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CLASS:- M.Sc. (Previous)

SUBJECT:- ORGANIC-CHEMISTRY

ROLL No:- 1285828

COLLEGE:- R. K. VIGYAN (PG) COLLEGE

KALVAR, JAIPUR

S. No.	Name of the Experiment	Page No.	Date of Experiment	Date of Submission	Remarks
(1.)	To prepare m-nitroaniline from nitro-benzene		Summed		
(2.)	To prepare sym-tribromo-benzene from aniline				
(3.)	To prepare benzoic acid from Benzoin				
(4.)	To prepare dibenzalacetone from the benzaldehyde				
(5.)	Separation of identity of the organic compound from given mixture of also suitable derivative.		Summed		
(6.)	Separation of identity of the organic cpd from given mixture of also suitable derivative.				
(7.)	Separation of identity of the organic cpd from given mixture of also suitable derivative.		Summed		

S. No.	Name of the Experiment	Page No.	Date of Experiment	Date of Submission	Remarks
(8.)	separate & identify the organic cpd from given mixture & also form suitable derivatives.		Sumera		
(9)	separate & identify the organic cpd from given mixture & also the suitable derivatives.				
(10.)	separate & identify the organic cpd from given mixture & also form suitable derivative.		Sumera		

Step = 1

Object:-

To prepare m-nitro aniline from nitro-benzene

Requirement:-

Nitrobenzene = 12ml

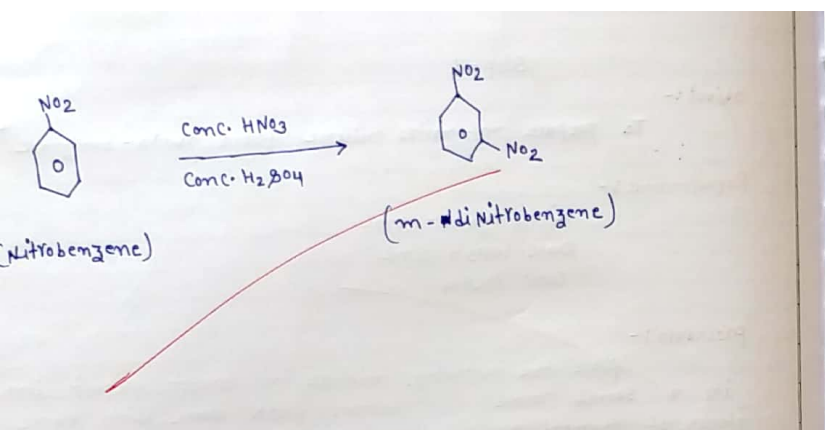
conc. HNO_3 = 15mlconc. H_2SO_4 **Procedure:-**

place the nitrating mixture in correct proportion in a 500ml round bottom flask along with some pieces of unglazed porcelain or glass to avoid bumping attach condenser and add 12ml of the nitro-benzene portion wise.

Add 15ml conc. H_2SO_4 , slowly with constant shaking through the condenser then add 5ml conc. HNO_3 with conc. stirring, reflux the mixture on a boiling water bath for one hour with occasional shaking.

Allow the mixture to cool pour it with constant stirring to 300ml cold water taken in a beaker filter the product, wash with the water and recrystallisation with the alcohol.

Teacher's Signature : _____



Result:- We get bright yellow crystals of the m-dinitrobenzene are 14.5211 gms. In weight and these melting point = -90°C

Sumed
23/4/18

Step = 2

Object :-

To prepare m-nitro aniline from the m-dinitrobenzene.

Requirements :-

m-dinitrobenzene = 6.7 gm

Sodium sulphide = 18 gm

Na_2CO_3 = 6 gm

powdered sulphur = 5 gm

Ice

Procedure :-

Take 15 gm of sodium sulphite dissolve in 30 ml of H_2O add 5 gm sodium bicarbonate add 40 ml methanol with constant stirring.

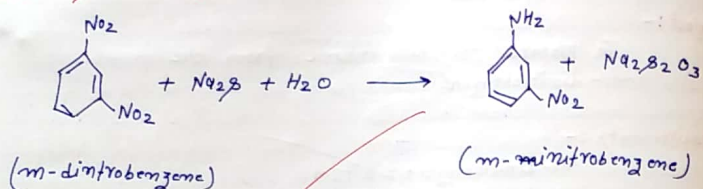
Suspend 6.7 gm of m-dinitrobenzene in 100 ml water in a 500 ml beaker heat it to boiling and then add dropwise with constant and vigorous stirring the sodium sulphite solution from the separating funnel.

Cool the contents for about 20 minutes, by adding ice. Filter the product at the pump and wash with cold water.

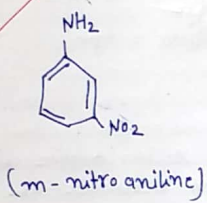
Transfer the product to a 500 ml beaker having about 15 ml of water and 18 ml of conc. HCl boil the contents for about 15 min. When the m-nitroaniline

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Chemicals reactions:-



Structure:-



dissolves leaving the unused sulphur and m-dinitrobenzene.

Filter the contents and precipitate out m-nitro aniline from the filtrate by adding excess of conc. aqeous ammonia solution filter the m-nitro aniline and recrystallize from boiling water.

Result:-

We get by this preparation bright yellow needle shaped crystals of m-nitro aniline which are 5.532 gm in weight.

crystals melting point is -114°C

S. Umesh
23/4/18

To prepare sym-tribromobenzene from aniline.

Step-1.

Object:-

To prepare 2,4,6-tribromoaniline from aniline

Requirements:-

Aniline = 5ml

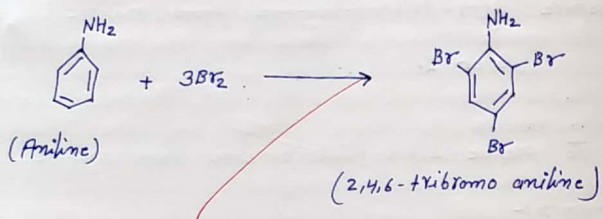
Glacial acetic acid = 19ml

Bromine in glacial acetic acid 8.4ml in 20ml of glacial acetic acid.

Procedure:-

Take 5ml of aniline and 19ml of the glacial acetic acid in a flask. place the flask in ice bath & add to it 8.4ml of bromine water in 20ml of glacial acetic acid dropwise with constant stirring through a burette or separating funnel, pour the reaction mixture solution to a beaker having excess of water filter the product wash with water and recrystallise from alcohol or rectified spirit.

Chemical Reaction:-



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Result:-

We get colourless shining long needles crystal which are 10.027 gms in weight and these are of sym-tribromobenzene these melting point is 120°C

Step-2

Object:-

To prepare sym-tribromobenzene from 2,4,6-tribromo aniline.

Requirements:-

sym-tribromoaniline = 10gm

Rectified spirit = 60gm

benzene = 15ml

conc. H_2SO_4 = 3.5ml

Sodium Nitrate = 120ml

decolourising carbon = 2.5gm

Ice

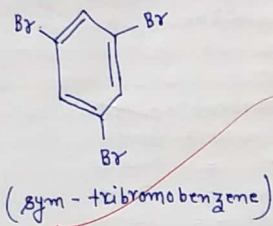
Procedure:-

Dissolve 10gm of sym-tribromoaniline in 60ml rectified spirit and 15ml benzene by heating on a water bath add about 3.5ml of conc. H_2SO_4 dropwise from the burette or a graduated pipette to the hot solution of tribromoaniline and ~~stir~~ the solid the liquid gently.

Heat the constant on water bath and using reflux water condenser until the clear solution boils.

Remove the condenser add about 1.75gms of sodium nitrate attach the condenser and shake the flask vigorously.

structure :-



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Remove the condenser and heat the flask boiling water bath till the gas is evolved. Allow the reaction mixture to stand for about 10 minutes.

Now immerse it on ice bath and then filter the product with solution on a buchner funnel wash with a little amount of alcohol followed by several water washings to remove all the sodium sulphate.

Result :-

We get colourless shining needle shaped crystals of sym-tribromobenzene.

which are 5.2460 gms in weight crystals melting point is 122°C.

Suvarat
23/4/18

To prepare benzoic acid from benzoin.

Step - 1.

Object:-

To prepare benzil from benzoin of HNO_3 .

Requirements:-

Benzoin = 8.5 gm

Glacial acetic acid = 40 ml

conc. HNO_3 = 20 ml

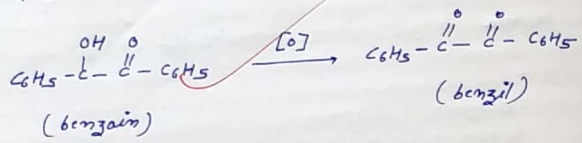
Ice.

Procedure:-

In a round bottom flask we take 0.5 gms of benzoin 40 ml of glacial acetic acid 20 ml of conc. HNO_3 swirl the flask vigorously and then heat the reaction mixture on a steam bath add 150 ml of water min. thoroughly and allow the yellow ppt. of benzil to settle filter the product with suction wash thoroughly with water & dry the product by pressing a crystals with a clean glass stopper.

Although again the benzil obtained is sufficiently pure for conversion to derivatives or to the benzoic acid it may be purified by recrystallisation.

Chemical reaction :-



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from methanol or 5% aqueous ethanol.

Result :- we get yellow crystals of benzoic acid which are 6.462 gms in weight crystals melting point is 95°C.

Suresh
23/4/18

Step-2.

Object:-

To prepared benzoic acid from benzil.

Requirements:-

benzil = 5gms
 KOH = 5gm
 Ethyl alcohol = 15ml
 conc. HCl
 Ice.

Procedure:-

dissolve 5gms of KOH filtrate in 10ml of H₂O in a ~~small~~ flask add 15ml of ethyl alcohol acid mixture will the solution by swirling.

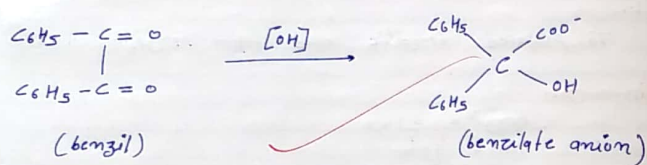
Add 5gms of pure benzil to the resulting solution fit a reflux water condenser to the flask of boil the contents on a water bath for 10-15 min.

Transfer the contents of the flask to a small beaker or a porcelain dish cover it with a watch glass, allow it to stand undisturbed for a several hours.

preferably overnight until potassium benzoate is completely crystallised filter the product and wash it with ice cold ethanol.

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Chemical reaction:-



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Dissolve the product in above 50ml of water and then add an HCl drop by drop till the ppt. of the free acid is complete.

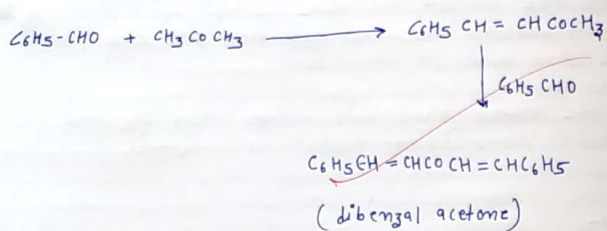
Filter the product wash with water press dry and recrystallise from boiling water or benzene.

Result:-

We get colourless shining crystals of the benzoic acid which are 9.261 gms in the weight

crystals m.p. is 150°C.

Chemical reaction:-



object:-

To prepare dibenzalacetone from the benzaldehyde.

Requirements:-

benzaldehyde = 10ml
Alcohol = 100ml
Acetone = 4ml
Sodium hydroxide = 10gms

procedure:-

Dissolve 10gms of NaOH in 80ml of H₂O and 100ml of alcohol in a flask, cool it in ice bath to about 20-25°C and then add 4ml of acetone.

slowly into installments with an interval of 15 min. but with constant stirring cork the flask or shake it vigorously for 30 min.

Filter the product wash with water or recrystallise from ethyl alcohol or hot ethyl acetate

Result:-

we get bright yellow crystals of dibenzalacetone which are 8.020gms in the weight.

Crystals m.p is 112°C.

Teacher's Signature : _____

(Separation of organic mixture)

Object:-

Separation of identify of the organic compound from given mixture also of also from suitable derivative.

Test for solid compound :-

Primary test :-

Physical state - solid

Colour - colourless

odour - odourless

Solubility - less soluble in water but soluble in organic solvents

Observation Table :-

Experiment	observation	inferences
(1) <u>Flame test</u> :- on burning the organic solid in burner flame	compound burns with smoky flame	compound is aromatic
(2) <u>litmus test</u> :- blue of red litmus paper take in the aqueous solution of the organic compound.	colour of blue litmus paper turned in red	nature of the organic compounds in basic

Teacher's Signature :

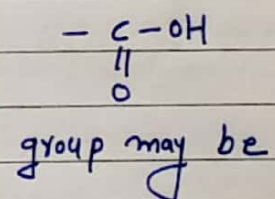
(3.) Element test :-

- | | | |
|--|---|--------------|
| (a) "N" Test - take L.S sol ⁿ + Freshly prepared FeSO ₄ sol ⁿ + conc H ₂ SO ₄ | blue / green ppt sol ⁿ is not obtained | "N" - absent |
| (b) "S" - Test :-
Sodium Nitro pryside test :- take L.S solution + add. Sodium Nitro - pryside | black colour solution or ppt is not obtained | "S" - absent |
| (c) "X" - test :-
L.S solution + conc HNO ₃ + AgNO ₃ | white ppt is not obtained | "X" - absent |

⇒ Test for functional group :-

organic mix. + conc. H₂SO₄
4-5 drop ethyl alcohol +
 $\Delta \longrightarrow$ CO₂

fruits like odour is
now evolved

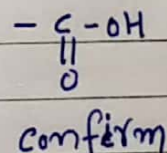
Iodate - colite Test :-

organic mix. + 2% KI
5 drop + 5 drop KIO₃ solⁿ

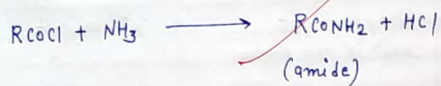
$\Delta \longrightarrow$ then cool +

1ml starch solution

blue colour is
obtained



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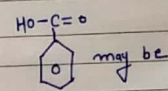


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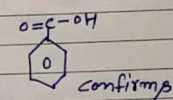
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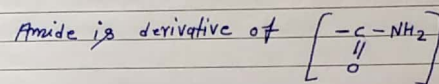
M.P of organic is $\approx 121^\circ\text{C}$



Confirmation test for functional group:-
neutral solution of organic mix. + FeCl_3 ppt is obtained



Formation of the derivative:-



carboxylic acid.

Method:-

- We take about 0.5gm organic compound in a round bottom flask.
- Now we add SOCl_2 & heat it about 20-30 mint. formation of a acylchloride take place by this process
- Now we add it in 10ml cool concentration ammonia solution with dropwise & vigorous reaction takes.
- Keep it in ice bath & cool it we obtained white crystalline ppt of amide.
- MP of $\begin{array}{c} -\text{C}-\text{NH}_2 \\ \parallel \\ \text{O} \end{array}$ is 128°C

Teacher's Signature :

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Test for liquid compounds :-primary test :-

physical state - liquid

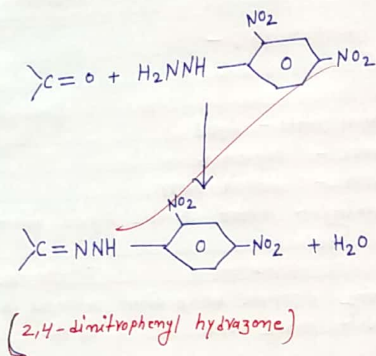
colour - colourless

odour - special odour

solubility - water soluble liquid solvent also

Experiment	observation	Inference
(1) Flame test :- on burning the organic compound in burner's flame	compound burns without smoky flame	compound is aliphatic
(2) Litmus test :- blue and red litmus paper take in the aqueous sol ⁿ of the organic compounds	colour of litmus paper not change	compound is neutral
(3) Element test :-		
(a) "N" - test :- Take L.S sol ⁿ + Freshly prepared FeSO ₄ sol ⁿ + conc. H ₂ SO ₄	blue/green ppt sol ⁿ is not obtained	"N" - absent
(b) "S" - test :- take L.S sol ⁿ + add sodium nitro-prusside	black colour sol ⁿ or ppt is not obtained	"S" - absent
(c) "X" - test :- L.S - sol ⁿ + conc. H ₂ SO ₄ + AgNO ₃	White ppt is not obtained	"X" - absent

Teacher's Signature : _____



Test for functional group:-

(a) Sodium bisulphite test - conc. + 2ml NaHSO ₃	White ppt is obtained	-C(=O)- may be
(b) 2,4-dinitrophenyl hydrazine test. Alcoholic sol ⁿ of the organic Cpd + 1ml 2,4-dinitrophenyl hydrazone $\xrightarrow{\Delta}$	yellow or orange colour ppt is obtained	-C(=O)- group is confirm
BP of organic compound	≈ 56°	CH ₃ -C(=O)-CH ₃ may be
Iodoform test - 1ml compound + 2ml I ₂ sol ⁿ + NaOH by dropping + Δ	yellow colour ppt is obtained	CH ₃ -C(=O)-CH ₃ confirm

Confirmative test for functional group:-

Iodoform test - 1ml compound + 2ml I ₂ sol ⁿ + NaOH by dropping + Δ	yellow colour ppt is obtained	CH ₃ -C(=O)-CH ₃ confirm
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Formation of derivative:-

Derivative of carbonyl group is 2,4-dinitrophenyl hydrazone.

Method:- All of first take 1ml compound and 4-5ml of 2,4-dinitrophenyl hydrazone in a test tube then mix up 2 drop of conc. H₂SO₄. Now heat it and then cool it from this process formation of 2,4-dinitrophenyl hydrazone yellow ppt takes place.

MP of 2,4-dinitrophenyl hydrazone is 128°C

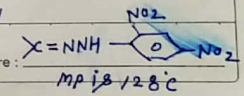
Result:- Solid & liquid compounds are

respectively present in given binary mixture.

Derivative of these compounds:-



derivate is -C(=O)NH₂ ⇒ MP is 128°C



Object:-

Separation and Identify of the organic compound from given mixture & also form the suitable derivative.

Test for solid compound:-**primary test:-**

Physical state - solid crystalline
 Colour - colourless
 odour - odourless
 Solubility - soluble in water & insoluble
 organic solution

observation table:-

S.No.	Experiment	observation	Inference
1.	Flame test:- on burning the organic solid in burner's flame	compound burns without smoky flame	compound is aliphatic
2.	litmus test:- blue & red litmus paper take in the aqueous solution of the organic compound	colour of litmus paper is not change in red or blue	Nature of organic compound is neutral
3.	Element test:-		
(a)	"N"-test:- Take L.S sol ⁿ + Freshly FeSO ₄ sol ⁿ + conc. H ₂ SO ₄	blue/green colour ppt sol ⁿ is obtained	"N"-present

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Expt. No.

(b)	<p>"S"-test :- Sodium Nitroprusside test - take L.S sol^m + add. Sodium Nitroprusside</p>	<p>black colour sol^m ppt is not obtained</p>	<p>"S"- absent</p>
(c)	<p>"X"-test :- L.S sol^m + conc. HNO₃ + AgNO₃</p>	<p>White ppt is not obtained</p>	<p>"X"- absent</p>
<p>Test for functional group:-</p>			
<p>organic cpd + NaOH solution $\xrightarrow{\Delta}$</p>		<p>Ammonia a like odour is evolved</p>	<p>$\begin{matrix} -C-NH_2 \\ \\ O \end{matrix}$ may be</p>
<p>above sol^m + conc. HCl</p>		<p>White ppt is obtained</p>	<p>$\begin{matrix} -C-NH_2 \\ \\ O \end{matrix}$ confirm</p>
<p>Melting point of organic compound</p>		<p>$\approx 132^\circ C$</p>	<p>$H_2N-\begin{matrix} C \\ \\ O \end{matrix}-NH_2$ may be</p>

Confirmative test for functional group:-

Byret test:-

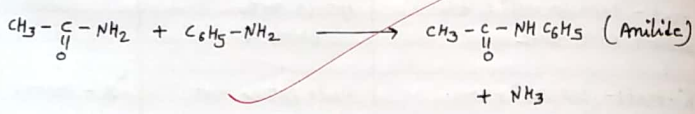
organic mix. + $\xrightarrow{\Delta}$ then
 $\uparrow NH_3 \longrightarrow$ cool \longrightarrow
 obtained

Violet colour sol^m ppt is obtained

White ppt + 2-3 drop H₂O + Cu₂SO₄ solution + NaOH

$NH_2-\begin{matrix} C \\ || \\ O \end{matrix}-NH_2$
 Confirms

Teacher's Signature : _____



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Formation of derivative of amide:- Amide reaction with aniline and formed $(\text{CH}_3-\overset{\text{O}}{\underset{\text{O}}{\parallel}}\text{C}-\text{NHC}_6\text{H}_5)$

Method:- Take 0.5gm compound and heat with 1ml aniline ammonia like odour is evolved white ppt is obtained.
MP of aniline is 250°C

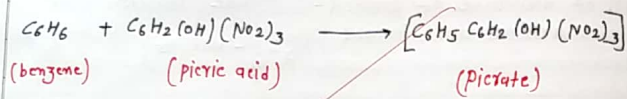
Test for liquid compounds:-

primary test:- physical state - liquid form
colour - colourless
odour - specific
solubility - less soluble in the water but soluble in organic solvent


Observation table:-

Experiment	Observation	Inference
1) Flame test:- on burning organic solid in the burner's flame	compound burns with smoky flame	Nature of compound is aromatic
2) Litmus test:- blue or red litmus paper take in aqueous sol ⁿ of the organic compound.	colour of blue/red litmus paper not changed	Nature of organic compound is neutral


Teacher's Signature : _____



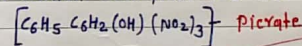
3) Element test:-

(a) "N"-test:- take L.S sol ⁿ + freshly FeSO ₄ sol ⁿ + conc. H ₂ SO ₄	blue/green ppt sol ⁿ is not obtained	"N"-absent
(b) "S"-test:- take L.S sol ⁿ + sodium nitroprusside	black colour sol ⁿ ppt is not obtained	"S"-absent
(c) "X"-test:- Compound + AlCl ₃ + Δ + CHCl ₃ BP	Red colour sol ⁿ /ppt is obtained ≈ 80°	Aromatic hydrocarbon is confirmed  may be


Determination test for functional group:-

organic mixture + conc. HNO ₃ + conc. H ₂ SO ₄	yellow colour sol ⁿ /ppt is obtained in only form	 confirm
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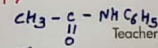
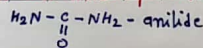
Formation of derivative of aromatic hydrocarbon cpd:- Aromatic hydrocarbon reaction with picric acid and formed picrate



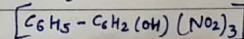
Method:- take 0.5gm organic cpd are mix. up 1ml picric acid & stirring constant we obtained solid picrate. We crystallize it with C₆H₆ mp of this is 90°C


Result:- Solid & liquid cpd are $H_2N-\overset{\text{O}}{\parallel}{C}-NH_2$ &  respectively present in given binary organic mix.

Derivative of these cpds:-



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 - picrate

MP = 90°C

object:- Separate & Identify the organic cpd from given mixture & also form suitable derivatives.

Test for solid compound:-

primary test:-

physical state - solid crystalline

colour - pink brown colour

odour - carbolic odour

solubility - less soluble in water
but solvents

Observation table:-

S.No	Experiment	Observation	Inference
(1)	Flame test:- on burning organic cpd in burner's flame	cpd burns with smoky flame	cpd is aromatic
2)	litmus test:- blue/red litmus paper take in the aqueous sol ⁿ of the organic cpd.	colour of litmus paper not change in red or blue	nature of cpd is neutral
3)	Element test:-		
(a)	"N" test:- Take L's sol ⁿ + Freshly FeSO ₄ + conc. H ₂ SO ₄	blue/green colour ppt is not obtained	"N" - absent
(b)	"S" test:- take L's sol ⁿ + add sodium nitroprusside	black colour ppt is not obtained	"S" - absent

Teacher's Signature : _____

(c) "X"-test :- $LS - soln + conc. HNO_3 + AgNO_3$ White ppt is not obtained "X"-absent

Test for functional group :-

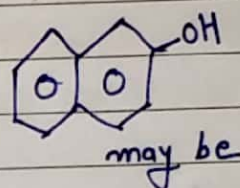
$soln$ organic cpd + sodium metal

H₂ gas is evolved

-OH group may be

MP of organic cpd.

≈ 123°C



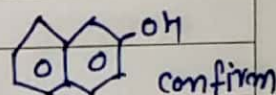
Confirmation test for functional group :-

organic mixture + NaOH

$soln + CHCl_3 +$

$\Delta \longrightarrow$

blue colour salt/ppt is obtained



Formation of derivative of phenolic group :-

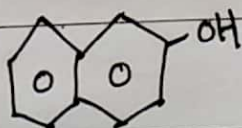
P-nitro benzal :-

Alcohol reaction with p-nitro benzal not chloride is formed p-nitro benzoate

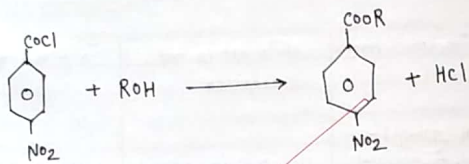
Method :- We take 10ml alcohol & 2-gram p-nitrobenzal chloride & kept in water bath about 15-20 minutes.

Then add 5% NaHCO₃ $soln$ & 10ml gradually then cool the mixture. We obtained solid p-nitro-benzoate & wash with cool water

Recrystallise with alcohol \rightarrow such we get p-nitrobenzoate derivative of β -naphthal.



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(Nitrobenzoyl chloride) \longrightarrow (p-nitro benzoate)

Test for liquid compounds:-

primary test:-

physical state - liquid

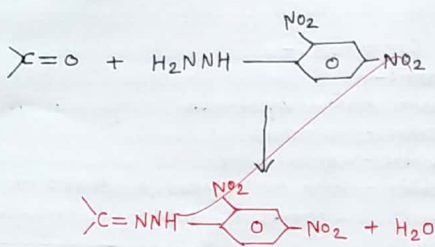
colour - colourless

odour - special odour

Solubility - Water soluble liquid & soluble in organic solvents.

S.No	Experiment	Observation	Inference
1)	Flame test:- on burning the organic cpd in burner's flame	cpd burns without smoky flame	cpd is an aliphatic
2)	Litmus test:- take red or blue litmus paper in aqueous sol ⁿ in organic cpd.	colour of litmus paper not change in blue/red	nature of cpd is neutral.
3)	Element test:-		
(a)	"N"-test:- take L/S sol ⁿ + freshly prepared FeSO ₄ sol ⁿ + conc. H ₂ SO ₄	green/blue coloured ppt sol ⁿ is not obtained	"N"- absent
(b)	"S"-test:- Take L/S sol ⁿ + sodium nitroprusside	black colour is not obtained	"S"- absent
(c)	"X"-test:- L/S sol ⁿ + conc. HNO ₃ + AgNO ₃	White ppt is not obtained	"X"- absent

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Test for functional group :-

Sodium bisulphite test :-

Compound + 2ml NaHSO_3	White ppt is obtained	$-\overset{\text{O}}{\parallel}{C}-$ group may be
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2,4-dinitrophenyl hydrazine test :-

Alcoholic sol ⁿ of organic cpd + 1ml 2,4-dinitro-phenyl hydrazine + Δ	yellow/orange colour ppt is obtained	$-\overset{\text{O}}{\parallel}{C}-$ group confirms
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BP of organic cpd -

$\approx 56^\circ\text{C}$

$\text{CH}_3-\overset{\text{O}}{\parallel}{C}-\text{CH}_3$ may be

Determination test for functional group :-

Iodoform test :-

1ml cpd + 2ml I_2 sol ⁿ + NaOH by dropping + Δ	yellow colour sol ⁿ is obtained	$\text{CH}_3-\overset{\text{O}}{\parallel}{C}-\text{CH}_3$ confirms
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Derivative :-

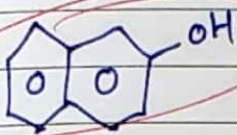
2,4-dinitrophenyl hydrazine react with acetone & formed 2,4-dinitro phenyl hydrazine

Method :-

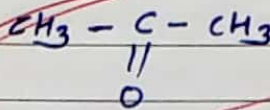
We take 1ml cpd & 4-5ml 2,4-dinitro-phenyl hydrazine in a test tube then add 2-drop conc H_2SO_4 & heat then cool hydrazine then crystalline it.

Result :-

The solid & liquid compounds are as

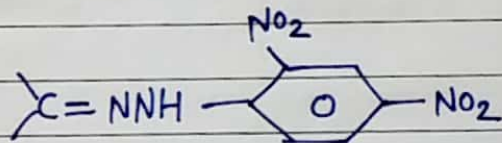
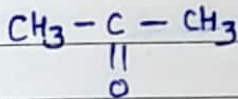
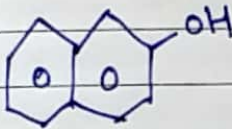


&



respectively in given

binary organic mixture.

Derivative :-

MP :- 128°C

~~Stemmed~~
2314118

object :-

separate and identify the organic compound from given mixture of also form suitable derivative.

Test for solid compounds :-**primary test :-**

physical state = solid crystals

colour = colourless

odour = odourless

Solubility - hot water soluble &
soluble in organic cpd.

Observation Table :-

S.No	Experiment	observation	Inference
1)	Flame test :- on burning of organic mix. on burner's flame.	Compound burns with smoky flame	Compound is aromatic
2)	Litmus test :- Red / blue litmus paper take in aqueous sol ⁿ of organic cpd.	colour of blue litmus paper turned into red	nature of cpd is acidic
3)	Element test :-		
(a)	"N"-test :- take L's sol ⁿ + freshly prepared Fe ₂ SO ₄ + conc. H ₂ SO ₄	blue/green coloured ppt is not obtained	"N" - absent.

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(b) "S"-Test:-

take l.s solⁿ +
sodium nitroprussideblack colour ppt is
not obtained

"S"- absent

(c) "X"-test:-

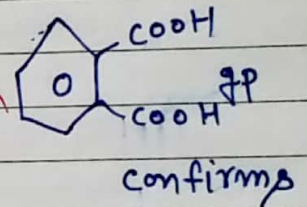
l.s solⁿ + conc. HNO₃
+ AgNO₃white ppt is
not obtained

"X"- absent

4.) Test for functional group:-

organic mix. + conc. H₂SO₄
+ 4-5 drop ethyl alcohol + Δ
→ then Coolfruit like odour is
evolved-C-OH grp
||
O may be

Confirmation test:-

0.2 organic mix. + 0.4 gm
phenol + 2 drops conc. H₂SO₄
+ Δ → then cool
+ 2ml H₂O, NaOHpink colour solⁿ is
obtained

Formation of derivative:-

MP 220°C acid react with SOCl₂
and formed acyl chloride which react with ammonia
+ formed amide derivative.

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Method:-

- Take 0.5gm organic compound in round bottom flask.
- Then acyl chloride is prepared.
 - Now add it in some conc. NH_3 solution by dropping a vigorous reaction takes place.
 - cool mixture on ice bath and achieved white crystalline amide.
 - Then filter and crystalline with water recrystalline with alcohol.
 - MP of $\text{R}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}_2$ is 220°C .

object :-

separate and identify the organic compounds from given mixture and also form suitable derivative.

Test for solid compounds :-**primary test :-**

physical state - solid crystals

colour - colourless or pink

odour - odourless

solubility - soluble in water

observation table :-

Experiment	Observation	Inference
1.) Flame test :- on burning the organic cpd in burners	compound burns with smoky	Compound is aromatic
2.) Litmus test :- Red/blue litmus paper take in the aqueous solution of the organic mixture.	No change in colour	Compound is neutral
3.) Element test :-		
(a) "N"-test :- take L.S sol ⁿ + freshly prepared FeSO ₄ sol ⁿ + conc. H ₂ SO ₄	Green/blue coloured ppt sol ⁿ is not obtained	"N"- absent

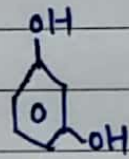
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c) "X" Test:- L.S. solution + conc. HNO ₃ + AgNO ₃	white ppt is not obtained	"X" - absent
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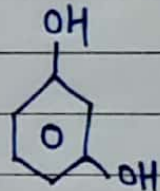
Functional group test:-

solution of organic mix. + sodium metal	Hg gas is evolved	-OH group may be
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ceric ammonium nitrate test:- organic mix. + 2-4 drop ceric	Red colour salt is obtained	-OH group confirm.
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MP of organic compound	≈ 110°C	 may be
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Conformative test for functional group:-

Aqueous solution of organic mixture + FeCl ₃ + ammonium nitrate	blue violet salt ppt is obtained	 confirm
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BP of organic compound	≈ 78°C	C ₂ H ₅ OH may be
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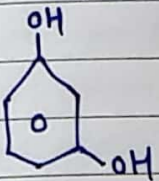
Determination test for functional group:-

Iodoform test:- 1ml mix. + 2ml I ₂ sol ⁿ + NaOH drop $\xrightarrow{\Delta}$ (gradually)	yellow coloured salt ppt is obtained	C ₂ H ₅ OH confirms
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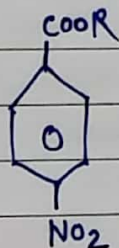
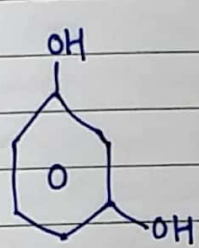
Result :-

The solid and liquid compound are



and C_2H_5OH respectively in given binary organic mixture.

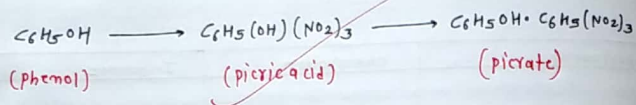
Derivative :-



MP = 182°C

Experiment	Observation	Inference
1) Flame test - on burning the organic solid on burners flame	Compound burns without smoky flame	Compound is aliphatic
2) Litmus test:- Red/blue litmus paper take in the aqueous sol ⁿ of organic compounds	colour of red & blue litmus does not change	Nature of compound is neutral.
3) Element test:- (a) "N"-test:- Take L.S solution + Freshly prepared FeSO ₄ sol ⁿ + conc H ₂ SO ₄	green/blue colour sol ⁿ ppt is not obtained	"N"-absent
(b) "S"-test:- Take L.S sol ⁿ + sodium nitroprusside	black ppt / sol ⁿ is not obtained	"S"-absent
(c) "X"-test:- L.S salt + conc. HNO ₃ + AgNO ₂	White ppt is not obtained	"X"-absent
Test for functional group:- salt of organic mix. + sodium metal	H ₂ gas evolved	-OH group may be
ceric ammonium nitrate test:- mix. + 2-7 drop ceric ammonium nitrate	Red colour sol ⁿ is obtained	-OH group confirms

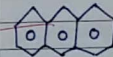
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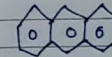
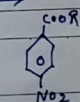
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3) Element test:-			
(a)	"N"-test - take L/s salt + freshly prepared FeSO ₄ sol ⁿ + conc. H ₂ SO ₄	green/blue coloured ppt/ sol ⁿ is not obtained	"N"- absent
(b)	"S"-test:- take L/s salt + sodium nitro prusside	black ppt/ salt is not obtained	"S"- absent
(c)	"X"-test:- take L/s salt + conc. HNO ₃ + AgNO ₃	white ppt is not obtained	"X"- absent

Test for functional group:-

compound + AlCl ₃ + Δ + CHCl ₃	Red colour is obtained	Aromatic hydrocarbon confirm
MP of organic compound	≈ 216°C	 may be

Confirmative test for functional group:-

organic mixture in saturated salt of picric acid in benzene + Δ	Red colour ppt/ salt is obtained	 confirms
C ₂ H ₅ OH → 	MP = 57°C	Teacher's Signature: _____

Object:-

separate and identify the organic compounds from given mixture and also form the suitable derivatives.

Test for solid compound:-

primary test:-

physical state - solid crystalline

colour - colourless or yellow

odour - specific odour

solubility - water insoluble but soluble in organic solution

observation table:-

S.No.	Experiment	observation	Inference
1.)	Flame test:- on burning the organic solid in burner flame	compound burns with smoky flame	compound is aromatic
2.)	litmus test:- Red and blue litmus paper take in aqueous solution of organic compounds.	No change in colour	compound compound is neutral

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Formation of derivative :-

melting point 138°C .
 derivative is picrate having

- phenol react with picric acid and is formed picrate.

Method :-

Take 0.5 gm mixture and add in 1ml of benzene and add 2ml picric acid and stirring it well.

- know we obtain yellow crystals and crystalline it with benzene.

Test for liquid compounds :-

primary test :-

physical state - liquid

colour - colourless

odour - sharp odour

solubility - soluble in water

Observation table :-

S.No.	Experiment	observation	Inference
1)	Flame test:- on burning of organic solid in burner's flame	compound burns without smoky flame	compound is aliphatic

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2.) Litmus test:-

Red / blue litmus papers
take in the aqueous
solution of organic

colour red / blue
litmus paper
coloured not change

nature of epd is
neutral

3.) Element test:-

(a) "N"-test:-

Take L.S solution + Freshly
prepared FeSO_4 solⁿ +
conc. H_2SO_4

green / blue ppt/
solⁿ is not
obtained

"N"- absent

(b) "S"-test:-

Take L.S solution + 2ml
sodium nitroprusside

black ppt / salt is
not obtained

"S"- absent

(c) "X"-test:-

L.S solution + conc. HNO_3
+ AgNO_3

white ppt is not
obtained

"X"- absent

Test for functional group:-

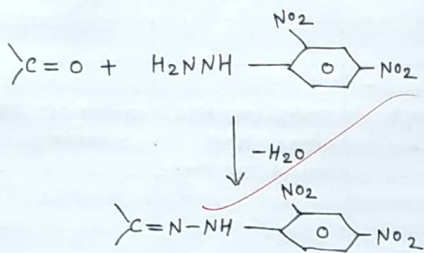
Sodium bisulphite test -

Compound + 2ml NaHSO_3

white ppt is
obtained

-C- group
may be

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2,4-dinitrophenyl hydrazone test -

Alcoholic salt of organic mixture + 1ml 2,4-dinitrophenyl hydrazene + Δ	yellow/orange sol ⁿ /ppt is obtained	-C- group confirm
Boiling point of organic compound	$\approx 20^\circ C$	CH ₃ -C-H O may be

confirmative test for functional group :-

Iodoform test :-

1ml organic mixture + 2ml of organic I ₂ solution + NaOH + Δ	yellow ppt / sol ⁿ is obtained	CH ₃ -C-H confirm
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Formation of derivatives :-

Derivative is of 2,4-dinitrophenyl hydrazone melting point is 168°C

- Aldehyde and keto react with 2,4-dinitrophenyl hydrazone with formed 2,4-dinitrophenyl hydrazone.

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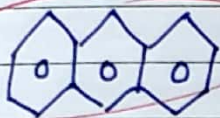
Method :-

Take 1ml compound and 4-5ml 2,4-dinitro phenyl ~~hydrazene~~ hydrazene.

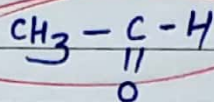
- Add 2-drop conc. H_2SO_4 + heat
- After cool the mixture we obtain yellow coloured ppt and crystalline with alcohol recrystalline is with benzene.

Result :-

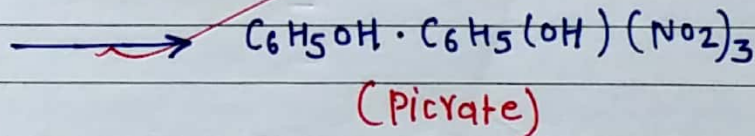
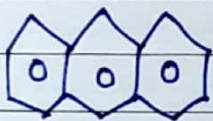
The solid and liquid compound are



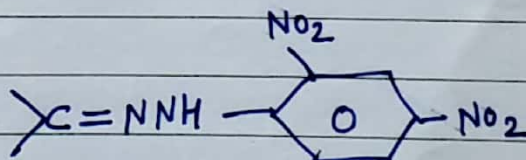
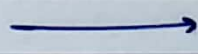
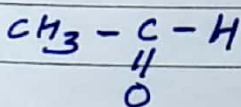
and



respectively in given binary organic mixture.

Derivative :-

Melting point - $138^\circ C$



Melting point = $168^\circ C$

Suvenil
23/4/18

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