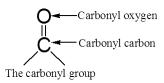
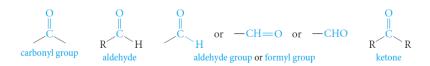


Structure of Aldehydes and Ketones

- Aldehydes and ketones are characterized by the presence of the **carbonyl group**.



- Aldehydes have at least one hydrogen atom attached to the carbonyl carbon atom.
 The remaining group may be another hydrogen atom or any aliphatic or aromatic organic group.
 The -CH=O group characteristic of aldehydes is often called a formyl group.
- In **ketones**, the carbonyl carbon atom is connected to two other carbon atoms.



Nomenclature of Aldehydes

IUPAC System

- Aliphatic aldehydes are named by dropping the suffix -e from the name of the hydrocarbon that has the same carbon skeleton as the aldehyde and replacing it with the suffix -al.

Alkane -e+al = Alkanal

- Substituted aldehydes, we number the chain starting with the aldehyde carbon.
 - -CH=O group is assigned the number 1 position.
 - Aldehyde group has priority over a double bond or hydroxyl group.

Nomenclature of Aldehydes

IUPAC System

- $Cyclic\ aldehydes$, the suffix —carbaldehyde is used.

- Aromatic aldehydes are usually designated as derivatives of the simplest aromatic aldehyde, benzaldehyde.

Nomenclature of Ketones

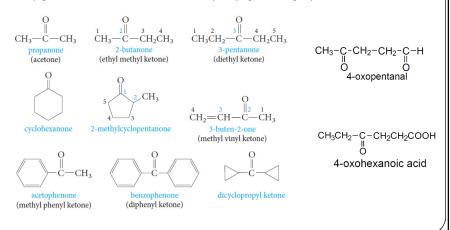
Common Names

- Common names of ketones are formed by adding the word ketone to the names of the alkyl or aryl groups attached to the carbonyl carbon. Alkyl ketone
- In still other cases, traditional names are used.

Nomenclature of Ketones

IUPAC Names

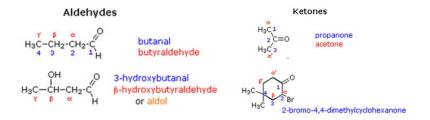
- In the IUPAC system, the ending for ketones is -one.
- The chain is numbered so that the carbonyl carbon has the lowest possible number.
- For cyclic ketones, numbering always starts from the C=O group.
- The prefix "oxo" is used when the ketone is not the principal functional group.



Nomenclature of Aldehydes Ketones

NOTES

- In common names carbon atoms near the carbonyl group are often designated by Greek letters.
- The atom adjacent to the function is *alpha* (α), the next removed is *beta* (β) and so on. Since ketones have two sets of neighboring atoms, one set is labeled α , β etc., and the other α ', β ' etc.

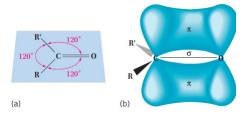


-The functional group priority order in nomenclature system is as following:

Acid and derivatives >aldehyde> ketone> alcoho > amine > alkene > alkyne > ether

The Carbonyl Group

- To best understand the reactions of aldehydes, ketones, and other carbonyl compounds, we
 must first appreciate <u>the structure and properties of the carbonyl group</u>.
- The carbon—oxygen double bond consists of a sigma bond and a pi bond.
- The carbon atom is sp²-hybridized. The three atoms attached to the carbonyl carbon lie in a plane with bond angles of 120°.
- The pi bond is formed by overlap of a p orbital on carbon with an oxygen p orbital.
- There are also two unshared electron pairs on the oxygen atom.
- The C=O bond distance is 1.24A, shorter than the C-O distance in alcohols and ethers (1.43A).



The Carbonyl Group

 Oxygen is much more electronegative than carbon. Therefore, the electrons in the C=O bond are attracted to the oxygen, producing a highly polarized bond.

- As a consequence of this polarization, *most carbonyl reactions involve* nucleophilic attack *at the carbonyl carbon*, often accompanied by addition of a proton to the oxygen (electron rich).

attack here by a
$$\longrightarrow$$
 $C = 0$ may react with a proton

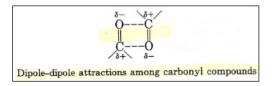
Physical Properties of Aldehydes and Ketones

- Boiling Points

 Carbonyl compounds boil at higher temperatures than hydrocarbons, but at lower temperatures than alcohols of comparable molecular weight.

$$\begin{array}{lll} \text{CH}_3(\text{CH}_2)_4\text{CH}_3 & \text{CH}_3(\text{CH}_2)_3\text{CH} \textcolor{red}{=}\text{O} & \text{CH}_3(\text{CH}_2)_3\text{CH}_2\text{OH} \\ \text{hexane (bp 69°C)} & \text{pentanal (bp 102°C)} & \text{pentanol (bp 118°C)} \end{array}$$

• This is due to the intermolecular forces of attraction, called <u>dipole-dipole interactions</u>, which is stronger than van der Waals attractions but not as strong as hydrogen bonds.



Physical Properties of Aldehydes and Ketones

- Solubility

- Carbonyl compounds as aldehydes and ketones have a C=O bond, but no O-H bond, cannot form hydrogen bonds with themselves.
- The polarity of the carbonyl group also affects the solubility properties of aldehydes and ketones.
- Carbonyl compounds with low molecular weights are soluble in water as they can form hydrogen bonds with O-H or N-H compounds.

$$C = 0 : \cdots H - 0$$

Preparation of Aldehydes and Ketones

1) Oxidation of Primary and Secondary Alcohols

- Chromium reagents, such as pyridinium chlorochromate (PCC), are commonly used in the laboratory.

N+—H CrO₃Cl[—]
pyridinium chlorochromate

- Oxidation of primary alcohols, under controlled conditions, yields aldehydes.

- Oxidation of secondary alcohols yields ketones.

Preparation of Aldehydes and Ketones

2) Hydration of Alkynes

- Hydration of acetylene yields acetaldehyde (catalyzed by acid and mercuric).

- Hydration of terminal alkynes EXCEPT acetylene yields ketones (catalyzed by acid and mercuric).

$$\begin{array}{c} O \\ \parallel \\ \text{CH}_3(\text{CH}_2)_5\text{C} \Longrightarrow \text{CH} \xrightarrow{\text{H}^+,\,\text{H}_2\text{O}} \text{CH}_3(\text{CH}_2)_5\text{CCH}_3 \\ \text{1-octyne} \end{array}$$

Preparation of Aldehydes and Ketones

3) Ozonolysis of Alkenes

- Product (aldehyde or ketone) depends on the structure of alkene.

$$\begin{array}{c} \text{H}_{3}\text{CH}_{2}\text{CHC} = & \begin{array}{c} \text{CHCH}_{3} & \begin{array}{c} \text{(1) O}_{3} \\ \text{(2) Zn, H}_{2}\text{O} \end{array} \end{array} \\ \text{H}_{3}\text{CH}_{2}\text{CC} = 0 \\ \text{H}_{3}\text{CC} = & \begin{array}{c} \text{CH}_{3} \\ \text{CCH}_{3} \end{array} \\ \begin{array}{c} \text{CH}_{3} \\ \text{(2) Zn, H}_{2}\text{O} \end{array} \end{array} \\ \text{H}_{3}\text{CC} = & \begin{array}{c} \text{CH}_{3} \\ \text{H}_{3}\text{CC} = 0 \\ \text{CCH}_{3} \end{array} \\ \begin{array}{c} \text{CH}_{3} \\ \text{H}_{3}\text{CC} = 0 \\ \text{CCH}_{3} \end{array} \\ \begin{array}{c} \text{CH}_{3} \\ \text{H}_{3}\text{CC} = 0 \\ \text{CCH}_{3} \end{array} \\ \begin{array}{c} \text{CH}_{3} \\ \text{CCH}_{3} \\ \text{CCH}_{3} \end{array} \\ \begin{array}{c} \text{CH}_{3} \\ \text{CH}_{3} \\ \text{CH}_{3} \end{array} \\ \\ \begin{array}{c} \text{CH}_{3} \\ \text{CH}_{3} \\$$

4) Friedel-Crafts Acylation

- Preparing ketones that contain an aromatic ring.

Reactions of Aldehydes and Ketones

A) Reduction of Carbonyl Compounds

- Aldehydes and ketones are easily reduced to primary and secondary alcohols, respectively.
- The most common metal hydrides used to reduce carbonyl compounds are lithium aluminum hydride (LiAlH₄) and sodium borohydride (NaBH₄).

- Example:

B) Oxidation of Carbonyl Compounds

- Oxidation of aldehydes gives a carboxylic acid with the same number of carbon atoms.
- Because the reaction occurs easily, many oxidizing agents, such as KmNO4, CrO3, Ag2O and peracids will work.

$$\begin{array}{c|c}
O & \text{oxidizing} \\
R - C - H & \xrightarrow{\text{agent}} & R - C - OH \\
\text{aldehyde} & \text{acid}
\end{array}$$

- Examples:

$$CH_3(CH_2)_5CH = O \xrightarrow{CrO_3, H^+} CH_3(CH_2)_5CO_2H$$

Reactions of Aldehydes and Ketones

C) Nucleophilic Addition Reactions

- Nucleophiles attack the carbon atom of a carbon-oxygen double bond because that carbon has a partial positive charge.
- The overall reaction involves addition of a nucleophile and a proton across the pi bond of the carbonyl group (when carried out in alcohol or water).

1) Addition of Grignard Reagents: Formation of Alcohols

- Grignard reagents act as carbon nucleophiles toward carbonyl compounds.
- The reaction of a Grignard reagent with a carbonyl compound provides a useful route to alcohols.

$$C = O \xrightarrow{1) \text{RMgX, dry ether}} R$$

$$C = O \xrightarrow{1) \text{RMgX, dry ether}} C - OH + Mg^{2+}X^{-}CI^{-}$$
an alcohol

-The type of carbonyl compound chosen determines the class of alcohol produced.

Reactions of Aldehydes and Ketones

- Formaldehyde gives primary alcohols.

$$R - MgX + H - C - H \longrightarrow R - C - OMgX \xrightarrow{H_2O} R - C - OH$$
formaldehyde
$$R - MgX + H - C - H \longrightarrow R - C - OH$$

$$R - MgX + H - C - OH$$

$$R - C - OH$$

- Other aldehydes give secondary alcohols

$$R - MgX + R' - C - H \longrightarrow R - C - OMgX \xrightarrow{H_2O} R - C - OH$$
aldehyde
$$R - MgX + R' - C - OH$$

$$R - C - OH$$

- Ketones give tertiary alcohols.

2) Addition of Hydrogen Cyanide: Formation of Cyanohydrins

- Hydrogen cyanide adds to the carbonyl group of aldehydes and ketones to form cyanohydrins, compounds with a hydroxyl and a cyano group attached to the same carbon.

$$C = O + HCN$$
 $C = O + HCN$
 $C = OH$

a cyanohydrin

- Example

$$\begin{array}{c} O \\ CH_3-C-CH_3 + HCN \xrightarrow{KOH} CH_3-C-CH_3 \\ \\ acetone \\ \end{array}$$

$$\begin{array}{c} CN \\ acetone \\ \\ \end{array}$$

$$\begin{array}{c} OH \\ CH_3-C-CH_3 \\ \\ CN \\ \end{array}$$

$$\begin{array}{c} CN \\ \\ \end{array}$$

Reactions of Aldehydes and Ketones

3) Addition of Alcohols: Formation of Hemiacetals and Acetals

- Alcohols add to the C=O bond, the OR group becoming attached to the carbon and the proton becoming attached to the oxygen.
- Aldehydes and ketones react with alcohols to form, first, hemiacetals and then, if excess alcohol is present, acetals.

- Hemiacetals; it contains both alcohol and ether functional groups on the same carbon atom.

$$\begin{array}{c} ROH + \begin{matrix} R' \\ H \end{matrix} C = O \xrightarrow{H^+} \begin{matrix} RO \\ R' \end{matrix} H$$

$$\begin{array}{c} ROH \\ R' \end{matrix} H$$

- Acetals have two ether functions at the same carbon atom.

3) Addition of Alcohols: Formation of Hemiacetals and Acetals

- The reverse of acetal formation, called acetal hydrolysis.
- Acetal can be hydrolyzed to its aldehyde or ketone and alcohol components by treatment with excess water in the presence of an acid catalyst.

$$\begin{array}{c} \text{OCH}_3 \\ \text{OCH}_3 \\ \text{OCH}_3 \end{array} \begin{array}{c} \text{H}_2\text{O} \\ \text{H}^+ \end{array} \\ \begin{array}{c} \text{CH=O} + 2 \text{ CH}_3\text{OH} \\ \end{array}$$

Reactions of Aldehydes and Ketones

4) Addition of Ammonia and Ammonia Derivatives

- The addition of nitrogen nucleophile, such as ammonia(NH_3) and substituted ammonia (NH_2 -Y).