Carboxylic Acids

SUMMARY

Section 19.1 Carboxylic acids take their names from the alkane that contains the same number of carbons as the longest continuous chain that contains the OCO₂H group. The *-e* ending is replaced by *-oic acid*. Numbering begins at the carbon of the OCO₂H group.



Section 19.2 Like the carbonyl group of aldehydes and ketones, the carbon of a CPO unit in a carboxylic acid is sp^2 -hybridized. Compared with the carbonyl group of an aldehyde or ketone, the CPO unit of a carboxylic acid receives an extra degree of stabilization from its attached OH group.



- Section 19.3 Hydrogen bonding in carboxylic acids raises their melting points and boiling points above those of comparably constituted alkanes, alcohols, aldehydes, and ketones.
- Section 19.4 Carboxylic acids are weak acids and, in the absence of electron-attracting substituents, have pK_a 's of approximately 5. Carboxylic acids are much stronger acids than alcohols because of the electron-withdrawing power of the carbonyl group (inductive effect) and its ability to delocalize negative charge in the carboxylate anion (resonance effect).



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Section 19.5 Although carboxylic acids dissociate to only a small extent in water, they are deprotonated almost completely in basic solution.



Sections Electronegative substituents, especially those within a few bonds of the carboxyl group, increase the acidity of carboxylic acids.



Trifluoroacetic acid $pK_a = 0.2$

Section 19.8 Dicarboxylic acids have separate pK_a values for their first and second ionizations.

Section 19.9 Carbon dioxide and carbonic acid are in equilibrium in water. Carbon dioxide is the major component.

$$O = C = O + H_2O \xrightarrow{0.3\%}_{99.7\%} HO \xrightarrow{O}_{OH}$$

 $pK_a = 0.6$

- Section 19.10 Several of the reactions introduced in earlier chapters can be used to prepare carboxylic acids (See Table 19.4).
- Section 19.11 Carboxylic acids can be prepared by the reaction of Grignard reagents with carbon dioxide.



Section 19.12 Nitriles, which can be prepared from primary and secondary alkyl halides by nucleophilic substitution with cyanide ion, can be converted to carboxylic acids by hydrolysis.



Likewise, the cyano group of a cyanohydrin can be hydrolyzed to OCO₂H.

Section 19.13 Among the reactions of carboxylic acids, their conversions to acyl chlorides, primary alcohols, and esters were introduced in earlier chapters and were reviewed in Table 19.5.

Section 19.14 The mechanism of acid-catalyzed esterification involves some key features that are fundamental to the chemistry of carboxylic acids and their derivatives.



Protonation of the carbonyl oxygen activates the carbonyl group toward nucleophilic addition. Addition of an alcohol gives a tetrahedral intermediate (shown in the box in the preceding equation), which has the capacity to revert to starting materials or to undergo dehydration to yield an ester.

Section 19.15 An intramolecular esterification can occur when a molecule contains both a hydroxyl and a carboxyl group. Cyclic esters are called *lactones* and are most stable when the ring is five- or six-membered.



4-Hydroxy-2methylpentanoic acid 2-Methyl-4-pentanolide

Section 19.16 Halogenation at the a-carbon atom of carboxylic acids can be accomplished by the *Hell–Volhard–Zelinsky reaction*. An acid is treated with chlorine or bromine in the presence of a catalytic quantity of phosphorus or a phosphorus trihalide:



This reaction is of synthetic value in that a-halo acids are reactive substrates in nucleophilic substitution reactions.

Section 19.17 1,1-Dicarboxylic acids (malonic acids) and b-keto acids undergo thermal decarboxylation by a mechanism in which a b-carbonyl group assists the departure of carbon dioxide.



Section 19.18 Carboxylic acids are readily identified by the presence of strong IR absorptions at 1700 cm²¹ (CPO) and between 2500 and 3500 cm²¹ (OH), a ¹H NMR signal for the hydroxyl proton at d 10–12, and a ¹³C signal for the carbonyl carbon near d 180.