl hydrazine

orange yellow  
crystalline  
precipitate is  
formed

The given organic  
compound contains  
aldehydic or a  
ketonic group

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## Schiff's test

organic compound in a test tube + Schiff's reagent

A pink colour is obtained

The given organic compound contains aldehydic group

## a) Tollen's test

organic compound in a test tube + Tollen's reagent keep in a boiling water bath

A black red precipitate or silver mirror is formed

The given organic compound confirms the presence of an aldehydic group

## d) Benedict's / Fehling's test :

organic compound + Benedict's / Fehling's solution & heated

A brick red precipitate is obtained

The given organic compound contains aldehydic group

5] Test for Ketonic ( $-CO-$ ) group

a) m-dinitrobenzene test  
organic compound in a test tube + m-dinitrobenzene (S) + dil. NaOH solution & shake well

A violet colour which disappears

The given organic compound contains Ketonic group

b) Sodium nitroprusside test



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organic compound  
in a test tube +  
Sodium nitroprusside  
solution + NaOH solution

A fine red  
colour is  
obtained

The given organic  
compound  
confirms the  
presence of a  
ketonic group

### 1] Test for carboxylic (-COOH) group

#### a) Litmus test

organic compound  
in a test tube +  
blue litmus paper

Blue litmus  
paper turns  
red

The given organic compound  
contains a carboxylic  
group

b) Sodium Bicarbonate test  
organic compound in a  
test tube + NaHCO<sub>3</sub>  
solution

Bubbles effusion  
of CO<sub>2</sub>

The given organic  
compound contains  
carboxylic group

#### c) Esterification

organic compound  
in a test tube +  
ethyl alcohol + 2  
drops of conc. H<sub>2</sub>SO<sub>4</sub>  
Keep it in a  
boiling water bath of  
5 minutes & cool pour  
into Na<sub>2</sub>CO<sub>3</sub>

A fruity  
smell

The given organic  
compound  
contains  
carboxylic  
group.

### 1] Test for amino group (-NH<sub>2</sub>) group.

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| Experiment                                                                                                                                                                                                                           | Observation                                       | Inference                                                                                                       |
|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---------------------------------------------------|-----------------------------------------------------------------------------------------------------------------|
| <p>a) Carbylamine test<br/>organic compound<br/>in a test tube +<br/>a few drops of<br/>chloroform + alcoholic<br/>potash - warm</p>                                                                                                 | <p>An intolerable<br/>Smell of<br/>isocyanide</p> | <p>The given organic<br/>compound is<br/>a primary amine<br/>&amp; confirms the<br/>presence of amino group</p> |
| <p>b) Diazotisation test<br/>for aromatic primary<br/>amine<br/>organic compound<br/>dissolved in dil HCl<br/>&amp; cooled in ice + ice<br/>cooled NaNO<sub>2</sub> solution<br/>+ ice cooled solution of<br/>β naphthol in NaOH</p> | <p>A red or<br/>orange dye<br/>is obtained</p>    | <p>Primary<br/>aromatic<br/>amine<br/>group is<br/>present.</p>                                                 |