

Daily Tutorial Sheet-6	Level-2

76.(C) Among α , β , γ hydroxy acids, β -hydroxy acids show dehydration to give α , β unsaturated acids. Hence option (C) is correct answer.

$$\begin{array}{c}
\beta \\
CH_2 - CH_2COOH \xrightarrow{H^+} CH_2 = CHCOOH \\
\downarrow \\
OH
\end{array}$$

 $\alpha\text{-hydroxy}$ acids $\begin{pmatrix} R \ CH - COOH \\ I \\ OH \end{pmatrix} \text{ form diesters (Lactides) under similar conditions.}$

$$2R\overset{\alpha}{\overset{\cap}{C}}H-COOH\overset{H^+}{\xrightarrow{\Delta}}\overset{R}{\overset{\circ}{\bigvee}}\underset{R}{\overset{\circ}{\bigvee}}$$
 Lactide

Options (A) : $CH_3\overset{\alpha}{C}H - COOH$ OH Lactic acid

(D) : $\overset{\alpha}{C}H_2OH$ Glycolic acid

will form Lactides

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Options (D) :
$$\begin{matrix} \gamma \\ \mathrm{CH_2} - \mathrm{CH_2}\mathrm{CH_2}\mathrm{COOH} & \xrightarrow{\mathrm{H}^+} & & \\ \mathrm{OH} & & & \\ \gamma\text{-hydroxy acid} & & & \gamma\text{-Lactone} \\ & & & & & \\ \text{(A cyclic ester)} \end{matrix}$$

77.(A) This involves Baeyer-Villiger oxidation of Ketones and Tischenko reaction of aldehydes.

$$\mathbf{CH_3} - \overset{\mathbf{O}}{\overset{||}{\mathbf{C}}} - \mathbf{C_2} \mathbf{H_5} \xrightarrow{\phantom{\mathbf{C_0 H_5 - C - O - O - H}}} \phantom{\mathbf{CH_3}} \bullet \overset{\mathbf{O}}{\overset{||}{\mathbf{C}}} - \mathbf{OC_2} \mathbf{H_5}$$

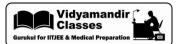
('O' atom enters towards C_2H_5 – side not methyl side)

The aldehyde giving same ester i.e. ethyl acetate has to be $\mathrm{CH_{3}CHO}$ via Tischenko reaction.

78.(C) NaBH $_4$ reduces only keto group, while LiAlH $_4$ reduces both keto and ester groups.

79.(A) Only HCOOH can undergoes oxidation as well as reduction.

80.(B)
$$CN \xrightarrow{H_3O^+} COOH \xrightarrow{COOH} \xrightarrow{\Delta} COOH \xrightarrow{A} COOH$$



81.(D) Visualise Cross-Claisen condensation.

$$\begin{array}{c} \text{CH}_3\text{CH}_2 - \text{C} - \text{OCH}_3 & \xrightarrow{\text{RO}^-} \text{CH}_3 - \overset{\circ}{\text{CH}} - \text{COOMe} & \xrightarrow{\text{H}^-\text{C}^-\text{OMe}} & \text{H}^-\text{C} - \text{CH} - \text{COOMe} \\ & \text{MeO}^-\text{CH}_3 & & \text{CH}_3 & & \text{CH}_3 \\ \end{array}$$

83.(A) Peroxyacetic acid is no doubt a weaker acid than acetic acid as its conjugate base is not resonance stabilised (an essential condition for acidity of an organic acid).

$$\begin{array}{c} \mathbf{O} \\ \mathbf{C} \\ \mathbf{H}_{3} \\ -\mathbf{C} \\ -\mathbf{O} \\ -\mathbf{O} \\ -\mathbf{H} \\ \longrightarrow \\ \mathbf{H}^{^{+}} \\ +\mathbf{C} \\ \mathbf{H}_{3} \\ -\mathbf{C} \\ -\mathbf{O} \\ -\mathbf{O}^{^{-}} \\ \end{array}$$

(– ve charge on $\hat{}$ O' is not in conjugation with $\hat{}$ π' electron)

Whereas in acetic acid, its conjugate base (CH₃COO⁻) is resonance stabilised.

$$\operatorname{CH}_3 - \operatorname{COOH} \longrightarrow \operatorname{H}^{^{\dagger}} + \operatorname{CH}_3 - \operatorname{C} \stackrel{\bigcirc}{\longrightarrow} \operatorname{O}$$
(Resonance stabilised)

84.(A) First visualise Rosenmund reduction of acid chloride to an aldehyde.

Then visualise reduction of aldehyde and ester by LiAlH₄ to form alcohols.

85.(C) Ester react with 1° amines to give 2° Amides : $R - COOR' + RNH_2 \longrightarrow R - C - NHR + R'OH$ $(2^{\circ} \text{ amide})$

Here cyclic ester will open up with 1° amine to form 2° amide as follows.

$$\begin{array}{c}
\text{Me} \\
\text{OH} \\
\text{NH - Me} \\
\text{2° Amide}
\end{array}$$