

Reactions due to H-atom of the carboxylic group Reaction due to Carbonyl group >C=O Reactions due to Carboxyl group

Reactions due to H-atom of the carboxylic group The OH group of carboxylic acid has ionisable hydrogen H⁺. **Acidic Nature** $RCOO^{-} + H_3O^{+}$ $RCOOH + H_2O$ **Evolution of Hydrogen** $2CH_3COOH + 2Na \longrightarrow$ $CH_3COONa + H_2$ $2CH_3COOH + Ca \longrightarrow$ $(CH_3COO)_2Ca + H_2$ Salt formation CH₃COOH +NaOH ----- \rightarrow CH₃COONa + H₂O $CH_{3}COOH + Na_{2}CO_{3} \rightarrow CH_{3}COONa + H_{2}O + CO_{2}$ $CH_3COOH + NH_3 \longrightarrow CH_3COONH_4$

Reaction due to Carbonyl group >C=O
Carboxylic acid undergo reduction in the presence of lithium aluminium hydrideto give pri. alcohols.
RCOOH _____ RCH₂OH + H₂O

Beauveault Blanc Reaction RCOOH + R'OH <u>++</u> RCOOR (Na/ alcohol) RCH₂OH + R'OH

Reactions due to Carboxyl group Decarboxylation (in presence of soda lime) CH₃COOH + NaOH \longrightarrow CH₃COONa + H₂O CH₃COONa + NaOH \longrightarrow CH₄ + Na₂CO₃ $C_6H_5COOH + NaOH$ $\longrightarrow C_6H_5COONa + H_2O$ $\longrightarrow C_6H_6 + Na_2CO_3$ C₆H₅COONa + NaOH Carboxylic acids containing electron withdrawing groups such as NO_2 , CH_3CO are decarboxylated by heating alone.

Hundsdiecker Reaction $RCH_2COOAg + Br_2 \longrightarrow RCH_2Br + CO_2 + AgBr$

Halogenation

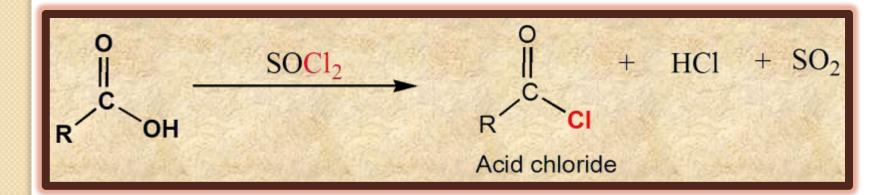
 α -hydrogen of carboxylic acids can be replaced by CI or Br to form α -halo acid in the presence of red phosphorus CH₃CH₂CH₂COOH + Br₂/P \longrightarrow

$CH_3CH_2CH(Br)COOH \longrightarrow CH_3CH_2CBr_2COOH$

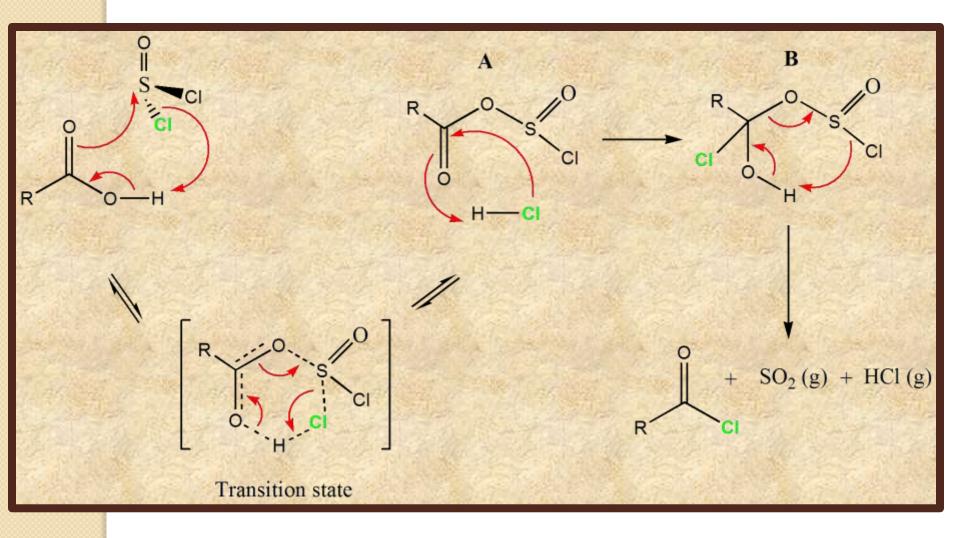
Carboxylic acids undergo reactions to produce derivatives of the acid. The most common derivatives formed are acid halides, esters, acid anhydrides and amides.

Acid chloride (ROCI)

Carboxylic acids react with phosphorous trichloride (PCI_3), phosphorous pentachloride (PCI_5), thionyl chloride ($SOCI_2$), and phosphorous tribromide (PBr_3) to form acyl halides. They are the most reactive derivatives of carboxylic acid.

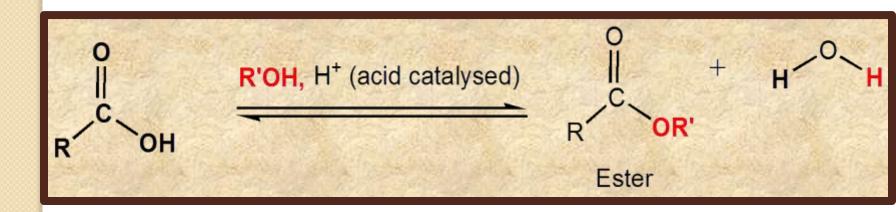


Mechanism of acid chloride formation with SOCI₂

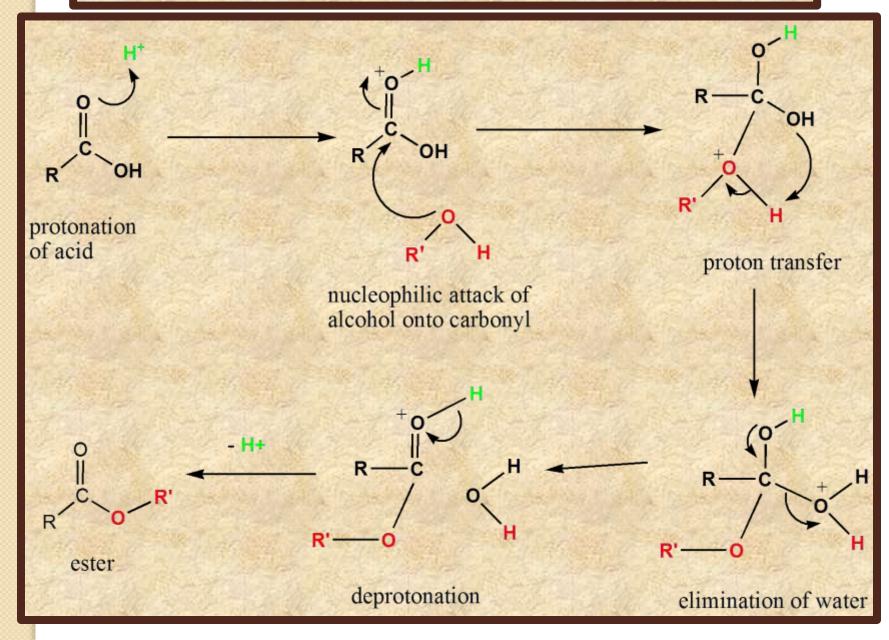


Esterification

Esters are compounds formed by the reaction of carboxylic acids with alcohols, and they have a general structural formula of RCOOR. Esters containing long alkyl chains (R) are main constituents of animal and vegetable fats and oils. Many esters containing small alkyl chains are fruity in smell, and are commonly used in fragrances. The acid-catalyzed esterification of carboxylic acids with alcohols to give esters is termed *Fischer Esterification*.



Mechanism of Fischer Esterification



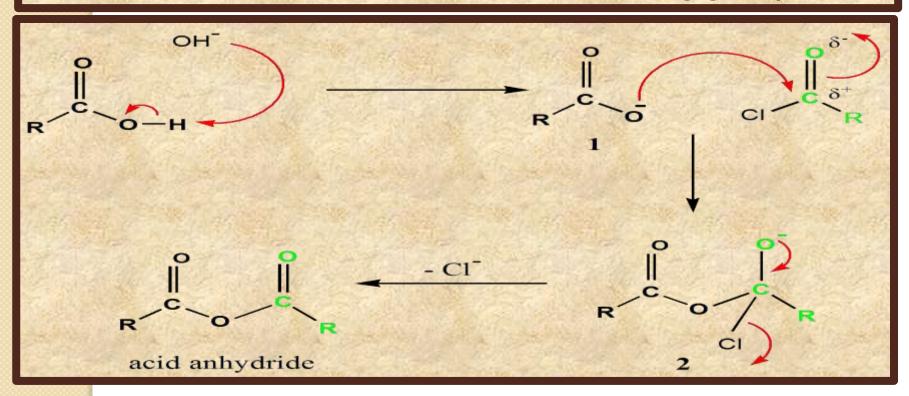
Thioester (RCOSR')

Thioesterification: A thioester is formed when a carboxylic acid reacts with a thiol (RSH) in the presence of an acid.

Thioesters are commonly found in biochemistry, the best-known example being *acetyl CoA*. The mechanism is the same as esterification (discussed above); only difference being that instead of an alcohol (R'OH), a thioalcohol (R'SH) is involved.



Acid anhydride is a compound that has two acyl groups (R-C=O) bonded to the same oxygen atom. These are formed when a carboxylic acid reacts with an acid chloride in the presence of a base. Similar to the Fischer esterification, this reaction follows an addition-elimination mechanism in which the chloride anion Cl^- is the leaving group.



Amide

Amides are compounds that contain the $RCONH_2$ group. The direct conversion of a carboxylic acid to an amide is difficult because amines are very basic and tend to convert carboxylic acids to their highly un-reactive carboxylate ions. Therefore, DCC (Dicyclohexylcarbodiimide) is used to drive this reaction.

Amides are ordinarily prepared by a reaction of acid chlorides with ammonia or amines.

