**11**.

## Alcohols, Phenols and Ethers

#### Section-I: Alcohols

#### 11.1 INTRODUCTION

**Definition of alcohol:** These are hydroxy derivatives of hydrocarbons obtained by replacing one of more hydrogen atoms from corresponding hydrocarbons by hydroxy groups.

e.g. 
$$CH_3 - OH$$
,  $C_2H_5 - OH$ 

#### 11.2 ISOMERISM IN ALCOHOLS

**Monohydric alcohols:** They shows chain, position, optical isomerism themselves and functional isomerism with ethers. Total number of isomeric alcohols can be calculated by formula,  $1 = 2^{n-2}$ .

**Ethers:** They shows metamerism, optical isomerism themselves and functional isomerism with alcohols.

Total number of isomeric ethers can be calculated by formula,  $I = 2^{n-1} - 1 - (2^{n-2})$  from  $C_3$  onwards. Total number of isomeric alcohols and ethers can be calculated by formula,  $1 = 2^{n-1} - 1$ .

- e.g. i)  $C_2H_6O$  (2-isomers)
  - a) CH<sub>3</sub>CH<sub>2</sub>OH ethyl alcohol
  - b) CH,OCH, dimethyl ether
  - ii) C<sub>3</sub>H<sub>5</sub>O (3–isomers. 2 alcohol, 1ether) Alcohols –
  - a) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>OH n-propyl alcohol
  - b) CH<sub>3</sub>CHOHCH<sub>3</sub> iso–propyl alcohol Ether –
  - a) CH<sub>2</sub>OC<sub>2</sub>H<sub>5</sub> ethyl methyl ether
  - iii)  $C_4H_{10}O$  (7-isomers.4 alcohols, 3 ethers)

Alcohols -

- a) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>OH n-butyl alcohol
- b) (CH<sub>2</sub>)<sub>2</sub>CHCH<sub>2</sub>OH isobutyl alcohol
- c) CH<sub>3</sub>CH<sub>2</sub>CHOHCH<sub>3</sub> 2° butyl alcohol
- d) (CH<sub>3</sub>)<sub>3</sub>COH 3° butyl alcohol

'a' and 'b' are chain isomers.

'a' and 'c', are position isomers. Only 'c' show optical isomerism.

Ethers – Following ethers are metamers.

- a) C<sub>2</sub>H<sub>5</sub>OC<sub>2</sub>H<sub>5</sub> diethyl ether
- b) CH<sub>2</sub>OCH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub> methyl n-propyl ether
- c) CH<sub>3</sub>OCH(CH<sub>3</sub>), methyl isopropyl ether

## 11.3 STRUCTURE OF -OH GROUP IN ALCOHOL

In alcohols, oxygen atom of the –OH group is attached to sp<sup>3</sup> hybridised carbon atom, by overlapping of sp<sup>3</sup> hybrid orbital of carbon and p–orbital of oxygen i.e sp<sup>3</sup> – p overlapping.

The bond angle C–O–H is slightly less than the tetrahedral angle. It is due to repulsion of lone pair electrons of the oxygen atom.



#### 11.4 PREPARATIO METHODS

1. Alkaline hydrolysis of alkyl halides:

$$R-X + NaOH \xrightarrow{aq.} R-OH + NaX$$

2. From hydration of alkene (Olefin) by using conc. H,SO<sub>4</sub>:

$$\begin{array}{lll} \mathrm{CH_2} = \mathrm{CH_2} + \mathrm{H_2SO_4} & \xrightarrow{\mathrm{cold}} & \mathrm{CH_3-CH_2-} \\ \mathrm{OSO_3H} & & & \\ \mathrm{ethylene} & & \mathrm{conc.} & \mathrm{ethyl} \, \mathrm{hydrogen} \, \mathrm{sulphate} \\ \mathrm{CH_3-CH_2-OSO_3H} + \mathrm{H_2O} & \xrightarrow{\mathrm{boil}} & \mathrm{CH_3-CH_2-} \\ \mathrm{OH} + \mathrm{H_2SO_{42}} & & \\ \mathrm{ethyl} \, \mathrm{alcohol} & & & \\ \end{array}$$

#### Mechanism:

The mechanism of reaction involves three steps.

**Step–(i):** Protonation of water followed by attack of electrophilic H<sub>2</sub>O<sup>+</sup> to form carbocation.

$$H_2O + H^+ \longrightarrow H_3O^+$$

$$C = C + H = C + H_2O$$

Step-(ii): Nucleophilic attack of water on carbocation.

$$\begin{array}{c|c} H & H & H \\ \downarrow C - C & \downarrow H_2 & \longleftarrow \\ \downarrow C - C - C - H_2 & \longleftarrow \\ \end{array}$$

3. By hydroboration oxidation of alkenes (HBO Reaction of alkenes):

e.g. i) 
$$2 \text{ CH}_2 = \text{CH}_2 + (\text{H} - \text{BH}_2)_2 \xrightarrow{\text{THF}}$$
  $2 \text{ CH}_3 - \text{CH}_2 - \text{BH}_2$  ethyl boron

$${\rm CH_3-CH_2-BH_2} + {\rm CH_2} = {\rm CH_2} \longrightarrow$$
  $({\rm CH_3-CH_2})_2 \ {\rm BH}$  diethyl boron

$$(CH_3 - CH_2)_2$$
 BH +  $CH_2 = CH_2$   $\longrightarrow$   
triethyl boron  $(CH_3 - CH_2)_2$  BH diethyl boron

$$(CH_3 - CH_2)_3 B + 3 HO \cdot OH \xrightarrow{NaOH}$$

$$3CH_3 - CH_2 - OH + B(OH)_3$$
ethyl alcohol

ii) 
$$CH_3 - C = CH_2$$
 i)  $\xrightarrow{I) B_2H_6 \text{ in THF}}$   $\downarrow$   $CH_3$ 

iso butylene  $CH_3$ – $CH(CH_3)CH_2$ –OH

#### Note:

- The overall process occurs with addition of H<sub>2</sub>O by anti Markownikoffs rule. Boron is electron deficient compound and behave as electrophiles.
- ii) The good yield of primary alcohols is obtained by this method.
- iii) In hydroboration oxidation (HBO) process, the carbocation are not formed hence no rearrangement takes place.
- iv) 1°, 2°, 3° alcohol can be obtained by this method.

$$\begin{array}{c} \operatorname{CH_3} \\ | \\ \operatorname{CH_3} - \operatorname{C} - \operatorname{CH} - \operatorname{CH_2} \\ | \\ \operatorname{CH_3} \end{array} \xrightarrow{i) \operatorname{B_2H_6, \, THF}} \\ \downarrow \\ \operatorname{CH_3} \end{array}$$

$$CH_3 - \begin{vmatrix} CH_3 \\ -C-CH_2-OH \\ -CH_3 \end{vmatrix}$$

- 4. From reduction of carbonyl compounds:
- I) By the reduction of aldehydes and ketones: Aldehydes on reduction gives 1° alcohols. While ketones on reduction gives 2° alcohols. These are reduced by two methods.
- i) Catalytic hydrogenation in the presence of nickel or raney nickel platinum or palladium as catalyst at 413K to 453K. Raney nickel is an alloy of aluminium and nickel (50 %) leached by NaOH solution.
- ii) Reduction by using Na Hg +  $\rm H_2O$  or NaBH $_4$  or Na + ethanol or Zn + HCl or LiAlH $_4$  or  $\rm B_2H_6$  as reducing agents.
- a) Catalytic hydrogenation by using Hz and Ni or Pt or Pd at 413 K: When aldehydes and ketones are reduced by passing H<sub>2</sub> gas in presence of Ni or raney Ni or Pt or Pd catalyst at 413K to 453K to give alcohols. Which affect unsaturation.

$$> C = O + H_2 \xrightarrow{Ni 413 \text{ K}} > CH - OH$$

b) Reduction by using NaHg + H<sub>2</sub>O (Bouveault- Blanc reduction): When aldehydes and ketones are reduced by Na.Hg + H<sub>2</sub>O or LiAlH<sub>4</sub> or NaBH<sub>4</sub> gives alcohols.

Na.Hg + H<sub>2</sub>O 
$$\longrightarrow$$
 NaOH + Hg + [H]  
> C = O + 2 [H]  $\xrightarrow{\text{NaHg + H}_2\text{O}}$  > CH - OH

Note: LiAlH<sub>4</sub> or B<sub>2</sub>H<sub>6</sub> or NaBH<sub>4</sub> does not reduce  $>C = C < -C \equiv C$  bonds.

II) By reduction of carboxylic acids:.

$$R$$
– $COOH + 4[H] \frac{LiAlH_4/Ether}{H_3O^+}$ 

$$R-CH_2-OH + H_2O$$

III) By reduction of esters:

Esters are reduced by LiAlH<sub>4</sub> or by passing hydrogen gas in the presence of catalyst like Ni or Pt or Pd or raney Ni to give mixture of alcohol. But it is easily reduced by LiAlH<sub>4</sub> in ether because it is more reactive.

$$\begin{array}{c} O \\ \parallel \\ R-C -OR'+2H_2 \xrightarrow{\quad \mathrm{Ni} \quad} R-CH_2-OH+R' \\ -OH \end{array}$$

From Grignards reagent (From action of R

 MgX on carbonyl compounds)

 Preparation of 1<sup>o</sup> alcohols:

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

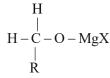
Mg-complex

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

$$R - CH_2 - OH + MgXOH$$
 primary alcohol

#### Preparation of secondary alcohols:

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



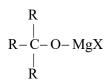
Mg-cornplex

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

sec. alcohol

#### **Preparation of tertiary alcohols:**

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



Mg-complex

ter. alcohol

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

#### 11.5 REACTIVITY OF ALCOHOLS

Alcohols under goes two type of reactions,

# a) Breaking of O-H bond (Acidity of alcohols) : The acidic character of alcohols is due to polar nature of -OH bond. An electron releasing groups (+I effect) increases electron density on oxygen leading to decrease the polarity of -O-H bond. This decrease the acidic strength. Therefore acidity of alcohol is, 1° > 2° > 3°.

The decreasing order of acidic strength of alcohol is

$$H_2O > R \rightarrow -CH_2 - OH > \frac{R}{R} \nearrow CH - OH >$$

$$\begin{array}{c}
R \\
\downarrow \\
R \to C - OH \\
\uparrow \\
R
\end{array}$$

Alcohols are weaker acidic than water

e.g. 
$$R - O^- + H - O - H \longrightarrow R - OH + OH^-$$

alkoxide base water acid conjugate acid conjugate base This reaction shows that, water is better proton

Alkoxide ion is better proton accepter than hydroxide ion, which suggest that sodium alkoxides are stronger base than NaOH.

b) Breaking of C – O bond: Electron releasing groups (+ I effect) increases electron density on carbon leading to increase the polarity of –C –O bond. This increase the reactivity of alcohols. Therefore reactivity of alcohol is, 3° > 2° > 1°.

#### 11.6 CHEMICAL PROPERTIES

donor than alcohols.

Alcohols undergo following types of reactions,

- 1] Reaction involving breaking of O–H bond.
- 2] Reaction involving breaking of C–O bond.
- 3] Reaction involving both alkyl and O–H group.

#### 11.6.1 Reaction involving breaking of O-H bond:

a) Reaction with metals:

$$2R-OH + 2Na \longrightarrow 2R-ONa + H_2$$
  
alcohol sod. alkoxide  
 $6R-OH + 2Al \longrightarrow 2(R-O)_3Al + 3H_2$   
alcohol aluminium alkoxide  
This reaction involving cleavage of oxygen

hydrogen bond (-O-H) such type of reaction follows the order, 1° alco.  $> 2^{\circ}$  alco.  $> 3^{\circ}$  alco. Tertiary alcohols are almost inert to sodium metal and aluminium metal.

b) Reaction with carboxylic acid, acid chloride and acid unhydride (ester formation):

$$R'-OH + R-COOH \xrightarrow{H^+} R-COO-R' + H_2O$$

$$R'-OH + (R-CO)_2O \xrightarrow{H^+} R-COO-R'$$

$$+ RCOOH$$

$$R'-OH + R-COC1 \xrightarrow{Pyridine} R-COO-R'$$

$$R'-OH + R-COCI \xrightarrow{-ignum} R-COO-R' + HCI$$

#### 11.6.2 Reaction involving breaking of C-O bond:

- 1. Reaction with cone. HX:
- a) Reaction with cone. HCl (Grooves process)
  Reaction with cone. HCl in the presence of anhydrous ZnCl<sub>2</sub> with alcohol is known as Lucas test. This test is used to distinguish between 1°,

2°, 3° alcohols. R–OH + 
$$\xrightarrow{\text{HCl Anhydrus ZnCl}_2}$$
  $\xrightarrow{\Delta}$  cone. R–Cl +  $\text{H}_2\text{O}$ 

b) Reaction with cone. HBr:

$$R - OH + HBr \xrightarrow{NaBr + conc.H_2SO_4} R - Br + H_2O$$

$$conc.$$

**Note:** In the case of HBr, a small amount of concentrated H<sub>2</sub>SO<sub>4</sub> is also added for the reaction with primary alcohols. But no catalyst is added with secondary and tertiary alcohols, which get dehydrated in the presence of concentrated H<sub>2</sub>SO<sub>4</sub>.

$$R - OH + HI \longrightarrow R - I + H_2O$$

#### Mechanism of action of HX:

**Step-I:** Formation of oxocation by protonation of alcohol.

$$\begin{array}{c}
H \\
| \\
R - O - H + H^+ \longrightarrow R - O - H \\
& \text{oxocation}
\end{array}$$

**Step-II:** Loss of water from oxocation gives carbocation

$$\begin{array}{c}
H \\
R - O - H
\end{array}
\longrightarrow R^{+}$$

**Step-III:** Action of X– on carbocation

$$R^+ + X^- \longrightarrow R - X$$

Dehydration of alcohols: Dehydration of alcohol is a β-elimination reaction in which H<sub>2</sub>O molecule is removed form adjacent carbon atoms (α, β carbon atoms). Dehydration of all types of alcohols leads to olefins.

Alcohols are dehydrated by cone.  $H2SO_4$  or  $Al_2O_3$  or  $H_3PO_4$  or  $P_2O_5$  or  $KHSO_4$  or  $ThO_2$  or Boric acid ( $H_3BO_3$ ). Ease of dehydration is,  $3^{\circ}>2^{\circ}>1^{\circ}$  alcohols.

a) Dehydration by using cone. H<sub>2</sub>SO<sub>4</sub> (liquid phase dehydration): Alcohol which do not contain β-Hatom on dehydration gives alkene through rearrangement reaction in the presence of acid only.

$$\begin{array}{c|c} CH_{3} & CH_{3} \\ \hline CH_{3} - C - CH_{2} - OH \\ \hline CH_{3} & CH_{3} - C = CH - CH_{3} + H_{2}O \\ \hline CH_{3} & 2 - methyl 2 - butene \end{array}$$

neo-pentyl alcohol

This reaction can be proceeding through following steps –

$$\begin{array}{c} \text{CH}_{3} \\ \text{CH}_{3} - \text{C} - \text{CH}_{2} - \text{OH} \\ \text{CH}_{3} \\ \text{CH}_{3} \end{array} \xrightarrow[]{\text{Dehydration}} \begin{array}{c} \text{CH}_{3} \\ \text{Dehydration} \\ \text{H}_{2}\text{SO}_{4} \end{array} \xrightarrow[]{\text{CH}_{3} - \text{C} - \text{CH}_{2} - \text{O}^{+} - \text{H}} \xrightarrow[]{\text{CH}_{3} - \text{C} - \text{CH}_{2}^{+}} \\ \text{CH}_{3} \\ \text{Oxonium ion} \end{array}$$

3° pentyl carbonium ion

2-methyl 2-butene

#### Mechanism of dehydration:

i) Formation of protonated alcohol  $(R - \overset{\oplus}{O} H_2)$ 

protonated alcohol (oxonium ion)

ii) Formation of carbonium ion by loss of water molecule

carbonium ion

iii) Formation of alkene by loss of proton

$$\begin{array}{c|c} -C & -C^{\oplus} \\ \downarrow & & & \downarrow \\ H & & & -H^{\oplus} \end{array} \begin{array}{c} -C = C - \\ \text{alkene} \end{array}$$

In all cases, reaction intermediate is carbocation, and in higher alcohols there may be 1–2 hydride shift or 1–2 methyl shift to form more stable carbocation.

e.g. 
$$\begin{array}{c}
CH_{3} \\
CH_{3} - C - CH_{2} - OH \\
CH_{3}
\end{array}$$

$$\begin{array}{c}
CH_{3} CH_{3} \\
CH_{3} - C = C - H \\
2 - methyl but - 2 - ene
\end{array}$$

$$\begin{array}{c}
CH_{3} CH_{3} \\
CH_{3} - C = C - H \\
2 - methyl but - 2 - ene$$

# 11.7 Oxidation of primary, secondary and tertiary alcohols OR Distinction of primary, secondary and tertiary alcohols by oxidation reaction:

The oxidation can be carried out by various oxidising agents such as

- i)  $K_2Cr_2O_7 + dil. H_2SO_4$
- ii) KMnO<sub>4</sub> + dil. H<sub>2</sub>SO<sub>4</sub>
- iii) Neutral and aqueous KMnO<sub>4</sub>
- iv) Oil. HNO<sub>3</sub>
- v) CrO<sub>3</sub> Chromium oxide
- vi) H<sub>2</sub>CrO<sub>4</sub> Chromic acid
- vii) Alkaline KMnO<sub>4</sub>
- viii) Pyridinium chloro chromate(PCC) or Pyridinium dichromate (PDC) C<sub>5</sub>H<sub>5</sub>NHCrO<sub>3</sub>Cl<sup>-</sup> Mild oxidising agent. It is also known as Collin's reagent. PCC is mixture of pyridine.CrO<sub>3</sub> and HCl in 1 : 1 : 1 ratio.

Ease of oxidation of alcohol is  $1^{\circ} > 2^{\circ} > 3^{\circ}$ 

#### a) Primary alcohols:

#### b) Secondary alcohols:

$$\begin{array}{c} \text{CH}_{3} \\ \text{CH}_{3} - \text{CH} - \text{OH} + [\text{O}] \xrightarrow{K_{2}\text{Cr}_{2}\text{O}_{7} + \text{dil}.\text{H}_{2}\text{SO}_{4}} & \text{CH}_{3} - \text{C} = \text{O} + \text{H}_{2}\text{O} \\ \text{2-propanol} & \text{acetone} \\ \\ \text{CH}_{3} \\ \text{CH}_{3} - \text{C} = \text{O} + 4[\text{O}] \xrightarrow{K_{2}\text{Cr}_{2}\text{O}_{7} + \text{dil}.\text{H}_{2}\text{SO}_{4}} & \text{CH}_{3} - \text{COOH} + \text{CO}_{2} + \text{H}_{2}\text{O} \\ & \text{acetic acid} \end{array}$$

#### c) Tertiary alcohols:

$$\begin{array}{c|c} \operatorname{CH_3} & & \operatorname{CH_3} \\ \operatorname{CH_3} - \operatorname{C} - \operatorname{OH} & & & \operatorname{CH_3} \\ | & & & & \operatorname{strong\ heat} \end{array} \rightarrow \begin{array}{c} \operatorname{CH_3} \\ \operatorname{CH_3} - \operatorname{C} - \operatorname{H_2} \operatorname{O} \\ | & & & & | \\ \operatorname{CH_3} - \operatorname{C} - \operatorname{H_2} \operatorname{O} \end{array}$$

$$\begin{array}{c}
CH_{3} \\
CH_{2} = C + 4[O] \\
CH_{3}
\end{array}
\xrightarrow{K_{2}Cr_{2}O_{7} + dil.H_{2}SO_{4} \atop drastic oxidation}$$

$$\begin{array}{c}
CH_{3} \\
CH_{3} - C = O + H_{2}O + CO_{2}
\end{array}$$

$$\begin{array}{c} \text{CH}_{3} \\ | \\ \text{CH}_{3} - \text{C} = \text{O} + 4[\text{O}] \end{array} \xrightarrow{\begin{array}{c} \text{K}_{2}\text{Cr}_{2}\text{O}_{7} + \text{dil.H}_{2}\text{SO}_{4} \\ \text{drastic oxidation} \end{array}} \begin{array}{c} \text{CH}_{3} - \text{COOH} + \text{CO}_{2} + \text{H}_{2}\text{O} \end{array}$$

Actually, formic acid is obtained along with acetic acid but it readily undergoes further oxidation gives CO<sub>2</sub> and water.

acetic acid

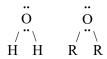
#### Section -II Ethers

#### 11.8 INTRODUCTION

Ethers are dialkyl derivatives of water obtained by replacing both hydrogen atoms from water by alkyl groups.

$$H-O-H \xrightarrow{-2H} R-O-R$$

Ethers are represented as, R–O–R, known as ether linkage. These are represented by general formula  $C_nH_{2n+2}O$ . In ethers, central oxygen atom is  $sp^3$ –hybridised state forming two sigma bonds with two alkyl groups leads to angular V shaped geometry like water, having bond angle 110°, bond angle in water is 104.5°. Thus ethers have bent structure and dipolar in natur.



#### 11.8.1 ISOMERISM

They shows optical, metamerism themselves and functional isomerism with monohydric alcohols.

#### Metamerism:

Metamerism is present in same class of compounds.

Organic compound having same molecular formula and different structural formula and different alkyl groups (nature and type) attached to either side of oxygen atom.

e.g. i) Molecular formula  $C_4H_{10}O$  have three ethers and all are metamers.

CH<sub>3</sub>-O-CH(CH<sub>3</sub>)<sub>2</sub> methyl iso propyl ether

ii) Methyl n-propyl ether and methyl iso propyl ether are position isomers

$$CH_3-O-CH_2-CH_2-CH_3$$
,  $CH_3-O-CH(CH_3)_2$ 

**Note:** Metamers are either chain or position isomers. If metamerism is mentioned never write chain or position isomerism.

#### 11.9 PREPARATION METHODS

1. Williamson's synthesis: When alkyl halides are heated with alcoholic sodium alkoxide to give ethers. This is the laboratory method used for preparation of simple and mixed aliphatic and aromatic ethers. But this method is conveniently used to prepare mixed ether

$$R-X+$$
  $R-ONa \xrightarrow{\Delta} R-O-R + NaX$ 

alkyl halide sod. alkoxide ether

e.g. When methyl chloride is heated with alc. sodium methoxide, gives dimethyl ether.

**Mechanism:** It is SN<sup>2</sup> reaction of R–X, in which –X is replace by RO<sup>-</sup>

Step-1 : 
$$R-ONa \longrightarrow RO^- + Na^+$$

Step-2:

Step- 3: 
$$Na^+ + X^- \longrightarrow NaX$$

**2. From diazomethane and alcohol:** This method is used to prepare simple and mixed ethers.

When alcohols are reacted with diazomethane in the presence of fluoroboric acid HBF<sub>4</sub> or BF<sub>3</sub> as catalyst to give ethers.

$$\text{R-OH} + \text{CH}_2\text{N}_2 \xrightarrow{\quad \text{HBF}_4 \quad} \text{R-O-CH}_3 + \text{N}_2 \uparrow$$

**Note:** This is also known methylation of alcohols.  $CH_2N_2$  acts as methylating agent. Ether produced is pure will always have methyl as one of the group. These are nucleophilic substitution reactions.

3. By continuous etherification process (Intermolecular dehydration of alcohols):

#### Step-I:

When ethyl alcohol is heated with cone. H<sub>2</sub>SO<sub>4</sub> at 413K, gives ethyl hydrogen sulphate.

$$C_2H_5$$
-OH +  $H_2SO_4$   $\xrightarrow{413K}$   $C_2H_5$ -OSO $_3$ H +  $H_2$ O ethanol conc. ethyl hydrogen sulphate

#### Step-II:

Ethyl hydrogen sulphate is reacted with another molecule of ethyl alcohol gives ethoxy ethane.

$$\begin{array}{c} {\rm C_2H_5\text{-}OSO_3H} + {\rm C_2H_5\text{-}OH} \xrightarrow{\phantom{-}413{\rm K}} \\ {\rm C_2H_5\text{-}O\text{-}C_2H_5} + {\rm H_2SO_4} \\ {\rm ethoxy\ ethane} \end{array}$$

The H<sub>2</sub>SO<sub>4</sub> produce in second step is used to convert more and more alcohol into ether. Hence this method is known as continuous etherification process.

#### **Purification:**

Ether obtained by this method contains  $H_2O$ , alcohol and  $H_2SO_4$ , To remove these impurities it is first shaken with NaOH, which neutralise  $H_2SO_4$  then add  $CaCl_2$  to remove alcohol. Then dried over fused  $CaCl_3$ .

#### Mechanism:

 Formation of ether is SN<sup>2</sup> reaction involving the attack of proton on alcohol to form protonated alcohol.

$$CH_3$$
- $CH_2$ - $\overset{\cdot \cdot \cdot}{O}$  - H + H<sup>+</sup>  $\longrightarrow$ 

protonated alcohol

ii) Attack of protonated alcohol on another alcohol to form oxocation.

$$CH_3 - CH_2 - \ddot{O} : + CH_3 - CH_2 \xrightarrow{CH_3} H \xrightarrow{Slow} H$$

$$CH_{3} - CH_{2} - \overset{\oplus}{O} + CH_{2} - CH_{3} + H_{2}O$$

iii) Loss of proton from oxocation to form ether

$$\begin{array}{c} \operatorname{CH_3} - \operatorname{CH_2} - \overset{\oplus}{\operatorname{O}} + \operatorname{CH_2} - \operatorname{CH_3} \\ \downarrow \\ \operatorname{H} \end{array} \xrightarrow{\operatorname{Fast}}$$

$$CH_3 - CH_2 - \ddot{O} - CH_2 - CH_3 + H^+$$

#### 11.10 CHEMICAL PROPERTIES

These are chemically less reactive (inert) towards most of the base, metals, oxidising agents, reducing agents and RMgX. Thus the properties of ethers are due to alkyl group, lone pair of electrons on oxygen and breaking of C–O bond. The C–O bond is not so stable as C–C bond. The C–O bond is ruptured in the presence of number of reagents like cone. HBr or cone. HI or dil. H<sub>2</sub>SO<sub>4</sub> and undergoes nucleophilic substitution reactions.

- I) Reactions due to breaking of C–O bond:
  Reaction with HX: The cleavage of ethers takes place with concentrated HI or HBr. The reactivity of HX is HI > HBr > HCl. The reaction takes place with only conc. HI and HBr because these are sufficiently acidic. Reaction does not takes place with conc. HCl because it is not sufficiently acidic.
- 1. Reaction with cold cone. or dry HI: When ethers are reacted with cold HI to give alcohols and alkyl iodides.

$$R - O - R + HI \xrightarrow{\text{cold}} R - I + R - OH$$
  
ether alkyl iodide alcohol  
e.g. When methoxy ethane is reacted with cold  
HI to give methyl iodide and ethyl alcohol.

$$C_2H_5$$
-O-CH $_3$  + HI  $\xrightarrow{cold}$   $CH_3$ -I +  $C_2H_5$ -OH methoxyethane methyl iodide ethanol

**Note:** The nucleophile 1– attacks –CH<sub>3</sub> group and not –CH<sub>2</sub> group of ethyl group, because the

positive charge on  $-CH_2$  is reduced by electron donating effect of  $-CH_3$  in ethyl group. Consequently, positive charge on  $-CH_3$  is more than on  $-CH_3$  of ethyl group.

#### Mechanism:

#### Step - I:

The reaction of ethyl methyl ether with cone HI starts with protonation of ether molecule to form oxonium ion.

$$CH_3 - \overset{\dots}{O} - C_2H_5 + H - I \Longrightarrow$$

$$H$$

$$CH_3 - \overset{\dots}{O} - C_2H_5 + I^-$$

$$Oxonium ion$$

#### Step – II:

Iodide is good nucleophile, it attack on least substituted carbon of oxonium ion and displace ethanol by SN<sup>2</sup> mechanism.

$$CH_{3} \stackrel{\stackrel{f}{\circ}}{-} C_{2}H_{5} \stackrel{\ominus}{+} \stackrel{\ominus}{I} \longrightarrow \begin{bmatrix} H \\ I - CH_{3} - O - C_{2}H_{5} \\ T.S. \end{bmatrix} \longrightarrow CH_{3} - I + C_{2}H_{5} - OH$$

#### **Branched ethers:**

If mixed branched ethers containing one the alkyl group is tertiary then halide formed is tertiary.

e.g. i) 
$$C_2H_5-O-CH(CH_3)_2 + HI \xrightarrow{cold}$$
  
2-ethoxy propane  $C_2H_5-I + (CH_3)_2CH-OH$   
ethyl iodide iso propyl alcohol

ii) 
$$CH_3$$
-O- $C(CH_3)_3 + HI \xrightarrow{cold}$ 
 $t$ -butyl methy ether
$$CH_3$$
-OH +  $(CH_3)_3$ C-I
methanol  $t$ -butyl iodide

#### Mechanism of t-butyl methy ether:

It is three step reaction follows  $SN^1$  mechanism and formation of  $(CH_3)_3$   $C^+$  carbocation.

**Step-I:** Protonation of ether to form oxonium ion

$$\begin{array}{c} \operatorname{CH_3} \\ \mid \\ \operatorname{CH_3} - \operatorname{C} - \operatorname{O} - \operatorname{CH_3} + \operatorname{HI} \\ \mid \\ \operatorname{CH_3} \end{array}$$

$$\begin{array}{c} \operatorname{CH_3} \\ | \\ \operatorname{CH_3} - \operatorname{C} - \operatorname{O} - \operatorname{CH_3} + \operatorname{I}^- \\ | \\ \operatorname{CH_3} \end{array}$$

**Step–II:** Dissociation of C – O bond and formation of stable t–butyl carbonium ion by SN¹ reaction.

$$(CH_3)_3C - \underset{\oplus}{\overset{H}{\circ}} - CH_3 \longrightarrow (CH_3)_3C^+ + CH_3 - OH$$

t-butyl carbonium ion

**Step-III:** Attack of nucleophile (I<sup>-</sup>) on t-butyl carbonium ion.

$$(CH_3)_3 C^+ + I^- \longrightarrow (CH_3)_3 C - I$$

t-butyl iodide

**Note:** This reaction is useful for detection of alkyl group in mixed ether.

#### **Aromatic ethers:**

e.g. i) When anisole is reacted with cold HI to give phenol and iodo methane.

$$C_6H_5$$
-O-CH<sub>3</sub> + HI  $\xrightarrow{\text{cold}}$   $C_6H_5$ -OH+CH<sub>3</sub>-I anisole

In case of anisole methyl phenyl oxonium ion is formed by protonation

The bond in  $\mathrm{CH_3}-\mathrm{O}$  is weaker than  $\mathrm{C_6H_5}-\mathrm{O}$  because carbon of phenyl group is  $\mathrm{sp^2}$  hybridised state and there is a partial double bond character. Therefore attack of lion breaks  $\mathrm{CH_3}-\mathrm{O}$  bond to form  $\mathrm{CH_3}-\mathrm{I}$ , and  $\mathrm{C_6H_5}-\mathrm{O^-}$  combine with  $\mathrm{H^+}$  to form  $\mathrm{C_6H_5}-\mathrm{OH}$ .

ii) 
$$\langle \bigcirc \rangle$$
 + HI  $\xrightarrow{\text{cold}}$ 

benzyl phenyl ether

$$\bigcirc$$
 OH +  $\bigcirc$  CH<sub>2</sub> – I

phenol benzyl iodide

II) Reaction with hot cone. or dry (excess) HI:

$$R - O - R + HI \xrightarrow{\Delta} 2R - I + H_2O$$

Note: Both alkyl group produces alkyl iodide.

III) Acid hydrolysis (Reaction with dil. H<sub>2</sub>SO<sub>4</sub>): When ethers are hydrolysed by dilute H<sub>2</sub>SO<sub>4</sub> under pressure to give alcohols.

$$R-O-R+H_2O \xrightarrow{\quad dil.\,H_2SO_4\Delta \quad \quad } R-OH+R-OH$$

ether

alcohol

Note: Both alkyl group produces alcohols.

#### 11.11 ELECTROPHILIC SUBSTITUTION REACTION OF AROMATIC ETHERS

The alkoxy group attached to benzene ring is ortho and para directing like phenol

1. **Halogenation:** Alkyl phenyl ether reacts with bromine in acetic acid in the absence of FeCl<sub>3</sub> or FeBr<sub>3</sub> catalyst to give ortho bromo alkyl phenyl ether (10%) and parabromo alkyl phenyl ether (90%).

e.g. 
$$CH_3COOH$$

$$CH_3COOH$$

$$CH_3COOH$$

$$CH_3COOH$$

$$CH_3COOH$$

$$CH_3COOH$$

$$CH_3COOH$$

$$CH_3COOH$$

$$CH_3COOH$$

$$TO = Br$$

$$TO = B$$

2. **Nitration:** Alkyl phenyl ether reacts with nitrating mixture (cone. HNO<sub>3</sub> + cone. H<sub>2</sub>SO<sub>4</sub>) to give a mixture of o–nitro alkyl phenyl ether (minor) and p–nitro alkyl phenyl ether (major).

- 3. Friedel Craft reactions:
  - a) Friedel Craft alkylation: In this reaction alkyl groups is attached to ortho and para position.

b) Friedel- Craft acylation: Acylation can be carried out by acylating agents like RCOCl, (RCO)<sub>2</sub>O, R-COOH

e.g. 
$$O-R$$
 $O-R$ 
 $O-R$ 

#### 11.12 CROWN ETHERS

Charles J. pederson discovered crown ether. It is macrocyclic polyether containing large ring of carbon and oxygen atoms. Crown ethers are named as n-crown-m. Where 'n' is the total carbon and

oxygen atoms and 'm' is the total oxygen atoms.

a) The first crown ether is 18-crown 6-ether



18-crown 6-ethers

The cavity inside is able to trap K<sup>+</sup> but not Na<sup>+</sup> and Li<sup>+</sup>

b) The another crown ether is 15–crown–5 ether, it is able to trap Na<sup>+</sup> ions.





15-crown 5-ether

crown ether sodium ion complex

In crown ether each oxygen atom is co-ordinatly bonded with sodium atom.

- c) Crown ether also form complex with NH<sub>4</sub><sup>+</sup> and R-NH<sub>3</sub><sup>+</sup>
- d) Stability of these complexes depends upon size of the metal ion and cavity available in the ring of perticular crown ether.
- e) Application:
  - i) It is used for increasing solubility of ionic salt in non-polar solvent
  - ii) They also act as catalyst in certain reactions involving organic salt.
  - iii) They are used to extract specific ion from mixtures and enrich isotopic mixtures.
  - iv) It is used to remove radiactive elements from radioactive waste.
  - vi) Specific crown ether 18–crown–6 is used to extract cesium and strontium.

#### Section -III: Phenols

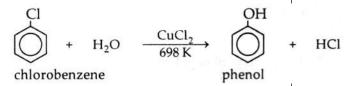
#### 11.1 INTRODUCTION

Phenols are aromatic hydroxy compounds in which one or more hydroxy groups are directly attached to benzene nucleus.

Simple phenol is hydroxy benzene or phenyl hydroxide or benzenol or phenol it self also known as carbolic acid. It is represented as Ar–OH or Ph–OH where Ar or Ph is  $C_6H_5$ – group.

#### 11.2 PREPARATION OF PHENOL (CARBOLIC ACID)

1. From chlorobenzene by Raschig's method (1943) Industrial method:



2. From chlorobenzene by Dow's method (1928) Industrial method:

3. From benzene sui phonic acid:

4. From oxidation of cumene (isopropyl benzene or 2-propyl benzene or 2-phenyl propane) :

$$\begin{array}{c} \text{CH}_{3} \\ \text{CH}_{3}\text{-C-H} \\ \\ \text{CH}_{3}\text{-C-OOH} \\ \\ \text{CH}_{3}\text{-C-OOH} \\ \\ \text{CH}_{3}\text{-C-OOH} \\ \\ \text{CH}_{3}\text{-C-OOH} \\ \\ \text{Cumene (2-propyl benzene)} \\ \end{array}$$

5. From diazotisation reaction:

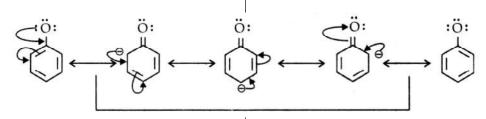
#### 11.3 ACIDITY OF PHENOLS

The reaction of phenol with sodium, potassium, aluminium metal, NaOH, KOH, indicates the acidic nature of phenol. It does not react with NaHCO<sub>3</sub>, Na<sub>2</sub>CO<sub>3</sub>.

In phenols – OH group is directly attached to  $sp^2$  hybridised carbon atom, this increase the polarity of O – H bond and increase the ionisation of phenols than alcohols.

$$C_6H_5OH + H_2O \Longrightarrow C_6H_5O^- + H_3O^+$$
  
phenoxide ion

Phenoxide ion is more stabilized than phenol by resonance. Hence phenol is acidic in nature.



Resonating structure of phenoxide ion

#### 11.4 EFFECT OF SUBSTITUENTS ON ACIDITY OF PHENOLS

The presence of electron attracting and electron donating groups affects the acidic strength of phenols.

1. Effect of electron withdrawing group (EWG) like -NO<sub>2</sub>, - CHO, - COOH, -CN etc on acidity of phenols:

Electron withdrawing grou ps or atoms stabilize the phenoxide ion and increase the acidic strength of phenols.

$$OH \longrightarrow O^{\ominus}$$

Acidic strength more W-stabilize the anion

But this effects is more when substituents is present on ortho and para position than meta position.

It follows the order p > o > m > Ph - OH

Greater the number of electron withdrawing groups at ortho and para position, more the acidity of phenol. Thus increasing order acidic strength of nitrophenol is p—nitrophenol > o—nitrophenol > m—nitrophenol > phenol.

2. Effect of electron donating groups (EDG) like R- OH, -OR,-NH<sub>2</sub>-, NR<sub>2</sub> etc. on acidity of phenols:

Electron donating groups destabilize the phenoxide ion by donating electron to benzene ring and decrease the acidic strength of phenol. Acidic nature is more when electron donating group is present on meta position.

Acidic strength less

D-destabilize the anion

Thus cresols are less acidic than phenols. The decreasing order of acidic strength of cresols is,

**Note:** Lower the pKa value stronger the acid, or higher the Ka value stronger the acid. The pka value of ethanol is 15.9 hence it very very less acidic than phenol.

Compounds	pKa value
Ethanol	15.9
p-amino phenol	10.5
p-cresol	10.2
o –cresol	10.14
m-cresol	10.1
Phenol	10.0
p-chloro phenol	9.4
m-chloro phenol	9.1
o-chloro phenol	8.6
m-nitro phenol	8.3
o-nitro phenol	7.2
p-nitro phenol	7.1

Following groups increases the acidity of phenol in decreasing order.

$$-SO_3H > -COOH > -NO_2 > F > Cl > Br > I > -OH > -OCH_3 > -CH_3 > -NH_2$$

#### 11.5 CHEMICAL PROPERTIES

Phenol undergoes two types of reactions by breaking of O–H bond and by breaking of C – O bond. Hence reactions are divided in to two groups.

#### 1. Reaction involving breaking of – OH bond (acidity of phenol)

i) Reaction with Na metal

ii) Reaction with NaOH

$$OH \qquad ONa \\ + NaOH \longrightarrow O + H_2O$$

# 11.6 ELECTROPHILIC SUBSTITUTION REACTION OF PHENOL (REACTION INVOLVING BREAKING OF C – O BOND)

- 1. Bromination in different solvent:
- a) Reaction with Br<sub>2</sub> gas in inert solvent non polar solvent (CS<sub>2</sub>, CHCl<sub>3</sub>, CCl<sub>4</sub>):

OH 
$$CS_2$$
  $CS_2$   $CS_2$ 

b) Reaction with bromine water:

- 2. Nitration at different condition:
- a) Reaction with dil. HNO<sub>3</sub>:

OH 
$$+$$
 HNO<sub>3</sub>  $\xrightarrow{R.T.}$  OH  $+$  H2O  $+$ 

b) Reaction with cone, nitric acid (nitrating mixture):

 $HNO_3 + 2H_2SO_4 \longrightarrow 2HSO_4 + H_3O_7 + NO_2$ 

- 3. Sui phonation at different temperature:
  - a) Reaction with cone. H<sub>2</sub>SO<sub>4</sub>, at R.T./300K:

phenol

ortho phenol sulphonic acid

b) Reaction with cone. H<sub>2</sub>SO<sub>4</sub> at high temperature (373K):

4. Kolbe's Schmidt reaction

sodium phenoxide

5. Reimer and Tiemann reaction:

OH OH CHO 
$$+$$
 CHCl<sub>3</sub> + 3 NaOH  $\xrightarrow{333\,\mathrm{k}}$  OH CHO  $+$  3NaCl + 2H<sub>2</sub>O

6. Reaction with Zn metal

$$\begin{array}{c}
OH \\
\hline
O + Zn \longrightarrow \hline
O + ZnO
\end{array}$$

Phenol

7. Oxidation of Phenol: When exposed to air or heating with 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>, it undergoes slow oxidation and turn pink to form phenoquinone or p-benzoquinone or conjugated diketone

$$OH \qquad O$$

$$+ O_2 \xrightarrow{air} O$$

$$+ H_2O$$

8. Colour test of phenols: In fact all compound containing enolic group (C = C-OH) gives FeCl<sub>3</sub> test, forming chelate compound. Phenol gives ferric phenoxide as chelate compound.

$$6 C_6 H_5 OH + FeCl_3 \longrightarrow [Fe (C_6 H_5 O)_6]^{3-} + 3H^{+} + 3 HCl$$



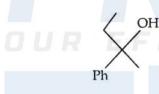
## **MULTIPLE CHOICE QUESTIONS**

# INTRODUCTION, NOMENCLATURE, CLASSIFICATION ISOMERISM

1. IUPAC name of following compound is



- a) pent-3-en-2-ol
- b) pent 2-en-4-o1
- c) but-3-en-2-ol
- d) but-2-en-4-o1
- 2. How many isomers of C<sub>5</sub>H<sub>11</sub> OH will be 1° alcohols?
  - a) Five
- b) Four
- c) Two
- d) Seven
- 3. Which of the following is trihydric alcohol?
  - a) Glycine
- b) Glycerol
- c) Glycol
- d) 2-heptanol
- 4. In glycerine,
  - a) one 1° OH group is present
  - b) one 2° OH group is present
  - c) two 2° OH groups are present
  - d) one 3° OH group is present
- 5. Correct IUPAC name of the following compound is



- a) 3-phenyl butan-3-o1
- b) 2-phenyl propan-2-o1
- c) 2-phenyl butan-2-o1
- d) 3-phenyl butan-2-o1
- 6. General representation of primary alcohol is
  - a) > C = O
- b) -CH2OH
- c) >CHOH
- d) ) COH
- 7. The general formula, which represent the homologous series of alcohol is
  - a) C<sub>n</sub>H<sub>2n</sub>O
- b)  $C_{n}H_{2n+1}O$
- c)  $C_n H_{2n} O_2$
- d)  $C_n H_{2n+2} O$
- 8. Grain spirit is
  - a) isopropyl alcohol
- b) isobutyl alcohol
- c) methyl alcohol
- d) ethyl alcohol
- 9. Which of the following has primary alocholic group?

- a) \rightarrow\_OH
- b) OH
- c) OH
- d) +ОН
- 10. The characteristic grouping of secondary alcohol is
  - a) > C = O
- b) -CH,OH
- c) >CHOH
- d) COH
- 11. Which of the following alcohol contain vinyl group
  - a) //OH







- 12. How many ethers are possible for formula  $C_4H_{10}O$ ?
  - a) 2
- b) 3
- c) 4
- d) 5
- 13. In allylic alcohol OH group is attached to
  - a) sp hybridised carbon atom
  - b) sp<sup>2</sup> hybridised carbon atom
  - c) sp<sup>3</sup> hybridised carbon atom
  - d) sp<sup>3</sup>–d–hybridised carbon atom
- 14. Which of the following is allylic alcohol







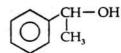


- 15. Butane -2 o 1 is
  - a) primary alcohol
- b) secondary alcohol
- c) tertiary alcohol
- d) aldehyde
- 16. Allylic allohols may be
  - a) 1° types
- b) 2° types
- c) 3° types
- d)  $1^{\circ}$ ,  $2^{\circ}$ ,  $3^{\circ}$  types
- 17. Which is a primary alcohol?
  - a) Butan-2-o1
- b) Butan-1-o1

- c) Propan–2–ol
- d) Isopropyl alcohol
- 18. Aralkyl alcohols are also named as a) allylic alcohol
  - b) vinylic alcohol
  - c) benzylic alcohol
- d) aryl alcohol
- 19. IUPAC name of the compound
  - CH,CH(C,H,)CH,CH(OH)CH, is
  - a) 2-methyl 3-hexanol
  - b) 4-methyl 2-hexanol
  - c) heptanol
  - d) all of these
- 20. Following alcohol can be named as



- a) 1-phenyl 1-methyl propan-2-ol
- b) 3-phenyl butan-2-ol
- c) 1-methyl 1-phenyl butan-2-vol
- d) 3-phenyl 3-methyl propan-2-ol
- 21. Wood alcohol is
  - a) phenol
- b) CH,OH
- c) C<sub>2</sub>H<sub>5</sub>OH
- d) CH<sub>2</sub>COOH
- 22. In vinyl alcohol OH group is attached to
  - a) sp hybridised carbon atom
  - b) sp<sup>2</sup> hybridised carbon atom
  - c) sp<sup>3</sup> hybridised carbon atom
  - d) sp<sup>2</sup> d–hybridised carbon atom
- 23. How many metamers are possible for  $C_4H_{10}O$ ?
  - a) 1
- c) 3
- d) 4
- 24. IUPAC name of the following compound is



- a) 1-rnethyl 1-phenyl methanol
- b) 2-methyl benzyl alcohol
- c) 2-phenyl ethan-2-ol
- d) 1-phenyl ethan-1-ol
- 25. Ethyl methyl carbinol is,
  - a) n-butyl alcohol
- b) t-butyl alcohol
- c) sec. butyl alcohol
  - d) isobutyl alcohol
- 26. Following alcohol is the type of



- a) allylic alcohol
- b) vinylic alcohol
- c) aralkyl alcohol
- d) benzylic alcohol
- 27. Methyl carbinol is
  - a) ethanol
- b) 2-propanol
- c) 1-propanol
- d) methanol
- 28. Vinyl carbinol is
  - a) HO-CH,-CH = CH,
  - b)  $CH_3 C = CH_3$ ÓН
  - c)  $CH_3$ –CH = CH OH
  - d)  $CH_{2} = CH OH$
- 29. C-OH group is present in
  - a) 1° R-OH
- b) 2° R-OH
- c) 3° R-OH
- d) 4° R-OH
- 30. Ethanol containing some methanol is called as
  - a) methylated spirit
- b) rectified spirit
- c) absolute spirit
- d) proof spirit
- 31. IUPAC name of secondary butyl alcohol is
  - a) 2-methyl propan-I-ol
  - b) bu tan-2-ol
  - c) 2-methyl prcpan-2-vol
  - d) 1-butanol
- 32. IUPAC name of the following compound is



- a) cyclohexa 1-3-dien-3-o1
- b) cyclohexa 1, 2-dien-I-ol
- c) cyclohexa 1, 3-dien-3-o1
- d) cyclohexa 1, 5-dien-1-ol
- 33. IUPAC name of following compoud is

$$\begin{array}{c} \operatorname{CH_3} \\ | \\ \operatorname{CH_3} - \operatorname{CH_2} - \operatorname{CH} - \operatorname{CH_2} \operatorname{CH_2} \operatorname{OH} \end{array}$$

- a) 5-ethyl 3-methyl pent-1-ol
- b) 3-methyl pent-1-ol
- c) 3-ethyl pent-1-ol
- d) 3, 5-diethyl pent-1-ol
- 34. IUPAC name of the following compound is

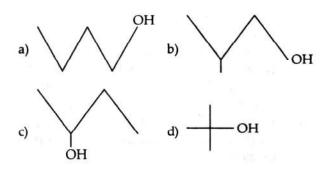


a) 2-chloro cyclohex-1-en-1 -ol

- b) 1-chloro 2-hydroxy cyclohex-2-en
- c) 6-chloro cyclohex-l-en-1-ol
- d) 2-chloro 1-hydroxy cyclohex-1-en
- 35. IUPAC name of t-butyl alochol is
  - a) 2-methyl propan-1-ol
  - b) 2-methyl butan-1-ol
  - c) 2-methyl propan-2-ol
  - d) 1-methyl propan-2-ol
- 36. IUPAC name of following compound



- a) 2, 5-dimethyl phenol
- b) 3–5–dimethyl phenol
- c) para-dimethyl phenol
- d) dimethylene phenol
- 37. The compound which is not isomeric with diethyl ether is
  - a) n-propyl methyl ether
  - b) butan-1-ol
  - c) 2-methyl propan-2-o1
  - d) butanone
- 38. How many compounds show optical isomerism of molecular formula  $C_5H_{12}O$ ?
  - a) 2
- b) 3
- c) 4
- d) 5
- 39. Molecular formula  $C_2H_6O$  represents
  - a) alcohols and acids
  - b) alcohols and ethers
  - c) only alcohols
  - d) only ethers
- 40. Which isomer of  $C_4H_{10}O$  is optically active?



- 41. Molecular formula C<sub>3</sub>H<sub>5</sub>O shows
  - a) chain and optical isomers
  - b) position and functional isomers

- c) functional and metamers
- d) chain and position isomers
- 42. IUPAC name of following compound is



- a) pent-2-yn-5-ol
- b) pent-3-yn-1 -ol
- c) pentane-2-yn-5-ol
- d) pentane-3-yn-1-o1
- 43. Alcohols exhibit
  - a) chain isomerism
- b) position isomerism
- c) optical isomerism
- d) all of these
- 44. Total number of isomers including structural and stereo isomers of molecular formula  $C_4H_{10}O$ 
  - a) 4

b) 7

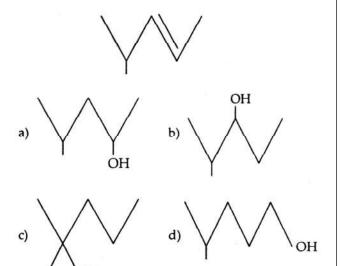
c) 8

- d) 10
- 45. What type of isomerism is exhibited by pentanol?
  - a) Position
- b) Chain
- c) Optical
- d) All of these
- 46. Number of isomeric ethers with molecular formula  $C_5H_{12}O$  are
  - a) 4
- b) 6
- c) 8
- d) 10
- 47.  $C_4H_{10}O$  represents
  - a) aldehydes
- b) ketones
- c) alcohols
- d) acids
- 48. Which of the following are functional isomers?
  - a) Alcohols and ethers
  - b) Alcohols and acids
  - c) Alcohols and aldehydes
  - d) Alcohols and ketones
- 49. How many 3° alcohols are possible for C<sub>3</sub>H<sub>5</sub>O?
  - a) 0
- b) 1
- c) 2
- d) 3

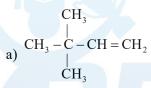
#### PREPARATION METHODS

- 50. Hydration of alkene produces
  - a) 1° alcohols or 2° alcohols
  - b) 1° alcohols or 3° alcohols
  - c) 1, 2° and 3° alcohols
  - d)  $1^{\circ}$  or  $2^{\circ}$  or  $3^{\circ}$  alcohols
- 51. Hydration:
  - a) takes place through carbocation
  - b) is a reversible process
  - c) follows electrophilic addition

- d) follows all of the above
- 52. Hydration of following compound gives



53. Which is hydrated to a maximum extent?



- b)  $CH_3$ – $CH = CH_2$
- c)  $CH_2 = CH_2$
- d) CH = CH
- 54. Which reagent can bring about following conversion,



- a) CrO<sub>3</sub>
- b) acidic KMnO<sub>4</sub>
- c) dil. H<sub>2</sub>SO<sub>4</sub>
- d) dil. KOH
- 55. Hydroboration oxidation of alkene give
  - a) acids
- b) aldehydes
- c) ketones
- d) alcohols
- 56. In hydration of alkene first step is
  - a) nucleophilic attack of water on carbocation
  - b) deprotonation of carbocation
  - c) protonation of alkene
  - d) attack of H<sub>2</sub>O+
- 57. In the following reaction most stable intermediate is

$$CH_3$$
 $|$ 
 $CH_3CHCH = CH_2 + H_2O \xrightarrow{H^+}$ 

a) 
$$CH_3$$
  $CH_3$   $CH_3$   $CH_3$   $CH_3$   $CH_3$   $CH_4$   $CH_5$   $CH_5$ 

$$\begin{array}{ccc} \text{CH}_3 & \text{CH}_3 \\ | & | \\ \text{c) CH}_3 \underset{\oplus}{\text{CCH}}_2 \text{CH}_3 & \text{d) CH}_3 \text{CHCH}_2 \overset{\Theta}{\text{CH}}_2 \end{array}$$

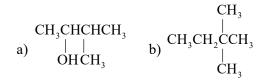
- 58. In the hydration of an alkene carbocation is formed from:
  - a) carbanion
- b) oxonium ion
- c) hydroxide ion
- d) hydride ion
- 59. Reaction intermediate in hydration of alkene is
  - a) carbanion
- b) carbon free radicals
- c) carbocation
- d) carbene

60. 
$$\bigcirc$$
OH  $\stackrel{B}{\longleftarrow}$ O $\stackrel{A}{\longrightarrow}$ OF

In above reaction A and B respectively

- a) H<sub>2</sub>/Pt, LiAlH<sub>4</sub>/H<sub>2</sub>O
- b) H<sub>2</sub>/Pt, H<sub>2</sub>/Pt
- c) LiAlH<sub>4</sub>/H<sub>2</sub>O, LiAlH<sub>4</sub>/H<sub>2</sub>O
- d) LiAlH<sub>4</sub>/H<sub>2</sub>O, H<sub>2</sub>/Pt
- 61. In hydroboration oxidation of alkene, the initial attack is
  - a) boron
- b) NaOH
- c) H<sub>2</sub>O<sub>2</sub>
- d) H<sup>+</sup>
- 62. Select incorrect statement about hydroboration oxidation:
  - a) Addition is against Markownikoff rule
  - b) Intermediate is a carbocation
  - c) It does not involve rearrangement
  - d) It is a addition reaction
- 63. Reagents used in hydroboraion oxidation reaction
  - a)  $B_2H_6 + CrO_3 + NaOH$
  - b)  $B_2H_6 + H_2O_2 + NaOH$ , THF
  - c) B<sub>2</sub>H<sub>6</sub> + acidic KMnO<sub>4</sub> + NaOH
  - d)  $B_2H_6$  + dil. HNO<sub>3</sub> + NaOH
- 64. A changes to ...... with hydroboration-oxidation

$$CH_3 \\ | \\ A: CH_3CH = CCH_3$$



## CH<sub>3</sub>CH<sub>2</sub>CHCH<sub>2</sub>OH

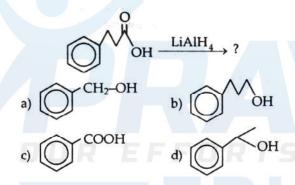
- c) CH,
- d) none of these
- 65. Alcohol containing least number of carbon which can be prepared using Grignard reagent is:
  - a) (CH<sub>3</sub>),CHOH
- b) (CH<sub>3</sub>)<sub>3</sub>COH
- c) CH<sub>3</sub>OH
- d) CH,CH,OH
- 66. Following alkene will give same product by any method out of hydration, hydroboration oxidation:
  - a)  $CH_3CH = CH_2$
- b)  $CH_3CH = CHCH_3$
- $CH_{3}CHCH = CH_{2}$   $CH_{3}$ 
  - $^{2}$  d)  $\bigcirc$  CH<sub>2</sub>
- 67. Find out (B) in the following reaction

$$C_3H_5 + Br_2 \xrightarrow{AlBr_3} (A) \xrightarrow{Aq. KOH} (B)$$

- a)  $CH_3$ – $CH = CH_2$
- b) CH<sub>3</sub>-CH<sub>2</sub>-CH<sub>2</sub>-OH
- c) CH<sub>3</sub>-CHOH-CH<sub>3</sub>
- d) CH<sub>3</sub>-O-C<sub>2</sub>H<sub>5</sub>
- 68. Which of the following reactions will yield 2–propanol?
  - 1)  $CH_2 = CH CH_3 + H_2O \xrightarrow{H^+}$
  - 2)  $CH_3CHO \xrightarrow{CH_3MgI} H_3O^+$
  - 3) HCHO  $\xrightarrow{C_2H_5MgI}$
  - 4)  $CH_3COCH_3 \xrightarrow{CH_3MgI}$
  - a) 1 and 2
- b) 2 and 3
- c) 3 and 1
- d) 2 and 4
- 69. Treatment of 1-butene with cone. H<sub>2</sub>SO<sub>4</sub> followed by treatment with water forms
  - a) 1-butanol
- b) 2-butanol
- c) 2-propanol
- d) 1–2–propan–diol
- 70. Acetone is treated with sodium amalgam and water gives,
  - a) (CH<sub>3</sub>)<sub>2</sub>CHOH
- b) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>OH
- c) C<sub>2</sub>H<sub>5</sub>OH
- d) CH, CH, COOH
- 71. 2-methy 12-pentanol is prepared from acetone and what?
  - a) C<sub>2</sub>H<sub>5</sub>MgI
  - b) (CH<sub>3</sub>)<sub>3</sub>CMgI
  - c) 2-molecules of CH<sub>2</sub>MgI
  - d) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>MgI

- 72. The only primary alcohol that can be prepared by the indirect hydration of alkene is
  - a) ethyl alcohol
- b) n-propyl alcohol
- c) isobutyl alcohol
- d) methyl alcohol
- 73.  $R_2CO \longrightarrow R_2CHOH$ . The conversion is,
  - a) reduction
- b) oxidation
- c) hydrolysis
- d) hydration
- 74. Propene can be converted into 2–propanol by hydration. Which of the following reagents is ideal to affect the conversion?
  - a) Alkaline KMnO<sub>4</sub>
  - b) Zn dust + H<sub>2</sub>O
  - c) Conc. H<sub>2</sub>SO<sub>4</sub>
  - d) Conc. HCl
- 75. Using CH<sub>3</sub>MgBr, which substrate would lead to (CH<sub>3</sub>)<sub>3</sub>COH?
  - a) Acetone
  - b) Acetyl chloride
  - c) Acetaldehyde
  - d) Isopropyl alcohol
- 76. Which of the following is IUPAC name of the compound formed from reduction of 2-butanone?
  - a) 1-butanol
- b) 2-butanol
- c) 1-butanal
- d) 2-butanal
- 77. Alcohols can be prepared by hydration of,
  - a) alkanes
- b) alkyl halides
- c) alkyl amines
- d) alkenes
- 78. CH<sub>3</sub>-CH<sub>2</sub>-CH<sub>2</sub>-CO-CH<sub>2</sub>-CH(CH<sub>3</sub>)<sub>2</sub>
  Catalytic hydrogenation of above compound in the presence of nickel catalyst gives
  - a) an optically inactive compound
  - b) an optically active compound
  - c) compound with plane of symmetry
  - d) an tertiary alcohol
- 79. Alkenes convert into alcohols by
  - a) hydrolysis by dill. H<sub>2</sub>SO<sub>4</sub>
  - b) hydration of alkene by cone. H<sub>2</sub>SO<sub>4</sub>
  - c) hydrolysis by water vapours and conc. H<sub>2</sub>SO<sub>4</sub>
  - d) hydration of alkene by aqueous KOH
- 80. Acid catalysed hydration of alkenes except ethene leads to the formation of
  - a) secondary or tertiary alcohol
  - b) primary alcohol
  - c) mixture of secondary and tertiary alcohols
  - d) mixture of primary and secondary alcohols
- 81. Following compound on reduction gives eft

- Which of the following is best reducing agent to convert - COOH to - CH, - OH
  - a) Fe + cone. HCl
- b) LiAlH<sub>4</sub>
- c) NaBH<sub>4</sub>
- d) Zn.Hg + HC1
- 83. Methanol is obtained by reduction of
  - a) CH<sub>3</sub>-CHO
- b) CH<sub>2</sub>-COOH
- c) H-CONH,
- d) H-COOH
- 84. Product of the following reaction will be



Ethanol is obtained by in which of the following reaction

1. 
$$\frac{\text{i) conc. H}_2\text{SO}_4}{\text{ii) H}_2\text{O}}$$

2. 
$$\bigvee_{H}^{O} + CH_3 - MgX \xrightarrow{dry \text{ ether} \atop H_3O +}$$

3. 
$$H \mapsto CH_3 - MgX \xrightarrow{dry \text{ ether}} H_3O + H_3O +$$

- $\xrightarrow{\text{LiAlH}_4}$
- a) 1, 3
- b) 1, 2, 4
- c) 3, 4
- d) 1, 3, 4
- 86. Which of the following is best reagent used for

following conversion?

$$\bigvee_{H} \xrightarrow{?} \bigvee_{OH}$$

- a) H2/Ni
- b) H,O+
- c) NaBH<sub>4</sub>
- d) Zn.Hg + HCl
- 87. Which one of the following will yield propan–z– ol, choose the right answer from a to d?

1) 
$$M_2O/H^+$$

2) 
$$\bigvee_{O}^{H} \xrightarrow{i) CH_3 MgX} ii) H_2O / H^+$$

3) 
$$CH_2O \xrightarrow{i) C_2H_5MgX}$$
  
 $ii) H_3O^+$ 

4) 
$$\frac{\text{Neutral}}{\text{KMnO}_4}$$

- a) 3, 4
- b) 2, 3
- c) 1, 3
- d) 1, 2
- When C<sub>2</sub>H<sub>5</sub>MgI react with acetone and the addition product is hydrolysed we get
  - a) 1° alcohol
- b) 2° alcohol
- c) 3° alcohol
- d) an aldehyde
- Identity (Z) in the following series of reaction.

$$\bigcap_{OH} \xrightarrow{PCl_5} (X) \xrightarrow{alc.KOH} (Y)$$



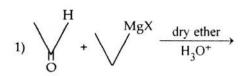
- 90. Benzyl alcohol is obtained by reduction of
  - a) benzoic acid
- b) acetophenone
- c) benzonitrile
- d) benzamide
- 91. To prepare butan-2-o1 from methyl magnesium iodide. The compound required is







92. Isobutyl alcohol can not be obtained by the reaction between



2) 
$$H + CH_3 - MgX \xrightarrow{dry \text{ ether}} H_3O^+$$

3) 
$$H \downarrow O H + \downarrow O H AgX \xrightarrow{dry ether} H_3O^+$$

4) 
$$V_{O} + CH_{3} - MgX \xrightarrow{dry \text{ ether}} H_{3}O^{+}$$

- a) 1, 2, 3
- b) 1, 2, 4
- c) 1, 3, 4
- d) 2, 3, 4
- 93. Which set of following reagent would you select to convert C<sub>6</sub>H<sub>5</sub>COCH<sub>3</sub> to following alcohol

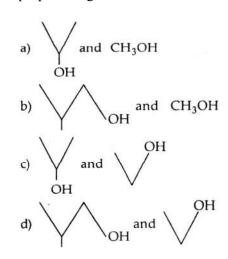


a)  $CH_3 - CH_2 - CH_2$  MgBr and hydrolysis

$$\begin{array}{c} \operatorname{CH_3} - \operatorname{CH} - \operatorname{CH_3}, \operatorname{AlCl_3} \\ \operatorname{b)} & \stackrel{|}{\operatorname{Br}} \end{array}$$

- c) (CH<sub>3</sub>)<sub>2</sub> CHMgBr and acid hydrolysis
- d) CH<sub>3</sub> CHOH CH<sub>3</sub>, Zn
- 94. Propan–1–ol may be prepared by the reaction of propene with
  - a) H<sub>3</sub>BO<sub>3</sub>
  - b)  $H_2SO_4RT / H_2O$
  - c)  $B_2H_6/$  THF,  $H_2O_2$  and NaOH
  - d) (CH<sub>3</sub>COO)<sub>2</sub> Hg/NaBH<sub>4</sub>
- 95. Styrene on hydroboration oxidation gives
  - a) 2-phenyl ethan-1-ol
  - b) I-phenyl ethan-1-ol
  - c) benzoic acid
  - d) benzaldehyde
- 96. Which of the following produces only one product on reduction with LiAIH<sub>4</sub>

- a) CH, COO C,H,
- b) C,H,COOC,H,
- c) C<sub>2</sub>H<sub>5</sub>COOCH<sub>3</sub>
- d) CH<sub>3</sub>COOCH<sub>2</sub>-CH<sub>2</sub>-CH<sub>3</sub>
- 97. Catalytic hydrogenation of methyl 2–methyl propanoate gives



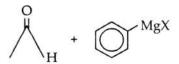
- 98. CH<sub>3</sub>-CH-C-O CH<sub>3</sub>, on reduction give,
  - a) l-phenyl propan-1-ol and methanol
  - b) 2-phenyl propan-1-ol and ethanol
  - c) 2-phenyl propan-1-ol and methanol
  - d) 2-phenyl propan-2-ol and methanol
- 99. Aldehydes and ketone reacts with Grignards reagent gives
  - a) mixture of 1°, 2°, 3° alcohols
  - b) 1° or 2° or 3° alcohols
  - c) 1° or 2° alcohols
  - d) 2° or 3° alcohols
- 100. Benzaldehyde and phenyl magnesium halide gives
  - a) Benzyl alcohol
- b) diphenyl ketone
- c) diphenyl ethanol
- d) diphenyl methanol
- 101. Which reagent can bring about

R-COOH  $\longrightarrow$  R-CH,-OH

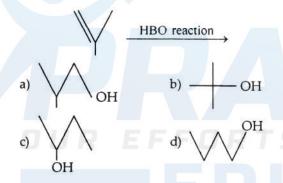
- a) Sn + HCl
- b) Na +  $C_2H_5OH$
- c) H<sub>2</sub> + Pt
- d) LiAlH<sub>4</sub>
- 102. When wine is put in air, it become sour due to
  - a) oxidation of C<sub>2</sub>H<sub>5</sub>-OH
  - b) reduction of C<sub>2</sub>H<sub>5</sub>-OH

- c) formation of C<sub>2</sub>H<sub>5</sub>-COOH
- d) dissolution of CO,
- 103. Reaction used to convert acid to 10 alcohol is
  - a) oxidation
- b) reduction
- c) polymerisation
- d) pyrolysis
- 104. Carboxylic acid on reduction produces
  - a) 1° alcohols
- b) 2° alcohols

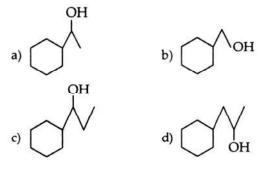
- c) aldehydes
- d) ketones
- 105. Product of the following reaction is



- a) 2-phenyl ethanol
- b) 1-phenyl ethanol
- c) phenyl methanol
- d) methyl phenyl ketone
- 106. Ethene is subjected to hydroboration oxidation reaction followed by treatment with PCC gives
  - a) ethyl alcohol
- b) acetaldehyde
- c) acetic acid
- d) acetone
- 107. Product of the following reaction is

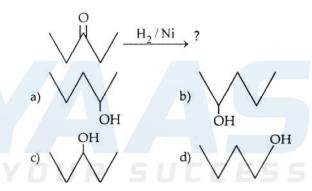


- 108. –CH<sub>2</sub>OH group is obtained by reduction of
  - a) R-CN
- b) R-NO<sub>2</sub>
- c) R-CO-R
- d) R-COOH
- 109. Cyclohexane carbaldehyde is reacted with ethyl magnesium halide in the presence of dry ether and product on acid hydrolysis gives

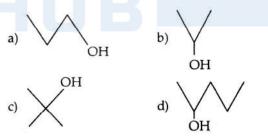


110. Product of the following reaction is

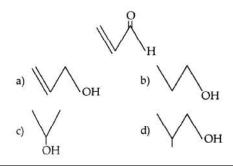
- 111. Hydroboration oxidation of 3–methyl but–1–ene gives
  - a) 3-methyl butan-2-ol
  - b) 2-methyl butan-2-ol
  - c) 3-methyl butan-l-ol
  - d) 2-methyl butan-l-ol
- 112. Product of the following reaction.



113. Acrolin on reduction by using H<sub>2</sub> / Ni gives

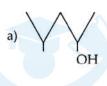


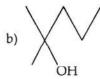
114. Following compound is reduced by NaBH<sub>4</sub> or LiAIH<sub>4</sub> gives



- 115. 2,2–dimethyl propan–1–ol is obtained from 2,2–dimethyl but–1–ene by using
  - a) dil. H<sub>2</sub>SO<sub>4</sub>
  - b) B<sub>2</sub>H<sub>6</sub>, H<sub>2</sub>O<sub>2</sub> and NaOH
  - c) (CH<sub>3</sub>COO)<sub>2</sub>Hg + NaBH<sub>4</sub>
  - d) H<sub>2</sub>/Ni
- 116. Benzaldehyde on reduction by using NaHg+H<sub>2</sub>O gives
  - a) benzyl alcohol
  - b) phenol
  - c) sodium benzoate
  - d) sodium phenoxide
- 117. following reaction is

$$\bigvee \sqrt{ \frac{B_2 H_6 / THF}{H_2 O_2 + OH^-}} ?$$

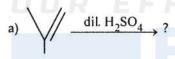






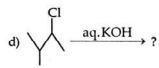


118. In which of the following reaction carbocation is not formed?

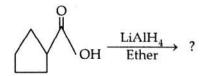




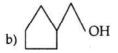


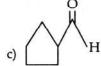


119. Product of the following reaction is



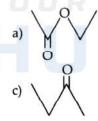


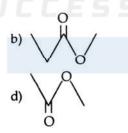






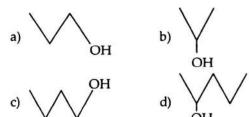
- 120. Alcohols are obtained by reduction of
  - 1. R-CHO
- 2. R-COR
- 3. R-COOH
- 4. RCOOR
- a) 1, 3
- b) 1, 3
- c) 3, 4
- d) 1, 2, 3, 4
- 121. Ester on reduction by using LiAlH<sub>4</sub> produces.
  - a) single aldehyde
  - b) single alcohol
  - c) mixture of aldehyde
  - d) mixture of alcohol
- 122. Ester are converted in to mixture of alcohol by
  - a) acid hydrolysis
  - b) alkaline hydrolysis
  - c) catalytic hydrogenation
  - d) oxidation
- 123. Ethyl formate on catalytic hydrogenation gives
  - a) CH<sub>3</sub>-OH
- b) CH<sub>3</sub>OH and C<sub>2</sub>H<sub>5</sub>OH
- c) C<sub>2</sub>H<sub>5</sub>-OH
- d) H-CHO and CH<sub>2</sub>-OH
- 124. Mixture of methanol and ethanol is obtained from catalytic hydrogenation of





- 125. 3-methyl but-1-ene on HBO reaction gives
  - a) 3-methyl butan-2-ol
  - b) 2-methyl butan-2-ol
  - c) 3-methyl butan-1-ol
  - d) 2-methyl butan-1-ol
- 126. HBO of but-2-ene produces
  - a) butan-1-ol
  - b) butan-2-ol
  - c) 2-methyl propan-2-ol
  - d) 2-methyl propan-1-ol
- 127. Reduction of aldehydes and ketones produces
  - a) 1º alcohols
- b) 2° alcohols

- c) 3º alcohols
- d) 1º or 2º alcohols
- 128. Hydroboration oxidation of propene produces



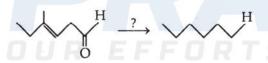
- 129. Molar concentration of alkene used up in hydrobonation oxidation reaction to form alcohol is
  - a) 2
- b) 4
- c) 3
- d) 5

#### PROPERTIES AND USES

- 130. Consider the following species
  - 1)  $RCH + CH_3$
- 2) RCH,CH,+
- 3) RCH<sub>2</sub>CH<sub>2</sub>O<sup>+</sup>H<sub>2</sub>

In the dehydration of straight chain 1° alcohols, the correct sequence of formation of the species involved is

- a) 2, 1
- b) 1, 2
- c) 3, 2
- d) 2, 3
- 131. Which of the following reducing agent is used for following conversion?



- a)  $H_2 + Ni$
- b) LiAlH,
- c) NaBH<sub>4</sub>
- d) Na.Hg + H<sub>2</sub>O
- 132. The correct order of increasing boiling points is
  - a) n-butane < 1-butanol < n-butyl chloride < isobutane
  - b) n-butane < isobutane < n-butyl chloride < 1-butanol
  - c) isobutane < n-butyl chloride < n-butane < 1-butanol
  - d) isobutane < n-butane < n-butyl chloride < 1-butanol.
- 133. List the class of alcohols in decreasing order of reactivity towards HX
  - a)  $3^{\circ} > 1^{\circ} > 2^{\circ} > \text{MeOH}$
  - b)  $3^{\circ} > 2^{\circ} > 1^{\circ} > \text{MeOH}$
  - c) MeOH>  $1^{\circ} > 2^{\circ} > 3^{\circ}$
  - d)  $2^{\circ} > 3^{\circ} > 1^{\circ} > \text{MeOH}$
- 134. List the hydrogen halide acids in decreasing order

of reactivity in the following reaction

$$R - OH + HX \longrightarrow RX + H,O$$

- a) HBr > HI> HCl > HF
- b) HI > HBr > HCl > HF
- c) HI > HF > HBr > HC1
- d) HI > HCl > HBr > HF
- 135. Which one of following is more reactive than the rest towards a Lucas reagent?
  - a) 1–butanol
- b) 2-butanol
- c) methanol
- d) 2-methyl 2-propanol
- 136. Sodium reacts with alcohol as given below

$$2R-OH + 2Na \longrightarrow 2R-ONa + H$$

Place the type of alcohol into decreasing order of reactivity towards sodium.

- a)  $3^{\circ} > 2^{\circ} > 1^{\circ}$
- b)  $1^{\circ} > 3^{\circ} > 2^{\circ}$
- c)  $2^{\circ} > 3^{\circ} > 1^{\circ}$
- d)  $1^{\circ} > 2^{\circ} > 3^{\circ}$
- 137. The main product of the reaction of  $(C_2H_5)_2$ CHCHOHCH<sub>3</sub> with conc.  $H_2SO_4$  is
  - a)  $(CH_3CH_2)_2CH-CH = CH_2$
  - b)  $CH_3-CH(C_3H_5)CH = CH-CH_3$
  - c)  $(C_2H_5)_2C = CH CH_3$
  - d) both 'a' and 'b'
- 138. Place the following alcohols in decreasing order of rate of dehydration with conc. H<sub>2</sub>SO<sub>4</sub>,
  - 1) CH,CH,CH(OH)CH,CH,CH,
  - 2)  $(CH_3)_2C(OH)CH_2CH_2CH_3$
  - 3) CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>CH<sub>4</sub>CH<sub>2</sub>OH
  - 4) (CH<sub>3</sub>)<sub>2</sub>C(OH)CH(CH<sub>3</sub>),
  - a) 4 > 2 > 1 > 3
- b) 1 > 2 > 3 > 4
- c) 4 > 3 > 2 > 1
- d) 4 > 3 > 1 > 2
- 139. An alcohol C<sub>4</sub>H<sub>9</sub>OH on dehydration gives an alkene, which on oxidation yield a acetone. The alcohol is
  - a) (CH<sub>3</sub>)<sub>3</sub>COH
  - b) CH<sub>3</sub>CH<sub>2</sub>CH(OH)CH<sub>3</sub>
  - c) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>OH
  - d) (CH<sub>3</sub>),CHCH<sub>2</sub>OH.
- 140. The most stable carbonium ion is
  - a) methyl carbonium ion
  - b) primary carbonium ion
  - c) secondary carbonium ion
  - d) tertiary carbonium ion.
- 141. The compound with highest boiling point is
  - a) CH<sub>4</sub>
- b) CH<sub>2</sub>OH
- c) CH<sub>2</sub>Cl
- d) CH<sub>2</sub>Br

- 142. Hydrogen bonding is maximum in
  - a) ethanol
- b) diethyl ether
- c) ethyl chloride
- d) triethylamine.
- 143. 1-butanol is treated with PCC gives,
  - a) CH, CH, COOH
  - b) CH, CH, CH, CHO
  - c) CH, COCH, CH,
  - d) CH<sub>2</sub>COOH
- 144. Lucas reagent is used to distinguish among primary, secondary and tertiary
  - a) alkyl halides
- b) alcohols
- c) aliphatic amines
- d) aromatic amines.
- 145. The compound which reacts faster with Lucas reagent at room temperature is
  - a) butan-1-ol
  - b) butan-2-ol
  - c) 2-methyl propan-1-ol
  - d) 2-methyl propan-2-ol
- 146. t-butyl alcohol is heated with Al<sub>2</sub>O<sub>3</sub> gives
  - a)  $CH_3CH = CHCH_3$  b)  $CH_3CH_3CH = CH_3$
  - c)  $(CH_{2})_{2}C = CH_{2}$ 
    - d) all of these
- 147. The compound which gives the most stable carbonium ion on dehydration is
  - a) CH<sub>3</sub>-CH(CH<sub>3</sub>)-CH<sub>2</sub>OH
  - b)  $(CH_3)_2C OH$
  - c) CH<sub>3</sub>-CH<sub>2</sub>-CH<sub>2</sub>-CH<sub>2</sub>OH
  - d) CH<sub>2</sub>CH(OH)CH<sub>2</sub>CH<sub>2</sub>
- 148. Maximum number of active hydrogens are present in
  - a) ethanoic acid
- b) ethyl alcohol
- c) ethylene glycol
- d) glycerol
- 149. When t-butyl alcohol is heated with Cu at 573 K, it forms
  - a) butanal
- b) prop anal
- c) ethyl methyl ketone d) 2–methyl prop–1–ene
- 150. 1-butanol is oxidised by acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>2</sub> give,
  - a) butanal
- b) butanoic acid
- c) butene
- d) butane
- 151. Alcohols gives alkyl halides, treatment with
  - a) PX<sub>3</sub>
- b) PX<sub>5</sub>
- c) HX
- d) all of these
- 152. When 2—methyl 2—butanol is a dehydrated to give an alkene, the preferred product is
  - a) 2-methyl 2-butene
  - b) 2-methyl 1-butene
  - c) 2-methyl 1-propene
  - d) n-hexene

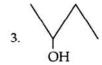
- 153. Consider the following compounds
  - 1) CH,CH,CH,CH,OH
  - 2) CH<sub>3</sub>CH<sub>2</sub>CHOHCH<sub>3</sub>
  - 3)  $(CH_3)_2COH$

These compounds are dehydrated by treatment with H<sub>2</sub>SO<sub>4</sub>. The correct sequence of increasing order of the reactivity of three compounds towards dehydration is

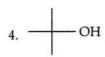
- a) 3 < 1 < 2
- b) 1 < 2 < 3
- c) 2 < 1 < 3
- d) 1 < 3 < 2
- 154. Which of the following alkene on acid catalysed hydration form propan–2–ol,
  - a)  $CH_3CH = CH_3$
- b)  $(CH_3)_{,}C = CH_{,}$
- c)  $CH_3CH = CHCH_3$  d)  $CH_3CH_2CH = CH_2$
- 155. In isomeric alcohols correct order of boiling point is,
  - a)  $2^{\circ} > 3^{\circ} > 1^{\circ}$
- b)  $3^{\circ} > 2^{\circ} > 1^{\circ}$
- c)  $1^{\circ} > 2^{\circ} > 3^{\circ}$
- d)  $2^{\circ} > 1^{\circ} > 3^{\circ}$
- 156. Which one of the following compound would not be oxidised by acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>?
  - a) CH<sub>3</sub>OH
- b) (CH<sub>3</sub>),CHOH
- c) (CH<sub>2</sub>),COH
- d) C<sub>2</sub>H<sub>5</sub>OH
- 157. Which of the following is expected to have highest boiling point?
  - a) (CH<sub>3</sub>)<sub>2</sub>CHCl
- b) (CH<sub>3</sub>)<sub>2</sub>CHOH
- c) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>OH
- d) CH,CH,CH,C1
- 158. Final product by the treatment of isobutyl alcohol with alumina is,
  - a) 2-methyl propene
  - b) 2-methyl but-l-ene
  - c) ethyl t-butyl ether
  - d) acetone and acetic acid
- 159. Ethyl alcohol is heated with SOCl, gives,
  - a)  $C_2H_5Cl + HCl$
  - b)  $C_2H_5Cl + SO_2$
  - c)  $C_2H_2Cl + HCl + SO_2$
  - d)  $CH_3OH + CH_3Cl + SO_2$
- 160. Isopropyl alcohol is oxidised by CrO<sub>3</sub> gives,
  - a) CH<sub>2</sub>COCH<sub>2</sub>
- b) CH<sub>2</sub>COOH
- c)  $CH_3CH = CH_2$
- d) CH<sub>3</sub>CHO
- 161. Which of the following is most acidic?
  - a) H,O
- b) CH<sub>3</sub>OCH<sub>3</sub>
- c) CH<sub>3</sub>OH
- d) C<sub>6</sub>H<sub>5</sub>OH
- 162. Which of the following is oxidised to form ethyl methyl ketone?
  - a) 2-propanol
- b) 2-butanol
- c) 1-butanol
- d) 1 -propanol

- 163. Which of the following alcohol is least soluble in water?
  - a) CH<sub>2</sub>OH
- b) C<sub>2</sub>H<sub>7</sub>OH
- c) C<sub>6</sub>H<sub>13</sub>OH
- d)  $C_{10}H_{21}OH$
- 164. Which of the following is more acidic alcohol?
  - a) C<sub>2</sub>H<sub>5</sub>OH
  - b) CH<sub>3</sub> CH CH<sub>3</sub> | OH
  - c) CH<sub>3</sub>-CH<sub>2</sub>-CH<sub>2</sub>-OH
  - d) (CH<sub>3</sub>)<sub>3</sub>-OH
- 165. Decreasing order of acidity of following alcohol is









- a) 1 > 2 > 4 > 3
- b) 3 > 1 > 4 > 2
- c) 2 > 3 > 1 > 4
- d) 2 > 1 > 3 > 4
- 166. Which of the following is most acidic
  - a) H<sub>2</sub>O
- b) CH<sub>3</sub>-OH
- c) C<sub>2</sub>H<sub>5</sub>-OH
- d)  $C_3H_7$ -OH
- 167. Which of the following is strong base
  - a) CH<sub>3</sub>ONa
- b) NaOH
- c) KOH
- d) Na<sub>2</sub>CO<sub>3</sub>
- 168. Reactivity of alcohol in breaking O-H bond is
  - a)  $1^{\circ} > 2^{\circ} > 3^{\circ}$
- b)  $3^{\circ} > 2^{\circ} > 1^{\circ}$
- c)  $2^{\circ} > 1^{\circ} > 3^{\circ}$
- d)  $1^{\circ} > 3^{\circ} > 2^{\circ}$
- 169. Reactivity of alcohol in breaking of C–O bond is
  - a)  $1^{\circ} > 2^{\circ} > 3^{\circ}$
- b)  $3^{\circ} > 2^{\circ} > 1^{\circ}$
- c)  $3^{\circ} > 1^{\circ} > 2^{\circ}$
- d)  $1^{\circ} > 3^{\circ} > 2^{\circ}$
- 170. Boiling points of alcohols are generally high. This is due to
  - a) hydrogen-bonding intermolecular attractions
    - b) dipole-dipole attractions
    - c) both of the above
    - d) none of the above
- 171. In the following dehydration of alcohol (A) leading to the formation of styrene (B)

$$\begin{array}{c}
CHCH_3 \xrightarrow{H^+} CHCH_2 \\
OH \\
(A)
\end{array}$$

$$CH = CH_2$$

$$(B)$$

- a) intermediate is carbocation
- b) carbocation is resonance-stabilised
- c) initial attack is H+
- d) all are correct statements
- 172. Which of the following functional groups can not be reduced by  $H_2$  / Ni
  - a) R-CHO
- b) R-COOH
- c) R-COO-R
- d) R-CO-R
- 173. 3-ethyl pentan-3-ol is obtained by C<sub>2</sub>H<sub>5</sub>MgBr and what?
  - a) pentan-2-one
  - b) pentan-3-one
  - c) pentanal
  - d) 3-methyl butan-2-one
- 174. Acidic character of alcohols depends up on
  - a) number of alkyl groups
  - b) polarity of –OH groups
  - c) types of alkyl groups
  - d) all of these
- 175. Order of acidity 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)  $3^{\circ} > 1^{\circ} > 2^{\circ}$
- 176. The B.P. of alcohols are much higher than the ethers of comparable molecular masses due to
  - a) interamolecular H bonding
  - b) intermolecular H bonding
  - c) dipole dipole attraction
  - d) Hitler London forces
- 177. Which one is not characteristic of alcohols?
  - a) They are lighter than water
  - b) Their B.P. rise uniformly with increasing molecular mass
  - c) Lower members are insoluble in water but solubility increases regularly
  - d) Lower members have pleasent odour and burning test
- 178. Consider following reactions,
  - I.  $CH_3CH_2CHCH_3 \xrightarrow{H^+} A \text{ (major)}$  OH

$$\begin{array}{c} \operatorname{CH_3} \\ | \\ \operatorname{II.} \ \operatorname{CH_3C} - \operatorname{CHCH_3} \xrightarrow{\operatorname{H}^+} \operatorname{B} \text{ (major)} \\ | & | \\ \operatorname{CH_3OH} \end{array}$$

A and B (both alkenes) are respectively:

- a)  $CH_3CH = CHCH_3$   $CH_3CCH = CH_2$   $CH_3$   $CH_3$
- b)  $CH_3CH_2CH = CH_2CH_3CCH = CH_2$  $CH_3$
- c)  $CH_3CH = CHCH_3 CH_3C = CCH_3$  $\begin{vmatrix} & & & \\ & & &$
- d)  $CH_3CH_2CH = CH_2CH_3C = CCH_3$ | | |  $CH_3CH_3$
- 179. Glycerol is more viscous than propan-l-ol due to
  - a) many hydrogen bonding per molecule
  - b) high B.P.
  - c) high molecular weight
  - d) more Lewis basic character
- 180.  $C_2H_5OH$  has higher B.P. than,  $C_2H_5-SH$  due to
  - a) association
  - b) dissociation
  - c) low molecular mass
  - d) two lone pair of electron on oxygen
- 181. When 1 mol of ethanol reacts with sodium metal liberate how many gram of hydrogen?
  - a) 1 gm of hydrogen b) ½ gm of hydrogen
  - c) 1.5 gm of hydrogen d) 2 gm of hydrogen
- 182. The dimer of methyl alcohol will have strecture

a) 
$$H - O - C - H - O - CH_3$$

b) 
$$CH_3 - O.....O - CH_3$$
  
 $\begin{vmatrix} & & | \\ & & | \\ & & H \end{vmatrix}$ 

c) 
$$H - O - CH_3 \dots CH_3 - O - H$$

183. Which of following has highest B.P.

- a) CH<sub>3</sub>-O-CH<sub>3</sub>
- b) C<sub>2</sub>H<sub>5</sub>-OH
- c) C<sub>2</sub>H<sub>5</sub>-C1
- d) CH<sub>2</sub>-CHO
- 184. Which of following is the most viscous liquid?
  - a) C<sub>2</sub>H<sub>5</sub>OH

b) 
$$CH_2 - CH_2$$
  
 $\mid$   $\mid$  OH OH

c) 
$$CH_2 - CH - CH_2$$
  
 $\mid \quad \mid \quad \mid$   
 $OH \quad OH \quad OH$ 

- d) HO-CH<sub>2</sub>(CHOH)<sub>4</sub>CH<sub>2</sub> OH
- 185. Which of the following exhibit highest B. P.?
- 186. Which of the following is most soluble in water
  - a) CH<sub>3</sub>-OH
- b)  $C_2H_5$ –Cl
- c) CH<sub>3</sub>-O-CH<sub>3</sub>
- d) C<sub>2</sub>H<sub>5</sub>-OH
- 187. Reason for excessive solubility of alcohol in water is due to
  - a) covalent bonding
- b) H-bonding with H<sub>2</sub>O
- c) ionic bonding
- d) Lewis base character
- 188. Alcohols of high molecular masses are
  - a) high B.P. and excessible solubility
  - b) low B.P. and excessive solubility
  - c) high B.P. and low solubility
  - d) low B.P. and low solubility
- 189. Compound with molecular formula C<sub>3</sub>H<sub>5</sub>O on vigorous oxidation produces an acid C<sub>3</sub>H<sub>6</sub>O<sub>2</sub>. It is,
  - a) 3° alcohol
- b) 2° alcohol
- c) 1° alcohol
- d) not necessary
- 190. Which of the following reacts less easily with sodium metal?
  - a) t-butyl alcohol
- b) isopropyl alcohol
- c) methyl alcohol
- d) ethyl alcohol
- 191. Acetic acid and n-propyl alcohols has same molecular mass (60). Out of these two, which have higher B.P.
  - a) Acetic acid
- b) n-propyl alcohol
- c) either
- d) neither
- 192. ZnCl, in Lucas reagent is
  - a) Lewis acid
- b) Lewis base
- c) both 'a' and 'b'
- d) none of these
- 193. Oxidation of 2-pentanone mainly produces
  - a) butyric acid and CO<sub>2</sub> + H<sub>2</sub>O
  - b) acetic acid and propionic acid
  - c) propionic acid +  $CO_2$  +  $H_2O$

d) acetic acid and butyric acid

- 194. The correct increasing order of acidic strength is
  - a)  $CH_3OH > H_2O > (CH_3)_2$  CH OH
  - b)  $H_2O > CH_3OH > (CH_3)_2CHOH$
  - c)  $(CH_3)_2$  CH OH >  $CH_3$ OH >  $H_2$ O
  - d)  $H_2O > (CH_3)$ ,  $CH OH > CH_3OH$

$$CH_3$$

- 195. An alkyl halide  $CH_3 \dot{C}Cl C_2H_5$  can be obtained by the action of HCl on which alcohol
  - a) (CH<sub>2</sub>)<sub>2</sub>CH–CH(OH)CH<sub>2</sub>
  - b) (CH<sub>3</sub>)<sub>2</sub>CH-CH<sub>2</sub>-CH<sub>2</sub>-OH

d) all of these

196. 
$$(CH_3)_2 - C - CH_2 - C(CH_3)_3 \xrightarrow{\text{acidic} \atop \text{dichromate}}$$

the main product is

a) 
$$CH_3$$
  $CH_3$   $CH_3$   $CH_3$   $CH_3$   $CH_4$   $CH_5$   $CH_5$ 

$$\begin{array}{cccc} CH_{3} & CH_{3} \\ & & | \\ CH_{2} = C - CH_{2} - C - CH_{3} \\ & & | \\ CH_{3} \end{array}$$

$$\begin{array}{ccc} CH_3 & CH_3 \\ \mid & \mid \\ c) & CH_3 - C = CH - C = CH_2 \end{array}$$

$$\begin{array}{c|c} CH_3 & CH_3 \\ & \mid & \mid \\ d) \ CH_2 = C - CH_2 - C = CH_2 \end{array}$$

- 197. Ethanol is soluble in water due to
  - a) ethyl group
- b) hydrogen bonding
- c) its neutral
- d) dissociation in water
- 198. The final product of the oxidation of ethyl alcohol is
  - a) ethane
- b) acetone
- c) acetaldehyde
- d) acetic acid
- 199. Which has maximum p K<sub>3</sub> value?

- a) OH at C<sub>2</sub> is more basic than that of at C<sub>5</sub>
- b) OH at C<sub>2</sub> is more acidic than at C<sub>5</sub>
- c) both behave as a base
- d) both behave as an acid
- 201. When ethyl alcohol is oxidised by copper, then which of the following aldehyde is formed?
  - a) Formaldehyde
- b) Acetyldehyde
- c) Benzaldehyde
- d) Crotonaldehyde
- 202. The boiling point of a compound is raised by
  - a) volatility of compound
    - b) non-polarity in the molecules
    - c) intermolecular hydrogen bonding
    - d) intramolecular hydrogen bonding
- 203. The alcohol, that is used as a beverage, is
  - a) propanol
- b) butanol
- c) ethanol
- d) methanol
- 204. Wood spirit is known as
  - a) methanol
- b) ethanol
- c) acetone
- d) benzene
- 205. Solubility of alcohols in water depends upon
  - a) acidic nature
  - b) basic nature
  - c) neutral nature
  - d) tendency to form hydrogen bonding
- 206. Consider the following substances
  - 1) 2–propanol,
- 2) propanone,
- 3) methyl amine

The correct sequence of increasing order of boiling point is

- a) 2 < 3 < 1
- b) 1 < 2 < 3
- c) 2 < 1 < 3
- d) 3 < 2 < 1
- 207. Alcohols are miscible with H<sub>2</sub>O because of their
  - a) acidic character
- b) H-bonding
- c) alkyl group
- d) dissociation
- 208. The reaction

$$CH_{3}CH_{2}OH \xrightarrow{95\% H_{2}SO_{4}} CH_{2} = CH_{2} + H_{2}O$$

is an example of

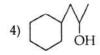
- a) dehydration
- b) dehydrogenation

- c) hydration
- d) decarboxylation
- 209. Tonics, generally contains,
  - a) ether
- b) methanol
- c) ethanol
- d) rectified spirit
- 210. Dehydration is most easy for
  - a) primary alcohols
- b) tertiary alcohols
- c) secondary alcohols d) ethanol
- 211. The alcohol, C<sub>4</sub>H<sub>9</sub>OH, when shaken with a mixture of anhydrous ZnCl<sub>2</sub> and cone. HCl give an immediate oil layer product. The alcohol is
  - a) H<sub>3</sub>C-(CH<sub>2</sub>)<sub>3</sub>-OH
  - b) H<sub>2</sub>C-CH(OH)CH<sub>2</sub>CH<sub>3</sub>
  - c) (CH<sub>3</sub>)<sub>2</sub>CHCH<sub>2</sub>-OH
  - d) (CH<sub>2</sub>)<sub>2</sub>C-OH
- 212. On oxidation of alcohol gives an acid having the same number of carbon atoms. The alcohol is,
  - a) 1° alcohol
- b) 2° alcohol
- c) 3° alcohol
- d) not necessary
- 213. Secondary butyl alcohol is dehydrated according to
  - a) Saytzeff rule
  - b) Markownikoff rule
  - c) Anti-Markownikoff rule
  - d) none of these
- 214. Olefins are obtained from alcohols by heating with
  - a) Al<sub>2</sub>O<sub>2</sub>
- b) LiAlH<sub>4</sub>
- c)  $B_2H_6$
- d) NaBH<sub>4</sub>
- 215. Cyclohexene is obtained from dehydration of









- a) 1
- b) 1, 2
- c) 1, 3
- d) 4
- 216. 1°, 2° and 3° alcohols are identified by
  - a) Lucas test
- b) oxidation test
- c) haloform test
- d) all of these
- 217. Following alcohol on dehydration gives



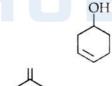








- 218. 3, 3-dimethyl butan-2-ol on dehydration gives
  - a) 3, 3-dimethyl but-2-ene
  - b) 2, 3-dimethyl but-2-ene
  - c) 2, 3-dimethyl but-l-ene
  - d) 3, 3-dimethyl but-l-ene
- 219. The compound on oxidation gives ketone, the original compound is
  - a) 1° alcohol
- b) 2° alcohol
- c) 3° alcohol
- d) carboxylic acid
- 220. A organic compound (A) has pleasent odour, on boiling (A) with conc.H<sub>2</sub>SO<sub>4</sub> at 443K produces colourless gas, which decolourises bromine water and Bayer's reagent. The original organic compound (A) is
  - a)  $C_2H_5-Cl$
- b) C<sub>2</sub>H<sub>5</sub>-COOCH<sub>3</sub>
- c) C<sub>2</sub>H<sub>5</sub>- OH
- d) C<sub>2</sub>H<sub>5</sub>- COOH
- 221. The most suitable reagent to convert primary alcohol to aldehyde
  - a) acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>
  - b) alkaline KMnO<sub>4</sub>
  - c) CrO<sub>3</sub>
  - d) pyridinium chlorochromate (PCC)
- 222. Following compound on dehydration gives

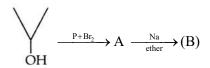






- c) [
- d) none of these
- 223. When compound (A) is oxidised by acidic  $K_2Cr_2O_7$  gave (B). Compound (B) on reduction with LiAIH<sub>4</sub> gave (A). The compound (A) and (B)are respectively.
  - a) CH<sub>3</sub> COCH<sub>3</sub> and CH<sub>3</sub>-COOH
  - b) C<sub>2</sub>H<sub>5</sub>OH and CH<sub>3</sub>-COCH<sub>3</sub>
  - c)  $C_2H_5$ -OH and  $CH_3$  COOH
  - d)  $CH_3$  CHO and  $CH_3$  CO  $CH_3$

224. The major product formed in following reaction is



The compound (B) is

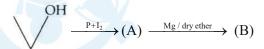




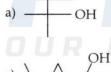




- 225. Which is best reagent to convert cyclohexanol to cyclohexene
  - a) cone. HCl
- b) conc. HBr
- c) cone. H<sub>2</sub>SO<sub>4</sub>
- d) Lucas reagent
- 226. In the following sequence of reaction



i) CH<sub>3</sub>COCH ii) dry ether iii)  $H_3O^+$  (C). The compound (C) is









227.  $C_2H_5COOH \xrightarrow{LiAlH_4} A$ 

$$A \xrightarrow{Al_2O_3} B + H_2O$$

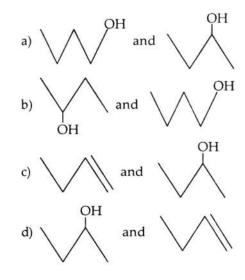
In above reaction A and B are respectively

- a)  $C_2H_5$ -OH and  $CH_2 = CH_2$
- b)  $C_2H_5$  CHO and  $C_2H_5OH$
- c)  $CH_3 CH_2 CH_2 OH$  and  $CH_3 CH = CH_2$
- d)  $CH_3 CH = CH_2$  and  $CH_3 CH_2 CH_2 OH$
- 228. In the sequence of reaction.

$$\mathrm{CH_3-CH_2-CH} = \mathrm{CH_2} \xrightarrow{\mathrm{i)} \, \mathrm{B_2H_6/THF}} \mathrm{ii) \, H_2O_2/NaOH}$$

$$A \xrightarrow{\text{conc. } H_2SO_4} B \xrightarrow{H_3O^+} C$$

In above reaction A and C are respectively

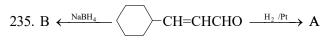


- 229. Dehydration of 3-phenyl butan-2-ol gives
  - a) 2-phenyl but-2-ene
  - b) 4-phenyl 2-methyl but-1-ene
  - c) 1-phenyl 3-methyl but-1-ene
  - d) 3-phenyl but-1-ene
- 230. In dehydration of alcohol 1st step is
  - a) formation of carbonium ion
  - b) formation of carbonion
  - c) loss of proton from carbonium ion
  - d) protonated of alcohol
- 231. Dehydration of alcohol produces alkene, the reaction intermediate is
  - a) carbonium ion
- b) carbanion
- c) carbon free radical d) carbene
- 232. Denatured spirit is mainly used as a
  - a) good fuel
  - b) drug
  - c) solvent in preparing varnishes
  - d) material in the preparation of oil
- 233. Acetylation is a process in which the hydrogen of O-H group is replaced by
  - a)  $-C \equiv C-H$
- b) -CH = CH
- c) -COCH<sub>3</sub>
- d)  $-COC_6H_5$

is

Br Br 
$$\mid$$
 a)  $(CH_3)_3CCHCH_3$  b)  $(CH_3)_2CCHCH_3$ 

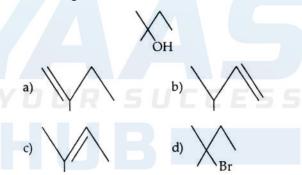
- c) both 'a' and 'b'
- d) none is correct



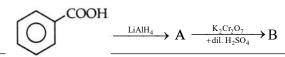
A and B are respectively:

- a) CH<sub>2</sub>CH<sub>2</sub>CHO,
  - CH=CHCH<sub>2</sub>OH
- b) CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>OH,
  - $\bigcirc$  -CH = CHCH<sub>2</sub>OH
- c)  $\sim$  CH = CHCH<sub>2</sub>OH in both cases
- d) CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>OH in both cases
- 236. When 2, 3 dimethyl 2-butanol under goes acid catalysed dehydration the minor product is,
  - a) 2, 3 dimethyl 1-butene
  - b) 2, 3 dimethyl 2-butene
  - c) 3, 3 dimethyl 1-butene
  - d) none of these
- 237. Lucas reagent is
  - a) anhydrous ZnCl, dissolved in cone. HNO,
  - b) hydrous ZnCl, dissolved in conc. HCl
  - c) anhydrous ZnCl, dissolved in cone. HCl
  - d) anhydrous ZnCl, dissolved in dil. HCl
- 238. Alcohols of low molecular weight are
  - a) insoluble in all solvents
  - b) insoluble in water
  - c) soluble in water at room temperature
  - d) soluble in water on heating
- 239. 6 mole of ethyl alcohol reacts with sodium metal. How many moles of hydrogen are liberated?
  - a) 2
- b) 3
- c) 4
- d) 6
- 240. Catalytic oxidation of benzyl alcohol gives
  - a) benzaldehyde
- b) benzoic acid
- c) toluene
- d) phenol
- 241. The ease of dehydration of alcohol is in the order
  - a)  $1^{\circ} > 2^{\circ} > 3^{\circ}$
- b)  $1^{\circ} > 3^{\circ} > 2^{\circ}$
- c)  $3^{\circ} > 2^{\circ} > 1^{\circ}$
- d)  $2^{\circ} > 1^{\circ} > 3^{\circ}$
- 242. Alkenes are obtained from alcohols by
  - a) oxidation
  - b) hydration

- c) intermolecular dehydration
- d) intramolecular dehydration
- 243. Methanol and ethanol are miscible in water due to
  - a) dissociation in water
  - b) their acidic nature
  - c) hydrogen bond with water
  - d) alkyl groups
- 244. When ethanol is treated with acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>, it forms acetic acid. It is an example of
  - a) hydrolysis
- b) oxidation
- c) reduction
- d) rearrangement
- 245. Which of the following compound react fastest with sodium metal.
  - a) H<sub>2</sub>O
- b) CH<sub>3</sub>-OH
- c) C<sub>2</sub>H<sub>5</sub>-OH
- d) CH<sub>3</sub>-CH<sub>2</sub>-CH<sub>2</sub>-OH
- 246. Cyclohexanol is reacted with Lucas reagent gives
  - a) 1-cyclohexyl chloromethane
  - b) chlorocyclohexane
  - c) 1-chlorocyclohexene
  - d) 1-chlorocyclohexyne
- 247. Following compound is reacted with conc. H<sub>2</sub>SO<sub>4</sub> at 363K gives



- 248. 1, 1–diphenyl methanol is reacted with, HI give
  - a) 1, 1-diphenyl iodomethane
  - b) 1, 1 -diphenyl iodoethane
  - c) diphenyl
  - d) none of these
- 249. Which of the following is not dehydrating agent?
  - a) H,SO<sub>4</sub>
- b) H<sub>3</sub>BO<sub>3</sub>
- c) ThO,
- d) NaBH,
- 250. Ethyl alcohol on heating with HI yield
  - a) ethane
- b) ethylene
- c) methane
- d) ethyl iodide
- 251. Product of the following reaction is



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

- a) benzaldehyde
- b) benzyl alcohol
- c) toluene
- d) benzoic acid
- 252. The decreasing order of basicity of alcohols are
  - a)  $3^{\circ} > 2^{\circ} > 1^{\circ}$
- b)  $3^{\circ} > 1^{\circ} > 2^{\circ}$
- c)  $2^{\circ} > 3^{\circ} > 1^{\circ}$
- d)  $1^{\circ} > 2^{\circ} > 3^{\circ}$
- 253. Which of the following alcohol reacts with HI by SN¹ reaction?
  - a) t-butyl alcohol
- b) methyl alcohol
- c) n-propyl alcohol
- d) ethyl alcohol
- 254. Dehydration of neophentyl alcohol gives
  - a) 2-methyl but-l-ene
  - b) 3-methyl but-l-ene
  - c) 2-methyl but-2-ene
  - d) no product will be formed
- 255. Which of the following can be used as dehydrating agent for alcohols?
  - a) H<sub>3</sub>PO<sub>4</sub>
- b) H<sub>2</sub>SO<sub>4</sub>
- c) Al<sub>2</sub>O<sub>3</sub>
- d) all of these
- 256. Which of the following compound will lose a molecule of water of treating with conc. H<sub>2</sub>SO<sub>4</sub>?
  - a) CH<sub>3</sub>COCH<sub>3</sub>
- b) CH<sub>3</sub>COOH
- c) CH<sub>3</sub>CH<sub>2</sub>OH
- d) CH<sub>3</sub>OCH<sub>3</sub>
- 257. Which of the following is an example of elimination reaction?
  - a) Dehydration of alcohol
  - b) Chlorination of CH,
  - c) Hydroxylation of C<sub>2</sub>H<sub>4</sub>
  - d) Nitration of benzene
- 258. 23 g of sodium reacts with 1 mole methyl alcohol to give
  - a) half mole of H,
- b) one mole of O,
- c) one mole of H,
- d) either 'b' or 'c'
- 259. Which of the following compound will give ketone on oxidation?
  - a) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>OH
  - b) CH<sub>3</sub>CH<sub>2</sub>CH(OH)CH<sub>3</sub>
  - c) (CH<sub>3</sub>),COH
  - d) (CH<sub>2</sub>)<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>OH
- 260. Which of the following is most acidic?
  - a) RCH,OH
- b) R,CHOH
- c) R,C(OH),
- d) CH,OH
- 261. Lucas test is positive with
  - a) CH<sub>3</sub>-O-CH<sub>3</sub>
- b) C<sub>2</sub>H<sub>5</sub>-OH
- c) CH<sub>2</sub>-Cl
- d) CH<sub>2</sub>-CHO
- 262. The first product of oxidation of primary alcohol is

- a) carboxylic acid b) ketone
- c) ester
- d) aldehyde
- 263. Sodium metal reacts readily with
  - a) R-CHO
- b) R-CH<sub>2</sub>OH
- c) ester
- d) R-NH,
- 264. KMnO<sub>4</sub> acts as oxidising agent in
  - a) acidic medium
- b) neutral medium
- c) alkaline medium
- d) all of the above
- 265. The group obtained by the removal of H-atom of the -OH group of ROH is called
  - a) alkyl group
- b) alkene
- c) alkoxy group
- d) all of these
- 266. Alcohols are
  - a) neutral
- b) strongly acidic
- c) basic
- d) amphoteric
- 267. Which one of the following on oxidation gives a ketone?
  - a) Primary alcohol
- b) Secondary alcohol
- c) Tertiary alcohol
- d) All of these
- 268. Correct order of increasing boiling points is
  - a) propane < n-butane < ethanol < water
  - b) propane < ethanol < n-butane < water
  - c) water < ethanol < propane < n-butane
- d) water < propane < n-butane < ethanol</li>
   269. The order of reactivity of following alcohols towards HCl is,
  - 1. CH<sub>2</sub>OH
- 2. CH, CH, CH, OH
- 3. (CH<sub>3</sub>)<sub>2</sub>CHOH
- 4. (CH<sub>3</sub>),COH
- a) 1 > 2 > 3 > 4
- b) 4 > 3 > 2 > 1
- c) 3 > 4 > 2 > 1 d) 2 > 4 > 1 > 3270. Tertiary alcohols are resistant to oxidation
  - a) they do not have a-hydrogen atom
  - b) due to large +1 effect of alkyl group
  - c) due to greater steric hindrance
  - d) all of these
- 271. Which of the following reactions shows acidic nature of alcohol?
  - a) ROH + HOOCR  $\rightarrow$  RCOOR + H<sub>2</sub>O
  - b)  $2ROH + 2Na \rightarrow 2RONa + H$ ,
  - c) ROH + CIOCR  $\rightarrow$  RCOOR + HCl
  - d) ROH + HCl  $\rightarrow$  RCl + H<sub>2</sub>O
- 272. Boiling point of alcohol is more than that of ether of corresponding molecular weight, because
  - a) alcohol bein.g more soluble in water
  - b) ethers are non-polar solvent

- c) hydrogen bonding exist between alcohol
- d) none of these
- 273. Use of methanol may causes
  - a) blindness and death due to its oxidation to CO,
  - b) blindness and death due to HCOOH
  - c) deficinency of calcium due to formation of salt
  - d) disorder of blood hormones
- 274. Which of the following compound does not react with Lucas reagent?
  - a) (CH<sub>3</sub>)<sub>2</sub>C-CHO
- b) (CH<sub>3</sub>),C-OH
- c)  $(CH_3)_2$  CH–OH
- d) CH<sub>2</sub>-OH
- 275. The final subsidiary product in the following reaction is, R-OH + PX<sub>3</sub>  $\xrightarrow{\Delta}$  R-X +?
  - a)  $H_3PO_3$
- b) HX
- c) H<sub>3</sub>PO<sub>4</sub>
- d) HPO<sub>4</sub>
- 276. Which one of the following process is used to distinguish between the three types of alcohols?
  - a) Reduction
- b) Hydrolysis
- c) Oxidation
- d) Hydrogenation
- 277. Ketone is the first oxidative product of,
  - a) 1° alcohol
- b) 3° alcohol
- c) 2° alcohol
- d) acid
- 278. The compound which liberates H<sub>2</sub> gas with sodium metal is,
  - a) aldehyde
- b) ethanol
- c) ether
- d) ketone
- 279. 1–propenol is converted into propene, which of the following agent is used?
  - a) ale. KOH
- b) dil. NaOH
- c) dil. HCl
- d) cone, H<sub>2</sub>SO<sub>4</sub>
- 280. Optical isomer of molcular formula  $C_4H_{10}O$  on  $\alpha$  –elimination gives
  - a) butanal
- b) 2–methyl propanal
- c) 2-butanone
- d) 2-methyl propene
- 281. Which of the following is associated liquids?
  - a) ROH
- b) H,O
- c) RNH,
- d) All of these
- 282. Methyl alcohol on oxidation with acidified  $K_2Cr_2O_7$  gives
  - a) CH<sub>2</sub>CHO
- b) HCOOH
- c) CH<sub>3</sub>COCH<sub>2</sub>
- d) CH<sub>3</sub>COOH
- 283. 2-butanol on dehydration mainly gives,
  - a)  $CH_3CH_2CH = CH_3$
  - b)  $CH_3CH = CHCH_3$
  - c)  $(CH_3)_3C = CH_2$
- d)  $CH_3CH = CH_2$
- 284. 1-propanol can be converted into 1-chloro

- propane by HCl in the presence of catalyst,
- a) hydrous ZnCl<sub>2</sub>
- b) unhydrous ZnCl,
- c) H<sub>3</sub>PO<sub>4</sub>
- d)  $P_2O_5$
- 285. Oxidation means,
  - a) addition of oxygen
  - b) increase in oxidation state
  - c) loss of electron
  - d) all of these
- 286. Which statement is not correct about the alcohols?
  - a) Alcohols involves H-bonding
  - b) Alcohols evaporates quickly than water
  - c) Alcohols of less number of carbon atoms are less soluble than more number of carbon atoms
  - d) All of these
- 287. Product 'B' of the following reaction is

$$CH_3$$
- $Cl + CH_3$  -  $COOAg \longrightarrow A \xrightarrow{H_2+Ni} B$ 

- a) CH<sub>3</sub>-OH and C<sub>2</sub>H<sub>5</sub>-OH
- b) CH,-CHO and CH,-OH
- c) CH<sub>2</sub>-CHO and CH<sub>2</sub>-CHO
- d) C<sub>2</sub>H<sub>5</sub>-OH and C<sub>2</sub>H<sub>5</sub>-OH
- 288. Find out correct reducing agent in following conversion.

$$CH_3$$
- $CH_2$ - $COOH \longrightarrow$ 

- a)  $H_2 + Ni$
- b) NaBH<sub>4</sub>
- c) NaHg + H<sub>2</sub>O
  - d) LiAlH
- 289. Compound A (ester) reacts with LiAIH<sub>4</sub> gives B and C. Compound B on oxidation gives acetic acid and compound C on oxidation gives formic acid. The compound 'A' is
  - a) C<sub>2</sub>H<sub>5</sub>-COOCH<sub>3</sub>
    - b) CH<sub>3</sub>COOC<sub>2</sub>H<sub>5</sub>
  - c) CH<sub>3</sub>-COOCH<sub>3</sub>
- d) CH<sub>3</sub>-CH<sub>2</sub>-COOH
- 290. Which of the following compound have covalent and ionic bond?
  - a) R-OH
- b) H-O-H
- c) R-X
- d) R-ONa

## OH

291. 
$$(CH_3)_2CH - CH - CH_3 \xrightarrow{Al_2O_3} X$$

Give the IUPAC name of major product formed in the reaction

- a) 3-methyl 2-butene b) isobutylene
- c) 2-methyl 2-butene d) 2-methyl 1-propene
- 293. Which of the following is / are correct?
  - a) Absolute alcohol is 100 % ethanol

- b) The alcohol sold in the market for polishing is known as methylated spirit
- c) Ordinary ethanol is known as rectified spirit
- d) All of these
- 294. Mixture of acetic acid and propionic acid is obtained from oxidation of
  - a) CH<sub>3</sub>-COCH<sub>3</sub>
  - b) CH<sub>3</sub>-COC<sub>2</sub>H<sub>5</sub>
  - c) CH<sub>3</sub>-CH<sub>2</sub>-COCH<sub>2</sub>-CH<sub>3</sub>
  - d) CH<sub>3</sub>-CH<sub>2</sub>-CHO
- 295. An organic compound (A) produces  $(CH_2)_2C = CH - CH_2$  on dehydration.

The compound A is

- a)  $(CH_2)_2CCH_2 OH$
- b) (CH<sub>3</sub>),COHCH, CH,
- c) (CH<sub>3</sub>)<sub>2</sub>CH–CHOH CH<sub>3</sub>
- d) all of these
- 296. The hydrogen bonding ability of 1°, 2° and 3° alcohols is of the order
  - a)  $3^{\circ} > 2^{\circ} > 1^{\circ}$
- b)  $1^{\circ} > 2^{\circ} > 3^{\circ}$
- c)  $3^{\circ} > 1^{\circ} > 2^{\circ}$  d)  $1^{\circ} > 3^{\circ} > 2^{\circ}$
- 297. When 3-methyl 3-pentanol is heated with alumina. The main product formed is,
  - a) 2-methyl 1-pentene
  - b) 3-methyl 2-pentene
  - c) 2-methyl 1-butene
  - d) 3-methyl 2-butene
- 298. An organic compound X reacts with sodium metal and evolve hydrogen gas, on oxidation of X by PCC give aldehyde. The formula of X could be
  - a) (CH<sub>2</sub>),CH–OH
- b) CH<sub>2</sub>-CH<sub>2</sub>-OH
- c) (CH<sub>3</sub>)C-OH
- d) CH<sub>3</sub>-CHOHC<sub>2</sub>H<sub>5</sub>
- 298. 2-butanol  $\xrightarrow{\text{conc. H}_2SO_4}$  1- butene + 2 butene

В

- which are correct statements?
- a) A is Saytzelf product B is not b) B is Saytzelf product A in not
- c) Either 'a' and 'b'
- d) Neither
- 299. Which of the following is maximum basic?
  - a) CH<sub>2</sub>CH<sub>2</sub>-CH<sub>2</sub>-CH<sub>2</sub>-OH
  - b) (CH<sub>3</sub>),CH–CH,–OH
  - c) CH<sub>2</sub>CHOHC<sub>2</sub>H<sub>5</sub>
  - d) (CH<sub>3</sub>)<sub>3</sub>C-OH
- 300. The C<sub>4</sub>H<sub>10</sub>O (alcohols) produces immediate terbidity with Lucas reagent the alcohol is

- a) CH<sub>3</sub>-CH<sub>2</sub>-CH<sub>2</sub>-CH<sub>3</sub>-OH
- b) (CH<sub>2</sub>),CH-CH<sub>2</sub>-OH
- c) CH<sub>2</sub>CHOHC<sub>2</sub>H<sub>2</sub>
- d) (CH<sub>3</sub>)<sub>3</sub>C-OH
- 301. Solubility of alcohol in water is due to
  - a) hydrophobic R-group
  - b) hydrophillic OH-group
  - c) hydrophobic OH-group
  - d) hydrophilic R-group
- 302. Boiling point of ethanol is greater than isomeric ether is due to
  - a) hydrogen bonds are much stronger than dipole dipole altraction
  - b) dipole-dipole attraction is much stronger than hydrogen bond
  - c) ether has two hydrophobic group while alcohol
  - d) ether has two hydrophilic group while alcohol has one
- 303. Lucas test is used to distingush between 1°, 2° and 3° alcohols. This shows that
  - a) R-OH behaves as base
  - b) greater the Pka value of alcohol, greater the reactivity of alcohol with HCl and thus sooner the formation of white turbidity
  - c) both are correct
  - d) none is correct
- 304. The most suitable reagent for convertion of  $R-CH_2-OH \rightarrow R-CHO$  is
  - a) neutral KMnO b) PCC
  - c) acidic K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> d) CrO,

305. A 
$$\xrightarrow{\text{K}_2\text{Cr}_2\text{O}_7}$$
 B  $\xrightarrow{\text{CH}_3\text{MgI}}$  CH<sub>3</sub>  $\xrightarrow{\text{CH}_3\text{MgI}}$  CH<sub>3</sub>  $\xrightarrow{\text{C}}$  CH<sub>4</sub>  $\xrightarrow{\text{C}}$ 

The reactant A is

- a) CH<sub>3</sub>CHOHCH<sub>3</sub>
- c) C,H,OH
- b) CH, COCH,
- d) CH<sub>2</sub>COOH
- 306. ZnXF or the reaction  $C_2H_5OH + HX \xrightarrow{ZnX_2}$  $C_2H_2X$ .

The order of reactivity is

- a) HBr > HI > HCl b) HI > HCl > HBr
- c) HI > HBr > HCl d) HCl > HBr > HI

- 307. Ethanol is converted into ethyl chloride by reacting with
  - a) POCl<sub>3</sub>
- b) SOCl,
- c) KCl
- d) NaCl
- 308. Which of following reducing agent is used to convert carboxylic acid into alcohol?
  - a) Na.Hg + H<sub>2</sub>O
- b) LiAIH,
- c) NaBH<sub>4</sub>
- d) Sn + HCl
- 309. Primary, secondary and tertiary alcohols may be distinguished by employing
  - a) Hoffmann's test
- b) Fehling solution test
- c) Lucas test
- d) None of the above
- 310. Oxidation of ethanol by chromic acid forms
  - a) ethanal
- b) methanol
- c) 2-propanone
- d) ethanoic acid
- 311. Conc.  $H_2SO_4$  reacts with  $C_2H_5OH$  at 443K to form
  - a) CH<sub>3</sub>COCH<sub>3</sub>
- b) CH,COOH
- c) CH<sub>2</sub>CHO
- d) C,H4
- 312. Distinction between primary, secondary and tertiary alcohol is done by
  - a) oxidation method
- b) Lucas test
- c) silver mirror test
- d) both 'a' and 'b'
- 313. Low molecular weight alcohols are
  - a) soluble in water
  - b) soluble on heating
  - c) insoluble in water
  - d) insoluble in all solvent
- 314. Isopropyl alcohol on oxidation forms
  - a) acetone
- b) ether
- c) ethylene
- d) acetaldehyde
- 315. Among the following compounds which can be dehydrated very easily is

a) 
$$CH_3$$
  $CH_3 - CH_2 - C - CH_2 - CH_3$   $CH_3 - CH_3 -$ 

d) 
$$CH_3 - CH_2 - CH - CH_2 - CH_2 - OH$$
  $CH_3$ 

316. Which of the following is not characteristic of

- alcohols?
- a) Lower alcohols are stronger and have bitter taste
- b) Higher alcohols are stronger and have bitter taste
- c) The boiling points of alcohols increase with increasing molecular mass
- d) The lower alcohol are soluble in water
- 317. In reaction of alcohols with alkali metal which of the following alcohols will react fastest
  - a) secondary
- b) tertiary
- c) primary
- d) all equal
- 318. The –OH group of methyl alcohol cannot be replaced by chlorine by the action of
  - a) chlorine
  - b) hydrogen chloride
  - c) phosphorus trichloride
  - d) phosphorus pentachloride
- 319. When ethanol is passed over red hot copper at 573K, the product formed is
  - a) CH<sub>3</sub>CHO
- b) CH<sub>3</sub>COCH<sub>3</sub>
- c) C<sub>2</sub>H<sub>4</sub>
- d) CH<sub>2</sub>COOH
- 320. A mixture of methanol vapours and air is passed over heated copper. The products are
  - a) carbon monoxide and hydrogen
  - b) formaldehyde and H, gas
  - c) formic acid and water vapour
  - d) carbon monoxide and water vapour
- 321. When ethyl alcohol reacts with thionyl chloride in the presence of pyridine, the product obtained is
  - a) CH<sub>3</sub>CH<sub>2</sub>Cl + HCl
  - b)  $C_2H_5Cl + HCl + SO_2$
  - c) CH<sub>3</sub>CH<sub>2</sub>Cl + H<sub>2</sub>O + SO,
  - d) H<sub>3</sub>CH<sub>2</sub>Cl + Cl<sub>2</sub> + SO,
- 322. Primary alcohols on dehydration give
  - a) alkenes
- b) ether
- c) alkane
- d) ester
- 323. Primary and secondary alcohols on action of reduced copper give
  - a) aldehydes and ketones respectively
  - b) ketones and aldehydes respectively
  - c) only aldehydes
  - d) only ketones
- 324. Methyl alcohol on oxidation with acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> gives
  - a) CH<sub>3</sub>COCH<sub>3</sub>
- b) CH<sub>3</sub>CHO

- c) HCOOH
- d) CH<sub>2</sub>COOH
- 325. Ethyl alcohol on oxidation with K<sub>2</sub>Cr<sub>2</sub>O<sub>3</sub> gives

  - a) acetic acid
- b) acetaldehyde
- c) formaldehyde
- d) formic acid
- 326. On heating ethanol with excess of cone. H<sub>2</sub>SO<sub>4</sub> at 443K, product obtained is
  - a) ethene
- b) ethane
- c) ethyne
- d) ethoxy ethane
- 327. When 3, 3-dimethyl 2-butanol is heated with  $H_2SO_4$  the major product obtained is
  - a) 2, 3-dimethyl 1-butene
  - b) 3, 3-dimethyl 1-butene
  - c) 2, 3-dimethyl 2-butene
  - d) cis and trans isomers of 2, 3-dimethyl 2-butene
- 328. Which of the following will give benzoic acid on oxidation?
  - a) Benzyl alcohol
- b) Benzaldehyde
- c) Acetophenone
- d) All of these
- 329. n-propyl alcohol and isopropyl alcohol can be chemically distinguished by which reagent
  - a) PCl<sub>5</sub>
  - b) reduction
  - c) oxidation with potassium dichromate
  - d) PCl,
- 330. Which of the following pairs of alcohols are distinguished by oxidation test, Lucas test and haloform test?
  - a) Methanol and ethanol
  - b) Ethanol and 3-pentanol
  - c) Ethanol and 2-propanol
  - d) 1–propanol and 3–pentanol
- 331. Consumption of alcohol by vehicle drivers is detected by
  - a) blow of mouth air in test tube containing acidic  $K_{2}Cr_{2}O_{7}$

- b) blow of mouth air in test tube containing alcoholic KOH
- c) blow of mouth air in test tube containing Cu<sub>2</sub>O
- d) blow of mouth air in test tube containing Schiff's reagent
- 332. Which of the following is pyridinium chlorochromate?
  - a) C<sub>6</sub>H<sub>5</sub>NH + CrO<sub>3</sub>Cl b) C<sub>5</sub>H<sub>5</sub>NH + CrO<sub>3</sub>Cl
  - c) C<sub>6</sub>H<sub>5</sub>SO<sub>5</sub>Cl
- d) C<sub>5</sub>H<sub>5</sub>SO<sub>5</sub>Cl
- 333. Separation of two layers are seen when Lucas reagent is treated with
  - a) CH<sub>2</sub>OH
- b) CH,Cl
- c) (CH<sub>2</sub>)<sub>2</sub>C-OH
- d) (CH<sub>2</sub>)<sub>2</sub>C-Cl
- 334. Aluminium metal reacts with alcohol gives
  - a) ROAl
- b) (RO),A1
- c) (RO), A1
- d) RCOAl
- 335. When six mole of alcohol reacts with aluminium metal. How many gram of hydrogen is liberated?
  - a) 2
- b) 4
- c) 5
- d) 6
- 336. Ethyl alcohol is reacted with acetyl chloride gives
  - a) ethyl acetate
- b) ethyl formate
- c) ethyl propanoate
- d) ethyl methyl ketone
- 337. Methyl acetate is formed from methyl alcohol and what?
  - a) Acetic acid
- b) Acetic unhydride
- c) Acetyl chloride
- d) All of these
- 338.  $C_2H_5OH + A \xrightarrow{H^+} C_2H_5COOC_2H_5 +$ C,H,COOH The compound 'A' is
  - a) C<sub>2</sub>H<sub>5</sub>COCl
- b) (C,H,CO),O
- c) C<sub>2</sub>H<sub>5</sub>COOH
- d) C,H,CHO

COC



# SECTION - I : ALCOHOLS

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





### **MULTIPLE CHOICE QUESTIONS**

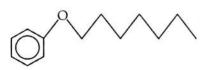
# INTRODUCTION, NOMENCLATURE, CLASSIFICATION, ISOMERISM

- 1. An ether is
  - a) R-O-R'
- b) RCOR
- b) RCHO
- d) RCOOR
- 2. The monovalent RO group is called
  - a) alkyl group
- b) alkoxy group
- c) alkenyl group
- d) all of these
- 3. Which of the following is a simple ether?
  - a) CH<sub>3</sub>OC<sub>2</sub>H<sub>5</sub>
- b) C,H,-OC,H,
- c) C<sub>2</sub>H<sub>5</sub>OC<sub>2</sub>H<sub>7</sub>
- d) CH<sub>3</sub>CH<sub>2</sub>OCH(CH<sub>3</sub>),
- 4. Which of the following is an unsymmetrical ether?
  - a) CH<sub>2</sub>-O-C<sub>2</sub>H<sub>7</sub>
- b) CH<sub>2</sub>-O-CH<sub>2</sub>
- c)  $C_2H_5-O-C_2H_5$
- d) All of these
- 5. An example of a compound with the functional group –O– is
  - a) acetic acid
- b) methyl alcohol
- c) diethyl ether
- d) acetone
- 6. In R'-O-R, the R' is a higher alkyl group, it is come from
  - a) alkane
- b) alcohol
- c) both 'a' and 'b'
- d) not predicted
- 7. The IUPAC name of C<sub>2</sub>H<sub>5</sub>OCH<sub>2</sub>CH(CH<sub>3</sub>)<sub>2</sub> is,
  - a) 3-ethoxy 2-methyl propane
  - b) 1-ethoxy 2-methyl propane
  - c) 1-ethoxy butane
  - d) 2-ethoxy butane
- 8. IUPAC name of the following compound (CH<sub>3</sub>)<sub>2</sub>CHOC(CH<sub>3</sub>)<sub>2</sub> is,
  - a) t-butyl isopropyl ether
  - b) 2, (2-propoxy) 2-methyl propane
  - c) 2-methyl 1-ethoxy 2-propane
  - d) 1-methyl 2-propoxy 2-propane
- 9. IUPAC name of ether is
  - a) alkyl alkanoate
- b) alkoxy alkane
- c) alkanamine
- d) alkyl acetate
- 10. IUPAC name of  $CH_3$ -O- $C_2H_5$  is
  - a) ethoxy methane
- b) methoxy methane
- c) methoxy ethane
- d) ethyl methyl ether
- 11 IUPAC name of methyl n-propyl ether is
  - a) propoxy methyl
  - b) 2-methoxy propane
  - c) 1-methoxy propane

- d) methyl propaonate
- 12. According to Lewis concept of acids and bases, ether is
  - a) acidic
- b) basic
- c) neutral
- d) amphoteric
- 13. What is IUPAC name of compound when divalent oxygen atom is attached to n-propyl group and iso-propyl group?
  - a) Propoxy 2-propane
  - b) Propoxy ethane
  - c) 1 (2-propoxy) propane
  - d) 1-ethoxy butane
- 14. Ethers have angular V-shaped geometry like
  - a) NH<sub>2</sub>
- b) H<sub>2</sub>O
- c) CH<sub>4</sub>
- d) CH<sub>2</sub>+
- 15. The IUPAC name of CH<sub>3</sub>OCH(CH<sub>3</sub>)<sub>2</sub> is,
  - a) 2-methoxy propane
  - b) 2-epoxy propane
  - c) 2-methoxy propane
  - d) 1-epoxy propane
- 16. IUPAC name of the following compound is



- a) 2-pentoxy benzene
- b) phenetol
- c) 2-phenoxy pentane
- d) isopentyl phenyl ether
- 17. IUPAC name of the following compound is



- a) 1-heptoxy benzene
- b) heptyl phenyl ether
- c) 1-phenoxy heptane
- d) 2-phenoxy heptane
- 18. Ethers are
  - a) Lewis acid
- b) acid
- c) Lewis base
- d) base
- 19. IUPACname of  $CH_3$ -O- $C(C_2H_5)_2$  is
  - a) 3-methoxy 2-ethyl pentane
  - b) 2-methoxy 2-ethyl pentane

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

- c) 3-methoxy 3-ethyl pentane
- d) 2-methoxy 3-ethyl pentane
- 20. How many metamers are possible for molecular formula  $C_4H_{10}O$ ?
  - a) 3
- b) 7
- c) 5
- d) 2
- 21. Following compounds are



- a) chain isomer
- b) linkage isomers
- c) position isomer
- d) functional isomers
- 22. IUPAC name of following compound is  $(CH_3)_3C-O-CH_2CH(CH_3)_2$ 
  - a) 2 (2-methyl 1-propoxy) 2-methyl propane
  - b) 1 (2–methyl 2–propoxy) 2–methyl propane
  - c) 1 (2-propoxy) 2-methyl propane
  - d) 2 (2-propoxy) 2-methyl propane
- 23. IUPAC name of ethyl t-butyl ether is
  - a) 2-ethoxy propane
  - b) 2-ethoxy 2-methyl propane
  - c) 2-ethoxy 2, 2-dimethyl ethane
  - d) 2-ethoxy 1-methyl propane
- 24. The compounds
  - CH<sub>3</sub>-O-C<sub>3</sub>H<sub>7</sub> and C<sub>2</sub>H<sub>5</sub>-O-C<sub>2</sub>H<sub>5</sub> exhibit
  - a) metamerism
- b) chain isomerism
- c) optical isomerism
- d) cis-trans isomerism
- 25. Molecular formula C<sub>4</sub>H<sub>10</sub>O has ...... isomeric ethers,
  - a) 4
- b) 3
- c) 7

- d) 5
- 26. The compound which is not isomeric with diethyl ether is
  - a) butan-1-ol
  - b) 2-methyl propan-2-ol
  - c) butanone
  - d) n-propyl methyl ether
- 27. Diethyl ether exhibits metamerism with
  - a) CH<sub>3</sub>OCH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>b) CH<sub>3</sub>OCH(CH<sub>3</sub>)<sub>2</sub>
  - c) both 'a' and 'b'
- d) CH,COC,H,
- 28. Ethers are isomeric with
  - a) aldehydes
- b) alcohols
- c) acids
- d) ketones
- 29. CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>OH is functional isomer of
  - a) ethyl methyl ether
  - b) ethyl n-propyl ether

- c) methyl n-propyl ether
- d) 2-propanol
- 30. The compound which is functional isomer of diethyl ether is,
  - a) 1-methoxy propane
  - b) 1-butanol
  - c) 2-methoxy propane
  - d) both 'a' and' c'
- 31. The compound is not isomeric with diethyl ether
  - a) 1-butanol
  - b) 2-propanol
  - c) 2-methyl 2-propanol
  - d) 2-butanol
- 32. 1-methoxy propane and 2-methoxy propane are
  - a) position isomers
- b) chain isomers
- c) metamers
- d) functional isomers
- 33. Dimethyl ether is associated with which one of the isomer?
  - a) Ethanol
- b) Methanol
- c) Formic acid
- d) Methyl formate
- 34. Molecular formula C<sub>4</sub>H<sub>10</sub>O exhibits
  - a) chain isomerism
- b) position isomerism
- c) metamerism
- d) all of these
- 35. Which of the following compound shows metamerism?
  - a) CH<sub>2</sub>OCH<sub>2</sub>
- b) CH<sub>2</sub>OC<sub>2</sub>H<sub>7</sub>
- c) CH<sub>2</sub>OC<sub>2</sub>H<sub>5</sub>
- d) CH, COC, H,
- 36. Molecular formula C<sub>2</sub>H<sub>6</sub>O shows
  - a) functional isomerism
  - b) metamerism
  - c) position isomerism
  - d) optical isomerism
- 37. Diethyl ether can be distinguished from butan–1 vol by
  - a) aq. FeCl<sub>2</sub>
- b) Na metal
- c) Tollens reagent
- d) Fehling reagent
- 38.  $C_6H_5$ -O-CH<sub>3</sub> can be named as
  - a) phenoxy methane
- b) phenetole
- c) methoxy phenyl
- d) methoxy benzene
- 39. Ethers and alcohols are
  - a) metamers
- b) functional isomers
- c) tautomers
- d) position isomers
- 40. Which isomerism is not possible in ethers?
  - a) Tautomerism
- b) Chain isomerism
- c) Metamerism
- d) Position isomerism
- 41. Which of the following compound show

metamerism?

- a) CH<sub>3</sub>-O-CH<sub>3</sub>
- b) CH<sub>3</sub>-CO-CH<sub>3</sub>
- c) CH<sub>3</sub>NHCH<sub>3</sub>
- d) CH<sub>3</sub>-O-C<sub>3</sub>H<sub>7</sub>
- 42. Metamerism is advance type of
  - a) optical isomerism
  - b) geometrical isomerism
  - c) chain and position isomerism
  - d) only chain isomerism
- 43. Geometry of ether is
  - a) linear
  - b) pyramidal
  - c) trigonal planar
  - d) octahedral
- 44. Following compounds are



- a) functional isomers b) metamers
- c) chain isomers
- d) position isomers
- 45. Metamerism is possible in
  - a) same polyvalent functional group
  - b) same monovalent functional group
  - c) different polyvalent functional group
  - d) different monovalent functional group
- 46. Which of the following compound does not show metamerism?

  - a)  $CH_3$ –O– $CH_3$  b)  $CH_3$ –O– $C_4$   $H_9$

  - c)  $CH_{2}-O-C_{2}H_{7}$  d)  $CH_{3}-O-C_{5}H_{11}$
- 47. Following compound can be named as

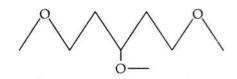
$$CH_3$$
-O- $(CH_2)_4CH_3$ 

- a) 3-methoxy pentaneb) 2-methoxy pentane
- c) 4-methoxy pentaned) 1-methoxy pentane
- 48. Following compounds are



- a) metamers
- b) position isomers
- c) chain isomers
- d) functional isomers
- 49. How many metamers are possible for molecular formula C<sub>5</sub>H<sub>12</sub>O
  - a) 4
- b) 6
- c) 8
- d) 10
- 50. How many ethers are possible for formula  $C_5H_{12}O$ 
  - a) 12
- b) 14
- c) 8
- d) 6

51. IUPAC name of following compound is



- a) 1, 3, 5-trimethoxy pentane
- b) 2, 4, 8 trioxa nonane
- c) 1, 3, 5 trimethoxy heptane
- d) 1, 3, 5 trimethoxy hexane
- 52. Hydride of ether is
  - a) aldehyde
- b) alcohol
- c) ketone
- d) carboxylic acid

### PREPARATION METHODS

- 53. The reaction of CH<sub>2</sub>CH<sub>2</sub>Br and (CH<sub>2</sub>)<sub>2</sub>CONa to form ether is called
  - a) Williamson reaction
  - b) Wurtz reaction
  - c) Cannizzaros reaction
  - d) Hoffmans reaction
- 54. Which of the following reaction does not form ether?
  - a) RX + aq. KOH
  - b) RX + RON a
  - c) CH,N, + ROH
  - d) ROH +  $H_2SO_4$  at 413 K
- 55. Reaction between sodium ethoxide and bromoethane forms
  - a) ethyl methyl ether b) diethyl ether
  - c) dimethyl ether
- d) acetic acid
- 56. Williamson's reaction is
  - a) SN<sup>1</sup> reaction of R-X
  - b) SN<sup>2</sup> reaction of R-X
  - c) SN<sup>2</sup> reaction of alkoxide
  - d) dehydration of R-X
- 57. In Williamson reaction intermediate formed is
  - a) carbocation
- b) free radical
- c) carbanion
- d) T.S
- 58. Select incorrect statement about following reaction of ether synthesis:

$$R-X + R' ONa \longrightarrow ROR' + NaX$$

- a) It follows S<sub>N</sub>2 mechanism
- b) Alkyl halide (RX) should be 2° or 3° while alkoxide (RO-Na<sup>+</sup>) should be 1°
- c) Alkyl halide should be 1° while alkoxide should be  $2^{\circ}$  or  $3^{\circ}$ .

- d) 2° and 3° alkyl halide may undergo E2 elimination in the presence of a strong base to form alkenes.
- Consider following reactions:

I. ONa + 
$$CH_3I$$

II. 
$$\bigcirc$$
 Br + CH<sub>3</sub>ONa  $\longrightarrow$ 

III. 
$$(CH_3)_3C-Cl + CH_3ONa \longrightarrow$$

$$IV.(CH_2)_2C-ONa + CH_2C1 \longrightarrow$$

Which of the above methods cannot be used in the synthesis of ether?

- a) I, IV
- b) I, III
- c) II, IV
- d) II, III
- 60. Catalytic dehydration of ethanol at 413 K gives
  - a) ethene
- b) ethoxy ethane
- c) ethane
- d) epoxy ethane
- 61. Preparation of ether from ethanol by continuous etherification process is
  - a) SN¹ reaction
- b) SN<sup>2</sup> reaction
- c) E<sup>1</sup> reaction
- d) E<sup>2</sup> reaction
- 62. Williamson's reaction of 30 alkyl halide is
  - a) SN¹ reaction
- b) Elimination reaction
- c) SN<sup>2</sup> reaction
- d) Reduction
- 63. Density of ether is
  - a) higher than water b) equal to water
  - c) lower than water
- d) can't be predicted
- 64. Sodium phenoxide is reacted with ethyl chloride gives
  - a) o-ethyl sodium phenoxide
  - b) p-ethyl sodium phenoxide
  - c) ethoxy benzene
  - d) m-ethyl sodium phenoxide
- 65. Methoxy benzene is prepared from CH<sub>2</sub>-Cl and what?
  - a) phenol
- b) sodium benzoate
- c) sodium phenoxide d) benzyl chloride
- 66. Which is the leaving group in the following reaction?

$$CH_3OH + CH_2N_2 \xrightarrow{HBF_4}$$

- a) H<sub>2</sub>O
- c) CH,
- d) H+

- 67. In the preparation of ether, one of the reactant is R-X another is
  - a) ale. R-ONa
- b) moist Ag<sub>2</sub>O
- c) CH,N,
- d) RCOONa
- 68. Select correct statement(s) about following reaction:

$$2R-OH \xrightarrow{H^+} R-O-R + H_2O$$

- a) It is an example of S<sub>N</sub> reaction in which protonated alcohol is the substrate and second molecule of alcohol is the nucleophile.
- b) It is intermolecular dehydration of alcohols
- c) This reaction can be S<sub>N</sub>2 if alcohol is 1°
- d) All of the above statements are correct.
- 69. Continuous etherification process based upon a) oxidation
  - b) intermolecular dehydration
  - c) intramolecular dehydration
  - d) reduction
- 70. In continuous etherification process 1st step is
  - a) formation of carbocation
  - b) protonation of alcohol
  - c) loss of proton from oxocation
  - d) cleavage of O-H bond in alcohol
- 71. Consider the following alkyl halides
  - 1) (CH<sub>3</sub>)<sub>3</sub>CBr
- 2) CH<sub>3</sub>Br
- 3) C<sub>2</sub>H<sub>5</sub>Br
- 4) CH, CHBrCH,

Arrange these alkyl halides in decreasing order of reactivity in Williamson reaction.

- a) 1 > 4 > 3 > 2
- b) 1 > 2 > 3 > 4
- c) 4 > 3 > 2 > 1
- d) 2 > 3 > 4 > 1
- 72. When ethyl hydrogen sulphate in heated with ethanol at 413 K, the product formed is
  - a) ethyne
- b) ethene
- c) diethyl ether
- d) diethyl sulphate
- 73. Diethyl ether is conveniently prepared in laboratory from
  - a) diazomethane
  - b) continuous etherification process
  - c) Williamson's synthesis
  - d) all of these
- 74. In Williamsons synthesis t–alkyl halide can not be used for preparation of alkyl t-butyl ether,
  - a) it is difficult to remove halogen atom
  - b) the reaction become reversible
  - c) it is not reactive

- d) it readily decompose to give olefin
- 75. Which of the following pair is used to prepare 2– ethoxy 2-methyl propane?
  - a)  $(CH_3)_3CC1 + C_2H_5ONa$
  - b)  $(CH_2)_2CONa + C_2H_5Cl$
  - c) CH<sub>2</sub>CH<sub>2</sub>ONa + C<sub>2</sub>H<sub>5</sub>Cl
  - d) CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>ONa + C<sub>2</sub>H<sub>2</sub>Cl
- 76. Ethyl iodide on treatment with sodium methoxide gives
  - a) CH<sub>3</sub>-O-CH<sub>3</sub>
- b) C,H,OCH,
- c) C<sub>2</sub>H<sub>5</sub>OC<sub>2</sub>H<sub>5</sub>
- d) C<sub>2</sub>H<sub>6</sub>
- 77. In the preparation of aromatic ether one of the reactant is sodium phenoxide, another is
  - a) R-OH
- b) R-Cl
- c) R-CHO
- d) R-COONa
- 78. In Williamson's synthesis
  - a) sodium alkoxide is treated with alkyl halide
  - b) sodium metal is treated with alkyl halide
  - c) an excess of alcohol is treated with conc. H<sub>2</sub>SO<sub>4</sub> at 413 K
  - d) vapours of alcohol are passed over heated Al<sub>2</sub>O<sub>3</sub> at 633 K
- 79. Excess of ethanol is heated with cone. H<sub>2</sub>SO<sub>4</sub> at 413 K. the compound that distills is
  - a) diethyl sulphate
  - b) diethyl ether
  - c) ethylene hydrogen sulphate
  - d) ethylene
- 80. 1-butanol is reacted with diazomethane to give,
  - a) 1-methoxy butane b) 2-methoxy butane

  - c) 1-ethoxy butane d) 2-ethoxy butane
- 81. Which of the following statement(s) is/are true about Williamsons synthesis?
  - a) It is desirable to use primary R-X
  - b) This method is particularly used for preparation of mixed ether
  - c) It is best to use the alkoxide of 2° and 3° alcohols
  - d) All of these
- 82. 2–ethoxy propane is effectively prepared from
  - a) isobutyl iodide
- b) isopropyl iodide
- c) ethyl iodide
- d) n-propyl iodide
- 83. The reaction between C<sub>2</sub>H<sub>5</sub>ONa + C<sub>2</sub>H<sub>5</sub>I to give C<sub>2</sub>H<sub>5</sub>OC<sub>2</sub>H<sub>5</sub> is called
  - a) Wurtz reaction
  - b) Kobles synthesis
  - c) Williamson synthesis
  - d) Hoffman reaction

- 84. The reaction of alkali alcoholate and monohalo alkane is called as,
  - a) Wurtz reaction
  - b) Cannizzaros reaction
  - c) Williamsons synthesis
  - d) Aldol condensation
- 85. Williamson's synthesis is used to prepare
  - a) diethyl ether
- b) amine
- c) ethanol
- d) ethanal
- 86. The intermediate product obtained during continuous etherification process is,
  - a) alkyl hydrogen sulphite
  - b) alkyl hydrogen sulphate
  - c) alkyl sulphate
  - d) alkyl sulphite
- 87. Reaction of t-butyl bromide with sodium methoxide produces
  - a) sodium tertiary butoxide
  - b) tertiary butyl methyl ether
  - c) isobutane
  - d) isobutylene
- 88. Excess of isopropyl alcohol is heated with cone, H<sub>2</sub>SO<sub>4</sub> at 413 K, gives
  - a) (CH<sub>3</sub>), CHOCH(CH<sub>3</sub>),
  - b)  $CH_{,}CH = CH_{,}$
  - c) both 'a' and 'b'
  - d) none of these
- 89. From Williamsons synthesis, which one of the following is most desirable to prepare ether?
  - a) 30 R-X and alkoxide of 1° alcohol
  - b) 3° R-X and alkoxide of 2° alcohol
  - c) 2° R-X and alkoxide of 1° alcohol
  - d) 1° R-X and alkoxide of 3° alcohol
- 90. Methoxy ethane is obtained by C<sub>2</sub>H<sub>5</sub>OH and what?
  - a) CH<sub>2</sub>Cl
- b) CH<sub>2</sub>ONa
- c) CH,N,
- d) CH,Cl,
- 91. Intermolecular dehydration of alcohol gives
  - a) Alkenes
- b) Ethers
- c) Alkynes
- d) Aldehydes
- 92. In the formation of ether, one of the compound is alcohol another is
  - a) R-ONa
- b) CH<sub>2</sub>N<sub>2</sub>
- c) R-OAg
- d) R-OK
- 93. The reaction given below is known as  $C_2H_5ONa + IC_2H_5 \rightarrow C_3H_5OC_3H_5 + NaI$

- a) Kolbe's synthesis
- b) Wurtz's synthesis
- c) Williamson's synthesis
- d) Grignard's synthesis
- 94. Ether is prepared by
  - a) Williamson's synthesis
  - b) Wurtz's reaction
  - c) Fridel-Craft's reaction
  - d) Hoffman bromide reaction
- 95. When an alkyl halide is allowed to react with a sodium alkoxide the product most likely?
  - a) An aldehyde
- b) A ketone
- c) An ether
- d) A carboxylic acid

### PROPERTIES AND USES

- 96. When ethyl methyl ether is reacted with cold cone. HI gives ethanol and methyl iodide. The reaction proceeding through
  - a) SN1
- b) SN<sup>2</sup>
- c) E1
- d)  $E^2$
- 97. When t-butyl methyl ether is reacted with cold cone. HI to gives t-butyl iodide and methyl alcohol. The reaction proceeding through
  - a) SN1
- b) SN<sup>2</sup>
- c) E<sup>1</sup>
- d)  $E^2$
- 98. Halogenation of anisole is carried in the presence of catalyst
  - a) FeCl<sub>2</sub>
- b) AlCl,
- c) BF<sub>3</sub>
- d) CH,COOH
- 99. Bromination of anisole gives major product
  - a) o-bromoanisole
  - b) p-bromoanisole
  - c) m-bromoanisole
  - d) di-orthobromoanisole
- 100. Nitration of phenyl alkyl ether gives
  - a) o-nitro alkyl phenyl ether
  - b) p-nitro alkyl phenyl ether
  - c) mixture of ortho and para nitro phenyl alkyl ether
  - d) m-nitro alkyl phenyl ether
- 101. Product of following reaction is

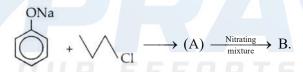


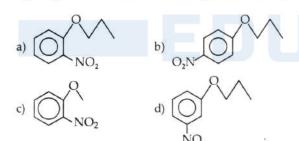
- b) CH, CHOHCH,
- c) C<sub>2</sub>H<sub>5</sub>OSO<sub>2</sub>H
- d) CH, CH, CH, OH
- 102. Which of the following ether produces methyl alcohol when reacts with cold HBr

- a) 2-Methyl 2-methoxy propane
- b) 2-methoxy propane
- c) 1-methoxy propane
- d) methoxy ethane
- 103. Two mole of alkyl iodide is formed when ether react with
  - a) Hot I,
- b) Cold HI
- c) Cold I,
- d) Hot HI
- 104. C-O bond in ether is not cleaved by
  - a) HCl
- b) Dil. H,SO4
- c) HBr
- d) HI
- 105. Acidic hydrolysis of ether gives
  - a) Two mole of aldehyde
  - b) One mole of alcohol
  - c) One mole of aldehyde
  - d) Two mole of alcohol
- 106. Reaction of dimethyl ether with cold HI is
  - a) E<sup>2</sup> reaction
- b) SN1 reaction
- c) E<sup>1</sup> reaction
- d) SN<sup>2</sup> reaction
- 107. Molecular formula(A)C<sub>4</sub>H<sub>10</sub>O on acid hydrolysis gives two mole of same alcohol. The compound A is
  - a) 2-methoxy propane
  - b) 1-methoxy propane
  - c) t-butyl alcohol
  - d) Oiethyl ether
- 108. Ethers are reacted with cold HI gives
  - a) One mole alcohol and one mole of alkyl iodide
  - b) Two mole alcohol
  - c) Two mole of alkyl iodide
  - d) Two mole alcohol and one mole of alkyl iodide
- 109. The reaction CH<sub>2</sub>OC<sub>2</sub>H<sub>5</sub> with cold HI gives

  - a)  $CH_2OH + C_2H_5I$  b)  $CH_2I + C_2H_5OH$
  - c)  $CH_{\lambda}I + C_{\lambda}H_{\xi}I$
- d) CH<sub>2</sub>OH + C<sub>2</sub>H<sub>5</sub>OH
- 110. Diethyl ether on heating with cone. HI gives two moles of
  - a) ethanol
- b) iodoform
- c) ethyl iodide
- d) methyl iodide.
- 111. Natalite is a mixtu re of,
  - a) diethyl ether and methanol
  - b) diethyl ether and ethanol
  - c) dimethyl ether and methanol
  - d) dimethyl ether and ethanol
- 112. Ethers are inactive because they do not contain
  - a) active atom
- b) active group
- c) multiple bond
- d) all of these

- 113. Diethyl ether with cold HI yields,
  - a) C<sub>2</sub>H<sub>5</sub>I
  - b) C<sub>2</sub>H<sub>5</sub>OH
  - c) C<sub>2</sub>H<sub>5</sub>I and C<sub>2</sub>H<sub>5</sub>OH
  - d) none of these
- 114. An hypothetical compound does not react with sodium metal. Which type of compound behave like this?
  - a) Alcohol
- b) Phenols
- c) Ethers
- d) Acid
- 115. The ethers heated with excess of HI gives three different products. The ether will be,
  - a) simple
  - b) mixed
  - c) either simple or mixed
  - d) unpredictable in nature
- 116. The compound which is mixed with alcohol to get a substitute for petrol is
  - a) ethanol
- b) diethyl ether
- c) acetaldehyde
- d) propanol
- 117. Oiethyl ether does not react with
  - a) dil.H,SO<sub>4</sub>
- b) HI
- c) CH,COOH
- d) PCl<sub>5</sub>
- 118. Find out major final product in following reaction.





119. The ether

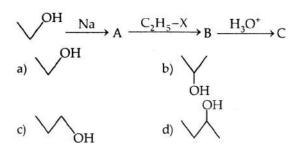
when reacted with cold HI gives







- a) mixture of 3 and 4 b) mixture of 1 and 2
- c) mixture of 2 and 3 d) mixture of 1 and 4
- 120. Diethyl ether can be regarded as unhydride of
  - a) CH<sub>3</sub>-OH
- b) C<sub>2</sub>H<sub>5</sub>-OH
- c) C<sub>2</sub>H<sub>5</sub>-COOH
- d) CH<sub>2</sub>COOH
- 121. Dimethyl ether can be decomposed by heating with
  - a) H<sub>2</sub>O
- b) NaOH
- c) KMnO<sub>4</sub>
- d) HI
- 122. The product (C) in the following reaction is



123. Following reaction is of the type

$$R-O-R' + HBr \xrightarrow{Cold} R-Br + R'-OH$$

If R' is 3° alkyl group and R is 1° alkyl group, then

- a) SN¹ with tertiary alkyl group
- b) SN<sup>2</sup> with primary alkyl group
- c) both of the above types
- d) none of the above types
- 124. The mixture of ethanol and H<sub>2</sub>SO<sub>4</sub> is distilled in distillation flask at 140°C. The flask would then contain
  - a) H<sub>2</sub>SO<sub>4</sub> and C<sub>2</sub>H<sub>5</sub>-O-C<sub>2</sub>H<sub>5</sub> only
  - b) H<sub>2</sub>O, H<sub>2</sub>SO<sub>4</sub> and C<sub>2</sub>H<sub>5</sub>OSO<sub>3</sub>H only
  - c) H<sub>2</sub>O, C<sub>2</sub>H<sub>5</sub>OSO<sub>3</sub>H, C<sub>2</sub>H<sub>5</sub>
  - d)  $C_2H_5$ –O– $C_2H_5$ ,  $C_2H_5OSO_3H$ ,  $H_2SO_4$  and  $H_2O$
- 125. Methoxy ethane does not react with
  - a) HI
- b) HBr
- c) dil. H<sub>2</sub>SO<sub>4</sub>
- d) Na
- 126. Which of the following will not form ether?

a) 
$$R-X + C_6H_5ONa \longrightarrow$$

- b)  $R X \xrightarrow{\text{moist Ag}_2O}$
- c)  $R-X + R ONa \longrightarrow$
- d)  $CH_2N_2 + R-OH \xrightarrow{HBF_3}$

- 127. Anisole is reacted with cold HI gives
  - a) Benzyl iodide and methanol
  - b) Phenol and methanol
  - c) Iodobenzene and iodomethane
  - d) Phenol and iodomethane
- 128. Reaction of CH, OCH, CH, is maximum with
  - a) HF
- b) HCl
- c) HBr
- d) HI
- 129. Ethyl phenyl ether is reacted with cold cone. HBr gives
  - a) phenol and ethane
  - b) bromobenzene and ethane
  - c) bromobenzene and bromoethane
  - d) phenol and bromoethane
- 130. HI cleaves ether (A) forming

$$A: \langle \bigcirc \rangle - CH_2 - O - CH_3$$

- CH,OH and CH,I
- CH,I and CH,OH
- CH,I and CH,I
- ⊢CH,OH and CH,OH
- 131. Anisole on treatement with Br<sub>2</sub> / CS<sub>2</sub> gives
  - a) bromobenzene
  - b) methyl 2-bromophenyl ether
  - c) o and p-bromoanisole
  - d) phenol
- 132. Which of the following is not cleaved by HI even at 525 K
  - a)  $C_6H_5-O-CH_3$  b)  $CH_3-O-C_2H_5$
  - c)  $C_6H_5 O C_2H_7$  d)  $C_6H_5 O C_6H_5$
- 133. Which is the most stable ion is formed in the protonation of (CH<sub>3</sub>)<sub>3</sub>C-O-CH<sub>2</sub>CH<sub>3</sub>

a) 
$$(CH_3)_3C - \overset{\oplus}{O} - CH_2CH_3$$
  
 $\overset{\parallel}{H}$ 

- b) (CH<sub>2</sub>),C<sup>+</sup>
- c) CH<sub>3</sub>-CH<sub>2</sub>+
- d) None of these
- 134. Which of the following reaction is not possible

a) 
$$C_2H_5$$
-O- $C_2H_5$   $\xrightarrow{Na}$ 

- b)  $R X + moist Ag_2O \longrightarrow$
- c)  $C_2H_5OH \xrightarrow{\text{cone. } H_2SO_4} \longrightarrow$
- d)  $C_2H_5$ -Cl  $\xrightarrow{\text{alc.KOH}}$
- 135. Methyl phenyl ether can be obtained by reacting
  - a) phenolate ions and methyl iodide
  - b) methoxide ion and bromobenzene
  - c) methanol and phenol
  - d) bromobenzene and methyl bromide
- 136. The major organic product in following reaction

$$\longrightarrow + \text{HI} \xrightarrow{\text{cold}}$$

- a) CH<sub>3</sub>I and
- b) CH<sub>3</sub>– I and
- c) CH<sub>2</sub>-OH and
- d) CH, OH and
- 137. Formation ether from ethanol based on
  - a) dehydration
- b) hydrogenation
- c) dehydrogenation
- d) hydration
- 138. In the reaction

which of the following compound will be formed?

- $\frac{1}{1}$  and and
- 139. Compound 'A' react with CH<sub>2</sub>-Cl gives B. B react with dil. H<sub>2</sub>SO<sub>4</sub> gives ethyl alcohol and CH<sub>3</sub>–OH.

The compound A is

- a)  $C_2H_5$ -OH
- b)  $CH_{2} = CH_{2}$
- c)  $CH \equiv CH$
- d) C<sub>2</sub>H<sub>5</sub>-ONa
- 140. In which of the following reaction product is t—butyl methyl ether

a) 
$$C_2H_5$$
-OH +  $C_2H_5$ -OH  $\xrightarrow{\text{conc. H}_2SO_4}$ 

b) 
$$(CH_3)_3 C - Br + CH_3ONa \longrightarrow$$

c) 
$$CH_3$$
-Br +  $(CH_3)_3$  C-ONa  $\longrightarrow$ 

d) 
$$(CH_3)_3$$
 C-Br +  $CH_3$ -OH  $\longrightarrow$ 

- 141. Which of the following compound when heated with HI gives two mole of different alkyl iodide?
  - a) C<sub>2</sub>H<sub>5</sub>-OH
- b) C<sub>2</sub>H<sub>5</sub>-O-C<sub>2</sub>H<sub>5</sub>
- c) CH<sub>3</sub>-O-C<sub>2</sub>H<sub>5</sub>
- d) CH<sub>3</sub>-O-CH<sub>3</sub>
- 142. Which of the following will gives good yield of ether?

a) 
$$\bigcirc$$
 OH  $\stackrel{i)}{\bigcirc}$  Na  $\stackrel{i)}{\bigcirc}$  OCH<sub>3</sub>

b) 
$$\bigcirc$$
 OH  $\xrightarrow{i)}$  Na  $\xrightarrow{ii)}$  CH<sub>3</sub>-1  $\bigcirc$  OCH<sub>3</sub>

c) 
$$\bigcirc$$
 OH  $\stackrel{i)}{\text{ii)}}$   $\stackrel{\text{Na}}{\text{CH}_3-\text{Cl}}$   $\bigcirc$  OCH<sub>3</sub>

d) 
$$\bigcirc$$
 OH  $\stackrel{\text{i) Na}}{\bigcirc}$   $\stackrel{\text{OCH}_3}{\bigcirc}$ 

- 143. Ethers can be prepared by
  - 1) heating alkyl halide with R-OH
  - 2) boiling alkyl halide with alc.KOH
  - 3) heating alkyl halide with sodium alkoxide
  - 4) reacting alcohol with diazomethane
  - a) 2,3,4
- b) 1,2,3
- c) 3,4
- d) 1,2
- 144. Product of following reaction will be









- 145. What is the function of diethyl ether in Grignards reagent preparation
  - 1) to act as a catalyst
  - 2) to act as a solvent
  - 3) to provide lone pair electron to co-ordination
  - 4) to act as an acid
  - a) 1, 2
- b) 2, 3
- c) 3, 4
- d) 2, 4
- 146. Phenetole react with cold HI gives

a) 
$$C_{5}H_{5} - I + C_{7}H_{5} - OH$$

b) 
$$C_{2}H_{5} - I + C_{6}H_{5} - OH$$

c) 
$$C_6H_5CH_2 - OH + C_2H_5 - I$$

d) 
$$C_6H_5 - OH + CH_3 - CH_2 - CH_2OH$$

- 147. Select the incorrect statement among the following
  - a) C-O-C bond angle in ether is 110°
  - b) ethoxy ethane is reacted with excess of HI gives ethyl iodide
  - c) ethers and alcohols are functional isomers
  - d) ethers are Lewis base hence do not react with Bronsted acid like H<sub>2</sub>SO<sub>4</sub>
- 148. Which of the following statement about ethers is/are incorrect?
  - 1) Ethers are very reactive
  - 2) Ethers are weakly acidic
  - 3) Ethers are Lewis base
  - 4) Ether form stable complex with Lewis acid
  - a) 1, 2, 3
- b) 1, 2
- c) 2, 3, 4
- $\frac{1}{2}$  4
- 149. CH<sub>3</sub>-O-C(CH<sub>3</sub>)<sub>3</sub> on reaction with dil. H<sub>2</sub>SO<sub>4</sub> under pressure gives (A) and (B). These are reacted with cold HI gives.

The compound A and B are respectively

- a)  $(CH_3)_3 C I$  and  $CH_3 I$
- b) (CH<sub>3</sub>)<sub>3</sub>C OH and CH<sub>3</sub>–I
- c)  $(CH_3)_3$  C I and  $CH_3$ –OH
- d) (CH<sub>3</sub>)<sub>3</sub> C -OH and CH<sub>3</sub>-OH
- 150. Conversion of alcohol to ether in the presence of conc. H<sub>2</sub>SO<sub>4</sub> is an example of
  - a) SN¹ reaction
- b) SN<sup>2</sup> reaction
- c) E<sup>1</sup> reaction
- d) E<sup>2</sup> reaction

151. 
$$CH_3 - C - CH_2O - CH_3$$

$$CH_3 - C - CH_2O - CH_3$$

prepared from Williamson reaction using

c) 
$$\sqrt{Cl}$$
 and  $ONa$ 

- d) both b and c
- 152. Which of the following is not usefull for the synthesis of ether?

a) 
$$\bigvee^X$$
 + R-ONa  $\longrightarrow$ 

b) 
$$CH_2N_2 + R-OH \xrightarrow{HBF_4}$$

c) 
$$+R-X \longrightarrow$$

- d) 2ROH \_\_\_\_cone. H<sub>2</sub>SO<sub>4</sub>/413K
- 153. Product of following reaction is

154. 
$$C_4H_{10}O + HI \xrightarrow{Hot} R_1 - I + R_2 - I$$

and

 $\rm R_1-I$  and  $\rm R_2$  –I on alkaline hydrolysis gives alcohols A and B respectively. The compound A and B gives haloform test. The original compounds is





- c) \
- d) OF
- 155. Which of the following can not be made hy Williamson's reaction?
  - a) ethoxy ethane
  - b) methoxy ethane
  - c) 1-methoxy propane
  - d) 2-(1, I-dimethyl ethoxy) propane
- 156. Some statements are given below about ethers,
  - 1. oxygen atom is sp<sup>3</sup>-hybridised
  - 2. they are liquids at room temperature
  - 3. they have higher boiling point than alcohols
  - 4. they are very active

Among the above, correct statement(s) is/are

- a) only 1
- b) only 3 and 4
- c) only 1 and 2
- d) all of these
- 157. Which one of the following is incorrect about dimethyl ether?
  - a) It has boiling point lower than alcohol
  - b) It is symmetrical ether
  - c) Its boiling point is more than ethanol
  - d) On treating with hot HI give single product
- 158. Diethyl ether finds its use in medicine as
  - a) anaesthetic
- b) antiseptic
- c) hypnotic
- d) pain killer
- 159. Ethers form co-ordination complexes with
  - a) BF,
- b) AlCl,
- c) ZnCl<sub>2</sub>
- d) all of these
- 160. Ethers on hydrolysis yield
  - a) ketone
- b) acid
- c) alcohol
- d) aldehyde
- 161. Ethers are basic in nature owing to the presence of
  - a) unshared electron pairs on oxygen
  - b) alkyl group
  - c) turn blue limus red
  - d) all of these
- 162. Ethers with cone. HI at low temperature form
  - a) alkyl iodide
- b) alcohol
- c) Both a and b
- d) oxonium salt
- 163. When ethyl isopropyl ether is reacted with HI in cold gives,
  - a) C<sub>2</sub>H<sub>5</sub>I and (CH<sub>2</sub>)<sub>2</sub>CHOH
  - b) C<sub>2</sub>H<sub>5</sub>I and (CH<sub>3</sub>)<sub>2</sub>CHI

- c) C<sub>2</sub>H<sub>5</sub>OH and (CH<sub>3</sub>)<sub>2</sub>CHI
- d) C<sub>2</sub>H<sub>5</sub>OH and (CH<sub>3</sub>)<sub>2</sub>CHOH
- 164. Some statements are given below about ethers,
  - 1. with strong acid, forms oxonium salt
  - 2. mixture of diethyl ether and ethanol called as nata lite
  - 3. t-butyl halide and sodium ethoxide give ethyl t-butyl ether
  - 4. these are acids

Among the above, false statement(s) is/are

- a) only 1 and 3
- b) only 2 and 4
- c) only 3 and 4
- d) only 4
- 165. The compound that does not react with sodium is
  - a) CH<sub>3</sub>OCH<sub>3</sub>
- b) CH<sub>2</sub>COOH
- c) CH,CHOHCH,
- d) C<sub>2</sub>H<sub>5</sub>OH
- 166. Which of the following ether will give two successive members of homologous series, on acid hydrolysis?
  - a) Oiethyl ether
  - b) Dimethyl ether
  - c) Ethyl methyl ether
  - d) Methyl n-propyl ether
- 167. The reaction of ether with HI is preferred than HBr because
  - a) HI is a stronger acid than HBr
  - b) HI gives a higher concentration of oxonium ion
  - c) I– is a better nucleophile in SN2 reaction than the Br– ion
  - d) all of these
- 168. Find out process involved in the following reaction,

$$R-O-R+H_2O \xrightarrow{\text{dil. } H_2SO_4 \Delta \atop \text{under press}} ROH + ROH$$

- a) hydrolysis
- b) oxidation
- c) hydration
- d) reduction
- 169. The central oxygen atom in ether is
  - a) sp<sup>2</sup>–hybridised
- b) sp<sup>3</sup>–hybridised
- c) sp-hybridised
- d) dsp<sup>2</sup>-hybridised
- 170. Which of the following statements is false in case of ethoxy ethane?
  - a) It is inflammable
  - b) It is simple ether
  - c) It react with Na metal
  - d) It is used as anaesthetic
- 171. Ether does not react with,
  - a) sodium metal
  - b) sodium hydroxide
  - c) phosphorus trichloride

- d) all of these
- 172. Isobutyl n–butyl ether is reacted with cold conc. HI gives,
  - a)  $CH_3CH_2CH_2CH_2I + (CH_3)_2CHCH_2OH$
  - b)  $CH_3CH_2CH_2CH_2OH + (CH_3)_2CHCH_2I$
  - c)  $CH_3CH_2CH_2CH_2I + (CH_3)_2CHCH_2I$
  - d)  $CH_3CH_2CH_2CH_2OH + (CH_3)_2CHCH_2OH$
- 173. The reaction of CH<sub>3</sub>OCH<sub>2</sub>CH<sub>3</sub> with hot excess HI gives
  - a) CH<sub>3</sub>OH + CH<sub>3</sub>CH<sub>2</sub>I
  - b) CH<sub>3</sub>I + CH<sub>3</sub>CH<sub>2</sub>OH
  - c) CH,OH + CH,CH,OH
  - d)  $CH_3I + CH_3CH_2I$
- 174. Ethers are mainly used as,
  - a) solvent
- b) cooling agent
- c) anaesthetic
- d) substitute of petrol
- 175. A temperature of -110°C can be obtained by using,
  - a) ether and CO,
  - b) ether and solid CO,
  - c) acetone and CO,
  - d) acetone and solid CO,
- 176. The cleavage of an ethyl methyl ether with cold hydrogen iodide will give
  - a) a molecule each of an methyl iodide and water
  - b) a molecule each of an ethyl iodide and water
  - c) a molecule each of ethanol and an methyl iodide
  - d) a molecule each of an ethyl iodide, methyl iodide and water
- 177. Which of the following is a gas at room temperature?
  - a) CH<sub>3</sub>OCH<sub>3</sub>
- b) HCHO
- c) Both 'a' and 'b'
- d) CH, COCH,
- 178. Some statements are given below about ethers,
  - 1. they are Lewis bases
  - 2. their boiling point increases with increasing molecular weight
  - 3. all are volatile liquids at room temperature
  - 4. with water dimethyl ether form hydrogen bond Among the above, false statement(s) is/are
  - a) only 4
- b) only 3 and 4
- c) only 3
- d) none of these
- 179. Ether  $\xrightarrow{\text{dil. H}_2\text{SO}_4}$  two products.

One of the product on oxidation give acetic acid, while the other on oxidation give acetone. The ether is,

- a) ethyl n-propyl ether
- b) ethyl methyl ether
- c) ethyl isopropyl ether
- d) methyl isopropyl ether
- 180. Some statements are given below about diethyl ether,
  - 1. its boiling point lower than J-butanol
  - 2. it is used as anaesthetic
  - 3. with dilute H<sub>2</sub>SO<sub>4</sub>, it give two homologue
  - 4. with cold HI give iodoethane

Among the above, correct statement(s) is/are

- a) only 1
- b) only 1 and 2
- c) only 2
- d) only 1, 2 and 4
- 181. The number of bond pair and lone pair on oxygen atom in ether are respectively
  - a) 1 and 2
- b) 2 and 1
- c) 2 and 2
- d) 1 and 3
- 182. Diethyl ether can be decomposed by heating with
  - a) NaOH
- b) KMnO<sub>4</sub> solution
- c) Water
- d) HI
- 183. Which of the following is most suitable reagent to distinguish ether from alcohol?
  - a) Na metal
- b) HI
- c) HBr
- d) All of these
- 184. Which of the following is used in the preparation of RMgX?
  - a) Dimethyl ether b) Diethyl ether
  - c) Ethyl methyl ether d) Ethanol
- 185. Which of the following give methyl alcohol with cold HBr
  - a) CH<sub>3</sub>-O-CH<sub>2</sub>-CH<sub>2</sub>-CH<sub>3</sub>-CH<sub>3</sub>

$$\begin{array}{c} CH_{3} \\ \text{b) } CH_{3} - O - \begin{array}{c} CH_{3} \\ - C - CH_{3} \\ \end{array} \\ CH_{3} \end{array}$$

$$\begin{array}{c} CH_3 \\ \downarrow \\ c) CH_3 - O - CH_2 - CH - CH_3 \end{array}$$

d) 
$$CH_3 - O - CH - CH_2 - CH_3$$
  
 $CH_3$ 

- 186. On heating diethyl ether with conc.HI, 2 moles of which of the following is formed?
  - a) Ethanol
- b) Iodoform
- c) Ethyl iodide
- d) Methyl iodide

- 187. By the action of CH<sub>2</sub>I on sodium ethoxide, we
  - a) CH<sub>2</sub>COOCH<sub>2</sub>
- b) CH, COC, H,
- c) CH<sub>3</sub>OC<sub>2</sub>H<sub>5</sub>
- d) Ethyl acetate
- 188. Ethyl chloride is converted into diethyl ether by
  - a) Wurtz synthesis
  - b) Grignard reaction
  - c) Perkin's reaction
  - d) Willaimson's synthesis
- 189. Consider the following reaction

A + CH<sub>3</sub>ONa 
$$\stackrel{\Delta}{\longrightarrow}$$
 B  $\stackrel{\text{cold HI}}{\longrightarrow}$  C<sub>2</sub>H<sub>5</sub>-OH + CH<sub>2</sub>-I.

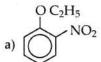
The compound A is

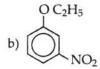
- a) C<sub>2</sub>H<sub>5</sub>-OH
- b) C<sub>2</sub>H<sub>5</sub>-CHO
- c)  $C_2H_5$ –Br
- d) C,H,
- 190. Which one of the following statement is not true regarding ether?
  - a) These are Lewis bases
  - b) They are highly inflammable.
  - c) They on acid hydrolysis gives alcohol.
  - d) These are Lewis acid
- 191. What are the product of following reaction?

$$C_2H_5OCH_3 \xrightarrow{Cold HI}$$

- a)  $C_2H_5OH + CH_3I$
- b)  $C_2H_3I + CH_3OH$
- c) C<sub>2</sub>H<sub>6</sub>OH + CH<sub>2</sub>OH
- d)  $C_2H_5I + CH_3I$
- 192. IUPAC name of ethyl isopropyl ether
  - a) 2-ethoxy propane
  - b) 1-ethoxy propane
  - c) 2-methyl 2-ethoxy propane
  - d) 1-methyl 2-ethoxy propane
- 193. Major product of following reaction is

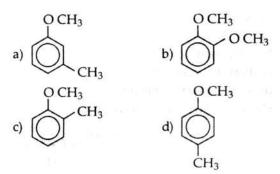
$$O-C_2H_5 + HNO_3 \xrightarrow{H_2SO_4}$$



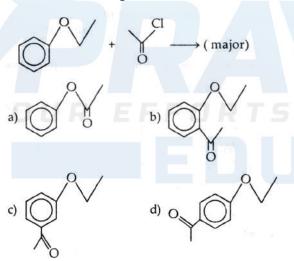




- 194. 4-alkoxy alkyl benzene is obtained from
  - a) Friedel Craft reaction
  - b) Ulmann reaction
  - c) Wurtz fittig reaction
  - d) Fittig reaction
- 195. Minor product obtained when anisole on methylation



- 196. Acylation of alkyl phenyl ether gives
  - a) 2-alkoxy alkyl phenyl ketone (major)
  - b) 4-alkoxy alkyl phenyl ketone (minor)
  - c) 4-alkoxy alkyl phenyl ketone (major)
  - d) 3-alkoxy alkyl phenyl ketone (major)
- 197. Product of following reaction is



198. 4-methoxy acetophenone is obtained from

- 199. Actully 18-crown-6 ether means,
  - a) 18-oxygen and 6 carbon
  - b) 12-carbon and 6-oxygen
  - c) 12-carbon and 12-oxygen
  - d) 6-oxygen and 6-carbon
- 200. 18-crown -6 ether is able to trap
  - a) K<sup>+</sup>
  - b) Na<sup>+</sup>
  - c) Li<sup>+</sup>
  - d) all of these
- 201. Which crown ether is used to extract cerium?
  - a) 15-crown-5
  - b) 18-crown-6
  - c) 12-crown -4
  - d) 10- crown-3
- 202. Diethyl ether is safe anesthetic agent. On administration it affect quickly to the central nerve system because
  - a) it is more soluble in fatty acid than water
  - b) it is more soluble in water than fatty acid
  - c) it is not soluble in fatty acid
  - d) it is highly inflammable
- 203. Which of the following is not used as anesthetic agent?
  - a) Diethyl ether
  - b) Nitrous oxide
  - c) Haloethane
  - d) Methanol
- 204. Lower molecular weight optically active ether is reacted with cold HI gives
  - a) butan-l-ol and iodomethane
  - b) butan-2-ol and iodomethane
  - c) propan-l-ol and iodoethane
  - d) propan-2-ol and iodoethane
- 205. Product of the following reaction is

$$\begin{array}{c} \mathrm{CH_3} - \mathrm{CH} - \mathrm{CH_2} - \mathrm{O} - \mathrm{C_2H_5} + \mathrm{HBr} \xrightarrow{\quad \Delta \quad} ? \\ \mathrm{OH} \end{array}$$

- a) C<sub>2</sub>H<sub>5</sub>Br and CH<sub>2</sub>CHBrCH<sub>2</sub>
- b) C<sub>2</sub>H<sub>e</sub>Br and CH<sub>2</sub>CHBrCH<sub>2</sub>Br
- c) C<sub>2</sub>H<sub>5</sub>Br and CH<sub>3</sub>CHOHCH<sub>2</sub>Br
- d) C<sub>2</sub>H<sub>5</sub>OH and CH<sub>3</sub>CHBrCH<sub>5</sub>Br
- 206. The reaction

$$\begin{array}{c} \operatorname{CH_3} \\ | \\ \operatorname{CH_3} - \operatorname{C} - \operatorname{ONa} + \operatorname{CH_3} \operatorname{CH_2} - \operatorname{Cl} \xrightarrow{-\operatorname{NaCl}} \\ | \\ \operatorname{CH_3} \end{array}$$

$$\begin{array}{c} \operatorname{CH_3} \\ \operatorname{CH_3} - \operatorname{C} - \operatorname{O} - \operatorname{CH_2} - \operatorname{CH_3} \\ | \\ \operatorname{CH_3} \end{array}$$

is called.

- a) Williamsons Synthesis
- b) Williamson continous etherification process
- c) Etard reaction
- d) Gatterman-Koch reaction

OOO

# **ANSWERS - MCQ'S**

# **SECTION - II : ETHERS**

1.	a)	2.	b)	3.	b)	4.	a)	5.	c)	6.	a)	7.	b)	8.	b)	9.	b)	10.	c)
11.	c)	12.	b)	13.	c)	14.	b)	15.	c)	16.	a)	17.	c)	18.	c)	19.	c)	20.	a)
21.	c)	22.	b)	23.	b)	24.	a)	25.	b)	26.	c)	27.	c)	28.	b)	29.	a)	30.	b)
31.	b)	32.	c)	33.	a)	34.	d)	35.	b)	36.	a)	37.	b)	38.	d)	39.	b)	40.	a)
41.	d)	42.	c)	43.	b)	44.	b)	45.	a)	46.	a)	47.	d)	48.	d)	49.	b)	50.	d)
51.	a)	52.	b)	53.	a)	54.	a)	55.	b)	56.	b)	57.	d)	58.	b)	59.	d)	60.	b)
61.	b)	62.	b)	63.	c)	64.	c)	65.	c)	66.	b)	67.	a)	68.	d)	69.	b)	70.	b)
71.	d)	72.	c)	73.	b)	74.	d)	75.	b)	76.	b)	77.	b)	78.	a)	79.	b)	80.	a)
81.	d)	82.	c)	83.	c)	84.	c)	85.	a)	86.	b)	87.	d)	88.	b)	89.	d)	90.	c)
91.	b)	92.	b)	93.	c)	94.	a)	95.	c)	96.	b)	97.	a)	98.	d)	99.	b)	100.	c)
101.	a)	102.	a)	103.	d)	104.	a)	105.	d)	106.	d)	107.	d)	108.	a)	109.	b)	110.	c)
111.	b)	112.	d)	113.	c)	114.	c)	115.	b)	116.	b)	117.	c)	118.	b)	119.	b)	120.	b)
121.	d)	122.	a)	123.	c)	124.	d)	125.	d)	126.	b)	127.	d)	128.	d)	129.	d)	130.	a)
131.	c)	132.	d)	133.	b)	134.	a)	135.	a)	136.	b)	137.	a)	138.	c)	139.	d)	140.	c)
141.	c)	142.	b)	143.	c)	144.	c)	145.	b)	146.	b)	147.	d)	148.	b)	149.	a)	150.	b)
151.	b)	152.	a)	153.	b)	154.	b)	155.	d)	156.	a)	157.	c)	158.	a)	159.	d)	160.	c)
161.	a)	162.	c)	163.	a)	164.	c)	165.	a)	166.	c)	167.	d)	168.	a)	169.	b)	170.	c)
171.	d)	172.	b)	173.	d)	174.	a)	175.	b)	176.	c)	177.	c)	178.	c)	179.	c)	180.	b)
181.	c)	182.	d)	183.	a)	184.	b)	185.	b)	186.	c)	187.	c)	188.	d)	189.	c)	190.	d)
191.	a)	192.	a)	193.	c)	194.	a)	195.	c)	196.	c)	197.	d)	198.	c)	199.	b)	200.	a)
201.	b)	202.	a)	203.	d)	204.	b)	205.	b)	206.	a)					l			



### MULTIPLE CHOICE QUESTIONS

### **Section - II: ETHERS**

- Fluoroboric acid is used as catalyst in preparation of ether from
  - a) sodium alkoxide
- b) diazomethane
- c) alkyl halide
- d) acetone
- Sodium iso-propoxide + A  $\xrightarrow{\Delta}$  2-ethoxy propane + NaCl

The compound A is

- a) C<sub>2</sub>H<sub>5</sub>ONa
- b) C<sub>2</sub>H<sub>5</sub>COCl
- c) C<sub>2</sub>H<sub>5</sub>Cl
- d) CH,ONa
- 3. Which of the following ether is not produced from methylation of alcohol?
  - a) CH<sub>3</sub>-O-CH<sub>3</sub>
- b) CH<sub>3</sub>OC<sub>2</sub>H<sub>5</sub>
- c) CH<sub>3</sub>-OCH(CH<sub>3</sub>), d) C<sub>2</sub>H<sub>5</sub>OC<sub>2</sub>H<sub>5</sub>
- 2-ethoxy propane is formed from ethyl bromide and what?
  - a) Sodium ethoxide
- b) Sodium iso-propoxide
- c) Iso-butyraldehyde d) Iso-propyl alcohol
- 5. CH<sub>2</sub>N<sub>2</sub> and 2-propanol gives

- a) (CH<sub>2</sub>)<sub>2</sub>CHOCH<sub>2</sub>
- b) (CH<sub>3</sub>),CHCOOH
- c) (CH<sub>3</sub>), NCH<sub>3</sub>
- d) (CH<sub>2</sub>), CH-H-CH<sub>2</sub>
- Which one of the following ether produces in higher yield by continuous etherification process?
  - a) C<sub>2</sub>H<sub>2</sub>OCH<sub>2</sub>
- b) C,H,OC,H,
- c) CH,OCH(CH,), d) C,H,OCH,CH,CH,
- Williamson's reaction is used in the preparation of
  - a) alcohols
- b) ethers
- c) aldehydes
- d) ketones
- $R-O-R' + HI \xrightarrow{cold} ROH + R' I$

If R contains three carbon atoms then how many carbon atoms are in R'?

- a) 2
- b) 4
- c) 5
- d) 6
- The reaction between alcohol and conc. H<sub>2</sub>SO<sub>4</sub> at 413 K gives
  - a) diethyl ether
- b) isopropyl alcohol
- c) diethyl alcohol
- d) ethene

COC

# ANSWERS - QUESTIONS ASKED IN MHT-CET

# **SECTION - II : ETHERS**

3. d) 7. b) 8. a) 9. a) 4. b) 5. a) 6. b)



### **MULTIPLE CHOICE QUESTIONS**

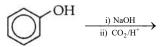
### Section - III: PHENOLS

- 1. In industry phenol is prepared from
  - 1. Raschig's method
  - 2. Dow's method
  - 3. Oxidation of cumene
  - 4. Oxidation of hexane
  - a) 1, 2
- b) 2, 3
- c) 3, 4
- d) 1, 2, 3
- 2. Replacement of N2<sup>+</sup> X<sup>-</sup> can be done by
  - a) H<sub>2</sub>O<sup>+</sup>
- b) aq. NaOH
- c) ale.KOH
- d) moist Ag<sub>2</sub>O
- 3. Carbolic acid is obtained from oxidation of
  - a) sodium salicylate
- b) salicylic acid
- c) toluene
- d) cumene
- 4. Cumene on air oxidation give
  - a) carbonic acid
- b) carbolic acid
- c) carboxylic acid
- d) oxalic acid
- 5. Phenol reacts with NaOH and followed by treatement with CH<sub>3</sub>–X gives
  - a) phenetole
- b) pheneyl acetate
- c) anisole
- d) toluene
- 6. Select incorrect statement
  - a) Reaction with Br<sub>2</sub> and water gives 2, 4, 6–tribromo phenol.
  - b) Reaction with dilute HNO<sub>3</sub> gives mixture of o-nitrophenol (minor) and p-nitrophenol (major).
  - c) Reaction with nitrating mixture gives picric acid.
  - d) Reaction with conc. H<sub>2</sub>SO<sub>4</sub> at 300 K gives ophenol sui phonic acid.
- 7. Consider the following species
  - 1) o-nitrophenol
  - 2) p-ni trophenol
  - 3) o-brorno phenol

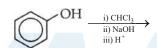
Intramolecular hydrogen bonding can takes place in

- a) only 3
- b) 1 and 3
- c) only 1
- d) 2 and 3
- 8. Which is steam volatile?
  - a) o-nitrophenol
- b) m-nitrophenol
- c) p-nitrophenol
- d) picric acid
- 9. When bromine is dissolved in CS<sub>2</sub> and treated with phenol gives major product
  - a) o-bromophenol

- b) p-brornophenol
- c) mixture of 'a' and 'b'
- d) 2, 4, 6-tribromophenol
- 10. Kolbes Schmidt reaction is used to prepare
  - a) salicylic acid
- b) salicylaldehyde
- c) phenyl acetate
- d) o-xylene
- 11. Reimer and Tiemann reaction is used to prepare
  - a) salicylic acid
- b) salicylaldehyde
- c) phenyl benzoate
- d) picric acid
- 12. Product of the following reaction is



- a) 2-hydroxy benzoic acid
- b) 2-hydroxy benzaldehyde
- c) sodium benzoate
- d) sodium phenoxide
- 13. Product of the following reaction is



- a) 2-hydroxy benzaldehyde
- b) 2-hydroxy benzoic acid·
- c) benzoic acid
- d) benzaldehyde
- 14. Benzene diazonium chloride is converted into phenol by
  - a) oxidation
- b) reduction
- c) neutralisation
- d) hydrolysis
- 15. Cresols have
  - a) 2-OH groups
- b) 4-OH groups
- c) 1-OH group
- d) 5-OH groups
- 16. Find out A in the following reaction

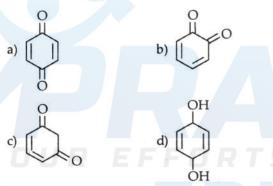
The compound A is

- a) o-bromo phenol
- b) 2,4,6 –tribrorno phenol
- c) p-bromo phenol
- d) m-bromo phenol
- 17. Benzene sulphonic acid is reacted with NaOH gives

- a) sodium phenoxide
- b) cumene
- c) cumene hydroperoxide
- d) sodium benzene sulphonate
- 18. Product obtained when steam is passed over chlorobenzene
  - a) sodium phenoxide
  - b) sodium benzene sulphonate
  - c) carbolic acid
  - d) benzene diazonium salt
- 19. Find out A and B in the following reaction respectively

$$Ph-Cl \xrightarrow{A} Ph-ONa \xrightarrow{B} Ph-OH$$

- a) NaOH and HCl
- b) H<sub>2</sub>O and HCl
- c) HCl and NaOH
- d) HCl and H<sub>2</sub>O
- 20. Phenol is
  - a) neutral
- b) amphoteric
- c) basic
- d) acidic
- 21. Phenol on oxidation by chromic acid gives



- 22. P-benzoquinone is obtained from phenol by
  - a) reduction
- b) oxidation
- c) acidic hydrolysis
- d) alkaline hydrolysis
- 23. Find out 'B' in the following reaction

$$+ H_2O \xrightarrow{SiO_2} A \xrightarrow{conc.H_2SO_4} B$$

$$SO_3H$$

$$b) \bigcirc OH$$

$$SO_3H$$

$$OH$$

$$OH$$

$$OH$$

$$OH$$

$$OOH$$

$$OOH$$

$$OOH$$

$$OOO_3H$$

- 24. Which of the following is/are steam volatile?
  - 1. p-nitrophenol
  - 2. o-nitrophenol
  - 3. o-hydroxy acetophenon
  - 4. p-hydroxy acetophenone
  - a) 1, 4
- b) 2, 3
- c) 1, 3
- d) 3, 4
- 25. The formation of salicylic acid from phenol using NaOH and CO<sub>2</sub> is known as
  - a) Friedel Craft reaction
  - b) Kolbe's-Schmidt reaction
  - c) Reimer and Tiemann reaction
  - d) Fittig reaction

26. Phenol 
$$\xrightarrow{Z_{\text{n}}}$$
 (A)  $\xrightarrow{\text{conc.H}_{2}\text{NO}_{3}}$  (B)

 $\xrightarrow{\text{Fe + HCl}}$  (C)

In above reaction, compound A, B, C are

- a) benzene, benzene sulphoric acid, aniline
- b) benzene, trinitrobenzene, aniline
- c) benzene dinitrobenzene, aniline
- d) benzene, nitrobenzene, aniline
- 27. In diazotisation reaction, carbolic acid is prepared from
  - a) cumene
- b) chlorobenzene
- c) aniline
- d) sod. phenoxide
- 28. Cumene is converted in phenol by
  - a) reduction and decomposition by acid
  - b) oxidation and decomposition by acid
  - c) reduction and decomposition by alkali
  - d) oxidation and decomposition by alkali
- Phenol on standing in air develop a red colour, due to formation of
  - a) cyclohexane
- b) phenoquinone
- c) resorcinol
- d) quinol
- 30. Benzene is obtained from phenol by using
  - a) Na metal
- b) Ca metal
- c) Zn metal
- d) NaOH
- 31. When phenol is reacted with CHCl<sub>3</sub> and NaOH, followed by treatment with LiAIH<sub>4</sub> gives
  - a) m-hydroxy methyl phenol
  - b) p-hydroxy methyl phenol
  - c) o-hydroxy methyl phenol
  - d) o-hydroxy methyl phenol
- 32. Reagent used in Reimer Tiemann reaction are
  - a) CH<sub>2</sub>Cl and aq.NaOH
  - b) CH<sub>3</sub>Cl and POCl<sub>3</sub>

- c) CHCl<sub>3</sub> and aq.NaOH
- d) CHCl, and alc.NaOH
- 33. Ka value of phenol is
  - a) More than carboxylic acid
  - b) Less than alcohol
  - c) More than alcohol
  - d) Less than water
- 34. Salicylic acid is prepared from phenol by the reaction known as
  - a) Wurtz reaction
  - b) Williamson reaction
  - c) Kolbes-Schmidt reaction
  - d) esterification
- 35. C–O bond length in phenol is less than c–o bond length in methyl alcohol because
  - a) partial double bond character due to resonance
  - b) partial double bond character due to inductive effect
  - c) more electronegativity of oxygen
  - d) oxygen contain two lone pair of electrons
- 36. The most suitable method of separation of ortho and para-nitrophenol mixed in the ratio of 1:1 is
  - a) steam distillation
- b) vapourisation
- c) crystallisation
- d) colour spectrum
- 37. The reaction of phenol with air. The product is
  - a) Anthraquinone
- b) Benzophenone
- c) Benzoquinone
- d) Propiophenone
- 38. Pieric acid contain
  - a) 2-nitro groups
- b) 3-nitro groups
- c) 2-nitrite groups
- d) 3-nitrite groups
- 39. Phenol is
  - a) a base weaker than ammonia
  - b) an acid stronger than carboxylic acid
  - c) an acid weaker than carboxylic acid
  - d) a neutral compound.
- 40. The synthesis of PhOH from PhCl is called
  - a) Cumene process
  - b) Dow's process
  - c) Williamson's synthesis
  - d) Wurtz synthesis
- 41. Phenol reacts with bromine in CS, at low temperature to give
  - a) m-bromophenol
  - b) p-bromophenol
  - c) o-and p-bromophenols
  - d) 2, 4, 6-tribromophenol

- 42. When phenol is treated with excess brominewater, it gives
  - a) m-bromophenol
  - b) o and p-brornophenols
  - c) 2, 4-dibromophenol
  - d) 2, 4, 6-tribromophenol
- 43. Which of the following reagents cannot be used to distinguish between phenol and alcohol?
  - a) Br<sub>2</sub>/CCl<sub>4</sub>
- b) NaOH
- c) NaHCO<sub>3</sub>
- d) neutral FeCl,
- 44. An organic compound with molecular formula C<sub>6</sub>H<sub>6</sub>O dissolves in NaOH and gives characteristics colour with neutral FeCl<sub>2</sub>. On treatment with bromine water, it gives tribromoderivative. The compound is,
  - a) alcohols
- b) ketones
- c) ethers
- d) phenol
- 45. In the nitration of phenol with a mixture of conc. HNO<sub>3</sub> and cone, H<sub>2</sub>SO<sub>4</sub>, the active species involved is
  - a) nitrite ion
- b) nitronium ion
- c) nitrate ion
- d) nitrogen peroxide
- 46. Carbolic acid is
  - a) C<sub>6</sub>H<sub>5</sub>CHO
- b)  $C_6H_6$
- c) C<sub>6</sub>H<sub>5</sub>COOH
- d) C<sub>6</sub>H<sub>5</sub>OH
- 47. Under different conditions nitration of phenol yields
  - a) o-nitrophenol
  - b) p-nitrophenol
  - c) 2, 4, 6-trinitro phenol
  - d) all of these
- 48. Picric acid is
  - a) a volatile liquid
  - b) trinitroaniline

  - c) 2, 4, 6–trinitrophenol
  - d) butyric acid
- 49. The end product in the following reaction is,

PhC1  $\xrightarrow{\text{H}_2\text{O} + \text{CuCl}_2}$  A  $\xrightarrow{\text{cone. HNO}_3}$  B

- a) PhOH
- b) PhBr
- c) PhNO,
- d) picric acid
- 50. Phenol is heated with conc. H<sub>2</sub>SO<sub>4</sub> at high temperature gives,
  - a) o-phenol sulphonic acid
  - b) p-phenol sui phonic acid
  - c) m-phenol sulphonic acid
  - d) all of these

- 51. Phenol gives violet colour with
  - a) neutral FeCl,
- b) neutral FeSO<sub>4</sub>
- c) acidic FeCl,
- d) acidic FeSO<sub>4</sub>
- 52. Picric acid is obtained by the nitration of
  - a) cumene
- b) phenol
- c) methanol
- d) ethanol
- 53. Acidic nature of phenol is due to
  - a) phenolic group
  - b) benzene group
  - c) hydrogen bonding
  - d) resonance stabilisation of phenoxide ion
- 54. Carbolic acid is reacted with conc. H<sub>2</sub>SO<sub>4</sub> at 300K gives,
  - a) 2-phenol sulphonic acid
  - b) 3-phenol sulphonic acid
  - c) 4-phenol sulphonic acid
  - d) 2 and 4-phenol sui phonic acid
- 55. Phenol reacts with Br, in CCl<sub>4</sub> at low temperature
  - a) o- and p- bromophenol
  - b) m-bromophenol
  - c) p-bromophenol
  - d) 2, 4, 6-tribromophenol
- 56. Which of the following is explosive?
  - a) Picric acid
- b) Methyl amine
- c) Cumene
- d) Ethanol
- 57. Nitrating mixture consists of
  - a) conc. HNO<sub>3</sub> + cone, HCl
  - b) cone. HNO<sub>3</sub> + cone. H<sub>2</sub>SO<sub>4</sub>
  - c) cone.  $H_2SO_4 + conc. H_3PO_4$
  - d) cone. HCl + conc. H<sub>2</sub>SO<sub>4</sub>
- 58. Cumene, is

  - a) phenyl n–propane b) 2–propyl benzene
  - c) chlorobenzene
- d) benzene
- 59. Sodium salt of benzene sulphonic acid on fusion with caustic soda and followed by treatment with HCl gives
  - a) acetic acid
- b) cumene
- c) phenol
- d) picric acid
- 60. Nitration of phenol is
  - a) nucleophilic substitution
  - b) electrophilic substitution
  - c) elimination
  - d) none of these
- 61. Phenol is ortho and para directing due to electron donating OH group, electron density increases

- at,
- a) ortho position
- b) para position
- c) meta position
- d) both 'a' and 'b'
- 62. 4-bromophenol is mainly formed, when phenol is reacted with,
  - a) Br<sub>2</sub>/water
- b) Br<sub>2</sub>/inert solvent
- c) HBr/water
- d) HBr/inert solvent
- 63. The number of a and n-bonds present in the molecule of carbolic acid are respectively
  - a) 7, 3
- b) 2, 3
- c) 4, 3
- d) 13, 3
- 64. During preparation of phenol from curnene, side product obtained is
  - a) acetone
- b) alcohol
- c) aldehyde
- d) acid
- 65. Suiphonation of phenol with conc. H<sub>2</sub>SO<sub>4</sub> at 288– 293 K gives
  - a) o-phenol sui phonic acid
  - b) p-phenol sulphonic acid
  - c) m-phenol sulphonic acid
  - d) all of these

66. 
$$\langle \bigcirc \rangle$$
 -CH(CH<sub>3</sub>)<sub>2</sub>  $\longrightarrow$   $\langle \bigcirc \rangle$  -OH

This change can be carried out by using

- a) reduction
- b) nascent oxygen
- c) air oxidation
- d) hydrolysis
- 67. OH A can be
  - a) SiO,
- b) SiO<sub>2</sub> / steam
- c) steam
- d) aq. KOH
- 68. For preparing monohalogen derivative of phenol, halogenation is carried out
  - a) at high temperature
  - b) at low temperature
  - c) inpresence of non-polar solvents
  - d) both 'b' and 'c'
- 69. 2-propyl benzene on air oxidation and followed by decomposition by dilute acid gives
  - a) phenol and propanal
  - b) phenol and propanone
  - c) phenol and propanol
  - d) phenol and propionic acid
- 70. The reaction

 $Ph-OH + dilute HNO_3 \longrightarrow ?$ 

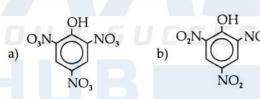
Gives predominately

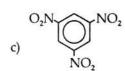
- a) 2-nitrocarbolic acid
- b) 4-nitrocarbolic acid
- c) 2-4-6 trinitro-carbolic acid
- d) 3-nitro carbolic acid
- 71. Some statements are given below about, carbolic acid
  - 1. it react with Na metal
  - 2. it gives violet colour with neutral FeCl,
  - 3. it forms only one monobrominated product
  - 4. it is acidic in nature.

Among the above, true statement(s) is / are

- a) only 4
- b) only 2 and 4
- c) only 1, 2 and 4
- d) all of these
- 72. During sulphonation, cone, H<sub>2</sub>SO<sub>4</sub> is used for
  - a) the introduction of -SO<sub>3</sub>H group in benzene
  - b) the introduction of -SO<sub>4</sub>H group in benzene
  - c) the introduction of -SO<sub>2</sub>H group in benzene
  - d) all of these
- 73. At different condition nitration of phenol gives
  - a) o-nitrophenol
- b) p-nitrophenol
- c) Picric acid
- d) All of these
- 74. How many O–H groups are present in phloroglucinol?
  - a) 3
- b) 2
- c) 4
- d) 5
- 75. Diazotisation reaction is used to prepare
  - a) alcohol
- b) phenol
- c) aldehyde
- d) ketone
- 76. The reaction of cone, HNO<sub>3</sub> and phenol forms
  - a) benzoic acid
  - b) salicylic acid
  - c) o- and p-nitrophenol
  - d) picric acid
- 77. At low temperature phenol reacts with  $\mathrm{Br_2}$  in  $\mathrm{CS_2}$  to form
  - a) m-bromophenol
  - b) o-and p-bromophenol
  - c) p-bromophenol
  - d) 2, 4, 6-tribromophenol
- 78. Picric acid is
  - a) trinitroaniline
- b) trinitrotoluene
- c) a volatile liquid
- d) 2, 4, 6–trinitrophenol
- 79. Chlorobenzene on fusing with solid NaOH gives
  - a) benzene
- b) benzoic acid
- c) phenol
- d) benzyl chloride
- 80. Bakelite plastic is formed, when phenol reacts

- with
- a) CH<sub>2</sub>CHO
- b) HCHO
- c) acetone
- d) HCOOH
- 81. Aromatic primary amine when treated with cold HNO<sub>2</sub> and HCl forms
  - a) benzene
- b) diazonium salt
- c) nitrobenzene
- d) benzyl alcohol
- 82. Phenol is treated with bromine water and shaken well. The white precipitate of which of the is formed
  - a) m-bromophenol
  - b) 2-4 dibromophenol
  - c) 2, 4, 6-tribromophenol
  - d) a mixture of o- and p-bromophenols
- 83. The bakelite is prepared by the reaction between
  - a) urea and formaldehyde
    - b) ethylene glycol
    - c) phenol and formaldehyde
    - d) tetramethylene glycol
- 84. Phenols are more acidic than aliphatic alcohols because
  - a) phenoxide ion is stabilised by resonance
  - b) phenols are more soluble in polar solvents
  - c) phenoxide ion do not have resonance
  - d) alcohols do not loose H-atom at all
- 85. Which of the following is explosive?





d) none of these

### ACIDIC NATURE

- 86. Phenol is less acidic than
  - a) p-nitro phenol
- b) cresol
- c) ethanol
- d) benzyl alcohol
- 87. Which of the following statement is correct?
  - 1. electron withdrawing groups stabilize the phenoxide ion and increase the acidic strength
  - 2. electron donating groups destabilise the phenoxide ion and decrease the acidic strength.
  - 3. -OH group in phenol is ortho and para

directing.

- 4. Intermolecular H–bonding is present in phenol
- a) 1, 3
- b) 1,2
- c)3,4
- d) 1,2,3,4
- 88. Phenol is
  - a) strong acidic
- b) weak acidic
- c) strong basic
- d) neutral
- 89. Which of the following group stabilise the phenoxide ion?
  - a) CH<sub>3</sub>
- b) OH
- c) OR
- d) NO,
- 90. Which of the following group destabalise the phenoxide ion?
  - a) NO,
- b) -CHO
- c) COR
- d) OH
- 91. Decreasing acidic nature of following compound is

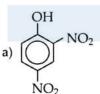




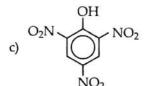




- a) 1 > 2 > 3 > 4
- b) 4 > 3 > 2 > 1
- c) 4 > 2 > 3 > 1
- d) 4 > 1 > 2 > 3
- 92. Which of the following is more acidic in nature?



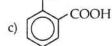




93. Which of the following has more pKa value?

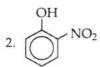


- c) OH CI
- d) OH
- 94. Which of the following has more Ka value?



- 95. Decreasing order of acidic nature of following compound is



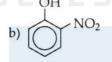






- a) 1 > 2 > 4 > 3
- b) 1 > 4 > 3 > 2
- c) 2 > 1 > 3 > 4
- d) 2 > 4 > 3 > 1
- 96. Which of the following is lowest Ka value?





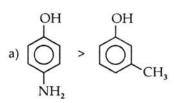




97. Decreasing order of acididy of substituted phenol is

- a) 3 > 1 > 4 > 2
- b) 4 > 3 > 1 > 2
- c) 4 > 3 > 2 > 1
- d) 3 > 1 > 2 > 4

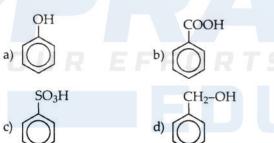
98. In which of the following first is more acidic than second?



c) 
$$OH$$
  $OH$   $OO_2$ 

d) 
$$\bigcirc$$
 >  $\bigcirc$  NO<sub>2</sub>

99. Which of the following is most acidic?



100. In the following compound order of acidity is



- 4. OH
- a) 4 > 2 > 3 > 1
- b) 4 > 3 > 1 > 2
- c) 4 > 3 > 2 > 1
- d) 4 > 1 > 2 > 3

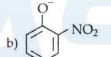
101. Which of the following is most acidic?

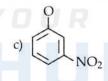


102. Which of the following is less acidic?

103. Which of the anion is most stable due to delocalisation?





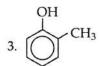




104. Which is the correct decreasing order of acidic strength of substituted phenol?









- a) 2 > 1 > 3 > 4
- b) 3 > 1 > 2 > 4
- c) 2 > 1 > 4 > 3
- d) 3 > 2 > 4 > 1

105. The correct acidic strength order of phenol i





a) 
$$1 > 2 > 4 > 3$$

b) 
$$3 > 1 > 4 > 2$$

c) 
$$1 > 2 > 3 > 4$$

d) 
$$3 > 1 > 2 > 4$$

106. Correct order of acidic nature i

a) 
$$1 > 3 > 2 > 4$$

b) 
$$3 > 4 > 2 > 1$$

c) 
$$1 > 3 > 4 > 2$$

d) 
$$3 > 4 > 1 > 2$$

107. Correct order of acidity of halogeneted phenol is where X–F, CI, Br, I

- a) 1 > 2 > 3 > 4
- b) 4 > 3 > 2 > 1
- c) 2 > 3 > 4 > 1
- d) 2 > 4 > 3 > 1

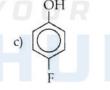
108. Which of the following is more acidic?

109. In which of the following first is more acidic than second?

a) 
$$\bigcirc$$
 OH OH NO<sub>2</sub> NO<sub>2</sub> NO<sub>2</sub>

c) 
$$NH_2$$
  $NH_2$ 

110. Which of the following is less acidic in nature?





 $\circ$ 

# **ANSWERS - MCQ'S**

# SECTION - III : PHENOLS

1.	d)	2.	a)	3.	d)	4.	b)	5.	c)	6.	b)	7.	c)	8.	a)	9.	b)	10.	a)
11.	b)	12.	a)	13.	a)	14.	d)	15.	c)	16.	b)	17.	d)	18.	c)	19.	a)	20.	d)
21.	a)	22.	b)	23.	c)	24.	b)	25.	b)	26.	d)	27.	c)	28.	b)	29.	b)	30.	c)
31.	c)	32.	c)	33.	c)	34.	c)	35.	a)	36.	a)	37.	c)	38.	b)	39.	c)	40.	b)
41.	c)	42.	d)	43.	c)	44.	d)	45.	b)	46.	d)	47.	d)	48.	c)	49.	d)	50.	b)
51.	a)	52.	b)	53.	d)	54.	a)	55.	a)	56.	a)	57.	b)	58.	b)	59.	c)	60.	b)
61.	d)	62.	b)	63.	d)	64.	a)	65.	a)	66.	c)	67.	b)	68.	d)	69.	b)	70.	a)
71.	c)	72.	a)	73.	d)	74.	a)	75.	b)	76.	d)	77.	b)	78.	d)	79.	c)	80.	b)
81.	b)	82.	c)	83.	c)	84.	a)	85.	b)	86.	a)	87.	d)	88.	b)	89.	d)	90.	d)
91.	b)	92.	c)	93.	a)	94.	c)	95.	c)	96.	d)	97.	a)	98.	d)	99.	c)	100.	b)
101.	a)	102.	b)	103.	d)	104.	c)	105.	b)	106.	c)	107.	c)	108.	c)	109.	b)	110.	d)



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### HINT & SOLUTIONS - MCQ'S

# Section I: Alcohols

- b) HOCH<sub>2</sub>CHOHCH<sub>2</sub>OH is a glycerol and trihydric alcohol.
- **4. b)** HOCH<sub>2</sub>ČH(OH)CH<sub>2</sub>OH is a glycerine. (\*) mark carbon atom is secondary carbon atom.
- 37. d) Ethers and alcohol are isomeric compounds. Molecular formula for diethyl ether and butanone are respectively C<sub>4</sub>H<sub>10</sub>O and C<sub>4</sub>H<sub>8</sub>O. Hence, butanone, a ketone, cannot be isomeric with ether.
- **39. b)** Molecular formula C<sub>2</sub>H<sub>6</sub>O are ethers and alcohols. i) C<sub>2</sub>H<sub>5</sub>OH, ii) CH<sub>3</sub>OCH<sub>3</sub>.
- 68. a) 1)  $CH_2=CH-CH_3+H_2O \xrightarrow{H^+} CH_3-CH(OH)-CH_3$  (Markownikoff addition)
  - 2)  $CH_3CHO \xrightarrow{CH_3MgI} CH_3-CH(OH)-CH_3$

Reactions (1) and (2) give 2-propanol.

**69. b)** Addition of water occurs according to Markownikoff's rule

$$CH_3CH_2CH = CH_2 \xrightarrow{H_2O / H_2SO_4} CH_3CH_2CHOHCH_3$$

- 70. a)  $CH_3COCH_3 + 2H \xrightarrow{NaHg + H_2O} (CH_3)_2CHOH$
- **71. d)** 2-methyl 2-pentanol is 3<sup>0</sup> alcohol prepared form acetone and n-propyl magnesium iodide.

$$\label{eq:h3COCH3+CH3CH2CH2MgI} \xrightarrow{\begin{subarray}{c} dry\ ether \\ $H_3O^+$\end{subarray}}$$
 
$$\begin{subarray}{c} (CH_3)_2C(OH)CH_2CH_2CH_3 + MgIOH \end{subarray}$$

- 72. a) CH<sub>2</sub>=CH<sub>2</sub>+H<sub>2</sub>SO<sub>4</sub> cold CH<sub>3</sub>CH<sub>2</sub>OSO<sub>3</sub>H
   CH<sub>3</sub>CH<sub>2</sub>OSO<sub>3</sub>H + HOH A CH<sub>3</sub>CH<sub>2</sub>OH + H<sub>2</sub>SO<sub>4</sub>
   Except ethanol, no other 1<sup>0</sup> alcohol can be prepared by this method as the addition of H<sub>2</sub>SO<sub>4</sub>, follows Markownikoffs of rule. Generaly 2<sup>0</sup> and 3<sup>0</sup> alcohols are obtained.
- 130. c)  $RCH_2CH_2OH \xrightarrow{H^+} RCH_2CH_2O^+H_2 \xrightarrow{RCH_2CH_2^+}$  protonation
- **132. d)** Increase in branching decreases surface area of the molecule and therefore, results in lesser

Van der Waal's forces. Consequently, isobutane has lesser boiling point than n-butane.

Alkyl chlorides have higher boiling than corresponding alkanes, because they are polar and stronger dipole-dipole force operate between the molecules of alkyl chloride.

Alcohols have higher boiling point than corresponding alkyl chloride, because alcohols molecules are associated through strong H-bonds.

- 133. b) More basic alcohol reacts faster with HX. Since basic nature decreases as 3<sup>0</sup>>2<sup>0</sup>>1<sup>0</sup>>MeOH. The same will be the decreasing order of reactivity of alcohol towards HX.
- 134. b) The relative rate follows same order as of acidity (HI > HBr >HCl>HF) and the nucleophilicity of their conjugate anion, X-(I<sup>-</sup>>Br<sup>-</sup>>Cl<sup>-</sup>>F<sup>-</sup>).
- 135. d) 2-methyl 2-propanol is tertiary alcohol, which forms tertiary carbocation (most stable carbocation)

$$(CH_3)_3C-OH+H^+ \longrightarrow (CH_3)_3C-O^+H_2 \longrightarrow (CH_3)_3C^++H_2O$$

 $(CH_3)_3C^+$   $\xrightarrow{Cl^-}$   $(CH_3)_3C$  -Cl

136. d) Reactivity of alcohol toward sodium depends upon the acidic nature of the alcohol. More the acidic nature of alcohol, the faster is the reaction between alcohol and sodium. The electron releasing alkyl groups decrease

the acidic nature of alcohol. Decreasing order of acidity of alcohol is  $CH_3OH > 1^0 > 2^0 > 3^0$ . The same is the order of reactivity towards sodium metal.

137. c) It is obtained according to Saytzeff rule

$$(C_2H_5)_2$$
CHCHOHCH $_3 \xrightarrow{\text{conc. } H_2SO_4} -H_2O \xrightarrow{}$ 

 $(C_2H_5)_2C=CH-CH_3$ 

138. a) The alcohol 4 and 2 are  $3^0$  alcohol but alcohol 4 gives more substituted alkene easily. Alcohol 1 is  $2^0$  alcohol and alcohol 3 is  $1^0$  alcohol. The rate of dehydration of alcohol with concentrated  $H_2SO_4$  follows the order  $3^0 > 2^0 > 1^0$ .

$$(CH_3)_2C=CH_2+4[O] \xrightarrow{K_2Cr_2O_7} CH_3COCH_3$$

 $+ CO_2 + H_2O$ 

- 140. d) The stability of carbonium ion depends on the nature of groups attached to the positively charged carbon atom.
  If the alkyl groups are attached to the positively charged carbon, the charge is dispersed by electron-donating inductive effect of alkyl group.
- 141. b) Alcohols have higher boiling points than non-polar hydrocarbon and polar haloalkane of comparable molecular masses, because alcohol molecules are associated through intermolecular hydrogen bonding.
   More the number of alkyl groups attached to the positively charged carbon, more the dispersal of positive charge and therefore, the greater is the stability of the carbonium ion. CH<sub>3</sub><sup>+</sup> < R<sub>2</sub>CH<sup>+</sup> < R<sub>3</sub>C<sup>+</sup>
- 142. a) In ethanol, H atoms is attached with oxygen; where as no hydrogen bonding occurs in diethyl ether, ethyl chloride and triethylamine, because no hydrogen is attached with oxygen or nitrogen.
- 143. b) 1-butanol is treated with alkaline KMnO<sub>4</sub> undergoes controlled oxidation gives butanal and when it is treated with strong oxidising agent it give directly butanoic acid.
- 144. b) Lucas reagent is the solution of anhydrous ZnCl<sub>2</sub> in conc. HCl.
   10 alcohols gives cloudiness not at all at room temperature
   20 alcohols gives cloudiness within five minutes.
   30 alcohols gives cloudiness immediately.
- **145. d)** It is a tertiary alcohol and hence, is most basic. Consequently, it reacts faster with HCl in presence of ZnCl<sub>2</sub> (Lucas reagent).
- **146.** c)  $(CH_3)_3COH \xrightarrow{Al_2O_3} (CH_3)_2C=CH_2+H_2O$
- **147. b)** Tertiary carbonium ion has maximum stability. The decreasing order of stability of carbonium ion is  $3^0 > 2^0 > 1^0$ .
- 148. d) Hydrogen attached to oxygen, nitrogen, sulphur, triply bonded-carbon etc. are called active hydrogen.
  Substance (a) and (b), contains 1 active hydrogen, (c) and (d) contains 2 and 3 active hydrogens respectively.

- 150. b) 1-butanol is treated with acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> oxidation gives butanoic acid and when treated with mild oxidising agent it give butanal.
- 151. d)  $3ROH + PX_3 \xrightarrow{\Delta} 3RX + H_3PO_3$   $ROH + PX_5 \xrightarrow{\Delta} RX + POX_3 + HX$  $ROH + HX \xrightarrow{\Delta} RX + H_2O$
- 152. a)  $(CH_3)_2C(OH)(C_2H_5) \xrightarrow{20\% H_2SO_4/363 K} (CH_3)_2C=CHCH_3+H_2O$
- 153. b) The increasing order of stabilities of carbonium ions is  $1^0 < 2^0 < 3^0$ . Since carbonium ion is formed as an intermediate in the dehydration of alcohol, so ease of dehydration in the increasing order is  $1^0 < 2^0 < 3^0$  alcohols.
- **154. a)** Addition of water occurs according to Markownikoff's rule

$$CH_3CH=CH_2 \xrightarrow{H_2O/H_2SO_4} CH_3CHOHCH_3$$

- 155. (c) Normal alcohols having higher boiling point than corresponding branched isomeric alcohols.
- 156. c) Because of absence of α-hydrogen atoms tertiary alcohols are resist oxidation. Compound (C) is t-butyl alcohol.
- of presence of hydrogen bonding. Normal alcohols have higher boiling point than isomeric branched alcohols. Compound (C) is normal propyl alcohol and compound (B) is branched propyl alcohol.
- 158. a)  $(CH_3)_2CHCH_2OH \xrightarrow{Al_2O_3} 425 \text{ K}$   $(CH_3)_2C=CH_2+H_2O$
- 159. c)  $C_2H_5OH + SOCl_2 \xrightarrow{pyridine} C_2H_5Cl + HCl + SO_2$
- 160. a)  $(CH_3)_2CHOH + [O] \xrightarrow{\text{oxidation}}$   $CH_3COCH_3 + H_2O$ Isopropyl alcohol first give acetone on
  - Isopropyl alcohol first give acetone on oxidation under drastic condition and then acetic acid.
- 161. d) Phenol is more acidic than all, because phenoxide ion is more stabilise by resonance than phenol.
- 162. b)  $CH_3CHOHC_2H_5 \xrightarrow{oxidation} CH_3COC_2H_5$
- **163. d)**. Solubility depends upon molecular weight. Solubility decreases with increasing molecular weight.

- carbon atoms, hence it is primary alcohol.

  190. a) Due to the three methyl groups which are electron donating, the carbon atom bearing OH group acquires partial negative charge, with the result it pushes the electron pair to the O-H bond towards hydrogen and thus, hydrogen atom is firmly attached with the oxygen atom, hence it does not replace easily by sodium metal.
- 191. a) Acetic acid exist as cyclic dimer due to intermolecular hydrogen bonding which requires high energy than hydrogen bonding present in alcohols.
- 192. a)  $ZnCl_2 + 2HCl \longrightarrow H_2ZnCl_4 \xrightarrow{ROH}$

Lewis acid

- 193. b) Major mode of C-C bond fission is according to Popoffs rule
- 194. b) Water is more acidic than alcohols because of minimum positive inductive effect of hydrogen atom.

 $R-O^+H_2 \xrightarrow{Cl^-} RCl + H_2O$ 

- **196.** a) It is 3<sup>0</sup> alcohol on dehydration gives alkene according to Saytzeff rule
- 197. b) We know that ethanol is soluble in water due to the hydrogen bonding. Therefore it is miscible with in all proportions.
- 198. d) We know that,
  CH<sub>3</sub>-CH<sub>2</sub>OH CH<sub>3</sub>CHO CH<sub>3</sub>COOH.
  Thus in this reaction, the final product is acetic acid (CH<sub>3</sub>COOH).
- 201. b) We know that,

CH<sub>3</sub>CH<sub>2</sub>OH  $\xrightarrow{\text{Cu}/573\,\text{K}}$  CH<sub>3</sub>CHO+H<sub>2</sub>. Thus primary alcohol which is ethyl alcohol (CH<sub>3</sub>CH<sub>2</sub>OH) is reduced by copper and forms acetaldehyde (CH<sub>3</sub>CHO).

- 202. c) We know that, the boiling point of a compound, evidently, is a consequence of strong intermolecular forces due to hydrogen bond. Therefore the boiling point of a compound is raised by intermolecular hydrogen bonding.
- 203. c) We know that ethanol is used as a beverage. It is known as spirit of wine or grain alcohol.
- 204. a) Methanol is also referred as wood alcohol or wood spirit or wood naphtha as the earliest method for its preparation was by destructive distillation of wood.

- to form hydrogen bond with water. More the tendency to form hydrogen bond with water more the solubility. In isomeric alcohols, tendency to form hydrogen bond with water is  $3^{0} > 2^{0} > 1^{0}$ .
- 206. a) The high boiling point of alcohols and amines is due to intermolecular hydrogen bonding. Alcohols have higher boiling point than amines, because of more electronegativity of oxygen than nitrogen. Carbonyl compounds have no hydrogen bonding. The dipole-dipole forces of carbonyl compounds causes them to have higher boiling points but less than compounds which have intermolecular hydrogen bonding.



## HINT & SOLUTIONS - MCQ'S

# Section - II: Ethers

- **24. a)** We know that metamerism is a type of isomerism exhibited by members of the same homologous series due to the difference in the nature of alkyl groups such as attached to the polyvalent atom of the functional group.
- 53. a)  $CH_3CH_2Br + (CH_3)_3CONa \xrightarrow{\Delta} CH_3CH_2OC(CH_3)_3 + NaBr$
- 54. a) RX + aq. KOH  $\xrightarrow{\Delta}$  ROH + KX RX + RONa  $\xrightarrow{\Delta}$  ROR + NaX CH<sub>2</sub>N<sub>2</sub> + ROH  $\xrightarrow{\Delta}$  ROCH<sub>3</sub> + N<sub>2</sub> 2ROH + H<sub>2</sub>SO<sub>4</sub>  $\xrightarrow{413K}$  ROR + H<sub>2</sub>SO<sub>4</sub> + H<sub>2</sub>O
- 55, **b)**  $C_2H_5ONa + C_2H_5Br \xrightarrow{\Delta} C_2H_5OC_2H_5 + NaBr$
- 71. d) The Williamson reaction involves nucleophilic substitution of alkoxide ion. It is strictly analogous to the preparation of alcohols by treatment of alkyl halide with aqueous NaOH. Alkyl halides undergoes substitution reaction most readily, t-butyl bromide are inert to nucleophilic substitution

because they show maximum steric hindrance to the approaching nucleophile. Thus, reactivity of alkyl halides towards Williamson reaction is,

 $1^0 RX > 2^0 RX > 3^0 RX$ .

- 72. c)  $CH_3CH_2OSO_2OH + HOCH_2CH_3 \longrightarrow CH_3CH_2OCH_2CH_3 + H_2SO_4$
- 73. b) Using dizomethane only methyl ethers are prepared, while using Williamson's synthesis simple and mixed ethers are prepared. However by using continuous etherification process only simple ether can be prepared.
- 110. c)  $CH_3CH_2OCH_2CH_3+2HI \longrightarrow 2CH_3CH_2I+H_2O$

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