# 12. Aldehydes, Ketones and Carboxylic Acids

#### Section-I: Aldehydes and Ketones

#### 12.1 INTRODUCTION

Aldehydes and ketones are carbonyl compounds, as they contain carbonyl (>C=O) group. General formula of aldehydes and ketones is  $C_nH_{2n}O$ .

#### **Difinitions:**

**Aldehydes:** These are first oxidative products of primary alcohols, in which carbonyl carbon atom is attached to, at least one H- atom. These are represented as R-CHO.

e.g. H-CHO CH<sub>3</sub> – CHO formaldehyde acetaldehyde

**Ketones:** These are first oxidative products of secondary alcohols, in which carbonyl carbon atom is attached to two same or different alkyl groups. These are represented as, R-CO-R.

e.g. CH<sub>3</sub> -CO-CH<sub>3</sub> CH<sub>3</sub> -CO-C<sub>2</sub>H<sub>5</sub> acetone ethyl methyl ketone

#### 12.2 ISOMERISM IN ALIPHATIC ALDEHYDFS AND KETONES:

Aldehydes: They shows chain. tautomerism, optical isomerism themselves and functional isomerism with ketones.

Ketones: They shows chain, tautomerism, optical isomerism, metamerism themselves and functional isomersism with aldehydes.

e.g. i)  $C_3H_6O$ 

CH,CH,CHO propanal

CH,COCH, propanone

 $C_{4}H_{5}O$ 

CH, CH, CH, CHO butanal a)

CH<sub>2</sub>CH(CH<sub>2</sub>)CHO 2-methyl propanal b)

CH,COCH,CH, 2-butanone c)

 $C_5H_{10}O$ 

CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CHO pentanal a)

CH, CH, CH(CH, )CHO 2-methyl butanal b)

CH,CH(CH,)CH,CHO 3-methyl butanal c)

(CH,),CCHO 2,2—dimethyl propanal d)

CH, CH, COCH, CH, 3-pentanone e)

CH, COCH, CH, CH, 2-pentanone

 $CH_3COCH(CH_3)$ , 3-methyl 2-bu tanone **Metamerism:** 

Metamerism is present in same class of compounds. It is shown by the compounds which have different alkyl groups( nature or type) attached to either side of polyvalent functional group.

e.g. pentan-2-one and pentan-3-one are me tamers, not position isomers.

CH, CH, COCH, CH, diethyl ketone

CH, COCH, CH, CH, methyl n-propyl ketone pentan-2-one and 3-methyl butan-2-one are me tamers not chain isomers

> CH<sub>3</sub>COCH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub> 2-pentanone CH<sub>3</sub>COCH(CH<sub>3</sub>), 3-methyl, 2-butanone

Note: Metarners are either chain or position isomers. If metamerism is mentioned then never write chain or positon isomerism.

#### **Tautomerism:**

**Definition:** It is one type of functional isomerism, in which a single compound exists in two or more readily interconvertible structures that differ in the relative position hydrogen.

Only  $\alpha$  –H atom involved in tautomerism

e.g. Acetone and prop-1-en-2-ol are tautomers

$$\begin{array}{ccc}
O & OH \\
\parallel & \parallel \\
CH, -C-CH, & \Longrightarrow CH, -C-CH,
\end{array}$$

#### 12.3 STRUCTURE OF CARBONYL GROUP

- Carbonyl group (> C = 0) consist of one sigma and one  $\pi$  bond.
- In carbonyl group carbonyl carbon atom is in sp<sup>2</sup> 2) hybridiscd state.
- The sigma bond is formed by sp<sup>2</sup> hybrid orbital of carbon and p-orbital of oxygen i.e. sp<sup>2</sup>-p overlapping.
- The  $\pi$  bond is formed by sideways overlapping of unhybridised p-orbital of carbon and p-orbital of oxygen i.e. p-p overlapping.
- Carbonyl carbon atom is joined to three atoms of sigma bond making a bond angle 120°, therefore,

carbonyl group has trigonal planer structure.

- 6) In carbonyl group, oxygen is more electronegative than carbon hence oxygen carries partial negative charge (nucleophilic or Lewis base centre) and carbon carries partial positive charge (electrophilic or Lewis acid centre).
- 7) The high polarity of carbonyl group is explained on the basis of resonance structure.

$$^{\prime}$$
C = O  $\longleftrightarrow$  >  $\overset{+}{C}$   $-\overline{O}$ 

neutral dipolar structure structure

## 12.4 PREPARATION OF ALDEHYDES AND KETONES

1. By oxidation of primary and secondary alcohols by using pyridinium chloro chromate( PCC) or pyridinium dichromate (PDC):

It oxidises only primary alcohol to aldehyde ane secondary alcohol to ketones. PCC is the mixture pyridine, CrO<sub>3</sub> and HCl in 1:1:1 ratio. This reagent does not attack >C=C< bond. It is also known as Collin's reagent.

e.g. 
$$CH_3 - OH + [O] \xrightarrow{PCC} H-CHO + H_2O$$

- 2. By dehydrogenaion of primary and secondary alcohols by using Cu or Fe or Ag metals at 573 K.
- a) Vapours of primary alcohols are passed over Cu or Fe metal at 573 K undergo dehydrogenation gives aldehydes. These are a–a elimination reactions.

$$-\text{CH}_2\text{OH} \xrightarrow{\text{Cu}/573\text{K}} -\text{CHO} + \text{H}_2$$

b) Vapours of secondary alcohols are passed over Cu or Fe metal at 573 K undergo dehydrogenation gives ketones.

$$>$$
CHOH  $\xrightarrow{Cu/573K}$   $>$ C = O + H<sub>2</sub>

**Note:** Tertiary alcohols are heated with copper metal at 573 K undergoes dehydration to give alkenes and water.

- 3. From hydrocarbon
- a) Ozonolysis of alkene:

$$\begin{array}{c} \mathrm{CH_2} = \mathrm{CH_2} + \mathrm{O_3} & \longrightarrow & \mathrm{CH_2} - \mathrm{O} - \mathrm{CH_2} \\ \text{ethylene} & & | & | \\ \mathrm{O} & \longrightarrow & \mathrm{O} \\ \text{ethylene ozonide} \end{array}$$

$$\begin{array}{ccc}
CH_2 - O - CH_2 &+ H_2O & \xrightarrow{Z_n} \\
 & & & \\
O - & & O
\end{array}$$

b) Hydration of alkyne

$$CH_{2} \equiv CH_{2} + H_{2}O \xrightarrow{40\%H_{2}SO_{4}} \rightarrow$$

$$CH_{2} = CHOH \xrightarrow{rearrangement} CH_{3} - CHO$$
vinyl alcohol acetaldehyde

- 4. From dry distillation (pyrolysis or decarboxylation) of calcium salt of acids
- a) Aldehydes:

$$(H-COO)_2Ca \xrightarrow{dry \text{ distilled}} H-CHO + CaCO_3$$
  
cal. formate formaldehyde

b) Ketones:

e.g. 
$$(C_2H_5-COO)_2Ca+(CH_3-COO)_2Ca$$
  
cal.propionate cal.acetate  $\xrightarrow{\text{dry distilled}} 2C_2H_5-CO-CH_3 + 2CaCO_3$   
ethyl methyl ketone

5. From alkaline hydrolysis of geminal dihalide:

The geminal dihalides are boiled with aqueous KOH or NaOH. It produces aldehydes and ketones.

$$\begin{array}{cccc}
 & & \downarrow & & \downarrow & \\
-C - X + 2KOH & \xrightarrow{\Delta} & -C - OH + 2KX & \\
\downarrow & & \downarrow & & \downarrow & \\
X & & & OH
\end{array}$$

geminal dihalide

unstable

$$\begin{array}{c} | \\ -C - OH \end{array} \longrightarrow > C = O + H_2O$$
 $\begin{array}{c} | \\ OH \end{array}$ 

#### 12.5 PREPARATION OF ONLY ALDEHYDES

1. Rosenmund reduction (Reduction of acyl chloride): Acid chloride is reduced by Lindlar catalyst to give aldehyde. Function of BaSO<sub>4</sub> is to poison the catalyst at aldehyde stage.

$$\begin{array}{c} O \\ R-C-Cl+H_2 \xrightarrow{Pd-BaSO_4} R-C-H+HCl \\ e.g. \ CH_3-COCl+H_2 \xrightarrow{Pd-BaSO_4} \\ acetyl chloride \end{array}$$

CH<sub>3</sub>COOH + HCl acetaldehyde

2] Stephen's reduction (Reduction of cyanide by SnCl, + HCI)

$$R-C \equiv N + 2 [H] \xrightarrow{SnCl_2 + HCl} R - CH = NH$$
  
imine

$$\xrightarrow{\text{H}_2\text{O}}$$
 R-CHO + NH<sub>3</sub>

e.g. 
$$CH_3-C \equiv N + 2[H] \xrightarrow{SnCl_2 + HCl}$$

$$CH_3-CH = NH \xrightarrow{H_2O} CH_3-CHO + NH_3$$
acetaldimine acetaldehyde

3] Reduction of cyanide by di-isobutyl aluminium hydride (iso-but)<sub>2</sub>-Al-H or [(CH<sub>3</sub>)<sub>2</sub>CHCH<sub>2</sub>]<sub>2</sub> Al-H or Di-iso-but-Al-H

$$R-C \equiv N + 2 [H] \xrightarrow{\text{(iso-but)}_2AI-H} R-CH = NH$$
 imine

$$\xrightarrow{\text{H}_2\text{O}}$$
 R-CHO + NH<sub>3</sub>

e. g. 
$$CH_3 - C \equiv N + 2[H] \xrightarrow{\text{(iso-but)}_2Al-H} \rightarrow$$

$$CH_3 - CH = NH \xrightarrow{H_2O} CH_3 - CHO + NH_3$$
acetaldimine acetaldehyde

4] Reduction of ester by (Iso-but),- Al-H

$$R - C - OR' + 2 [H] \xrightarrow{\text{(iso-but)}_2A1-H} \text{dry ether}$$

$$R - C - H + R' - OH$$

e.g.

$$\begin{array}{c} O\\ \parallel\\ CH_3-(CH_2)_9-C\\ -OC_2H_5+2[H] \xrightarrow{(iso-but)_2A1-H} \\ \text{ethyl undecanoate} \end{array}$$

$$CH_3 - (CH_2)_9 - C -H + C_2H_5OH$$
  
undecanal

5 From hydrocarbon :

Aromatic aldehydes and its derivatives are prepared from aromatic hydrocarban by following methods.

Strong oxidising agent oxidises hydrocarbon in to acids. It is possible to stop the oxidation at aldehyde stage using suitable reagent, that convert

methyl group into intermediate which is difficult to oxidised further.

i) Etard reaction (oxidation of toulene by chromyl chloride (CrO,Cl,)

When toulene is oxidised by  $CrO_2Cl_2$  first give chromium complex which on acid hydrolysis gives benzaldehyde

$$CH_3 + 2CrO_2Cl_2 \xrightarrow{CS_2}$$

$$CH \xrightarrow{OCrOHCl_2} \xrightarrow{H_3O^+} CHO$$

$$benzaldehyde$$

$$+ 2CrO_2Cl_2 + 2H_2$$

ii) Oxidation of toluene by CrO<sub>3</sub> (chromium oxide) in acetic anhydride

$$CH_{3} + 2[O] + (CH_{3}CO)_{2}O \xrightarrow{CrO_{3}/273k} -H_{2}O$$

$$CH \xrightarrow{OCOCH_{3}} \xrightarrow{H_{3}O^{+}} CHO$$

$$CH \xrightarrow{OCOCH_{3}} CHO$$

benzylidene diacetate

+2 CH<sub>3</sub>COOH

iii) By side chain chlorination followed by hydrolysis (commercial method).

$$CH_{3} + Cl_{2} \xrightarrow{UVL}$$

$$CHCl_{2} + 2KOH \xrightarrow{hydrolysis} CHO$$

$$+ 2KCl + 2H_{2}O$$

iv) By Gatterman - Koch reaction: In this reaction - H in benzene ring is converted into - CHO

#### 12.6 PREPARATIO OF ONLY KETONES:

From acyl chloride and dialkyl cadmium:
 Recently dialkyl cadmium is used for preparation of ketones from acyl halide in the presence of

RMgX. Since ketone formed in reaction do not react further to preduce 3° alcohol. It may be noted that R–MgX can not be used in place of dialkyl cadmium, this is because ketone formed further react with R–MgX to give 3° alcohol.

Dialkyl cadmium is obtained by reacting R–MgX with cadmium chloride

$$2R - MgX + CdCl_2 \longrightarrow R_2Cd + 2MgXCl$$

When acyl halide reacts with R<sub>2</sub>Cd in the presence of dry ether gives ketones.

$$2R - COX + R_2Cd \xrightarrow{dry \text{ ether}} 2R - COR + CdX_2$$
e.g. 
$$2CH_3COCl + (CH_3)_2 Cd \xrightarrow{dry \text{ ether}} 2CH_3COCH_3 + CdCl_3$$

2. From cyanide or nitrile and RMgX: When alkyl cyanide is reacted with RMgX in the presence of dry ether, gives addition product, which on acid hydrolysis, gives ketone

$$R-C = N + R-MgX \xrightarrow{dry \text{ ether}} R - C = N - MgX$$
alkane nitrile
$$Mg \text{ complex}$$

$$R = \frac{R}{R - C} = N - MgX + 2H_2O \xrightarrow{H^+} R - CO - R + NH_3 + MgXOH$$
ketone

3. From Friedel – Craft acylation reaction:

$$\bigcirc + R - C - Cl \xrightarrow{Anhy. AlCl_3} \bigcirc C - R + HCl$$

alkyl phenyl ketone

#### 12.7 CHEMICAL PROPERTIES:

## Mechanism in Nucleophilic addition reactions:

The carbonyl group has two active centre which can be easily attacked. The positively changed carbon is easily attacked by nucleophile and negatively charged oxygen is easily attacked by electrophile.

During addition reaction nucleophile attack first and followed by electrophile, because anion produced is more stable than cation.

During the nucleophilic addition reaction hybridisation of carbonyl carbon atom changes

from  $sp^2$  to  $sp^3$ .

carbonyl group

tetrahedral alkoxideanion

additionproduct

**Note:** If attacking nucleophile is weak i.e. ammonia and its derivatives the reaction is carried in weakly acidic medium.

Reactivity: Aldehydes are more reactive than ketone in nucleophilic addition reaction due to steric and electronic reasons (inductive effect). Aliphatic aldehydes and ketones are more reactive than aromatic aldehydes and ketones

i) Inductive effect: Reactivity of carbonyl group depends upon magnitude of positive charge on carbonyl carbon atom, more the positive charge on carbonyl carbon atom more will be the reactivity of carbonyl compounds.

Electron donating group decreases the magnitude of positive charge on carbonyl carbon atom, hence reactivity decreases. Electron withdrawing groups increases the reactivity of carbonyl compounds by increasing positive charge on carbonyl carbon atom.

$$+1$$
  $\delta + \delta -$   
 $C = 0$ 

positive charge on carbonyl carbon atom decreases

$$-1 + \delta + \delta - C = 0$$

positive charge on carbonyl carbon atom increases.

Thus reactivity is

$$\begin{split} \text{H-CHO} > \text{CH}_3 - \text{CHO} > \text{C}_2\text{H}_5 - \text{CHO} > \\ \text{CH}_3\text{COCH}_3 > \text{C}_6\text{H}_5\text{CHO} > \text{C}_6\text{H}_5\text{COC}_6\text{H}_5 \\ \text{CCl}_3 - \text{CHO} > \text{CHCl}_2 - \text{CHO} > \text{CH}_2\text{Cl-CHO} > \\ \text{CH}_3 - \text{CHO} \end{split}$$

ii) Steric effect: In aldehydes only one alkyl group is attached to carbonyl carbon atom hence attack is relatively more easy. In ketones two alkyl groups are attached to carbonyl carbon atom. Which hinders the approach of nucleophile, hence attack, is relatively difficult. This factor is also called as steric hinderence. In other word, more the steirc hinderence less the reactivity of carbonyl compounds.

$$CH_{3}CHO > CH_{3} - C - CH_{3} > CH_{3} - C - C_{2}H_{5} >$$

$$C_{2}H_{5} - C - C_{3}H_{5} = C - C_{5}H_{5}$$

- Il Addition reactions:
- 1. Addition of HCN (Prude add): When aldehydes and ketones are reacted with HCN in presence of base gives corresponding cyanohydrins. In this reaction oxygen atom in carbonyl group is changed to OH group.

Practically, HCN need for the reaction is generated in situ from dil. HCl and NaCN. The amount of acid is insufficient to react whole NaCN. Hence, solution remain alkaline and which catalysed the reaction.

$$-C = O + HCN \xrightarrow{base} -C - OH$$

carbonyl compounds cyanohydrine

2. Addition of sodium bisulphite (NaHSO<sub>3</sub>): When aldehydes and ketones are reacted with 40 % aqueous saturated solution of NaHSO<sub>3</sub>, gives corresponding crystalline bisulphite addition compounds. The reaction involves the nucleophilic attack of sulphite ion (SO3<sup>--</sup>).

carbonyl compounds

#### Note:

- i) Some ketones like diethyl ketone, acetophenone do not undergo this reaction.
- ii) The bisulphite addition product of aldehyde and ketone are crystalline solid and therefore this reaction is used in the separation of aldehydes and ketones from non carbonyl compounds.
- iii) As the reaction is reversible the addition product can be dissolved by dilute mineral acid or aqueous alkalies to regenerate original aldehydes and ketones. Hence this reaction used to purification of aldehydes and ketones.
- 3. Addition of RMgX: When carbonyl compounds are reacted with Grignards reagent in the presence of dry ether give addition product which

on acid hydrolysis gives alcohols (may be  $1^{\circ}$  or  $2^{\circ}$  or  $3^{\circ}$  alcohols).

$$\begin{array}{c}
-C - O - MgX \\
\mid R \\
+ H_2O \xrightarrow{H^+} \\
-C - OH - MgXOH \\
\mid R
\end{array}$$

a) On formaldehyde: (In this reaction > C = O group is converted into -CH<sub>2</sub>OH primary alcohol)

$$\begin{array}{c} H \\ H - C = O + RMgX \xrightarrow{dry \text{ ether}} \end{array}$$

$$\begin{array}{c} H \\ H - C - O - MgX \\ R \\ Mg-complex \end{array}$$

$$\begin{array}{c} H \\ R \\ Mg-complex \end{array}$$

$$\begin{array}{c} H \\ H - C - O - MgX + H_2O \xrightarrow{H^+} \end{array}$$

Mg–complex  $R-CH_s-OH+Mg$ 

R-CH<sub>2</sub> - OH + MgXOH primary alcohol

b) On aldehyde (RCHO): When aldehydes (except HCHO) are reacted with RMgX in the presence of dry ether, gives Mg complex. Which on acid hydrolysis, gives secondary alcohols.
 (In this reaction >C=O group is converted into >CH-OH sec. alcohol.)

$$\begin{array}{c} H \\ R-C=O+RMgX & \xrightarrow{dry \text{ ether}} \end{array}$$

$$\begin{array}{c} H \\ H-C & -O-MgX \\ R \\ \\ Mg-complex \end{array}$$

$$\begin{array}{c} H \\ | \\ H - C \\ | \\ R \end{array} - O - MgX + H_2O \xrightarrow{H^+}$$

secondary alcohol

c) On ketone: When ketones are reacted with RMgX in the presence of dry ether, gives Mg complex. Which on acid hydrolysis, gives tertiary alcohols.

(In this reaction >C=O group is converted into C-OH tertiary alcohol.)

$$\begin{array}{c} R \\ | \\ R - C \end{array} = O + RMgX \xrightarrow{dry \text{ ether}}$$

Mg-complex

$$\begin{array}{c|c} R & \\ R - C & - O - MgX + H_2O \xrightarrow{H^+} \\ R & \end{array}$$

**Note:** Using RMgX carbonyl compounds are convert into following.

- 1) Formaldehyde into -CH<sub>2</sub>-OH (primary alcohols)
- 2) Aldehyde into > CH–OH (secondary alcohols)
- 3) Ketone into C–OH (tertiary alcohols)
- **4. Addition of R–OH:** Aldehydes or ketones are reacts with alcohols to form acetal or ketals. The acetal is used as table sugar, wooden shop, cotton fibrics etc
- a) Aldehyde: Aldehyde reacts with alcohol in the presence of presence of dry HCl gas to give alkoxy alcohol (hemiacetal) or half acetal, which reacts with another molecule to form acetal.

$$R \to C = O + R - OH \xrightarrow{dry HCl} R \to C - OH$$

alkoxy alcohol (hemiacetal) or half acetal

b) Ketones: Ketones reacts with alcohol in the presence of presence of dry HCl gas to give hemiketal or half ketal, which reacts with another molecule to form ketal.

$$R - C = O + R - OH \xrightarrow{\text{dry HCl}}$$
ketone
$$R = R$$

$$\begin{array}{c|c} R & R \\ | & R - C - OR \end{array} \xrightarrow[R \ OH]{} R-OH & R - C - OR + H_2O \\ | & OR & OR \end{array}$$

hemiketal (half ketal) ketal

#### Note:

- i) Dry hydrogen chloride protonates the oxygen of carbonyl compound and increase the electrophilicity of carbonyl carbon atom and increase the rate of reaction.
- ii) Acetal and ketal are hydrolysed by ag. HCl to give corresponding aldehydes and ketones.
- **5.** Addition of ammonia: Ammonia reacts differently with formaldehyde, acetaldehyde and acetone forming.

e.g.

i) 
$$CH_3 - C = O + NH_3 \xrightarrow{dry \text{ ether}}$$
 acetaldehyde

$$\begin{array}{c} H \\ | \\ CH_3 - C - OH \\ | \\ NH_2 \\ \text{acetaldehyde amine} \end{array} \xrightarrow{-H_2O} CH_3 - CH = NH \\ \text{aldimine} \end{array}$$

$$3CH_{3}-CH=NH\xrightarrow{trimerisation} CH_{3}-CH$$

$$CH_{3}-CH$$

aldimine

ii) 6HCHO + 4NH<sub>3</sub>  $\xrightarrow{\text{dry ether}} \Delta$  methanal

$$(CH_2)_6N_4 + 6H_2O$$
  
urotropine

**Note:** Urotropine is condensation product of methanal and ammonia, it is not addition product.

$$\begin{array}{ccc} & CH_3 & O \\ & & | \\ iii) & CH_3 - C & = O + H_2 - CH - C & - CH_3 \\ & \text{acetone} \end{array}$$

$$\xrightarrow{\text{NH}_3/\Delta} \begin{array}{c} \text{CH}_3 & \text{O} \\ & \parallel \\ \text{CH}_3 - \text{C} & = \text{CH} - \text{C} - \text{CH}_3 + \text{H}_2\text{O} \\ \text{mesityl oxide} \end{array}$$

$$\begin{array}{ccc}
CH_3 & O \\
& \parallel \\
CH_3 - C & = CH - C - CH_3 + NH_3 & \xrightarrow{M.R.}
\end{array}$$

$$\begin{array}{c|cccc} \mathbf{CH_3} & \mathbf{CH_3} & \mathbf{O} & \mathbf{O} \\ \mathbf{CH_3} - \mathbf{C} & -\mathbf{CH_2} - \mathbf{C} & -\mathbf{CH_3} \\ \mathbf{NH_2} & \mathbf{NH_2} \end{array}$$

diacetone amine

II] CONDENSATION REACTIONS OR ADDITION FOLLOWED BY ELIMINATION OR REPLACEMENT OXYGEN:

$$> C = O + H_2N - Z \xrightarrow{acid} -C - OH$$
 $NHZ$ 

$$\longrightarrow > C = N - Z + H_2O$$
Where  $Z = R-$ ,  $C_6H_5-$ ,  $-OH$ ,  $-NH_2$ ,  $C_6H_5NH-$ ,

$$O_2N$$
 $-HN-NO_2$  (2,4-DNP),  $-NH-CONH_2$ 

(semicarbazide).

1. Reaction with R-NH<sub>2</sub>:

$$> C = O + R - NH_2 \xrightarrow{\text{acid}} > C = NR + H_2O$$

substituted imine

2. Reaction with Ph-NH,

$$\begin{array}{c} -\stackrel{|}{C}=O+H_2N \xrightarrow{acid} \\ -\stackrel{|}{C}=N \xrightarrow{acid} +H_2O \end{array}$$

phenyl imine (Schiffs base)

3. Reaction with hydroxyl amine:

$$-\overset{\mid}{C} = O + NH_2 - OH \xrightarrow{\text{acid}} -\overset{\mid}{C} = NOH + H_2O$$
oxime

4. Reaction with hydrazine: When aldehydes and ketones are heated with hydrazine in the presence of dilute acid undergoes condensation, gives corresponding crystalline red or orange colour hydrazone.

In this reaction >C = O group is converted into >C = NNH<sub>2</sub>.

$$-\overset{\mid}{C} = O + H_2 N - NH_2 \xrightarrow{\text{acid}} -\overset{\mid}{C} = N \cdot NH_2$$
hydrozone
$$+ H_2 O$$

5. Reaction with phenyl hydrazine: When aldehydes and ketones are heated with phenyl hydrazine in the presence of dilute acid undergoes condensation, gives corresponding crystalline red or orange colour phenyl hydrazone. In this reaction > C = O group is converted into > C = NNHC<sub>6</sub>H<sub>5</sub>.

$$\begin{array}{c} \stackrel{|}{-}\mathrm{C} = \mathrm{O} \ + \mathrm{H_2N} - \mathrm{NHC_6H_5} & \stackrel{\mathrm{acid}}{\longrightarrow} \\ & \stackrel{|}{-}\mathrm{C} = \ \mathrm{N} \cdot \mathrm{NHC_6H_5} \\ & \mathrm{phenyl} \ \mathrm{hydrazone} \end{array}$$

6. Reaction with H<sub>2</sub>NNH-CO-NH<sub>2</sub> (semicarbzide)

$$\begin{array}{c}
-C = O + H_2NNHCONH_2 \xrightarrow{\text{acid}} \\
-C = NNHCONH_2 \\
\text{semicarbazone}
\end{array}$$

7. Reaction with 2, 4-dinitrophenyl hydrazine (2,4-DNP)

$$\begin{array}{c|c}
 & NO_2 \\
-C = O + H_2N HN \\
\hline
 & NO_2 \\
-C = N HN \\
\hline
 & NO_2 \\
-NO_2 + H_2O \\
\hline
 & 2, 4, dinitrophenyl hydrozone (yellow)
\end{array}$$

#### **III] REDUCTION:**

Catalytic hydrogenation or reduction by using NaBH<sub>4</sub> or LiAlH<sub>4</sub> OR Na.Hg + H<sub>2</sub>O:

- 1. Catalytic hydrogenation: When aldehydes and ketones are reduced by passing hydrogen gas in the presence of Ni or Pt. or Pd. at 413K to 453K, gives 1° or 2° alcohols.
- a) Aldehydes on reduction gives primary alcohols.

$$\begin{array}{ccc} \text{RCHO} + \text{H}_2 & \xrightarrow{\text{Ni 413K}} & \text{R-CH}_2\text{OH} \\ & \text{aldehyde} & \text{primary alcohol} \end{array}$$

b) Ketones on reduction gives secondary alcohols.

$$\begin{array}{c} R \\ | \\ R-C = O+H_2 \end{array} \xrightarrow{\text{Ni 413K}} \begin{array}{c} R \\ | \\ C H-OH \end{array}$$
 ketones sec. alcohols

2. Reduction by using NaHg + H<sub>2</sub>O: When aldehydes and ketones are reduced by Na.Hg + H<sub>2</sub>O or ZnHg + CH<sub>3</sub>COOH or LiAIH<sub>4</sub>, NaBH<sub>4</sub> gives 1° or 2° alcohols. Note that NaBH<sub>4</sub> does not reduce ketone group.

$$Na.Hg + H_2O \longrightarrow NaOH + Hg + [H]$$

a) Aldehydes on reduction gives primary alcohols.

$$R-CHO + 2 [H] \xrightarrow{NaHg + H_2O} RCH_2OH$$
 aldehyde primary alcohol

b) Ketones on reduction gives secondary alcohols.

**Note:** Unsaturated aldehydes or ketones can be reduced to unsaturated alcohols without affecting >C=C< by using LiAIH<sub>4</sub> or NaBH<sub>4</sub>.

**3.** Clemmenson reduction (deoxygenation): In this reaction >C=O group is converted into >CH<sub>2</sub> (methylene) group.

e.g. When methanal is reduced by Zn. Hg + conc. HCl gives methane.

R-CHO + 4 [H] 
$$\xrightarrow{ZnHg + conc.HCl}$$
  $\xrightarrow{\Delta}$  R-H + H<sub>2</sub>O aldehyde alkane

4. Wolf – Kishner reduction: In this reaction > C = O group is reduced to > CH<sub>2</sub> group.

$$-C = O + NH_2 - NH_2 \xrightarrow{\text{acid}} -C = N \cdot NH_2 + H_2O$$
hydrazone

$$\begin{array}{c} \stackrel{|}{-C} = N \cdot NH_2 \xrightarrow{KOH + \text{ ethylene}} \stackrel{|}{-C}H_2 + N_2 \uparrow \end{array}$$

5. Reduction by Mg metal only ketone (Only for ketones): Only ketone give this reaction. When acetone is reacted with Mg metal in the presence of benzene gives Mg–complex, which on acid hydrolysis produces pinacol (2, 3–dimethyl butane 2, 3–diol).

$$(CH_3)_2CO$$
 + Mg Benzene + Mg

$$(CH_3)_2C-O$$
 $(CH_3)_2C-O$ 
 $(CH_3)_2C-OH$ 
 $(CH_3)_2C-OH$ 
 $(CH_3)_2C-OH$ 
 $(CH_3)_2C-OH$ 
pinacol

When pinacol is heated with acid undergoes rearrangement gives pinacolone. This is known as pinacol-pinacolone rearrangement.

$$(CH_3)_2C-OH \xrightarrow{Acid/\Delta} CH_3 - C - C - CH_3 + H_2O$$

$$(CH_3)_2C-OH \xrightarrow{pinacol} Dinacolone$$

#### IV| Oxidation reactions:

Oxidation by strong oxidising agent like K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> + dil. H<sub>2</sub>SO<sub>4</sub>

$$- \text{CHO} + [\text{O}] \xrightarrow{\text{K}_2\text{Cr}_2\text{O}_7 + \text{dil H}_2\text{SO}_4} \rightarrow -\text{COOH}$$
aldehyde acid

$$\begin{array}{c|c}
O \\
\parallel \\
-C -+[O] \xrightarrow{K_2Cr_2O_7 + \text{dil H}_2SO_4}
\end{array}$$

ketone

2. Oxidation by mild oxidising agents like Tollen's reagent, Fehling solution, Schiffs

'reagent. or Reducing properties of aldehydes or characteristic test of aldehydes or distinction test of aldehydes from ketones or reducing reaction of aldehydes.

Aldehydes and ketones are distinguished by Tollen's reagent, Fehling solution, Schiffs reagent.

i) Reaction with Tollens reagent:

R-CHO + 2[Ag (NH<sub>3</sub>)<sub>2</sub>]<sup>+</sup> OH<sup>-</sup> 
$$\xrightarrow{\Delta}$$
 aldehyde

$$RCOOH + 2Ag + 4NH_3 + H_2O$$
 acid

ii) Reaction with Fehling solution:

R-CHO + 2Cu<sup>++</sup> + 4OH<sup>-</sup> 
$$\xrightarrow{\Delta}$$
 aldehyde

R-COOH + 
$$Cu_2O \downarrow + 2H_2O$$
  
acid red ppt.

CHO + 
$$2Cu^{++}$$
 +  $4OH^{-}$   $\stackrel{\Delta}{\longrightarrow}$  No

reaction

benzaldehyde

iii) Reaction with Schiffs reagent: Schiffs reagent (colourless) is prepared by passing SO<sub>2</sub> gas in magenta or pink dye of para-rosaniline hydrochloride (Fuschin). When decolourised schiffs reagent is treated with aldehyde in cold original pink or magenta colour is developed.

R-CHO + Schiffs reagent 
$$\xrightarrow{\text{cold}}$$
 Pink

or magenta or bright crimson colour.

**Note:** Ketones does not give these reactions because of absence of H-atom on carbonyl carbon atom. Glucose, formic acid and enolic form of fructose gives above test.

3. Oxidation of methyl ketone by haloform reaction (Haloform test):

Haloform test means preparation of haloform i.e CHCl<sub>3</sub> (colourlss liquid), CHl<sub>3</sub> (yellow solid) and CHBr<sub>3</sub> (brown liquid). This test is given by some alcohols, some aldehydes and some ketones.

**Alcohols:** Alcohols that produces CH<sub>3</sub>–CO group (methyl ketone or 2–one group) on oxidation gives haloform test or oxidative product of alcohols contain 3–a–H atoms atoms on single carbon atom gives haloform test.

 $CH_3 - OH \xrightarrow{PCC} H$ -CHO Negative halo form

test

$$CH_3$$
- $CH_2$ -OH  $\xrightarrow{PCC}$   $CH_3$ -CHO Positive haloform test

**Aldehydes and ketones:** They contain CH<sub>3</sub>–CO group (methyl ketone or 2–one group) or 3–a–H atoms on single carbon atom.

$$\begin{array}{lll} \mbox{H-CHO} & \mbox{Negative haloform test} \\ \mbox{CH}_3\mbox{-CHO} & \mbox{Positive haloform test} \\ \mbox{CH}_3\mbox{-CH}_2\mbox{-CHO} & \mbox{Negative haloform test} \\ \mbox{CH}_3\mbox{-CH}_2\mbox{-CO-CH}_2\mbox{-CH}_3 & \mbox{Negative haloform} \\ \mbox{test} \end{array}$$

O
$$\parallel$$
R-C-CH<sub>3</sub> + 3NaOX  $\xrightarrow{\text{NaOH} + \text{X}_2}$ 
R-COONa + CHX<sub>3</sub> + 2NaOH
O
 $\parallel$ 

e.g. 
$$H-C-CH_3 + 3NaOC1 \xrightarrow{NaOH + Cl_2}$$

$$H-COONa + CHCl_3 + 2NaOH$$

**Note:** The importance of this reaction is, if >C=C< present in molecule is not attacked by NaOX.

$$CH_{3}-CH=CH-C-CH_{3}+3NaOC1 \xrightarrow{NaOH+Cl_{2}} CH_{3}-CH=CH-COONa+CHCl_{3}+2NaOH$$

- V Reaction due to  $\alpha$  –H atom
- 1) Acidity of  $\alpha$  –H atom in aldehydes and ketones:

Aldehydes and ketones undergoes number of reactions due to acidic nature of  $\alpha$  –H atom. The acidity of  $\alpha$  –H depends upon electron withdrawing group (–I effect) of carbonyl group and resonance stabilisation of conjugate base.

When  $\alpha$  –H atom is attached to only carbonyl group, then more acidic is  $\alpha$  –H atom.

$$\begin{matrix} O \\ \parallel \\ CH_3-C-H - \text{more acidic } \alpha-H \end{matrix}$$

When  $\alpha$  –H atom is attached to carbonyl group and alkyl group, then less acidic will be the  $\alpha$  –H atom.

$$$^{\text{O}}_{\text{}}$$$
  $$^{\alpha}_{\text{}}$$   $$^{\text{CH}}_3-\overset{\alpha}{\text{C}}$}$$   $+$  CH  $_3$  – C  $+$  H  $-$  less acidic  $\alpha$  –H

$$\begin{array}{ccc} O \\ \parallel \\ CH_3-C-\overset{\alpha}{C}H_3 \end{array} & - \mbox{more acidic } \alpha -\!H \end{array}$$

$$CH_3$$
  $C - CH_2 - CH_3$   $CH_3$   $CH_$ 

**Note:** In ald ol condensation reaction more acidic  $\alpha$  –H atom involved in condensation reaction.

- VI] Aldol condensation reaction or self condensation or reaction with dilute alkali or 10% alkali or weak alkali:
- Simple aldol condensation reaction: It is the reaction between two same aldehydes or two same ketones.
  - e.g. When acetaldehyde is warmed with dil. (10%) NaOH, gives acetaldol (3-hydroxy butanal).
  - e.g. Which on heating with acid loses water molecule and gives crotonaldehyde ( $\alpha \beta$  unsaturated aldehyde).

$$CH_3 - CH_2 - CH_2 - CH_2 - CH_2 - CH_3 -$$

 $_{2}H_{5}$   $-\overset{\parallel}{C}$  +  $\overset{\parallel}{H}$  -  $\overset{\parallel}{C}$   $-\overset{\parallel}{H}$  acetaldol

$$\begin{array}{c|c} OH & O \\ & | & | \\ CH_3 - C - CH_2 - C - H \\ & | & \xrightarrow{Acid} \\ H \end{array} \longrightarrow \begin{array}{c} Acid \\ heat \end{array}$$

$$O$$
 $\parallel$ 
 $CH_3 - CH = CH - C - H + H_2O$ 
crotonaldehyde (2-butenal)

2. Cross aldol condensa'tion reactions:

$$\begin{array}{c} O & O \\ \parallel & \parallel \\ C_2H_5-C+H-CH_2-C-H \end{array} \xrightarrow{\begin{array}{c} 20\% \text{NaOH} \\ \text{Warm} \end{array}} \\ \text{propanal} \end{array}$$

$$\begin{array}{c} O & O \\ \parallel & \parallel \\ C_2H_5-C+H-CH_2-C-H \\ \parallel \\ H \end{array}$$

3-hydroxy pentanal

$$\begin{array}{c} O & O \\ \parallel & \parallel \\ C_2H_5-C+H-CH_2-C-H \\ \parallel & \\ H \end{array} \xrightarrow{\Delta/H^+}$$

3-hydroxy pentanal

$$C_2H_5 - CH = CH - CHO + H_2O$$

Mechanism of aldol condensation reaction:

**Step-I:** Base  $OH^-$  takes  $\alpha$  -H-atom from aldehyde to form carbanion (enolate ion).

$$\begin{matrix} O & & O \\ \parallel & \parallel & \parallel \\ CH_3-C-H+OH^- & \longrightarrow CH_2-C-H+H_2O \end{matrix}$$

carbanion

**Step-II:** The carbanion attack electrophilic carbonyl carbon of second aldehyde molecule to form alkoxide ion.

$$CH_{3} - C - H + \overline{C}H_{2} - C - H \rightarrow$$

$$CH_{3} - C - CH_{2} - C - H$$

$$CH_{3} - C - CH_{2} - C - H$$

alkoxide ion

**Step-III:** Alkoxide ion takes hydrogen ion from water to form aldol. Base OH<sup>-</sup> ion is generated.

$$\begin{array}{c|c} \overline{O} & O \\ | & \parallel \\ CH_3 - C - CH_2 - C - H \\ | & + H - OH \end{array} \longrightarrow$$

$$\begin{array}{c|c} \text{OH} & \text{O} \\ & \parallel \\ \text{CH}_3 - \text{C} - \text{CH}_2 - \text{C} - \text{H} \\ & \parallel \\ \text{H} \end{array} + \text{OH}^-$$
 aldol

**Note:** Formaldehyde, benzaldehyde, trimethyl acetaldehyde, benzoquinone,chloral, iodal etc does not give this reaction because of absence of  $\alpha$  –hydrogen atoms.

#### VII] CANNIZZARO'S REACTION OR SELF REDOX REACTION OR BASE CATALYSED SELF OXIDATION REDUCTION REACTION:

**Condition:** It is a characteristics reaction of aldehydes and ketones which does not contain  $\alpha$  –H atom. It is carried by strong alkali.

e.g.

i) When formaldehyde is heated with cone, (50%) NaOH, gives sodium formate and methyl alcohol.

$$\begin{array}{ccc}
O & O \\
\parallel & \parallel \\
H-C & -H+H-C & -H+NaOH \longrightarrow
\end{array}$$

HCOONa + CH<sub>3</sub>-OH sod. formate methyl alcohol

ii) When benzaldehyde (benzene carbaldehyde) is heated with strong KOH, gives pot. benzoate and benzyl alcohol.

$$C_6H_5-C_9-H_5+C_6H_5-C_9-H_9+KOH_9$$
benzaldehyde (strong) cone.
$$C_6H_5COOK_9+C_6H_5-CH_2-OH_9$$
pot. benzoate benzyl alcohol

iii) Cross Cannizzaro's reaction: When benzaldehyde and formaldehyde are heated with strong NaOH e.g gives benzyl alcohol and sodium formate. In this reaction more reactive aldehydes give oxidative product in crossed Cannizzaro's reaction.

$$C_6H_5CHO + HCHO + NaOH \longrightarrow$$

$$C_6H_5CH_2OH + HCOONa$$
benzyl alcohol sod. formate

**Note :** Isobutyraldehyde gives Cannizzaro's reaction eventhough presence of  $\alpha$  –H atom. The exception behavior of isobutyraldehyde is due to +I effect of two alkyl groups.

e.g. 
$$(CH_3)_2CH$$
– $CHO + (CH_3)_2CH$ – $CHO + NaOH$  conc.

$$\longrightarrow$$
 (CH<sub>3</sub>(<sub>2</sub>CH – COONa + (CH<sub>3</sub>)<sub>2</sub>CH–CH<sub>2</sub>OH.  
sod. iso–butyrate iso–butyl alcohol

## VIII] ELECTROPHILIC SUBSTITUTION REACTION OF AROMATIC ALDEHYDES AND KETONES:

Note that -CHO and > C = O group are meta directing

#### 1. Halogenation:

i) 
$$CHO \rightarrow CHO \rightarrow C$$

m-chlorobenzaldehyde

ii) 
$$\longleftrightarrow$$
 + Br<sub>2</sub>  $\xrightarrow{\text{FeBr}_3}$   $\longleftrightarrow$   $\overset{\text{O}}{\longleftrightarrow}$  +HBr

m-bromoacetophenone

m-nitro acetophenone

#### 2. Nitration

i) 
$$+HNO_3$$
  $\xrightarrow{conc.H_2SO_4}$   $\xrightarrow{273K}$   $\xrightarrow{CHO}$   $+H_2O$   $\xrightarrow{NO_2}$   $\xrightarrow{m-nitro benzaldehyde}$   $+HNO_3$   $\xrightarrow{conc.H_2SO_4}$   $\xrightarrow{273K}$   $\xrightarrow{1- phenyl ethan-1-one}$   $\xrightarrow{COCH_3}$ 

#### 3. Suiphonation:

i) 
$$+H_2SO_4 \xrightarrow{Fuming} CHO$$

$$CHO CHO$$

$$SO_3H + H_2O$$
m-formyl benzene sulphonic acid

ii) 
$$\longleftrightarrow$$
 + H<sub>2</sub>SO<sub>4</sub>  $\xrightarrow{Fuming}$   $\xrightarrow{COCH_3}$   $\longleftrightarrow$  SO<sub>3</sub>H m-oxo methyl benzene sulphonic acid

#### Section-II: Carboxylic Acids

#### 12.1 INTRODUCTION

**Definition**: These are carboxyl derivatives of alkane obtained by replacing hydrogen atom from alkane by carboxyl (-COOH) group.

#### 12.2 ISOMERISM

Carboxylic acids: They shows chain, optical isomerism themselves and functional isomerism with esters. Total number of isomeric acids can be calculated by formula,  $I = 2^{n-3}$ .

Esters: They shows chain, metamerism, optical isomerism themselves and functional isomerism with carboxylic acids.

Total number of isomeric esters can be calculated by formula,  $I = 2^{n-1} - (n-2) - 2^{n-3}$ from C<sub>3</sub> onwards.

Total number of isomeric acids and esters can be calculated by formula,  $I = 2^{n-1} - (n-2)$ .

- e.g. i)  $C_1H_4O_2$ , (2-isomers)
  - a) CH, COOH acetic acid
  - methyl formate b) HCOOCH,
  - ii)  $C_3H_6O_2$  (3-isomers. 1 acid, 2 esters)
  - a) CH<sub>2</sub>CH<sub>2</sub>COOH propionic acid
  - HCOOC,H, ethyl formate
  - c) CH,COOCH, methyl acetate
  - iii)  $C_4H_5O_2$  (6-isomers. 2 acids, 4 esters) Acids-
  - a) CH,CH,CH,COOH n-butyric acid
  - (CH,),CHCOOH isobutyric acid b)

#### Esters-

b)

- a) HCOOCH, CH, CH, n-propyl formate
- HCOOCH(CH,), isopropyl formate
- CH,COOCH,CH, ethyl acetate c)
- C,H,COOCH, methyl propionate d)

#### 12.3 STRUCTURE OF -COOH GROUP

The carboxylic carbon is less electrophilic than carbonyl carbon atom in aldehydes and ketones because of resonance stabilisation of COOH group.

The carbon atom is sp<sup>2</sup> hybridised state. The O-C-O bond angle is 120°.

#### 12.4 PREPARATION METHODS

- From oxidation of 1° alcohols, aldehydes and ketones.
- From primary alcohols: Primary alcohols on a) controlled oxidation by acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> gives carboxylic acid.

$$-\text{CH}_2$$
 $-\text{OH} + 2[\text{O}] \xrightarrow{K_2\text{CrO}_2 + \text{dil. H}_2\text{SO}_4}$ 

COOH + H<sub>2</sub>O

From aldehydes: Aldehydes on oxidation by b) acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> gives carboxylic acid.

$$-CHO + 2 [O] \xrightarrow{K_2CrO_2 + dil. H_2SO_4} -COOH$$

From ketones: Ketones on controlled oxidation by acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> gives carboxylic acid.

>C = O + 2 [O] 
$$\xrightarrow{K_2CrO_2 + dil. H_2SO_4}$$
  $\rightarrow$   $-COOH + H_2O + CO_2$ 

From acid hydrolysis of cyanide or nitrile:

$$R-C \equiv N + H_2O \xrightarrow{H^+} R-CONH_2 \xrightarrow{H_2O}$$

alkyl cyanide

acid

Note: Mild reaction conditions are used to stop reaction at amide stage.

$$-C \equiv N + H - OH \xrightarrow{H^+} C = NH = NH$$
imine

$$\xrightarrow{\text{isomenst}} \begin{array}{c} O \\ \parallel \\ -C - NH_2 \end{array}$$

amide

From oxidation of alkyl benzene or phenyl alkene: Alkyl benzene are oxidised by strong oxidising agent like chromic acid or acidic KMnO<sub>4</sub> or alkaline KMnO<sub>4</sub>. The entire side chain is converted into carboxyl group irrespective of length of the side chain. Tertiary alkyl group donot

under goes oxidation because of absence of benzylic hydrogen atom.

e.g. 
$$CH_3 \xrightarrow{KMnO_4 + KOH}$$

$$\begin{array}{c}
COOK \\
\xrightarrow{\text{H}_3O^+}
\end{array}$$
benzoic acid

4. From dry ice and R-MgX:

$$O = C = O + RMgX \xrightarrow{\text{dry ether}} O = C - OMgX$$

$$\mid$$

$$R$$

dry ice

addition product

$$\begin{array}{c}
O = C - OMgX + H - OH \xrightarrow{H^+} \\
R
\end{array}$$

- 5. From hydrolysis of acid halide and acid anhydride:
- a) Acid halide is hydrolysed with water gives corresponding acids.

$$R-COCl + H-OH \longrightarrow R-COOH + HCl$$

**Note:** Acid chloride are more readily hydrolysed by aqueous base to give carboxylate ion, which on acidification gives corresponding acids.

$$R-COCI + OH^{-} \longrightarrow R-COO^{-} + HCI^{-}$$

$$R-COO^{-} \xrightarrow{H^{+}} R-COOH$$

b) Acid anhydride is hydrolysed by water gives acids

$$(R-CO)_2O + H_2O \longrightarrow 2R-COOH$$

6. Acid hydrolysis of esters by using dil. HCl or dil. H<sub>2</sub>SO<sub>4</sub>:

$$\begin{array}{c} \text{R-COO-R} + \text{H}_2\text{O} \xrightarrow{\text{cone. H}_2\text{SO}_4} \\ & \\ \text{ester} \end{array}$$

**Note:** These are reverse of esterification reaction. It is auto catalysed reaction because acid formed in this reaction acts as catalyst.

7. Alkaline hydrolysis of ester (Saponification):

R-COO-R + NaOH 
$$\stackrel{\Delta}{\longrightarrow}$$
 R-COONa + R-OH ester aq. sod. salt of acid alcohol R-COONa + H-OH  $\stackrel{H^+}{\longrightarrow}$  R-COOH + NaOH

#### Note:

- i) Acid hydrolysis and alkaline hydrolysis of esters are nucleophilic substitution reaction.
- ii) Hydrolysis of ester by dilute alkali is irreversible and faster reaction than acid hydrolysis.
- iii) Alkaline hydrolysis of ester is preferred over acid hydrolysis because the acid salt formed can not force backward reaction.
- iv) Reactions of ester with RMgX are nucleophilic addition reactions.
- 8. Oxidation of alkene by strong oxidising agent (vigrous oxidation):

$$>C=C$$
  $\xrightarrow{\text{vigrous}}$   $>C=O + O=C$   $>C=O + O=C$ 

e.g.

i) 
$$CH_2 = CH_2 + 4[O] \xrightarrow{KMnO_4 + \atop H_2SO_4 \Delta} 2H-COOH$$

ii) 
$$+4 [O] \xrightarrow{KMnO_4+} COOH$$
 cyclohexene adipic acid

#### 12.5 ACIDIC NATURE OF ALIPHATIC ACID

- Acidic nature of carboxylic acid can be best explained on the basis of resonance and Lowry Bronsted theory.
- 2. Carboxylic acid are weak acids and turns blue litmus to red. In water they are partially dissociated and equilibrium is shifted in left side. Thus, aqueous solution of carboxylic acid contains more undissociated acid molecules. Carboxylic acids are stronger acid than R–OH, C<sub>6</sub>H<sub>5</sub>–OH water and weaker than mineral acids.

$$\mathsf{RCOOH} + \mathsf{H_2O} \ \Longleftrightarrow \ \mathsf{RCOO^-} + \mathsf{H_3O^+} + \mathsf{RCOOH}$$

3. Carboxylate ion is best stabilised by resonance hence these are acidic in nature.

$$R - C \longrightarrow R - C \bigcirc = R - C \bigcirc \frac{0^{-\frac{1}{2}}}{0^{-\frac{1}{2}}}$$

$$I \qquad II$$

4. Acidic strength can also be explain on the basis of Ka and pKa values. More the Ka value or less the pKa value more the acidic strength and vise versa.

#### Acidic nature

Elelctron donating groups (+I) effect i.e. NH<sub>2</sub> > OH > OCH<sub>3</sub> > CH<sub>3</sub> destabilize the anion, by intensify negative charge and thus decrease the acidity of acid.

$$EDG \longrightarrow -C \stackrel{O^{\delta-}}{\stackrel{\circ}{\circ}}^{\delta}$$

2. Elelctron withdrawing groups (–I) effect. i.e.  $CF_3 > NO_2 > CN > F > Cl > Br > I > C_6H_5$ stabilize the anion, by dispersing negative charge and thus increase the acidity of acid.

EWG 
$$\leftarrow$$
 - C  $\stackrel{\circ}{\sim}_{O^{\delta-}}$ 

#### Which is depends up on following factors:

- The number of electron attracting groups or atoms (-I effect) increases, the strength of acid increases. The order of acidity
- e.g.  $CCl_3COOH > CHCl_2COOH > CH_2CICOOH$ >  $CH_3COOH$
- b) The power electron attracting groups or atoms increases, the acidity increases.
- e.g. CH,FCOOH > CH,ClCOOH > CH,BrCOOH
- c) The distance of electron attracting groups or atoms from carboxylic group increases, acidity decreases. Therefore, decreasing acidity order in n-butyric acid is  $\alpha > \beta > \gamma > \delta$ . Inductive effect is stronger at  $\alpha$  -position than  $\beta$ -position, similarly at  $\beta$ -position it is more stronger than  $\gamma$ -position.

- e.g. CH<sub>3</sub>CH<sub>2</sub>CHClCOOH > CH<sub>3</sub>CHClCH<sub>2</sub>COOH > CH<sub>2</sub>ClCH<sub>2</sub>CH<sub>2</sub>COOH > CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>COOH
- d) More the power of electron withdrawing goup, strength of carboxylic acid increases.
- e.g.  $CH_3-CH_2-COOH < H-CH_2-COOH < C_6H_5CH_2-COOH < HO-CH_2-COOH < Cl-CH_2-COOH$
- e) Direct attachment of groups such as phenyl  $(C_6H_5-)$  or vinyl  $(CH_2 = CH-)$  or acetylenic  $(CH \equiv C-)$  to -COOH group, increases the

acidity of corresponding carboxylic acid. This is due to more electronegative Sp<sup>2</sup>- and sp-hybridised carbon atom to which –COOH group is attached.

$$H_2C \xrightarrow{C} C$$
 $OH$ 
 $OH$ 
 $OH$ 
 $OH$ 

e.g.

- i)  $H-COOH > C_6H_5-COOH > C_6H_5CH_2$  $COOH > CH_3COOH > C_2H_5COOH$
- ii)  $CH \equiv C-COOH > C_6H_5COOH > CH_2 = CH-COOH (C_6H_5 group is more electron withdrawing than <math>CH_2 = CH-$ ).

#### 12.6 ACIDIC NATURE IN AROMATIC ACID

In aromatic carboxylic acid electron donating group decrease the acidic strength while electron withdrawing group increase the acidic strength

i) Electron donating groups decrease the acidic strength of acids by destabilizing benzoate ion.
 It follows the order, Ph-COOH > o > m > p

ii) Electron withdrawing groups increase the acidic strength of acids by stabilizing benzoate ion.It follows the order, o > p > m > Ph—COOH

## 12.7 REACTION SHOWS THE ACIDIC NATURE OF CARBOXYLIC ACIDS

1. Reaction with metal: (Na, K, Ca, Zn)



$$2R$$
-COOH +  $2Na \longrightarrow 2R$ -COONa +  $H_2$ 

b) Reaction with potassium metal:

$$2R$$
-COOH +  $2K$   $\longrightarrow$   $2R$ -COOK +  $H$ ,

c) Reaction with calcium metal:

$$2R-COOH + Ca \longrightarrow (R-COO)_2Ca + H_2$$

d) Reaction with Zn metal:

$$2R-COOH + Zn \longrightarrow (R-COO), Zn + H,$$

- 2. Reaction with strong alkali (NaOH, KOH)
- a) Reaction with NaOH:

$$R-COOH + NaOH \longrightarrow R-COONa + H_2O$$

b) Reaction with KOH:

$$R$$
-COOH +KOH  $\longrightarrow$   $R$ -COOK +  $H_2O$  pot. formate

- 3. Reaction with weak alkali (NaHCO<sub>3</sub>) NaZCO<sub>3</sub>, NH<sub>4</sub>OH):
- a) Reaction with NaHCO<sub>3</sub> (sodium bicarbonate or baking soda):

$$R-COOH + NaHCO_3 \xrightarrow{Fast}$$

b) Reaction with Na<sub>2</sub>CO<sub>3</sub> (Sodium carbonate):

$$2R$$
-COOH + Na<sub>2</sub>CO<sub>3</sub>  $\xrightarrow{\text{Slow}}$ 

$$2R$$
– $COONa + CO2 + H2O$ 

c) Reaction with NH<sub>4</sub>OH (ammonium hydroxide):

$$R-COOH + NH_4OH \longrightarrow R-COONH_4 + H_2O$$

- 4. Reaction involving breaking of C-O bond (Nu-SR)
- a) Formation of anhydride (Reaction with cone. H,SO<sub>4</sub> or P,O<sub>5</sub>)
- 1) From  $P_2O_5$ :

$$\begin{array}{ccc} R-COOH & & R-CO \setminus \\ + & + & \xrightarrow{P_2O_5} & R-CO / \\ R-COOH & & & R-CO / \end{array}$$

acid

$$P_2O_5 + H_2O \longrightarrow 2HPO_3$$

meta phosphoric acid

**Note**: Formic acid does not form formic unhhydride (H–CO)<sub>2</sub>O. When it is heated with  $P_2O_5$ , undergoes dehydration forming CO.

$$H$$
-COOH  $\xrightarrow{P_2O_5}$  CO +  $H_2O$ 

2) From acyl chloride and sodium salt of carboxylic acids:

$$R-COCl + R-COONa \xrightarrow{Pyridine}$$

b) Esterification (Fischer esterification) OR Alcoholysis of acids OR Acylation of alcohols.

$$R$$
–COOH +  $R$ '–OH  $\leftarrow \Delta$ 

acid alcohol

Note:

- i) These reactions are reversible in nature.
- ii) The yield of ester may increased by using excess of alcohol or removal of water by distillation or excess of acid. Water may be removed by using concentrated H<sub>2</sub>SO<sub>4</sub> or dry HCl or anhydrous ZnCl<sub>2</sub>:
- iii) The concentrated H<sub>2</sub>SO<sub>4</sub> or dry HCl or anhydrous ZnCl<sub>2</sub> acts as mainly dehydrating agent and it also act as catalyst in the esterification reaction.
- iv) These are nucleophilic substitution reaction.
- v) The reaction is shifted to the right by using excess of alcohol or removal of water by distillation.
- vi) The greater the bulk of the substituents near the OH group of alcohol and -COOH group of acid, the slower is the rate of esterification due to stearic hindrance.

Thus, the reactivity of alcohol towards esterification is –

$$CH_3OH > C_2H_5OH > C_3H_7OH > C_4H_9OH$$
  
Similarly the reactivity of carboxylic–acid increase

$$HCOOH > CH_3COOH > C_2H_5COOH > (CH_3)_2CHCOOH$$

- vii) The reverse of esterification is known as hydrolysis which may effected by acids or alkalies.
- viii) The mechanism of esterification is detected by tracer technique by taking isotopic oxygen in alcohol. After esterification isotopic oxygen become in ester. This indicate that during esterification the cleavage of C-O bond of carboxylic acid and O-H bond of alcohol.

$$\begin{array}{c}
O \\
R - C - OH + R - O - H \xrightarrow{\text{conc.}} \\
\downarrow \\
18
\end{array}$$

$$\begin{array}{c} O \\ \parallel \\ R - C - O - R + H_2O \\ \downarrow \\ 18 \end{array}$$

c) Reaction with PCl<sub>3</sub>, PCl<sub>5</sub> and SOCl,

$$3RCOOH + PCl_3 \longrightarrow 3RCOCl + H_3PO_3$$

$$RCOOH + PCl_5 \longrightarrow RCOC1 + POCl_3 + HCl$$

$$\mathsf{RCOOH} + \mathsf{SOCl}_2 \xrightarrow{\quad \mathsf{pyridine} \quad} \mathsf{RCOCl} + \mathsf{SO}_2 + \mathsf{HCl}$$

When acid chloride is heated with ammonia gives acid amide.

$$RCOCl + NH_3 \longrightarrow RCONH_2 + HCl$$

d) Reaction with ammonia:

When carboxylic acid is reacted with ammonia, gives ammonium salt of acid. Which on heating loses water molecule and gives amide.

$$R-COOH + NH_3 \longrightarrow R-COONH_4$$
  
 $R-COONH_4 \xrightarrow{\Delta} R-CONH_2 + H_2O$ 

3) Reaction involving -COOH group

Reduction by using LiAIH<sub>4</sub> or diborane ( $B_2H_6$ ) as reducing agent or hydrogen in the presence of copper chromite (CuO, CuCr<sub>2</sub>O<sub>4</sub>) as catalyst: Carboxylic acid are reduced by using LiAIH<sub>4</sub> as catalyst gives primary alcohols.The >C=O group is reduced to >CH<sub>2</sub> group.

$$\begin{array}{c} \text{R-COOH} + 4[\text{H}] \xrightarrow{\quad \text{i) LiAlH}_4/\text{ether}} \\ & \text{R-CH}_2\text{OH} + \text{H}_2\text{O} \\ \\ \text{CH}_3\text{-COOH} + 4[\text{H}] \xrightarrow{\quad \text{i) LiAlH}_4/\text{ether}} \\ & \text{CH}_3\text{-CH}_2\text{OH} + \text{H}_2\text{O} \end{array}$$

Note:

Sodium boronhydride (NaBH<sub>4</sub>) does not reduces carboxyl group.

- ii) Decarboxylation:
- a) Decarboxylation by using soda lime: Sodium salt of carboxylic acid is heated with soda lime (NaOH + CaO 3 : 1 ratio) gives alkane

$$\text{R-COONa} + \text{NaOH} \xrightarrow{\quad \text{CaO} \quad \quad } \text{R-H} + \text{Na}_2\text{CO}_3$$

4) Halogenation (HeIl-Vohlard-Zelinsky reaction) or HVZ reaction:

Carboxylic acids having  $\alpha$  –H atom undergoes halogenation at only  $\alpha$  –position in the presence of red phosphorus to give  $\alpha$  –halo carboxylic acids.

$$R - \overset{\alpha}{C} H_2 - COOH + X_2 \xrightarrow{i) \text{ Red.P.}}$$

$$X = C1 \text{ Rr}$$

α -halo carboxylic acid

e.g. i) 
$$CH_3$$
-COOH +  $Cl_2 \xrightarrow{i) Red.P.}$   $CH_2Cl$ -COOH +  $HCl$  chloroacetic acid

ii) 
$$CH_2$$
-COOH +  $Br_2$   $ii)$   $Red.P$   $iii)$   $H_2O$ 

1-bromo 1-phenyl ethanoic acid

The importance of this reaction is that only  $\alpha$  –H atom is substituted.

e.g.

i) 
$$CH_3$$
-COOH  $\xrightarrow{Br_2/Red P}$ 

$$Br-CH_2-COOH \xrightarrow{Br_2/Red P}$$

$$Br_2CH-COOH \xrightarrow{Br_2/Red P} Br_3C-COOH$$

ii) 
$$CH_3$$
-CHBr-COOH  $\xrightarrow{Br_2/Red P}$ 

$$CH_3$$
-CHBr-COOH  $\xrightarrow{Br_2/Red P}$ 

$$CH_3$$
- $CBr_2$ - $COOH \xrightarrow{Br_2/Red P}$  No reaction

## 12.8 ELECTROPHILIC SUBSTITUTION REACTION OF CARBOXYLIC ACIDS

Aromatic carboxylic acid undergoes electrophilic substitution reaction. Carboxyl group is meta directing.

**Note:** Aromatic carboxylic acid never undergoes Friedel –Craft reaction because only, AlCl,,

(Lewis acid) get bonded to carboxyl group. Other Lewis acid does not bonded to carboxyl group.

#### 1. Halogenation

e.g.

i) 
$$X = CI$$
, Br,  $X = COOH$ 

COOH

TOOH

2. Nitration

e.g.

COOH
$$+ \text{HNO}_3 \xrightarrow{\text{conc.H}_2\text{SO}_4} \longrightarrow \bigcirc \text{NO}_2 + \text{H}_2\text{O}$$

$$\text{m-nitro benzoic acid}$$

OOO





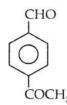
#### **MULTIPLE CHOICE QUESTIONS**

#### SECTION - I: ALDEHYDES AND KETONES

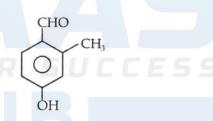
#### INTRODUCTION, CLASSIFICATION, NOMENCLATURE, ISOM'ERISM, STRUCTURE OF > C = O GROUP

- Aldehydes are characterised by the general formula
  - a)  $C_n H_{2n} O$
- b) C<sub>n</sub>H<sub>2n</sub>
- c)  $C_n H_{2n} + OH$
- d)  $C_n H_{2n+2}$ , O
- 2. Hybridisation of carbon in -CHO group is
  - a) sp
- b)  $sp^2$
- c)  $sp^3$
- d) none of these
- 3. Formalin is a
  - a) 100% solution of HCHO
  - b) 40% solution of HCHO
  - c) 60% solution of HCHO
  - d) 40% solution of CH3COOH
- An aldehyde group can be present
  - a) in between carbon chain
  - b) at any position in carbon atom
  - c) only at the end of carbon chain
  - d) at the second carbon atom of the carbon chain
- Butanal is an example of
  - a) primary alcohol
- b) secondary alcohol
- c) aliphatic aldehyde d) aliphatic ketone
- 6. In IUPAC system aldehydes are called
  - a) alkanes
- b) alkenes
- c) alkanals
- d) alkanols
- IUPAC name of mesityl oxide is
  - a) 4-methyl penta-3-none
  - b) 4-methyl pente-3-none
  - c) 4-methyl pent-3-en 2-one
  - d) 4-methyl pent-3-en 'l-one
- 8. IUPAC name of pinacol is
  - a) 2,3–dimethyl butan 2,3–diol
  - b) 2,3-dimethyl butane 2,3-diol
  - c) 2,3–dimethyl 2–butanol
  - d) 2,3-dimethyl 3-butanol
- IUPAC name of diacetone amine is
  - a) 2-methyl 4-keto 2-pentanamine
  - b) 4-amino 4-methyl 2-pentanone
  - c) 4-amino 2-pentanone
  - d) 4-amino 3-pentanone
- 10. The IUPAC name of crotonaldehyde is

- a) pentenal
- b) bu t-2-en-1-al
- c) bu t-2-an-1-al
- d) bu t-2-en-1-ol
- 11. IUPAC name of the following compound is



- a) 4-formyl acetophenone
- b) 4-ketomethyl benzaldehyde
- c) 1-formyl 4-ketomethyl benzene
- d) 4-formyl I-ketomethyl benzene
- 12. IUPAC name of ketone is
  - a) alkanol
- b) alkanal
- c) alkanone
- d) alkyl alkanoate
- 13. Tautomerism is possible in
  - 1) CH,-CHO
- 3) H-CHO
- 2) CH<sub>2</sub>-CO-CH,
- 4) (CH<sub>3</sub>)<sub>3</sub>C-CHO
- a) 1, 2
- b) 2, 3
- c) 3, 4
- d) 1, 2, 4
- 14. IUPAC name of following compound will be



- a) 4-formyl 3-methyl 1-hydroxy benzene
- b) 4–formyl 3–methyl phenol
- c) 4-hydroxy 2-methyl benzaldehyde
- d) 4-hydroxy 2-methyl carbaldehyde
- 15. IUPAC name following compound is



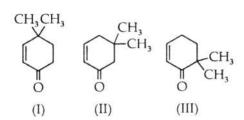
- a) 4-oxo pent-2-ene b) pent-2-en-4-one
- c) 2-oxo-pent-3-ene d) pent-3-en-2-one
- 16. Which of the following is
  - 3-phenylprop-2-en-1-al?
  - a)  $C_6H_5$ -CH = CH-CH<sub>2</sub>-CHO
  - b)  $C_{\varepsilon}H_{\varepsilon}$ –CH = CH–CHO
  - c) C<sub>6</sub>H<sub>5</sub>-CH<sub>2</sub>-CHO
  - d)  $C_6H_5$ -CH = CH-C $H_2$ -CHO

- 17. IUPAC name of benzophenone is
  - a) 2-phenyl propenal
  - b) 3-phenyl butenal
  - c) diphenyl methanone
  - d) diphenyl ethanone
- 18. IUPAC name of the following compound is



- a) 2-(2-propenyl) butanal
- b) 2–(1–propeny l)bu tanal
- c) 4-formyl 4-ethyl but-2-ene
- d) 2-ethyl pent-3-en-I-al
- 19. Aldehyde group can occur,
  - a) any where in carbon chain
  - b) in the middle of carbon chain
  - c) at only second carbon atom of carbon chain
  - d) only at end of the carbon chain
- 20. The carbon atom of carbonyl group is
  - a) sp<sup>3</sup>–hybridised
- b) sp<sup>2</sup>-hybridised
- c) sp-hybridised
- d) sp<sup>3</sup>-d-hybridised
- 21. C<sub>n</sub>H<sub>2</sub>O is general formula for
  - a) aldehydes
  - b) ketones
  - c) all carbonyl compounds
  - d) aldehyde and ketones
- 22. IUPAC name of β-methyl butyraldehyde

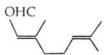
  - a) 2-methyl butanal b) 3-methyl butanal
  - c) 2-methyl propanal d) 3-methyl pentanal
- 23. Given:



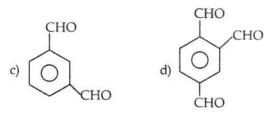
Which of the given compunds can exhibit tautomerism?

- a) I and II
- b) I and III
- c) II and III
- d) I, II and III
- 24. IUPAC name of isobutyraldehyde is
  - a) 2-methyl propanone
  - b) 2-methyl propanal
  - c) 2-methyl propanone

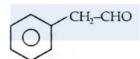
- d) methyl ethanoate
- 25. Which of the following is trimethyl acetaldehyde?
  - a) (CH<sub>2</sub>),C-CH<sub>2</sub>-CHO
  - b) (CH<sub>3</sub>)<sub>3</sub>C-CH<sub>2</sub>-CH<sub>2</sub>-CHO
  - c) (CH<sub>2</sub>),C-CHO
  - d) (CH<sub>2</sub>)<sub>2</sub>CH–CH(CH<sub>2</sub>)CHO
- 26. IUPAC name of the following compound is



- a) 2-methyl nona 2,6-dien-1-al
- b) 3-methyl nona 2,6-dien-1-al
- c) 4,6-dimethyl hepta 3,5-dien-1-al
- d) 3,7-dimethyl octa 2,6-dien-1-al
- 27. Aldehyde shows functional isomerism with
  - a) aliphatic ether
  - b) ketones
  - c) dimer of ethanoic acid
  - d) thio cyclic ether
- 28. Pentan-2-one and 3-methyl butan-2-one are
  - a) chain isomers
- b) position isomers
- c) metamers
- d) tautomers
- 29. Tautomer of ethyl methyl ketone is
  - a) bu t-2-en-2-ol
  - b) bu t-2-en-l-ol
  - c) but-2-en-1-oic acid
  - d) prop-I-en-2-ol
- 30. Which of the following is acrolein?
  - a)  $CH_{3}$ –CH = CH–CHO
  - b) CH<sub>2</sub> = CH-CHO
  - c)  $CH_2 = CH_2 CH_2$
  - d)  $CH_{2} = CH_{2} CH_{2} CH_{3}$
- 31. What is the IUPAC name of compound when -CHO group is attached to neoamyl group?
  - a) 2-methyl propanal
  - b) 3-methyl propanal
  - c) 2, 2–dimethyl butanal
  - d) 3, 3-dimethyl butanal
- 32. Which of the following is isophthaldehyde?



- 33. What is the IUPAC name of compound when carbonyl carbon atom is attached to phenyl group and ethyl group
  - a) propanone benzene
  - b) 1-phenyl propan-1-one
  - c) 2-phenyl propan-1-one
  - d) propiophenone
- 34. What is IUPAC name of compound when ketone group is attached to one isopropyl group and one t-butyl group?
  - a) 2, 4, 4-trimethyl 3-pentanone
  - b) 2, 2, 4–trimethyl 3–pentanone 45.
  - c) 2, 4–dimethyl 3–pentanone
  - d) 2, 2-dimethyl 3-pentanone
- 35. IUPAC name of ethyl isopropyl ketone is
  - a) 4-methyl pent-3-one 46.
  - b) 2-methyl pent-3-one
  - c) 4-methyl pent-2-one
  - d) 2-methyl pent-2-one
- 36. IUPAC name of acrolein is
  - a) but-2-enal
    - b) prop-2-enal
  - c) pentanal
- d) 2-methyl propanal
- 37. IUPAC name of the following compound is



- a) Phenyl acetaldehyde
- b) 1– Phenyl formyl methane
- c) 2- Phenyl ethanal
- d) 1- Phenyl methanal
- 38. Following compound is named as

- a) 4- methyl cyclohexane carbaldehyde
- b) δ-methyl cyclohexane carbaldehyde
- c) γ –methyl cyclohexane carbaldehyde
- d) both a and b

- 39. Keto-enol tautomerism is observed in
  - a) CH,OH
- b) CH, COCH,
- c) CH,COOH
- d) CH, COOCH,
- 40. Ketones are isomeric with
  - a) alcohols
- b) aldehydes
- c) ester
- d) acetic acid
- 41. The structural formula of the compound isomeric with acetone is
  - a) CH, CH, CHO
- b) CH<sub>2</sub>CHO
- c) CH<sub>3</sub>CH<sub>2</sub>OH
- d) none of these
- 42.  $CH_3 CO CH_3$  and  $CH_2 = COH CH_3$  are
  - a) metamers
  - b) tautomers
  - c) geometrical isomers
  - d) optical isomers
- 43. Aldehydes and ketone are
  - a) tautomers
- b) chian isomers
- c) functional isomers d) position isomers
- 44. Total number of isomeric aldehydes can be calculated by formula
  - a)  $I = 2^{n-2}$
- b)  $I = 2^{n-3}$
- c)  $I = 2^{n-2} 1$
- d)  $I = 2^{n}$
- 45. Total number of isomeric ketones can be calculated by formula
  - a)  $I = 2^{n-3} 1$
- b)  $I = 2^{n-2}$
- c)  $I = 2^{n-2} 1$
- d)  $I = 2^{n}$
- 46. Molecular formula C<sub>3</sub>H<sub>6</sub>O represents
  - a) aldehydes
  - b) ketones
  - c) both 'a' and 'b'
  - d) aldehydes and alcohols
- 47. C<sub>2</sub>H<sub>4</sub>O shows
  - a) chain isomerism
- b) position isomerism
- c) metamerism
- d) functional isomerism
- 48. How many aldehydes are possible for molecular formula C<sub>4</sub>H<sub>6</sub>O?
  - a) 2
- b) 3
- d) 5
- 49. How many ketones are possible for molecular formula C<sub>4</sub>H<sub>o</sub>O?
  - a) 1
- b) 2

c) 3

- d) 4
- 50. 2-pentanone and 3-pentanone are
  - a) chain isomers
- b) position isomers
- c) metamers
- d) functional isomers
- 51. Ketone never show

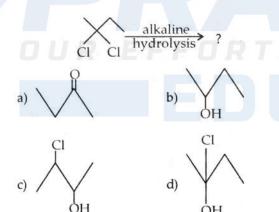
- a) chain isomerism
- b) position isomerism
- c) geometrical isomerism
- d) optical isomerism
- 52. In > C = O group sigma bond is formed by
  - a) sp<sup>2</sup>-p-overlapping b) sp<sup>3</sup>-p-overlapping
  - c) sp-p-overlapping d) s-p-overlapping
- 53. The  $\pi$  -bond in carbonyl group is formed by
  - a) s–s–overlapping b) p-p-overlapping
- c) s-p-overlapping d) p-d-overlapping
- 54. IUPAC name of following compound is



- a) heptane 2,6-dione b) octan-6-one
- c) octane 2,7–dione d) hexane-2,5-dione
- 55. Butanone and butanal are
  - a) chain isomers b) position isomers
  - c) functional isomers d) tautomers
- 56. Metamerism is present in
  - a) CH<sub>2</sub>-O-CH<sub>2</sub>
- b) CH<sub>2</sub>-CO-CH<sub>2</sub>
- c)  $CH_3-CO-C_2H_5$  d)  $CH_3-CO-C_2H_7$

#### PREPARATION METHODS

57. Product of following reaction is

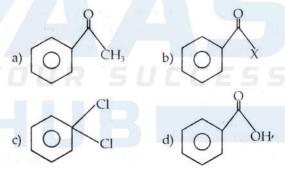


- 58. Calcium formate, on dry heating, produces
  - a) acetone
- b) acetic acid
- c) acetaldehyde
- d) formaldehyde
- 59. The reaction of RCN with RMgX followed by hydrolysis gives
  - a) an aldehydes
- b) an ketones
- c) 20 alcohols
- d) 30 alcohols
- 60. An isomer of C<sub>2</sub>H<sub>2</sub>Cl<sub>2</sub> on boiling with aq. KOH give acetone. Hence, the isomer is,
  - a) 2, 2-dichloro propane

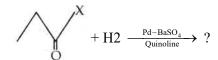
- b) 1, 2-dichloro propane
- c) 1, 1-dichloro propane
- d) 1, 3-dichloro propane
- 61. 2, 2-dichloro butane on boiling with aq. potash gives,
  - a) butanal
- b) 2-butanone
- c) 2-butanol
- d) butanoic acid
- 62. Propanone can be obtained from oxidation of what?
  - a) 2-propanol
- b) 1-propanol
- c) 2-butanol
- d) 2 methyl, 1-butanol
- 63. The reaction

$$CH_3COC1 + H_2 \xrightarrow{Pd-BaSO_4} CH3-C-H+HC1$$
 is called

- a) Cannizzaro's reaction
- b) Stephen's reduction
- c) Haloform reaction
- d) Rosenmund's reduction
- 64. Rosenmund's reduction is used to prepare
  - a) alcohol
- b) carboxylic acid
- c) aldehyde
- d) ketones
- 65. Benzaldehyde is obtained from Rosenmund's reduction of



- 66. 1,1–dichlorocyclohexane on alkaline hydrolysis
  - a) cyclohexane carbaldehyde
  - b) cyclohexanone
  - c) benzophenone
  - d) cyclohexane carboxylic acid
- 67. Product of the following reaction is IYx



- a) propanone
- b) propanal
- c) propionic acid
- d) propane
- 68. Rosenmund's reduction convert

- a) carboxylic acid to aldehyde
- b) ketone to 2°-alcohol
- c) acyl halide to ketone
- d) acyl halide to aldehyde
- 69. An optical active alcohol of formula C<sub>4</sub>H<sub>10</sub>O on oxidation gives which of the following compound?
  - a) (CH<sub>2</sub>)<sub>2</sub>CHCHO
- b)  $(CH_{3})_{2}C = CH_{3}$
- c) CH<sub>3</sub>COC<sub>2</sub>H<sub>5</sub>
- d) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CHO
- 70. A single compound of the structure

$$\begin{array}{ccc} CH_3 & CH_3 \\ & | & | \\ OHC - CH_2 - CH & - CH_2 - CH & = O \end{array}$$

is obtainable from ozonolysis of which of the following cyclic comounds?

a) 
$$H_3C$$
  $CH_3$ 





- 71. 2-methyl propanal is formed from isopropyl magnesium halide and what?
  - a) CH<sub>3</sub>CN
- b) HCHO
- c) HCN
- d) CH,COC
- 72. Calcium acetate on heating gives
  - a) acetone
- b) formic acid
- c) acetal dehyde
- d) methane
- 73. Which of the following is Collin's reagent?
  - a) pyridine
  - b) chromium oxide
  - c) pyridinium chlorochromate
  - d) KOH + KMnO<sub>4</sub>
- 74. Dry distillation of a mixture of calcium acetate and calcium formate can form
  - a) formaldehyde
- b) acetaldehyde
- c) acetone
- d) all of these
- 75. Which of the following is not used in the preparation of ketone?
  - a) Oxidation of secondary alcohols
  - b) Dehydrogenation of 2° alcohol
  - c) Pyrolysis of calcium acetate
  - d) Acid hydrolysis of alkyl cyanide
- 76. Which of the following compounds is oxidised to prepare ethyl methyl ketone?

- a) 1–butanol
- b) t-butyl alcohol
- c) 2-butanol
- d) 2-propanol
- 77. Product formed when but–1–yne is reacted with dil.H<sub>2</sub>SO<sub>4</sub> and HgSO<sub>4</sub>





- 78. Isopropyl alcohol on oxidation forms
  - a) acetone
- b) ether
- c) acetaldehyde
- d) ester
- 79. Ethylidene chloride on hydrolysis with aq . NaOH gives
  - a) CH<sub>3</sub>CHO
- b) CH,COC,H,
- c) C<sub>2</sub>H<sub>5</sub>OH
- d) CH<sub>3</sub>CH(OH),
- 80. Acetone is prepared by
  - a) oxidation of acetic acid
  - b) pyrolysis of acetic acid
  - c) pyrolysis of calcium acetate
  - d) oxidation of n-propyl alcohol
- 81. Formation of acetaldehyde from ethanol is known as
  - a) oxidation
- b) substitution
- c) addition
- d) reduction
- 82. Ketones can be obtained in one step by,
  - a) hydrolysis of esters
  - b) oxidation of primary alcohols
  - c) oxidation of secondary alcohols
  - d) reaction of acid unhydride and alcohol.
- 83. 2—butanone can be obtained by heating a mixture of calcium salt of
  - a) formic acid and butyric acid
  - b) propionic acid and formic acid
  - c) propionic acid and acetic acid
  - d) acetic acid and formic acid
- 84. Ozonolysis following compound in the presence of Zn gives



- On heating calcium acetate and calcium formate, the product formed is
  - a) CH, COCH,
- b) CH,CHO
- c) HCHO + CaCO<sub>3</sub> d) CH<sub>3</sub>CHO + CaCO<sub>3</sub>
- 86. Which of the following compound gives a ketone with Grignard reagent?
  - a) Formaldehyde
- b) Ethyl alcohol
- c) Methyl cyanide
- d) Methyl iodide
- 87. Which one is not synthesized by Grignard reagent?
  - a) Primary alcohol
- b) Secondary alcohol
- c) A ketone
- d) An ester
- 88. Ethylidene chloride on treatment with aqueous KOH gives?
  - a) Acetaldehyde
- b) Ethylene glycol
- c) Formaldehyde
- d) None
- 89. Find out reducing agent in following conversion.



- a) Na.Hg + H,O
- b) H, + Ni
- c) Pd.BaSO<sub>4</sub> + quinoline
- d) SnCl<sub>2</sub> + HCl
- 90. Which poison the catalyst (Pd) at aldehyde stage in the reduction of acyl halide
  - a) PCC
- b) BaSO,
- c) SnCl,
- d) CuSO<sub>4</sub>
- 91. Cyclohexanoyl chloride on reduction by poisoned catalyst gives
  - a) benzaldehyde
- b) cyclohexanol
- c) cyclohexanone
- d) cyclohexyl methanal
- 92. Stephen's reduction convert
  - a) acyl chloride to aldehyde
  - b) cyanide to aldehyde
  - c) cyanide to carboxylic acid
  - d) cyanide to ketones
- 93. The reaction

$$R-C \equiv N \xrightarrow{SnCl_2+HCl \atop H_3O^+} R-CHO+NH_3$$

- a) Rosenmund's reduction
- b) Stephen's reduction

- c) Clemmenson's reduction
- d) Cannizzaro's reaction
- 94. Formonitrile is reduced by SnCi, + HCl and product on acid hydrolysis gives
  - a) formic acid
- b) methanal
- c) acetic acid
- d) acetone
- 95. Product of the following reaction is

- 96. Rosenmund's reduction carried out by using
  - a) H<sub>2</sub>/Ni
- b) Na.Hg + H<sub>2</sub>O
- c) LiAlH,
- d) Pd-BaSO<sub>4</sub> + quinoline
- Product of the following reaction is

c) 
$$CN$$
 $+2(H)$ 
 $SnCl_2+HCl$ 
 $H_3O$ 
 $CHO$ 
 $CHO$ 

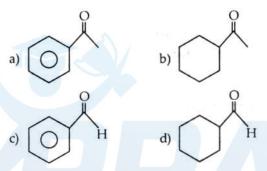
- 98. Conversion of ester to aldehyde which of the following reagent is used?
  - a) SnCl<sub>2</sub> + HCl
- b) Pd.BaSO<sub>4</sub> + quinoline
- c) Cr<sub>2</sub>O<sub>2</sub>
- d)  $(isobut)_2Al H$
- 99. The product of following reaction is

 $C_2H_5COOCH_3 \xrightarrow{(iso-but)_2 Al-H}$ 

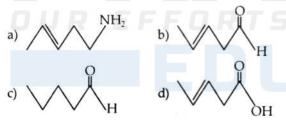
- a) CH,-CHO
- b) C<sub>2</sub>H<sub>2</sub>CHO

c)  $CH_3 - \overset{\parallel}{C} - C_2H_5$  d)  $C_2H_5 - COOH$ 

- 100. Compound A is reacted with KCN and followed by reduction using SnCl<sub>2</sub> + HCl and successive hydrolysis gives propanal. The compound A is
  - a)  $C_2H_5-X$
- b) C<sub>2</sub>H<sub>5</sub>-OH
- c) CH<sub>3</sub>-CH<sub>2</sub>-CH<sub>2</sub>-X d) C<sub>2</sub>H<sub>5</sub>-CN
- 101. Acetaldehyde is
  - a) oxidising agent
  - b) reducing agent
  - c) both oxidising and reducing agent
  - d) none of these
- 102. Ethyl cyanide is reduced by di–isobutyl aluminium hydride gives
  - a) propanal
- b) propanoic acid
- c) propan-1-amine
- d) propane
- 103. Methyl benzoate is reduced by di-isobut aluminium hydride gives



104. Pent–3–ene–1–nitrile on reduction by di–isobutyl aluminium hydride gives



- 105. Etard oxidation convert toluene to
  - a) benzoic acid
- b) acetophenone
- c) benzaldehyde
- d) benzoyl chloride
- 106. Oxidising agent in Etard oxidation is
  - a) K<sub>2</sub>Cr<sub>2</sub>O<sub>2</sub> + dil.H<sub>2</sub>SO<sub>4</sub>
  - b) KOH + KMNO,
  - c) dil. HNO<sub>3</sub>
  - d) CrO<sub>2</sub>Cl, in CS<sub>2</sub>
- 107. Oxidation of toluene by chromium oxide in acetic anhydride gives
  - a) benzaldehyde
- b) benzoic acid
- c) benzophenone
- d) acetophenone
- 108. p–nitro toluene convert p–nitro benzaldehyde by using

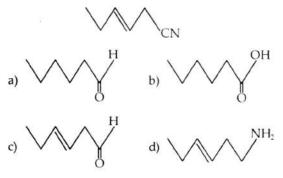
- a) KOH + KMnO<sub>4</sub>
- b) CrO<sub>3</sub> in (CH<sub>3</sub>CO)<sub>2</sub>O
- c) SnCl<sub>2</sub> + HCl
- d) Pd BaSO<sub>4</sub> + guinoline
- 109. Find out B in the reaction

- a) Acetophenone
- b) benzaldehyde
- c) cyclohexyl carbaldehyde
- d) benzoic acid
- 110. Find out C in the following reaction

- a) benzaldehyde
- b) 1-phenyl ethanal
- c) Acetophenone
- d) benzophenone
- 111. The reaction

$$\bigcirc + CO + HCI \xrightarrow{CuCl} \bigcirc CHO$$

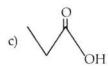
- a) Rosenmund's reaction
- b) Stephen's reaction
- c) Cannizzaro's reaction
- d) Gatterman-Koch reaction
- 112. Toluene is subjected to Gatterman–Koch reaction produces
  - a) benzaldehyde
  - b) benzoic acid
  - c) 2-methyl benzaldehyde
  - d) acetophenone
- 113. Following compound on reduction by using (iso-but), Al-H gives



114. Ethyl acetate is reduced by di–isobutyl aluminium hydride gives

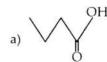




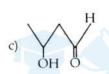




115. Compound A is reacted with C<sub>2</sub>H<sub>5</sub>-X gives B. The compound B is reduced by di-isobutyl aluminium hydride gives butanal. The compound A is









- 116. Benzal chloride on hydrolysis gives
  - a) benzoic acid
- b) benzaldehyde
- c) benzo chloride
- d) benzyl chloride
- 117. Toluene on side chain chlorination and followed by hydrolysis gives
  - a) Tolyl chloride
- b) benzaldehyde
- c) p-chloro toluene
- d) acetophenone
- 118. Side chain chlorination of alkyl benzene followed by hydrolysis gives
  - a) aromatic aldehyde
  - b) aromatic ketones
  - c) aromatic carboxylic acid
  - d) both a and b

#### PROPERTIES AND USES

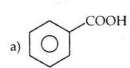
- 119. A reagent which reacts differently with CH<sub>2</sub>O, C<sub>2</sub>H<sub>4</sub>O, C<sub>2</sub>H<sub>5</sub>O is
  - a) NH<sub>2</sub>OH
- b) C<sub>6</sub>H<sub>5</sub>NHNH<sub>2</sub>
- c) HCN
- d) NH,
- 120. Which of the following has maximum reactivity towards HCN?
  - a) HCHO
- b) CH<sub>3</sub>CHO
- c) CH<sub>3</sub>COCH<sub>3</sub>
- d) CH<sub>3</sub>COC<sub>2</sub>H<sub>5</sub>
- 121. Consider the following substances
  - 1) HCHO
- 2) CH<sub>3</sub>CHO
- 3) CH<sub>3</sub>COCH<sub>3</sub>
- 4) CH<sub>3</sub>COC<sub>2</sub>H<sub>5</sub>

The correct order of reactivity towards

- nucleophilic addition reaction is
- a) 1 > 2 > 3 > 4
- b) 1 > 3 > 2 > 4
- c) 1 > 2 > 4 > 3
- d) 1 > 4 > 2 > 3
- 122. The reagent with which both acetaldehyde and ketone react easily is
  - a) Fehling solution
  - b) Grignard's reagent
  - c) Schiff's reagent
  - d) Tollen's reagent
- 123. Inpresence of cone. alkali formaldehyde undergoes
  - a) Aldol condensation
  - b) Cannizzaro's reaction
  - c) Esterification
  - d) Wurtz reaction
- 124. The Cannizzaro's reaction is not given by
  - a) benzaldehyde
- b) chloral
- c) methanal
- d) ethanal
- 125. Treatment of butanal with dilute NaOH solution gives
  - a) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>COOCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>
  - b) CH,CH,CH,CHOHCH,CH,CH,CHO
  - c) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CHOHCH(C<sub>3</sub>H<sub>5</sub>)CHO
  - b) CH,CH,CH,COCH,CH,CH,CHO
- 126. The products of the reaction of two molecules of HCHO with strong potash is
  - a) CH,OH and HCOOH
  - b) CH<sub>3</sub>OH and HCOONa
  - c) CH<sub>3</sub>OH and HCOOK
  - d) C<sub>2</sub>H<sub>2</sub>OH
- 127. In the reaction of NaHSO<sub>3</sub> with carbonyl compounds to form bisulphite product, the nucleophile is
  - a) HSO<sub>3</sub>
- b) SO<sub>3</sub>Na
- c) SO<sub>2</sub>--
- d) none of the above
- 128. The formation of cyanohydrin from a ketone is an example of
  - a) electrophilic addition
  - b) nucleophilic addition
  - c) electrophilic substitution
  - d) nucleophilic substitution
- 129. Acetone on treatment with CH<sub>3</sub>MgI on further hydrolysis gives,
  - a) (CH<sub>3</sub>),CHCH<sub>2</sub>OH
  - b) (CH<sub>3</sub>)<sub>3</sub>COH
  - c) CH<sub>3</sub>CH(OH)CH<sub>2</sub>CH<sub>3</sub>
  - d) CH<sub>3</sub>CH<sub>2</sub>CH(OH)CH<sub>2</sub>CH<sub>3</sub>

- 130. Base catalysed aldol condensation occurs with
  - a) propanal
- b) butanal
- c) ethanal
- d) all of these
- 131. Treatment of propanal with dilute NaOH solution gives
  - a) CH<sub>3</sub>CH<sub>2</sub>COOCH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>
  - b) CH<sub>2</sub>CH<sub>2</sub>CHOHCH(CH<sub>2</sub>)CHO
  - c) CH,CH,CHOHCH,CH,CHO
  - b) CH,CH,COCH,CH,CHO
- 132. Ethanal and propanone undergoes aldol condensation reaction inpresence of dilute alkali to form
  - a) CH<sub>2</sub>C(OH)(CH<sub>2</sub>)CH<sub>2</sub>CHO
  - b) CH,CH(OH)CH,COCH,
  - c) CH<sub>3</sub>COC(OH)(CH<sub>3</sub>),
  - d) CH<sub>3</sub>COCH(CH<sub>3</sub>)CH<sub>2</sub>OH
- 133. Iodoform test is not given by
  - a) ethanol
- b) ethanal
- c) acetone
- d) 3-pentanone
- 134. Methyl ketones are usually characterised through
  - a) the Tollen's reagent
  - b) the Schiffs reagent
  - c) the iodoform test
  - d) Cannizzaro's reaction
- 135. Grignards reagent adds to
  - a) > C = O
- b) > C = S
- c)  $-C \equiv N$
- d) all of the above
- 136. An organic compound of formula, C<sub>3</sub>H<sub>6</sub>O forms phenyl hydrazone, but gives negative Tollen's test. The compound is
  - a)  $CH_2 = CHOCH_3$
- b) CH<sub>3</sub>CH<sub>2</sub>CHO
- c) CH<sub>3</sub>COCH<sub>3</sub>
- d)  $CH_2 = CHCH_2OH$
- 137. Which does not react with Fehling solution?
  - a) Ethanal
- b) Glucose
- c) Formic acid
- d) Benzaldehyde
- 138. The reaction of RMgX with a ketone, followed by treatment with H<sub>3</sub>O<sup>+</sup> forms
  - a) 1° alcohol
- b) 2° alcohol
- c) 3° alcohol
- d) alkane.
- 139. When acetone is heated with ammonia gives diacetone amine, the intermediate compound formed in this reaction is,
  - a)  $(CH_3)_2C = CHCOCH_3$
  - b)  $CH_3CH_2CH = CH_3COCH_3$
  - c)  $(CH_3)_2C = CHCH_2CHO$
  - d)  $(CH_2)_2C = CH_2CH_2COCH_2$

- 140. >C = O is converted into -CH<sub>2</sub>- by
  - a) Clemmenson reduction
  - b) H<sub>2</sub>/Ni
  - c) Mendius reduction
  - d) NaHg + H<sub>2</sub>O reaction.
- 141. Which of the following give self redox reaction?
  - a) Methanal
- b) Ethanal
- c) Butanal
- d) Methanol
- 142. Which of the following reagent react with CH<sub>2</sub>O forming urotropine?
  - a) NH<sub>2</sub>OH
- b) NH<sub>3</sub>
- c) HCN
- d) RMgX
- 143. Propanal and prop an one undergoes condensation reaction in presence of dil.KOH to form,
  - a) CH<sub>3</sub>CH<sub>2</sub>CH(OH)CH<sub>2</sub>COCH<sub>3</sub>
  - b) CH<sub>3</sub>C(OH)(CH<sub>3</sub>)CH(CH<sub>3</sub>)CHO
  - c) CH<sub>3</sub>CH(OH)(CH<sub>3</sub>)CH<sub>2</sub>CH<sub>2</sub>CHO
  - d) none of these
- 144. Which one of the following reaction yield diacetone alcohol from carbonyl compound?
  - a) Cannizzaro's reaction
  - b) Catalytic hydrogenation
  - c) Aldol condensation
  - d) Oxidation
- 145. When H–CHO reacts with NH<sub>3</sub>, urotropin is formed. In this molecule how many C–C bonds are present?
  - a) 4
- b) 2
- c) 0
- d) 6
- 146. Acetone shows similarity with acetaldehyde in reacting to
  - a) Tollen's reagent
  - b) Schiff's reagent
  - c) Fehling solution
  - d) Grignard reagent
- 147. Cyanohydrine of the following compound on hydrolysis give optically active compound.
  - a) CH<sub>2</sub>-COCH<sub>2</sub>
- b) H–CHO
- c) CH<sub>2</sub>-CHO
- d) All of these
- 148. When ethanal is heated with Fehling's solution, it gives a precipitate of
  - a) Cu
- b) CuO
- c) Cu,O
- d)  $Cu_2O + CuO$
- 149. Benzene is reacted with carbon monoxide in HCl in the presence of catalyst, cupric chloride gives



- 150. Which of the following is used to prepare ketone from acyl chloride
  - a) R-MgX
- b) R,Cd
- c) CO + HC1
- d) CrO<sub>3</sub>
- 151. Dimethyl cadmium and acetyl chloride produces
  - a) acetone
- b) t-butyl alcohol
- c) 2°-propyl alcohol d) n-propyl alcohol
- 152. Product of the following reaction is

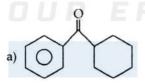
$$CH_{3}COC1 + CH_{3}MgX \xrightarrow{dry \text{ ehter} \atop H_{3}O^{+}}$$



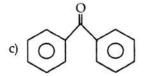


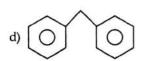


153. Benzonitrile and phenyl magnesium bromide produces

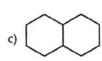


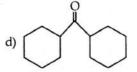






154. Product of the following reaction is





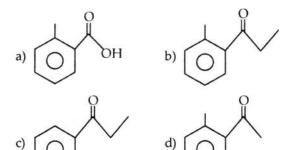
- 155. Friedel Craft acetylation of benzene gives
  - a) benzophenone
- b) acetophenone
- c) alkyl benzene
- d) diphenyl
- 156. Diphenyl methanone is obtained from

$$a) \bigcirc + \bigcirc \stackrel{O}{\longleftarrow} \stackrel{Unhy.AlCl_3}{\longrightarrow}$$

b) 
$$O \longrightarrow + Cl_2 \xrightarrow{U.V.L} A \xrightarrow{H_2O}$$

$$d) \overbrace{\hspace{1cm}}^{\text{CH}_3} \xrightarrow{\text{(isobutyl)}_2 \text{Al-H}}$$

157. Product of the following reaction is

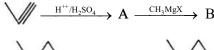


158. In the following sequence of reaction end product

$$CaC2 \xrightarrow{H_2O} A \xrightarrow{Hg^+/H_2SO_4} B \xrightarrow{[O]} C$$

$$\xrightarrow{Ca} D \xrightarrow{dry distillation} E$$

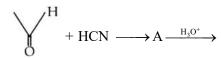
- a) acetaldehyde
- b) propanone
- c) methanal
- d) ethanoic acid
- 159. In the following sequence of reaction end product







- 160. The product 'B' in the following reaction...











- 161. The correct order of solubility of following compound is
  - 1) H-CHO
- 2) CH<sub>3</sub>-CHO
- 3) CH<sub>3</sub>-CH<sub>2</sub>-CHO 4) CH<sub>3</sub> COCH<sub>3</sub>
- a) 4 > 3 > 2 > 1
- b) 1 > 2 > 4 > 3
- c) 1 > 2 > 3 > 4
- d) 1 > 4 > 2 > 3
- 162. Acetylene on hydration gives compound A, which on Clemmenson's reduction gives?
  - a) Ethane
- b) Ethyl alcohol
- c) Acetaldehyde
- d) Ethene
- 163. Compound 'A' has molecular formula C<sub>5</sub>H<sub>10</sub>O reduce Tollen's reagent on oxidation gives monocarboxylic acid with same number of carbon atoms. The compound 'A' is
  - a) CH<sub>3</sub>COCH(CH<sub>3</sub>), b) CH<sub>3</sub>(CH<sub>3</sub>)<sub>3</sub>-CHO
  - c) C,H,COC,H,
- d) CH<sub>3</sub>COCH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>
- 164. Wolf-Kishner reduction is
  - a) reduction of carbonyl compound in to alcohol
  - b) reduction of carbonyl compound in to alkene
  - c) reduction of carboxyl compound in to alkane
  - d) reduction of nitro compound into aniline
- 165. Among the following compound the reactivity order is

2) CH<sub>2</sub>-CHO 1) H-CHO

O

- 3)  $CH_2 \overset{\parallel}{C} CH$  4)  $C_2H_5 COCH_2$
- a) 1 > 2 > 4 > 3
- b) 4 > 3 > 2 > 1
- c) 1 > 2 > 3 > 4
- d) 1 > 3 > 4 > 2
- 166. Less reactivity of ketone is due to
  - a) +I inductive effect decrease positive charge on carbonyl carbon atom
  - b) steric effect of two bulky alkyl groups
  - c) sp<sup>2</sup> hybridised carbon atom of carbonyl carbon atom
  - d) both a and b
- 167. To distinguish between acetone and iso-propyl alcohol, which of the following reagent can help
  - a) NaCl
- b) NaOH
- c) Na<sub>2</sub>CO<sub>3</sub>
- d) NaHSO,
- 168. Which of the following statement is wrong?
  - a) aldehyde and ketones are reducing agents
  - b) aldehyde and ketones are nonpolar compounds
  - c) aldehydes are more reactive than ketones
  - d) aldehydes and ketones are reduced to alcohol
- 169. Which of the following statement is correct regarding the aldol condensation?
  - a) All aldehydes give this reaction
  - b) Ketones do not give this reaction
  - c) This reaction proceeds in presence of strong alkali
  - d) Ketones, in which a-hydrogen atom is present, give this reaction
- 170. If formaldehyde and KOH are heated then we get
  - a) methane
- b) acetylene
- c) ethyl formate
- d) methyl alcohol
- 171. Benzaldehyde is converted to benzyl alcohol by
  - a) Wurtz reaction
  - b) Fitting reaction
  - c) Wurtz fittings reaction
  - d) Cannizzaro's reaction
- 172. Aldehydes are oxidised to acids by
  - a) potassium dicromate
  - b) Tollen's reagent
  - c) Fehling's solution
  - d) all of these
- 173. Formaldehyde + ammonia  $\rightarrow$  Y, the product Y is a) methanol

- b) formamide
- c) para-formaldehyde
- d) hexamethylene tetraamine
- 174. Acetaldehyde and acetone can be identified by
  - a) Schiff's test
- b) Lucas test
- c) Iodoform test
- d) Bromoform test
- 175. Ethanal reacts with alkali to give 3-hydroxy butanal. The reaction is
  - a) Aldol condensation
  - b) Claisen condensation
  - c) Cannizzaro's reaction
  - d) Clemmenson reduction
- 176. Which of the following undergoes haloform reaction?
  - a) HmO
- b) (CH<sub>2</sub>)<sub>2</sub>CO
- c) C<sub>2</sub>H<sub>5</sub>Cl
- d) CH<sub>3</sub>-O-CH<sub>3</sub>
- 177. Which of the following cation is involved In Tollens reagent?
  - a)  $[Ag (NH_3)_2]^+$
- b) 20H-
- c)  $[Ag (NH_3)_3]^+$
- d) AgNO<sub>2</sub>
- 178. Diethyl ketone and a dimethyl ketone can be distinguished with
  - a) Tollen's reagent
- b) Fehling's solution
- c) Schiffs reagent
- d) Haloform test
- 179. Tollen's reagent is
  - a) ammonical CuSO<sub>4</sub>
  - b) ammonical AgO,
  - c) alkaline solution containing complex of copper nitrate
  - d) none of these
- 180. Reduction of a keto group to a methylene group is converted by using
  - a) ZnHg + cone, HCl b) NaHg + water
  - c) Sn + cone, HCl
- d) Zn + CH, COOH
- 181. Which statement is false regarding acetaldehyde and acetone?
  - a) Both reduce ammonical silver nitrate to silver
  - b) Both react with hydroxylamine to form oximes
  - c) Both react with hydrazine to form hydrazone derivative
  - d) Both react with sodium bisulphite to form addition product
- 182. Which of following give haloform test?
  - a) CH<sub>3</sub>-CHO
- b) C<sub>2</sub>H<sub>2</sub>-CHO
- c) H-CHO
- d) C<sub>2</sub>H<sub>2</sub>-CO-C<sub>2</sub>H<sub>2</sub>
- 183. Which of the following compounds does not

- contain an -OH group?
- a) Alcohol
- b) Phenol
- c) Aldehyde
- d) Carboxylic acid
- 184. HCHO and CH<sub>3</sub>CHO differ from each other towards
  - a) Schiff's reagent
- b) Fehling solution
- c) ammonia
- d) ammonical AgNO,
- 185. The most active carbonyl compound is
  - a) HCHO
- b) CH,CHO
- c) CH, COCH,
- d) C,H,CHO
- 186. Out of
  - 1) butane
- 2) butan-1-o1
- 3) butanal
- 4) butanone.
- a) 1 > 2 > 3 > 4
- The decreasing order of their B.P is
- c) 2 > 3 > 4 > 1
- b) 2 > 4 > 3 > 1d) 4 > 2 > 3 > 1
- 187. Acetone is converted into propane by Wolff-Kishner reduction by using reagent
  - a) LiAIH,
- b) Zn.Hg + cone HCl
- c) H<sub>2</sub> + pd BaSO<sub>4</sub> d) NH<sub>2</sub>–NH<sub>2</sub> and KOH
- 188. A neutral compound C<sub>4</sub>H<sub>5</sub>O<sub>5</sub>, reduce Fehling's solution, and liberate H<sub>2</sub> gas when treated with sodium metal and give positive iodoform test. The com pound is
  - a) CH<sub>3</sub>-CHOHCH<sub>3</sub>-CHO
  - b) HO-CH<sub>2</sub>-CH<sub>2</sub>-CHO
  - c) CH<sub>3</sub>-CO-CH<sub>2</sub>-CHO
  - d) CH,COCH,-CH,-OH



189. Compound of general formula

are called

- a) diester
- b) acid anhydride
- c) hemiacetal
- d) acetal
- 190. In the reaction

$$CH_3CH = O + A \xrightarrow{dry HC1} CH_3CH(OC_2H_5)_2 + H_2O$$

The compound A is

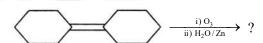
- a) propan-1-o1
- b) ethanol
- c) methanol
- d) C,H,-O-C,H,
- 191. During reduction of aldehyde with hydrazine and KOH, the first is the formation of
  - a) R CH = NH
- b) R CH = N NH
- c) RCONH,
- d) R C = N
- 192. Product of following reaction will be

$$\begin{array}{c|c} & & & & \\ & & & \\ & &$$

193. Compound of general formula are called

- a) acetal
- b) hemiketal
- c) ketal
- d) hemiacetal
- 194. Acetals are
  - a) aldehyde
- b) ketones
- c) ethers
- d) diether
- 195. Pentan–3–en–2–one is reduced by LiAlH<sub>4</sub> gives
  - a) pent-4-en-2-ol
- b) pent-3-en-l-ol
- c) pent-2-ol
- d) pent-3-en-2-ol
- 196. Which is best solvent
  - a) H-CHO
- b) CH, COCH,
- c) CH<sub>3</sub>-CHO
- d) CH, COOH
- 197. What is the structure of carbonyl compounds that will give following compound?

- 198. Prop 1,3-diphenyl 2-en-l-one is crossed condensation product. Which is obtained from
  - a) benzophenone
  - b) Acetophenone
  - c) benzaldehyde
  - d) Acetophenone and benzaldehyde
- 199. In mechanism of aldol condensation reaction the second step is
  - a) abstraction of  $\alpha$  –H atom carbon of aldehyde by base to form carbanion
  - b) The attack of carbanion on carbonyl carbon atom of another molecule to form alkoxide ion
  - c) The attack of carbanion on carbonyl carbon atom of another molecule to form oxocation
  - d) Alkoxide ion take proton from water to form
  - β-hydroxy aldehyde
- 200. Formaldehyde is used as
  - a) solvent
  - b) antiseptic
  - c) disinfectant
  - d) disinfectant and preservative
- 201. Carbonyl group in aromatic aldehyde and ketone
  - a) o-directing
- b) p-directing
- c) o-and p-directing d) m-directing
- 202. Benzaldehyde on nitration gives
- - a) o-nitrobenzaldehyde
  - b) p-nitrobenzaldehyde
  - c) mixture of a and b
  - d) m-nitrobenzaldehyde
- 203. Product of the following reaction will be



- a) cyclohexanal
- b) cyclohexanone
- c) cyclohexane carbaIdehyde
- d) cyclohexane carboxylic acid
- 204. But–2–en–l–ol on oxidation by using PCC gives
  - a) Butan-l-ol
- b) butanal
- c) but-2-en-al
- d) butan-2-one
- 205. Following compound on oxidation by K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>+ dil.H,SO, gives

$$O_2N$$
—OH

206. Product of the following reaction is

- a) acetophenone
- b) diphenyl methanone
- c) 2-diphenyl ethanone
- d) 1-phenyl ethan one
- 207. Methyl ketone group is identified by
  - a) Lucas test
- b) Haloform test
- c) Hinsberg test
- d) Millon's test
- 208. Match the List–I and II and select the correct answer

#### List-I

#### List-II

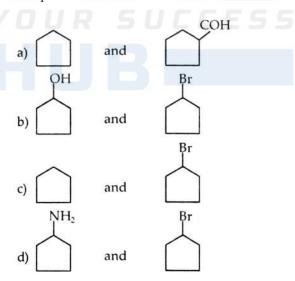
- 1. Cannizzaro's reaction
- A) SnCl<sub>2</sub> + HCl
- 2. Stephen's reaction B) 50% NaOH
- 3. Clemmenson
  - C) Pd-BaSO<sub>4</sub> + quinoline
- reduction
  4. Rosenmund's reduction
- D) Zn.Hg + Conc. HCl
- a) 1-B, 2-D, 3-A, 4-C
- b) 1-B, 2-C, 3-D, 4-A
- c) 1-B, 2-A, 3-D, 4-C
- d) 1-D, 2-A, 3-C, 4-B
- 209. Which of the following is not correct reaction?

a) 6H–CHO + 4 NH
$$_3$$
  $\longrightarrow$  (CH $_2$ ) $_6$ N $_4$  + 6 H $_2$ O

210. What is product formed when cyclohexane carbaldehyde undergoes aldol condensation followed by heating

211. 
$$\xrightarrow{NH_2-NH_2} A \xrightarrow{Br_2} B$$

The product A and B are



212. A carbonyl compound 'A' react with hydrogen cyanide to give cyanohydrin 'B' which on acid hydrolysis gives optically active α –hydroxy propanoic acid 'C'. Compounds A, B, C respectively are,

b) 
$$CH_3$$
 –  $CHO$ ,  $CH_3$  –  $C$  –  $OH$  ,  $CH_3$  –  $C$  –  $OH$  | COOH

c) 
$$C_2H_5$$
–CHO,  $C_2H_5$  –C –OH ,  $C_2H_5$  –C –OH ,  $C_2H_5$  –C –OH , COOH

d) 
$$CH_3 - C = O$$
,  $CH_3 - C - OH$ ,  $CH_3 - C - OH$ 

213. Acetone + A  $\longrightarrow$  Oxime of acetone

Acetone + B  $\longrightarrow$  Propane

Acetone + C  $\longrightarrow$  Pinacoi.

A, B and C are

- a) NH<sub>2</sub> OH, Mg, Zn.Hg + conc. HCl
- b) NH, OH, Zn.Hg + conc. HCl, Mg
- c) Mg, NH<sub>2</sub> OH, Zn.Hg + cone, HC
- d) NH, OH, Na.Hg + H,O, Mg

214. 
$$\mid$$
 + NaOH  $\xrightarrow{50\%}$  Product (s) . COOH

Identify the products

b) 
$$\begin{array}{c} \text{CH}_2-\text{OH} \\ \text{COONa} \end{array}$$

- c) both a and b
- **CHO**
- 215. Structure C<sub>5</sub>H<sub>5</sub>Cl<sub>7</sub> an alkaline hydrolysis gives a product which does not give iodoform test but give silver mirror test, is

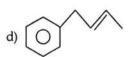
a) 
$$CCl_2-C_2H_5$$
 b)  $Cl_2-CH_3$ 

c) 
$$CH_2-CI$$
  $CH_2-CH CI_2$   $CH_3-CH CI_2$ 

- 216. Which of the following reagent to get sp<sup>3</sup>hybridised carbon atom from sp<sup>2</sup>-hybridised carbon atom?
  - a) NH<sub>2</sub>-NH<sub>2</sub>
- b) NH<sub>2</sub>-CO-NH-NH<sub>2</sub>
- c) NH<sub>2</sub>-NH<sub>2</sub>/KOH d) NH<sub>2</sub>OH
- 217. An alkene (A) on ozonolysis gives (B) as one of the product, (B) can gives Cannizzaro's reaction. The compound (A) is







218. Which of the following alkane cannot be prepared by Clernmenson's reduction of ketones?





- 219. Reaction of aldehyde with NH<sub>2</sub>-OH will be fastest at which of the following PH
  - a) 1
- b) 6
- c) 7
- d) 8
- 220. In the reaction.

$$C = O \xrightarrow{NH_2 - NH_2 + KOH} A$$

$$Na \cdot Hg + H_2O \qquad C$$

- a) A = aldehyde, B = ketone, C = alcohol
- b) A, B, C are alkene
- c) A, B are alkane and C = alcohol
- d) A, B are alkane, C = alkene
- 221. In Cannizzaro's reaction of formaldehydeinvolves
  - a) Intramolecular shift of proton

- b) Intramolecular shift of hydride
- c) Intermolecular shift of proton
- d) Intermolecular shift of hydride
- 222. Consider two aldehydes
  - 1) CH<sub>3</sub>-CHO
- 2) H-CHO

The correct statement is/are

- a) both undergoes aldol condensation
- b) both donot undergoes aldol condensation
- c) only (1) undergoes aldol condensation
- d) only (2) undergoes aldol condensation
- 223. Final product 'B in the following sequence of reaction

$$\begin{array}{c} \text{C}_{6}\text{H}_{5}\text{--CH}_{2}\text{OH} \xrightarrow{\quad \text{Cu} \quad } \text{A} \xrightarrow{\quad \text{NaOH} \quad } \text{B} + \\ \\ \text{C}_{6}\text{H}_{5}\text{CH}_{2}\text{--OH} \end{array}$$

- a) benzaldehyde
- b) acetophenone
- c) sodium phenoxide d) sodium benzoate
- 224. The final product 'C' in the following reaction.....

$$H-CHO + CH_3 - CH_2-CHO \xrightarrow{OH^-} A$$

$$\xrightarrow{\Delta/H^+} B \xrightarrow{NaBH_4} C$$

a) 
$$CH_2 = C$$
 -CHO
$$CH_3$$

b) 
$$CH_3$$
- $CH$  - $CH_2$ - $OH$   $CH_3$ 

c) 
$$CH_2 = C - CH_2 - OH$$
  
 $CH_3$ 

d) 
$$CH_2 = CH - CH_2 - CH_2 - OH$$

225. Which of the following four hydrogens which is most acidic?

- a) 1
- b) 2
- c) 3
- d) 4
- 226. Find out A, B, C, D in following reaction

1) 
$$CH_3$$
- $CH = CH$ - $CHO \xrightarrow{A}$ 

$$CH_3$$
- $CH = CH - COOH$ 

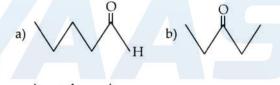
2) 
$$CH_3$$
– $CH = CH - CHO \xrightarrow{B}$ 

$$CH_3$$
- $CH = CH - CH_2 - OH$ 

3) R-COOH 
$$\stackrel{c}{\longrightarrow}$$
 R - CH<sub>2</sub> - OH

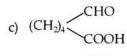
4) 
$$R - COC1 \xrightarrow{D} R - CHO$$

- a) 1 = H<sub>2</sub>.Pd.BaSO<sub>4</sub> 2 = Ammoniacal AgNO<sub>3</sub>
  - $3 = LiAIH_4$  4 =
    - $4 = LiAIH_4$
- b)  $1 = LiAIH_4$   $2 = LiAIH_4$ 
  - $3 = H_2.Pd.BaSO_4$   $4 = Ammoniacal AgNO_3$
- c)  $1 = \text{Ammoniacal AgN0}_3 2 = \text{LiAlH}_4$ 
  - $3 = LiAlH_{4}$
- $4 = H_2Pd + BaSO_4$
- d)  $1 = LiAlH_4$
- $2 = LiAlH_4$
- $3 = Ammoniacal AgNO_3$   $4 = H_2Pd + BaSO_4$
- 227. Which among the following is most reactive in nucleophilic addition?
  - a) FCH<sub>2</sub>-CHO
- b) Cl-CH,-CHO
- c) Br–CH<sub>2</sub>–CHO
- d) I–CH<sub>2</sub>–CHO
- 228. Which of the following carbonyl compound is most polar?
  - a) CH<sub>3</sub>COC<sub>2</sub>H<sub>5</sub>
- b) CH, COCH,
- c) CH<sub>2</sub>CHO
- d) H CHO
- 229. Which of the following isomeric compounds is most reactive?

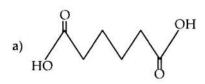


- c) \\
- d) all are equally reacive
- 230. The reagent which oxidise aldehyde and ketone is
  - a) Tollen's reagent
- b) Fehling's solution
- c) Schiff's reagent
- d)  $K_2Cr_2O_7 + H_2SO_4$
- 231. The product of given reaction is

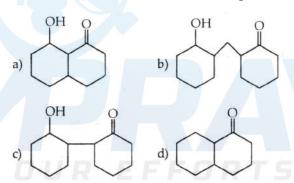
- a)  $CH_2 = CH(CH_2)_3 COOH$
- b) CH<sub>3</sub> –(CH<sub>2</sub>)<sub>4</sub> COOH



- d) (CH<sub>2</sub>)<sub>4</sub> CHO
- 232. 3-methyl cyclohexene on oxidation will give



233. Cyclohexanone is treated with Ba(OH), gives



- 234. Mixture of ethanal and propanal is subjected to aldol condensation, the product formed are
  - a) 1
- b) 2
- c) 3
- d) 4
- 235. Aldehyde which shows Cannizzaro's reaction is
  - a) H-CHO
- b) C<sub>6</sub>H<sub>5</sub>-CHO
- c) CCl,-CHO
- d) all of these
- 236. The aldehyde having  $\alpha$  –H atom gives Cannizzaro's reaction is

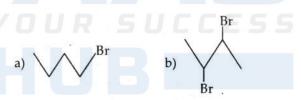
a) 
$$CH_3 - CH_2 - CHO$$

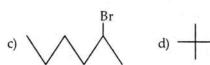
b) 
$$CH_3 - CH - CHO$$

c) 
$$CH_3 - CH_2 - CH_2 - CHO$$

237. Mixture of H-CHO and C<sub>6</sub>H<sub>5</sub>-CHO is treated

- with cone, NaOH then self redox reaction involves
- 1) oxidation of H-CHO
- 2) oxidation of C<sub>6</sub>H<sub>5</sub>-CHO
- 3) reduction of H-CHO
- 4) reduction of C<sub>6</sub>H<sub>5</sub>-CHO
- a) 1, 3
- b) 1, 2
- c) 1, 4
- d) 2, 3
- 238. When ethanoyl chloride is reduced with hydrogen using Pd deposited over BaSO<sub>4</sub> catalyst it forms
  - a) ethane
- b) chioro ethane
- c) ethanal
- d) ethanol
- 239. Crotonaldehyde is easily oxidised to crotonic acid using
  - a) alkaline KMnO<sub>4</sub>
- b) Tollen's reagent
- c) acidic KMnO<sub>4</sub>
- d) CrO<sub>3</sub>
- 240. Which of the following aldehyde forms stable hydrate?
  - a) Trichloroacetaldehyde
  - b) acetaldehyde
  - c) formaldehyde
  - d) prop anal
- 241. An unknown alkyl halide 'A' reacts with alcoholic KOH to produce C<sub>4</sub>Hg. Ozonolysis of the hydrocarbon gives one mole of prop anal and one mole of methanal the correct structure of 'A' is





- 242.  $(CH_3)_2 C = CHCOCH_3$  can be oxidised to  $(CH_3)_2 C = CH-COONa$  by
  - a) NaOH + KOH
- b) NaOH
- c) NaOI
- d) KMnO<sub>4</sub>/H<sup>+</sup>
- 243. For the following conversion which can be used

$$R_2C = O \longrightarrow R_2CH$$

- a) Clemmenson's reaction
- b) Wolff Kishner reaction
- c) Zn + cone. HCl
- d) both 'a' and 'b'

- 244. Benzaldehyde can be prepared by hydrolysis of
  - a) Benzyl chloride
- b) Benzal chloride
- c) Benzotrichloride
- d) benzonitrile
- 245. 1-phenyl ethan-1-ol is prepared by reaction of benzaldehyde with
  - a) methyl bromide
  - b) ethyl magnesium iodide
  - c) methyl magnesium bromide
  - d) methyl iodide and zinc metal
- 246. Oxidation of toluene to benzaldehyde by chromyl chloride is called
  - a) Rosenmund reaction
  - b) Wurtz reaction
  - c) Etard reaction
  - d) Fittig reaction
- 247. Which of the following reagent convert

$$C_6H_5CHO \longrightarrow C_6H_5COOH$$

- a) Fehling's solution
  - b) PCC
- c) Tollen's reagent
- d) H<sub>2</sub> + Ni
- 248. The product of following reaction will be



- a) m-hydroxy benzaldehyde
- b) potassium m-chlorobenzoate and benzyl alcohol
- c) potassium m-chlorobenzoate m-chlorobenzyl alcohol
- d) m-cresol
- 249. Hydrogenation of benzoyl chloride in the presence of Pd on BaSO<sub>4</sub> gives
  - a) benzyl alcohol
- b) benzoic acid
- c) benzaldehyde
- d) phenol
- 250. An organic compound does not give a precipitate with 2,4–DNP and does not react with sodium metal. It could be
  - a) CH<sub>3</sub>-CH<sub>2</sub>-CHO
- b) CH<sub>3</sub>COCH<sub>3</sub>
- c) CH<sub>2</sub>-CHO
- d) CH<sub>3</sub>OC<sub>2</sub>H<sub>5</sub>
- 251. What is 'B' in the following reaction?

$$CH_{3}CN \xrightarrow{\quad i) \, SnCl_{2} \, + \, HCl \quad} A \xrightarrow{\quad Na_{2}CO_{3} \quad} B$$

- b)  $CH_3$ -CH- $CH_2$ -CHOOH
- c) CH<sub>3</sub>-CH<sub>2</sub>-COONa
- d) CH<sub>2</sub>-CH<sub>2</sub>-ONa
- 252. Under Wolff–Kishner reduction the conversion may be brought about are
  - a) benzophenone to diphenyl methane
  - b) benzaldehyde to benzyl alcohol
  - c) cyclohexanone to cyclohexene
  - d) cyclohexanone to cyclohexanol
- 253. When formaldehyde react with ethyl amine, it gives
  - a) formaldehyde imine
  - b) formaldehyde ethyl amine
  - c) dimethyl amine
  - d) diethylamine
- 254. Imine derivatives of aldehyde and ketone is called as
  - a) Schiff's reagent
- b) Fehling's reagent
- c) Schiff's base
- d) Schiff's acid
- 255. Acetone on condensation gives
  - a) mesitylene
- b) mesityl oxide
- c) prop anal
- d) di-isopropyl ether
- 256. A carbonyl compound with molecular weight 86, does not reduce Fehling's solution, but form crystalline bisulphite derivatives and gives iodoform test. The possible compound can be
  - a) Pentan—2-one and pentan-3-one
  - b) Pentan-2-one and 3-methyl butan-Z-one
  - c) Pentanal
  - d) pent-3-one
- 257.  $CH_3 CH = CH CHO$  is oxidised to  $CH_3 CH = CH COOH$  using oxidising agent a) alkaline KMnO<sub>4</sub>
  - b)  $K_2Cr_2O_7 + conc.H_2SO_4$
  - c) dil.HNO<sub>3</sub>
  - d) ammoniacal AgNO,

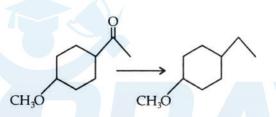
O ||

- 258. Ketones  $R \stackrel{||}{C} R'$  where R = R' alkyl group can be obtained in one step by
  - a) oxidation of 1° alcohol
  - b) hydrolysis of ester
  - c) hydrolysis of acid anhydride
  - d) oxidation of 3° alcohol

- 259. Among the given compound the most susceptible to nucleophilic attack at carbonyl group is
  - a) HCHO
- b) MeCOOMe
- c) MeCHO
- d) MeCOO COme
- 260. The reaction given below is

The slowest step is

- a) Wurtz's reaction
- b) Cannizzaro's reaction
- c) Fittig's reaction
- d) Etard oxidation
- 261. Which of the following does not used to convert ketone to secondary alcohol?
  - a) LiAlH<sub>4</sub>
- b) NaBH<sub>4</sub>
- c) H<sub>2</sub>/Pd-BaSO<sub>4</sub> d) Na + C<sub>2</sub>H<sub>5</sub>OH
- 262. The appropriate reagent for following transformation.



- a) Zn.Hg + HCl
- b) NH,NH,, KOH
- c)  $H_2 + Ni$
- d) both a and b
- 263. In the following sequence of reaction,

Toluene 
$$\xrightarrow{\text{KMnO}_4}$$
 A  $\xrightarrow{\text{SOCl}_2}$  B  $\xrightarrow{\text{H}_2 + \text{Pd}}$  C

The product C is

- a) C<sub>6</sub>H<sub>5</sub>-COOH
- b)  $C_6H_5-CH_3$
- c) C<sub>6</sub>H<sub>5</sub>-CHO
- d) C<sub>6</sub>H<sub>5</sub>-CH<sub>2</sub>-OH
- 264. Which of the following is correct?
  - a) reduction of any aldehyde gives 2° alcohol
  - b) reduction of any ketone gives tertiary alcohols
  - c) Wolff–Kishner reduction convert > C = O to -CH<sub>2</sub>-OH group
  - d) Ozonolysis of alkene in the presence of zinc gives aldehydes or ketones/mixture of aldehyde and ketones
- 265. Ozonolysis of C<sub>7</sub>H<sub>14</sub> give methanal and 2–methyl pentan-3-one. The  $C_7H_{14}$  is
  - a) 2-ethyl 3-methyl but-1-ene
  - b) 3-ethyl-2-methyl but-3-ene
  - c) hept-1-ene
  - d) hept-2-ene
- 266. In the reaction

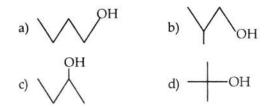
$$C_{2}H_{5}$$

$$CH_{3} - C = O + HCN \longrightarrow A \xrightarrow{H_{3}O^{+}} CH_{3} - C - OH$$

$$COOH$$

The acid obtained will be

- a) L-isomer
- b) D-isomer
- c) 20% 0 + 80% L-isomer
- d) 50% 0 + 50% L isomer
- 267. A substance C<sub>4</sub>H<sub>10</sub>O yield on oxidation C<sub>4</sub>H<sub>5</sub>O which gives oxime and positive iodoform test. The original substance on treatment with conc. H<sub>2</sub>SO<sub>4</sub> gives C<sub>4</sub>H<sub>5</sub>. The structure of compound is



- 268. Which of the following react with NaOH produces acid and alcohol?
  - a) C<sub>6</sub>H<sub>5</sub>CHO
- b) CH<sub>3</sub>-CHO
- c) CH<sub>2</sub>COOH
- d) C.H.COOH
- 269. Which of the following is not correctly matched?

$$a) > C = O \xrightarrow{\text{Clemmenson's}} > CH_2$$

$$(b) > C = O \xrightarrow{\text{Wolff-Kishner}} > CH - OH$$

c) 
$$-\text{COC1} \xrightarrow{\text{Rosenmund's}} - \text{CHO}$$

d) 
$$-C \equiv N \xrightarrow{\text{Stephen's}} - CHO$$

270. Compound A of formula C<sub>3</sub>H<sub>5</sub>O is treated with acidic KMnO<sub>4</sub> to form product B of formula C<sub>3</sub>H<sub>6</sub>O, which form shining silver mirror on warming with ammoniacal AgNO<sub>3</sub>, when B is. treated with NH, CONHNH, in HCl and sodium acetate gives the product C. Identity the structure of C.

a) 
$$CH_3 - C = N - NH - C - NH_2$$

$$CH_3$$

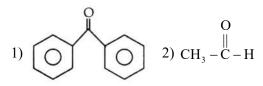
b) 
$$CH_3 - C = N - C - NH - NH_2$$

$$CH_3$$

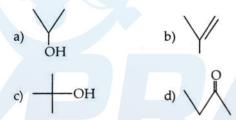
$$\begin{array}{c} & O \\ \parallel \\ c) \ CH_3 - CH_2 - CH = N.NH-C - NH_2 \end{array}$$

d) 
$$CH_3 - CH_2 - CH = N - C - NH - NH_2$$

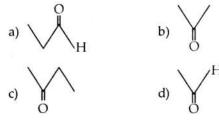
271. The correct order of reactivity of Ph-MgCl with



- a) 1 > 2 > 3
- b) 3 > 2 > 1
- c) 2 > 3 > 1
- d) 2 > 1 > 3
- 272. Which one of the following oxidised to corresponding carbonyl compound?



- 273. 2,2—diethoxy butane is obtained from ethanol and what?
  - a) CH<sub>2</sub> CH<sub>2</sub> CH<sub>2</sub> CHO
  - b)  $CH_3 COCH_2 CH_3$
  - c)  $CH_3 CH CH_2 CH_3$ OH
  - d)  $CH_3 CH_2 CH_2 CH_2 I$
- 274. Which reagent is used to convert butan–2–one to propionic acid?
  - a) Tollen's reagent
  - b) Fehling's solution
  - c) acidic KMnO<sub>4</sub>
  - d) PCC
- 275. The compounds HCHO and CH<sub>3</sub>CHO on intermolecular aldol condensation gives.
  - a) HO-CH<sub>2</sub>-CH<sub>2</sub>-CHO
  - b) CH<sub>3</sub>-CHOH-CH<sub>2</sub>OH
  - c) CH<sub>3</sub>-CHOH-CH<sub>3</sub>
  - d)  $CH_3$ -CHOH- $CH_2$ - $CH_2OH$
- 276. Nucleophilic addition reaction is most favored in



- 277. The smallest ketone and it's functional isomer are treated with NH<sub>2</sub>–OH to form oxime
  - a) two different oxime are formed
  - b) three different oxime are formed
  - c) two oximes are optically active
  - d) all oximes are optically active

278. R-CO-R' 
$$\xrightarrow{\text{HCN}}$$
 A  $\xrightarrow{\text{B}}$  R - C - R'  $\xrightarrow{\text{CH}_2 - \text{NH}_2}$ 

A and B are

a) 
$$A = R - C - R'$$
  $B = Na + C_2H_5OH$ 

$$CN$$

b) 
$$A = \begin{pmatrix} OH \\ | \\ | \\ COOH \end{pmatrix}$$
  $B = H_3O^+$ 

$$\begin{array}{c} OH \\ | \\ C) A = R - C - R' \qquad B = NH_3 \\ | \\ CN \end{array}$$

$$R'$$

$$\downarrow$$

$$d) A = R - C H - CN B = NaOH$$

- 279. On reaction with ketone with hydroxyl amine give ketoxime which on reduction produces
  - a) carboxylic acid
- b) 1° amine
- c) 2°-amine
- d) amide
- 280. Aldehyde with NH<sub>2</sub>-NH<sub>2</sub> forms
  - a) aniline
- b) nitrobenzene
- c) hydrazine
- d) hydrazone
- 281.  $C_3H_6O \xrightarrow{OH^-} Mesityloxide$ .

The C<sub>3</sub>H<sub>6</sub>O is

- a) CH<sub>3</sub>-CH<sub>2</sub>-CHO
- b) CH, COCH,
- c)  $CH_2 = CH CH_2 OH$
- d)  $CH_3 C = CH_3$
- 282. 3–hydroxy 2–methyl pentanal is formed when X react with Y in dilute Z solution. What are X,Y,Z
  - X
- Y
- Z
- a) CH<sub>2</sub>-CHO CH<sub>2</sub>-CHO NaOH
- b) C<sub>2</sub>H<sub>5</sub>-CHO CH<sub>2</sub>-CHO NaOH
- c) C,H,-CHO C,H,-CHO NaOH
- d) C<sub>2</sub>H<sub>5</sub>-CHO C<sub>2</sub>H<sub>5</sub>-CHO NaCl
- 283. A base can absract an  $\alpha$  H atom from
  - a)  $CH_{2} = CH_{2}$
- b)  $CH \equiv CH$
- c) CH<sub>2</sub> CHO
- d)  $C_6H_5$  CHO
- 284. Which of the following pathways produces 2pentanone
  - 1. pent-1-yne is treated with H<sub>2</sub>SO<sub>4</sub> + HgSO<sub>4</sub> and water
    - 2. 3-methyl hex-2- ene is treated with O, followed by hydrolysis
    - 3. n-butylmagnesium halide is reacted with formaldehyde
    - 4. hydroboration oxidation of pent-1-ene
    - a) 1, 3
- b) 1, 2
- c) 2, 4
- d) 1, 3, 4
- 285. Which of the following reagent react with same manner with

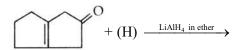


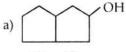
H - CHO, CH<sub>3</sub> - CHO,

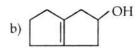
CH, COCH,

- a) Fehling's solution
- b) NH<sub>3</sub>
- c) HCN
- d) NaOH
- 286. Which statement is correct with ethanal and propanone
  - 1) Both react with NaHSO,
  - 2) Both gives iodoform test
  - 3) Both can be reduced into alcohols
  - 4) Both undergoes aldol condensation
  - a) 1, 3
- b) 2, 3
- c) 2, 4
- d) all of these
- 287. Which of the following react same manner with H-CHO and C<sub>6</sub>H<sub>5</sub>-CHO?
  - a) Fehling's solution b) CH<sub>3</sub>-MgX

- c) NH<sub>2</sub>
- d) dil. NaOH
- 288. Acetone gives addition elimination reaction with
  - 1) NH<sub>2</sub>-OH
- 2) NH<sub>2</sub>-CONH-NH<sub>3</sub>
- 3) NaHSO<sub>3</sub>
- 4) CH<sub>3</sub>-NH<sub>2</sub>
- a) 2,4
- b) 3,4
- c) 1, 2, 4
- d) 1,2,3
- 289. An carbonyl compounds gives nucleophilic addition reaction with
  - 1) HCN
- 2) NaSHO,
- 3) R-MgX
- 4) NH,-NH,
- a) 1, 2
- b) 2, 3
- c) 1, 2, 4
- d) 1, 2, 3
- 290. Product of the following reaction is









- 291. The product formed when HCHO is heated with conc.KOH
  - a) CH,CHO
- b) CH,OH
- c) C,H,
- d) CH<sub>4</sub>
- 292. Formaldehyde reacts with NH, to form urotropine which has the composition
  - a)  $(CH_{2})_{5}N_{5}$
- b) (CH<sub>2</sub>)<sub>5</sub>N<sub>5</sub>
- c)  $(CH_{2})_{4}N_{6}$
- d)  $(CH_{2})_{6}N_{4}$
- 293. Pinacol is converted in to pinacolone by
  - a) rearrangment
- b) oxidation
- c) reduction
- d) hydrolysis
- 294. Aldehydes with Tollen's reagent are
  - a) reducing agent
- b) oxidising agents
- c) hydrating agents
- d) bleaching agents
- 295. Low reactivity of ketones with respect to aldehydes is due to
  - a) greater +I effect of alkyl group
  - b) greater steric hindrance of alkyl group
  - c) both 'a' and 'b'
  - d) less steric hindrance of alkyl group
- 296. Aldehydes can be distinguished from ketones by using
  - a) dil. NaOH
- b) R-MgX
- c) Na<sub>2</sub>CO<sub>2</sub>
- d) Schiff's reagent
- 297. Which of the following are generally used for

preparing derivatives of aldehydes and ketones?

- a) Hydrocyanic acid b) Hydroxylamine
- c) Phenyl hydrazine d) All of these
- 298. Which of the following is an example of aldol condensation?
  - a)  $2CH_{3}CHO \xrightarrow{OH^{-}} CH_{3}CH(OH)CH_{2}CHO$
  - b)  $C_6H_5CHO \xrightarrow{OH} C_6H_5CH_2OH$
  - c) HCHO  $\stackrel{\text{OH}^-}{\longrightarrow}$  HCH<sub>2</sub>OH
  - d)  $C_6H_5CHO + HCHO \xrightarrow{OH} C_6H_5CH_9OH$
- 299. Which of the following can be used to differentiate between acetaldehyde and prop anal?
  - a) Ammonical AgNO,
  - b) CuSO<sub>4</sub> + Na-K tartarate in NaOH
  - c) I<sub>2</sub> in the presence of base
  - d) Decolourised Fuchin
- 300. Molecular formula 'A' ( $C_4H_5O$ ) reacts with  $CH_3Mg1$  gives 3—methyl 2—butanol. The structure of A is
  - a) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CHO b) C<sub>2</sub>H<sub>5</sub>COCH<sub>3</sub>
  - c) (CH<sub>3</sub>)<sub>2</sub>CHCHO
- d) C<sub>2</sub>H<sub>5</sub>OCH<sub>3</sub>
- 301. When acetone reacts with Ba(OH), it gives
  - a) acetoxime
- b) urotropine
- c) t-butyl alcohol
- d) diacetone alcohol
- 302. Which of the following aldehydes does not reduce Fehling solution readily?
  - a) Acetaldehyde
- b) Formaldehyde
- c) Propanal
- d) Benzaldehyde
- 303. 1–propanol is obtained from  $C_2H_5$  MgI and what?
  - a) CH<sub>3</sub>CHO
- b) HCHO
- c) HCN
- d) C<sub>2</sub>H<sub>5</sub>CHO
- 304. Treatment of propionaldehyde with dil. NaOH solution gives
  - a) CH<sub>3</sub>CH<sub>2</sub>COOCH<sub>2</sub>C<sub>2</sub>H<sub>2</sub>
  - b) CH<sub>3</sub>CH<sub>2</sub>CHOHCH<sub>2</sub>CH<sub>2</sub>CHO
  - c) CH<sub>3</sub>CH<sub>2</sub>CH(OH)CH(CH<sub>3</sub>)CHO
  - d) CH<sub>3</sub>CH<sub>2</sub>COCH<sub>2</sub>CH<sub>2</sub>CHO
- 305. Oxidation number of carbon in HCHO is,
  - a) 0
- b) 2
- c) 3
- d) 4
- 306. Which of the following compound does not undergo aldol condensation?
  - a) Acetaldehyde
- b) Prop anal
- c) Ethyl methyl ketoned) Benzaldehyde

- 307. Which of the following compound undergoes aldol condensation?
  - a) CH<sub>3</sub>CHO
- b) C,H,CHO
- c) CH<sub>3</sub>COCH<sub>3</sub>
- d) All
- 308. The reaction,

CH<sub>3</sub>CH(OH)CH<sub>2</sub>CHO

represents

- a) Cannizzaro reaction
- b) Aldol condensation
- c) Wurtz reaction
- d) Mendius reaction
- 309. Mesityl oxide is formed by the condensation of
  - a) acetaldehyde
- b) acetone
- c) prop anal
- d) formaldehyde
- 310. The most probable compound whose molecular formula is C<sub>3</sub>H<sub>6</sub>O and which can give Tollen's reagent test
  - a) CH,CH,CHO
- b) CH,COCH,
- c) CH,OCH,CH,
- d) CH<sub>2</sub> = CHCHPH
- 311. When two molecules of HCHO react in presence of base to produce CH<sub>3</sub>OH and HCOONa, the reaction is called
  - a) Wurtz reaction
- b) Cannizzaro reaction
- c) Aldol condensation d) Hoffman reaction
- 312. Aldol condensation of acetaldehyde results in the formation of
  - a) CH<sub>3</sub>COCHOHCH<sub>3</sub>
  - b) СН, СНОНСН, СНО
  - c) CH<sub>2</sub>CH<sub>2</sub>CHOHCHO
  - d) CH<sub>3</sub>CH<sub>2</sub>OH + CH<sub>3</sub>COOH
- 313. Which of the following reagents can not be used to distinguish between pentanal and 2-pentanone?
  - a) I, in NaOH
- b) Fehling solution
- c) Na metal
- d) Tollen's reagent
- 314. Acetaldehyde reacts with NH<sub>2</sub>OH to give
  - a) acetal amine
- b) acetal oxide
- c) acetaldoxime
- d) amino acetal
- 315. Which of the following statement is true?
  - a) Aldehydes are less susceptible to oxidation than ketones
  - b) All aldehydes undergo Cannizzaro reaction
  - c) Aldehydes are more susceptible to oxidation than ketones
  - d) Formaldehyde does not react with ammonia
- 316. Silver mirror test is given by

### **Alcohols, Phenols and Ethers**

- a) aldehyde
- b) ketones
- c) amines
- d) ethers
- 317. Which of the following reagent is used to convert  $> C = O \text{ to } > CH_2$ ?
  - a)  $ZnHg + H_2O$
- b) ZnHg + conc.HCl
- c) NaHg + H<sub>2</sub>O
- d) Sn + conc. HC1
- 318. The formation of acetone sodium bisulphite from a acetone is an example of
  - a) electrophilic addition
  - b) nucleophilic addition
  - c) nucleophilic substitution
  - d) electrophilic substitution
- 319. In which reaction new C–C bond does not form
  - a) Aldol
- b) Cannizzaro's
- c) Wurtz
- d) HCN + RMgX
- 320. Mesityl oxide is
  - a) ketone
- b) aldehyde
- c) ester
- d) ethers
- 321. Schiff's reagent is obtained by passing
  - a) SO<sub>2</sub> gas in aq. solution of rosaniline
  - b) NO, gas in aq. solution of rosaniline
  - c) O, gas in aq. solution of rosaniline
  - d) CO, gas in aq. sol. of rosaniline
- 322. Which of the following can reduce Tollen's reagent but not Fehling solution?
  - a) C<sub>6</sub>H<sub>5</sub>CHO
- b) CH, CHO
- c) C<sub>2</sub>H<sub>2</sub>CHO
- d) CH, COCH,
- 323. An aldehyde on oxidation gives
  - a) an alcohol
- b) an acid
- c) an ether
- d) a ketone
- 324. Magenta is
  - a) ale. phenolphthalein
  - b) p-rosaniline hydrochloride
  - c) methyl red
  - d) red litmus
- 325. Aldehyde on reaction with Grignard reagent and subsequent hydrolysis yields
  - a) primary alcohol
- b) tertiary alcohol
- c) secondary alcohol d) dihydroxy alcohol
- 326. Which of the following does not have alpha hydrogen?
  - a) Formaldehyde
  - b) Acetaldehyde
  - c) Dimethyl acetaldehyde
  - d) Acetone
- 327. Rochelle salt is
  - a) ammonium tartarate

- b) sodium potassium tartarate
- c) potassium ammonium tartarate
- d) sodium tartarate
- 328. Fuschin is,
  - a) pink dye of para–rosa–aniline hydrochloride
  - b) Schiff's reagent
  - c) Tollen's reagent
  - d) Fehling solution
- 329. Which of the following is a gas at room temperature?
  - a) CH<sub>2</sub>CHO
- b) HCHO
- c) CH,COCH,
- d) C<sub>2</sub>H<sub>5</sub>OC<sub>2</sub>H<sub>5</sub>
- 330. Clemmenson reduction is carried out with
  - a) H<sub>2</sub>/Pd
  - b) NH<sub>2</sub>NH<sub>2</sub>, followed by treatment with glycolic
  - c) LiAlH<sub>4</sub> in ether
  - d) Zn(Hg)/HCl
- 331. Aldehydes having no  $\alpha$  –hydrogen atom undergo
  - a) Hoffman reaction
  - b) Aldol condensation
  - c) Cannizzaro reaction
  - d) Wurtz reaction
- 332. Which of the following does not react with phenyl hydrazine?
  - a) Ethanol
- b) Ethanal
- c) Acetone
- d) 2 pentanone
- 333. Which of the following compound does not undergo Cannizzaro reaction?
  - a) Ethanol
  - b) Methanal
  - c) Benzaldehyde
  - d) 2,2,4,4—tetra methyl 3—pentanone
- 334. Reduction in presence of amalgamated zinc and conc. HCl is known as
  - a) Mendius reduction
  - b) Wurtz reaction
  - c) Hoffmanns reaction
  - d) Clemmenson's reduction
- 335. Aldehyde having no α-hydrogen undergo disproportionation in presence of strong potash to give a corresponding salt of acid and alcohol.

The reaction is known as,

- a) Wurtz reaction
- b) Aldol condensation
- c) Cannizzaro's reaction
- d) Esterification

- 336. An organic compound contains  $\rm H_2O$  and single carbon. It responds positive to Tollen's reagent. The compound is
  - a) HCHO
- b) CH<sub>2</sub>OH
- c) CH,CHO
- d) none of these
- 337. Formaldehyde gives an addition product with methyl magnesium iodide which on acid hydrolysis gives
  - a) CH<sub>3</sub>OH
- b) C<sub>2</sub>H<sub>5</sub>OH
- c) (CH<sub>2</sub>),CHOH
- d) CH, CHOHCH,
- 338. Formaldehyde and formic acid are distinguished by treating with
  - a) Tollen's reagent
- b) NaHCO,
- c) Fehling solution
- d) Schiff's reagent
- 339. Pick up the correct statements from the following
- a) both aldehydes and ketones use sp<sup>2</sup>-hybrid carbon atoms for their addition reaction
  - b) acetic acid neither reacts with Fehling's solution nor with Tollen's reagent
  - c) chloral is aldehyde
  - d) all are correct
- 340. Butanal is heated with ammonical silver nitrate. The product formed is,
  - a) CH, CH, CH, COOH
  - b) (CH<sub>2</sub>)<sub>2</sub>CHCOOH
  - c) CH<sub>3</sub>COOH + CH<sub>3</sub>OH
  - d) HCOOH + C<sub>2</sub>H<sub>5</sub>COOH
- 341. Fehling solution is
  - a) acidified CuSO<sub>4</sub> solution
  - b) ammonical AgNO<sub>3</sub> solution
  - c) copper sulphate, sodium hydroxide and Rochelle salt
  - d) none of these
- 342. Union of two or more molecules of the same or different compound with elimination of water to form a new substance is known as
  - a) synthesis
- b) polymerisation
- c) condensation
- d) none of these
- 343. Which of the following compound will undergo self aldol condensation inpresence of dilute alkali?
  - a) C<sub>6</sub>H<sub>5</sub>CHO
- b) (CH<sub>2</sub>)<sub>2</sub>CCHO
- c) C,H,CHO
- d) CCl<sub>3</sub>CHO
- 344. When a ketone is condensed into an ketol, the reagent used is
  - a) Na<sub>2</sub>CO<sub>2</sub>
- b) NaHCO,
- c) Br, water
- d) Cl<sub>2</sub>
- 345. Benzaldehyde undergoes auto-oxidation and reduction in presence of

- a) cone, NaOH
- b) Na,CO,
- c) NaHCO<sub>2</sub>
- d) dil. NaOH
- 346. Nucleophilic attack on carbonyl carbon changed its hydridisation from
  - a) sp to  $sp^2$
- b)  $sp^2$  to  $sp^3$
- c) sp<sup>3</sup> to sp<sup>2</sup>
- d) sp to sp<sup>3</sup>
- 347. Which of the following reaction is used for detecting the presence of carbonyl group in aldehydes and ketones?
  - a) Reaction with hydroxylamine
  - b) Reaction with phenyl hydrazine
  - c) Reaction with hydrazine
  - d) All of these
- 348. Which of the following reagent form oxime with carbonyl compound?
  - a) NH,OH
- b) NH,OH
- c) NaOH
- d) CH,N,
- 349. Ketonic form of acetone contains,
  - a) 8-a bonds and 2-n bonds
  - b) 9-a bonds and I-n bond
  - c) 7-a bonds and 3-n bonds
  - d) 4-a bonds and 5-n bonds
- 350. 11-hydroxy butyraldehyde is an example of
  - a) aldol
- b) ketol
- c) ester
- d) alcohol
- 351. Which is most difficult to oxidise?
  - a) HCHO
- b) CH, COCH,
- c) CH<sub>3</sub>CHO
- d) C<sub>2</sub>H<sub>5</sub>CHO
- 352. Aldehydes or ketones having atleast one ahydrogen undergo condensation reaction inpresence of dilute base. This reaction is called
  - a) Aldol condensation
  - b) Cannizzaro's reaction
  - c) Hoffmann's reaction
  - d) Mendius reaction
- 353. The reagent with which both aldehyde and acetone react easily is
  - a) Fehling's reagent
- b) Grignard reagent
- c) Schiff's reagent
- d) Tollen's reagent
- 354. When, acetaldehyde is heated with Tollen's reagent, following is obtained
  - a) Methyl alcohol
- b) Silver acetate
- c) Silver mirror
- d) Formaldehyde
- 355. Which of the following compounds would undergo Cannizzaro's reaction
  - a) Propionaldehyde
- b) Benzaldehyde
- c) Bromobenzene
- d) Acetaldehyde

- 356. Schiff's reagent gives pink colour with
  - a) Aldehydes
- b) Ethers
- c) Ketones
- d) Carboxylic acid
- 357. An aldehyde on oxidation gives
  - a) An alcohol
- b) An acid
- c) A ketone
- d) An ether
- 358. Schiff's reagent is
  - a) Magneta solution decolourised with SO,
  - b) Ammonical cobalt chloride solution
  - c) Ammonical manganese sulphate solution
  - d) Magneta soluation decolourised with chlorine
- 359. Acetone and acetaldehyde are differentiated by
  - a) NaOH + I,
- b)  $Ag(NH_3)_2^+$
- c) HNO,
- d) I,
- 360. Hexamethylene tetramine is used as
  - a) analgesic
- b) antipyretic
- c) urinary antiseptic
- d) all the above
- 361. Benzyl alcohol is obtained from benzaldehyde by
  - a) Fittg's reaction
- b) Cannizaro's reaction
- c) Kolbe's reaction
- d) Wurtz's reaction
- 362. Which of the following gives difference between aldehyde and ketone?
  - a) Fehling's solution
- b) Tollen's reagent
- c) Schiff's reagent
- d) All of the above
- 363. Formaldehyde gives an additive product with ethyl magnesium iodide which on aqueous hydrolysis gives
  - a) isopropyl alcohol
- b) ethyl alcohol
- c) methyl alcohol
- d) n-propyl alcohol
- 364. The reagent which easily reacts with ethanal and propanal is
  - a) Fehling solution
- b) Grignard reagent
- c) Schiff's reagent
- d) Tollen's reagent
- 365. Magneta is
  - a) alkaline phenolphthalein
  - b) methyl red
  - c) p-rosaniline hydrochloride
  - d) red litmus
- 366. Grignard reagent on reaction with ketones forms
  - a) tertiary alcohol
- b) secondary alcohol
- c) acetic acid
- d) acetaldehyde
- 367. The compound which reacts with Fehling solution is
  - a) C<sub>6</sub>H<sub>5</sub>COOH
- b) HCOOH
- c) C<sub>6</sub>H<sub>5</sub>CHO
- d) CH, ClCH,
- 368. Acetone gives test with

- a) Phenyl hydrazine b) Fehling solution
- c) Schiff's reagent
- d) All the above
- 369. An aldehyde can undergo the aldol condensation having
  - a) an aromatic ring
  - b) no alpha H atom
  - c) at least one  $\alpha$  –H atom
  - d) at least one  $\beta$ -H atom
- 370. An organic compund 'X' having molecular formula C<sub>5</sub>H<sub>10</sub>O yields phenyl hydrazone and gives negative response to the iodoform test and Tollen's test. It produces n–pentane on reduction 'X' could be:
  - a) pentanal
- b) 2–pentanone
- c) 3–pentanone
- d) n-amyl alcohol
- 371. Aldol condensation will not take place in
  - a) HCHO
- b) CH<sub>3</sub>CHO
- c) CH<sub>3</sub>COCH<sub>3</sub>
- d) CH<sub>3</sub>CH<sub>2</sub>CHO
- 372. CH<sub>3</sub>COCH<sub>3</sub> can be converted to CH<sub>3</sub>CH<sub>2</sub>CH<sub>3</sub> by the action of
  - a) HIO<sub>3</sub>
- b) Zn Hg + HC1
- c) HNO<sub>2</sub>
- d) H<sub>3</sub>PO<sub>3</sub>
- 373. The compound most suitable for the preparation of cyanohydrin is
  - a) C<sub>2</sub>H<sub>5</sub>COOH
- b) C,H,COC,H,
- c)  $C_6H_5NH_2$
- d)  $C_2H_5-C_2H_5$
- 374. Baking powder is
  - a) NaHCO<sub>3</sub> c) Na<sub>2</sub>CO<sub>3</sub>
- b) NaHCO<sub>3</sub>.6H<sub>2</sub>O d) Na<sub>2</sub>CO<sub>3</sub>.10H<sub>2</sub>O
- 375. Which one of the following undergoes reaction with 50% sodium hydroxide solution to give the corresponding alcohol and acid?
  - a) Butanal
- b) Formaldehyde
- c) Phenol
- d) Benzoic acid
- 376. Which of the following compounds can be regenrated to carbonyl compounds?
  - a) Urotropin
- b) Bisulphite complex
- c) Cynohydrine
- d) Diacetone amine
- 377. What is 'C' in the following reaction?

$$C_6H_5CN + C_6H_4MgI \xrightarrow{i) dry ether} \rightarrow$$

$$A \xrightarrow{NH_2OH} B \xrightarrow{LiAlH_4} C$$

- a) diphenyl methane
- b) diphenyl methanamine
- c) nitrodiphenyl methane
- d) diphenyl

### **ANSWERS - MCQ'S**

### SECTION - I : ALDEHYDES AND KETONES

			_				_		_						_				
1.	a)	2.	b)	3.	b)	4.	c)	5.	c)	6.	c)	7.	c)	8.	b)	9.	b)	10.	b)
11.	b)	12.	c)	13.	a)	14.	c)	15.	d)	16.	b)	17.	c)	18.	d)	19.	d)	20.	b)
21.	d)	22.	b)	23.	d)	24.	b)	25.	c)	26.	d)	27.	b)	28.	c)	29.	a)	30.	b)
31.	d)	32.	c)	33.	b)	34.	b)	35.	b)	36.	b)	37.	c)	38.	d)	39.	b)	40.	b)
41.	a)	42.	b)	43.	c)	44.	b)	45.	a)	46.	c)	47.	d)	48.	a)	49.	a)	50.	c)
51.	c)	52.	a)	53.	b)	54.	c)	55.	c)	56.	d)	57.	a)	58.	d)	59.	b)	60.	a)
61.	b)	62.	a)	63.	d)	64.	c)	65,	b)	66.	b)	67.	b)	68.	d)	69.	c)	70.	a)
71.	c)	72.	a)	73.	c)	74.	b)	75.	d)	76.	c)	77.	b)	78.	a)	79.	a)	80.	c)
81.	a)	82.	c)	83.	c)	84.	a)	85.	d)	86.	c)	87.	d)	88.	a)	89.	c)	90.	b)
91.	d)	92.	b)	93.	b)	94.	b)	95.	d)	96.	d)	97.	b)	98.	d)	99.	b)	100.	a)
101.	c)	102.	a)	103.	c)	104.	<b>b</b> )	105.	c)	106.	d)	107.	a)	108.	b)	109.	b)	110.	c)
111.	d)	112.	c)	113.	c)	114.	a)	115.	b)	116.	b)	117.	b)	118.	d)	119.	d)	120.	a)
121.	a)	122.	b)	123.	b)	124.	d)	125.	c)	126.	c)	127.	c)	128.	b)	129.	b)	130.	d)
131.	b)	132.	b)	133.	d)	134.	c)	135.	d)	136.	c)	137.	d)	138.	c)	139.	a)	140.	a)
141.	a)	142.	b)	143.	a)	144.	c)	145.	c)	146.	d)	147.	c)	148.	c)	149.	c)	150.	b)
151.	a)	152.	b)	153.	a)	154.	d)	155.	b)	156.	a)	157.	b)	158.	b)	159.	c)	. 160.	d)
161.	c)	162.	a)	163.	b)	164.	c)	165.	c)	166.	d)	167.	d)	168.	b)	169.	d)	170.	d)
171.	d)	172.	d)	173.	d)	174.	a)	175.	a)	176.	b)	177.	a)	178.	d)	179.	b)	180.	a)
181.	a)	182.	a)	183.	c)	184.	c)	185.	a)	186.	c)	187.	d)	188.	a)	189.	d)	190.	b)
191.	b)	192.	c)	193.	b)	194.	d)	195.	d)	196.	b)	197.	b)	198.	d)	199.	b)	200.	d)
201.	d)	202.	d)	203.	b)	204.	c)	205.	d)	206.	ь)	207.	b)	208.	c)	209.	b)	210.	b)
211.		212.	b)	213.	b)	214.	b)	215.	d)	216.	c)	217.	b)	218.	a)	219.	b)	220.	c)
221.	b)	222.	c)	223.	d)	224.	c)	225.	b)	226.	c)	227.	a)	228.	d)	229.	a)	230.	d)
231.		232.	d)	233.	c)	234.	d)	235.	d)	236.	b)	237.	c)	238.	c)	239.	b)	240.	a)
241.	a)	242.	c)	243.	d)	244.	b)	245.	c)	246.	c)	247.	c)	248.	c)	249.	c)	250.	d)
251.		252.	a)	253.	b)	254.	c)	255.	b)	256.	b)	257.	d)	258.	d)	259.	a)	260.	b)
261.		262.	d)	263.	c)	264.	d)	265.	a)	266.	d)	267.	c)	268.	a)	269.	b)	270.	c)
271.	c)	272.	a)	273.	b)	274.	c)	275.	a)	276.	d)	277.	a)	278.	a)	279.	b)	280.	d)
281.	b)	282.	c)	283.	c)	284.	b)	285.	c)	286.	d)	287.	b)	288.	c)	289.	d)	290.	b)
291.	5=35	292.	d)	293.	a)	294.	a)	295.	c)	296.	d)	297.	d)	298.	a)	299.	c)	300.	c)
301.		302.	d)	303.		304.	0.00	305.	a)	306.	d)	307.	d)	308.	b)	309.	b)	310.	a)
311.		312.		313.	c)	314.	c)	315.		316.	a)	317.	b)	318.	b)	319.	b)	320.	a)
321.		322.		323.	b)	324.	b)	325.		326.	a)	327.	b)	328.	a)	329.	b)	330.	d)
331.		332.	a)	333.	a)	334.	d)	335.	c)	336.	a)	337.	b)	338.	b)	339.	d)	340.	a)
341.		342.	c)	343.	c)	344.	a)	345.		346.	b)	347.	d)	348.	b)	349.	b)	350.	a)
351.		352.		353.	b)	354.	c)	355.	b)	356.	a)	357.	ь)	358.	a)	359.	b)	360.	c)
361.		362.		363.		364.		365.	c)	366.	a)	367.	b)	368.	a)	369.	c)	370.	c)
371.	a)	372.	b)	373.	b)	374.	a)	375.	b)	376.	b)	377.	b)						
													1	7	33	51	,	Si .	)



### **MULTIPLE CHOICE QUESTIONS**

### SECTION - II: CARBOXYLIC ACIDS

### INTRODUCTION, NOMENCLATURE, **ISOMERISM**

- 1. General formula for carboxylic acid is
  - a) C<sub>n</sub>H<sub>n</sub>O<sub>2</sub>
- b)  $C_n H_{2n} O_2$
- c) C<sub>n</sub>H<sub>2n</sub>O
- d)  $C_{n}H_{2n+2}O_{2}$
- 2. The general formula of an ester, where R represents an alkyl group, is
  - a) ROH
- b) RCOOH
- c) RCOOR
- d) ROR
- Vinegar contains acetic acid nearly
  - a) 6–10%
- b) 30%
- c) 12–15%
- d) 20%
- The bite of honey—bee inject into our body 4.
  - a) acetic acid
- b) formic acid
- c) butyric acid
- d) carbolic acid
- Glacial acetic acid at low temperature is a
  - a) thin liquid
- b) viscous liquid
- c) ice like solid
- d) semi solid
- The suffix used in carboxylic acid is 6.
  - a) -oic
- b) -ane
- c) ene
- d) -al
- Formic acid owes its origin to
  - a) milk
- b) butter
- c) red ants
- d) vinegar
- Vinegar obtained from canesugar contains
  - a) Citric acid
- b) Lactic acid
- c) Acetic acid
- d) Palmitic acid
- Write the IUPAC name of CH<sub>3</sub>CH<sub>2</sub>COOH.
  - a) ethyl formic acid
  - b) ethyl carboxylic acid
  - c) ethane methanoic acid
  - d) propanoic acid
- 10. IUPAC name m-toluic acid is
  - a) 3-methyl benzoic acid
  - b) 2-methyl benzoic acid
  - c) 4-methyl benzoic acid
  - d) 2,4-dimethyl benzoic acid
- 11. IUPAC name of carboxylic acid is
  - a) alkinoic acid
- b) alkanoic acid
- c) alkenoic acid
- d) alkyl alkanoate
- 12. IUPAC name of ester is
  - a) alkoxy alkane
- b) alkoxyalkene

- d) alkyl alkanoate c) alkane alkanoate
- 13. IUPAC name of isobutyric acid is
  - a) 3-methyl propionic acid
  - b) 2-methyl propanoic acid
  - c) 3-methyl butanoic acid
  - d) isobutanoic acid
- 14. IUPAC name of following compound is (CH<sub>2</sub>),CHCOOC,H<sub>5</sub>
  - a) ethyl isobutanoate
  - b) ethyl isobutyrate
  - c) ethyl 2–methyl propanoate
  - d) ethyl 1-methyl ethanoate
- 15. IUPAC name of following compound is
  - $(CH_3)_3C-CH_2-COOH$
  - a) 2, 2-dimethyl butanoic acid
  - b) 3, 3-dimethyl butanoic acid
  - c) 2, 3-dimethyl propanoic acid
  - d) 2, 3-dimethyl butanoic acid
- 16. IUPAC name of following compound is  $C_{2}H_{3}$ -COOCH(CH<sub>3</sub>),
  - a) isopropyl propanoate
  - b) isopropyl propionate
  - c) 2-propyl propanoate
  - d) 2-propyl ethanoate
- 17. Structure of ethyl ethanoate is
  - a) CH<sub>3</sub>COOC<sub>2</sub>H<sub>5</sub> b) C<sub>2</sub>H<sub>5</sub>COOCH<sub>3</sub>
  - c) H-COOC<sub>2</sub>H<sub>5</sub>
- d) H-COOCH,
- 18. An organic compound having the molecular formula C<sub>2</sub>H<sub>4</sub>O<sub>2</sub> is
  - a) formic acid
- b) acetic acid
- c) ethyl acetate
- d) propionic acid
- 19. Aliphatic carboxylic acids shows isomer of type,
  - a) position
- b) chain
- c) functional
- d) all of these
- 20. Carboxylic acid is isomeric with
  - a) aldehydes
- b) alcohols
- c) esters
- d) saturated ketones
- 21. Acids and esters are
  - a) chain isomers
- b) position isomers
- c) functional isomers d) metamers
- 22. Molecular formula C<sub>2</sub>H<sub>4</sub>O<sub>2</sub> represents a) acids
  - b) esters
  - c) aldehydes
- d) both 'a' and 'b'
- 23. Molecular formula C<sub>2</sub>H<sub>4</sub>O<sub>2</sub> shows

- a) chain isomerism
- b) position isomerism
- c) functional isomerism
- d) metamerism
- 24. IUPAC name of following compound is
  - a) 2-carboxy phenol
  - b) 2-carboxy benzyl alcohol
  - c) 2-hydroxy benzoic acid
  - d) 2-hydroxy benzene carboxylic acid
- 25. How many carboxylic acids are possible for  $C_4H_5O_7$ ?
  - a) 2
- b) 3
- c) 4
- d) 5
- 26. Total number of isomeric acids can be calculate by formula
  - a)  $I = 2^{n-1}$
- b)  $I = 2^{n-3}$
- c)  $I = 2^{n}$
- d)  $I = 2^{n-2} 1$
- 27. Citric acid is
  - a) monocarboxylic acid
  - b) dicarboxylic acid
  - c) tricarboxylic acid
  - d) fatty acid
- 28. The following structure is
  - a) oxalic acid
- b) phthalic acid
- c) fumaric acid
- d) succinic acid
- 29. The acid which is obtained from plant
  - a) propionic acid
- b) valeric acid
- c) fumaric acid
- d) malic acid
- 30. Lemon is sour due to
  - a) oxalic acid
- b) tartaric acid
- c) citric acid
- d) malic acid
- 31. Saturated monocarboxylic acid is second oxidative product of
  - a) 1° alcohol
- b) 2° alcohol
- c) both a & b
- d) ketones
- 32. Monocarboxylic acid shows functional isomerism with
  - 1) Ester
- 2) aldehydes
- 3) ketones
- 4) ethers
- a) 1, 2
- b) 1
- c) 2, 3
- d) 3, 4
- 33. IUPAC name of malic acid is
  - a) 3-hydroxy butanoic acid
  - b) 2-hydroxy butanedioic acid

- c) 2-hydroxy butanoic acid
- d) 2-hydroxy pentanedioic acid
- 34. IUPAC name of  $\beta$ -methyl butyric acid is
  - a) 3-methyl butanoic acid
  - b) 4-methyl pentanoic acid
  - c) 2-methyl butanoic acid
  - d) 2-methyl propanoic acid
- 35. IUPAC name of following compound is
  - a) 4-methyl cyclohexane l-carboxyltc acid
  - b) 4-carboxyl I-methyl cyclohexane
  - c) m-Toluic acid
  - d) o-Toluic acid
- 36. IUPAC name of following compound is
  - a) 2-oxo butane dioic acid
  - b) 2-oxo propane 1, 3-dicarboxylic acid
  - c) 2-oxo 1, 5-pentane dioic acid
  - d) Hexane dioic acid
- 37. The IUPAC name of caproic acid is
  - a) pentanoic acid
  - b) 2-phenyl ethanoic acid
  - c) 2-phenyl propanoic acid
  - d) hexanoic acid
- 38. Which of the following is isobutyric acid
- 39. Which of the following is 2-phenyl pentanoic acid?
- 40. n-valeric acid is functional isomer of
  - a) methyl propionate b) n-butyl formate
  - c) propyl isobutyrate d) ethyl isobutyrate
- 41. Butyric acid and isobutyric acid are
  - a) chain isomers
- b) position isomers
- c) metamers
- d) tautomers
- 42. Which isomers of  $C_5H_{10}O_2$  shows optical isomerism?
  - a) 2-methyl butanoic acid
  - b) valeric acid
  - c) neo-valeric acid
  - d) 3-methyl butanoic acid
- 43. How many carboxylic acids are possible for molecular formula  $C_5H_{10}O_2$ ?
  - a) 3

b) 4

c) 5

- d) 6
- 44. Molecular formula C<sub>4</sub>H<sub>5</sub>O<sub>5</sub> shows

- 1) chain isomerism
- 2) metamerism
- 3) functional isomerism
- 4) optical isomerism
- a) 1, 2, 3
- b) 3, 4
- c) 2, 3, 4
- d) all of these
- 45. Common name of the following compound is
  - a) o-Toluic acid
- b) m-Toluic acid
- c) p-Toluic acid
- d) Phthalic acid
- 46. IUPAC name of malonic acid is
  - a) propane dioic acid b) ethane dioic acid
  - c) butane dioic acid
- d) pentane dioic acid
- 47. Carbon atom in carboxyl group is
  - a) sp-hybridised state
  - b) sp<sup>2</sup> hybridised state
  - c) sp<sup>3</sup> hybridised state
  - d) sp<sup>3</sup> d hybridised state
- 48. Which of the following is 3–oxo cyclohex–4–ene –1–carboxylic acid
- 49. Which of the following is not fatty acid?
  - a) propionic acid
- b) butyric acid
- c) iso-butyric acid
- d) iso-phthalic acid
- 50. IUPAC name of following acid is
  - a) isophthalic acid
  - b) benzene 1, 3-dicarboxylic acid
  - c) phenyl 1, 3-dicarboxylic acid
  - d) terephthalic acid
- 51. Carboxylic acid donot give characteristic property of
  - a) R group
- b) COOH group
- c) > C = O group
- d) O H group

### PREPARATION METHODS

52. Consider the following sequence of reactions and identify the final product (Y)

$$CH_3CH_2CH_2Br \xrightarrow{Mg} X \xrightarrow{CO_2/H_3O^+} Y$$

- a) CH, CH, COOH
- b) (CH<sub>3</sub>),CHCOOH
- c) CH,CH,CH,COOH
- d) CH,CH,CH,COOH
- 53. Solid carbondioxide and RMgX produces
  - a) alkanoic acid
- b) alkanal
- c) alkanone
- d) alkyl alkanoate
- 54. Carbonation of CH<sub>3</sub>MgI gives an organic compound. This compound is also obtained by

- a) hydrolysis of acetonitrile by a mineral acid
- b) oxidation of methyl alcohol
- c) hydrolysis of isoacetonitrile by a mineral acid
- d) hydrolysis of methyl formate with dilute mineral acid
- 55. Propyl propionate is prepared from which of the following
  - a) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>OH, CH<sub>3</sub>CH<sub>2</sub>COOH
  - b) CH<sub>3</sub>COOH, CH<sub>3</sub>(CH<sub>2</sub>)<sub>2</sub>COOH
  - c) CH<sub>3</sub>CHOHCH<sub>3</sub>, CH<sub>3</sub>CH<sub>2</sub>COOH
  - d) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>OH, (CH<sub>3</sub>)<sub>2</sub>CHCOOH
- 56. A mixture of carboxylic acid (A) and alcohol (B) on heating give and ester (C) having molecular mass 74. What is (C)?
  - a) HCOOC<sub>2</sub>H<sub>5</sub>
- b) CH<sub>3</sub>COOCH<sub>3</sub>
- c) Both a and b
- d) None of these
- 57. CH<sub>3</sub>-CH<sub>2</sub>-COOH is can be obtained in the following synthesis
  - a)  $CH_3 CH_2 CN \xrightarrow{OH^-}$
  - b) CH<sub>2</sub> CH<sub>2</sub> CN  $\xrightarrow{\text{H}_3\text{O}^+}$
  - c)  $CH_3 CH_2 CN \xrightarrow{N_2 + Ni}$
  - d)  $CH_3 CH_2 CN \xrightarrow{CH_3MgX}$

O

- 58. CH<sub>3</sub>-C -CH<sub>3</sub> can be converted to CH<sub>3</sub>-COOH by following method.
  - 1)  $CH_3 COCH_3 \xrightarrow{I_2 + NaOH}$
  - 2)  $CH_3 COCH_3 \xrightarrow{K_2Cr_2O_7 + H^+}$
  - 3)  $CH_3 COCH_3 \xrightarrow{Ag(NH_3)_2^+}$
  - a) only 2
- b) 1 and 2
- c) 2 and 3
- d) 1 and 3
- 59. Which is leaving group in the hydrolysis of CH<sub>3</sub>COOC<sub>2</sub>H<sub>5</sub>?
  - a)  $C_2H_5^+$
- b) C<sub>2</sub>H<sub>5</sub>O
- c) CH<sub>3</sub>COO-
- d) CH<sub>2</sub>
- 60. For the preparation of isopropyl acetate from esterification. The compounds used as,
  - a) CH<sub>3</sub>COOH and CH<sub>3</sub>CH<sub>2</sub>OH
  - b) (CH<sub>3</sub>)<sub>3</sub>CHCOOH and CH<sub>3</sub>OH
  - c) CH<sub>2</sub>COOH and (CH<sub>2</sub>)<sub>2</sub>CHOH
  - d) C<sub>2</sub>H<sub>5</sub>COOH and CH<sub>2</sub>OH
- 61. Find out X in the following reaction

$$X+CH_3OH \xrightarrow{conc.H_2SO_4} C_2H_5COOCH_3 + H_2O$$

- a)  $C_2H_5-X$
- b) (CH,CO),O
- c) CH,COOH
- d) C<sub>2</sub>H<sub>5</sub>COOH
- 62. Ethyl propionate is formed from ethyl iodide and what?
  - a) C<sub>2</sub>H<sub>5</sub>COOH
- b) C<sub>2</sub>H<sub>5</sub>COOAg
- c) (C,H,CO),O
- d) All of these
- 63. Propanoic acid is the oxidative product of,
  - a) ethanol
- b) 3-pentanone
- c) 1-butanol
- d) 2-propanol
- 64. Reaction

 $CH_3COOH + C_2H_5OH \rightarrow CH_3COOC_2H + H_2O$  is an illustration of

- a) dehydration
- b) esterification
- c) neutralization
- d) dehydrogenation
- 65. R-CH<sub>2</sub>-CH<sub>2</sub>OH can be converted into RCH<sub>2</sub>COOH. The reaction is
  - a) reduction
- b) hydrolysis
- c) oxidation
- d) decarboxylation
- 66. If hydrolysis of ester is carried out inpresence o an acid, using water containing radioactive oxygen atom, the product most likely to be radioactive is,
  - a) alcohol
- b) ester
- c) acid
- d) both 'a' and 'c'
- 67. If acetic acid reacts with methyl alcohol containing labelled oxygen atom. Inpresence of dry HCl, the labelled oxygen atom, at the complete reaction will be found in
  - a) methyl acetate
- b) water
- c) may be 'a' and 'b'
  - d) not predictable
- 68. Propionic acid is obtained from dry ice and what?
  - a) CH<sub>2</sub>MgI
- b) C<sub>2</sub>H<sub>2</sub>MgBr
- c) C<sub>2</sub>H<sub>2</sub>MgBr
- d) None of these
- 69. Carbonation of Grignard reagent followed by hydrolysis gives
  - a) aldehyde
- b) acid
- c) ketone
- d) primary alcohol
- 70. Which of one the esters is formed by esterification of propan–2–ol with acetic acid?
  - a) (CH<sub>3</sub>),CHCOOCH<sub>3</sub>
  - b) CH<sub>3</sub>COOCH(CH<sub>3</sub>),
  - c) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>COOCH<sub>3</sub>
  - d) (CH<sub>3</sub>)<sub>2</sub>CHCOOCH<sub>2</sub>CH<sub>3</sub>
- 71. CH<sub>3</sub>CO CH<sub>3</sub> can be converted into CH<sub>3</sub>COOH by
  - a) reduction
- b) hydrolysis

- c) decarboxylation d) oxidation
- 72. Which reaction does not have the correct oxidising agent?
  - a)  $CH_3$ -CO  $CH_3 \xrightarrow{PCC} CH_3COOH$
  - b)  $CH_3$ -CHO  $\xrightarrow{Ag(NH_3)^+}$   $CH_3COOH$
  - c)  $CH_3$ -CHO  $\xrightarrow{Cr_2O_7^2/H^+}$   $CH_3COOH$
  - d)  $CH_3CO CH_3 \xrightarrow{Ct_2O_7^{2-}/H^+} CH_3COOH$
- 73. For the following reaction

$$R-CN \xrightarrow{H_3O^+} R-COOH$$

- a) there is protonation of electronegative nitrogen
- b) an amide is formed as an intermediate
- c) nitrogen atom is expelled as ammonia
- d) all are correct
- 74. Acetylation is the introduction of
  - a) CH,COOH group b) R-CO group
  - c) CH,CH,O-
- d) CH<sub>3</sub>-CO
- 75. 2-methyl propane nitrile on acid hydrolysis give
  - a) butyric acid
- b) isobutyric acid
- c) propionic acid
- d) pentanoic acid
- 76. In esterification reaction the correct order of reactivity of alcohol is,
  - a)  $1^{\circ} > 2^{\circ} > 3^{\circ}$
- b)  $3^{\circ} > 2^{\circ} > 1^{\circ}$
- c)  $1^{\circ} > 3^{\circ} > 2^{\circ}$
- d)  $3^{\circ} > 1^{\circ} > 2^{\circ}$
- 77. Monocarboxylic acid are regarded as oxidation product of aldehyde
  - a) first
- b) second
- c) third
- d) fourth
- 78. Formic acid is obtained when
  - a) Acetaldehyde is oxidised by acidic KMnO<sub>4</sub>
  - b) Calcium formate is heated with calcium acetate
  - c) Methanol is oxidised by PCC
  - d) Methanol is oxidised by acidic dichromate
- 79. Which of the following acid containing aldehyde group?
  - a) CH<sub>2</sub>-COOH
- b) H-COOH
- c) HOOC-COOH
- d) (CH<sub>2</sub>)<sub>2</sub>CH–COOH
- 80. Formic acid is not representative member of carboxylic acid because
  - a) it is first member of the series
  - b) it does not contain alkyl group
  - c) itisagas
  - d) it reduced Tollen's reagent
- 81. Molecular formula C<sub>4</sub>H<sub>5</sub>O on oxidation gives

acetic acid. The structure of  $C_4H_5O_2$  will be

- 82. Mixture of acetic acid and propionic acid is obtained by oxidation of
- 83. Isobutyric acid is obtained by oxidation of
  - a) butanal
  - b) 2-methyl propanal
  - c) 2-methyl propan -1-o1
  - d) both b and c
- 84. 1-phenyl ethanone on oxidation gives
  - a) salicylic acid
- b) benzoic acid
- c) succinic acid
- d) phthalic acid
- 85. Cyclohexane carboxylic acid is obtained by oxidation of
  - d) both b and c
- 86. Following compound on oxidation by acidic KMnO<sub>4</sub> gives o
  - a) Adipic acid
- b) succinic acid
- c) fumaric acid
- d) maleic acid
- 87. Benzoic acid is obtained by oxidation of
  - 1) benzaldehyde
- 2) benzyl alcohol
- 3) Acetophenone
- a) only 1
- b) 2, 3
- c) 1, 3
- d) 1,2,3
- 88. Find out A in the following reaction

$$A \xrightarrow{aq.KOH} B \xrightarrow{PCC} propanal$$

- a) n-propyl alcohol b)
- b) n-propyl halide
- c) n-propyl amine
- d) diethyl ether
- 89. Acid hydrolysis of cyanide may produces
  - a) carboxylic acid
- b) mineral acid
- c) aldehyde
- d) amine
- 90. Molecular formula  $C_4H_7$  on acid hydrolysis gives 2—methyl propanoic acid. The possible structure of  $C_4H_7$  will be
- 91. Benzoic acid is obtained by acid hydrolysis of
  - a) benzonitrile
  - b) 2-phenyl ethane nitrile
  - c) phenyl isocyanide
  - d) 2-phenyl ethane isonitrile
- 92. Find out final product in following reaction
- 93. Compound A an hydrohalogenation gives B, which

- is treated with alc.KCN and followed by acid hydrolysis gives 2-methyl butanoic acid. The compound A will be
- 94. Hexane dioic acid in obtained by oxidation of
  - a) benzene
  - b) cyclohexane
  - c) cyclohexene
  - d) cyclohexane carbaldehyde
- 95. Find out missing compound in following reaction

$$CO_2 + ? \xrightarrow{\text{dry ether} \atop H_3O^+} R - COOH$$

- a) R-X
- b) R-Li
- c) R-MgX
- d) R-ONa
- 96. Dry ice and benzyl magnesium halide produces
  - a) benzoic acid
  - b) phthalic acid
  - c) 2-phenyl ethanoic acid
  - d) 2-phenyl propanoic acid
- 97. Product of the following reaction is
- 98. 2—methyl propanoic acid is obtained from dry ice and what
- 99. Find out final product 'E' in the following reaction series

$$\xrightarrow{Zn} A \xrightarrow{CH_3-Cl} B \xrightarrow{Cl_2/U.V.L} C$$

$$\xrightarrow{Mg/Ehter} D \xrightarrow{CO_2/Ehter} B$$

- 100. Hydrolysis of acyl halide gives
  - a) aldehydes
- b) ketones
- c) carboxylic acids
- d) esters
- 101. Following compound on hydrolysis gives

$$+ H.OH \longrightarrow ?$$

- 102. Benzoic acid is obtained by hydrolysis of
- 103. 1–phenyl ethanoyl chloride on hydrolysis gives
  - a) ethanoic acid
  - b) 1-phenyl ethanoic acid
  - c) 2-phenyl ethanoic acid
  - d) benzoic acid
- 104. Hydrolysis of acid anhydride gives
  - a) aldehydes
- b) ketones
- c) acid amides
- d) carboxylic acids

- 105. Hydrolysis of ethanoic anhydride gives
  - a) methanoic acid
- b) propanoic acid
- c) ethanoic acid
- d) oxalic acid
- 106. Following compound on hydrolysis gives
- 107. Product of the following reaction is
- 108. Which of the following compound does not gives benzoic acid on oxidation
- 109. Benzoic acetic anhydride on hydrolysis gives
  - a) benzoic acid
  - b) acetic acid
  - c) mixture of benzoic acid and acetic acid
  - d) benzyl alcohol and acetic acid
- 110. Product 'B' in the following reaction

$$CH_3$$
- $CH_2$ - $COX$ 
 $\xrightarrow{Pd. BaSO_4 + quinoline} A \xrightarrow{[O]} B$ 

a) CH<sub>3</sub>-CH<sub>2</sub>-COOH b) CH<sub>3</sub>- COOH



- c)  $CH_3$ - $CH_2$ -CHO d)  $CH_3$ -C- $CH_3$
- 111. Acid hydrolysis of ester produces
  - a) carboxylic acid
  - b) alcohol
  - c) carboxylic acid and alcohol
  - d) aldehyde and ketone
- 112. Molecular formula C<sub>4</sub>H<sub>5</sub>O<sub>5</sub> on acid hydrolysis gives acetic acid and ethanol. The structures of  $C_4H_5O_7$  will be
- 113. A + H.OH  $\xrightarrow{H^+}$

The compound A is

- 114. Molecular formula C<sub>3</sub>H<sub>6</sub>O<sub>2</sub> on acid hydrolysis gives A and B. The compound 'A' reduce Tollens -reagent. The possible structure of C<sub>3</sub>H<sub>6</sub>O<sub>2</sub> will
  - a) CH<sub>3</sub>-CH<sub>2</sub>-COOH b) CH<sub>3</sub>-COOCH<sub>3</sub>

c) H-COOC,H,

- d) CH<sub>3</sub>- C CH<sub>3</sub>
- 115. Which of the following method cannot be used in the preparation of acid

a) 
$$R - COCl + H_2O \longrightarrow$$

b)

d) 
$$R - C = N \xrightarrow{Na + ethanol}$$

- 116. Vigorous oxidation by acidic permagnet solution of  $(CH_3)_{,C} = CH - CH_{,CH_3}$  gives
- 117. A set of reaction yielded a product D.

$$\begin{array}{cccc} C_2H_5COOH & \xrightarrow{SOCl_2} & A & \xrightarrow{C_6H_6} & B \\ & \xrightarrow{HCN} & C & \xrightarrow{H_3O^+} & D \end{array}$$

118. Product of the following reaction

$$\xrightarrow{\text{HBr}} A \xrightarrow{\text{Mg/ether}} B \xrightarrow{\text{CO}_2} C$$

- 119. Alkyl benzene on oxidation by alkaline KMnO<sub>4</sub> gives
  - a) aldehyde
- b) ketone
- c) carboxylic acid
  - d) esters
- 120. Product of the reaction

$$\xrightarrow{\text{KOH} + \text{KMnO}_4}$$

- 121. o-xylene on oxidation gives
  - a) phthalic acid
- b) isophthalic acid
- c) terephthalic acid
  - d) citric acid
- 122. Following compound is obtained by oxidation is
  - a) 1, 3
- c) 2, 4
- b) 2, 3 d) 1, 2, 3, 4
- 123. Phenyl ethene on oxidation produces
  - a) phthalic acid
  - b) p-ethenyl benzoic acid
  - c) benzoic acid
  - d) salicylic acid
- 124. Find out final product of the reaction

+ CH<sub>3</sub>COCl 
$$\xrightarrow{\text{unhy.AlCl}_3}$$
 A  $\xrightarrow{\text{KOH + KMnO}_4}$  B

- 125. Potassium salt of isophthalic acid on acid hydrolysis gives
  - a) phthalic acid
- b) isophthalic acid
- c) terephthalic acid
- d) benzoic acid
- 126. Carbonation of organometallic reagent
  - a) is a one carbon homologation
  - b) involves its carbanionic attacks on electron deficient carbonyl carbon
  - c) both are correct

- d) none is correct
- 127. Hydrolysis of nitriles with alkaline solution gives
  - a) salt of acid
- b) amides
- c) acid
- d) esters
- 128. The mechanism of carbonation of organometallic compound is analogous to that of addition to
  - a) aldehyde and ketones
  - b) nitriles
  - c) alcohols
  - d) amines
- 129. Dry ice reacts with n-propyl magnesium iodide to give
  - a) propyl acetic acid b) propionic acid
  - c) butanoic acid
- d) none of these
- 130. Esterification reaction is
  - a) nucleophilic substitution
  - b) electrophilic substitution
  - c) electrophilic addition
  - d) dehydration
- 131. In which case R-COOH is not product?
  - a) R– $CN + RMgX \rightarrow$
  - b) R-CN  $\xrightarrow{\text{H}_3\text{O}^+}$
  - c) R-MgX  $\xrightarrow{CO_2}$
  - d) RCOR \_\_\_oxidation\_\_\_
- 132. In esterification reaction concentrated H<sub>2</sub>SO<sub>4</sub> act as
  - a) dehydrating agent b) catalyst
  - c) reducing agent
- d) oxidising agent
- 133. After completion of esterification, excess of alcohol is removed by using
  - a) CaCl,
- b) CaOCl,
- c) Na<sub>2</sub>CO<sub>2</sub>
- d) NaOH
- 134. Synthesis of ester involves the reaction of alcohol with
  - a) a ketone
- b) an amide
- c) CH<sub>2</sub>MgBr
- d) RCOOH
- 135. Which of the following reagent produces 2—methyl propanoic acid with dry ice?
  - a) CH<sub>3</sub>MgI
- b) C<sub>2</sub>H<sub>5</sub>MgI
- c) (CH<sub>3</sub>)<sub>2</sub>CHMgI
- d) C<sub>3</sub>H<sub>7</sub>MgI
- 136. 1-propanol on oxidation give
  - a) propanal
- b) propionic acid
- c) propanone
- d) ethanal
- 137. -CN group is converted into -COOH group by,

- a) hydrolysis b) oxidation
- c) reduction
- d) esterification
- 138. Esters are formed from acid by the replacement of
  - a) non-ionisable H atom by alkyl group
  - b) ionisable H atom by alkyl group
  - c) OH group by RO group
  - d) both 'b' and 'c'
- 139. Which one of the following correctly represents esterification reaction
  - a) RCO  $\overline{OH+H}$  O-R'  $\rightarrow$  R-COOR' + H<sub>2</sub>O
  - b) RCOO $\overline{\text{H} + \text{HO}} \text{R'} \rightarrow \text{R-COOR'} + \text{H}_2\text{O}$
  - c) RCO  $\overline{\text{OH+H}}$  O-R'  $\rightarrow$  R'-COOR + H<sub>2</sub>O
  - d) RCOO $\overline{\text{H} + \text{HO}} \text{R'} \rightarrow \text{R'-COOR} + \text{H}_2\text{O}$
- 140. The compound formed when propyl magnesium bromide is treated with carbon dioxide is
  - a) C<sub>2</sub>H<sub>7</sub>COOH
- b) C,H,COOH
- c) C<sub>3</sub>H<sub>7</sub>CHO
- d) C<sub>2</sub>H<sub>7</sub>OCH<sub>2</sub>
- 141. The alkyl cyanides when hydrolysed to the corresponding acid, the gas evolved is
  - a)  $N_2$
- b) O,
- c) NH<sub>3</sub>
- d) CO<sub>2</sub>
- 142. In esterification of an acid, the other reagent is
  - a) Alcohol
- b) An aldehyde
- c) Amine
- d) Water
- 143. When methyl cyanide is hydrolysed in presence of alkali and followed by treatment with HCl. The product is
  - a) Acetamide
- b) Methane
- c) Formic acid
- d) Acetic acid
- 144. Which product is formed, when acetonitrile is hydrolysed partially with dil. HCl?
  - a) Acetic acid
- b) Acetamide
- c) Methyl cyanide
- d) Acetic anhydrides
- 145. Carboxylic acid is formed by the hydrolysis of an addition compound formed by reaction of Grignard's reagent with
  - a) ethanal
- b) methanal
- c) carbon dioxide
- d) methyl cyanide
- 146. Which is formed, when benzonitrile is hydrolysed by dil. HCl?
  - a) Benzoic acid
- b) Benzaldehyde
- c) Benzamide
- d) Benzoic unhydride

ACIDIC NATURE

- 147. The acidic nature of carboxylic acid is due to
  - a) high degree of ionisation of acid
  - b) greater resonance stabilization of the acid
  - c) greater resonance stabilization of its anion
  - d) all of the above
- 148. Acidic character of carboxylic acid with an molecular mass increases
  - a) sometime increases some time decreases
  - b) decreases
  - c) increases
  - d) none of the above
- 149. Among the acid
  - 1) HC = C-COOH
  - 2)  $H_2C = CH COOH$
  - 3) CH<sub>3</sub>-CH<sub>2</sub>-COOH

The acidic strength follows the order

- a) 3 < 2 < 1
- b) 3 = 2 < 1
- c) 1 < 2 < 3
- d) 1 < 2 = 3
- 150. Which of the following it the most stable acid?
  - a) CH<sub>2</sub>(F)COOH
- b) CH<sub>2</sub>(Br)COOH
- c) CH<sub>2</sub>(Cl)COOH
- d) CH<sub>2</sub>(I)COOH
- 151. Which of the following acid is strongest
  - a) CH<sub>3</sub>-COOH
- b) Cl<sub>2</sub>-CH<sub>2</sub>-COOH
- c) Cl<sub>2</sub>CH-COOH
- d) CCl<sub>3</sub>-COOH
- 152. Which of the following is weak acid?
- 153. Which of the following is strongest acid?
  - a) CH,CH,CH,COOH
  - b) CH<sub>3</sub>CH<sub>2</sub>CHBrCOOH
  - c) CH<sub>3</sub>CHBrCH<sub>2</sub>COOH
  - d) CH<sub>2</sub>BrCH<sub>2</sub>CH<sub>2</sub>COOH
- 154. Which of the following has highest Pka value?
  - a) C<sub>6</sub>H<sub>5</sub>-COOH
- b) CH<sub>2</sub>-OH
- c) C<sub>6</sub>H<sub>5</sub>- OH
- d) CH<sub>3</sub> COOH
- 155. Which of the following is strong acid?
  - a) CF<sub>3</sub>COOH
- b) CCl<sub>3</sub>COOH
- c) CBr<sub>3</sub>COOH
- d) Cl<sub>2</sub>COOH
- 156. Which of the following is strongest acid?
  - a) H–COOH
- b) C<sub>6</sub>H<sub>5</sub>-COOH
- c) CH<sub>3</sub> COOH
- d) CH<sub>2</sub>-CH<sub>2</sub>-COOH
- 157. Weakest acid among the following is
  - a) H-COOH
  - b) C<sub>6</sub>H<sub>5</sub>- COOH
  - c)  $o-NO_2-C_6H_4$  COOH
  - d) CH<sub>3</sub>-COOH
- 158. Benzoic acid is more stronger than all aliphatic

- carboxylic acid except H COOH, why?
- a) C<sub>6</sub>H<sub>5</sub>- COOH is aromatic acid
- b) In  $C_6H_5$  COOH, the COOH group is attached to sp<sup>2</sup>–hydridised carbon atom
- c) resonance stabilization of benzene ring
- d) resonance stabilization of benzoate ion.
- 159. The aqueous solution of RCOOH contains,
  - a)  $RCOO^- + H^+$
  - b)  $RCO^+ + OH^-$
  - c)  $RCOO^- + H_3O^+$
  - d) RCOOH + RCOO $^-$  +  $H_3O^+$
- 160. Which one of the following would be expected to be most highly ionised in water
  - a) Cl-CH,-CH,-CH,-COOH
  - b) CH, CHClCH, -COOH
  - c) CH<sub>3</sub>-CH<sub>2</sub>-CHCl-COOH
  - d) CH<sub>3</sub>-CH<sub>2</sub>-CH<sub>2</sub>-COOH
- 161. Between CH<sub>3</sub>COOH and H-COOH, the HCOOH will be
  - a) less acidic
- b) more acidic
- c) equally acidic
- d) non-acidic
- 162. Formic acid and acetic acid may be distinguished by the reaction with
  - a) sodium metal
- b) Tollen's reagent
- c) sodium ethoxide
- d) NaOH
- 163. Which is tribasic acid
  - a) adipic acid
- b) succinic acid
- c) Tartaric acid
- d) citric acid
- 164. Consider the following acids
  - 1) CH<sub>3</sub>COOH
- 2) Cl,CHCOOH
- 3) ClCH,COOH
- 4) Cl<sub>3</sub>CCOOH
- The correct sequence of acidity is
- a) 1 > 4 > 3 > 2 c) 1 > 2 > 3 > 4
- b) 1 > 3 > 2 > 4 d) 4 > 2 > 3 > 1
- 165. Which of the following order of relative strength of acid is correct?
  - a) F-CH<sub>2</sub>-COOH > Cl CH<sub>2</sub> COOH >

$$Br-CH_{2} - COOH > CH_{3} COOH$$

b)  $CH_3 COOH > F - CH_2 - COOH >$ 

 $C1 CH_{2} - COOH > CH_{2} Br - CH_{2}COOH$ 

c) Br CH, COOH > Cl-CH, COOH >

 $F-CH_2$  -COOH >  $CH_3$ COOH

d) ClCH,COOH > F CH<sub>2</sub>-COOH >

Br CH<sub>2</sub>-COOH > CH<sub>2</sub>COOH

166. Which of the following is weakest acid

- 167. Which of the following acid has smallest dissociation constant?
  - a) CH<sub>2</sub>-CHFCOOH
  - b) BrCH,CH,-COOH
  - c) F-CH,-CH,-COOH
  - d) CH<sub>3</sub> CHBr COOH
- 168. Which among the following is strongest acid?
- 169. Among the acid which have lowest PKa value.
  - a) CH, COOH
  - b) CH<sub>3</sub>-CH<sub>2</sub>-COOH
  - c) CH<sub>3</sub> CH COOH NO,
  - d) CH<sub>3</sub> CH COOH COOH
- 170. The correct order of increasing strength of compounds
  - 1) CH<sub>2</sub> COOH
- 2) NO, CH, COOH
- 3) CF<sub>3</sub>COOH
- 4) C<sub>2</sub>H<sub>5</sub>-COOH
- a) 4 > 3 > 2 > 1
- b) 3 > 2 > 1 > 4
- c) 3 > 1 > 2 > 4
- d) 3 > 4 > 2 > 1
- 171. Among the following acidic strength is
  - 1) CH<sub>3</sub>-CH<sub>2</sub>-COOH
  - 2) CH<sub>3</sub>-CH-COOH
  - 3)  $CH_3 CH COOH$
  - 4) CH<sub>3</sub> CH COOH  $NO_{2}$
  - a) 4 > 1 > 2 > 3
- b) 3 > 4 > 2 > 1
- c) 4 > 2 > 3 > 1
- d) 4 > 3 > 2 > 1

### PROPERTIES AND USES

- 172.  $CH_3COOH + CH_3 \cdot H \xrightarrow{H^+} (ester) + B$ (water) Alcohol contain isotopic oxygen {•}. The isotopic oxygen will be in
  - a) CH<sub>3</sub>CO·CH<sub>3</sub> + H<sub>2</sub>O
  - b) CH<sub>3</sub>COOCH<sub>3</sub> + H<sub>3</sub>•
  - c) both are correct
  - d) none is correct

- 173. Select incorrect statement
  - a) NaOH is strong enough to remove the proton from phenols and carboxylic acids
  - b) NaHCO<sub>3</sub> is only strong enough to remove the proton of carboxylic acids
  - c) Mixture of phenol and carboxylic acid can be separated using NaHCO,
  - d) Mixture of phenol and carboxylic acid can be separated using NaOH
- 174. What is the main reason for the fact that carboxylic acid can undergo ionization?
  - a) Absence of alpha–hydrogen
  - b) Resonance stabilization of the carboxylate ion
  - c) High reactivity of alpha-hydrogen
  - d) Less reactivity of a-H atom
- 175. Tollen's reagent is reduced by
  - a) HCOOH
- b) CH<sub>3</sub>COOH
- c) CH<sub>2</sub>ClCOOH
- d) CCl,COOH
- 176. The end product of following sequence of reaction is

$$\text{CH}_3\text{--CHO} \xrightarrow{ \ [\text{O}\ ] \ } \text{A} \xrightarrow{ \ \text{NH}_3 \ } \text{B} \xrightarrow{ \ \Delta \ } \text{C}$$

- a) CH<sub>3</sub>COONH<sub>4</sub> b) CH<sub>3</sub>CONH<sub>2</sub> c) CH<sub>3</sub>COOH d) CH<sub>3</sub>-OH + N d)  $CH_3$ -OH +  $N_2$
- 177. Toluene  $\xrightarrow{KOH + KMnO_4} A \xrightarrow{PCl_5} B$

Product 'C' is

- a) benzoic ethanoic unhydride
- b) benzophenone
- c) acetophenone
- d) benzoquinone
- 178. Propionic acid on bromination in the presence of red phosphorus gives 2-bromo propionic acids it shows
  - a) optical isomers
- b) tautomerism
- c) cis-trans isomers d) metamerism
- 179. Product 'B' following reaction is OCN
- 180. Which has maximum value of equilibrium constant?
  - a)  $CF_3COOH + H_2O \rightleftharpoons CF_3COO^- + H_3O^+$
  - b) HCOOH + H<sub>2</sub>O  $\rightleftharpoons$  HCOO- + H<sub>3</sub>O+
  - c) CH<sub>3</sub>COOH + H<sub>2</sub>O CH<sub>3</sub>-COO<sup>-</sup> + H<sub>3</sub>O<sup>+</sup>
  - d)  $C_6H_5COOH + H_2O \rightleftharpoons C_6H_5-COO^- + H_3O^+$

- 181. When propionic acid is heated with ammonia give
  - a) CH<sub>3</sub>CONH<sub>2</sub>
- b) C<sub>2</sub>H<sub>5</sub>COONH<sub>4</sub>
- c) C<sub>2</sub>H<sub>5</sub>CONH<sub>2</sub>
- d) CH<sub>3</sub>COONH<sub>4</sub>
- 182. Product 'A' and 'B' in the following reaction is

$$CH_3CH = CH_2 \xrightarrow{K_2Cr_2O_7 + dil. H_2SO_4} A + B$$

- a) acetone
- b) acetaldehyde and formaldehyde
- c) propanoic acid
- d) acetic acid and formic acid
- 183. Sodium acetate reacts with acetyl chloride to give
  - a) acetic acid
- b) acetic anhydride
- c) methyl acetate
- d) ethyl acetate
- 184. Acetic acid react with PCl<sub>5</sub> gives,
  - a) CH<sub>2</sub>COCl
- b) ClCH,CH,Cl
- c) CH<sub>3</sub>COOCl
- d) CH<sub>2</sub>Cl
- 185. Which of the following compound reacts with NaHCO<sub>3</sub> solution to give sodium salt and carbon dioxide?
  - a) Phenol
- b) 1-propanol
- c) Acetic acid
- d) Nitric acid
- 186. An optically active compound 'X' evolve CO<sub>2</sub> with NaHCO<sub>3</sub>. The 'X' react with LiAlH<sub>4</sub> gives chiral compound. The compound 'X' is
  - a) CH<sub>3</sub>-CH<sub>2</sub>-CH<sub>2</sub>-COOH
  - b)  $CH_3 CH CH_2 CH_3$ COOH
  - c) CH<sub>3</sub> CH<sub>2</sub> CH CH<sub>2</sub> CH<sub>3</sub>
  - d)  $CH_3 CH CH_2 CH_3$ COOH
- 187. An propanoyl chloride is formed when PCl<sub>5</sub> reacts with an
  - a) propanoic acid
- b) alcohol
- c) acetic acid
- d) ester
- 188. Which of the following substance will give amide, when it reacts with NH<sub>2</sub>?
  - a) CH,X
- b) CH<sub>3</sub>NH<sub>3</sub>
- c) CH,COCl
- d) (CH<sub>3</sub>CO)<sub>2</sub>O
- 189. Formic acid is a stronger acid than acetic acid. This is due to the fact that
  - a) formic acid is reducing agent

- b) formic acid molecule is of smaller size
- c) there is no alkyl group on  $\alpha$  –carbon in formic acid
- d) formic acid does not undergo association
- 190. What are the product in the following reaction?

$$\begin{array}{c} \text{CH}_{3} \\ \mid \\ \text{CH}_{3} - \text{C} = \text{CH}_{2} \end{array} \xrightarrow{\begin{array}{c} \text{K}_{2}\text{Cro}_{7}^{+} \\ \text{dil. H}_{2}\text{SO}_{4} \end{array}} \rightarrow \end{array}$$

- a) CH<sub>3</sub>-COCH<sub>3</sub> and H-COOH
- b) CH<sub>3</sub>-CH<sub>2</sub>CHO and CO,
- c) CH<sub>2</sub>-COOH and H-COOH
- d) CH<sub>2</sub>-CHO and H-COOH
- 191. Product of the following reaction is

$$CH_3COCH_3 \xrightarrow{[O]} A \xrightarrow{PCl_3} B$$



- a) acetic unhydride b) propanoic unhydride
- c) ethyl propanoate
- d) isopropyl alcohol
- 192. Which of the following does not under go HVZ reaction?
  - a) CH<sub>3</sub>-COOH
  - b) CH<sub>3</sub>-CH<sub>2</sub>-COOH

d) (CH<sub>2</sub>), C-COOH

- 193. During formation of unhydride from P<sub>2</sub>O<sub>5</sub> and acid. The acid molecule under goes
  - a) intermolecular dehydration
  - b) interamolecular dehydration
  - c) oxidation
  - d) hydration
- 194. Acetic anhydride is obtained by the reaction of
  - a) sodium and acetic acid
  - b) ammonia and acetic acid
  - c) ethanol and acetic acid
  - d) P<sub>2</sub>O<sub>5</sub> and acetic acid.
- 195. An ester is subjected to hydrolyse. Product of hydrolysis will be tested for
  - a) carboxylic acid and alcoholic group
  - b) carboxylic acid and ketonic group
  - c) carboxylic acid and aldehyde group
  - d) aldehyde and ketonic group
- 196. Carboxylic acid reacts with potassium metal. The amount of potassium used up and hydrogen liberated are in the molar ratio of

- a) 2:1
- b) 1:1
- c) 3:2
- d) 2:3
- 197. HVZ reaction is characteristic reaction of
  - a)  $\alpha$  –H atom
- b)  $\beta$ -H atom
- c)  $\gamma$  –H atom
- d)  $\delta$ -H atom
- 198. Carboxylic acid reacts with calcium metal. The amount of calcium used up and hydrogen liberated are in the molar ratio of
  - a) 2:1
- b) 1:1
- c) 3:2
- d) 2:3
- 199. Two molecules of acetic acid are heated with P<sub>2</sub>O<sub>5</sub>. The product formed is
  - a) two moles of ethanol
  - b) two moles of methyl cyanide
  - c) acetic anhydride
  - d) formic acid
- 200. Carboxylic acid can
  - a) decompose carbonate and evolve CO,
  - b) reacts with metal forming H, gas
  - c) neutralise ammonium hydroxide form salt
  - d) all of the above
- 201. Which of the following group will increase more acidity of acetic acid?
  - a)  $-NO_2$
- b) –CH,
- c) CH,O-
- d) -NH<sub>2</sub>
- 202. P<sub>2</sub>O<sub>5</sub> is anhydride of
  - a) H<sub>3</sub>PO<sub>3</sub>
- b) HPO,
- c) H<sub>3</sub>PO<sub>4</sub>
- d) HClO
- 203. The solvent that can dissolve all the carboxylic acids is
  - a) water
- b) dil. HCl
- c) dil. NaOH
- d) conc. H,SO<sub>4</sub>
- 204. Acetyl chloride can be prepared
  - a) by the action of CH<sub>2</sub>COOH with chloroform
  - b) by the action of PCl, on acetic acid
  - c) by the action of Cl, on acetic acid
  - d) by the action of CCl<sub>4</sub> on acetic acid
- 205. Of the following four reactions, formic and acetic acid differ in which respect?
  - a) Formation of ester with alcohol
  - b) Replacement of hydrogen by sodium
  - c) Reduction of Fehling solution
  - d) Blue litmus reaction
- 206. Arrange the following carboxylic acid in their decreasing acidity

- COOH
- ĊООН
- Oxalic acid
- 2) HOOC-CH<sub>2</sub>-COOH Malonic acid

$$CH_2 - COOH$$

- Succinic acid
- a) 3 > 2 > 1
- b) 1 > 2 > 3
- c) 2 > 3 > 1
- d) 2 > 1 > 3
- 207. Among the following which compound will react with Na<sub>2</sub>CO<sub>3</sub> solution to give sodium salt and CO,?
  - a) Phenol
- b) 1-hexanol
- c) Formic acid
- d) Ethanal
- 208. Identify the correct order of boiling point of following compound?
  - a) 1 > 3 > 2 > 1
- b) 2 > 3 > 1 > 4
- c) 3 > 1 > 2 > 4
- d) 3 > 1 > 4 > 2
- 209. The compounds formed by the acid hydrolysis of ethyl acetate are
  - a) formic acid and propanol
  - b) acetic acid and ethanol
  - c) acetone and ethanol
  - d) acetone and methanol
- 210. -COOH group of a compound does not react with NaHSO<sub>3</sub> even though it has >C = O group because of
  - a) cyclic structure
- b) acidic character
- c) linear structure
- d) resonance
- 211. Final product of following reaction is

$$CO_2 + (CH_3)_3C-MgBr \xrightarrow{i) dry ether} A \xrightarrow{Br_2} B$$

a) 
$$CH_3$$
  $CH_3$   $CH_3$ 

b) 
$$CH_3 - C - CH - COOH$$

$$CH_3 - Br$$

c) 
$$CH_3 - CH_2 - CH - COOH$$
|
Br

- d) No product will be formed.
- 212. Acids having higher B.P. than alcohols because of
  - a) intra molecular HB
  - b) dimerisation of acid
  - c) dipole-dipole attraction
  - d) Vander Waals force of attraction
- 213. Product of the following reaction is

$$CH_3$$
- $Cl + C_2H_5$ - $COOAg \longrightarrow A \xrightarrow{H_3O^+} B$ 

- a)  $CH_3$ -COOH +  $C_2H_5$ -OH
- b) CH<sub>3</sub>-CHO + C<sub>2</sub>H<sub>5</sub>-OH
- c)  $C_2H_5$ -COOH +  $CH_3$ -OH
- d)  $C_2H_5$ -COOH +  $C_2H_5$ -OH
- 214. Adipic acid is formed by oxidation of
  - a) cyclohexane
- b) cyclohexene
- c) hex-1-ene
- d) hex-2-ene
- 215. Product 'B' of the following

$$CH_3$$
- $CH_2$ - $COOH +  $Cl_2$   $\xrightarrow{Red P}$   $A$$ 

 $\xrightarrow{\text{KOH alc.}} B$ 

- a) CH<sub>3</sub>-CHCl-COOH
- b) CH<sub>3</sub>-CHOH-COOH
- c) CH<sub>2</sub> = CH-COOH
- d) CH = C-COOH
- 216. Which of the following does not give HVZ reaction?
  - a) Acetic acid
- b) Propionic acid
- c) Isobutyric acid
- d) Neovaleric acid
- 217. Product 'B' of the following

$$\mathrm{CH_3-CH_2-CH_2-COOH}\,+\,\mathrm{Br_2}\,\stackrel{\mathrm{Red}\,\mathrm{P}}{--\!\!\!\!-}\,\mathrm{A}$$

 $\xrightarrow{\text{Aq.KOH}} B$ 

- a) CH<sub>3</sub>-CH<sub>2</sub>-CHOH-COOH
- b) CH<sub>3</sub>-CHOH-CH<sub>3</sub>-COOH
- c) HO-CH,-CH,-CH,-COOH
- d)  $CH_3$ -CH = CH-COOH
- 218. Terephthalic acid is heated with ammonia gives
  - a) phthalamide
- b) phthalimide
- c) isophthalic
- d) ammonium phthalate

OOO







### HINI & SULUTIONS - MCQ 5

### **SECTION - I: ALDEHYDES AND KETONES**

**39. b)** Ketone also shows tautomerism by transfer of hydrogen.

$$CH_3 - CO - CH_2H \longrightarrow CH_3 - C = CH_2$$

ketoform (stable) enolform (less stable)

- 58. d) We know that  $(HCOO)_2Ca \xrightarrow{\Delta} HCHO + CaCO_3.$  Thus in this reaction, formaldehyde (HCHO) is produced.
- 59. **b)** RCN + RMgX  $\xrightarrow{\text{dryether}}$

- 60. a)  $CH_3CCl_2CH_3 + 2KOH \xrightarrow{aq.} CH_3COCH_3 + 2KCl$
- 61. **b)**  $CH_3CCl_2CH_2CH_3 + 2KOH \xrightarrow{aq.} CH_3COCH_2CH_3 + 2KCI$
- 62. a) CH<sub>3</sub>CHOHCH<sub>3</sub> oxidation CH<sub>3</sub>COCH<sub>3</sub>+H<sub>2</sub>O
- 69. c) Molecular formula C<sub>4</sub>H<sub>10</sub>O has four isomers i.e. n-butyl alcohol, isobutyl alcohol, sec. butyl alcohol, t-butyl alcohol.Sec. butyl alcohol is optically active and on oxidation gives ethyl methyl ketone.

$$CH_3CHOHC_2H_5$$
  $\xrightarrow{\text{oxidation}}$   $CH_3COC_2H_5 + H_2O$ 

71. c) HCN + (CH<sub>3</sub>)<sub>2</sub>CHMgX 
$$\xrightarrow{\text{dry ether}}$$

$$(CH_3)_2CH$$
-CHO + MgXOH

- 119. d) The reagents a, b, c reacts with the carbonyl compounds in a same manner and only NH<sub>3</sub> react differently. Formaldehyde is reacted with ammonia give urotropine. Acetaldehyde reacts ammonia give acetaldehyde amine and acetone reacts ammonia give diacetone amine.
- 120. a) Positive charge on carbonyl carbon is maximum in case of HCHO. It is because no alkyl group (+I effect) is attached with it and therefore, positive charge is not dispersed. Hence formaldehyde is more reactive than other aldehydes.

- 121. a) Aldehydes are more reactive than ketones towards nucleophilic addition reactions, due to the electron repelling inductive effect (+ I) of alkyl groups. The alkyl groups at the carbonyl carbon in ketones reduces the positive charge over it and therefore, the tendency of a nucleophilic attack is reduced. The bulky alkyl groups also decrease the reactivity of >C=O group, due to steric hindrance.
- 122. b) Aldehydes and ketones both have carbonyl group >C=O, which reacts with Grignard's reagent. Fehling's solution gives red precipitate or coloration with aldehydes only, except aromatic aldehydes. Tollen's reagent gives silver mirror only with aldehydes. Only aldehydes turn Schiff's reagent pink.
- 123. b) Aldehydes lacking an  $\alpha$  hydrogen undergoes self oxidation reduction (disproportionation reaction), also known as Cannizzaro's reaction.
- 124. d) Cannizzaro's reaction is given by those aldehydes which have no α hydrogen atom. Compound (a) has no α hydrogen atoms, so it shows Cannizzaro's reaction. Compound (b) has no α-hydrogen atom but does not show Cannizzaro's reaction, because of –I effect.
- 125. c) This is aldol condensation reaction CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CHO + CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CHO

$$\xrightarrow{\text{dil. NaOH}}$$
 CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CHOHCH(C<sub>2</sub>H<sub>5</sub>)CHO  
2-ethyl 3-hydroxy hexanal

- 126. c) 2HCHO + KOH → CH<sub>3</sub>OH + HCOOK
- 127. c) The nucleophile is  $SO_3^{--}$  not  $HSO_3^{-}$ ,  $SO_3Na$
- **128.** b) The nucleophile is  $CN^-$  attack on  $> C^{\delta+}$

129. b) 
$$CH_3COCH_3 + CH_3MgI \xrightarrow{dryether} H_3O^+$$

(CH<sub>3</sub>)<sub>3</sub>COH + MgIOH

130. d) Aldol condensation reaction shown by those aldehydes and ketones containing α hydrogen atoms or aldehydes and ketones having methyl or methylene group in α position. Compounds a, b, c have α-hydrogen atoms.

2-methyl 3-hydroxy pentanal

132. b) When aldehydes condenses with ketone, produces four condenstion products. It is the α-hydrogen atom of the ketone which is involved in the condensation reaction and produces major product.

$$CH_3CHO + HCH_2COCH_3 \xrightarrow{dil. NaOH}$$
 $CH_3CH(OH)CH_2COCH_3$ 
 $4$ -hydroxy 2-pentanone

- 133. d) Iodoform test is given by organic compounds which contains CH<sub>3</sub>CO group or CH<sub>3</sub>CHOH grouping. 3-pentanone does not contains any one of such grouping.
- 134. c) Methyl ketones react with iodine and aqueous alkali to form yellow compound, iodoform (CHI<sub>3</sub>)
- **135. d)** Compounds a, b, c undergoes nucleophilic addition reactions. Hence, RMgX adds in above compounds.
- **136. c)** Ketones do not respond to Tollen's test. Aldehydes respond to Tollen's test.
- 137. d) Fehling solution is weak oxidising agent and benzaldehyde is weak reducing agent. Hence, it does not react with Fehling solution. Formic acid, glucose, ethanal contains CHO-group. Hence, they reduce Fehling solution.

138. c) RCOR + RMgX 
$$\frac{\text{dry ether}}{\text{H}_3\text{O}^+}$$
 R<sub>3</sub>C-OH + MgXOH

139. a) 
$$(CH_3)_2C = O + H_2CHCOCH_3 \xrightarrow{NH_3} (CH_3)_2C = CHCOCH_3 + H_2O$$

140. a) >C=O + 4H 
$$\xrightarrow{\text{ZnHg} + \text{HCl}}$$
 >CH<sub>2</sub> + H<sub>2</sub>O

- 141. a) Self redox reaction is Cannizzaros not given by those aldehydes which contain α-hydrogen atoms. Methanal does not have α-hydrogen atoms, hence, it give self redox reaction.
- 142. b) 6HCHO +  $4NH_3 \rightarrow (CH_2)_6N_4 + 6H_2O$
- 143. a) When aldehydes condenses with ketone, gives four condensation products. It is the α-hydrogen atom of the ketone which is involved in the condensation reaction and produces major product.

CH<sub>3</sub>CH<sub>2</sub>CH(OH)CH<sub>2</sub>COCH<sub>3</sub> 4 -hydroxy 2-hexanone

144. c) This is aldol condensation reaction,

$$CH_3COCH_3 + CH_3COCH_3 \xrightarrow{Ba(OH)_2}$$
  
 $CH_3C(OH)(CH_3)CH_2COCH_3$ 

144. c) This is aldol condensation reaction,

$$CH_3COCH_3 + CH_3COCH_3 \xrightarrow{Ba(OH)_2}$$
 $CH_3C(OH)(CH_3)CH_2COCH_3$ 

145. c) Structure of urotropine is,

146. d) Acetone and acetaldehyde, both contains >C=O group. The Grignards adds in >C=O group. These are nucleophilic substitution reaction.

147. c) 
$$CH_3$$
-CHO + HCN  $\longrightarrow$   $CH_3$  -  $C$  - H

$$\frac{\text{hydrolysis}}{\text{CH}_3 - \text{C} - \text{H}}$$
COOH

lactic acid (optically active acid)

- 148. c) We know that, 2CuO + CH<sub>3</sub>CHO → Cu<sub>2</sub>O↓ + CH<sub>3</sub>COOH. Thus when ethanal (CH<sub>3</sub>CHO) is heated with Fehling's solution, it gives a red precipitate of cuprous oxide (Cu<sub>2</sub>O).
- **169. d)** We know that in aldol condensation,  $\alpha$ -hydrogen atoms undergo self-addition between two molecules to form  $\beta$ -hydroxy ketone. Therefore ketone in which  $\alpha$ -hydrogen atom is present, give this reaction.
- **170. d)** We know that,

2HCHO + KOH  $\stackrel{\Delta}{\longrightarrow}$  HCOOK + CH<sub>3</sub>OH. Thus in this reaction, methyl alcohol (CH<sub>3</sub>OH) is produced. This reaction is known as Cannizaro's reaction.

- 171. d) We know that benzaldehyde which does not contain α-hydrogen atom, undergoes self oxidation-reduction when reacted with 50% aqueous alkali. In the process of converting benzaldehyde to benzly alcohol, one molecule is oxidesed to sodium benzoate at the expense of other which is reduced to benzyl alcohol. This reaction is know as Cannizzaro's reaction.
- 172. d) We know that aldehyde can be oxidised with potassium dicromate (K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>) in acidic medium to form carboxylic acid, when Tollens reagent is used to oxidise an aldehyde, the silver ion is reduced to metallic from if the reaction is carried out in a clean test tube. We also know that when Fehling's solution is used to oxidise an aldehyde, the complexe cupric ion is reduced to cuprous oxide. Therefore all of these options are correct.
- 173. d) We know that,
   6HCH = O + 4NH<sub>3</sub> → (CH<sub>2</sub>)<sub>6</sub>N<sub>4</sub> + 6H<sub>2</sub>O.
   Formaldehyde Hexamethylene tetramine
   Thus in this reaction, the product Y is hexamethylene tetramine [(CH<sub>2</sub>)<sub>6</sub>N<sub>4</sub>]
- 174. a) We know that, aldehyde restores the colour of Schiff's reagent in the cold, while ketones do not give this test.

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### **SECTION - II: CARBOXYLIC ACIDS**

- 18. b) C<sub>2</sub>H<sub>4</sub>O<sub>2</sub> have two isomers,i) CH<sub>3</sub>COOH, ii) HCOOCH<sub>3</sub>
- 52. d)  $CH_3CH_2CH_2Br \xrightarrow{Mg} CH_3CH_2CH_2MgBr$

$$\frac{\text{CO}_2/\text{H}_3\text{O}^+}{\text{dry ether}} \rightarrow \text{CH}_3\text{CH}_2\text{CH}_2\text{COOH}$$

- 53. a)  $CO_2 + RMgX \xrightarrow{dry \text{ ether}} RCOOH + MgXOH$
- 54. a) Carbonation of CH<sub>3</sub>MgI forms CH<sub>3</sub>COOH. Hydrolysis of acetonitrile by a mineral acid also forms CH<sub>3</sub>COOH.
- 55. a)  $CH_3CH_2CH_2OH + CH_3CH_2COOH \xrightarrow{conc.H_2SO_4} CH_3CH_2COOCH_2CH_2CH_3$  propyl propionate
- 56. c)  $(CH_3CO)_2O + CH_3OH \xrightarrow{\Delta}$   $CH_3COOH + CH_3COOCH_3$ mol. mass 74  $(HCO)_2O+C_2H_5OH \xrightarrow{\Delta} HCOOH + HCOOC_2H_5$ mol. mass 74
- 60. c)  $CH_3COOH + (CH_3)_2CHOH \xrightarrow{conc. H_2SO_4} CH_3COOCH(CH_3)_2 + H_2O$
- 61. a)  $C_2H_5COOH+CH_3OH \xrightarrow{pyridine} C_2H_5COOCH_3 + H_2O$
- 62. **b)**  $C_2H_5COOAg + C_2H_5I \xrightarrow{\Delta} AgI + C_2H_5COOC_2H_5$  ethyl propionate
- 64. b) We know that,
   CH<sub>3</sub>COOH + C<sub>2</sub>H<sub>5</sub>OH CH<sub>3</sub>COOC<sub>2</sub>H<sub>5</sub> + H<sub>2</sub>O.
   Thus in this reaction, an ester (CH<sub>3</sub>COOHC<sub>2</sub>H<sub>5</sub>) is formed. This process is called esterfication.
- 65. c) We know that,  $RCH_2CH_2OH \xrightarrow{Oxidation} RCH_2COOH.$
- 66. c) RCOOR' + H<sub>2</sub>O  $\frac{\text{conc. H}_2\text{SO}_4}{\Delta}$

RCOOH + R'OH

The radioactive oxygen must be in acid. This technique is known as tracer technique.

**150. d)** Electron withdrawing groups or atoms increase the strength of acid. More the electron withdrawing power, more the strength of acid.

In halogens electron withdrawing power is, F > Cl > Br > I.

Hence, CH2ICOOH is stable acid.

- 153. b) Strength of acid depends upon distance between electron withdrawing group and COOH group. Distance between electron withdrawing group and COOH group increases, the strength of the acid decreases.
- 164. d) Inductive effect diminishes with increases in the number of σ-bonds through which the effect operates. This makes CICH<sub>2</sub>COOH more acidic than CICH<sub>2</sub>CH<sub>2</sub>COOH. An electronegative atom Cl is an electron withdrawing group and acid strengthener. Therefore, CICH<sub>2</sub>COOH is a stronger acid than CH<sub>3</sub>COOH. Two chlorine atoms are more electron withdrawing than single Cl atom. So, Cl<sub>2</sub>CHCOOH stronger acid than CICH<sub>2</sub>COOH.
- 172. a) Molecular formula C<sub>4</sub>H<sub>8</sub>O<sub>2</sub> is acids and esters. The compound react with CH<sub>3</sub>MgI means it must be ester. The ester gives 2-propanol with excess CH<sub>3</sub>MgI, it means that ester is formate ester.
- 174. b) RCOOH + H<sub>2</sub>O → RCOO<sup>-</sup> + H<sub>3</sub>O<sup>+</sup>
   Because of resonance stabilization of carboxylate ion, the equilibrium shifts in the forward direction, thereby forming more and more of hydronium ions.
- 175. a) Formic acid has aldehydic group H-C=O, therefore reduces Tollen's reagent to give silver mirror test.
- 176. b)  $HCOOC_2H_5 + CH_3MgI \xrightarrow{dry \, ether} H_3O^+ \rightarrow$   $CH_3CHO + MgIOH + C_2H_5OH$   $CH_3CHO + CH_3MgI \xrightarrow{dry \, ether} H_3O^+ \rightarrow$

MgIOH + CH<sub>3</sub>CHOHCH<sub>3</sub>

- 177. d) Carboxylic acids are weak acids, they dissociate lesser extent in water and equilibrium is shifted to left. Hence, aqueous solution of carboxylic acids contains undissociated acid molecules.
- 178. a)  $CH_3-CH_2-COOH + Br_2 \xrightarrow{Red P} CH_3-CHBr-COOH$ It shows optical isomerism.

 $C_2H_5COONH_4 \xrightarrow{\Delta} C_2H_5CONH_2 + H_2O$ 183. b) CH<sub>3</sub>COONa + CH<sub>3</sub>COCl sod. acetate acetyl chloride (CH<sub>3</sub>CO)<sub>2</sub>O + NaCl184. a) CH<sub>3</sub>COOH + PCl<sub>5</sub> -CH<sub>3</sub>COCl + POCl<sub>3</sub> + HCl 185. c) We know that, CH<sub>3</sub>COOH + NaHCO<sub>3</sub> →  $CH_3COONa + CO_2 + H_2O.$ Thus in this reaction acetic acid (CH<sub>3</sub>COOH) reacts with NaHCO<sub>3</sub> to give sodium salt and carbon dioxide. 187. a) We know that, 3CH<sub>3</sub>COOH + PCl<sub>5</sub> --- $3CH_3COCl + POCl_3 + HCl.$ Thus in this reaction PCl<sub>5</sub> react with an acetic acid (CH<sub>3</sub>COOH) to produce acetyl chloride. 188. c) We know that,  $CH_3COCI + 2NH_3 \longrightarrow CH_3CONH_2 + HCI.$ Thus in this reaction, acetyl chloride (CH<sub>3</sub>COCl) when heated with ammonia (NH<sub>3</sub>)

### produces acetamide (CH<sub>3</sub>CONH<sub>2</sub>). 189. c) Electron donating alkyl groups decrease the strength of acid. That is why formic acid is strongest acid than acetic acid.

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