

# Organic Reactions & Reaction Mechanisms

- **Types of organic reactions and their mechanisms**
  - **Hydrocarbon derivatives:**  
**(alcohols and carbonyl compounds)**


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*Reactions are like Words,*  
*Mechanisms are like Grammar*  
**to “speak” Organic Chemistry, you need both**

# Classification of Organic reaction

There are so many types of organic reactions:

- **Addition Reactions**
- **Elimination Reactions**
- **Substitution Reactions**
- **Oxidation – Reduction Reaction**
- **Rearrangement Reaction**



In this lecture, we're going to focus on these only

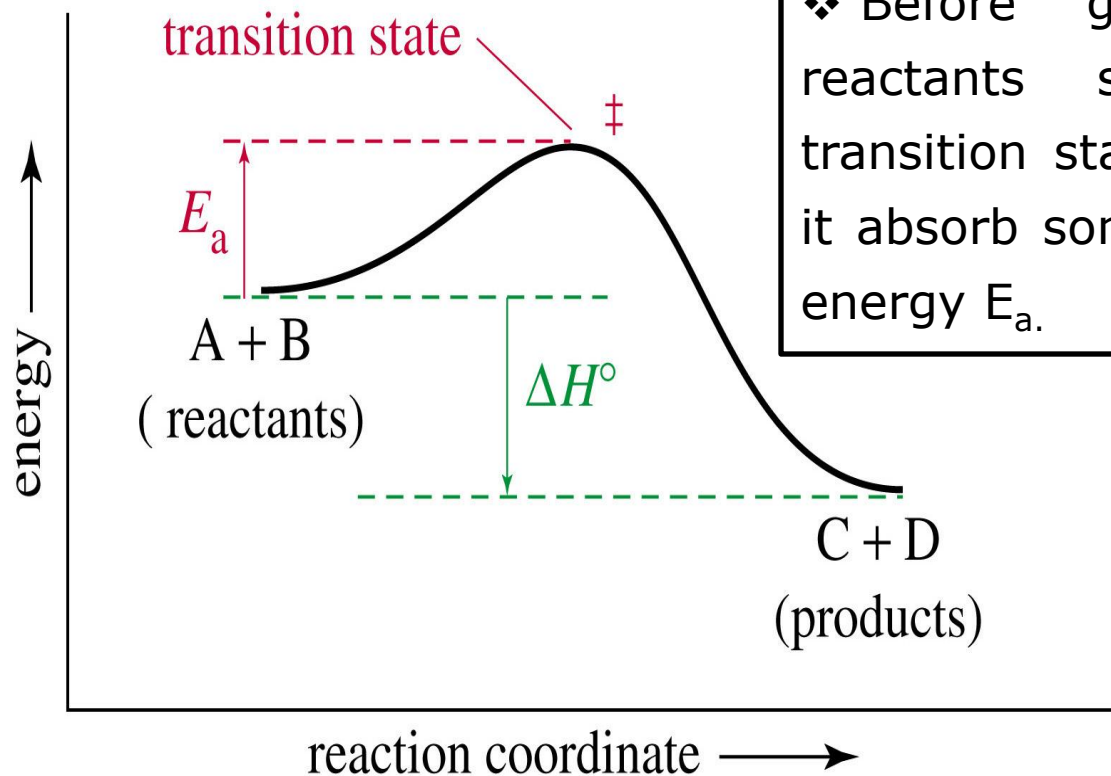
We will also study:

## **Reaction Mechanisms**

- ✓ Bond Dissociation
- ✓ Bond Formation

# Let's start with Reaction Profile (Exothermic)

If we have the following reaction:



❖ In this reaction, the reactants have higher energy than products. So, they lose energy equal to  $\Delta H$  to form products.

❖ Before going to the products, reactants should transform to a transition state (an intermediate), thus it absorb some energy called activation energy  $E_a$ .

# Some Definitions

- **Reaction Mechanism:** sequence of reaction steps that must occur to go from reactants to products.

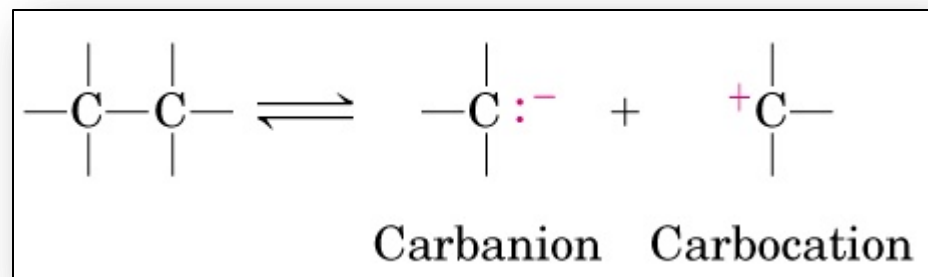
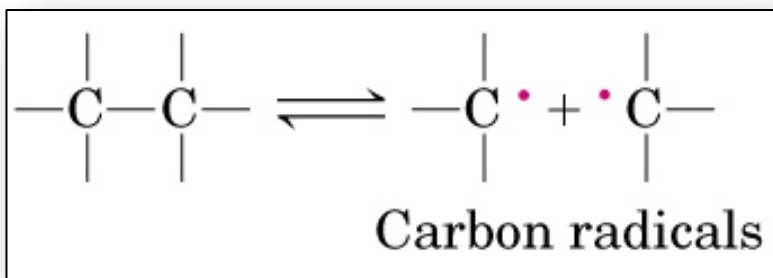
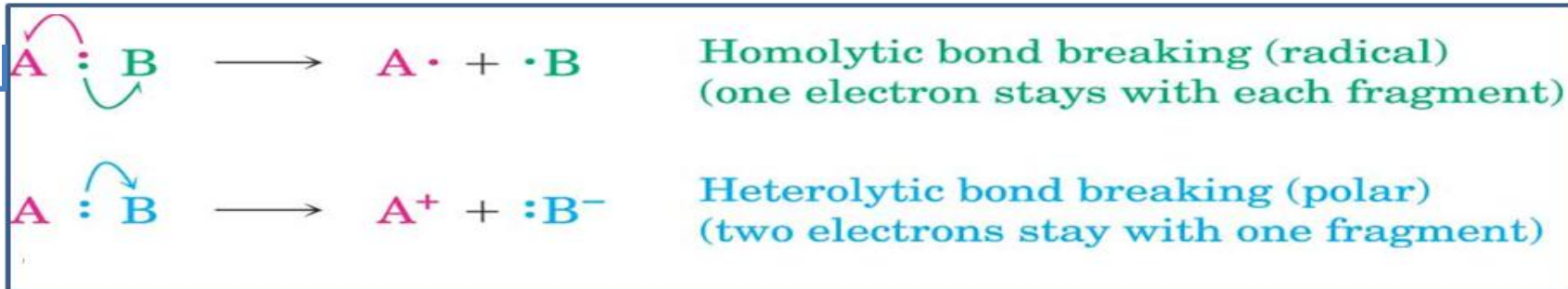
*Each step include dissociation of certain bond, or formation of new one.*

- **Thermodynamics:** The study of the energy changes that occur in chemical transformations. It shows us the stability of reactants compared to products.
- **Kinetics:** The study of reaction rates:
  - determining which product is formed rapidly.
  - predicting the factors affecting the reaction rate.
- **Transition state:** unstable species that has short life time, and convert rapidly to the final product.

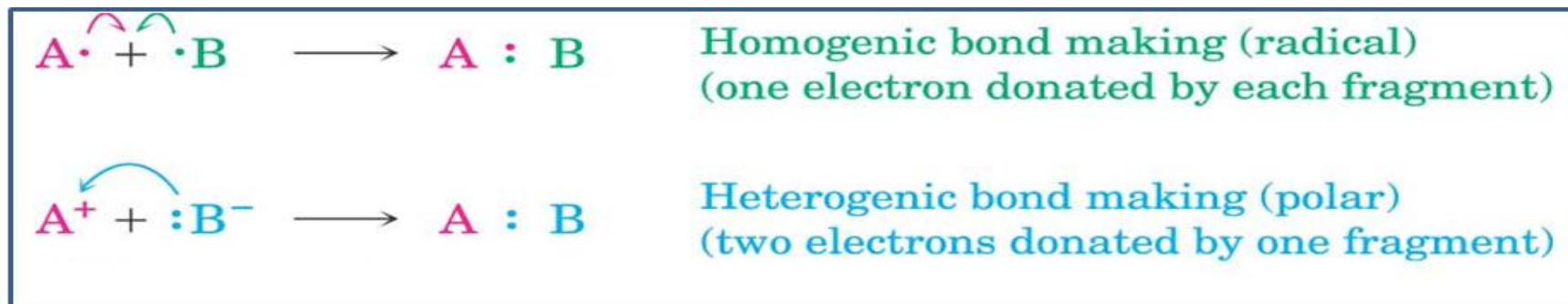
# Bond Breaking and Formation

## Polar and Non Polar

### Bond Breaking:

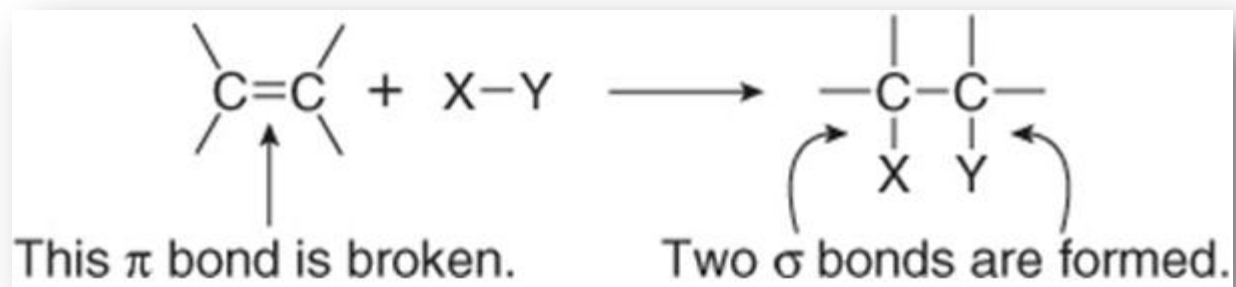


### Bond Formation:



# I- Addition Reactions

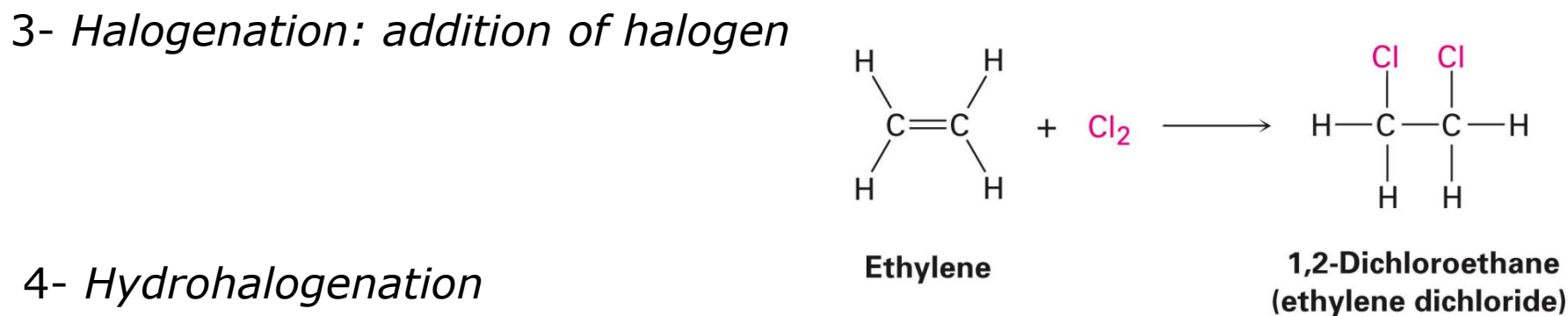
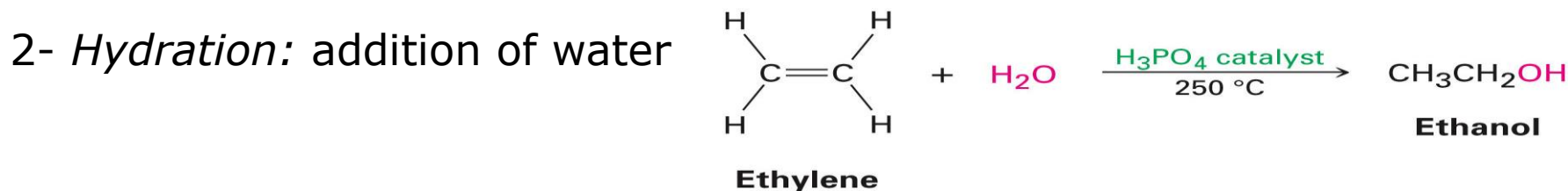
**Addition reaction:** two molecules combine to give one molecule.



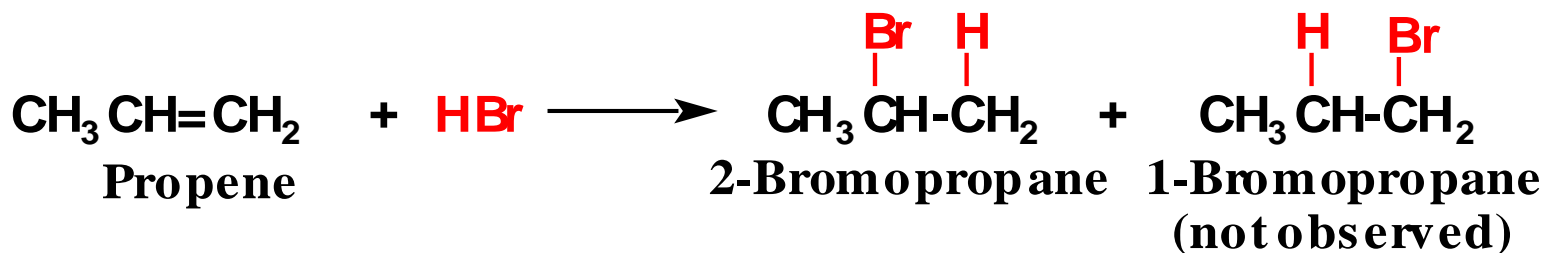
- It Occurs in alkene & alkyne.
- The double or triple bond is easily broken (highly reactive).
- Reactants are added to the carbon atoms in these bonds.

Reactant Added	Name of Addition Reaction
H <sub>2</sub>	Hydrogenation
Cl <sub>2</sub> , Br <sub>2</sub>	Halogenation
HCl, HBr, HI	Hydrohalogenation
HOH	Hydration

# Examples of Addition Reactions

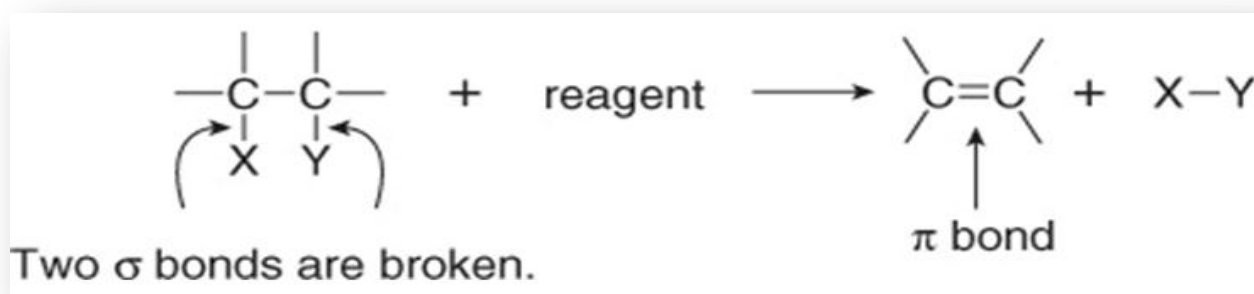


4- *Hydrohalogenation*



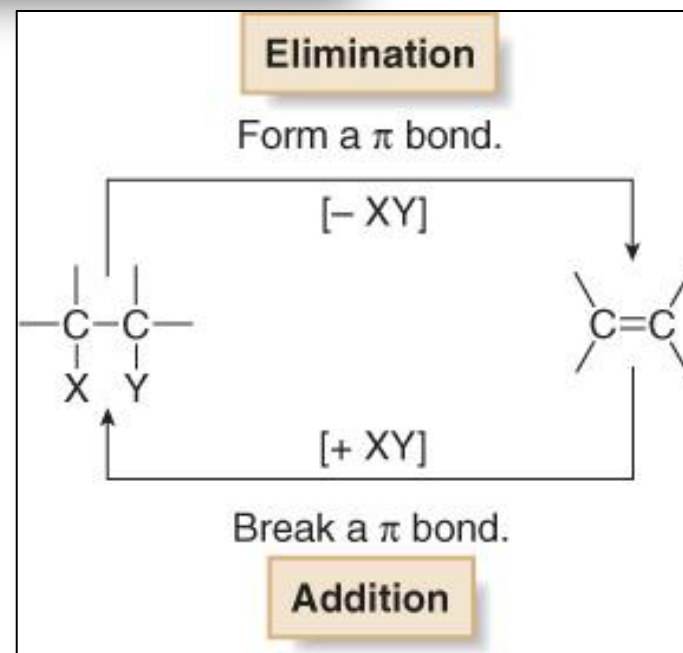
# II- Elimination Reactions

**Elimination Reaction:** removal of a molecule from two adjacent carbon atoms. Ex. Preparation of alkene or alkyne



- Two groups X and Y are removed from a starting material.
- Two  $\sigma$  bonds are broken, and a  $\pi$  bond is formed between adjacent atoms.

**Addition and elimination reactions are exactly opposite.**

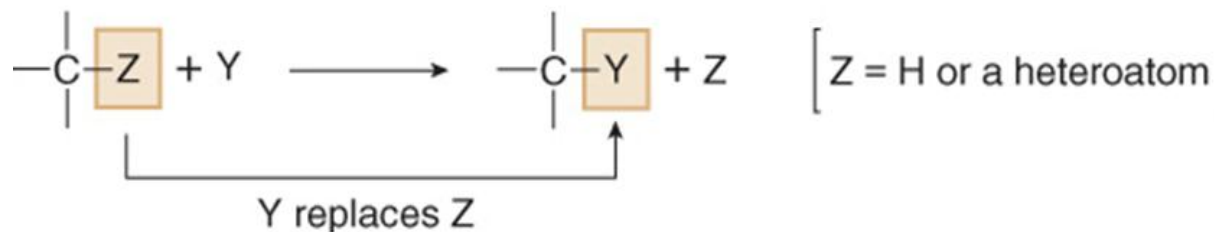




# III- Substitution Reactions

**Substitution Reaction:** a reaction in which an atom or a group of atoms is replaced by another atom or group of atoms.

- It occurs on  $\sigma$  bonds and at the same carbon atom: one  $\sigma$  bond breaks and another forms



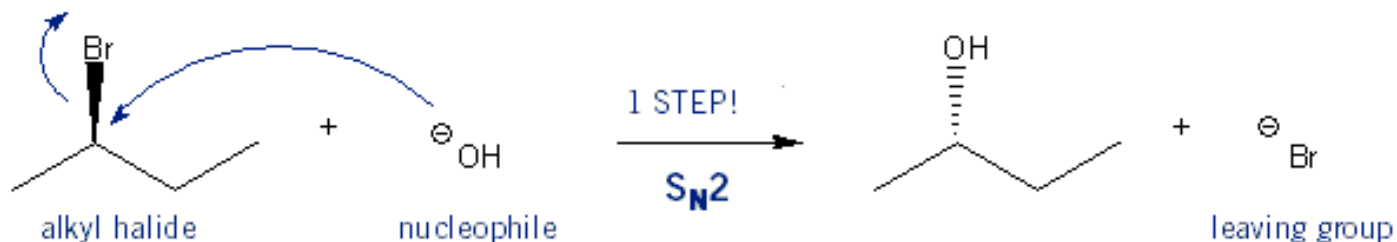
In a general substitution, Y (nucleophile) replaces Z on a carbon atom. Its mechanism can be classified into: (SN<sup>1</sup>) or (SN<sup>2</sup>) depending on timing of bond breaking and bond forming steps and the strength of Nu.

**Nucleophile:** a molecule or ion that donates a pair of electrons to another molecule or ion to form a new covalent bond.

## **S<sub>N</sub>2 Reaction:** Bimolecular Nucleophilic Substitution Reaction

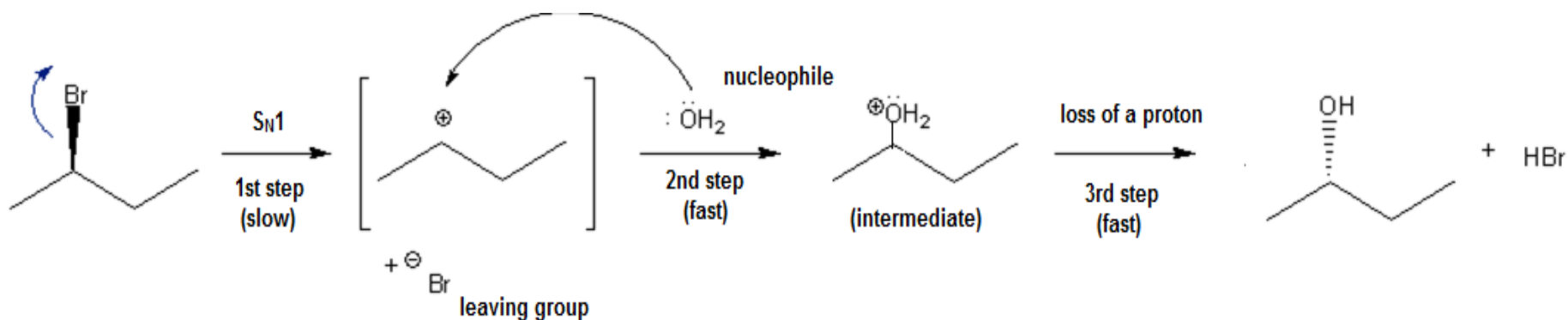
It takes place in a single step without intermediate

### S<sub>N</sub>2 Reaction



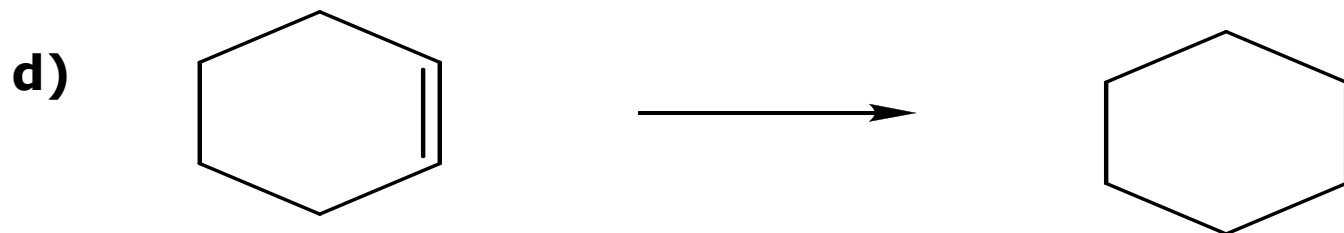
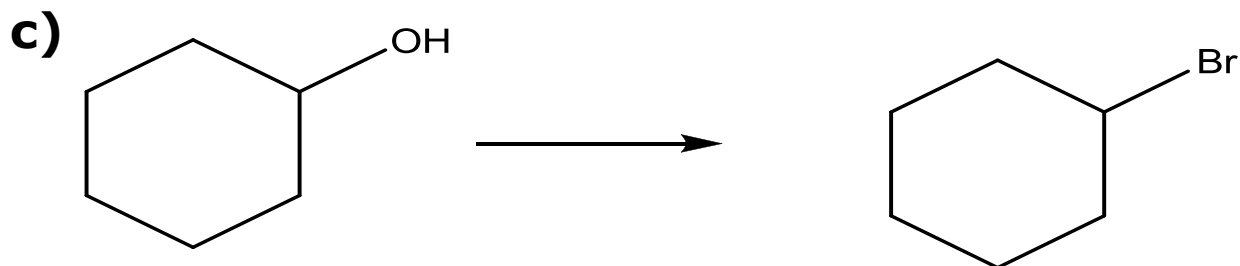
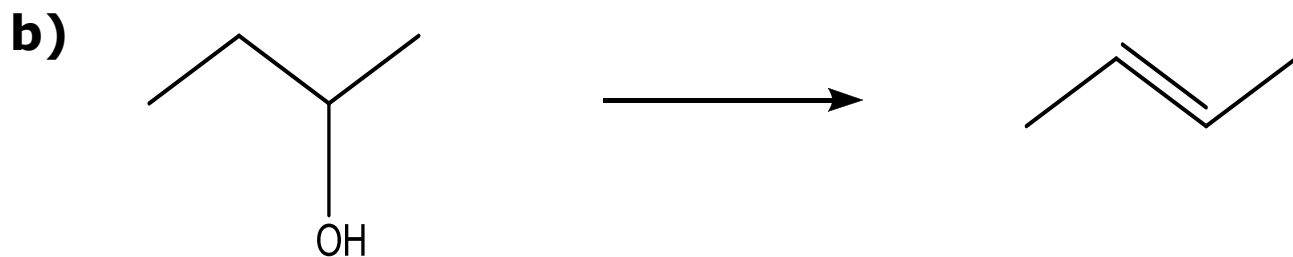
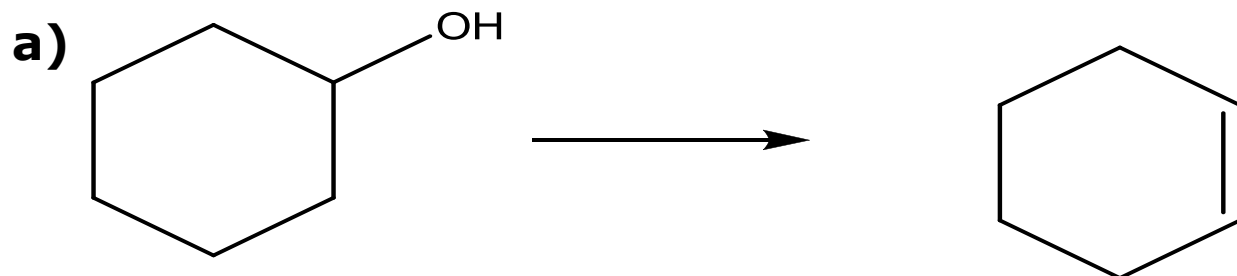
## **S<sub>N</sub>1 Reaction:** Unimolecular Nucleophilic Substitution Reaction

It takes place in a three steps and involve formation of intermediate



## Practice Exercises:

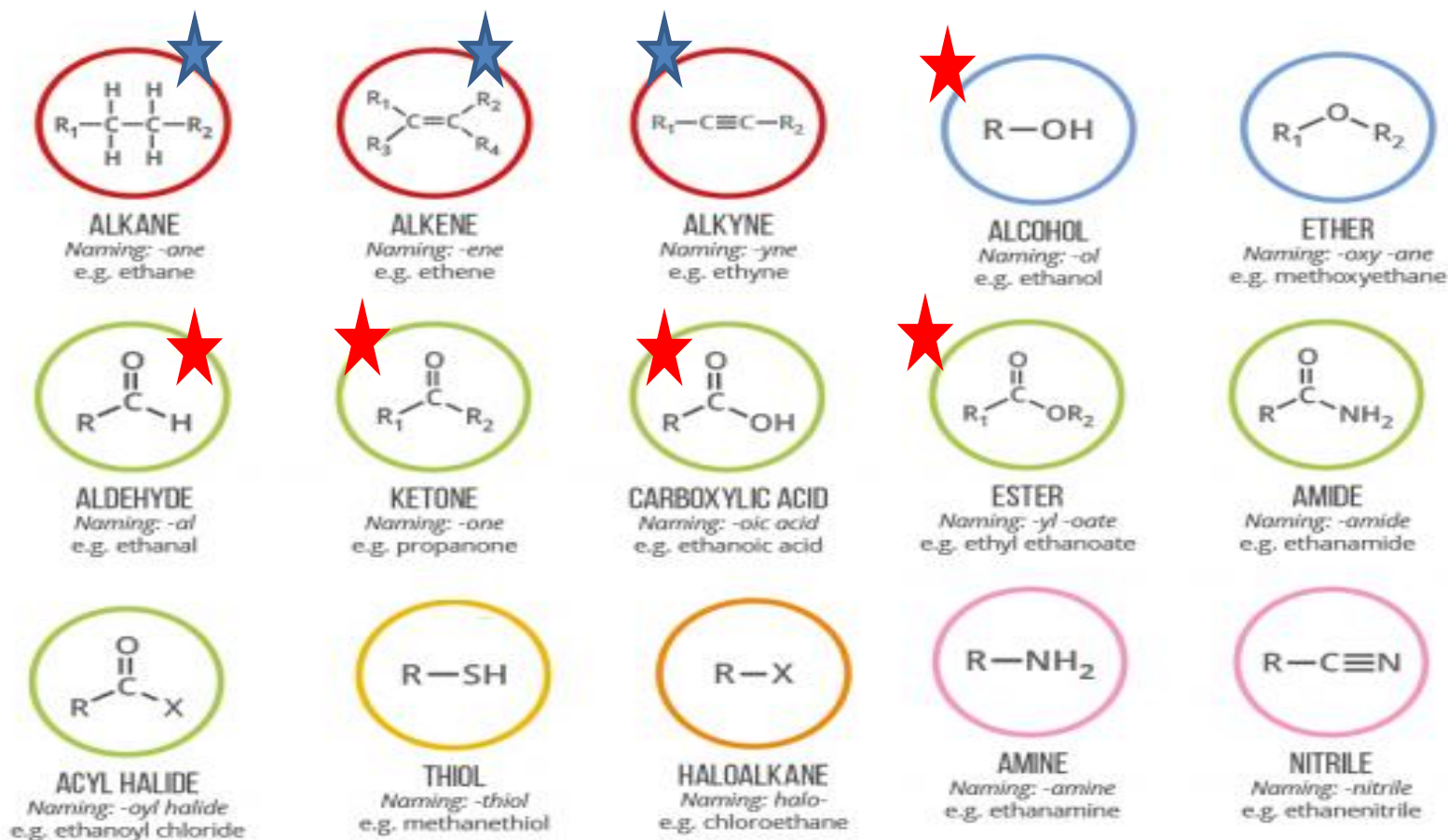
Classify the following reactions as substitution, elimination, or addition.



Types of reactions	Example
<p><b>Addition:</b> two molecules combine to give one molecule. Occurs in alkene &amp; alkyne</p>	$\text{H}_2\text{C}=\text{CH}_2 + \text{H}_2 \xrightarrow{\text{Pt}} \begin{array}{c} \text{H} \quad \text{H} \\   \quad   \\ \text{H}_2\text{C}-\text{CH}_2 \end{array}$
<p><b>Substitution:</b> one atom, ion or group is replaced (substituted) by another (<math>S_N1</math>, <math>S_N2</math>) Usually occurs in saturated compounds such as alkanes.</p>	$\text{CH}_4 + \text{Cl}_2 \xrightarrow{\text{light}} \text{CH}_3\text{Cl} + \text{HCl}$
<p><b>Elimination:</b> removal of a molecule from two adjacent carbon atoms.</p>	$\text{CH}_3-\text{CH}_2\text{Cl} \xrightarrow{\text{strong base}} \text{CH}_2=\text{CH}_2 + \text{HCl}$
<p><b>Oxidation –Reduction:</b> <b>Oxidation:</b> gain of O, loss of H, or both <b>Reduction:</b> gain of H, loss of O, or both</p>	$\text{R-CH}_3 \begin{array}{c} \xrightarrow{\text{oxidation}} \\ \xleftarrow{\text{reduction}} \end{array} \text{RCH}_2\text{OH} \begin{array}{c} \xrightarrow{\text{oxidation}} \\ \xleftarrow{\text{reduction}} \end{array} \text{RCHO}$
<p><b>Rearrangement:</b> molecule undergoes changes to be converted to another isomer.</p>	$\begin{array}{c} \text{CH}_3\text{CH}_2 \\   \quad   \\ \text{C}=\text{C} \\   \quad   \\ \text{H} \quad \text{H} \\ \text{1-Butene} \end{array} \xrightleftharpoons{\text{Acid catalyst}} \begin{array}{c} \text{H}_3\text{C} \quad \text{H} \\   \quad   \\ \text{C}=\text{C} \\   \quad   \\ \text{H} \quad \text{CH}_3 \\ \text{2-Butene} \end{array}$

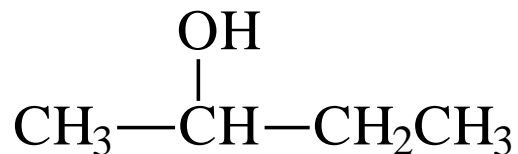
# Function groups in organic compounds

**Function group:** It is a group of atoms in a molecule which is responsible for its chemical reactions and behavior.

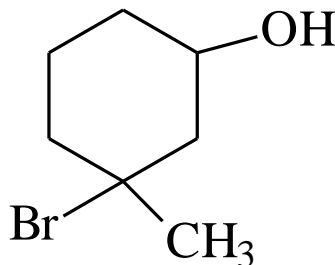


# Alcohols: Preparation – properties - uses

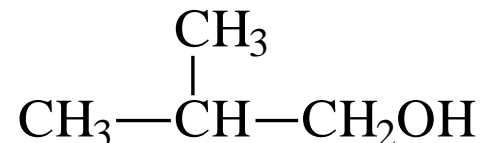
- **Alcohol:** organic compounds in which the hydroxyl functional group (-OH) is bonded to a saturated carbon atom.
- **Nomenclature:** add **"ol"** to the longest chain after removing "e" from alkane



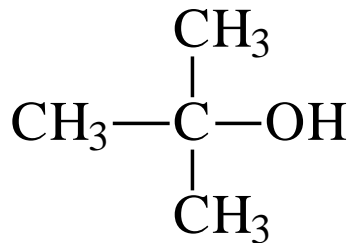
2-butanol or butyl alcohol



3-bromo-3-methylcyclohexanol



2-methyl-1-propanol  
Or isobutyl alcohol



2-methyl-2-propanol

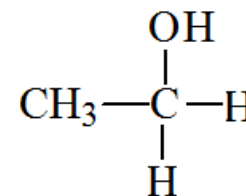


4-penten-2-ol

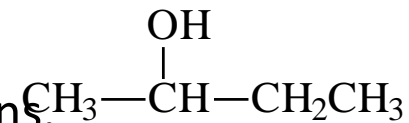
# Classifications

I) according to kind of the C-atom attached to OH:

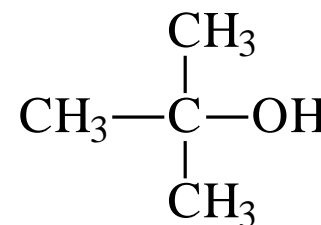
1- Primary (1°): carbon with -OH bonded to one other carbon.



2- Secondary (2°): carbon with -OH bonded to two other carbons.

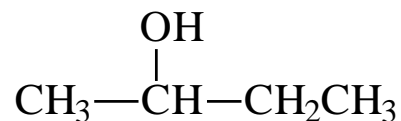


3- Tertiary (3°): carbon with -OH bonded to three other carbons.

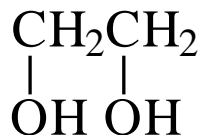


II) according to numbers of OH:

1- Monohydric alcohol

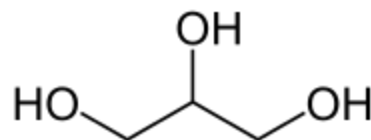


2- Dihydric alcohol



1,2-ethanediol (ethylene glycol)

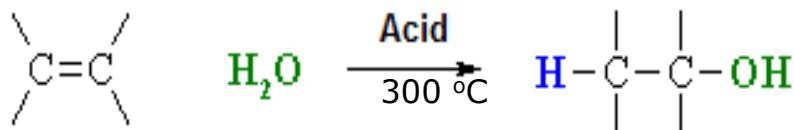
2- Trihydric alcohol



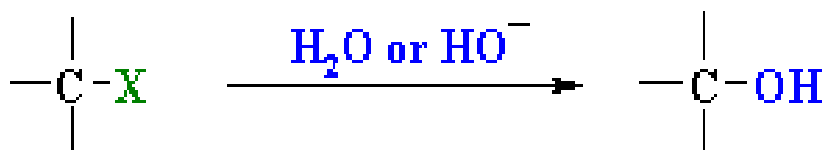
1,2,3-propanetriol (glycerol)

# Preparation of alcohols

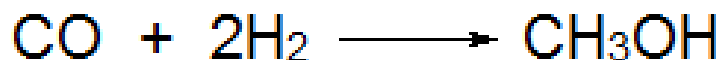
- Hydration of alkene:



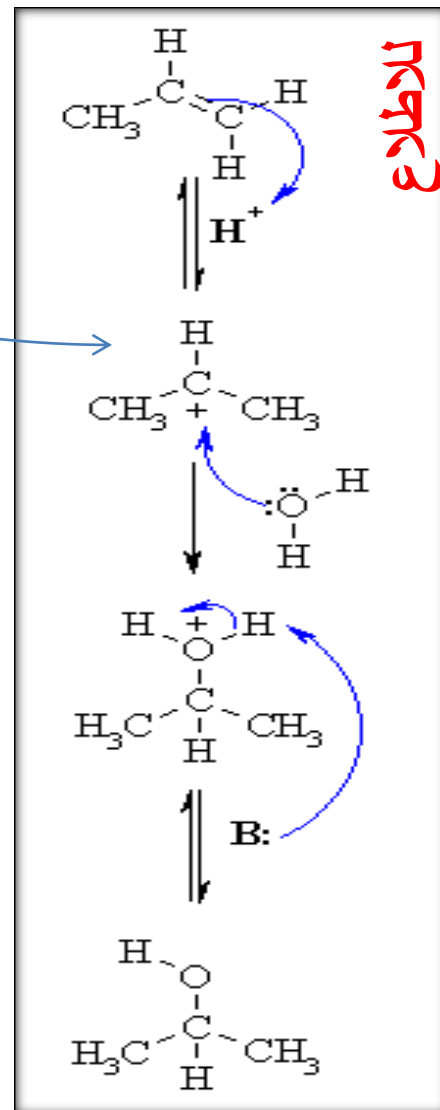
- Hydrolysis of alkyl halides



- In industry:** preparation of methanol from hydrogenation of Carbon monoxide CO



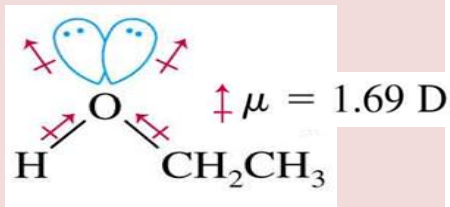
Reaction mechanism



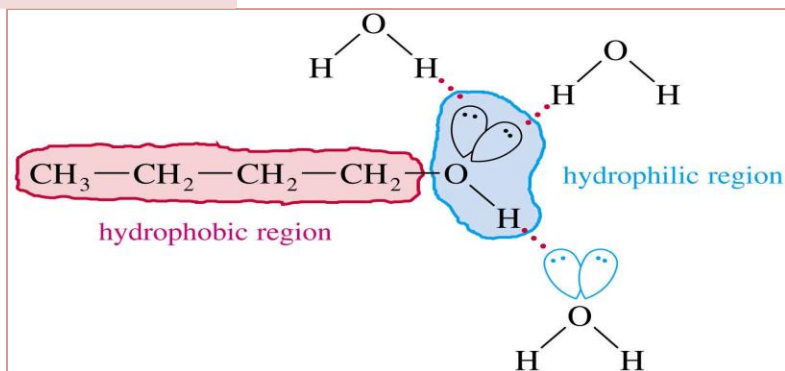
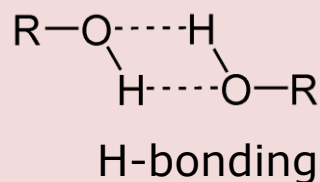


# Physical properties

- ❑ Alcohols are **Polar** compounds, they are the **third** in terms of polarity.
- ❑ They have higher boiling points (than their corresponding alkanes) due to hydrogen bonding between molecules.
- ❑ B.p of  $1^\circ$  alcohol  $>$   $2^\circ$   $>$   $3^\circ$
- ❑ Small alcohols are miscible in water, but solubility decreases as the size of the alkyl group increases.
- ❑ Solubility of  $1^\circ$  alcohol  $>$   $2^\circ$   $>$   $3^\circ$



Polarity



Solubility of alcohols at 25 °C

<i>Alcohol</i>	<i>Solubility in Water</i>
methyl	miscible
ethyl	miscible
<i>n</i> -propyl	miscible
<i>t</i> -butyl	miscible
isobutyl	10.0%
<i>n</i> -butyl	9.1%
<i>n</i> -pentyl	2.7%
cyclohexyl	3.6%
<i>n</i> -hexyl	0.6%
phenol	9.3%
hexane-1,6-diol	miscible

# Uses of alcohols

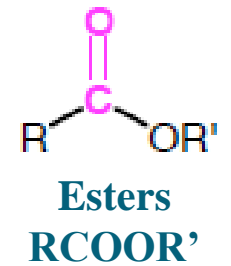
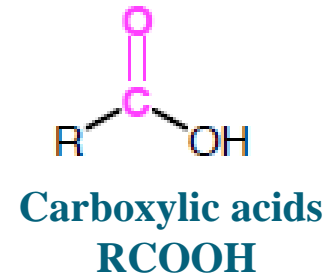
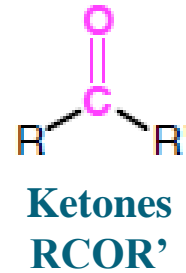
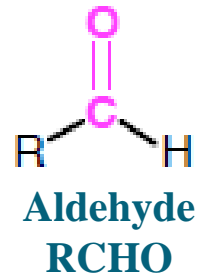
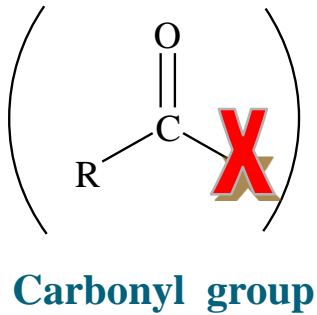
## **As a fuel**

- Methanol and ethanol burns to give  $\text{CO}_2$  and water.
- They can be used as a fuel alone, or in mixtures with petrol (gasoline). "Gasohol" is a petrol / ethanol mixture containing about 10 - 20% ethanol.
- Some countries can produced ethanol by fermentation to replace the fossil fuel (to reduce imports of petrol).
- **As a solvent**
- Ethanol and methanol are widely used as a solvent.
- Ethanol is relatively safe than methanol, and can be used to dissolve many organic compounds which are insoluble in water.
- They are used in manufacturing many perfumes and cosmetics.

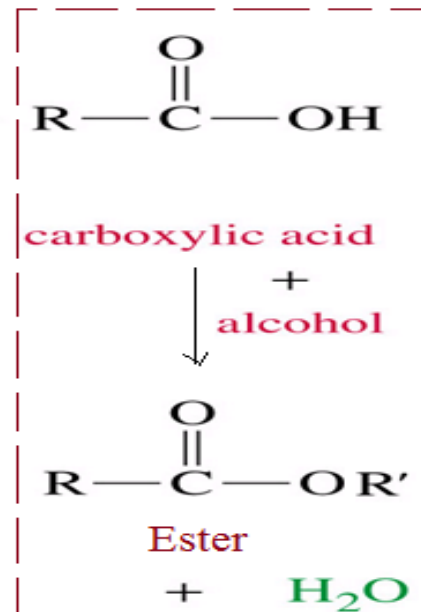
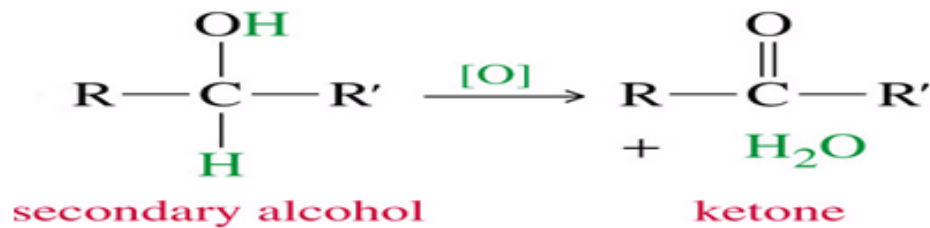
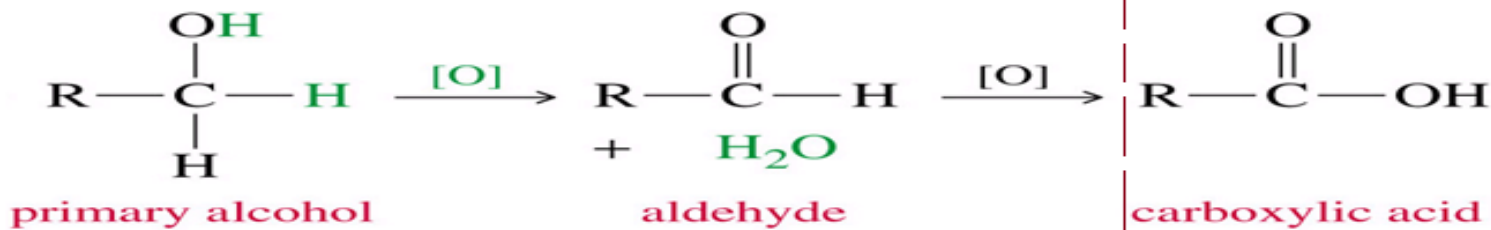
# Hydrocarbon derivatives (**carbonyl compounds**)

**Their preparation – properties – reactions –uses**

# Hydrocarbon derivatives containing carbonyl groups



OXIDATION

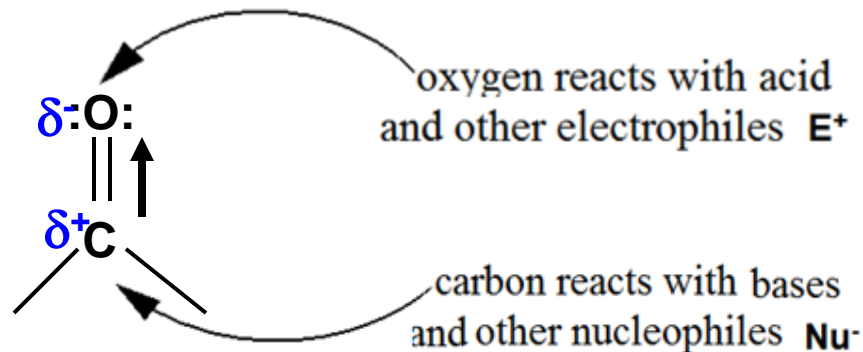


Hydration (addition of H<sub>2</sub>O)

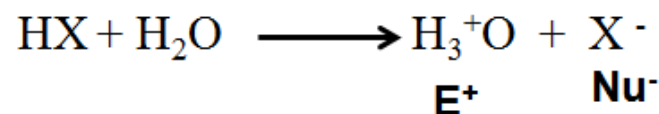
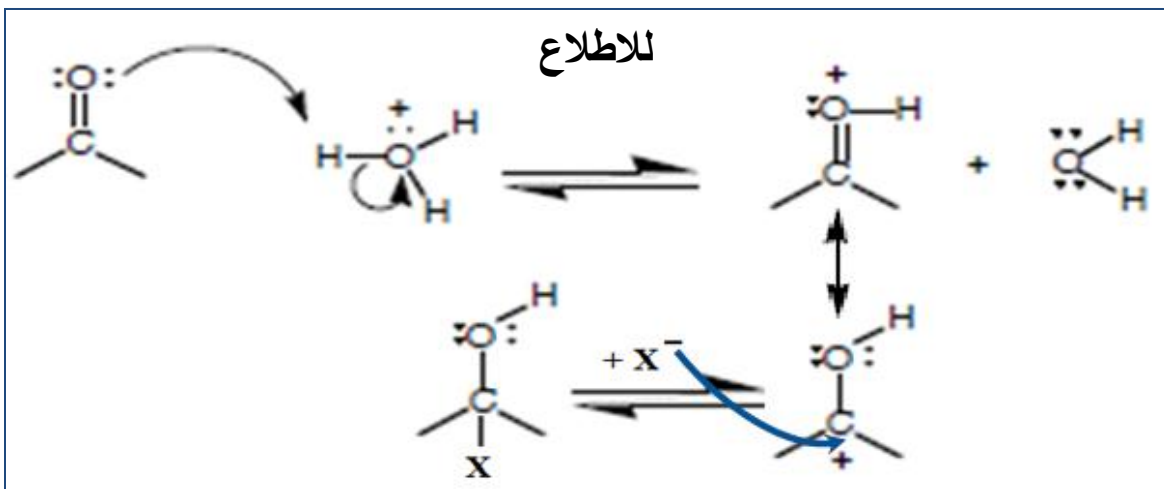
REDUCTION

# Reactions occur in carbonyl group $\text{>C=O}$

$\text{C=O}$  bond of the carbonyl group is polarized. This polarization is responsible for the characteristic reactions of carbonyl compounds.

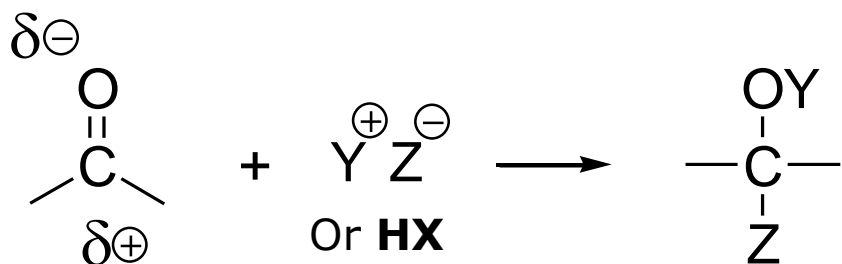


Ex. Reaction of carbonyl compound with an acid:



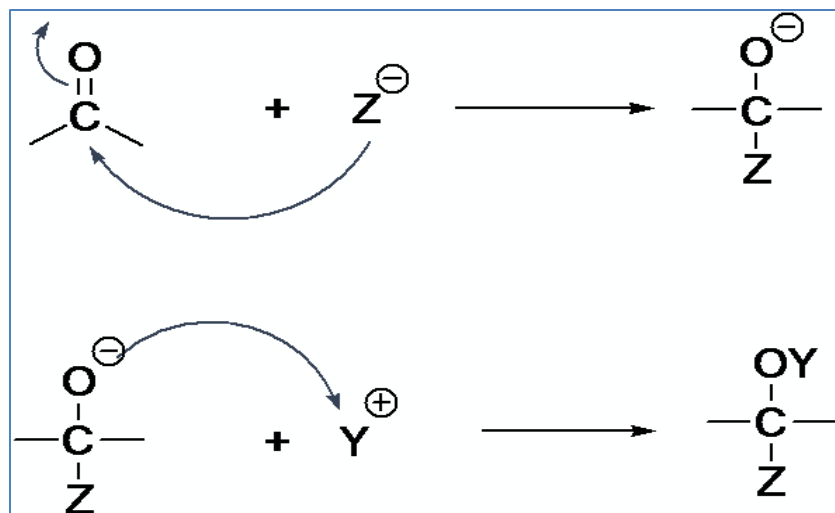
# General Reactions in carbonyl compounds

## 1- Nucleophilic Addition Reactions

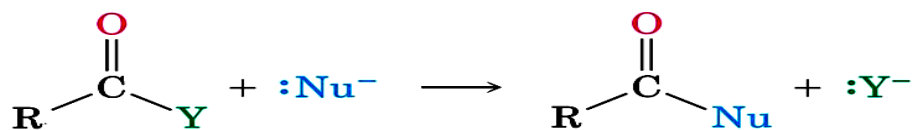


Occur in aldehyde and ketones

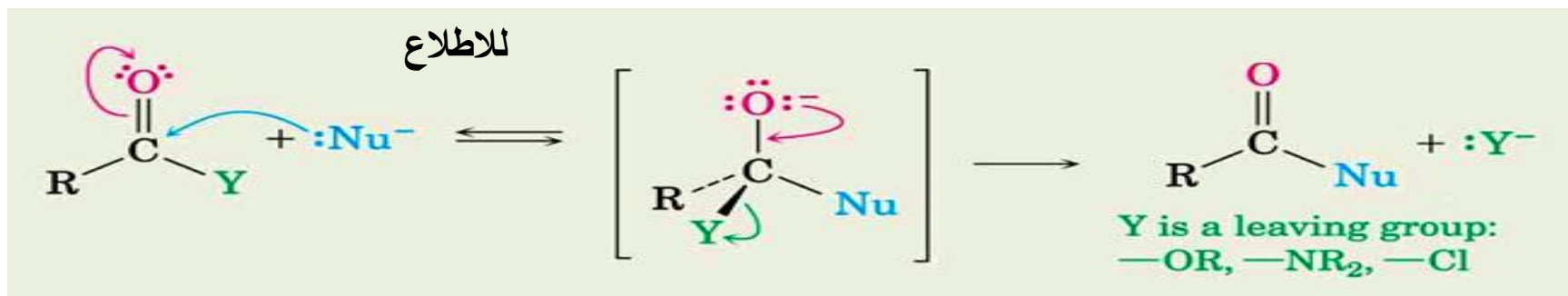
### Mechanism



## 2- Nucleophilic Substitution Reactions:



These type of reactions occur in carboxylic acids and esters.



## Aldehydes & ketones

➤ They are carbonyl compounds that contain C=O group.

They are similar in most properties such as:

- 1) They are polar molecules, so they have higher boiling points than alkenes of similar molecular weight but have lower boiling points than alcohols of similar molecular weight.
- 2) They undergo nucleophilic addition reactions.

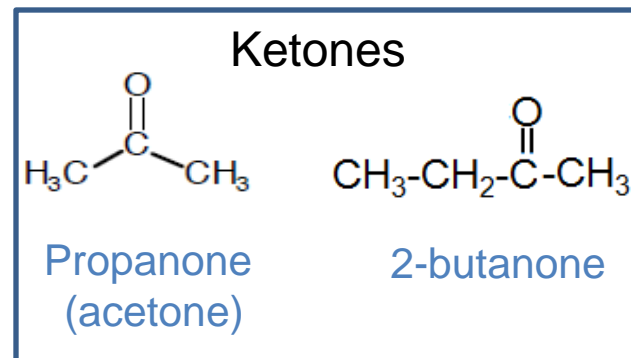
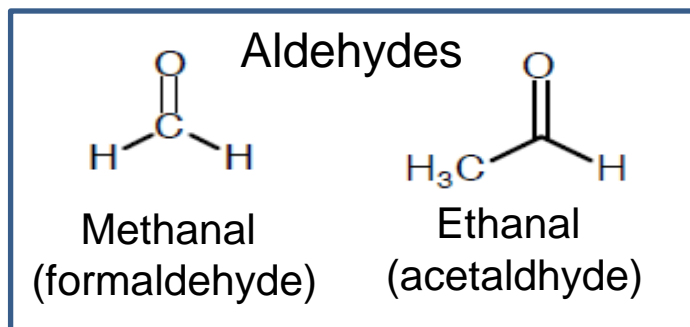
➤ But because aldehydes contain H atom attached to the C=O, there are some differences between them such as:

- 1) Aldehydes are quite easily oxidized, but ketones are oxidized with difficulty.
- 2) Aldehydes are more reactive than ketones toward nucleophilic addition.

# Aldehydes & ketones

## ➤ Nomenclature:

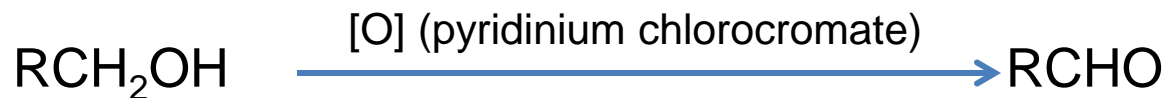
- In aldehydes: replace the (e) in alkane by (al), but in ketones, replace it by (one)



## ➤ preparation:

1- *From oxidation of alcohols:*

primary alcohol gives aldehyde, secondary alcohol gives ketones:



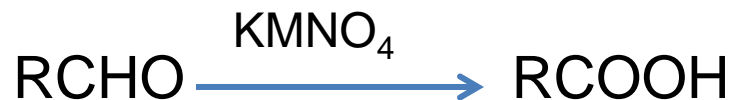
2- *From reduction of carboxylic acid:*





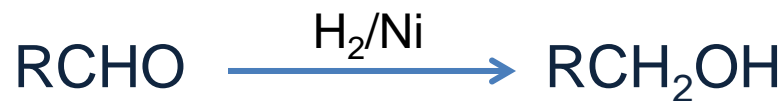
## Chemical Reactions of aldehydes & ketones

➤ **Oxidation:**, aldehydes are oxidized to carboxylic acids by mild oxidizing agents, but ketones are not:



- **Reduction:**, by reducing agents,

❖ Aldehydes are reduced to primary alcohols:



❖ Ketones are reduced to secondary alcohols:



# Uses of Aldehydes and Ketones:

## ➤ Uses of Aldehydes:

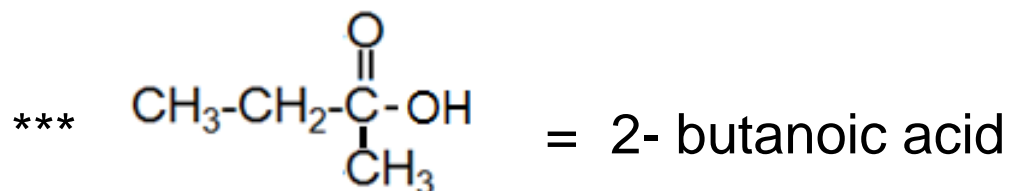
- Around 6 millions tons of formaldehyde produces every year. It is mostly used in the formation of resins, when mixed with melamine, urea, etc.
- 2.5 millions tons butyraldehyde are produce every year. It is mainly used as a plasticizer.
- Some other aldehydes are used as ingredients in flavors and deodorants.

## ➤ Uses of Ketone:

- Acetone, and cyclohexanone, are the most important ketones.
- Ketones are produced at very high scale to be used in medicine ,solvents, or in polymers synthesis.

## II) Carboxylic Acids

- Organic compounds having one or more carboxylic groups.  $\begin{array}{c} \text{O} \\ \parallel \\ -\text{C}-\text{OH} \end{array}$
  - This group is composed of two functional groups:  
carbonyl group  $-\text{C}=\text{O}$ , and the hydroxyl group  $-\text{OH}$
  - They are not strong acids as inorganic acids (HCl, HNO<sub>3</sub>...)
  - Their acid strength increases as the # of (COOH) increases.
  - Their IUPAC name is by replacing the letter (e) in the equivalent alkane, by the suffix (oic):
- \*  $\text{HCOOH}$  = methanoic acid (formic or ants acid),
- \*\*  $\text{CH}_3\text{COOH}$  = ethanoic acid (vinegar or acetic acid).



# Preparation of Carboxylic Acids

## 1- Oxidation of primary alcohol:



## 2- Hydrolysis of nitriles:

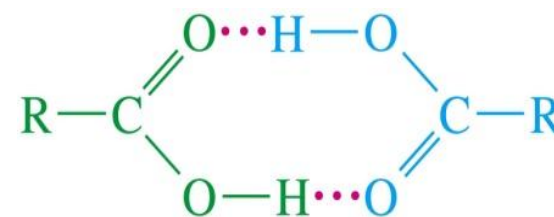


## Their physical properties:

1- First members are liquids, mild members are oily, and the highest members are solid.

2- Their solubility in water decreases with the length of the carbon chain.

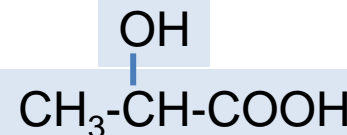
3- They have higher boiling points than similar alcohols, due to dimer formation.



hydrogen-bonded acid dimer

## Uses of Carboxylic Acid

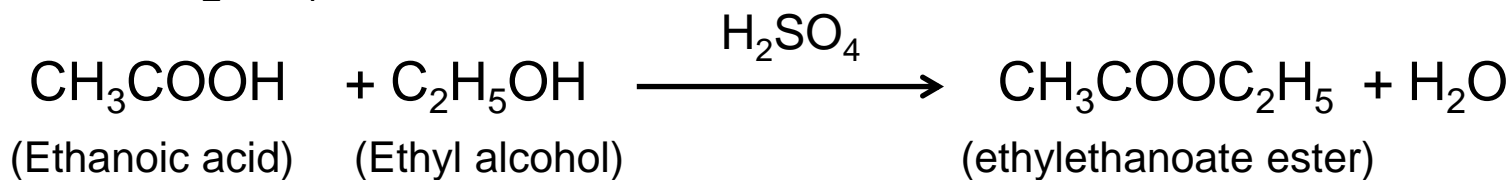
- ❖ Most of carboxylic acids are produced on a large scale for industrial purpose.
- ❖ In industry, carboxylic acids are used as additives or solvents in food production, drugs, and polymers, and some also used as a food preservative, chelating agent.
- ❖ Formic acid is used in manufacturing of dyes, insecticides, drug and plastic.
- ❖ Acetic acid is used in home as vinegar (4%), synthetic silk, dyes, and food additives.
- ❖ Lactic acid (found in milk) generated in human body as a result of hard effort, and causes a construction of muscles.
- ❖ Salysilic acid is used in the manufacture of Cosmetics and aspirin



Lactic acid

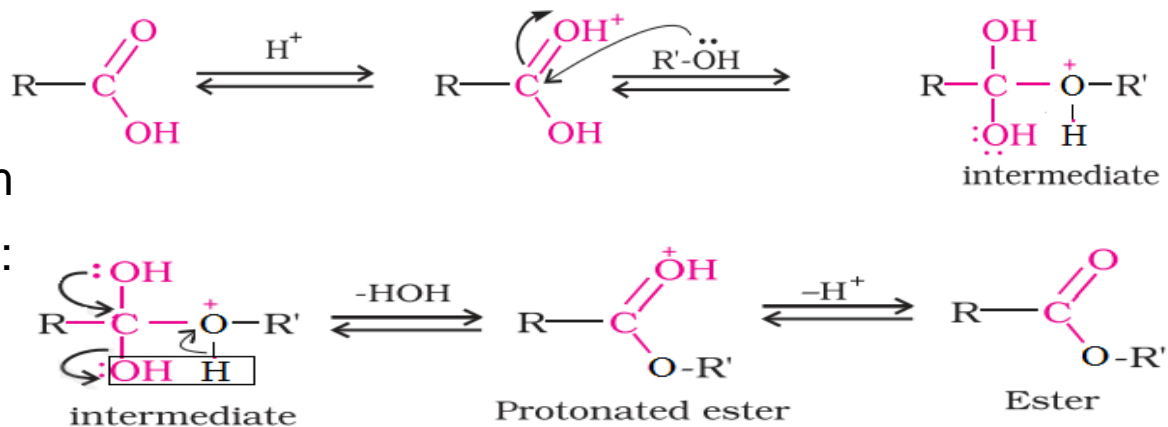
### III) Organic Esters

➤ Organic compound produced from reacting carboxylic acids with alcohols in presence of conc  $\text{H}_2\text{SO}_4$ :



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The reaction mechanism of the ester formation is:



➤ Their names are derived from the name of acid and alkyl group of alcohol:

\*  $\text{HCOOCH}_3$  = **methyl** methanoate ester,

\*\*  $\text{CH}_3\text{COOC}_2\text{H}_5$  = **ethyl**ethanoate ester

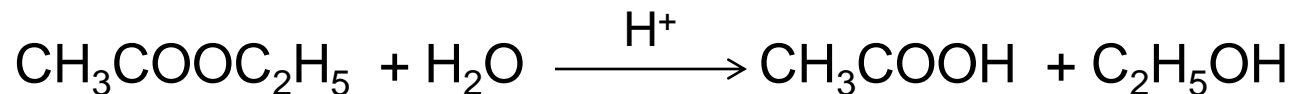
# Properties of Organic Esters

## Their physical properties:

- 1- Their B.P is lower than that of carboxylic acids or alcohol due to the absence of H-bonding.
- 2- Their odor is pleasant, so they are used in preparation of perfumes & flavors.
- 3- They also used in producing polyesters, dacron, and drugs such as Aspirin.

## Their chemical properties:

### 1- Acid hydrolysis:



### 2- Base hydrolysis (saponification):

