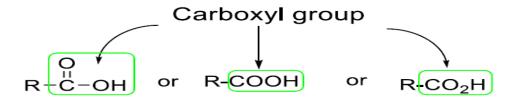
## **Chapter 6**

# **Carboxylic Acids**

#### 1. Structure

Carboxylic acids are organic compounds containing the carboxyl group (-COOH), where in the hydroxyl group (-OH) is directly attached to the carbonyl (C=O) group.



Three representations of a carboxylic acid

# **Nomenclature of Carboxylic Acids**

The common names of some basic carboxylic acids are derived from Latin names that indicate the first original natural source of the carboxylic acid.

Structure of Acid	Natural Source	Common
		Name
O H-C-OH	Ants (Formica)	Formic acid
О СН₃-Ё-ОН	Vinegar (Acetum)	Acetic acid
O CH₃CH₂−C-OH	Basic Fat (Propio)	Propionic acid
О СН <sub>3</sub> СН <sub>2</sub> СН <sub>2</sub> -С-ОН	Rancid butter (Butyrum)	Butyric acid
CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> -C-OH	Present in aValerian herb	Valeric acid
CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> -C-OH	Goat (Caper)	Caproic acid

# **Common Names of Carboxylic Acids**

The common name of a carboxylic acid (R-COOH) is derived by adding the suffix—ic acid to a prefix representing the chain length of the carboxylic acid.

# of Carbons	Prefix	Common Name of Acid
1	Form-	Formic acid
2	Acet-	Acetic acid
3	Propion-	Propionic acid
4	Butyr-	Butyric acid
5	Valer-	Valeric acid
6	Capro-	Caproic acid
Aromatic acid	Benzo-	Benzoic acid

# **IUPAC Nomenclature of Aliphatic Carboxylic Acids**

IUPAC names of straight chain aliphatic carboxylic acids are derived by adding the suffix **–oic acid** to the systematic name of the parent hydrocarbon. They are named as alkanoic acids.

# of Carbons	Structure & IUPAC	Structure & IUPAC
	Name of Alkane	Name of Acid
1	H H-C-H H	H-C-OH
	Methane	Methanoic acid
2	CH <sub>3</sub> -CH <sub>3</sub>	о сн <sub>3</sub> —С-он
	Ethane	Ethanoic acid
3	CH <sub>3</sub> CH <sub>2</sub> -CH <sub>3</sub>	О СН₃СН₂—С-ОН
	Propane	Pronanoic acid

# **Systematic Nomenclature of Substituted Carboxylic Acids**

The systematic names of substituted aliphatic carboxylic acids are derived by:

- (i) First identifying the parent chain that contains most, if not all, the carboxyl groups.
- (ii) Number the parent chain from the carbon of the carboxyl group i.e the carboxyl carbon is C-1.
- (iii) Identify the substituents and assign each substituent a locator/address number (2,3,4...etc.) consistent with the numbering in the parent chain.
- (iv) Arrange the names of the substituents in alphabetical order in the systematic name of the poly-substituted carboxylic acid.

# **Physical properties of Carboxylic Acids**

The physical properties of carboxylic acids can be explained from the perspective of the bond polarization in the carboxyl group and its capacity to engage in hydrogen-bonding. Carboxylic acids boil at considerably higher temperatures than alcohols, ketones, or aldehydes of similar molecular weight.

Strong intermolecular attractive forces hold the acid molecules together

The high boiling point of carboxylic acids is attributed to their capacity to readily form stable, hydrogen-bonded dimers.

# **Preparation of Carboxylic Acids**

# 1. Oxidation of Primary Alcohols

The synthesis of carboxylic acids requires the generation or incorporation of the carboxyl group in a substrate.

The best conditions for the oxidation of primary alcohols to carboxylic acids is under the basic conditions employing potassium permanganate.

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$$R-CH_2OH \xrightarrow{KMnO_4} R-CO_2 \ominus K \oplus \xrightarrow{H^{\oplus}} R-CO_2H$$

# Example

## 2. Oxidation of Aldehydes

Aldehydes can be oxidized to carboxylic acids by a variety of oxidizing agents. Both strong and mild oxidizing agents may be employed successfully.

# Strong Oxidizing Agents

- (a) CrO<sub>3</sub>
   Oxidations in water or dilute mineral acid
- (b) Na<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> or K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>
   Oxidation in dilute mineral acid
- (c) KMnO<sub>4</sub>
  Oxidations in basic media in the presence of KOH

# Mild Oxidizing Agents

(a) Ag<sub>2</sub>O

Note that Ag<sub>2</sub>O does not oxidize alcohols

CHO 
$$\xrightarrow{\text{KMnO}_4}$$
  $\xrightarrow{\text{NaOH, H}_2\text{O}}$   $\xrightarrow{\text{CO}_2 \cdot \text{Na}^+}$   $\xrightarrow{\text{H}_3 \circ \text{O}}$   $\xrightarrow{\text{CO}_2 \cdot \text{H}}$   $\xrightarrow{\text{CO}_2 \cdot \text{Na}^+}$   $\xrightarrow{\text{H}_3 \circ \text{O}}$   $\xrightarrow{\text{CO}_2 \cdot \text{H}}$   $\xrightarrow{\text{CO}_2 \cdot \text{Na}^+}$   $\xrightarrow{\text{CO}_2 \cdot \text{Na$ 

#### 3. Carboxylation of Grignard Reagents

R-Br 
$$\xrightarrow{\text{(1)}} \text{Mg}$$

$$(2) \text{ CO}_2$$

$$(3) \text{ H}_3\text{O} \xrightarrow{\oplus}$$
R-MgBr  $\xrightarrow{\text{CO}_2}$  R-CO<sub>2</sub>MgBr  $\xrightarrow{\text{H}_3\text{O}} \xrightarrow{\oplus}$  R-CO<sub>2</sub>H
$$\xrightarrow{\text{CO}_3} \text{R-CO}_2\text{MgBr} \xrightarrow{\text{CO}_4} \text{R-CO}_2\text{MgBr} \xrightarrow{\text{CO}_4} \text{R-CO}_2\text{HgBr} \xrightarrow{\text{CO}_4} \text{R-CO}_2\text{HgBr}} \xrightarrow{\text{CH}_3\text{CH}_2\text{-CH-OH}} \xrightarrow{\text{CH}_3\text{CH}_2\text{-CH-Br}} \xrightarrow{\text{CH}_3\text{CH}_2\text{-CH-MgBr}} \xrightarrow{\text{CH}_3\text{CH}_2\text{-CH-MgBr}} \xrightarrow{\text{CH}_3\text{CH}_2\text{-CH-CO}_2\text{HgBr}} \xrightarrow{\text{CH}_3\text{CH}_2\text{-CH-CO}_2\text{HgBr}} \xrightarrow{\text{CH}_3\text{CH}_2\text{-CH-CO}_2\text{HgBr}} \xrightarrow{\text{CH}_3\text{CH}_2\text{-CH-CO}_2\text{HgCl}} \xrightarrow{\text{CH}_3\text{CH}_2\text{-CH-CO}_2\text{MgCl}} \xrightarrow{\text{CH}_3\text{CH}_2\text{-CH-CO}_2\text{MgCl}} \xrightarrow{\text{Ph-CO}_2\text{HgBr}} \xrightarrow{\text{Ph-CO}$$

# 4. Hydrolysis of Nitriles

Acid-catalyzed hydrolysis of nitriles provides carboxylic acids.

$$R-C\equiv N \xrightarrow{H_3O^{\oplus}} R-CO_2H + NH_4^{\oplus}$$

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R-Br NaCN R-C
$$\equiv$$
N + NaBr

Alkylhalide Alkylnitrile

Example

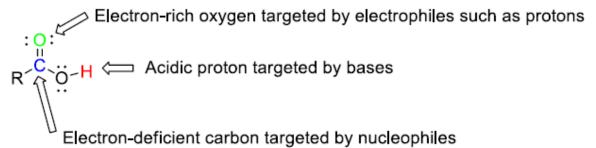
BrCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>Br NaCN NCCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CN H<sub>2</sub>O HO<sub>2</sub>CCH<sub>2</sub>CH<sub>2</sub>CO<sub>2</sub>H

1,3-Dibromopropane Pentanedinitrile Glutaric acid (Pentanedioic acid)

# **Reactions of Carboxylic Acids**

The reactions of carboxylic acids can be directed to various sites on the carboxyl group.

#### Reaction sites on the carboxyl group of Carboxylic Acids



Reactions of carboxylic acids can be placed into four categories:

- (1) Reactions at the acidic hydrogen on the carboxyl group.
- (2) Reactions at the carbonyl group
- (3) Reactions at the carboxylate oxygen
- (4) Reactions that lead to loss of the carboxyl group as CO<sub>2</sub>

# a) Reaction of Carboxylic Acids with Sodium Bicarbonate

The most reliable test for carboxylic acids employs NaHCO<sub>3</sub> leading to evolution of CO<sub>2</sub>. This is commonly called the bicarbonate test for carboxylic acids.

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# Example

## b) Reactions of Carboxylic Acids with Strong Bases

Bases such as metal hydroxides (NaOH and KOH) and amines abstract the acidic proton on carboxylic acids to form carboxylate salts.

# Mechanism

$$R-\overset{O}{C}-\overset{O}{O}-\overset{O}{H}$$
 +  $\overset{O}{\rightarrow}$  OH  $\xrightarrow{\qquad}$   $R-\overset{O}{C}-O\ominus$  +  $H-O-H$ 

# Example

# c) Acid-Catalyzed Esterification of Carboxylic Acids

The traditional method for converting carboxylic acids to esters is through an acidcatalyzed esterification in the presence of an alcohol: Commonly referred to as the Fischer esterification.

The acid-catalyst can be provided by strong mineral acids such as  $H_2SO_4$ , HCl and  $H_3PO_4$  or organic acids such as benzenesulphonic acid or p-toluenesulphonic acid.

#### Example

# Mechanism of the Acid-Catalysed Esterification of Carboxylic Acids

**Step 1:** Protonation of the carbonyl oxygen of the carboxylic acid (activation of the carbonyl carbon).

Step 2: Nucleophilic attack of the alcohol to the activated carbonyl and proton tranfer

**Step 3:** Loss of water to give the conjugate acid of the ester and regeneration of acid catalyst.

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#### d) Reaction of Carboxylic Acids with Amines

Amines, being organic bases, react with carboxylic acids to form ammonium salts.

Mechanism

# Example

# e) Reduction of Carboxylic Acids to Primary Alcohols

Carboxylic acids are reduced to primary alcohols when treated with a strong reducing agent such as LiAlH<sub>4</sub>.

R-CO<sub>2</sub>H 
$$\frac{(1) \text{ LiAIH}_4}{(2) \text{ H}_2\text{O or Dilute acid}}$$
 R CH<sub>2</sub>OH

Example

Ph-CO<sub>2</sub>H 
$$(1)$$
 LiAllH<sub>4</sub> Ph OH  $(2)$  H<sub>2</sub>O

Medium strength reducing agents such as sodium borohydride (NaBH<sub>4</sub>) that reduce aldehydes and ketones are not sufficiently strong to reduce carboxylic acids.

$$H \longrightarrow OH \longrightarrow HO \longrightarrow OH$$