

Class 12 Chapter 11 - Alcohols, Phenols and Ethers Important Questions with Answers

Short Answer Type Questions

Q1. What is the structure and IUPAC name of glycerol?

Answer:

Glycerol is trihydric alcohol. Structure of glycerol is

So, IUPAC name is — propan- 1, 2, 3-triol.

Q2. Write the IUPAC name of the following compounds.

Answer:

The IUPAC name for the compounds is given below.

- (a) 3-ethyl-5-methylhexane-2,4-diol
- (b) 1-methoxy-3-nitrocyclohexane
- **Q3.** Write the IUPAC name of the compound given below.



The IUPAC name of the compound given below is 3-methylpent-2-ene-1, 2-diol.

$$H_3C^5 - C^4H_2 - C^3 = C - OH$$
 $CH_3 CH_2 OH$

Q4. Name the factors responsible for the solubility of alcohols in water.

Answer:

- *Hydrogen bonds* Because of their propensity to create hydrogen bonds with water molecules or intermolecular hydrogen bonding, alcohols are soluble in water.
- Size of the alkyl or aryl groups The solubility of alcohols in water diminishes as the size of the alkyl or aryl groups grows larger. The presence of intermolecular hydrogen bonding between alcohol molecules makes low molecular mass alcohols liquid in water.
- Molecular mass of the Alcohols It reduces the effect of the polar character of the –OH group of alcohol with an increase in the alkyl group of alcohol or in the case of large molecular mass alcohols. As a result, as the molecular size of an alcohol increases, its solubility drops, therefore alcohols with lower molecular masses are more soluble than alcohols with larger molecular masses.

Q5. What is denatured alcohol?

Answer:

Denatured alcohol is ethanol (ethyl alcohol) that has been made unfit for human consumption by adding another chemical (mostly ethanol) to it. Denaturing refers to removing a property from the alcohol (being able to drink it), not to chemically altering or decomposing it. There are hundreds of ways ethanol is denatured. Denatured alcohol that is intended for use as a fuel or solvent typically contains 5% or more methanol.

Q6. Suggest a reagent for the following conversion.



$$\rightarrow$$

The given reactant is H₃C-CH=CH-CH(OH)-CH₃. It is a secondary alcohol. Secondary alcohol gives ketone when oxidised by CrO₃ or pyridinium chlorochromate without carrying out oxidation at the double bond.

OH
$$\begin{array}{c}
OH \\
4 \\
5
\end{array}$$
Pent-3-en-2-ol
$$\begin{array}{c}
OH \\
(C_6H_5N),CrO_3 \\
HCl
\end{array}$$
Pent-3-en-2-one

Q7. Out of 2-chloroethanol and ethanol which is more acidic and why?

Answer:

The acidity of 2-chloroethanol is higher than that of ethanol. The Cl-electron atom's density in the O-H bond reduces due to the -I effect (electron withdrawing group). As a result, the O-H bond of 2-chloroethanol weakens compared to the O-H bond of ethanol. As a result, 2-chloroethanol is acidic in comparison to ethanol.

$$\mathsf{CI} \leftarrow \mathsf{CH}_2 \leftarrow \mathsf{CH}_2 \leftarrow \mathsf{O} \leftarrow \mathsf{H} > \mathsf{CH}_3 \rightarrow \mathsf{CH}_2 \rightarrow \mathsf{O} \rightarrow \mathsf{H}$$

Stronger acid due to the -I effect of CI.

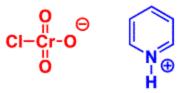
Q8. Suggest a reagent for conversion of ethanol to ethanal.

Answer:

Ethanol can be oxidized into ethanal by using pyridinium chlorochromate.







PCC (Pyridinium Chlorochromate)

Q9. Suggest a reagent for conversion of ethanol to ethanoic acid.

Answer:

Acidified KMnO₄ or $K_2Cr_2O_7$ can be used to convert ethanol to ethanoic acid. Both KMnO₄ and $K_2Cr_2O_7$ are powerful oxidizers.

Q10. Out of o-nitrophenol and p-nitrophenol, which is more volatile? Explain.

Answer:

o-nitrophenol is more volatile than p-nitrophenol due to presence of intramolecular hydrogen bonding. In para nitrophenol intermolecular hydrogen bonding is present. This intermolecular hydrogen bonding causes the association of molecules.

Q11. Out of o-nitrophenol and o-cresol which is more acidic?

Answer:

The presence of an electron-withdrawing group ($-NO_2$) in an ortho position relative to the -OH group increases the acidic strength of the chemical by stabilising the phenoxide ion, allowing o-nitrophenol to easily release a proton.

Because of the presence of an electron releasing group, o-cresol is less acidic (alkyl group). They prevent the production of the phenoxide ion. As a result, the compound's acidic strength is reduced.

Q12. When phenol is treated with bromine water, white precipitate is obtained. Give the structure and the name of the compound formed.



When phenol is treated with bromine water, white ppt, of 2,4,6- tribromophenol is obtained.

2,4,6-tribromophenol

Q13. Arrange the following compounds in increasing order of acidity and give a suitable explanation. Phenol, o-nitrophenol, o-cresol.

Answer:

Increasing order of acidity: o-cresol < phenol < o-nitrophenol

O-nitrophenol is a stronger acid than phenol due to the -I and -R effects of the NO_2 group, but o-cresol is a weaker acid due to the +I effect of the CH_3 group.

Q14. Alcohols react with active metals e.g. Na, K etc. to give corresponding alkoxides. Write down the decreasing order of reactivity of sodium metal towards primary, secondary and tertiary alcohols.

Answer:

The +I effect is produced by an electron releasing group, which raises the electron density on oxygen while decreasing the polarity of the O-H bond. The electron density on the O-H bond increases as the number of alkyl groups grows from one to three alcohols.

When going from one to three alcohols, the polarity will decrease and the strength of the O-H bond will grow.

As we all know, sodium metal is basic in nature, whereas alcohols are acidic. As a result, as the acidic strength lowers, the reactivity of alcohol with sodium metal reduces. As a result, the right sequence is 1>2>3.

Q15. What happens when benzene diazonium chloride is heated with water?



The phenol is formed. When benzene diazonium chloride is heated with water, it produces phenol as well as nitrogen gas and hydrochloric acid as by-products.

$$N = N - Cl$$

$$+ H_2O \xrightarrow{\text{dil.}} + N_2 + HCl$$

This process is frequently used to produce phenol from aniline. Diazonium salt is generated when aniline is treated with NaNO₂/HCI.

Q16. Arrange the following compounds in decreasing order of acidity. H₂O, ROH, HC=CH

Answer:

A stronger acid takes away the salt of a weaker acid. ROH is formed when RONa is handled with H_2O , hence water is a stronger acid than ROH. Acetylene is produced when sodium ethynide is handled with water and alcohol in the same way.

As a result, water and alcohol are more acidic than ethyne, and the decreasing order of acidity of given compounds is as follows:

Q17. Name the enzymes and write the reactions involved in the preparation of ethanol from sucrose by fermentation.

Answer:

Invertase and zymase are enzymes that are utilised in the fermentation process to convert sucrose to ethanol.

Sucrose is transformed to glucose and fructose by the enzyme invertase, and then glucose and fructose are turned to ethanol in the presence of zymase. The following is the response that occurs during the process:



$$C_{12}H_{22}O_{11} + H_2O \rightarrow C_6H_{12}O_6 + C_6H_{12}O_6$$

 $C_6H_{12}O_6 \rightarrow 2CH_3CH_2OH(Ethanol) + 2CO_2$

Q18. How can propan-2-one be converted into tert-butyl alcohol?

Answer:

Propan-2-one is a ketone when treated with Grignard reagents that give tertiary alcohols.



$$\begin{array}{c} \text{CH}_{3} \\ | \\ \text{CH}_{3} - \text{C} = \text{O} + \text{CH}_{3} \text{MgBr} \longrightarrow \\ \text{Acetone} \end{array} \xrightarrow{\begin{array}{c} \text{CH}_{3} \\ | \\ \text{CH}_{3} - \text{C} - \text{MgBr} \xrightarrow{\text{H'/H}_{2}\text{O}} \\ | \\ \text{CH}_{3} \\ \text{Addition product} \end{array} \xrightarrow{\begin{array}{c} \text{CH}_{3} \\ | \\ \text{CH}_{3} \\ \text{CH}_{3} \\ \text{OH} \end{array}} \xrightarrow{\text{OH}} \begin{array}{c} \text{OH} \\ \text{OH} \\$$

Q19. Write the structures of the isomers of alcohols with molecular formula $C_4H_{10}O$. Which of these exhibits optical activity?

Answer:

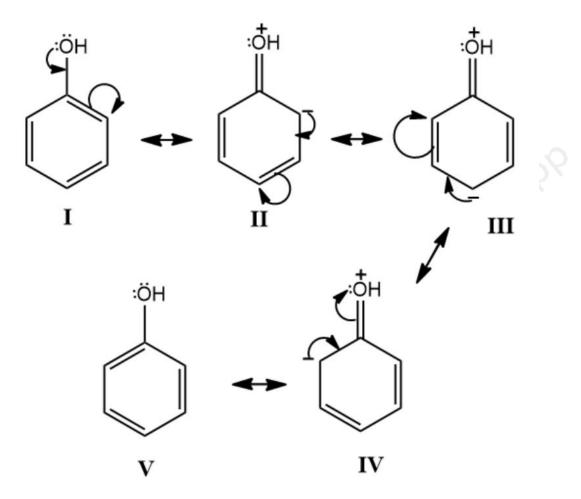
Four isomers of alcohols with molecular formula $C_4H_{10}O$ can be obtained. These isomers exhibit optical activity. They are

- 1. Butan-1-ol
- 2. 2-Methylpropan-1-ol
- 3. 2-methylpropan-2-ol
- 4. Butan-2-ol

Q20. Explain why the OH group in phenols is more strongly held as compared to OH group in alcohols.



The phenol –OH group is directly attached to the benzene ring's sp² hybridized carbon atom. The carbon-oxygen bond length in phenol is shorter than that in alkyl alcohol, which is owing to the partial double bond character of phenol or to the resonance and charge distribution in phenol. As a result, the OH group in phenols is retained more strongly than the OH group in alcohols.



Q21. Explain why nucleophilic substitution reactions are not very common in phenols.

Answer:

The OH group in phenols is a powerful electron donor. As a result, the benzene ring's electron density is quite high, and it repels nucleophiles. To put it another way, nucleophiles are unable to approach the benzene ring, hence phenols rarely undergo nucleophilic substitution reactions.

Q22. Preparation of alcohols from alkenes involves the electrophilic attack on alkene carbon atoms. Explain its mechanism.



To prepare alcohols from alkene in the presence of sulphuric acid through the process of hydration of alkenes.

Direct hydration takes place by adding water in the presence of a catalyst.

$$\begin{array}{c} \text{CH}_3 \\ \text{I} \\ \text{CH}_3 - \text{C} = \text{CH}_2 + \text{H}_2\text{O} \xrightarrow{\text{H}^{\oplus}} \text{(CH}_3)_3\text{C} - \overset{\oplus}{\text{OH}}_2 \xrightarrow{-\text{H}^{\oplus}} \text{(CH}_3)_3 - \text{C} - \text{OH} \\ \text{2-Methylpropane} \end{array}$$

Indirect hydration is achieved by the addition of sulphuric acid to alkane followed by hydrolysis of the alkyl hydrogen sulphate.

$$\begin{array}{cccc} \text{CH}_2 = \text{CH}_2 + \text{H}_2 \text{SO}_4 & \longrightarrow & \text{CH}_3 \text{CH}_2 \text{OSO}_2 \text{OH} & \xrightarrow{\text{H}_2 \text{O}} & \text{CH}_3 \text{CH}_2 \text{OH} \\ & & \text{Ethanol} & & & \text{Ethanol} \end{array}$$

Q23. Explain why is O=C=O nonpolar while R-O-R is polar.

Answer:

The dipole moments of the two C=O bonds are exactly equal and opposite, making O=C=O nonpolar. As a result, they cancel each other out, resulting in a net dipole moment of zero for O=C=O. Because the form of this compound is non-linear, the net dipole moment of R—O—R is not equal to zero, and hence R—O—R is polar in nature.

Q24. Why is the reactivity of all the three classes of alcohols with conc.HCl and ZnCl₂ (Lucas reagent) different?



Alcohols react with Lucas reagent (conc. HCl and ZnCl₂) using the SN1 mechanism. The stability of carbocations is required for the SN1 mechanism to occur (intermediate). The alcohol is more reactive when the intermediate carbocation is more stable.

Tertiary carbocations are most stable among the three classes of carbocations and the order of the stability of carbocation is 3°>2°>1°. This order, intrun, reflects the order of reactivity of three classes of alcohols i.e., 3°>2°>1°.

As a result of the differences in carbocation stability, the reactivity of all three types of alcohols with Lucas reagent varies.

Q25. Write steps to carry out the conversion of phenol to aspirin.

Answer:

The phenoxide ion is created by treating phenol with NaOH. Phenoxide ion then undergoes electrophilic substitution with CO2 to yield salicylic acid as the major product. This is called Kolbe's reaction. After that, the acetylation of salicylic acid yields aspirin, which is a significant product.

Q26. Nitration is an example of aromatic electrophilic substitution and its rate depends upon the group already present in the benzene ring. Out of benzene and phenol, which one is more easily nitrated and why?

Answer:

Nitration of benzene and phenol is an electrophilic substitution reaction. During nitration NO2 (nitronium ion) is produced as an intermediate as follows.



$$HO_3SO + HONO_2 \longrightarrow HO_2 + HSO_2$$
 $H + OO_2 \longrightarrow HOO_2 + HSO_2$
 $H + OO_2 \longrightarrow H_2O + OO_2$

(Protonated nitric acid) (Nitronium ion) (electrophile)

Phenol is more easily nitrated than benzene as the presence of — OH group in phenol increases the electron density at ortho and para positions in the benzene ring by + R effect. Since the electron density is more in phenol than in benzene, therefore, phenol is more easily nitrated than benzene.

Q27. In Kolbe's reaction, instead of phenol, phenoxide ion is treated with carbon dioxide. Why?

Answer:

Phenoxide ion is more reactive than phenol towards electrophilic aromatic substitution and hence undergo electrophilic substitution with carbon dioxide which is a weak electrophile.

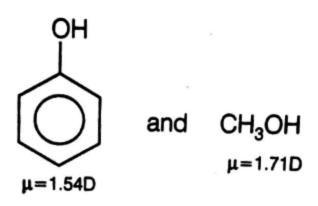
Q28. Dipole moment of phenol is smaller than that of methanol. Why?

Answer:

In phenol carbon is sp² hybridised and due to this reason the benzene ring is producing electron withdrawing effect.

On the other hand, carbon of methanol is sp³ hybridised and produces electron releasing effect (+ effect) Thus. C - O bond in phenol is less polar than the C - O bond in methanol and therefore, the dipole moment of phenol (1.54 D) is smaller than that of methanol (1.71 D).





Q29. Ethers can be prepared by Williamson synthesis in which an alkyl halide is reacted with sodium alkoxide. Di-tert-butyl ether can't be prepared by this method. Explain.

Answer:

In order to prepare di-tert- butyl ether, sodium tert-butoxide must be reacted with tert - butyl bromide. Alkoxides are not only nucleophiles but they are strong bases as well. They react with 3° alkyl halides leading to the elimination reaction.

When tert - butyl- bromide reacts with sodium tert-butoxide instead of substitution, elimination takes place. As a result of this elimination reaction, Isobutylene is formed instead of di - tert butyl ether.

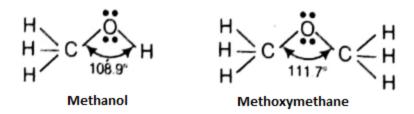
$$\begin{array}{c} \text{CH}_2-\text{H} & \text{CH}_3 & \text{CH}_2 & \text{CH}_3 \\ \text{H}_3\text{C}-\text{C}-\text{Br} & + \text{Na}-\text{O}-\text{C}-\text{CH}_3 & \text{Elimination} \\ \text{CH}_3 & \text{CH}_3 & \text{H}_3\text{C}-\text{C}-\text{CH}_3 + \text{H}_3\text{C}-\text{C}-\text{OH} + \text{NaBr} \\ \text{Iso-butylene} & \text{CH}_3 & \text{CH}_3 \\ \text{Tert-butyl bromide} & \text{Sodium tert butoxide} & \text{2--methylpropan -2-oi} \\ \\ \text{CH}_2-\text{H} & \text{CH}_3 & \text{CH}_3 & \text{Substitution} \\ \text{CH}_3 & \text{CH}_3 & \text{Substitution} \\ \text{CH}_3 & \text{CH}_3 & \text{CH}_3 \\ \end{array}$$

Q30. Why is the C-O-H bond angle in alcohols slightly less than the tetrahedral angle whereas the C-O-C bond angle ether is slightly greater?



The bond angle in C-:O:-H in alcohols is slightly less than tetrahedral angle (108.9). It is due to the repulsion between the unshared electron pairs of oxygen. In alcohols, two lone pairs of electrons are present. Therefore, there is comparatively more repulsion and less bond angle.

The C-O-C bond angle (111.7°) in ether is slightly greater than the tetrahedral angle due to the repulsive interaction between the two bulky (- R) groups.



Q31. Explain why low molecular mass alcohols are soluble in water.

Answer:

The solubility of alcohols and phenols in water is due to their ability to form hydrogen bonds with the water molecule. The hydrocarbon part methoxymethane (i.e., R group) tends to prevent the formation of hydrogen bonds.

Alcohols with lower molar mass will have smaller hydrocarbon parts and therefore tend to form hydrogen bonding more and they are more soluble in water.

Q32. Explain why p-nitrophenol is more acidic than phenol.

Answer:

Nitro group of phenol produces - I and - R effect. Because of these two effects -NO2 group is electron withdrawing in nature. So, the electron density in the O-H bond of p - nitrophenol decreases relative to the O-H bond of phenol.

The decrease in electron density of the O - H bond of p-nitrophenol, the polarity of O - H bond decreases. The electron withdrawing group (-NO), withdraws electrons and disperses the negative charge. Therefore, -NO group stabilises the phenoxide ion. Hence p-nitrophenol is more acidic than phenol.

Q33. Explain why alcohols and ethers of comparable molecular mass have different boiling points?



Because the ethers have a low polarity, they do not display any intermolecular hydrogen bonding.

Because breaking weak dipole-dipole forces in ethers requires less energy than breaking strong hydrogen bonds in alcohol.

$$CH_3CH_2 - O - H \cdots O - CH_2 - CH_3$$

Hydrogen Bond in alcohols.

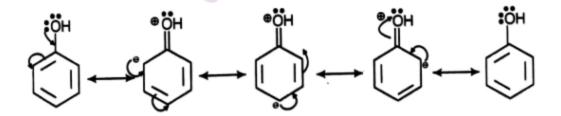
As a result, ethers have low boiling points, which are lower than isomeric alcohols and nearly equal to alkanes of comparable molecular weights. Alcohols and ethers have vastly different boiling points due to the presence of hydrogen bonding in alcohols.

Q34. The carbon-oxygen bond in phenol is slightly stronger than that in methanol. Why?

Answer:

The conjugation of unshared electron pairs over oxygen with the aromatic ring in phenol produces a partial double bond character in the C–O bond. This type of conjugation (resonance) is not feasible in methanol.

Oxygen is connected to sp² hybridised carbon in phenol, while oxygen is attached to sp³ hybridised carbon in methanol. The sp² hybridised carbon atom is more electronegative than the sp3 hybridised carbon atom (because of the higher 5-character). As a result, the connection formed by oxygen and sp² hybridised carbon is more stable than the one formed by oxygen and sp3 hybridised orbital.



The C - O bond of phenol gains some partial double bond character from the aforesaid resonating structure, whereas the C - O bond of methanol is strictly single bond.

Therefore, the carbon-oxygen bond in phenol is slightly stronger than that in methanol.



Q35. Arrange water, ethanol and phenol in increasing order of acidity and give reason for your answer.

Answer:

After removing a proton, the phenoxide ion is resonance stabilised in the following way.

The ethoxide ion formed after removing a proton, on the other hand, is not stabilised but destabilised due to the + I action of the $-C_2H_5$ group. As a result, phenol is a more powerful acid than ethanol.

$$C_2H_5 \rightarrow O-H$$
 $H-\ddot{O}-H$
Stabilised by resonance

Because the electron-releasing ethyl group increases the ethanol density of oxygen, ethanol is a weaker acid than water. As a result, the proton is not easily released. + In ethanol, the CH_3CH_2 group increases the electron density on the O - H bond. Ethanol has a higher electron density on its O - H bond than water. In the case of water, there is no such effect.

Thus, the increasing order of acidity is **ethanol < water < phenol**.

Long Answer Questions

Q1. Write the mechanism of the reaction of HI with methoxybenzene.

Answer:

In case of alkyl aryl ethers, the products are always phenol and an alkyl halide because due to resonance C6H5-O bond has partial double bond character. The mechanism is given below.

Mechanism Protonation of anisole gives methyl phenyl oxonium ion.



$$C_6H_5 - \overset{..}{O}CH_3 + H^+ \rightarrow C_6H_5 - \overset{+}{O}: -CH_3$$

In this ion, the bond between O-CH3 is weaker than the bond between

O-C6H5 which has a partial double bond character. This partial double bond character is due to the resonance between the lone pair of electrons on the 0 -atom and the sp2 hybridised carbon atom of the phenyl group. Therefore, attack by I- ion exclusively breaks the weaker O-CH3, bond forming methyl iodide and phenol.

- Q2. (a) Name the starting material used in the industrial preparation of phenol.
- (b) Write a complete reaction for the bromination of phenol in aqueous and non aqueous medium.
- (c) Explain why Lewis acid is not required in bromination of phenol?

Answer:

(a) The starting material used in the industrial preparation of phenol is cumene.

(b) Phenols when treated with bromine water give polyhalogen derivatives in which all the hydrogen atoms present at ortho and para positions with respect to the -OH group are replaced by bromine atoms.



Phenol

4-bromophenol

In aqueous solution, phenol ionises to form phenoxide ions. This ion activates the benzene ring to a very large extent and hence the substitution of halogen takes place at all three positions.

2-bromophenol

(c) In bromination of benzene, Lewis acid is used to polarise Br_2 to form the reactive electrophile, Br^+ In case of phenol, Lewis acid is not required because the O-atom of phenol itself polarises the Br_2 molecule to form Br^+ ions. further, +R-effect of OH group makes phenol highly activated towards electrophilic substitution reactions.

Mechanism of bromination of phenol.

$$OH$$
 OH
 $+Br-Br$
 OH
 Br
 $+Br$
 $+Br$
 $+Br$

Mechanism of bromination of benzene:

$$AlBr_3 + Br - Br \rightarrow [AlBr_4]^- + Br^+$$



$$H$$
 + Br⁺ H Br + [AlBr₄]⁻ H + HBr + AlBr₃

Q3. How can phenol be converted to aspirin?

Answer:

Allowing sodium phenoxide to absorb carbon dioxide and then heating the result to 400 K and 4-7 atm pressure is the typical procedure. The first unstable intermediate is generated, and it undergoes a proton shift to become sodium salicylate. Salicylic acid is produced as a result of the acidification of sodium salicylate.

As a result, phenol is transformed into salicylic acid. Acetic anhydride is used to treat this salicylic acid. Aspirin (acetylsalicylic acid) is generated when salicylic acid is acetylated.

Q4. Explain a process in which a biocatalyst is used in industrial preparation of a compound known to you.



Biocatalysts are enzymes. These biocatalysts (enzymes) are utilised in the production of ethanol in industry. Ethanol is made by fermenting molasses, a dark brown coloured syrup left behind after sugar crystallisation that still contains around 40% sugar.

Fermentation is the process of breaking down big molecules into smaller ones in the presence of enzymes. Yeast is the source of these enzymes. The numerous reactions that occur during carbohydrate fermentation are listed below.

$$C_{12}H_{22}O_{11} + H_2O$$
 Invertase $C_6H_{12}O_6 + C_6H_{12}O_6$
Sucrose Glucose Fructose $C_6H_{12}O_6$ $Zymase$ $2C_2H_5OH + 2CO_2$
Glucose or Ethanol carbon-di-oxide

Grapes are used to make wine because they contain sugars and yeast. The amount of sugar in grapes increases as they ripen, and yeast forms on the outer peel. When the sugar and enzyme in the grapes come into touch, fermentation begins. Fermentation occurs in anaerobic conditions, that is, without the presence of oxygen. During fermentation, CO₂ gas is produced.

Once the percentage of alcohol generated exceeds 14 percent, the action of zymase is blocked. If air is introduced into the fermenting mixture, oxygen oxidises ethanol to ethanoic acid, destroying the flavour of alcoholic beverages.