SUBJECT :- CHEMISTRY

JEE MAIN ANSWER KEY & SOLUTIONS

CLAS	S :- 12°						PAPER CODE :- CWT-8							
CHAF	CHAPTER :- ALCOHOL PHENOL ETHER													
	ANSWER KEY													
1.	(B)	2.	(C)	3.	(A)	4.	(A)	5.	(D)	6.	(C)	7.	(A)	
8.	(D)	9.	(B)	10.	(B)	11.	(B)	12.	(C)	13.	(D)	14.	(A)	
15.	(D)	16.	(D)	17.	(B)	18.	(B)	19.	(A)	20.	(D)	21.	58	
22.	2	23.	3	24.	3	25.	2	26.	4	27.	8	28.	6	
29.	4	30.	3											
SOLUTIONS														

1. (B)

Sol. (A) Fact



2. (C)

Sol. HBO \rightarrow Hydroboration oxidation OMDM \rightarrow Oxymercuration Demercuration.



3. (A)

Sol. Alcohols are soluble in water. This is due to the hydroxyl group in the alcohol which is able to form hydrogen bond with water molecules. Alcohol with a smaller hydrocarbon chains are very soluble. As the length of the hydrocarbon chain increases, the solubility in water decreases.



4. (A)

Sol. $CH_{3}-CH_{2}-CH_{2}-OH \xrightarrow{\text{oxidation}} CH_{3}-CH_{2}-COOH$ $(C_{3}H_{8}O) \qquad (C_{3}H_{6}O_{2})$ (Primary alcohol)

PRERNA EDUCATION

(D)



Sol.

More stabilised by intramolecular hydrogen bonding.



More strong intermolecular forces increase the boiling point.

6. (C)

Sol. The ether has a non-descript alkyl group 'R' on the one side and a triphenylmethyl group on the other side. The oxygen gets protonated and ROH is an excellent leaving group under these conditions. This is because the triphenylmethylium cation is super stable



Any factor that would improve the stability of the carbocation will make the hydrolysis faster! If any one of the phenyl groups is replaced by a methyl group, the loss of a phenyl group will be destabilizing the intermediate carbocation. This is because the methyl group's hyperconjugation is no match for the highly stabilizing phenyl ring (via conjugation).

Also, replacing a phenyl ring with a para-methoxyphenyl group will have a greater overall stabilizing effect on the intermediate carbocation. This is because the progressive increase in electron density through the carbon chain via the conjugation or resonance of the methoxy oxygen atom. Two such rings will make the reaction go fastest among the given options.





16. (D)
Sol.
$$(i) H_3PO_4, 150^{\circ}C$$
 $(i) C_{e}H_5-CO_3H$ $(i) C_{e}H_5-CO_3H$
17. (B)
1° Alcohol $\xrightarrow{PCC \text{ in } CH_2Cl_2}_{Mild \text{ oxidising agent}}$ Aldehyde
Sol. $(sarett reagent)$

18. (B)

Sol. The Williamson ether synthesis is an organicreaction used to convent an alcohol and an alkylhalide to ether using a base such as NaOH. Themechanism begins with the base abstracting theproton from the alcohol to form an alkoxideintermediate. The alkoxide then attacks thealkyl halide in a nucleophilic substitutionreaction (S_N2), which results in the formation of the final ether product and a metal halide by product.

Sol.
$$OCH_3 HO H$$

20. (D)

$$\begin{array}{ll} \mbox{Sol.} & A \rightarrow Q, S, T ; B \rightarrow S, T ; C \rightarrow P ; D \rightarrow R \\ & (A) CH_3 - CH_2 - CH_2 - CN \\ & \left[\begin{array}{c} Pd - C/H_2 \rightarrow CH_3 - CH_2 - CH_2 - CH_2 - NH_2 \\ \hline Sn Cl_2 / HCl \rightarrow CH_3 - CH_2 - CH_2 - CHO \\ \hline DIBAL - H \rightarrow CH_3 - CH_2 - CH_2 - CHO \\ \hline OH^- \rightarrow CH_3 - CH_2 - CH_2 - CHO \\ \hline OH^- \rightarrow CH_3 - CH_2 - CH_2 - OH \\ \hline DIBAL - H \rightarrow CH_3 - CH_2 - OH \\ \hline DIBAL - H \rightarrow CH_3 - CHO \\ \hline OH^- \rightarrow CH_3 - CHO \\ \hline OH^- \rightarrow CH_3 - CHO \\ \hline OH^- \rightarrow CH_3 - COO^- \\ & (C) CH_3 - CH = CH - CH_2 - OH \\ \hline DD CH_3 - CH_2 - CH_2 - NH_2 \\ \hline CHCl_3 \rightarrow CH_3 - CH_2 - CH_2 - CH_2 - OH \\ \hline CHCl_3 \rightarrow CH_3 - CH_2 - CH_2 - CH_2 - NC \\ \end{array}$$

21. 58

Sol. 58.00 g/mole









Br

(A)

26.

4



 H_2O



(X) Major product

The number of sp² hybridised carbon atom is 8

