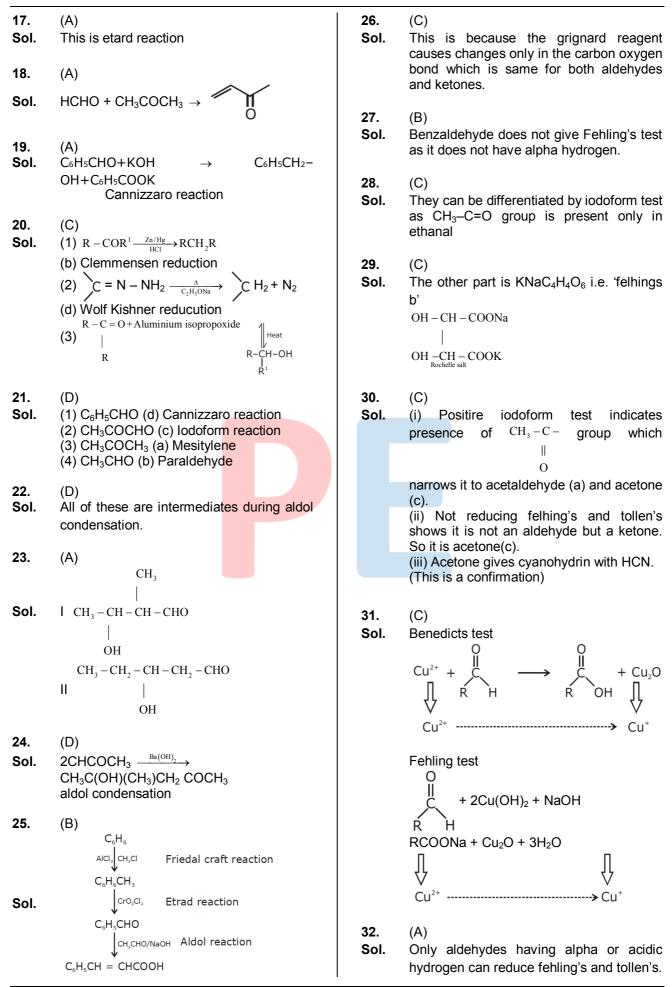
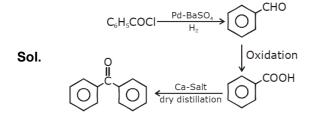
## **NEET ANSWER KEY & SOLUTIONS**

SUBJECT :- CHEMISTRY CLASS :- 12 <sup>th</sup>									PAPER CODE :- CWT-9					
CHAPTER :- ALDEHYDE & KETONE ,CARBOXYLIC ACID ANSWER KEY														
1.	(C)	2.	(C)	3.	(B)	4.	(C)	5.	(B)	6.	(A)	7.	(C)	
8.	(C)	9.	(C)	10.	(C)	11.	(A)	12.	(D)	13.	(A)	14.	(B)	
15.	(C)	16.	(D)	17.	(A)	18.	(A)	19.	(A)	20.	(C)	21.	(D)	
22.	(D)	23.	(A)	24.	(D)	25.	(B)	26.	(C)	27.	(B)	28.	(C)	
29.	(C)	30.	(C)	31.	(C)	32.	(A)	33.	(A)	34.	(A)	35.	(A)	
36.	(A)	37.	(A)	38.	(C)	39.	(A)	40.	(C)	41.	(D)	42.	(D)	
43.	(D)	44.	(C)	45.	(A)	46.	(B)	47.	(C)	48.	(B)	49.	(C)	
50.	(C)													

SOLUTIONS								
1. Sol.	SECTION-A (C) Aldehydes and ketones have elevated boiling	8. Sol.	(C) Protonation is the fastest step.					
2.	points due to dipole-dipole attraction.	9. Sol.	(C) This is due to increase in electron releasing or +I effect.					
Sol.	Aldehydes are easily oxidized to carboxylic acid but ketones are difficult to oxidise. Most	10. Sol.	(C) CH <sub>3</sub> –OH + HCOONa are formed.					
	books say that the hydrogen directly bonded to the C=O. in the aldehyde is what aids the oxidation process	11. Sol.	(A) It is known that basic need for the existance of Keto-enol tautomers is the presence of at least one hydrogen atom					
3. Sol.	(B) Carbonyl compounds are best purified by hydrolysis of sodium bisulphite adducts.	12.	at adjacent sp <sup>3</sup> carbon of carbonyl carbon. (D)					
4. Sol.	(C) $CH_3 - CH = CH - CH_3 \xrightarrow{(i)O_3} 2CH_3 - CHO$ $CH_3 - CH = CH - CH_3 \xrightarrow{(i)H_2/Zn} 2CH_3 - CHO$	Sol.	$   CH_{3}C \equiv CH \xrightarrow{H^{\oplus}/Hg^{2+}} CH_{3} - C - CH_{3}$					
	$CH_{3} - C \equiv CH \xrightarrow{Hg^{2} \cdot H^{\oplus}} $ $CH_{3} - \stackrel{OH}{C} = CH_{2} \leftrightarrow CH_{3} - \stackrel{OH}{C} = CH_{3}$	13.	(A) 0					
5. Sol.	(B) CH <sub>3</sub> MgI + CH <sub>3</sub> CN $\xrightarrow{H^{\oplus}}$ CH <sub>3</sub> - CO - CH <sub>3</sub>	Sol.	$   C_7H_{14} + O_3 \longrightarrow C_3H_6O + CH_3 - CH_3 - C - CH_3$					
6. Sol.	(A) $C_{6}H_{6}COCI \xrightarrow{Pd-BaSO_{4}} C_{6}H_{6}CHO$	14. Sol.	(B) $(CH_3)_2 C = O \xrightarrow{NaCN}_{HCl} (CH_3)_2 C(OH) CN \xrightarrow{H^{\oplus}} (CH_3)_2 C(OH) CO_2 H.$					
	(i) SnCl <sub>2</sub> /HCl (ii) H <sub>2</sub> O	15. Sol.	(C) Due to high polarity of C–O bond in O Ⅱ H−C−H					
7. Sol.	<ul> <li>(C)</li> <li>Aldehydes are more reactive than ketones</li> <li>and +I effect of alkyl groups and steric</li> <li>factor decreases rate.</li> </ul>	16. Sol.	(D) $HC \equiv CH \xrightarrow{H_2SO_4} HCOCH_3 \xrightarrow{NaOH} HCOCH_3 \rightarrow CH_3CH(OH)CH_2CHO.$					



- 33. (A)
  Sol. For more reactivity in nucleophilic addition, the carbonyl carbon should be more electron deficient. Presence of "R" group / +1 groups decrease the electron deficiency. More these groups present, less is the reactivity. Though Ar group are -1 groups they reduce reactivity due to steric hindrance.
- **34**. (A)



- 35. (A)
- **Sol.** O +  $(CH_3CO)_2O \xrightarrow{CH_5COONa} O$

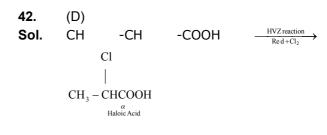
## **SECTION-B**

- 36. (A) Sol.  $C_6H_5CHO + Cl_2 \rightarrow C_6H_5COCI + HCI$ Schotten-Baumann reaction (a)  $R - C + H_2N - R \xrightarrow{NaOH} R - V + HCI$ (b)  $R - C + H_2N - R \xrightarrow{NaOH} R - V + HCI$
- 37. (A) СНО + нСНО -Sol.
  - СH<sub>2</sub>-ОН + НСООН
- **38.** (C) Sol.  $\bigcirc$  CHO  $\xrightarrow{CH_3COONa}{(CH_3CO)_2O} C_9H_8O_2$

**41**. (D)

Sol.  $CH_3$ -COOH + NaHCO<sub>3</sub>  $\longrightarrow$  CH<sub>3</sub>COONa + CO<sub>2</sub> + H<sub>2</sub>O Fehling solution test, tollen reagent are

used to test presence of aldehydic group.



43. (D) Sol.  $CH_3 - C-COOH + Br_2 \xrightarrow{Red P} No Reaction Only -H containing CH_3 - Only Oxidising Carboxylic acid give H.V.Z. Agent Agent Only Oxidiate the second second$ 

Sol. 
$$(HCOO)_2$$
 Ca  $\longrightarrow$  H - C - H  $\frac{NaOH}{Cannizzaro}$   
reaction  
2 HCOONA  $\stackrel{\Delta}{\longrightarrow}$  COONa + H<sub>2</sub>

COONa

## **45.** (A)

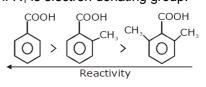
**Sol.**  $\beta$  -Keto acid decarboxylate easily.

## **46.** (B)

**Sol.** Rate of reactivity of acid, when attached with more electron withdrawing group

 $(R_{1}) = (R_{1})$  (R<sub>1</sub> is more electron withdrawing)

 $(R_2 - CH_2OH)$  while its reactivity decreases if  $R_1$  is electron donating group.



47. (C)  
Sol. 
$$\bigcup_{\substack{\text{Br} \\ (i) \text{ Mg } (ii) \text{ CO}_2 \\ (iii) \text{ H}^*}} \bigcup_{\substack{\text{COOH} \\ \text{SOCI}_3 \\ \text{SOCI}_3}} \bigcup_{\substack{\text{COH} \\ (\text{CH}_3 - \text{CH}_2)_2 \text{ Cd} \\ \text{CH}_3 - \text{CH}_3 \\ \text{CH}_3 \\ \text{CH}_3 - \text{CH}_3 \\ \text{CH}_3 - \text{CH}_3 \\ \text{CH}_3 - \text{CH}_3 \\ \text{CH}_3 - \text{CH}_3 \\ \text{CH}_3 \\ \text{CH}_3 \\ \text{CH}_3 - \text{CH}_3 \\ \text{CH}_3 \\$$

- **48.** (B)
- Rate of reactivity carboxylic acid  $\downarrow$  as Sol. groups attached to  $\alpha$  –C are electron donating. –CH<sub>3</sub> is electron donating via +I effect. Order of reactivity of esterification of given carboxylic acid with ethanol is HCOOH > CH<sub>3</sub>–COOH > CH<sub>3</sub> CH<sub>3</sub>  $\mathrm{CH}_3-\mathrm{C}-\mathrm{COOH}>\mathrm{CH}_3-\mathrm{C}-\mathrm{COOH}$

CH<sub>3</sub>

Η

49. (C) HOOC Sol.  $HOOC = C < H \xrightarrow{COOH} Br_2 + H_2O$  Anhydride Cis form

**50.** (C)

**Sol.**  $CH_3 - CH_2 - C \equiv N \xrightarrow{H_2O} CH_3 - CH_2 - COOH$ 

