

NEET ANSWER KEY & SOLUTIONS

SUBJECT :- CHEMISTRY

CLASS :- 12<sup>th</sup>

CHAPTER :- AI KYI HAI IDE

PAPER CODE :- CWT-7

SOLUTIONS

SECTION-A

- 1.** (D)

**Sol.**

$$\text{C}_2\text{H}_5\text{OH} \xrightarrow{\text{Red P} + \text{Br}_2} \text{C}_2\text{H}_5\text{Br}$$
$$\text{C}_2\text{H}_5\text{OH} \xrightarrow{\text{SOCl}_2} \text{C}_2\text{H}_5\text{Cl}$$
$$\text{C}_2\text{H}_5\text{OH} \xrightarrow{\text{KBr} + \text{Conc. H}_2\text{SO}_4} \text{C}_2\text{H}_5\text{Br}$$

All of the above leads to formation of an alkyl halide.

- 2. Sol.** (C) Darzen's reaction is

$$\text{ROH} + \text{SOCl}_2 \xrightarrow{\text{Pyridine}} \text{RCl} + \text{SO}_2 + \text{HCl}$$

- 3.** (A)

**Sol.**

$$\begin{array}{ccc}
 \text{COOAg} & & \text{Br} \\
 | & & | \\
 \text{C}_6\text{H}_5\text{O} & \xrightarrow[\text{Acetone}]{\text{Br}_2} & \text{C}_6\text{H}_5\text{OBr}
 \end{array}$$

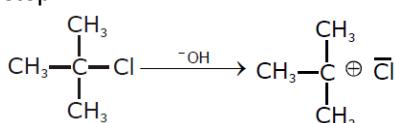
- 4.** (B)  
**Sol.** Polarity order of alkyl halides  
RF > RCl > RBr > RI because  
electronegative difference is decreasing  
F > Cl > Br > I

- 5.** (B)  
**Sol.** Finkelstein Reaction

$$\text{CH}_3\text{CH}_2\text{Br} + \text{NaI} \xrightarrow{\text{Acetone}} \text{CH}_3\text{CH}_2\text{I} + \text{NaBr}$$

- 6.** (B)  
**Sol.**  $\text{CH}_3\text{CH}_3\text{Cl} + \text{AgF} \rightarrow \text{CH}_3\text{CH}_2\text{F} + \text{AgCl}$   
This is Swarts reaction

- Sol.** In  $S_N1$ , formation of carbocation is the first step:



8. (D)

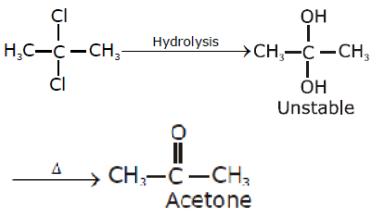
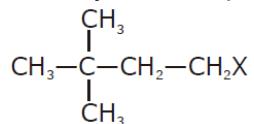
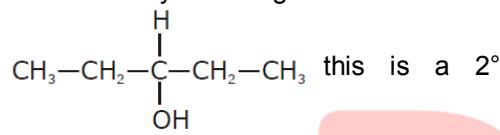
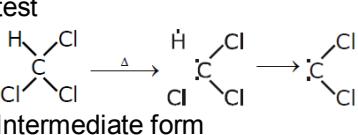
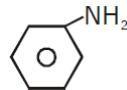
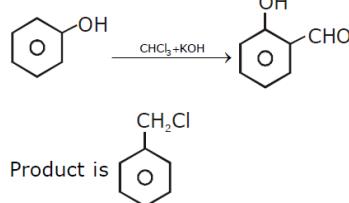
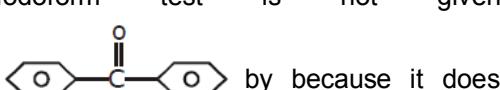
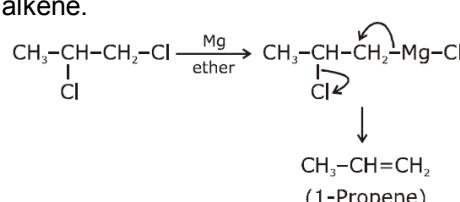
**Sol.** Chlorobenzene  is less reactive than  because substitution of Cl from chlorobenzene is difficult as compared to substitution of Cl from benzylchloride due to the formation of stable benzyl carbocation.

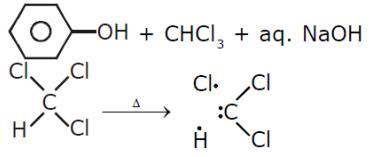
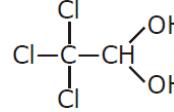
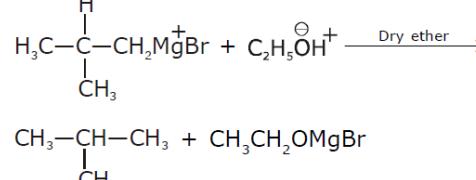
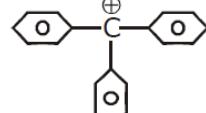
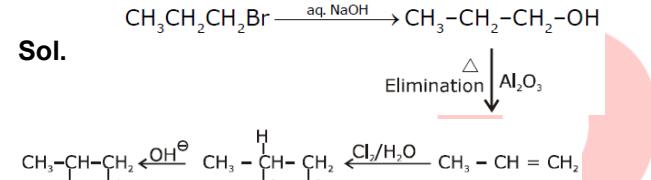
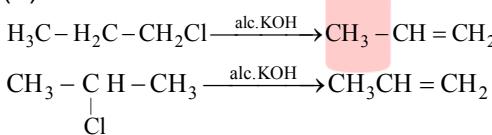
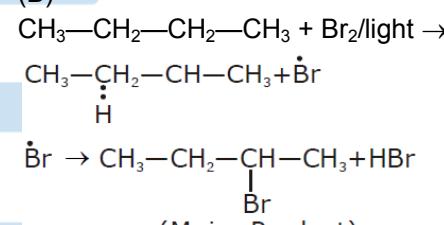
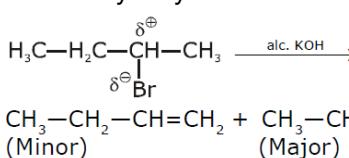
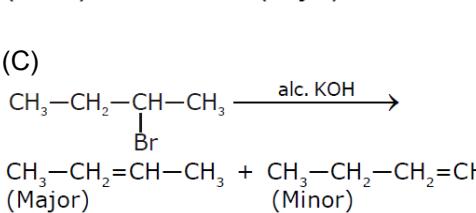
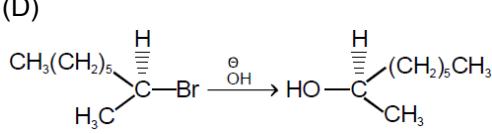
- 9.** (D)  
**Sol.** Vinyl halide are unreactive towards nucleophilic substitution because C-X bond develops a partial double bond character.





- Sol.**  $\text{CH}_3\text{CH}_3 \xrightarrow[\text{light}]{\text{Cl}_2} \text{CH}_3\text{CH}_2\text{OH} \xrightarrow{\text{oxidation}} \text{CH}_3\text{CHO}$

<p><b>12.</b> (B)  <b>Sol.</b> <math>\text{C}_2\text{H}_5\text{B} \xrightarrow[\text{H}_2\text{O}]{\text{moist Ag}_2\text{O}} \text{C}_2\text{H}_5\text{OH}</math> formed</p>	<p><b>20.</b> (B)  <b>Sol.</b> Catalyst used for preparation of <math>\text{CCl}_2\text{F}_2</math> dichlorodifluoromethane is <math>\text{SbCl}_5 + \text{HF}</math>.</p>
<p><b>13.</b> (A)  <b>Sol.</b> 2,2-Dichloropropane</p> 	<p><b>21.</b> (D)  <b>Sol.</b> Neohexyl halide is primary halide</p> 
<p><b>14.</b> (D)  <b>Sol.</b> Both geminal and vicinal dihalides when heated with zinc gives alkenes by losing both the halogen atoms. It is a dehalogenation reaction.</p>	<p><b>22.</b> (D)  <b>Sol.</b> Reactivity order of alcohols towards H-X will be II &gt; IV &gt; III &gt; I because II allylic, IV is 3° &amp; III is 1° while <math>\text{CH}_2 = \text{CH}_2 - \text{OH}</math> is vinylic which is least reactive</p>
<p><b>15.</b> (B)  <b>Sol.</b> Wrong Statement  * All secondary alcohol gives iodoform test</p>  <p>alcohol but it does not give iodoform test so it is wrong statement.</p>	<p><b>23.</b> (B)  <b>Sol.</b> Alkane isonitrile, <math>\text{RNC}</math>  <math>\text{RX} + \text{AgCN} \rightarrow \text{RNC}</math></p> <p><b>24.</b> (D)  <b>Sol.</b> <math>4\text{CH}_3 - \text{CH}_2 - \text{Cl} + 4\text{Na} + \text{Pb} \rightarrow \text{Pb}(\text{CH}_3\text{CH}_2)_4</math>, Tetra ethyl lead used as antiknocking agent.</p>
<p><b>16.</b> (B)  <b>Sol.</b> Ethanol is added to prevent oxidation of chloroform. Ethanol also converts phosgene to ethyl carbonate which is harmless.</p>	<p><b>25.</b> (B)  <b>Sol.</b> <math>\text{CaOCl}_2</math> (Bleaching Powder) <math>\xrightarrow{\text{distilled acetone}}</math> <math>\text{CHCl}_2</math> (Chloroform)</p>
<p><b>17.</b> (A)  <b>Sol.</b> Isocyanide reaction involve: <math>\text{CCl}_2</math> intermediate formation.  <math>1^\circ</math> amine + <math>\text{CHCl}_3 + \text{KOH} \rightarrow</math> Isocyanide test</p> 	<p><b>26.</b> (C)  <b>Sol.</b> Carbylamine reaction is used for test of both aromatic &amp; aliphatic primary amine.</p> 
<p><b>18.</b> (B)  <b>Sol.</b> <math>\text{C}_6\text{H}_5\text{NH}_2 + \text{CHCl}_3 + \text{alc. KOH} \rightarrow</math> <math>\text{C}_6\text{H}_5\text{NC}</math> (phenyl isocyanide)</p> <p>This is called as Hoffmann's carbylamine test, which is also called isocyanide test.</p>	<p><b>27.</b> (D)  <b>Sol.</b> Reimer-Tiemann's reaction</p> 
<p><b>19.</b> (A)  <b>Sol.</b> Iodoform test is not given</p>  <p>by because it does not contain <math>-\text{CH}_3</math> bonded to <math>-\text{C}-</math> group on terminal</p>	<p><b>28.</b> (B)  <b>Sol.</b> Grignard Reagent <math>\text{RMgX}</math>.  <math>\text{RX} + \text{Mg} \xrightarrow{\text{Ether}} \text{RMgX}</math> (Grignard Reagent)</p> <p>In case of vicinal dihalide, reaction with Mg leads to elimination reaction and gives alkene.</p> 

<p><b>29.</b> (B)  <b>Sol.</b> Carbylamine test (isocyanide test) Primary amine + <math>\text{CHCl}_3</math> or <math>\text{CHX}_3</math> + alc. KOH <math>\rightarrow</math> isocyanide carbylamine</p>	<p><b>38.</b> (B)  <b>Sol.</b> <math>\text{C}_2\text{H}_5\text{Br} + \text{KCN} \rightarrow \text{C}_2\text{H}_5\text{CN} + \text{KBr}</math>      This reaction is nucleophilic substitution reaction.</p>
<p><b>30.</b> (C)  <b>Sol.</b>       Intermediate formed   </p>	<p><b>39.</b> (C)  <b>Sol.</b> Stable compound due to more <math>-I</math> effect  </p>
<p><b>31.</b> (D)  <b>Sol.</b>  </p>	<p><b>40.</b> (C)  <b>Sol.</b> Most stable carbocation formed        because resonance is more.   </p>
<p><b>32.</b> (B)  <b>Sol.</b>  </p>	<p><b>41.</b> (D)  <b>Sol.</b> <math>\text{RX} + \text{aq. NaOH} \rightarrow \text{ROH} + \text{NaX}</math>      This is nucleophilic substitution reaction</p>
<p><b>33.</b> (A)  <b>Sol.</b>  </p>	<p><b>42.</b> (B)  <b>Sol.</b>       2-Bromo butane is the major product because secondary radical is stable than primary radical.   </p>
<p><b>34.</b> (B)  <b>Sol.</b> <math>\text{C}_2\text{H}_5\text{OH} + \text{HX} \xrightarrow{\text{ZnX}_2} \text{C}_2\text{H}_5\text{X}</math> order of reaction <math>\text{HI} &gt; \text{HBr} &gt; \text{HCl}</math> (<math>\text{I}</math> is the most reactive hydrogen halide)</p>	<p><b>43.</b> (A)  <b>Sol.</b> Bromination of alkane occurs at slower rate because it is less electronegative than chloride.</p>
<p><b>35.</b> (B)  <b>Sol.</b> <math>\text{CH}_3\text{Br} + \text{OH}^- \rightarrow \text{CH}_3\text{OH} + \text{Br}^-</math>      Mechanism <math>\text{S}_{\text{N}}2</math></p>	<p><b>44.</b> (C)  <b>Sol.</b> Secondary butyl bromide  </p>
<p><b>SECTION-B</b></p> <p><b>36.</b> (D)  <b>Sol.</b> <math>\text{CH}_3\text{CH}_2\text{CH}_2\text{Cl} + \text{KCN} \rightarrow \text{CH}_3\text{CH}_2\text{CH}_2\text{CN}</math> (Butyronitrile)</p>	<p><b>45.</b> (C)  <b>Sol.</b> </p>
<p><b>37.</b> (D)  <b>Sol.</b>       Inversion takes place in <math>\text{S}_{\text{N}}2</math> mechanism   </p>	

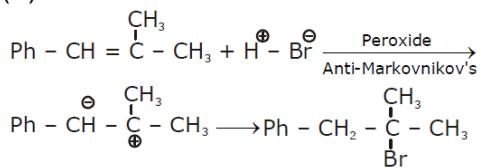
**46.** (D)

**Sol.**



**47.** (B)

**Sol.**



**48.** (B)

**Sol.**

Chlorethane is used as hypnotics.

**49.** (B)

**Sol.** Pyrene is used as fire extinguisher.

**50.** (D)

**Sol.** Freon's are used as (a) Coolant (b) Propellant (c) Solvent.

P  
E