



# R. K. GROUP OF COLLEGE

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**CHEMISTRY**





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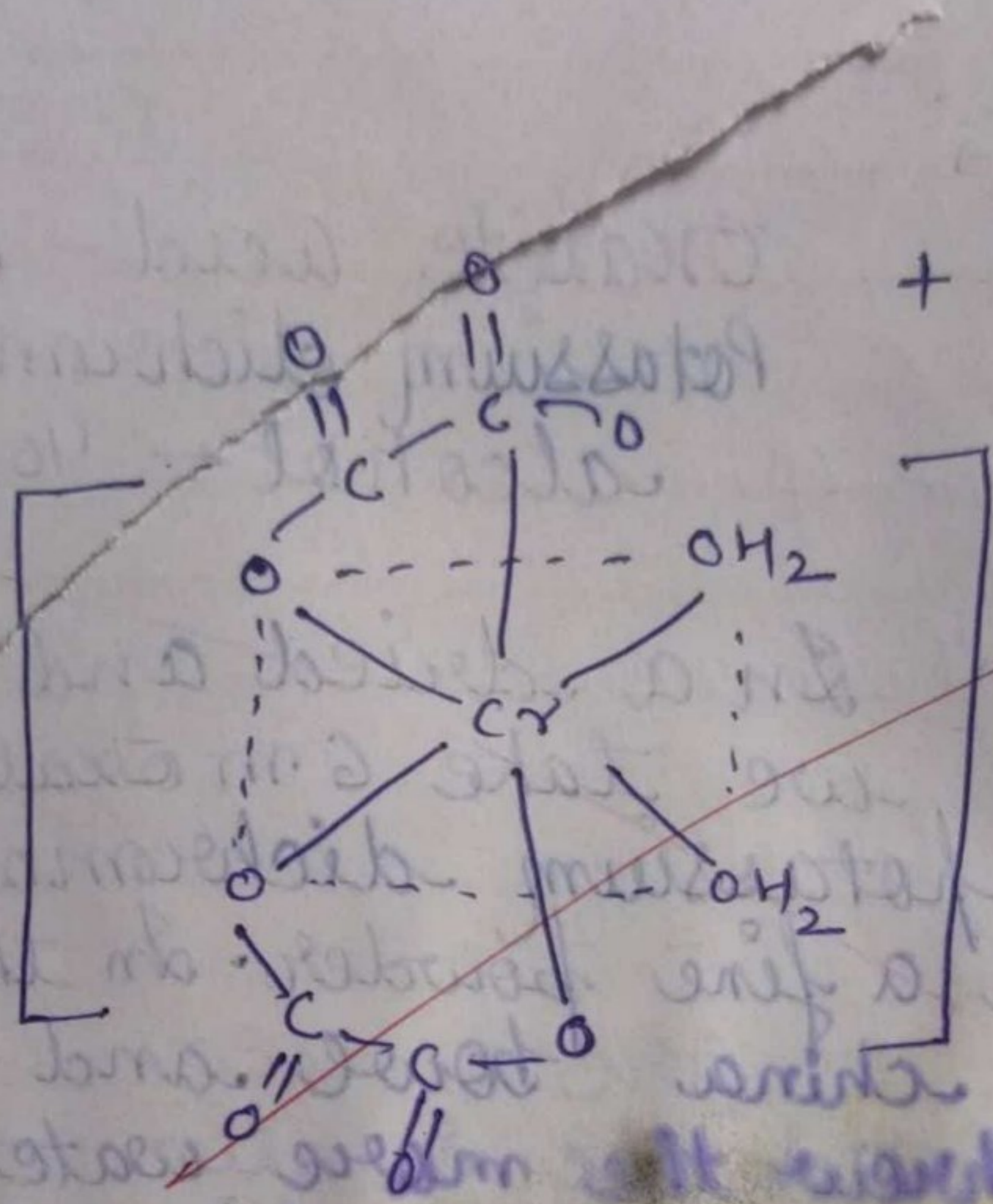
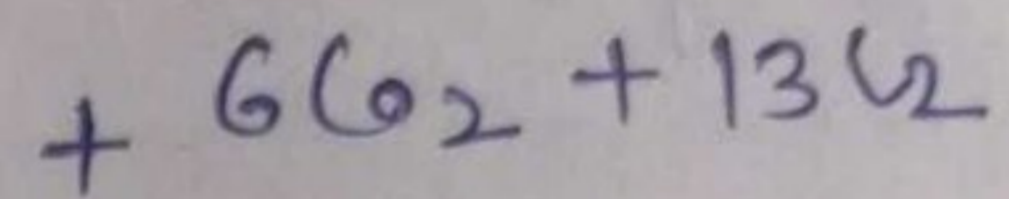
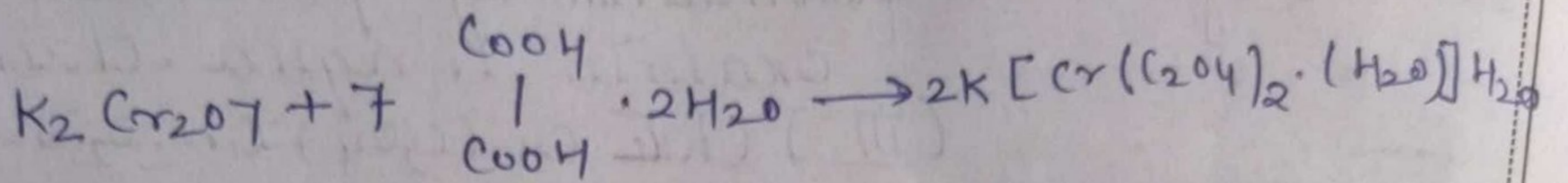
Object:- formation of potassium di-oxalato di aqua chromate (III)  $(K_2Cr(C_2O_4)_2 \cdot 2H_2O)$ .

Reagent:->

Oxalite acid - 6gm  
Potassium dichromate - 2gm  
alcohol - 40g/ml.

Method:-> In a dried and clean beaker we take 6m oxalate and 2gm potassium dichromate and crushed to make a fine powder. In this powder take a china bowl and wet this and throw the more water. now this water cover with watch glass and heat it with low. when it substance heat then reaction is fastly. and with high smell the  $CO_2$  gas removed. and this mixture converted into liquid dark colour. and this put out to burner and mix. 20ml alcohol the Spetulla then to move, it makes a solid substance. if solid substances are not formed then this maked liquid filtered and remaining solution put 20ml alcohol and mixed it in Spetulla. For this substance make solid. and this moved and heat it.







In which crystals are formed. This crystals are dark pink black coloured.

Result :-

By above method the potassium oxalato diaqua chromate (III) obtained.

→ This is black coloured solution.

3.2 gm produced

Precautions :- Beaker and flask both are dried and clean.

→ and heated the solution of china dish slowly, by which it does not blast.

→ In hot solution the alcohol mixed time the burner should be closed by which there is no risk of burn.

Sunand  
12/04/2021



Object: →

Opposition potassium di-oxalato aqua chromate (III)  
 $K_2[Cr(C_2O_4)_2(H_2O)_2] \cdot 2H_2O$  formation.

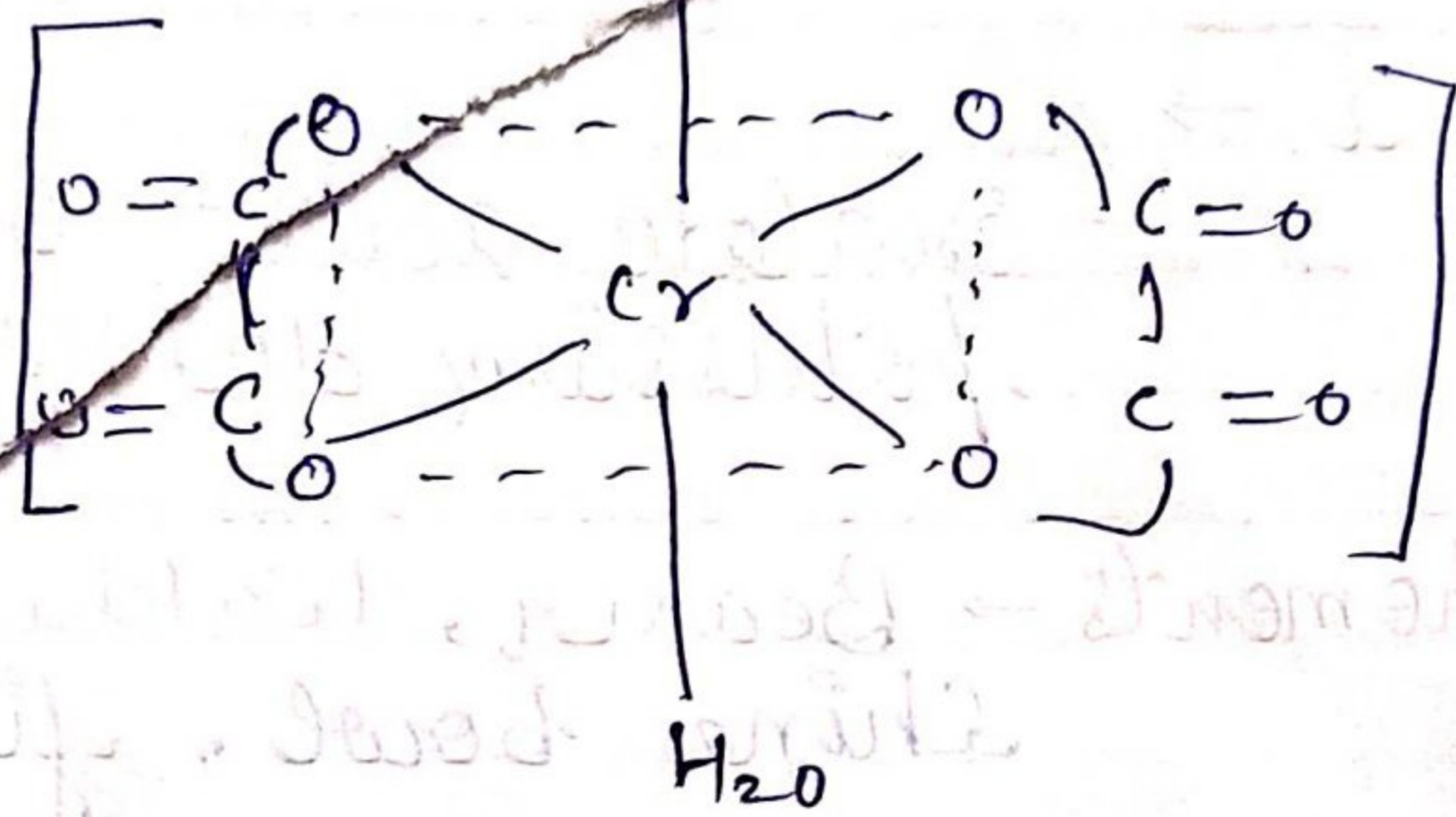
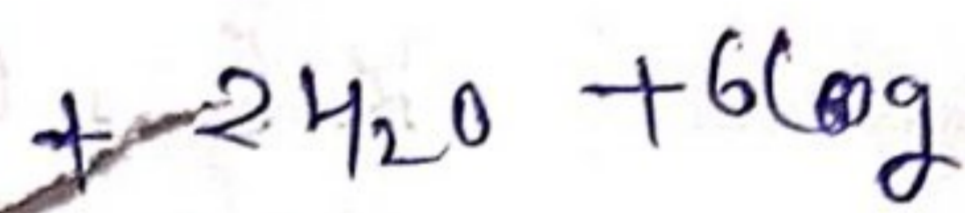
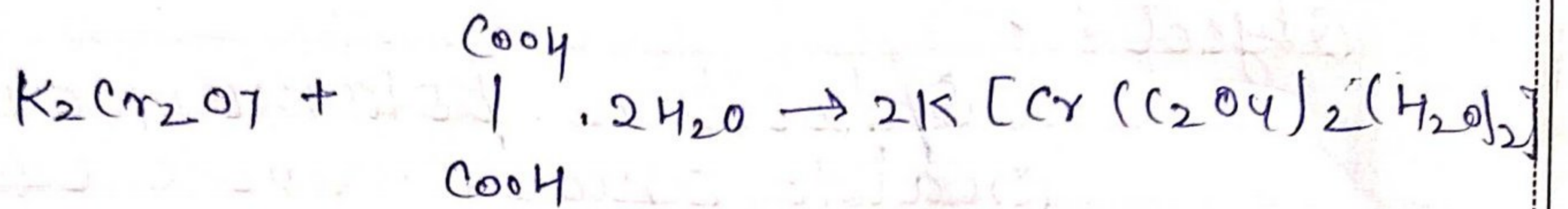
Reagent →

Oxalate acid - 6gm  
potassium dichromate - 2gm

Equipments → Beaker, watch glass,  
China bowl, filtered pipe.

Method: → In a 250gm beaker mixed 6gm Oxalic acid and mix it with lower amount and heat it. and by this method in filtered tube we take 2gm potassium dichromate put in and heat it. now this solution put it in beaker and cover with watch glass. because reaction is highly. now both solution then mixed and make dark colour liquid. and put it on china bowl then in one third heat it. and cold it on room temperature. and crystals are formed in china bowl. and wash this crystals with cold water. and in last alcohol







washed and crystals are formed of pink colour.

Result: → above method the opposite potassium di oxalate di-aqua chromate (III) produced which colour is pink and light brown.

Wt: → 5.3 gm

Precaution: → beaker is cleaned and dried.

→ china bowl substance heated with slow stir.



Object:  $\rightarrow$  tetra-ammine copper sulphate  
 $[\text{Cu}(\text{NH}_3)_4\text{SO}_4]$  formed.

Necessary Equipment  $\rightarrow$

- (1) copper sulphate - 5 gm
- (2) ammonia beaker - 10 ml
- (3) alcohol - 40 ml
- (4) dilute  $\text{H}_2\text{SO}_4$  - 1-2 drops

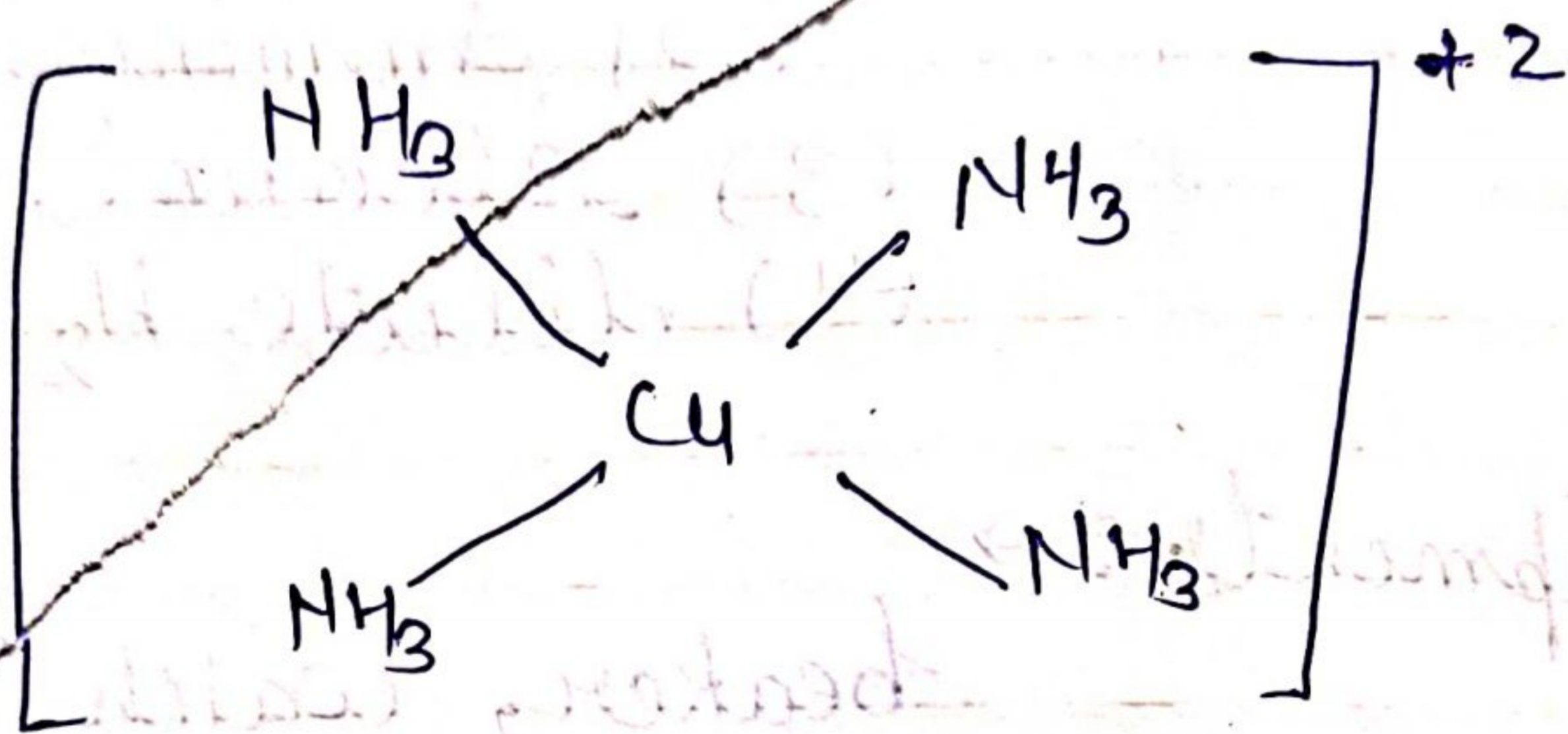
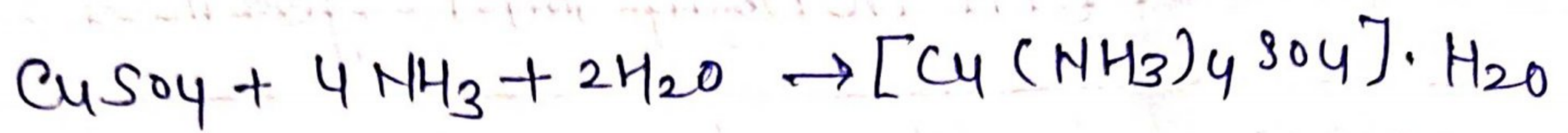
Equipments:  $\rightarrow$

beaker, watch glass.

Method:  $\rightarrow$  3 gm copper sulphate powder  
take in beaker and dissolved  
in low amount of water. and  
some drops mixed of  $\text{H}_2\text{SO}_4$ .

- $\rightarrow$  Now, in this  $\text{NH}_3$  solution ( $1:1$ ) moved  
regularly because cupric hydroxide  
blue the solution becomes dark blue  
coloured. and smell of  $\text{NH}_3$  comes.
- $\rightarrow$  Now, this blue colour solution moved  
regularly and put in beaker. after  
this heat for 10-15 minute on  $60^\circ\text{C}$ -  
 $70^\circ\text{C}$ . beaker cool and in this needle  
shaped crystals are separated.
- $\rightarrow$  and dried in the filter paper  
and separated.





tetraammine Copper (II)



Result: →

By above method diamine copper sulphate (II) obtained: which colour is blue.

Yield: → 4 gm.

Precaution: → beaker should be clean and dried.



Object :- nickel dimethylglyoxime  
Complex  $[Ni(DMG)_2]$

Necessary Equipment :-

- (i) Nickel Chloride - 6 gm
- (ii) Di-methylglyoxime - 10 ml
- (iii) Ammonia solution - 1:1

Equipments :- Beaker, watch glass

Method :-

One 400 ml. Beaker given nickel sulphate 6 gm Heat at  $72-80^\circ C$  and this solution put in 42-50 ml. Di-methylglyoxime. After this in solution. 1:1 ammonium solution drop by drop in watch glass. In Beaker the water. Heat at  $20-30^\circ C$  the down dropped read clear sum Di-methylglyoxime reagent mixed and test it. that the solution is complete or not. Beaker put for Half-Hour. this is separated by funnel and this is washed with water. two three time. and in last the solution of washed 5 ml. alcohol







And dried - it.

Result: -

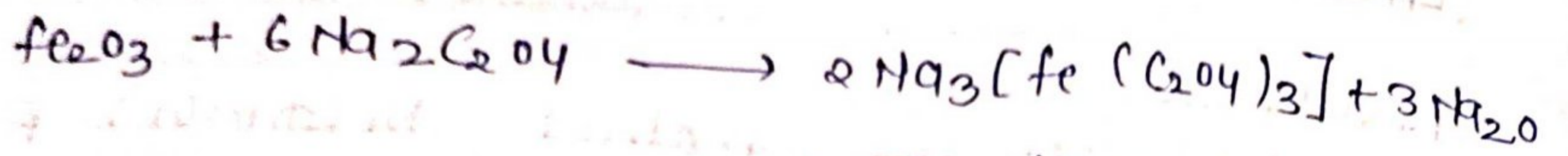
above method Ni (OH)<sub>2</sub> prod  
used Red Colour Solution.

~~gained :-~~

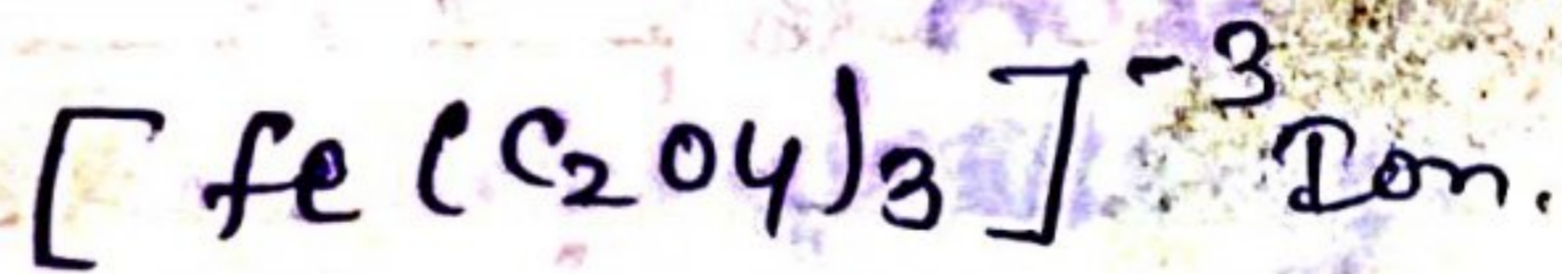
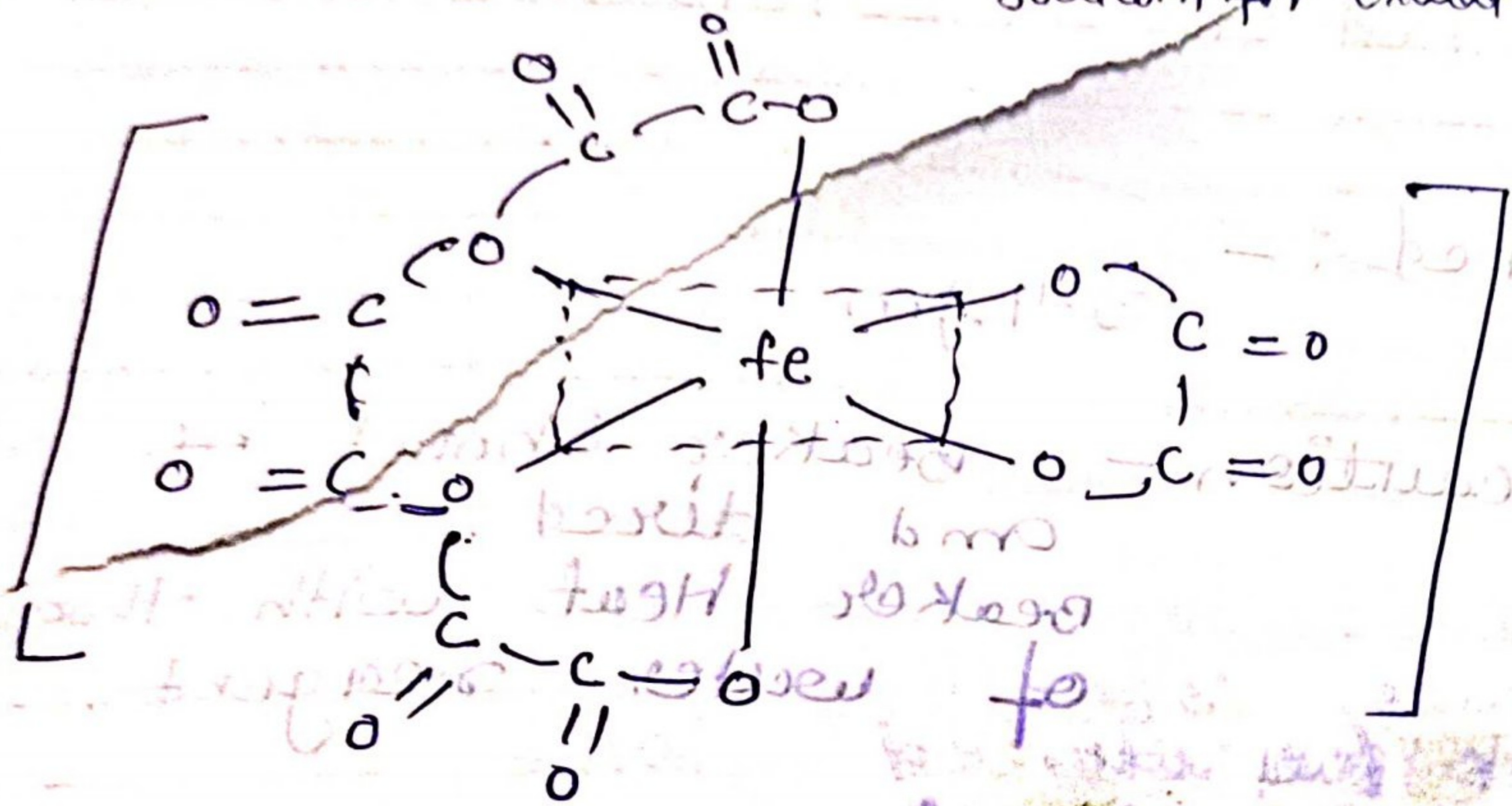
3-4 gm

precaution - beaker should be clean  
and dried.  
beaker Heat with the help  
of water reagent.





Sodium ferrioxalate





Object:- Sodium dioxaferit (III)  
 $\text{Na}_3[\text{Fe}(\text{C}_2\text{O}_4)_3]$  formation

Necessary equipment :-

- ① ferric chloride 10 gm
- ② NaOH - 12 gm
- ③ oxalic acid - 12 gm

Necessary equipment :- Beaker or watch glass.

Method:-> In a beaker 10 gm ferric chloride 10 ml water 3 gm NaOH 5 ml  $\text{H}_2\text{O}$  mixed now ferric chloride solution mixed in few amount of NaOH solution then ferric hydroxide is produce. Now one another Beaker mixed 12 gm oxalic acid in this solution. Now mixed 9 gm NaOH oxalate solution produce. Now with the help of glass rod the water solution mixed in ferric hydroxide solution and heat it. Now this solution filtered and cooled and filtered. the filtered cooled water and collected put for 2-3 days.







Result :-

Above method the crystal of Saccharindioxiuloferat green colour produce.

Yield :- 15 gm

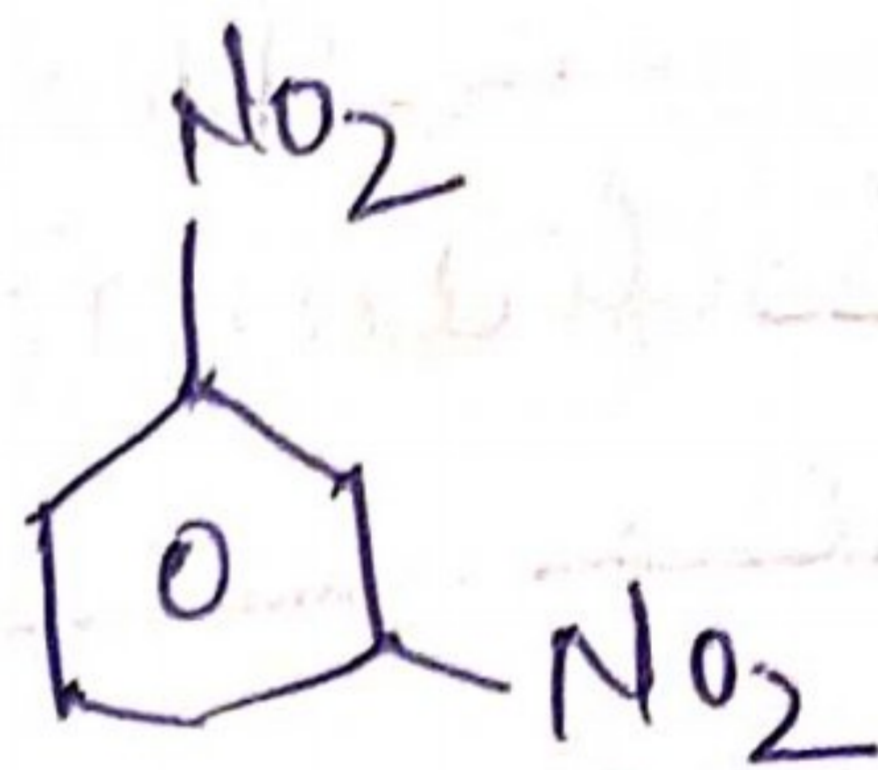
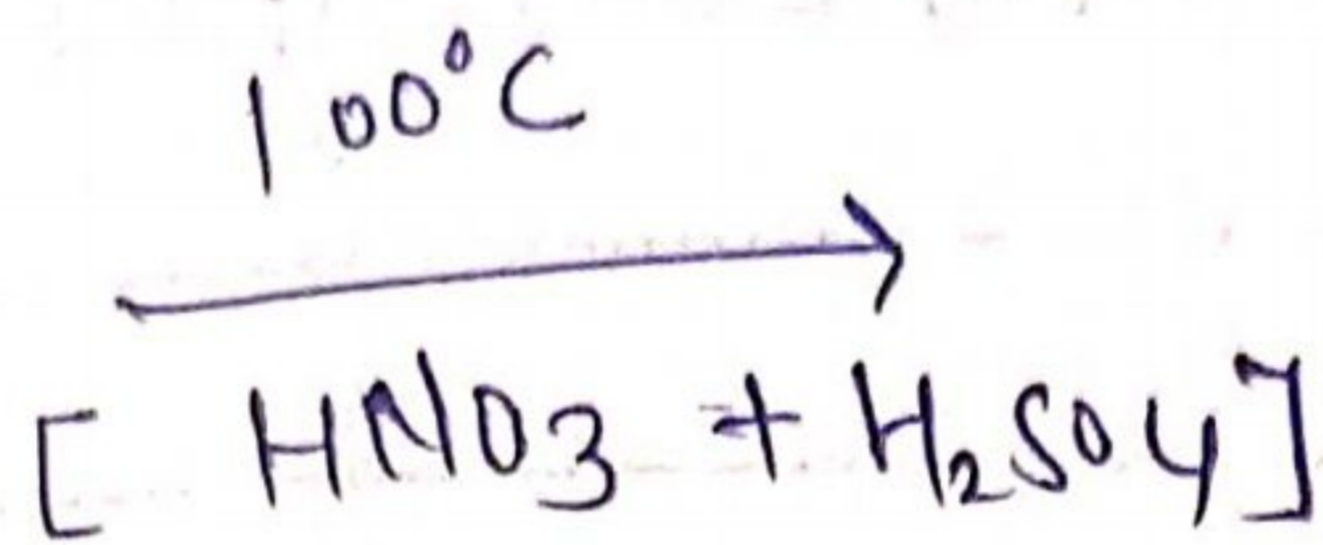
precaution :-

The Beaker should be cleaned properly.





[Nitro Benzen]



[m-Dinitro Benzen]



Object :-

preparation of benzaldehyde from aniline.

Necessary reagent :-

- ① aniline - 5 ml
- ② 10% NaOH solution 15 ml
- ③ 100 ml Benzoyl chloride

Equipment :- 250 ml conical flask, Beaker keep.

Method :-

250 ml conical flask mixed 5 ml aniline and 15 ml 10% NaOH solution and 100 ml Benzoyl chloride. mixed on the flask the clock move it for 15-20 min. Benzaldehyde washed with distilled water and dried the product. after it heat  $100^{\circ}\text{C}$  and mixed alcohol crystals. By this method pure benzaldehyde be separated.

Result :- By this method Benzaldehyde produced.

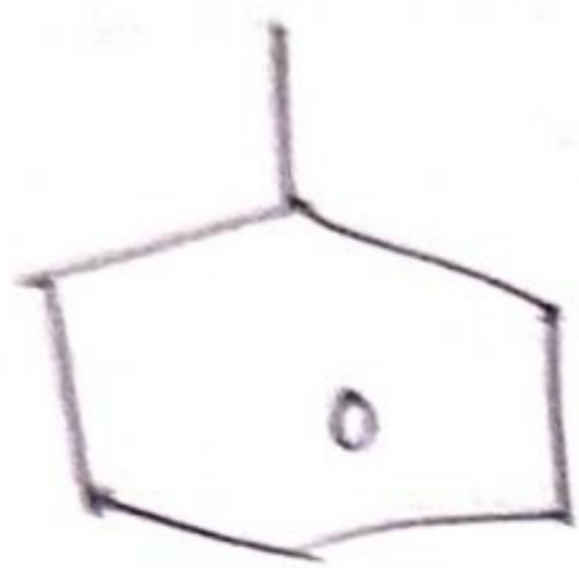
physical state :- white crystal solid substance

yield :- 8 gm

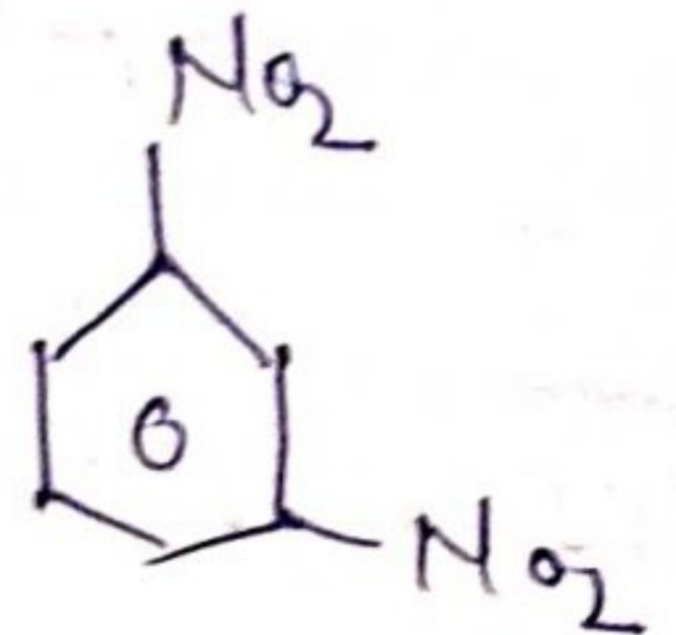
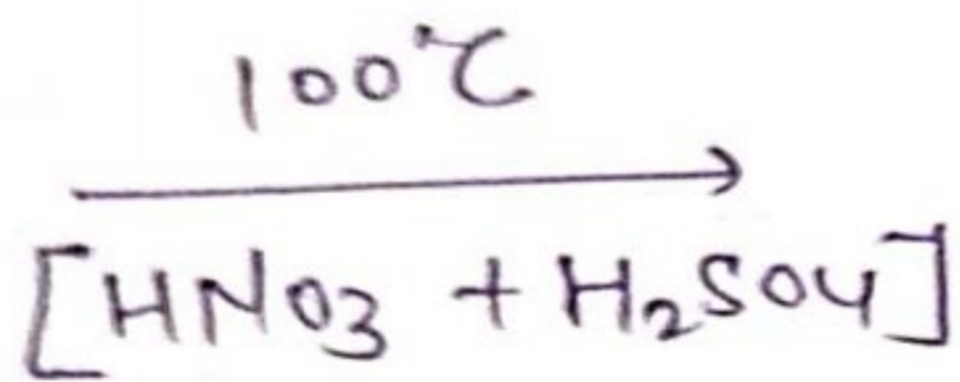
melting point  $\sim 165^{\circ}\text{C}$

Teacher's Signature .....





[Nitro-Benzene]



[m-Dinitro-Benzene]



Object:- preparation of m-dinitro benzene from Nitrobenzene

Useful Reagent:-

Nitro-Benzene 5ml  
 Conc  $H_2SO_4$  - 15ml  
 Conc  $HNO_3$  - 15ml  
 Alcohol

Equipment :- 10ml Round steel flask  
 glass air mixed and  
 250ml beaker

Method :- In 100ml round flask shaped beaker 15ml Conc  $HNO_3$  in this 15ml Conc  $H_2SO_4$  mixed and moved and the flask be cold in this nitration mixture the 5ml Nitrobenzene mixed in flask. after this in flask 4-5 small fresh of water available mixed in 1 hour. Now flask mixture be cold and one to 250ml beaker put sum ice. m-dinitrobenzene is a yellow solid. and it dried with the help of filter paper and test melting-point



Result:- By above method M-Dinitro  
Benzene produced.

gained:- 6.0 gm

M.p :- 90°C







Object :- preparation of iodoform from acetone

Useful Reagent :- ① acetone - 20ml  
 ② KI - 6gm - 10ml  
 ③ 5% Na-Hypochlorite - Solid

Equipment :- 250ml flask, keep

Method :- In a 250ml flask take 20ml acetone, 6gm potassium Iodide put and it mixed in 80-100ml water. Now this mixed continue and 5% Na-Hypochlorite - solution of Iodoform on this mixed iodoform make yellow crystal. in which Hypochlorite solution mix and iodoform yellow crystal be separated the flask put for 15-20 mint. Now iodoform be separated by funnel and washed with water and dried with filter paper this solid yellow substance crystallized by ethanol methylated spirit.

Result :- About reagent make formation of Iodoform.

physical state :- solid yellow colour solid crystal.

gained :- 3-4gm

M.P  $\rightarrow$  120-122  $^{\circ}$ C



Object:- preparation of methyl orange from Sulphonic acid.

Useful Reagent:-

- ① Sulphonic acid - 5 gm
- ② dehydrated Na<sup>+</sup>
- ③ Sodium nitrite - 2 gm
- ④ dimethyl aniline - 3 ml
- ⑤ Conc - HCl
- ⑥ Dil - HCl - 15 ml
- ⑦ Sodium chloride
- ⑧ 20% NaOH solution
- ⑨ ice
- ⑩ useful

Useful equipment :-

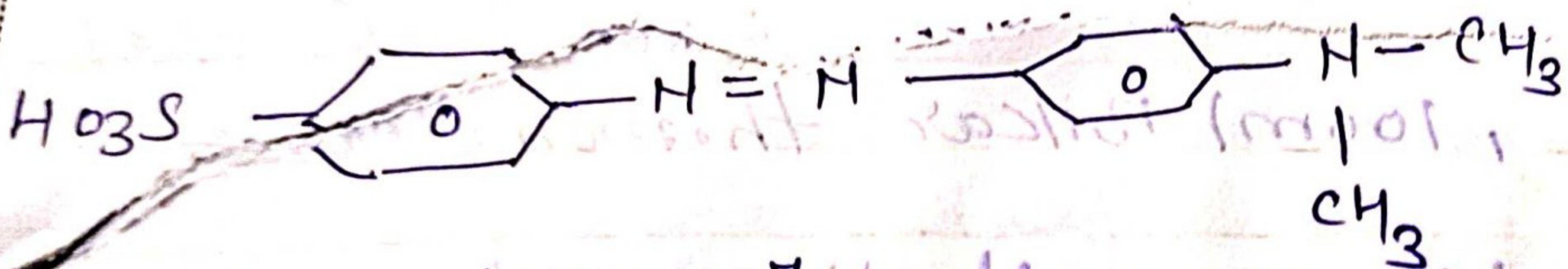
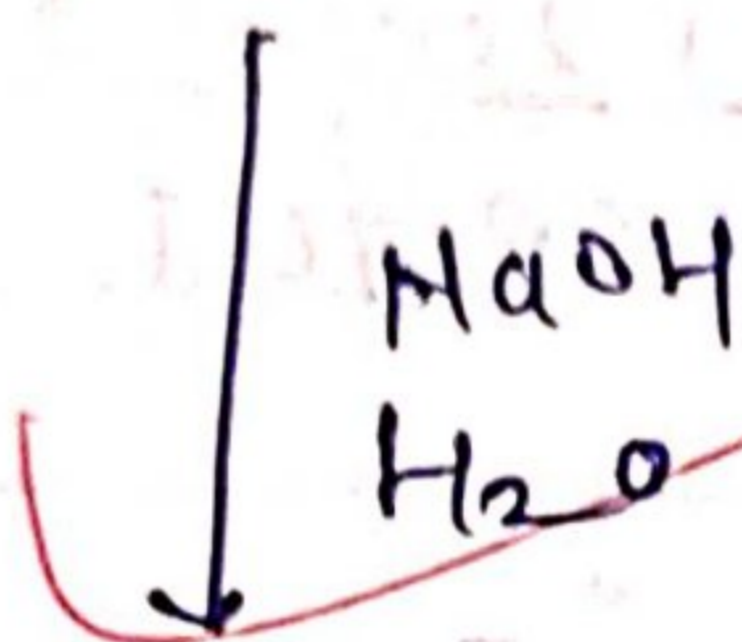
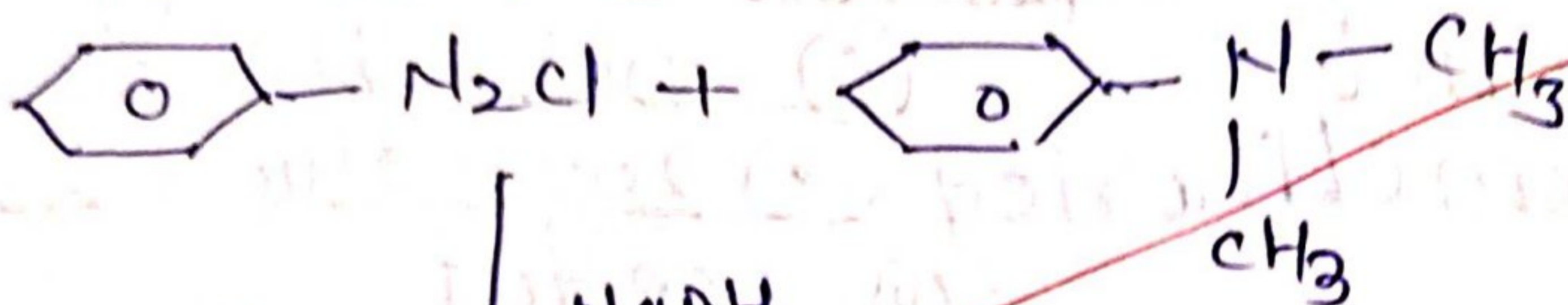
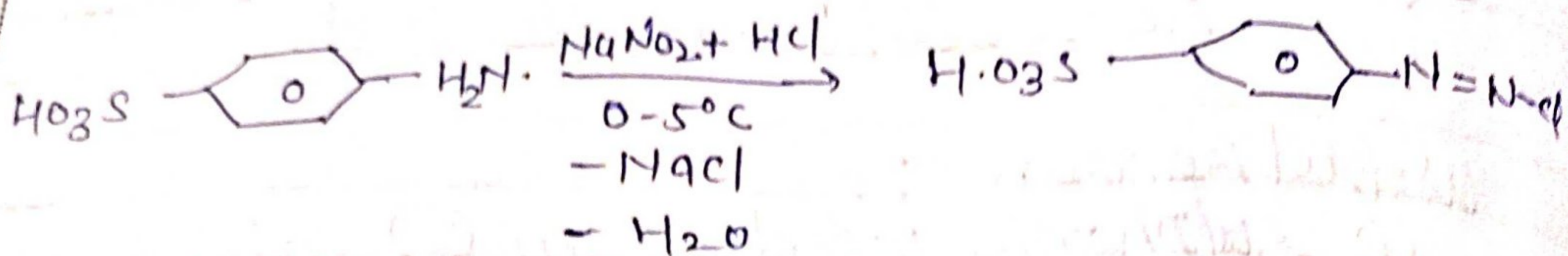
200 ml Conical flask, 100 ml Bicker thermometer

Method:- 5 gm methyl orange and 2 gm D-Hydroed NaOH put in flask and make a solution in 5 ml water, in this 2 gm Na nitrite and 10 ml solution in water. By this method solution produced at 25°C cold and put in flask in this 8 ml dil HCl mixed stalic. the yellow color temperature 10°C. Not be increased.

- One another Bicker 3 ml dimethyl aniline and 15 ml conc HCl solution and cobalt

- This solution the above mixed movement is stalic and mixture moved four 10 ml





[methylorange]



In this 25 ml 20% NaOH solution 6 gm NaCl mixed and moved, solid solution filter and wash with cool water this solid substance on crystallization the hot water produced orange color crystals produced which are filter and dry with the help of filter paper by which methel orange is a salt and this mp is not confirmed

Result:- By above method methel orange produced.

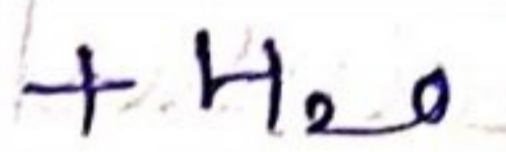
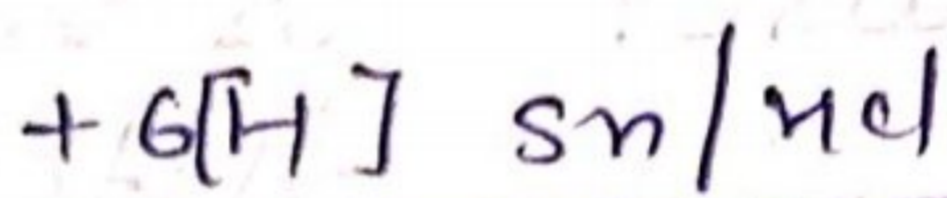
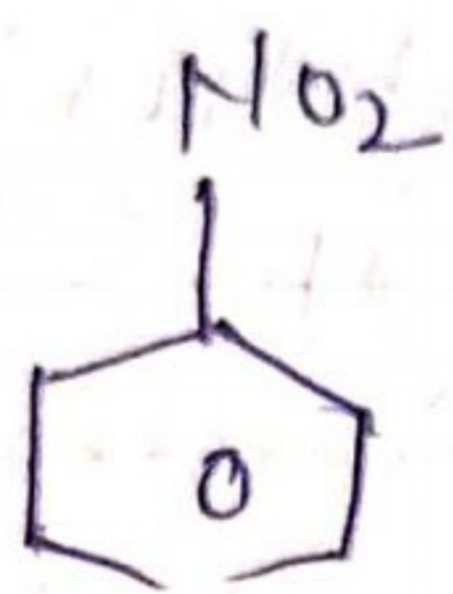
physical state :->

Dark orange red color crystal

gained :- 6.0 gm.



Redn:



[Nitrobenzene]

Aniline



Object :- preparation of Aniline from Nitrobenzene.

Useful Reagent :- (i) Nitrobenzene - 5 ml

(ii) Tartine 10 gm

(iii) Conc-HCl - 20 ml

(iv) Na Hydroxide - 15 gm

Equipment :- Round Conical flask 250 ml  
point kep watear and other  
necessary equipment.

Method :-

In a flask 5 ml Nitrobenzene and 10 gm ~~to~~ Crystal tin took. Now in this move with fastley sum 20 ml Conc-HCl mixed and cool it abter this mixture this Heat with one Hotte. it H<sub>2</sub>Had while the smail of NitroBenzene coming. Now in this NaOH solution and cool-it and mixed wateer move it by which the aniline droplets Be sepreted the point sepreted by keep or Aniline also sepreted.

Result :-

above method Aniline produced.

physical state :-

Colourless liquid.

gained :-

0.4 ml

M.p :- 183 °C



Object :->

In given organic mixture separated the organic compounds and identified the compound A and B and derived.

1. Starting Test :-

Physical state - Base  
 Colour - colourless  
 Smell - odourless smell  
 Solubility - insoluble in water.

Testing Table :- Test for compound 'A'

S.No	Experiment	Test	Result
2.	Flame Test :- To burn the take small amount of substance	It burns with smog flame.	This is aromatic compound.
3.	Litmus Test :- Component + red litmus paper	no change	Compound is indifferent.

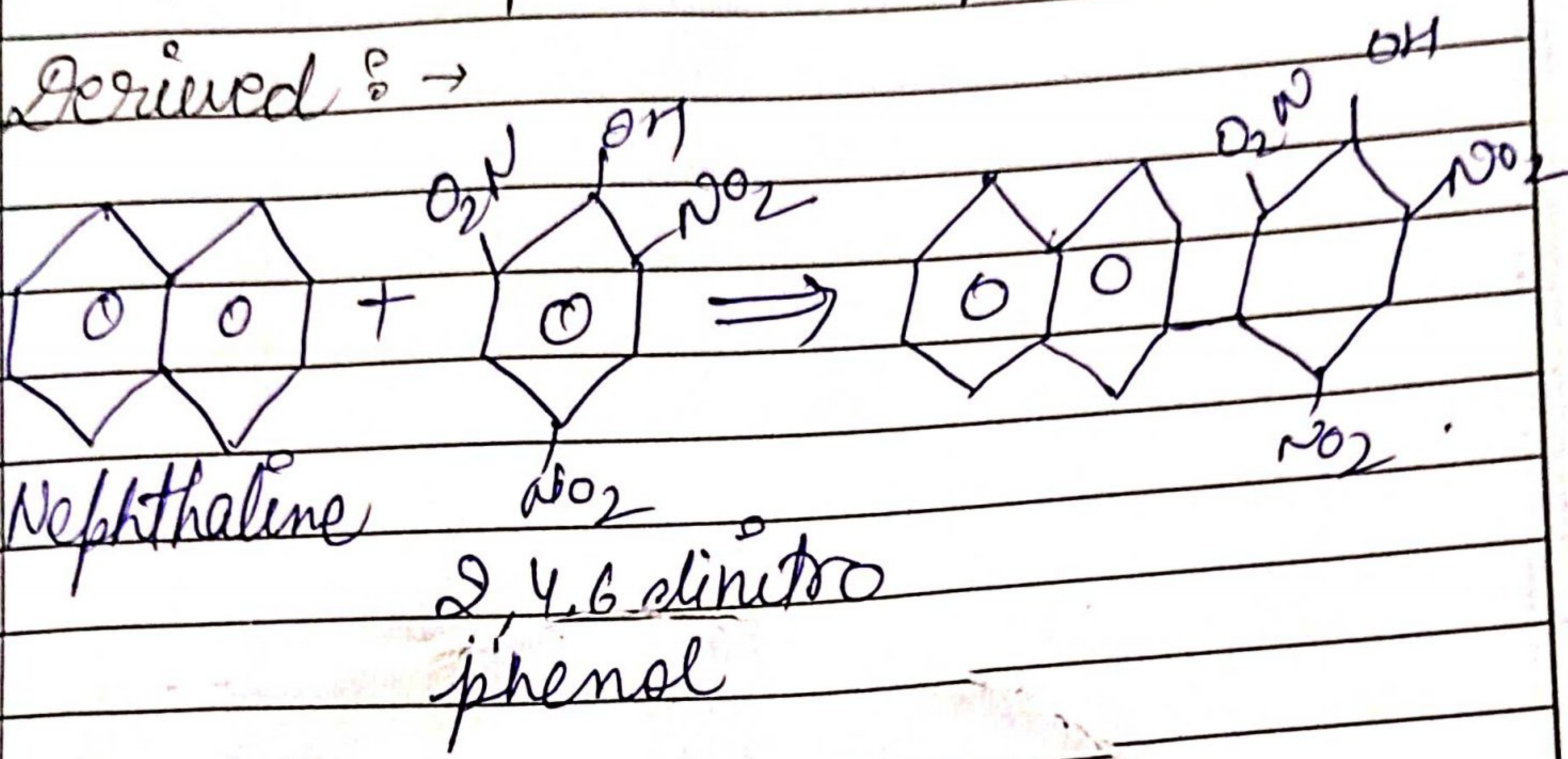


4.	Element Test:-		
(i)	Red acetate benzene solution + $(CH_3COO)_2Pb$ . (lead acetate).	Black colour is not pro- duced.	Sulphur is absent.
(ii)	Sodium nitroide + benzene solution + $Na_2[Fe(CN)_2NO]$	Violet colour is not produced.	S- Absent
(C)	Halogen Test: - benzene solution + dil $HNO_3$ + and cool $HgNO_3$ .	Yellow & white precipitate is not prod.	X - Absent
5.	Functional group Test: - Compound + dehydrate $AlCl_3$ + dried $CHCl_3 + \Delta$ .	green yell colour obtained.	$C_{10}H_8$ may be Naphthalene.
6.	M.P. ....	79°C obtain	$C_{10}H_8$ maybe
7.	Special test of Compound: -		
(1)	Compound + $KMnO_4$ solid + $Con \cdot H_2SO_4$ + $\Delta$ + Resorcinol + $\Delta$ + $NaOH$ + sol <sup>n</sup> .	green precipitate produced	May be confirmed $C_{10}H_8$ .



(111)	Compound + $CHCl_3 + AlCl_3$ dehydrate	green colour present	$C_{10}H_8$ confirmed.
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Derived :-  $\rightarrow$



Test for compound 'B'.

1. Primary Test :-

- Physical state - Solid
- Smell - Smellless
- Colour - colourless
- Solubility - Soluble in water.

Testing table :-

S.No.	Experiment	Test	Result.
2.	Flame Test :- The small amount of	Organic comp. burn with smokeless flame	Organic compound is aliphatic

Subst. take in the test & heat it

Teacher's Signature .....



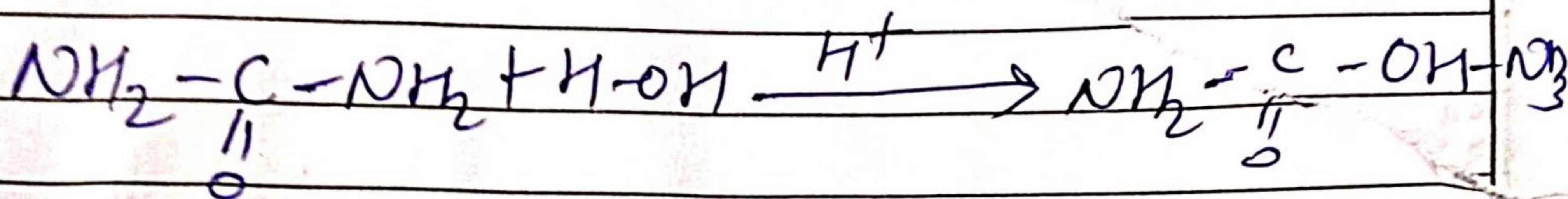
3.	Litmus Test → comp. + Red litmus in Blue litmus	No changes	Compound is Indiff- erent.
4.	Element Tests		
a)	N <sub>2</sub> Test → lamane sol. + FeSO <sub>4</sub> (Pure) + FeCl <sub>3</sub> + di Hcl.	produced blue colour solution	Nitrogen absent.
b)	Sulphur Test: - CH <sub>3</sub> COOH + lamane sol. + (CH <sub>3</sub> CO) <sub>2</sub> Pb	black preci- -pitate obtained.	Sulphur is absent
c)	Halogen Test: - lamane sol. + dil HNO <sub>3</sub> + Δ + cool + AgNO <sub>3</sub>	white & yellow preci. obtained.	
5.	Functional group Test: - Main subst. + dil HCl + NaN <sub>2</sub> on mixed solution	more smell produced.	$\text{NH}_2 - \underset{\text{O}}{\underset{ }{\text{C}}} - \text{NH}_2$ group.



6.	Melting Point	130°C	$\text{NH}_2 - \underset{\text{O}}{\underset{  }{\text{C}}} - \text{NH}_2$
7.	Compound Test		
	Compound + NaOH + Δ + dil CuSO <sub>4</sub>	purple colour	$\text{NH}_2 - \underset{\text{O}}{\underset{  }{\text{C}}} - \text{NH}_2$ confirmed.

Derivate of Urea:-

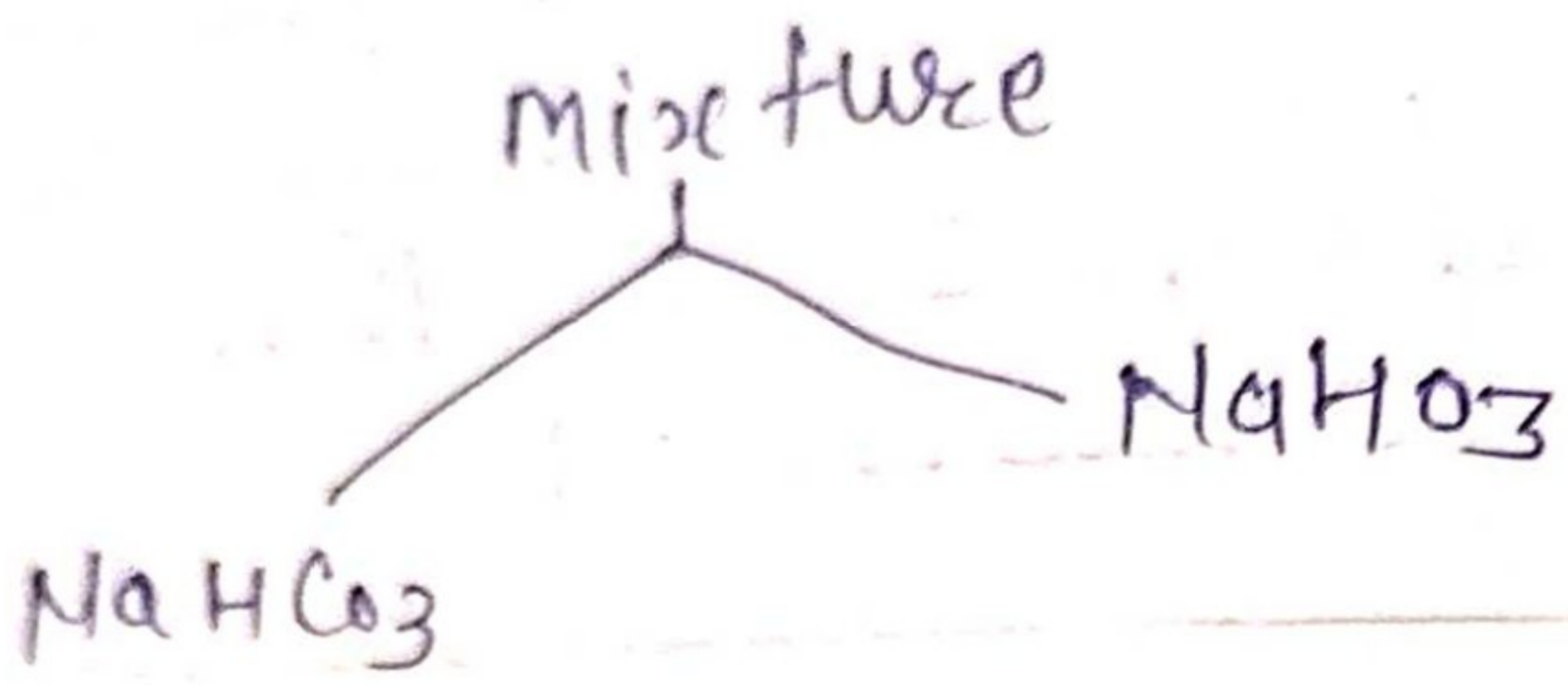
On water decomposition produced derivate:-



Result:-> In given organic mixture to separate organic compound the group A - Nephthaline B - Urea produced.

Melting Point:- 79°C and 13°C produced and derivate of Nephthaline is private.





[A]

[B]



Object: → In given organic mixture be separate the organic compound and identify the compound A and B and derivate one out of two.

Test for compound 'A'.

1. Primary State: -

Physical state - solid

Colour - colourless

Smell - odourless smell

Solubility → Insoluble in water

Table: -

S.No.	Experiment	Test	Result
1.	Flame Test: - Take compound on spatula and burn it.	compound burn with smoke flame.	Aromatic compound
2.	Litmus Test: - compound + Red / blue litmus paper	No Reaction	Compound is derived.

Teacher's Signature.....

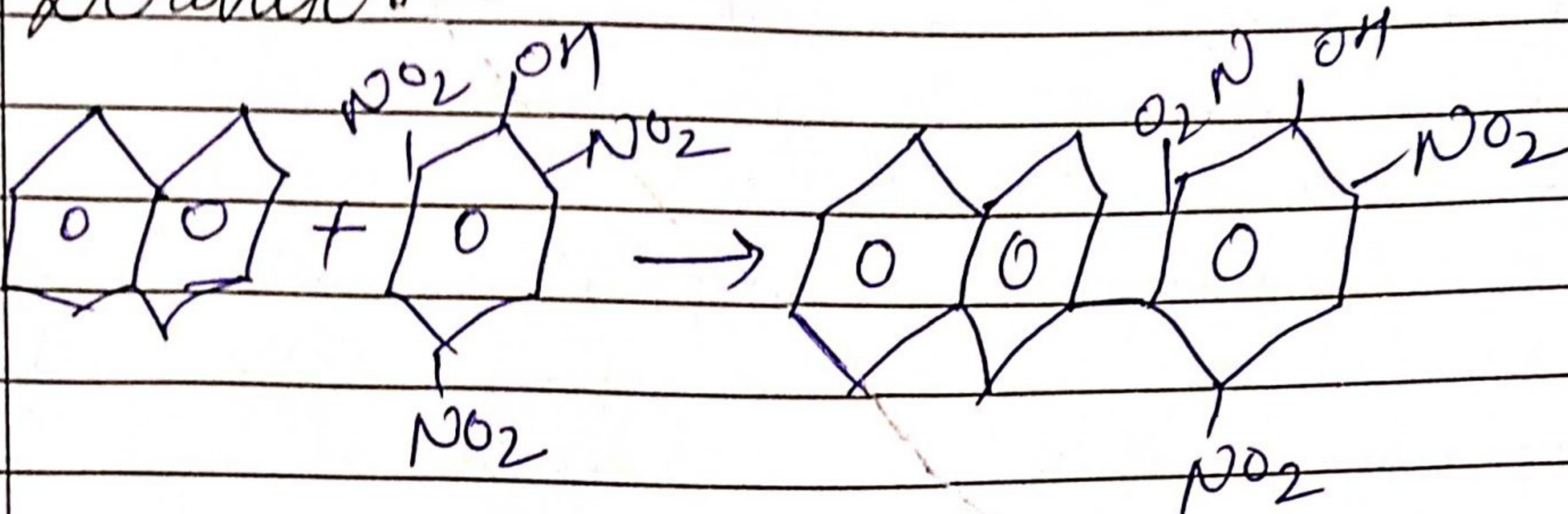


4.	Elementary Test		
(a)	$N_2$ -test + Lem -ane solution + $FeCl_3$ + dil HCl.	no Reaction	Nitrogen absent.
(b)	Sulphur test/ Lead acetate test $CH_3COOH$ + Lemane solution + $(CH_3COO)_2 Pb$	no Reaction	Sulphur absent.
(c)	Halogen Test? - Lemane sol. + dil $HNO_3$ + cool + $AgNO_3$	no Rx <sup>n</sup> .	X - Absent.
5.	Functional group Test :- Compound + dehy. $AlCl_3$ + $CHCl_3$ .	green/yell -one colour.	may be Nephthalin
6.	Special Test of compound :- Compound + $KMnO_4$ (solid) + con. $H_2SO_4$ + $\Delta$ + $NaOH$ solution.	green precipitate obtained.	May be $C_{10}H_8$ .



ii	Compound + $\text{CHCl}_3 + \text{FeCl}_3$	green colour produced.	$\text{C}_{10}\text{H}_8$ confirmed.
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8. Derivate :-  $\rightarrow$



Test for compound - "K"

1. Primary Test :-  $\rightarrow$

Physical state - Solid  
 Smell - odourless  
 colour - colourless  
 Solubility - soluble in  $\text{NaHCO}_3$ .

Table :-

S.No.	Flame Test :-	Test	Result.
	To burn Compound on Spetulla	Compound burn with smoke flame.	compound nature is aromatic.



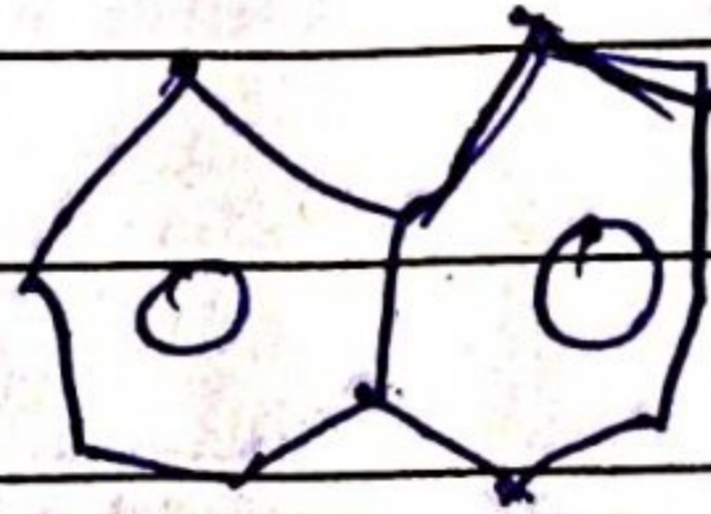
Litmus Test - Comp. + Red/Blue litmus paper.	Turns blue litmus into Red litmus.	may be carboxylic acid.
Elementary Test:-		
4. Nitrogen Test → Lamane sol. + pure FeSO <sub>4</sub> + FeCl <sub>3</sub> + dil HCl.	No Rx <sup>n</sup>	Nitrogen is absent.
b) Sulphur Test:- CH <sub>3</sub> COOH + Lamane solution + (CH <sub>3</sub> CO) <sub>2</sub> Pb.	No Rx <sup>n</sup> .	Sulphur absent.
c) Helogen Test:- Lamane sol. + dil HNO <sub>3</sub> + Cool it + mixed AgNO <sub>3</sub> .	No Rx <sup>n</sup> .	X - Absent.
5. Functional group Test:- Comp. + ethyl alcohol + Conc. H <sub>2</sub> SO <sub>4</sub> + Δ + Cool + H <sub>2</sub> O	Smell like fruits	may be carboxylic acid.



6.	Melting Point	159°C	
7.	Special test of compound →		
	Comp. + H <sub>2</sub> O + FeCl <sub>3</sub>	dark violet colour obt.	HOC <sub>6</sub> H <sub>4</sub> COOH may be.
→	Compound + CH <sub>3</sub> -OH + Conc. H <sub>2</sub> SO <sub>4</sub> + Δ + cooled + water.	Smell like wintergreen oil.	HOC <sub>6</sub> H <sub>4</sub> COOH is confirmed. (red colour).

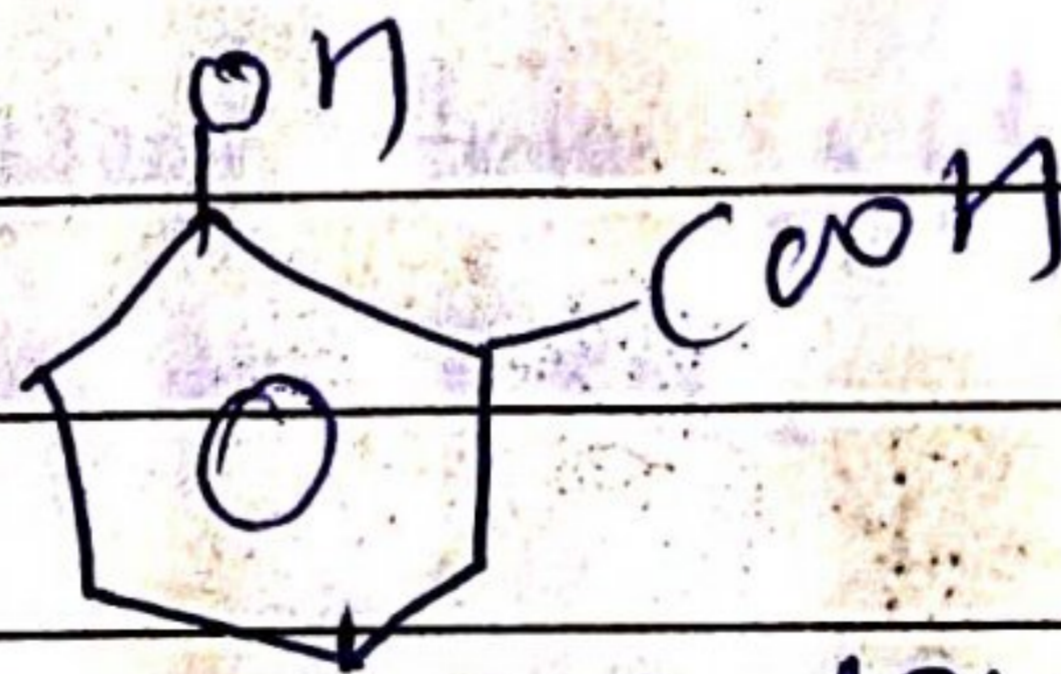
Result: → In given organic mixture the compound A and B is obtained.

Compound A →



Naphthalene

Compound B →



(Saliphatie acid).



Object? →

In given mixture separate organic compound and identify A and B compound and derivate one out of two compound - "A"

Primary test - physical state - Solid

Colour - Colourless

Smell - Smellless

Solubility - NaOH solubial

Test table:-

S.N.	Expt.	Test	Result
2.	Flame test		
	Comp. take in spe. tube and heat in fire.	flame with smoke.	Aromatic Compound.
3.	litmus test		
	Comp + Red/blue litmus paper.	Blue litmus change into Red.	Compound is Acidic.
4.	Element test		
	(a) Nitrogen test		
	Lamara salt + purple FeSO <sub>4</sub> + FeCl <sub>3</sub> salt + dil + HCl	Blue and green colour not obtain	N - Absent



Halogen test

Remana Sol<sup>n</sup> Dil + AgNO<sub>3</sub>  
+ Heat + Cool + AgNO<sub>3</sub>  
mixed

Whit / Yello  
precip. is Not  
obtain.

x<sup>o</sup> Absent

5. Functional group test

(1) Test of Alcohol

Alcoholic Solution +  
FeCl<sub>3</sub>

Violet Colour  
is produced.

OH group  
Present.

→ Comp + Hytamic acid  
+ Conc. H<sub>2</sub>SO<sub>4</sub>

green Colour  
produced

β-Nephtaline  
may be.

→ Comp + HNO<sub>3</sub> + A + dil  
H<sub>2</sub>SO<sub>4</sub>

green Colour  
produced

β-Nephtaline  
may be.

6. m.p

123° C

β-Nep. may be

7. Special test of Comp.

Comp + H<sub>2</sub>O + FeCl<sub>3</sub>

Comp + KOH + CHCl<sub>3</sub>

Comp + I<sub>2</sub> + NaOH + Sol<sup>n</sup>

milky / Whit

Blue Colour

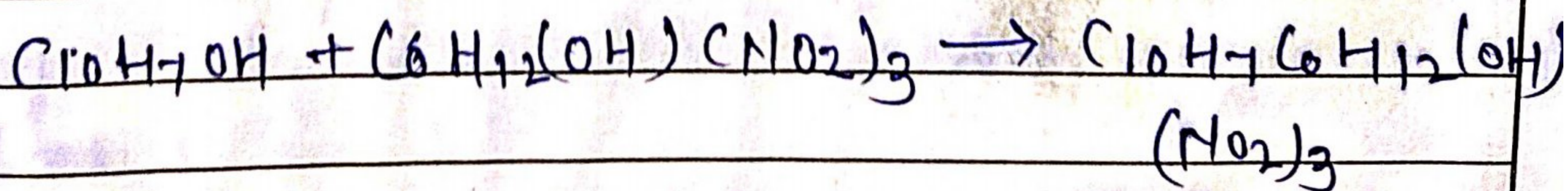
violet Colour

β-Nephtaline

is confirmed

β-Nephtalene

8. Derivate





Test Compound - "B"

1. primary test

physical state - Solid

Colour - Colourless

Smell - Alcoholic

Solubility - NaOH Soluble

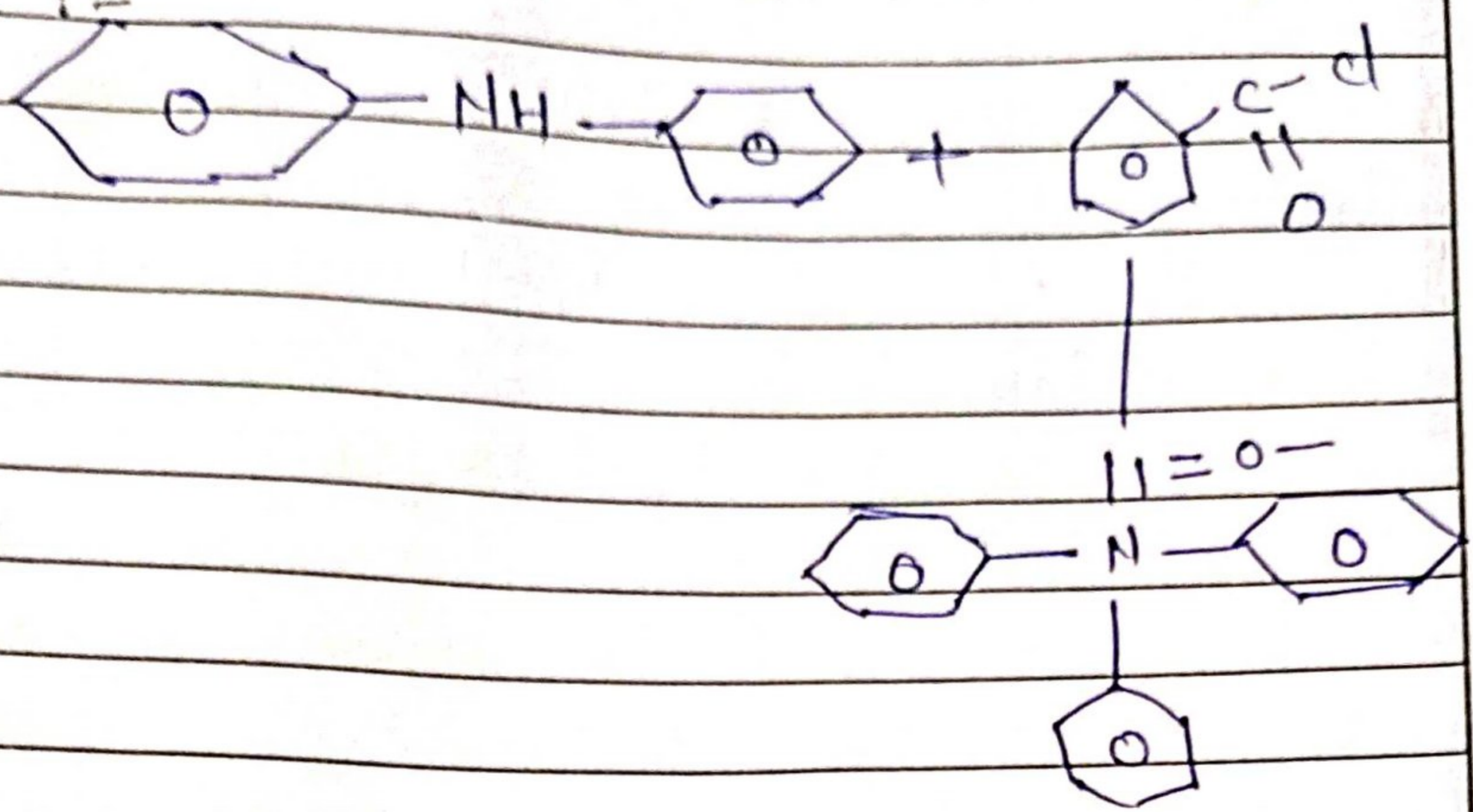
Table

S.No.	EXPT.	Test	Result.
	Comp. tests in Spectula and Heat.	Burnt with smoky flame.	Aromatic Comp.
3.	litmus test Comp + Red + Blue litmus paper	Red litmus Change in Blue.	Comp. is Basic
4.	Elementary test (a) N- test Lemana sol <sup>m</sup> + pale green + FeCl <sub>3</sub> + Dil HCl	Blue colour obtain	N- present
	(b) Sulphur test Lead acetate test CH <sub>3</sub> COOH + Lemana sol <sup>m</sup> CH <sub>3</sub> COOXPB	Black precip. Not obtain	S - Absent.
	M.P	536°C	
6.	Spe. test of compound Comp + CONC. H <sub>2</sub> SO <sub>4</sub> + Ag. NaNO <sub>2</sub>	Dark Blue colour sol <sup>m</sup> + is produced	Pi-phenyl Amine confirmed

Teacher's Signature.....

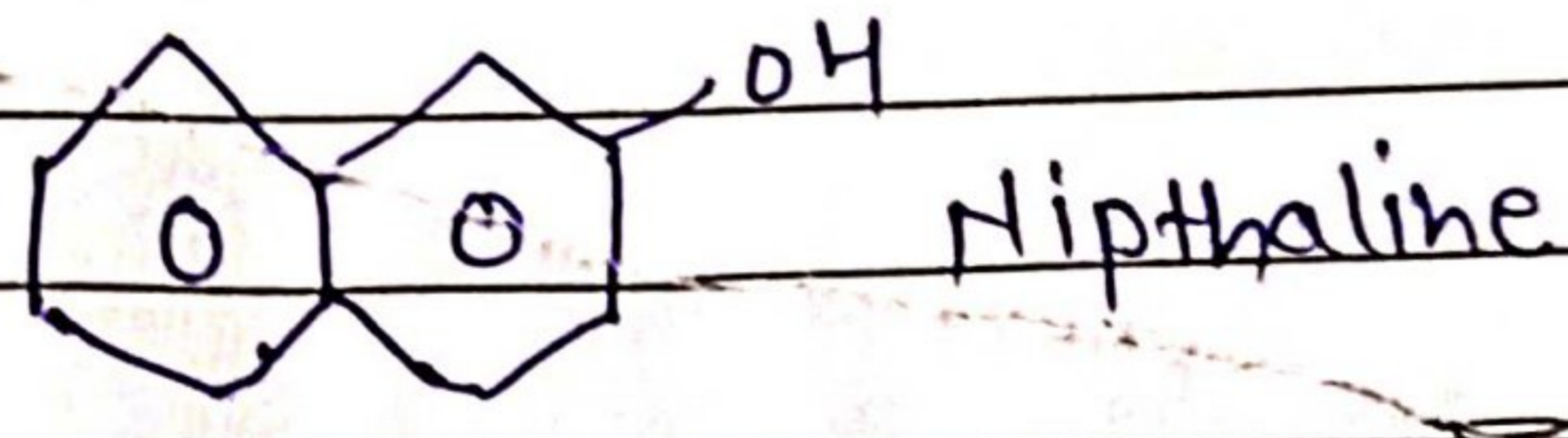


Derivate 1-

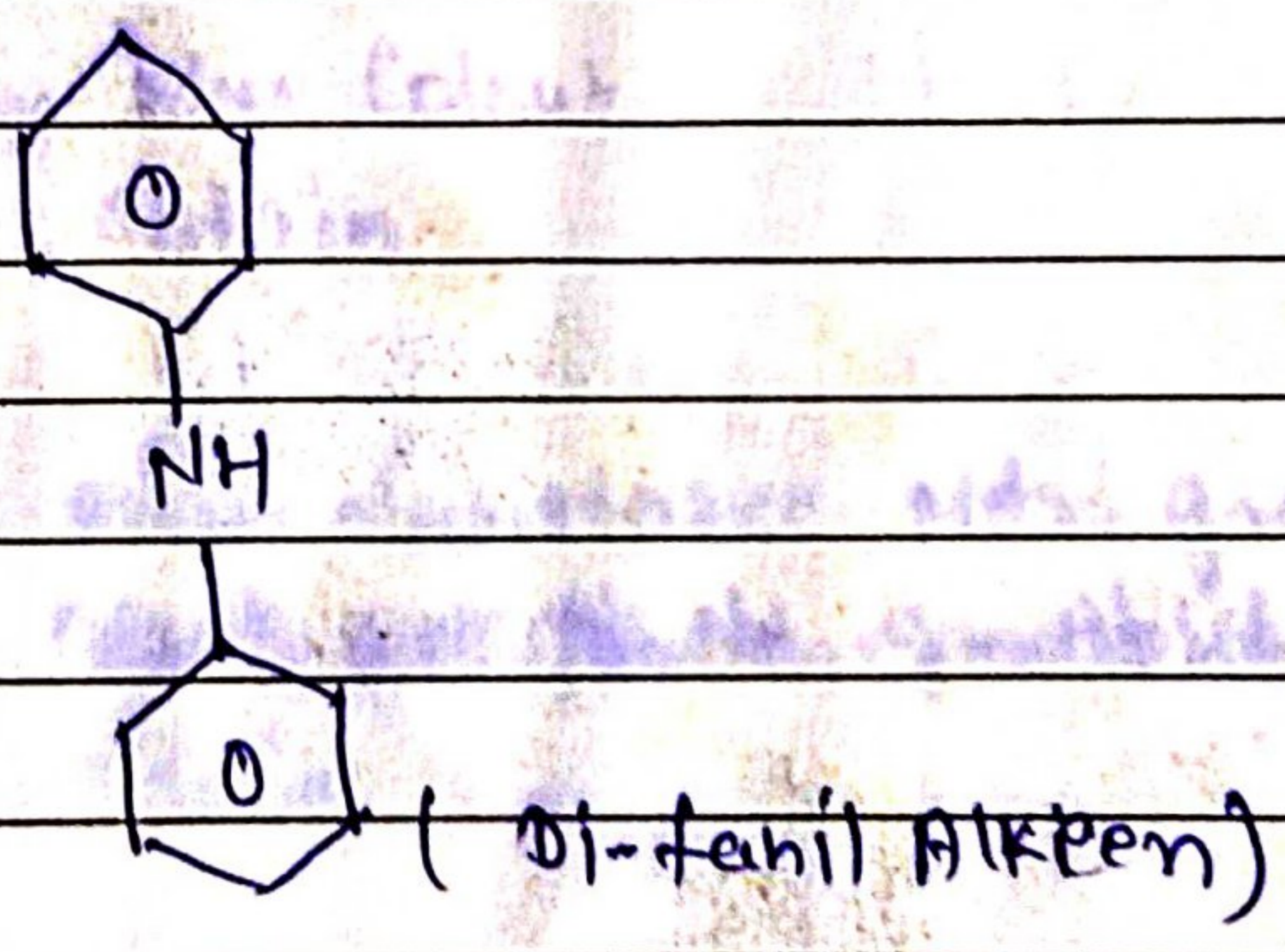


Result :-

Compound A -



Compound B -





Object :- In given mixture find organic compound.

Equipement :-

Test tube, Beker, Holder, Conical flask etc.

Comp "A" →

Colour - light yellow colour

Smell - Bitter Almond like sm

Nature - Derivate

flametest Burn with smoke so

it is aromatic com

Expto table

S.No.	Expt.	Test	Result
1.	Nitrogen Expt. Lamen Soln + Es peranto Burn FeSO <sub>4</sub> + NaOH + HCl	Persian blue Colour Not formed	N - Absent
2.	functional test Nitro compound test two ml Alcohol took. in compound + some drop of 10% CaCl <sub>2</sub> + Heat + Cool + Reagent	main substance is black / brown Colour	Nitro group is present.
3.	M.p.	114°C	m - Di-Nitro Benzen may be.

Teacher's Signature.....





organic Comp-test

→ Comp + NaOH + Heat      Red Brown Colour      m-Di Nitro  
 - Comp + Dil NaOH + Heat      produced      Benzene Comp  
 glucose      violet colour      m-Di Nitro Ben

Comp - B

primary stete

Colour - Colourless

Smell - Smellless

Nature - parvate

flame test - Burn Not with sm  
 So it is Aliphatic Co

Table Test

S.No	Expt.	Test	Result
1	Nitrogen group test ammonia soln + Esperan to burn + fesoy solubel + NaOH HCl	persian Blue Colour Not formed.	N - absent.
2.	functional test		
(a)	-COOH Group test l.s soln + Esperan Burn fesoy soln + NaOH + HCl S/. NaOH Co <sub>2</sub>	Cog gas Not Removed	-COOH group Absent.
(b)	final group test Comp. Caustic soln. + FeCl <sub>3</sub> include	Violet Colour not produced	OH Group Absent.

Teacher's Signature.....







Object: - In given mixture identified organic compound.

Equipment: -

Test tube, Bikaner, Holders, Conical flask etc.

Comp "A"

1. primary test: -

physical state - Solid  
 Colour - Cream Colour  
 Smell - Smell less  
 Nature - Acidic  
 flame test - fuming smoke  
 Burn so it is

Table Test

aromatic comp.

Expt. Test

Test

Result

N-Test

lemans sol<sup>n</sup> +

Blue/green

N - Absent

FeSO<sub>4</sub> + NaOH + CHCl<sub>3</sub> +

prec. produced

Conc. H<sub>2</sub>SO<sub>4</sub>

Not

2. functional Group test

Comp + NaHCO<sub>3</sub> water soln

Smell - COOH group  
 CO<sub>2</sub> gas removed are detectable

Astear test

Comp + Conc. H<sub>2</sub>SO<sub>4</sub> +

fruit like

COOH group

C<sub>2</sub>H<sub>5</sub>OH + A

Smell

Absent



m.p	133°C	Monamine Acid present
Acetic 2ml derivat Soln + 2ml residue KMnO <sub>4</sub> milking	Brown pre cipitate produced	Mepste Acid are present
- Comp. derivat Soln. KMnO <sub>4</sub> + Δ + filter cold	white kristal	menamic acid

Comp "B"

primary Expt -

Physical State - Solid

Colour - Colourless

Smell - like smell

Solubility -

Table Test

S.No	Expt.	Test	Result
1.	Element observation Lomana Soln + FeSO <sub>4</sub> + NaOH + FeCl <sub>3</sub> + Conc. H <sub>2</sub> SO <sub>4</sub>	Blue/green Colour not prod.	N-Absent.
2	functional test		HydroCarbon are present
(a)	Comp. + Conc. H <sub>2</sub> SO <sub>4</sub>	Not Soluble	

Teacher's Signature.....





b. Bayer Test  
 Comp + 2ml 2H<sub>2</sub>O + KMnO<sub>4</sub>  
 (Bayer Test)  
 are soln mixed.

3. m.p

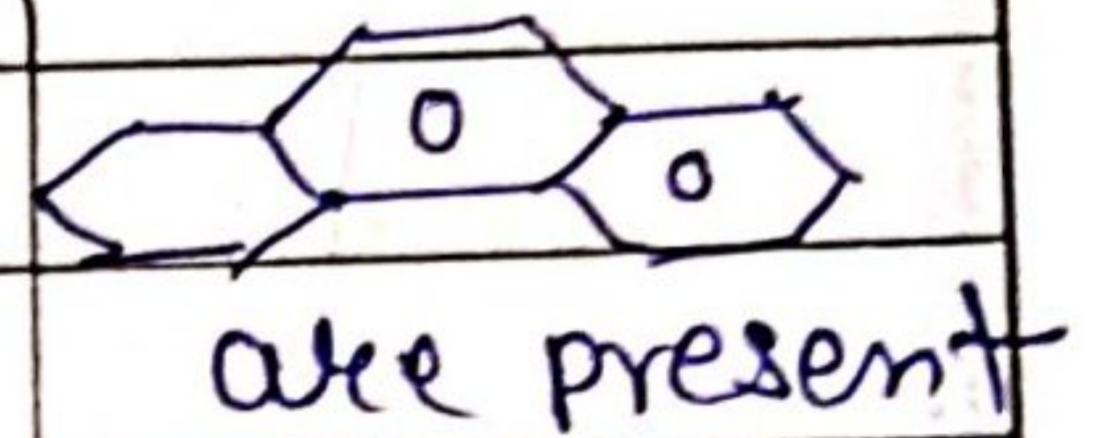
143°C



Special Test

Comp + Benzen are  
 mixed

Blue colour  
 are produced

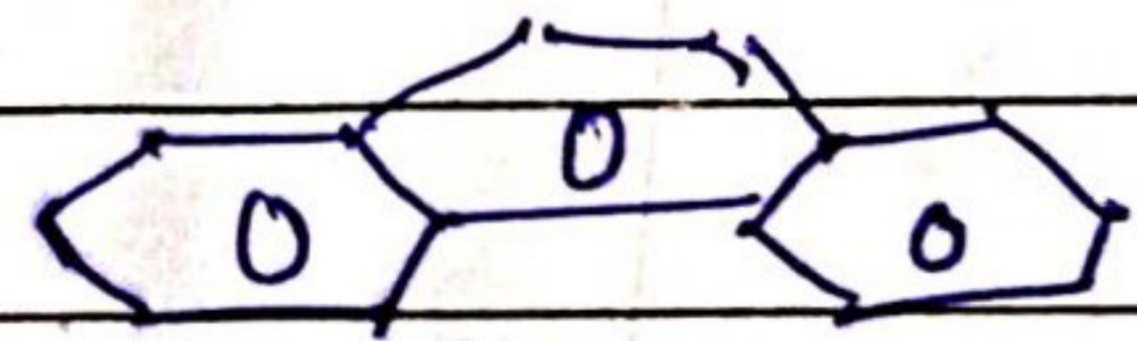
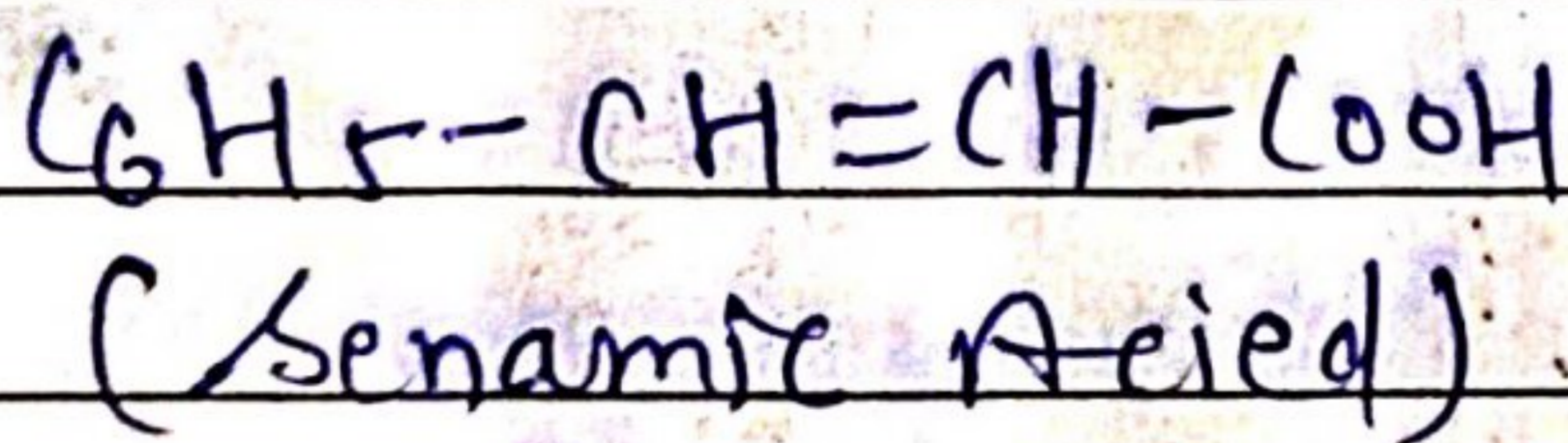


Result :-

In given mixture

Comp "A"

Comp "B"





Object :- In given mixture identified organic compound.

Equipment :- Test tube, Bika, Holder, Conical flask

Comp I

physical state - Solid

Colour - Colourless

Smell - Smellless

Nature - Acidic

flame test - Not Burn with

Smoke. So it is

Aliphatic Comp.

Table Test

Expt.

Test

Result

1. N-Test - L.S. Soln.

fresh FeSO<sub>4</sub>

Soln + NaOH + HCl

Persian Blue  
Colour Not  
produced

N - Absent

2. functional group Test

Carboxylic Group

Test. Comp. water

Soln + 5% NaHSO<sub>3</sub>

Soln mixed

CO<sub>2</sub> gas  
are produced

- COOH group  
present

m.p

101°C

oxalic acid

Special Test

Comp + dil. H<sub>2</sub>SO<sub>4</sub> +

CO + CO<sub>2</sub> gas  
are produced are present

Heat

Teacher's Signature.....



• Comp. 2 ml peric. sol<sup>n</sup> + CaCl<sub>2</sub> sol<sup>n</sup>      white colour precipitate      oxalic acid are present

Comp II

physical state - solid  
 colour - colourless  
 smell -

flame Test - Not Burn with smoke  
 So it Aromatic Comp.

Nature - Acidic.

Table Test

Expt.	Test	Result
1. N-Test	perision Blue colour Not produced.	N-absent
2. functional group Test		
(a) Carboxylic Group		
Test. - Comp. water sol <sup>n</sup> + 8% NaHCO <sub>3</sub> sol <sup>n</sup> are mixed.	CO <sub>2</sub> gas are produced	- COOH group are absent.
3. phenol group Test		
Comp. Acidic sol <sup>n</sup> + some FeCl <sub>3</sub> sol <sup>n</sup>	violet colour produced.	phenol group are present
4. Carbanic Comp. Test		
m.p.	94°C	



2-Naphthol comp +  
H<sub>2</sub>O + FeCl<sub>3</sub>

white precip.  
obtain

α-Nepthalin  
present

Comp + water NaOH  
+ Cl<sub>2</sub> + one drop of  
powder + A

Blue color  
obtain

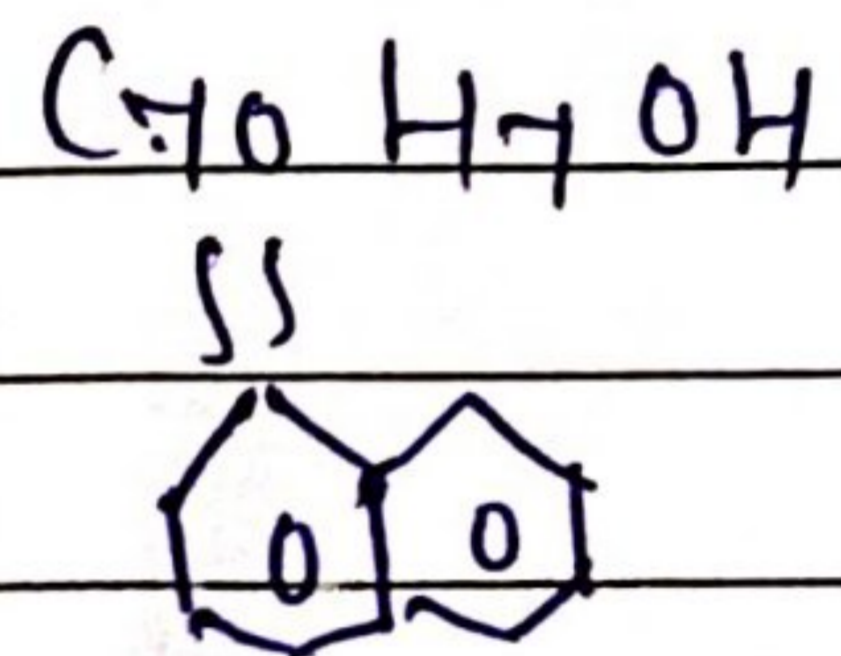
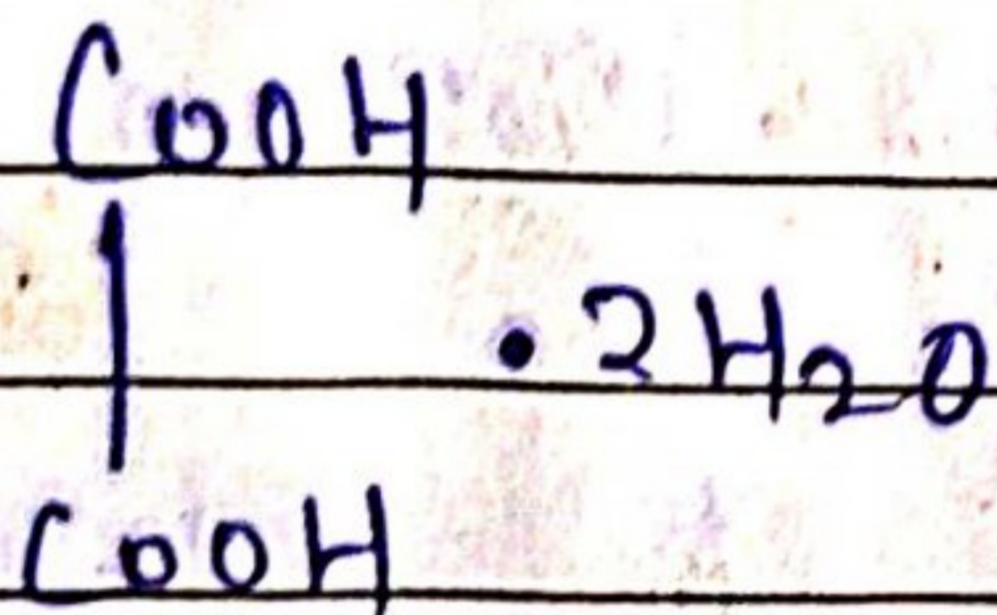
α-Nepthol  
present

Result :-

In given mixture oxalic <sup>acid</sup> comp  
α-Nepthaline are present.

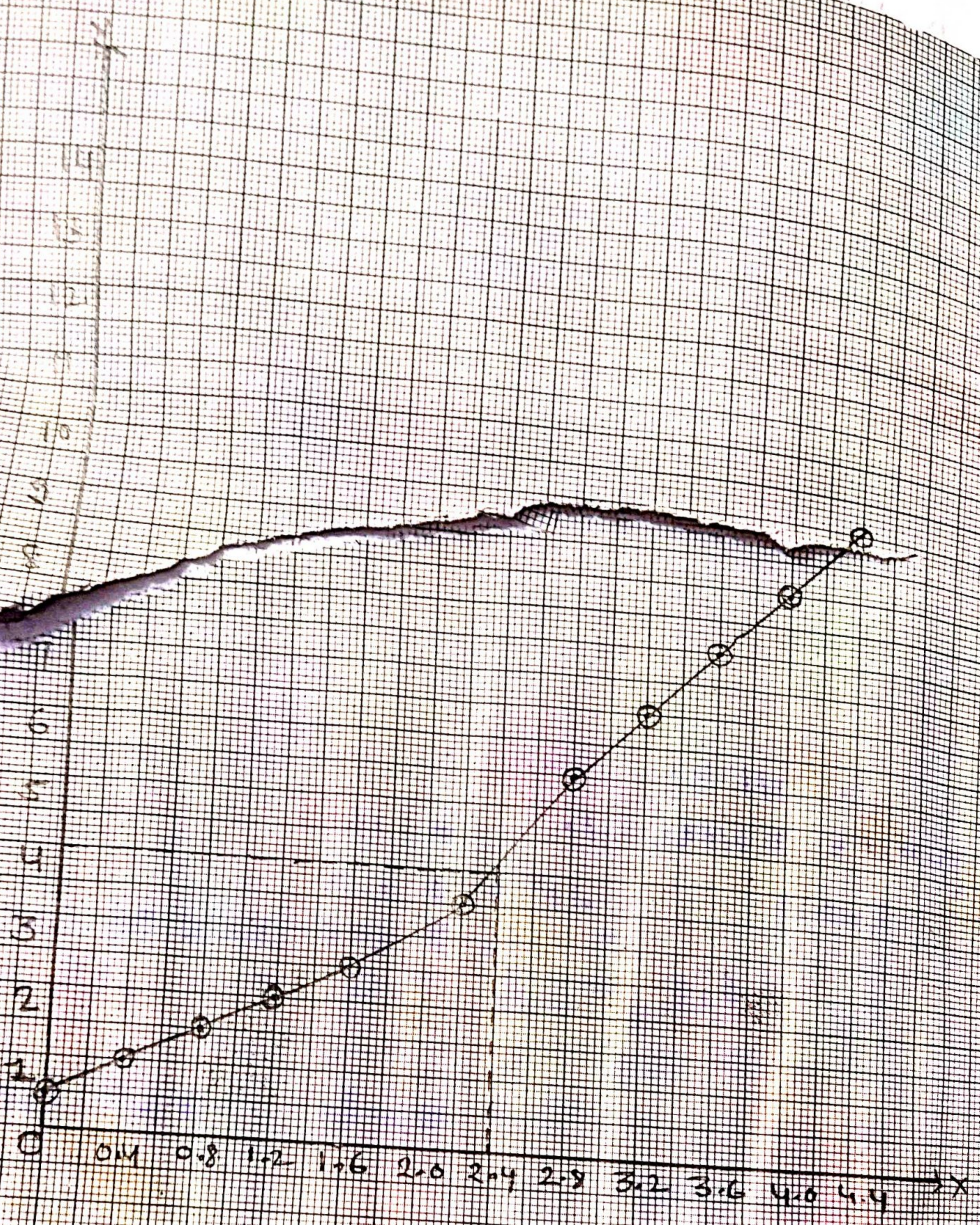
Comp - I

Comp - II



(α-Nepthol)







Object: - NaOH masked in HCl and calculate concentration.

Equipment: - Measured conductivity, measured cell, pipette, Burette, Beaker.

Chemical: - (i) NaOH solution.  
(ii) HCl solution.

Principle: -  $N_1 V_1 = N_2 V_2$   $N_1 =$  Normality of Acid sol<sup>n</sup>  
 $V_1 =$  Volume of Acid sol<sup>n</sup>  
 $N_2 =$  Normality of Basic sol<sup>n</sup>  
 $V_2 =$  Volume of Basic sol<sup>n</sup>.

$N_1 V_1 = 0.1 N \times \text{Volume of base at mazer point}$

$N_1 V_1 = 0.1 N \times \text{Volume of base at mazer point}$   
 and concentration.

$= N_1 \times \text{measured weight}$

Test: -

(i) measurement of HCl = 36.5 gm

(ii) Volume of Acid sol<sup>n</sup> = 10ml



Calculation: -

$$N_1 V_1 = N_2 V_2 \quad N_1 = ?$$

$$V_1 = 10 \text{ mL}$$

$$V_2 = 7 \text{ mL}$$

$$N_2 = 0.1 \text{ N}$$

$$N_1 \times 10 = 0.1 \text{ N} \times \text{Sol}^n \text{ of Base at mesurement}$$

$$N_1 = \frac{0.1 \text{ N} \times \text{Sol}^n \text{ of Base of measure}}{10}$$

$$N_1 = \frac{1 \times 7}{100}$$

$$N_1 = 0.07 \text{ N}$$

HCl measure weight of HCl acid

$$\text{Concentration} = N_1 \times \text{measure weight}$$

$$= 0.07 \times 36.5$$

$$= 2.555 \text{ gm/liter}$$



Expt. Table		
S.No	NaOH Volume of Sol <sup>n</sup>	Conductivity of Sol <sup>n</sup>
1	0.0	26.60
2	1	24.00
3	2	20.00
4	3	17.210
5	4	14.210
6	5	11.210
7	6	8.20
8	7	5.40
9	8	6.00
10	9	7.10
11	10	7.80
12	11	8.40
13	12	8.80
14	13	9.40
15	14	10.00
16	15	10.00
17	16	10.00
18	17	11.60
19	18	12.20
20	19	12.80
21	20	13.40
22	21	13.80
Result :- gm given acid concentration is 2.56 gm/liter.		

Teacher's Signature.....



Object:- In given Acid Sol<sup>n</sup> Salt identify the Sol<sup>n</sup> by Conductivity measurement method.

Equipement:-

measured Conductivity, measured solid

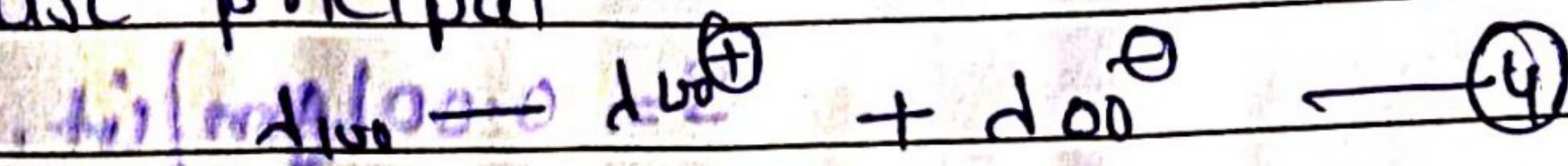
In given Sol<sup>n</sup>, the measured conductivity can be given  $\kappa = k \times V$  — (1)

$$d\kappa = \frac{k \times 100}{C} \text{ — (2)}$$

The acidic Sol<sup>n</sup>, Have the amount of salt Sol<sup>n</sup> is low, so it is fully in-soluble & in soluble salt

$$d\kappa = 100 \text{ — (3)}$$

Kolouse principal



$$C = \frac{k \times 100}{d\kappa^{\oplus} + d\kappa^{\ominus}}$$

While the in soluble salt solubility is less soluble so

$$\kappa_{\text{Sol}^n} = \kappa_{\text{Sol}^n} - \kappa_{\text{H}_2\text{O}} \text{ — (5)}$$

So solubility in less soluble salt

$$C = \frac{(\kappa_{\text{Sol}^n} - \kappa_{\text{H}_2\text{O}}) \times 100 \text{ gm/liter}}{d\kappa^{\oplus} + d\kappa^{\ominus}}$$



Calculation:-

$$K = K_{sol} - K_{H_2O}$$

$$K = 1.200 - 0.954$$

$$K = 0.246 \Omega^{-1} m^{-1}$$

But room on Conductivity

$$K = 0.00246 \Omega^{-1} m^{-1}$$

~~So!  $S = \frac{0.00246 \times 1000}{143.4 \times 200} \times 116.70$~~

$$S = \frac{0.246 \times 116.70}{143.4 \times 200}$$

$\frac{0.00143}{400} + 400$   $\frac{0.00143}{400} + 400$

$$C = \frac{0.00143}{400} + 400$$

2296.  $\frac{0.00143}{400} + 400$

$$C = \frac{0.00143}{400} + 400$$

2296.  $\frac{0.00143}{400} + 400$

$$C = \frac{0.00143}{400} + 400$$

$$C = \frac{0.00143}{400} + 400$$



Test Table :-

Temperature = 20°C

Cell measured = 1 cm<sup>2</sup>

$$\text{Conductivity of water} = \frac{0.93 \times 0.947 \times 0.976}{3}$$

$$= 0.954$$

$$\text{Conductivity of specific water} = 0.954 \times 1$$

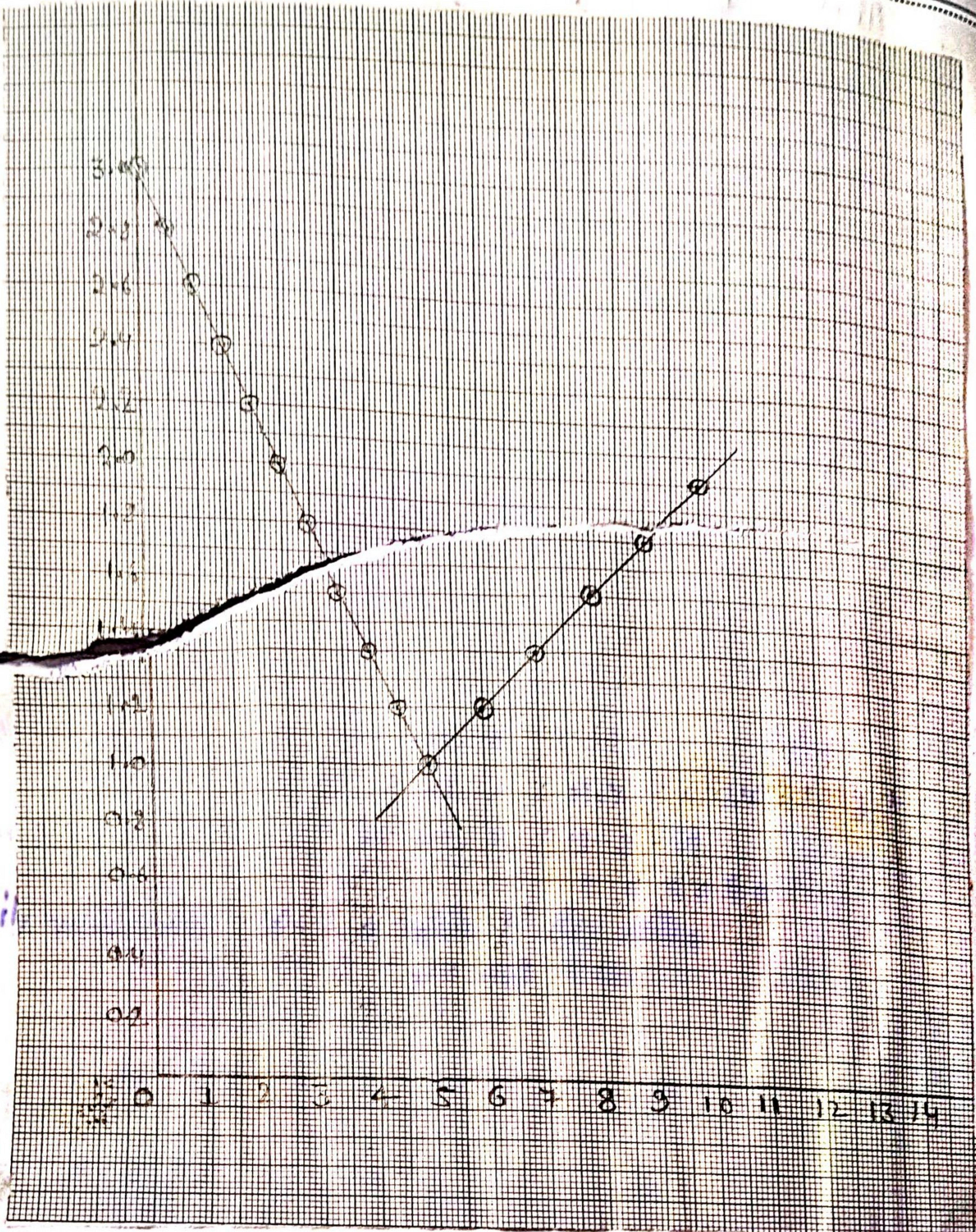
$$= 0.954$$

S.No	Basoy Sol <sup>n</sup> Test	Basoy Specific Conductivity	medium $\mu\text{m}^{-1}$
1.	1.194	$1 \times 1.194 \times 1.194$	
2.	1.200	$1 \times 1.200 \times 1.200$	1.200
3.	1.206	$1 \times 1.206 \times 1.206$	

Result :-

On 20°C Tem. The less solubility of salt Basoy = 0.001 gm/lt.





5th



Object:-

In given Conductivity measure  
weak acid ( $\text{CH}_3\text{COOH}$ ) Calculate Concentration  
of  $\text{NaOH}$  (Base)

useful Equipement:- measured Conductivity  
measure cell pipette, microtube, Beaker

Chemical:-  $\text{NaOH}$ , (i)  $\text{CH}_3\text{COOH}$

Principle:-  $N_1 V_1 = N_2 V_2$

$N_1$  = Normality of acid sol<sup>n</sup>.

$N_2$  = Normality of Base sol<sup>n</sup>.

$V_1$  = Volume of acidic sol<sup>n</sup>.

$V_2$  = Volume of Base sol<sup>n</sup>.

$N_1 V_1 = 0.1 N \times \text{Volume of Base on measure point}$

$N = \frac{0.1 N \times \text{Volume of Base on measure point}}{V}$

$N_1 = \text{Concentration} = N \times \text{measure weight}$

Test:-

(i)  $\text{CH}_3\text{COOH}$  measure weight of

(ii) Volume of acid sol<sup>n</sup>, = 10 ml



Calculation :-

$$N_1 V_1 = N_2 V_2$$

So

$$V_1 = 10 \text{ mL}$$

$$V_2 = 2.4 \text{ mL}$$

$$N_2 = 0.1 \text{ N}$$

So  $N_1 \times 10 = 0.1 \text{ N} \times \text{Volume of Base at measure point}$

$$N_1 = \frac{0.1 \text{ N} \times \text{Volume of Base at measure point}}{100}$$

$$N_1 = \frac{2.4}{100} = 0.024$$

$\text{CH}_3\text{COOH}$  Required measure weight = 60.06 g

Concentration =  $N_1 \times \text{measure weight}$

$$= 0.024 \times 60.06$$

$$= 1.441 \text{ gm/liter}$$



Table Test		
S.N	Volume of mixed Basic Sol <sup>n</sup> (ml)	Conductivity of Sol <sup>n</sup>
1	0.0	0.42
2	0.4	0.82
3	0.8	1.24
4	1.2	1.68
5	1.6	2.12
6	2.0	2.48
7	2.4	4.01
8	2.8	5.50
9	3.2	6.51
10	4.0	7.42
11	4.4	8.62
12	4.8	9.42
13	4.8	10.91

Result: -

In given Sol<sup>n</sup> Concentration is  
1.44 gm/liter



Object :-

In a water sol<sup>n</sup> By the method of Himant measured substance by atomic weight

Equipment :- Thermometer, Bika.

Chemical :- Ice, salt pure of solvent.

principle :- Any unevaporation the solubility of substance soluble w<sub>gm</sub> which show in  $\Delta T_f$

$$\Delta T_f = \frac{1000 \times K_f \times W}{m \times W} \quad \text{OR} \quad \frac{1000 \times K_f \times W}{\Delta T_f \times W}$$

$T_f$  = Cold measurement

$W$  = weight of sol<sup>n</sup>

$W$  = weight of soluble

$K_f$  = Cold measured

$m$  = atomic weight of sol<sup>n</sup>.

Test :-

(i) Volume of water sol<sup>n</sup>

(ii) Density on Room Temp. of water  
= 1 gm/lit.

(iii) given weight of water (V x d)  
15 x 1 = 15 gm

(iv) Cooling point of water



Calculation:- Cooling point of water soln.

$$T_0 = \frac{T_1 + T_2 + T_3}{3}$$

$$T_0 = \frac{(-6) + (-6) + (-6)}{3} = -6^\circ\text{C}$$

Q. Cooling Study of soln

$$T = \frac{(-4) + (-4) + (-4)}{3} = -4^\circ\text{C}$$

3. Cooling points

$$m = \frac{K_f \times 1000 \times W}{\Delta T \times W}$$

$$m = \frac{1.86 \times 1 \times 1000}{15 \times 2} = \frac{1860}{30}$$

$$m = 62 \text{ gm.}$$



Testing Table:-

S.No.	Weight of soln	Temp. of soln.	medium Temp of soln.	Temp of soln.	Freezing point	atomic weight
				10°C	$T_f(T_0^{-1})$	weight (m)
1	1	-4				
2	1	-4	-4	-6	$-4 - (-6)$	62 gm
3	1	-4			= 2	

Result :-

gn given Subsequent of atomic weight is 62 gm.



Object:- By conductivity method given HCl sol<sup>n</sup>. Calculate conduct concentration.

Equipment:- measure conductivity, conductivity cell, pipette, microburet, Biker etc.

Chemical:- (i)  $\text{NH}_4\text{OH}$  sol<sup>n</sup> HCl  
(ii) HCl sol<sup>n</sup>.

Principle:- In HCl sol<sup>n</sup> mixed  $\text{NH}_4\text{OH}$  sol<sup>n</sup> the last point conductivity is decreased. on after this  $\text{NH}_4\text{OH}$  have no change.

$$N_1 V_1 = N_2 V_2$$

where  $N_1$  = normality of acid sol<sup>n</sup>

$V_1$  = volume of acid sol<sup>n</sup>.

$N_2$  = normality of basic sol<sup>n</sup>.

$V_2$  = volume of basic sol<sup>n</sup>.

So  $N_1 V_1 = \frac{0.1 N \times \text{volume of basic at m.e.p.}}{N_1}$

Concentration = mix measure weight

Test.  $\rightarrow$  Above HCl sol<sup>n</sup> the const. is 40.0 ml



X-AXIS → 1 UNIT  
Y-AXIS → 0.1 UNIT

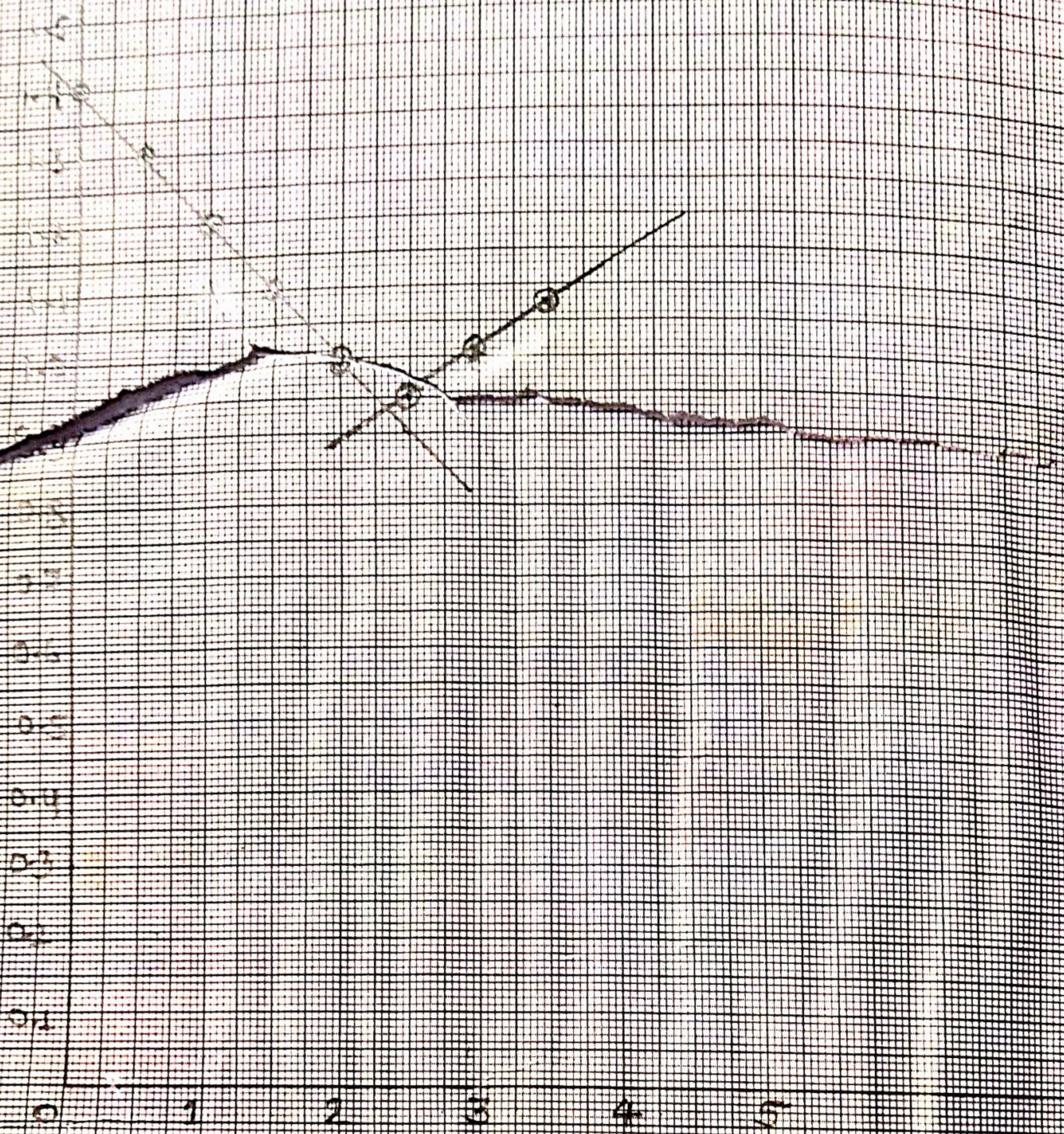




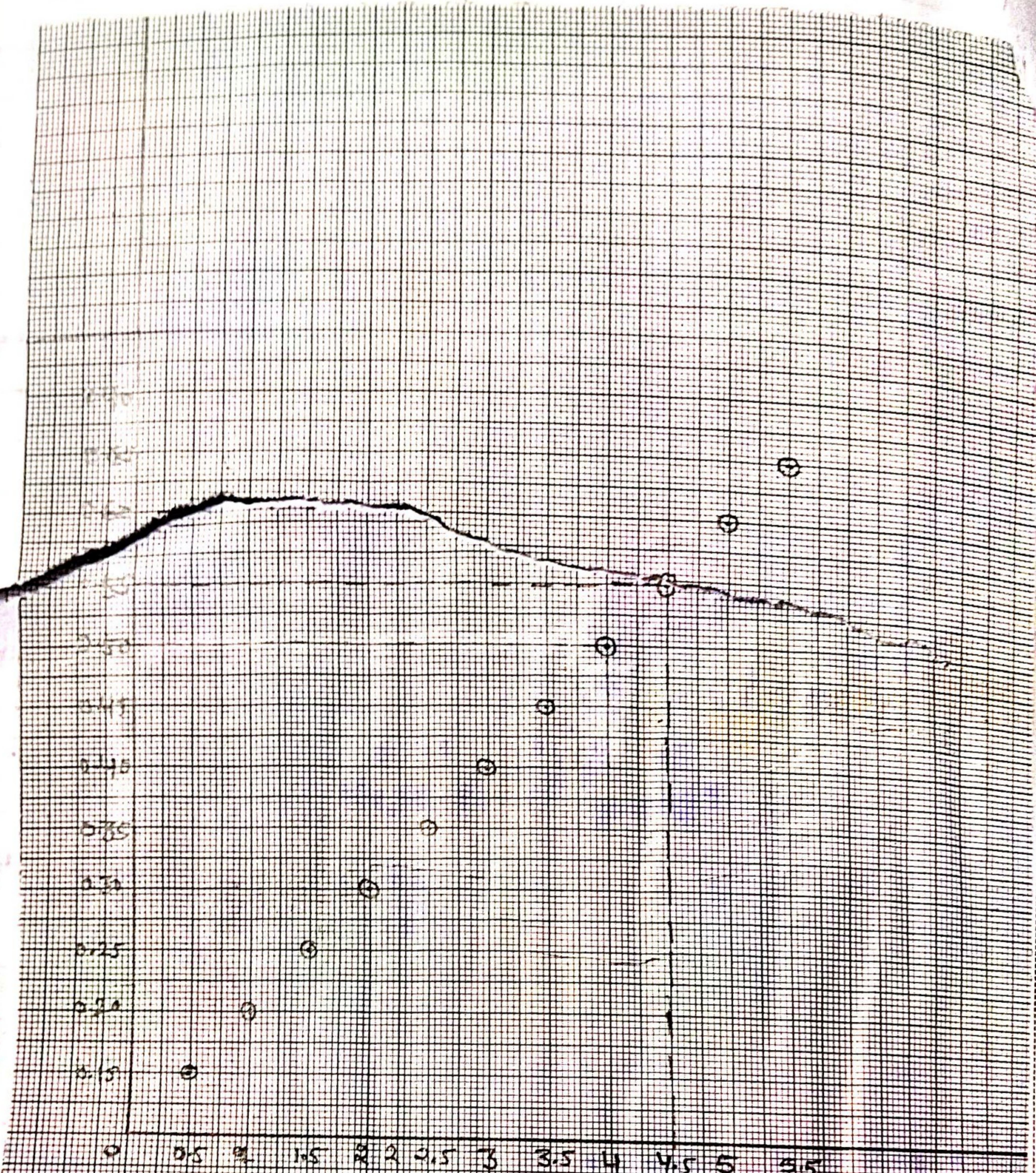
Table Test		
S.No	NaOH mixed volume (ml)	Conductivity
1	0.0	1.48
2	0.5	1.30
3	1.0	1.20
4	1.5	1.12
5	2.0	1.04
6	2.5	0.98
7	3.0	0.97
8	3.5	0.986
9	4.0	0.986
10	4.5	1.986
11	5.0	1.00
12	5.5	1.00

Result:-

In given unknown HCl sol<sup>n</sup>.  
Conc. is 0.2824 gm/lit.



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$\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$  (5%)



Object:- By the method of Kellary to Calculate Construction of unknown known soln.

Equipment :- flask, Burette, Buret, test Buret.

Soln. :- potassium permagnate 0.001N soln.

pricipal:- The pink colour of potassium permagnate by the water present of  $Mn^{+2}$  ion. So with the help of colour be calculate conc. of  $Mn^{+2}$  in soln. In this different conc. of magng ions are used (all soln. & max. ca or density) (0 measured or the graf of this us tell the conc. of soln.)

Test:- Temp. = 25°C



Calculation:-

$$(1) N_1 = 0.001 \quad V_1 = 1 \text{ ml} \quad V_2 = 10 \text{ ml} \quad N_2 = ?$$

$$N_2 = N_1 V_1 = N_2 V_2$$

$$N_2 = \frac{N_1 V_1}{V_2} = \frac{0.001 \times 1}{10} = 1 \times 10^{-4} \text{ N}$$

$$N_2 = 0.001, \quad V_1 = 2 \text{ ml}, \quad V_2 = 10 \text{ ml} \quad N_2 = ?$$

$$N_2 = \frac{0.001 \times 2}{10} = 2 \times 10^{-4} \text{ N}$$

$$3) \quad N_2 = \frac{0.001 \times 3}{10} = 3 \times 10^{-4} \text{ N}$$

$$4) \quad N_2 = \frac{0.001 \times 4}{10} = 4 \times 10^{-4} \text{ N}$$

$$5) \quad N_2 = \frac{0.001 \times 5}{10} = 5 \times 10^{-4} \text{ N}$$

$$6) \quad N_2 = \frac{0.001 \times 6}{10} = 6 \times 10^{-4} \text{ N}$$

$$7) \quad N_2 = \frac{0.001 \times 7}{10} = 7 \times 10^{-4} \text{ N}$$

$$8) \quad \frac{0.001 \times 8}{10} = 8 \times 10^{-4} \text{ N}$$

$$9) \quad N_2 = \frac{0.001 \times 9}{10} = 9 \times 10^{-4} \text{ N}$$

$$10) \quad N_2 = \frac{0.001 \times 10}{10} = 10 \times 10^{-4} \text{ N}$$

$\text{KMnO}_4$

= 158

weight = 31.6

Molarity = Normality  $\times$  measured w

$$= 3 \times 31.6 \times 10^{-4}$$

$$= 94.8 \times 10^{-4} \text{ m/lit.}$$



S.No.	Conc. of soln	Absorption
1	$1 \times 10^{-4}$	1.19
2	$2 \times 10^{-4}$	1.09
3	$3 \times 10^{-4}$	1.05
4	$4 \times 10^{-4}$	0.96
5	$5 \times 10^{-4}$	0.73
6	$6 \times 10^{-4}$	0.84
7	$7 \times 10^{-4}$	0.76
8	$8 \times 10^{-4}$	0.61
9	$9 \times 10^{-4}$	0.92
10	$10 \times 10^{-4}$	0.55
11	$11 \times 10^{-4}$	

Result: -

In given unknown  $Km^{-1}$  solution the conc. of is 94.8 m/lit.



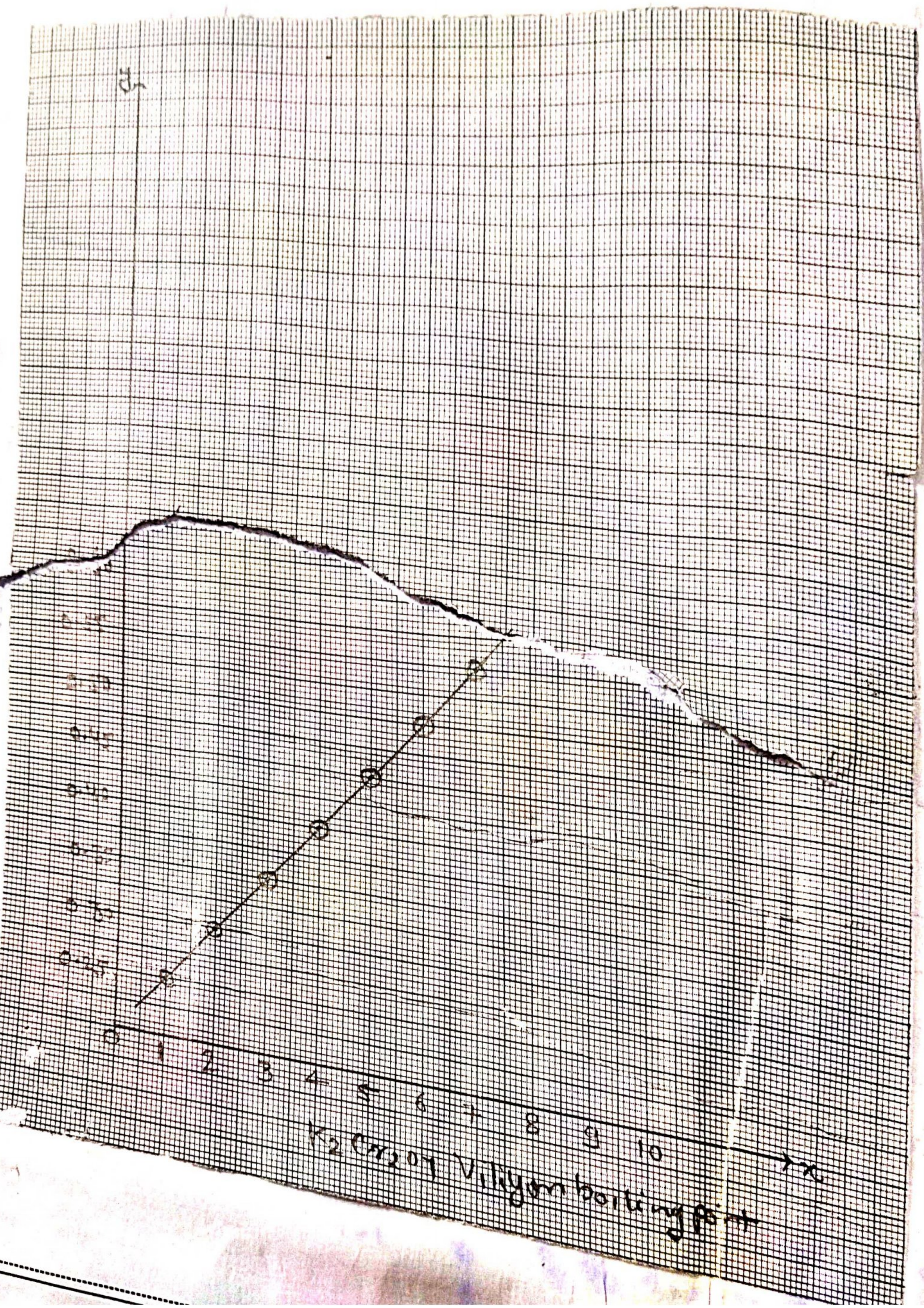
Faint handwritten text at the top of the page, possibly including a date or title.

Handwritten label 'd' located in the upper left corner of the graph area.

Vertical axis labels: 0.25, 0.5, 0.75, 1.0, 1.25, 1.5, 1.75, 2.0, 2.25, 2.5, 2.75, 3.0, 3.25, 3.5, 3.75, 4.0, 4.25, 4.5, 4.75, 5.0, 5.25, 5.5, 5.75, 6.0, 6.25, 6.5, 6.75, 7.0, 7.25, 7.5, 7.75, 8.0, 8.25, 8.5, 8.75, 9.0, 9.25, 9.5, 9.75, 10.0

Horizontal axis labels: 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10

Handwritten text at the bottom of the graph:  $K_2Cr_2O_7$  Volumetric analysis





PAGE NO. ....

Object :- By the method of Kellary calculate the conc of  $K_2Cr_2O_7$  soln.

Equality :- meter, flask, Buret, Burette Test tube.

Solution :- 0.1N soln. of  $K_2Cr_2O_7$

principle :-

The orange colour of  $K_2Cr_2O_7$  because of di-hydrated water of  $Cr_2O_7$  gas so with the help of colour we calculate conc of soln.  $Cr_2O_7^{2-}$  in this different conc of Di Chromet from the soln of is confirmed  $\lambda$ -maxim.

(0.00 measured) A or B after this we draw a line calculate conc. of unknown soln.



Calculate -

(i)  $M_1 = 0.1M$   $V_1 = 1ml$   $V_2 = 10ml$   $M_2 = ?$

$$M_2 = \frac{M_1 V_1}{V_2} = \frac{0.1 \times 1}{10} = 1 \times 10^{-2} M$$

2)  $M_2 = \frac{0.1 \times 2}{10} = 2 \times 10^{-2} M$

3)  $M_2 = \frac{0.1 \times 3}{10} = 3 \times 10^{-2} M$

4)  $M_2 = \frac{0.1 \times 4}{10} = 4 \times 10^{-2} M$

5)  $M_2 = \frac{0.1 \times 5}{10} = 5 \times 10^{-2} M$

6)  $M_2 = \frac{0.1 \times 6}{10} = 6 \times 10^{-2} M$

7)  $M_2 = \frac{0.1 \times 7}{10} = 7 \times 10^{-2} M$

8)  $M_2 = \frac{0.1 \times 8}{10} = 8 \times 10^{-2} M$

9)  $M_2 = \frac{0.1 \times 9}{10} = 9 \times 10^{-2} M$

10)  $M_2 = \frac{0.1 \times 10}{10} = 10 \times 10^{-2} M$

weight of  $K_2Cr_2O_7 = 294$

measured weight =  $\frac{294}{6} = 49$

molarity = measured weight  $\times$  normality

$$= 49 \times 5 \times 10^{-2}$$

$$= 2.45 ml$$



S. NO.

0.1N Conc. of  $K_2Cr_2O_7$  Soln.

absorption (A)

1.	$1 \times 10^{-1}$	0.25
2.	$1 \times 10^{-2}$	0.28
3.	$1 \times 10^{-3}$	0.31
4.	$1 \times 10^{-4}$	0.38
5.	$1 \times 10^{-5}$	0.39
6.	$1 \times 10^{-6}$	0.42
7.	$1 \times 10^{-7}$	0.43
8.	$1 \times 10^{-8}$	0.44
9.	$1 \times 10^{-9}$	0.47
10.	$1 \times 10^{-10}$	0.48
11.	$1 \times 10^{-11}$ Unknown	

Result :-

The conc. of unknown soln  $K_2Cr_2O_7$  is 0.48 ml.

*Suman*  
12/04/2021