

# Fundamentals of Organic Chemistry

## CHEM 109

*For Students of Health Colleges*

Credit hrs.: (2+1)





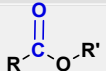
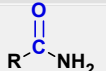
*King Saud University*

College of Science, Chemistry Department

### CHAPTER 5: Aldehydes and Ketones

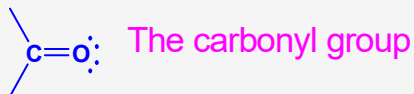
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## Common Classes of Carbonyl Compounds

Class	General Formula	Class	General Formula
Ketones		Aldehydes	
Carboxylic acids		Acid Chlorides	
Esters		Amides	

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## The Carbonyl Group



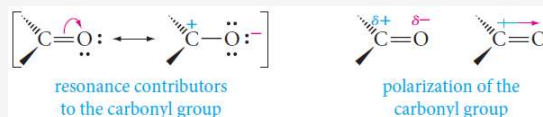
- The carbon–oxygen double bond consists of a sigma bond and a pi bond.
- The carbon atom is  $sp^2$ -hybridized.
- The three atoms attached to the carbonyl carbon lie in a plane with bond angles of  $120^\circ$ .
- The pi bond is formed by overlap of a  $p$  orbital on carbon with an oxygen  $p$  orbital.
- There are also two unshared electron pairs on the oxygen atom.
- The C=O bond distance is 1.24 Å, shorter than the C–O distance in alcohols and ethers (1.43 Å).

	length	energy
ketone C=O bond	1.23 Å	178 kcal/mol (745 kJ/mol)
alkene C=C bond	1.34 Å	146 kcal/mol (611 kJ/mol)

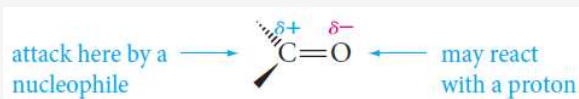
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## The Carbonyl Group

- 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**.



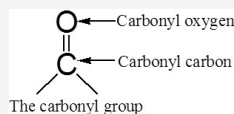
- As a consequence of this polarization, *most carbonyl reactions involve nucleophilic attack at the carbonyl carbon*, often accompanied by addition of a proton to the oxygen (electron rich).



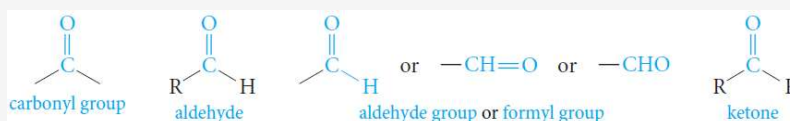
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## Structure of Aldehydes and Ketones

- 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  $\text{-CH=O}$  group characteristic of aldehydes is often called a formyl group.
- In ketones, the carbonyl carbon atom is connected to two other carbon atoms.



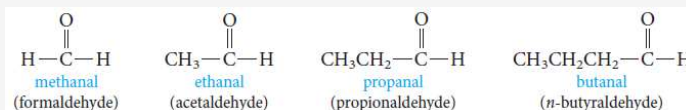
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## IUPAC System

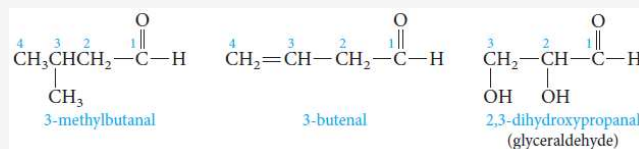
## Nomenclature of Aldehydes

- 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  $-e + al = \text{Alkanal}$



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

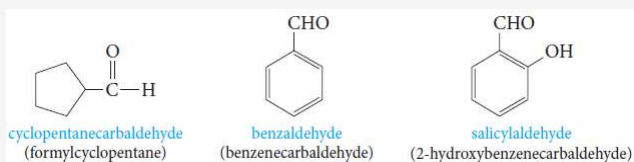


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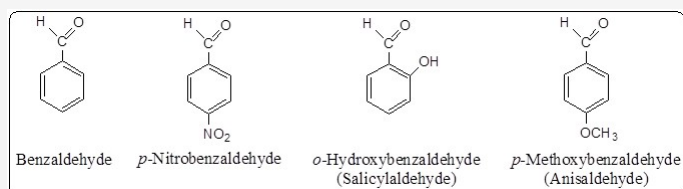
## IUPAC System

## Nomenclature of Aldehydes

- Cyclic aldehydes, the suffix *-carbaldehyde* is used.



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

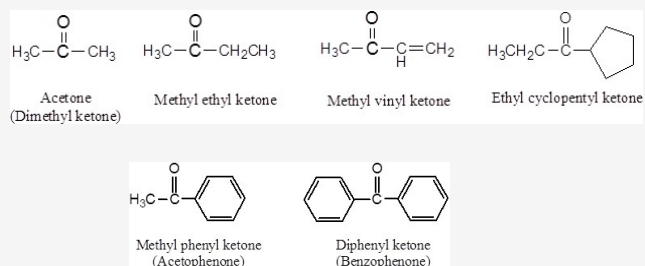


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## Common Names

## Nomenclature of Ketones

- 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*.

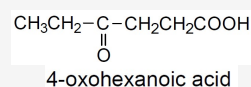
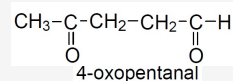
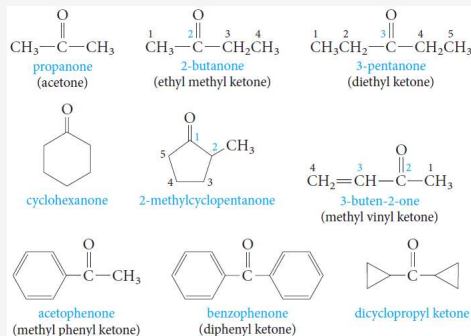


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## IUPAC System

## Nomenclature of Ketones

- 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.

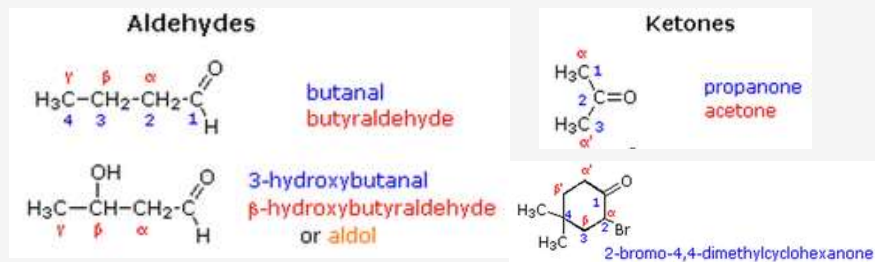


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## NOTES

## Nomenclature of Aldehydes and Ketones

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



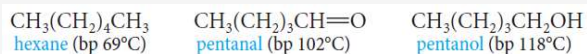
- The **functional group priority order in nomenclature system** is as following:  
 Acid and derivatives > aldehyde > ketone > alcohol > amine > alkene > alkyne > ether

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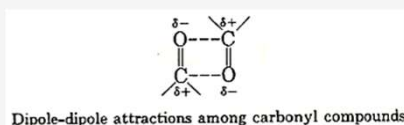
## Physical Properties of Aldehydes and Ketones

### Boiling Points

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



- This is due to the intermolecular forces of attraction, called [dipole-dipole interactions](#), which is stronger than van der Waals attractions but not as strong as hydrogen bonds.

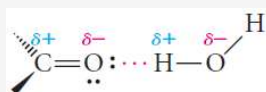


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## Physical Properties of Aldehydes and Ketones

### Solubility

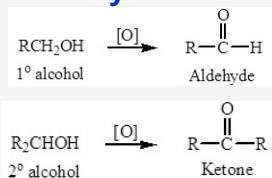
- Carbonyl compounds as aldehydes and ketones have a C=O bond, but no O-H bond, cannot form hydrogen bonds with themselves.
- The polarity of the carbonyl group also affects the solubility properties of aldehydes and ketones.
- Carbonyl compounds with low molecular weights are soluble in water as they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



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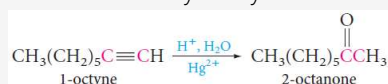
## Preparation of Aldehydes and Ketones

### 1) Oxidation of Primary and Secondary Alcohols

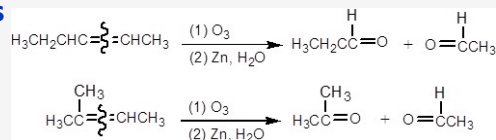


### 2) Hydration of Alkynes

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



### 3) Ozonolysis of Alkenes

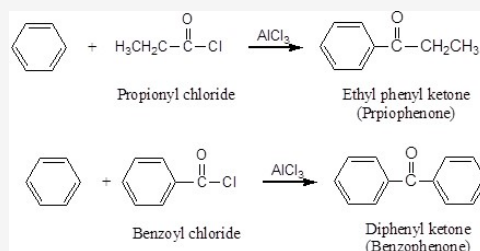


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## Preparation of Aldehydes and Ketones

### 4) Friedel-Crafts Acylation

Preparing ketones that contain an aromatic ring.

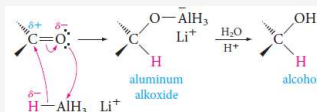


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## Reactions of Aldehydes and Ketones

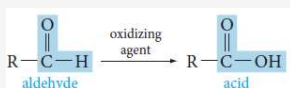
### A) Reduction of Carbonyl Compounds

- 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 ( $\text{LiAlH}_4$ ) and sodium borohydride ( $\text{NaBH}_4$ ).



### 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  $\text{KMnO}_4$ ,  $\text{CrO}_3$ ,  $\text{Ag}_2\text{O}$  and peracids will work.

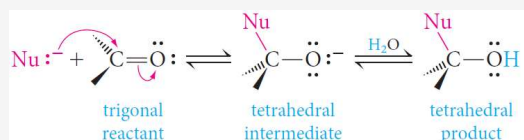


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## Reactions of Aldehydes and Ketones

### 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).



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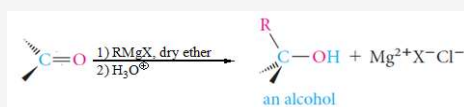
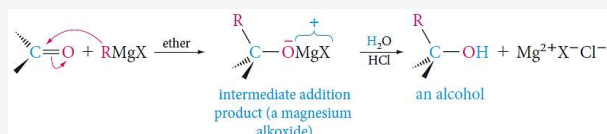


## C) Nucleophilic Addition Reactions

## Reactions of Aldehydes and Ketones

### 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.



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

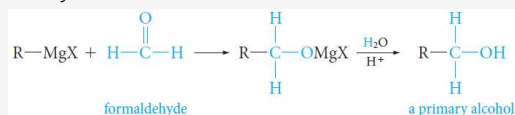
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## C) Nucleophilic Addition Reactions

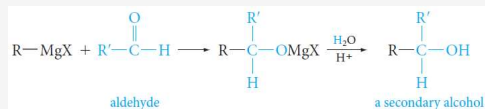
## Reactions of Aldehydes and Ketones

### 1) Addition of Grignard Reagents: Formation of Alcohols

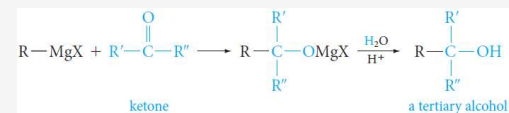
- Formaldehyde gives primary alcohols.



- Other aldehydes give secondary alcohols



- Ketones give tertiary alcohols.



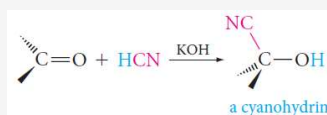
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## C) Nucleophilic Addition Reactions

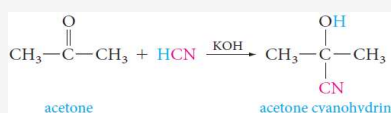
## Reactions of Aldehydes and Ketones

### 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.



#### Example



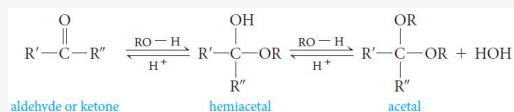
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## C) Nucleophilic Addition Reactions

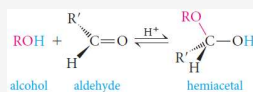
## Reactions of Aldehydes and Ketones

### 3) Addition of Alcohols: Formation of Hemiacetals and Acetals

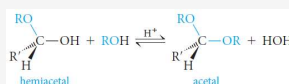
- Alcohols add to the C=O bond, the OR group becoming attached to the carbon and the proton becoming attached to the oxygen.
- Aldehydes and ketones react with alcohols to form, first, [hemiacetals](#) and then, if excess alcohol is present, [acetals](#).



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



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



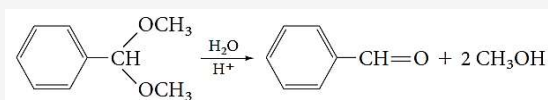
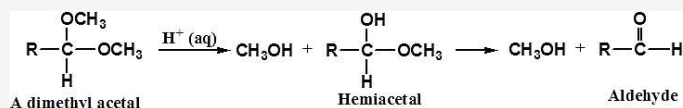
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## C) Nucleophilic Addition Reactions

## Reactions of Aldehydes and Ketones

### 3) Addition of Alcohols: Formation of Hemiacetals and Acetals

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



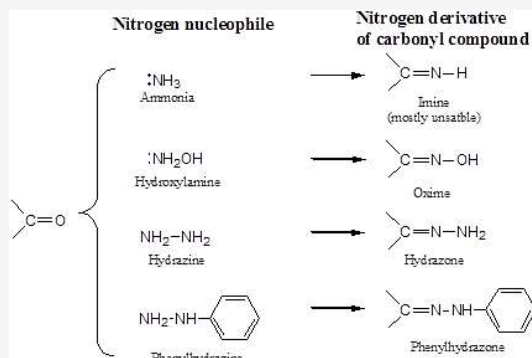
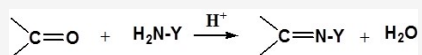
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## C) Nucleophilic Addition Reactions

## Reactions of Aldehydes and Ketones

### 4) Addition of Ammonia and Ammonia Derivatives

The addition of nitrogen nucleophile, such as ammonia ( $\text{NH}_3$ ) and substituted ammonia ( $\text{NH}_2\text{-Y}$ ).



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