



Alcohols and Phenols

Chapter 7

1432-2011

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Chapter Head Lines

- ▶ Introduction
- ▶ Types and Classifications.
- ▶ Nomenclature of Alcohols and Phenols.
- ▶ Physical Properties.
- ▶ Acidity of Alcohols and Phenols.
- ▶ Preparation of Alcohols and Phenols:
 - A. Preparation of Alcohols
 - 1- Hydration of Alkenes
 - 2- Hydroboration-Oxidation “ **anti-Markovnikov**”
 - B. Preparation of Phenols

► Reactions of Alcohols and Phenols:

I. Alcohols and Phenols as acids: Salt Formation

II. Oxidation

A. Oxidation of Alcohols

B. Oxidation of Phenols

III. Reactions Involving Carbon-Hydroxyl Bond Breaking

A. Formation of ester

B. Replacement of the OH Group by Halide: Alkyl Halides

1- Reaction with Hydrogen halides HX

2- Reaction thionylhalide SOX_2

3- Reaction with phosphorus trihalide PX_3 or PX_5

C. Dehydration of Alcohols: Formation of Alkenes

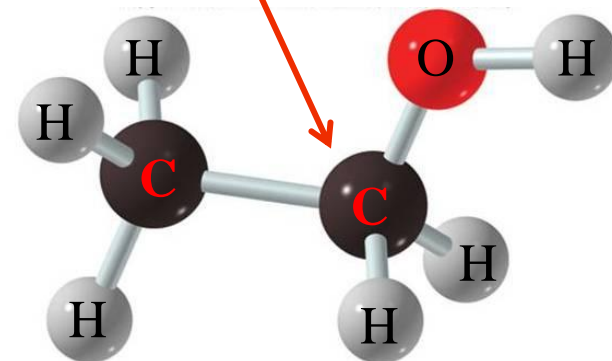
Alcohols

Introduction

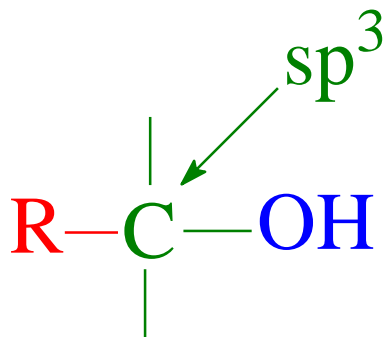
-Alcohols are characterized by *the hydroxyl group* -OH

-The general formula for Alcohols is $\text{R}-\text{OH}$

Carbinol carbon



The C-OH group is called *the carbinol group*, and the carbon of this group is called “*carbinol carbon*”.

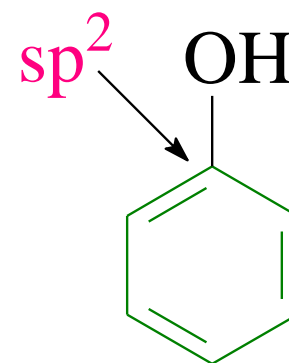
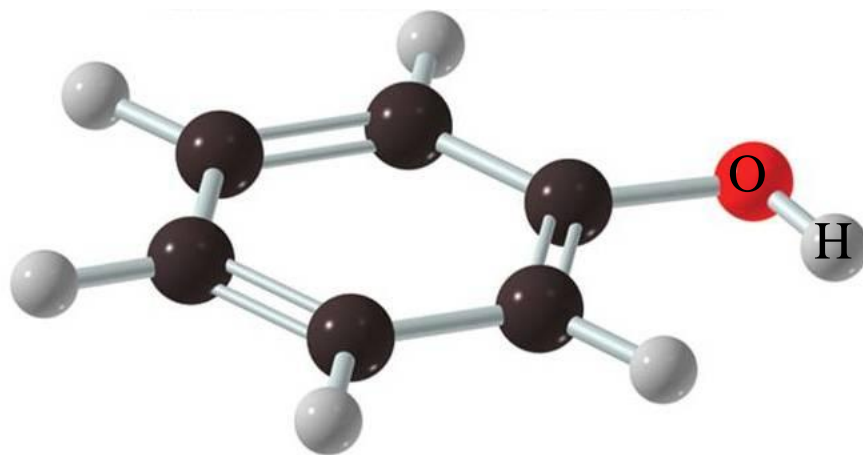


As all alcohols are the compounds containing hydroxyl group (-OH) attached to *the alkyl group*, so the *carbinol carbon* hybridization is sp^3

Phenols

Or, Aryl alcohols

- are hydroxyl derivatives of aromatic hydrocarbons, which are derived by replacing hydrogen atom attached to sp^2 hybridized carbon atom(s) of benzene ring by hydroxyl group.

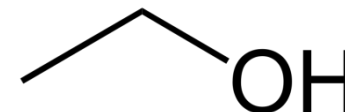
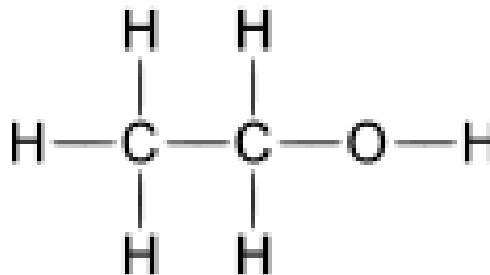


Phenols , ArOH

Types Of Alcohols

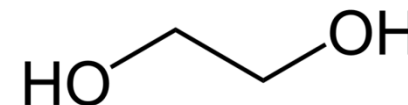
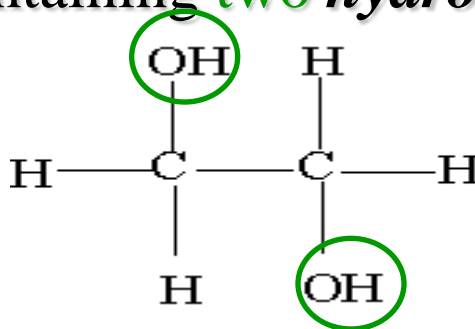
1. Monohydroxyls: containing **one hydroxyl group**.

Example; ethanol (C_2H_5OH)



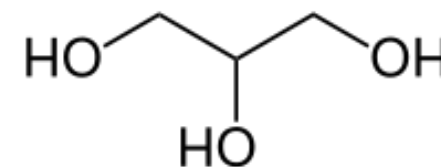
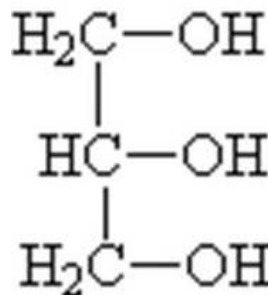
2. Dihydroxyls (glycols): containing **two hydroxyl groups** connected by different carbon atoms

Example; Ethylene glycol (CH_2OH-CH_2OH).



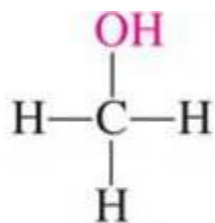
3. Polyhydroxyls: containing **more than two hydroxyl groups** on different carbon atoms

Example; 1,2,3-propanetriol ($CH_2OH-CHOH-CH_2OH$).

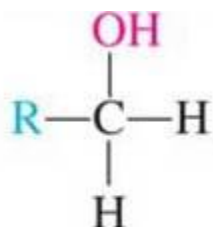


Classification of Monohydroxyl Alcohols

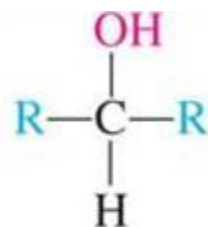
➤ *The mono hydroxyl alcohols* can be classified into three types according to the type of the carbon atom connected to *the hydroxyl group*:



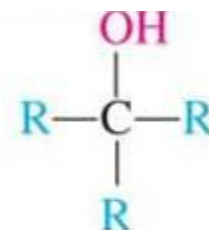
Methyl alcohol



1° Alcohol



2° Alcohol



3° Alcohol



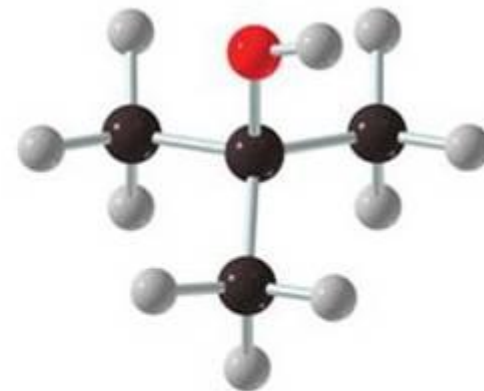
Methanol
(methyl alcohol)



Ethanol
(1° alcohol)



2-Propanol
(2° alcohol)



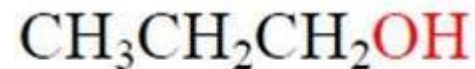
2-Methyl-2-propanol
(3° alcohol)

Nomenclature

1) Common Nomenclature (Alkyl + alcohol)

- You can use both **the common** and **IUPAC systems** to name **alcohols**.
- In ***the common system***, you **name an alcohol** by listing **the alkyl group** and adding the word ***alcohol***.

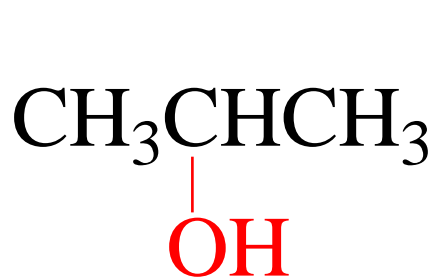
-Some examples of alcohols and their common names:



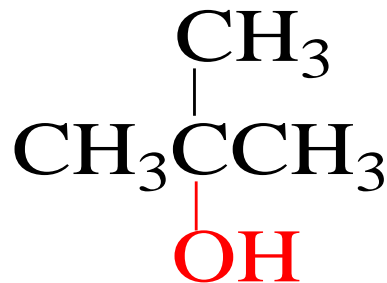
Methyl alcohol

Ethyl alcohol

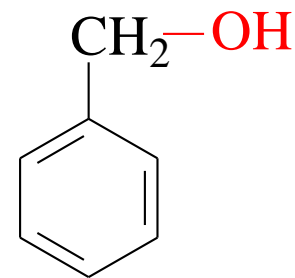
propyl alcohol



Isopropyl alcohol



t-butyl alcohol



Benzyl alcohol



Vinyl alcohol

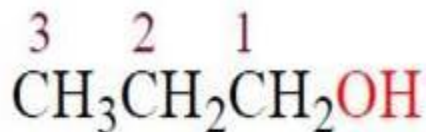


Allyl alcohol

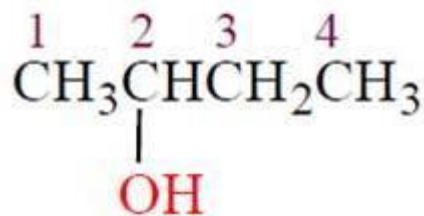
2) IUPAC Nomenclature

- 1) Select the **longest continuous carbon chain** *to which the hydroxyl is directly attached*.
- 2) Change the name of the **alkane** corresponding to this chain by dropping the final **-e** and adding the *suffix -ol*
- 3) Number the longest continuous carbon chain so as to give the carbon atom bearing the hydroxyl group the lower number.

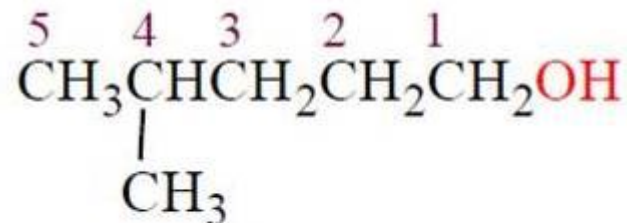
Examples



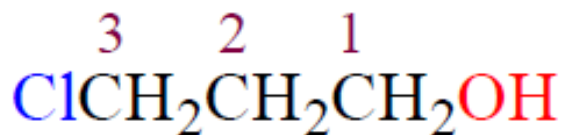
1-Propanol



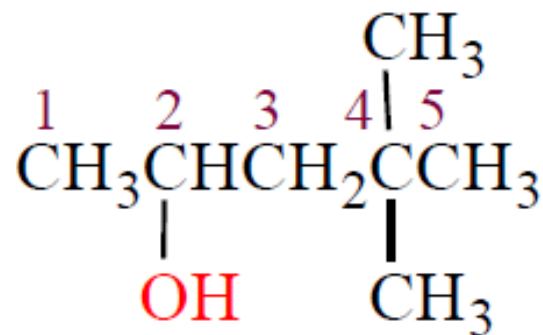
2-Butanol



4-Methyl-1-pentanol
(not 2-methyl-5-pentanol)

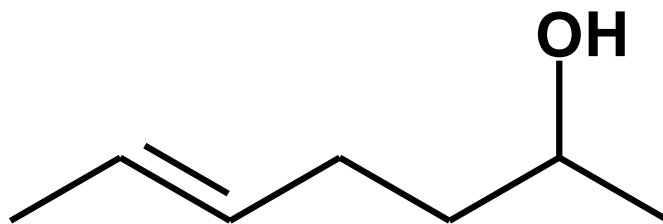


3-Chloro-1-propanol



4,4-Dimethyl-2-pentanol

4) **OH group** is preferred over the **double** or **triple bond** in numbering.

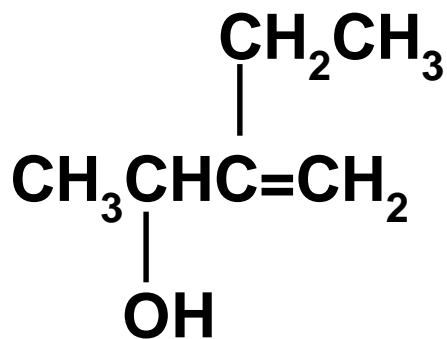


5-Hepten-2-ol

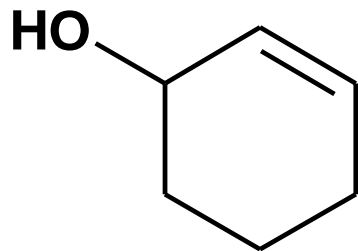
notice the removal of the (e)

5) If a compound contains both **OH** and a **double** or **triple bond**, choose the chain that *include them both* even if this is not the longest chain.

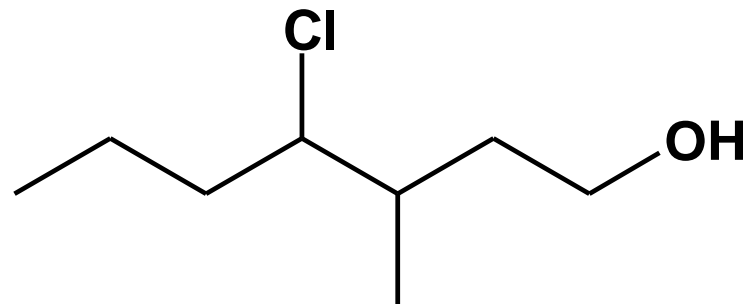
e.g.



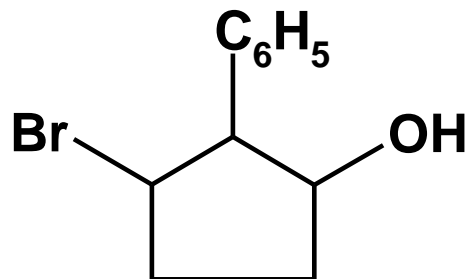
3-Ethyl-3-buten-2-ol



2-Cyclohexenol



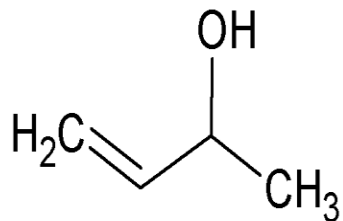
4-Chloro-3-methyl-heptan-1-ol



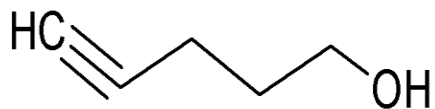
3-Bromo-2-phenyl cyclopentanol



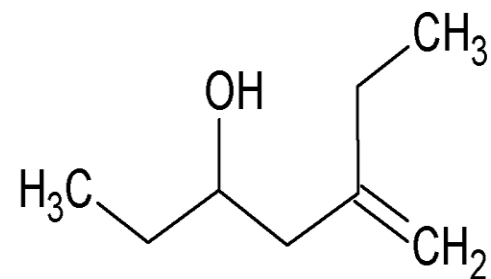
3-Butyn-2-ol



3-buten-2-ol

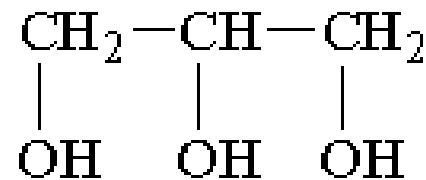
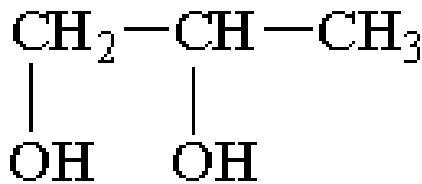
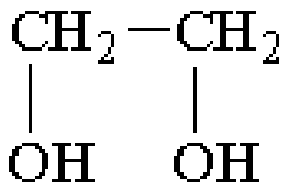


4-Pentyn-1-ol



5-Ethyl-5-hexen-3-ol

- In the IUPAC system, the suffix **diol** is added to the name of the parent hydrocarbon when **two hydroxyl groups** are present, and the suffix **triol** is added when there are **three OH** groups.
- Common names, **two OH groups** on adjacent carbons are known as **1,2-glycols**.



IUPAC **1,2-Ethandiol**

1,2-Propanediol

1,2,3-Propanetriol

Common **Ethylene glycol**

propylene glycol

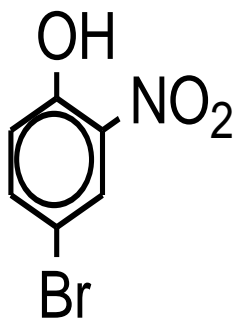
glycerol or glycerin

Nomenclature Of Phenols

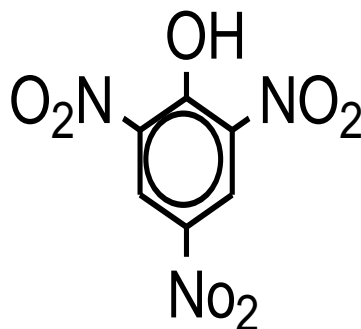
- Compounds that have a **hydroxyl group** attached directly to a **benzene ring** are called **phenols**.
- The **ortho**, **meta**, **para** system is used in common names.
- While the numbering system is employed in IUPAC names and in this case numbering of the ring begins at the hydroxyl-substituted carbon and proceeds in the direction of the next substituted carbon that possesses the lower number.



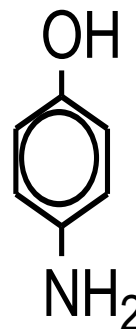
Phenol



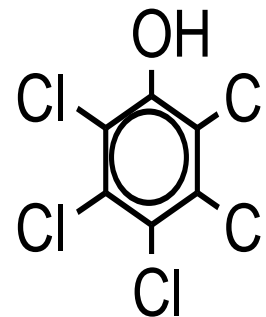
4-Bromo-2-nitrophenol



2,4,6-Trinitrophenol



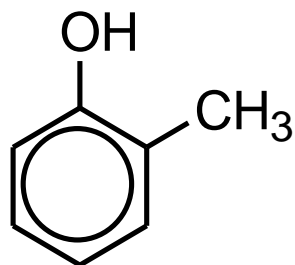
4-Aminophenol



2,3,4,5,6-Pentachlorophenol

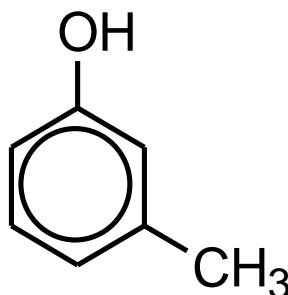
Common name: Picric acid

➤ Some phenols have **common names** as shown in the following examples



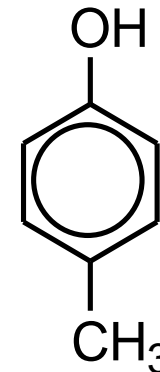
IUPAC: 2-Methyl-phenol

Common: *o*-Cresol



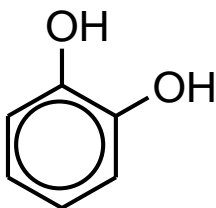
IUPAC: 3-Methyl-phenol

Common: *m*-Cresol



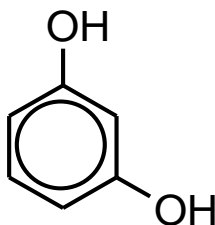
IUPAC: 4-Methyl-phenol

Common: *p*-Cresol



IUPAC: 2-Hydroxyphenol
or 1,2-Benzenediol

Common: Catechol



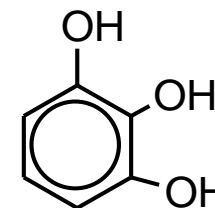
IUPAC: 3-Hydroxyphenol

Common: Resorcinol



IUPAC: 4-Hydroxyphenol

Common: Hydroquinone



IUPAC: 2,3-Dihydroxyphenol

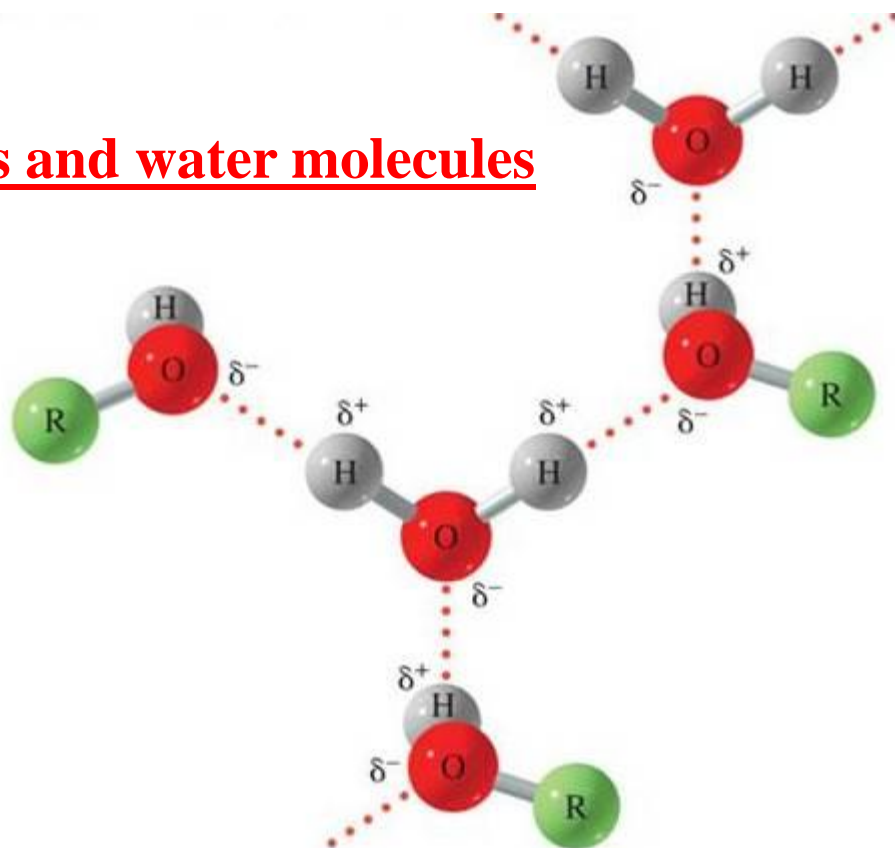
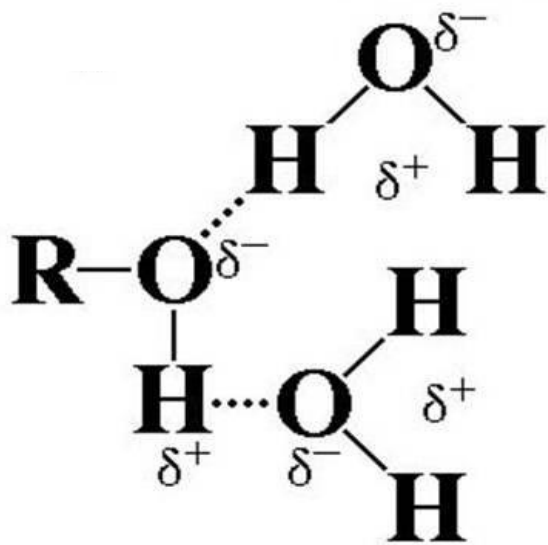
Common: Pyrogallol

Physical Properties of Alcohols & Phenols

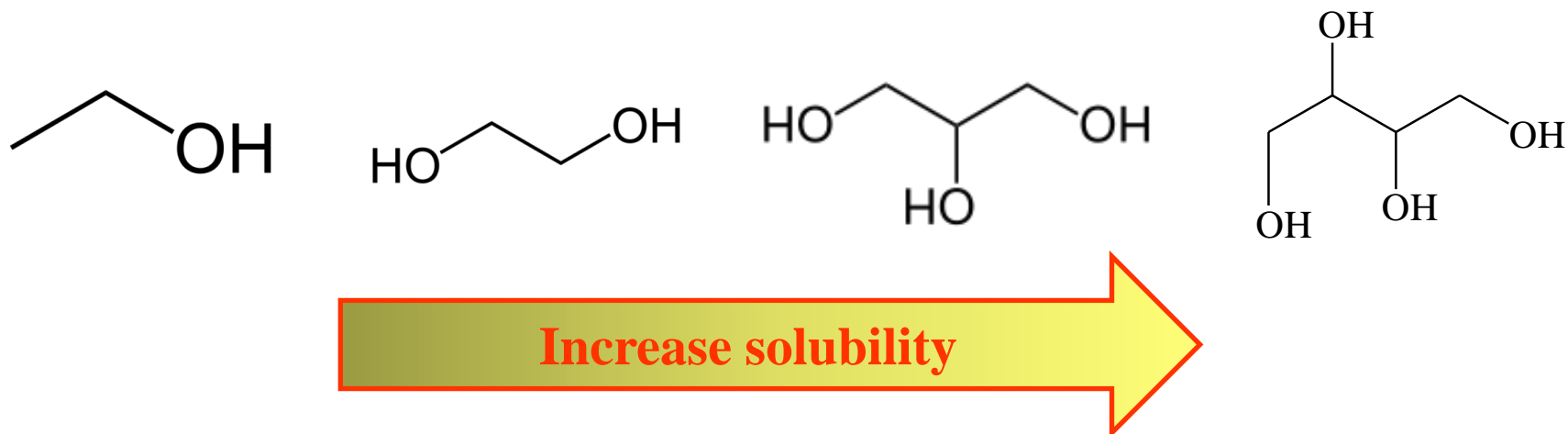
- The lower members of **alcohols** are colorless, **volatile liquids** with a **characteristic alcoholic smell** and **burning taste** whereas **higher alcohols** are **odorless** and **tasteless**.
Higher alcohols having 12 or more carbon atoms are colorless waxy solids.
- **Phenols** are **colorless**, **crystalline solids** or **liquids**.
- **Solubility of alcohols** The first three members are completely **miscible** with **water**. The solubility rapidly **decreases** with **increase in molecular mass**. The **higher members** are almost **insoluble** in **water** but are soluble in organic solvents like **benzene**, ether etc.

•The solubility of lower alcohols is due to the existence of **hydrogen bonds** between **water** and **polar -OH group** of alcohol molecules. **Phenols** too are sparingly soluble in **water**. The **-OH group** in **alcohols** and **phenols** contain a hydrogen bonded to an electronegative oxygen atom. Thus they form hydrogen bonds with water molecules.

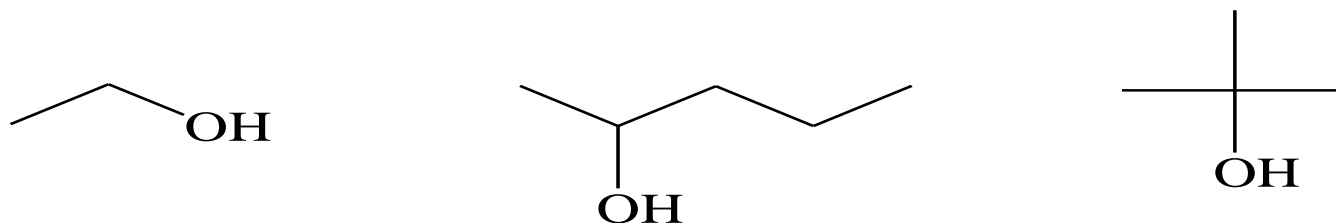
Hydrogen bonding between alcohols and water molecules



- The number of **hydroxyl groups** increases the solubility.



- The solubility **increases** with **branching of chain**.



- **Phenols** are sparingly soluble in **water** but readily **soluble in organic solvents** ..

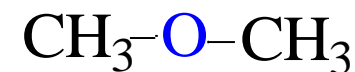
- **Boiling points of alcohols** Boiling point of alcohols are **much higher** than those of **alkenes**, **halo alkenes** or **ethers** of comparable molecular masses.



Eyhanol



n-propane



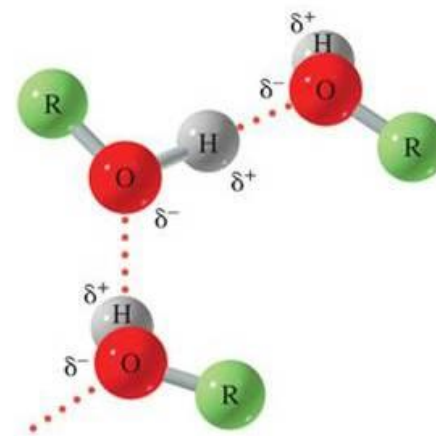
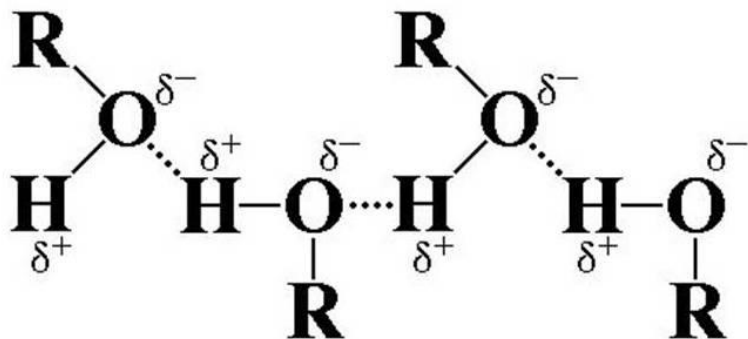
Dimethylether

Mol wt = 46; bp= 78°C

Mol wt = 44; bp= -42°C

Mol wt = 46; bp= -24°C

- This is because in **alcohols intermolecular hydrogen bonding** exists due to which a **large amount of energy** is required to break these bonds.

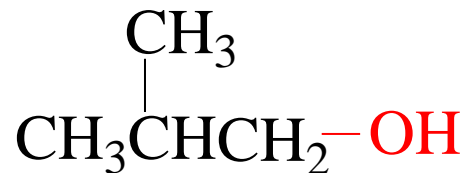


- Among isomeric alcohols, the boiling point decreases with increase in branching in the alkyl group.



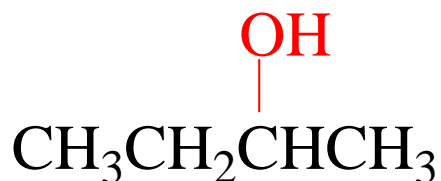
1-Butanol

(mol wt = 74; bp = 118°C)



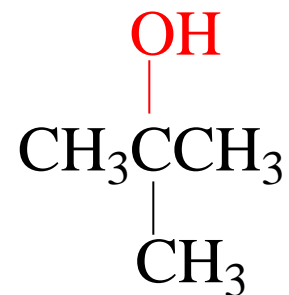
2-Methyl-1-propanol

(mol wt = 74; bp = 108°C)



2-Butanol

(mol wt = 74; bp = 99.5°C)



2-Methyl-2-propanol

(mol wt = 74; bp = 83°C)

- Boiling points of 1° alcohol > 2° alcohol > 3° alcohol
- boiling points increase with the increase of number of hydroxyl groups .

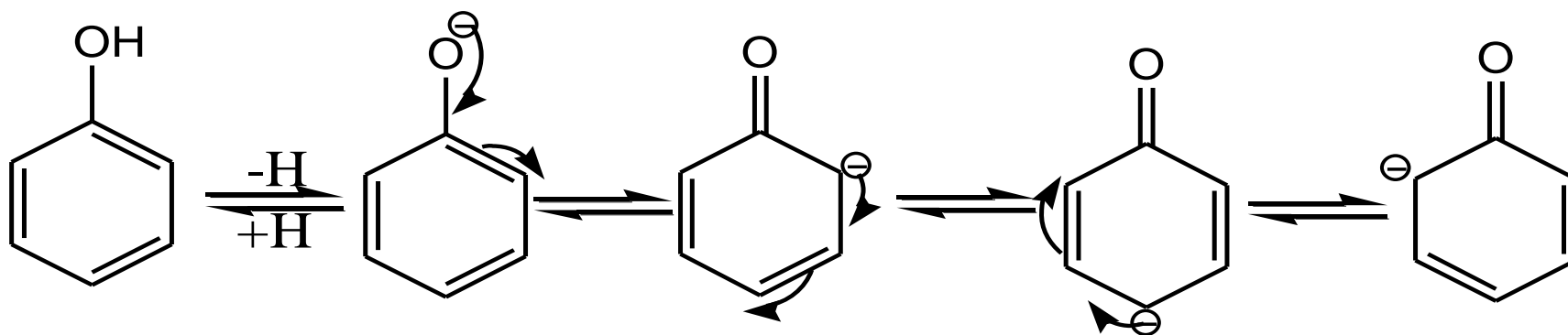
Acidity of Alcohols & Phenols

- Due to the electronegativity of the O atoms, alcohols are *slightly acidic* (pKa 16-18)
- The anion derived by the deprotonation of an alcohol is the **alkoxide**.
- Alcohols and phenols have weak acidic properties.

- Phenols are **much stronger acids** than alcohols.

Why ??

Because the negative charge in oxygen is dispersed by resonance through the benzene ring.

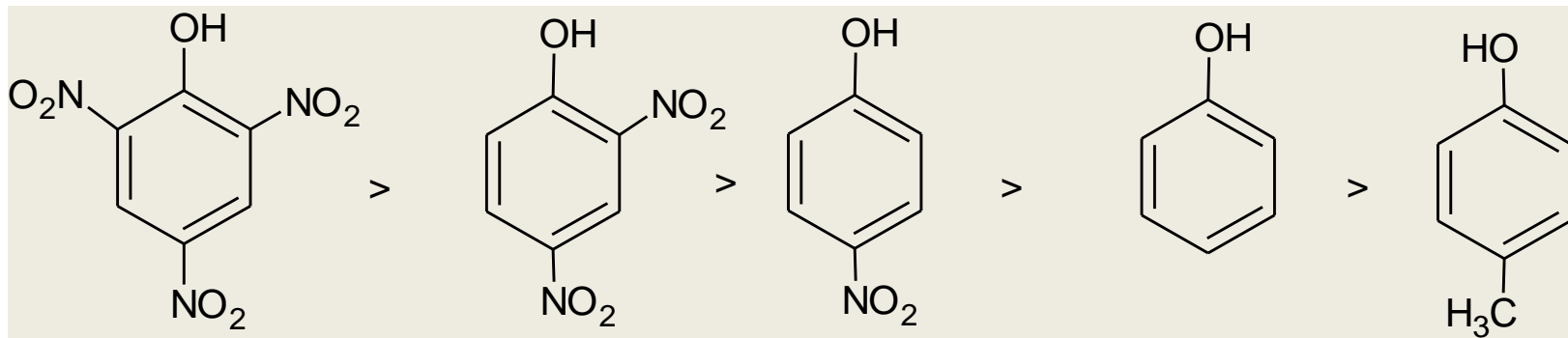


Resonance structures of phenoxide anion

Effect of substituents on the acidity of phenols

- Introduction of **electron-withdrawing groups (EWG)**, such as **NO₂** or **CN**, **X** on the ring **increases the acidity** of phenol.
- Also, introducing **electron-donating groups (EDG)**, such as **NH₂**, **R**, **OR** **decrease the acidity** of phenols.
- The electron **donating** or **withdrawing** effect is generally more pronounced if **the substituted** is **present at o- and p- position**.
- The **greater the number of electron withdrawing at o- and p- position**, **more in the acidic character** of phenol.

Acidity order



Example: In each of the following pairs of compounds, indicate which is more acidic.

(a) *p*-chlorophenol or *p*-nitrophenol

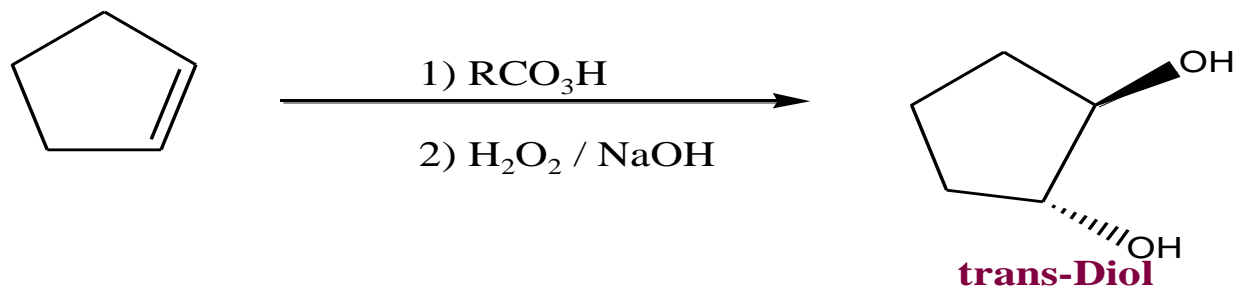
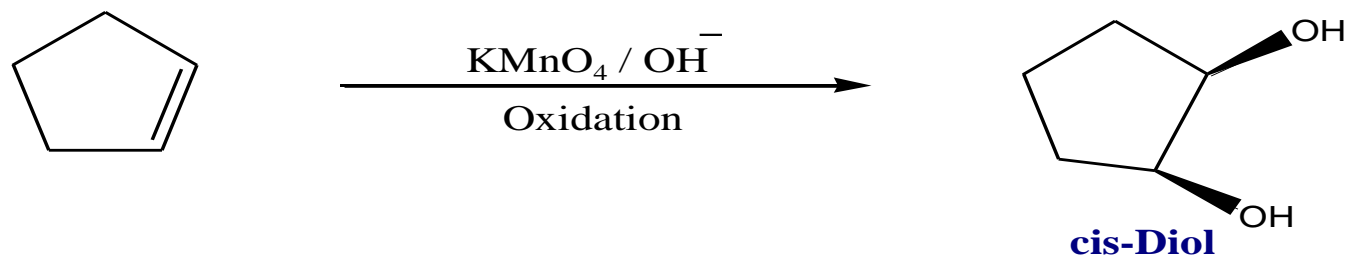
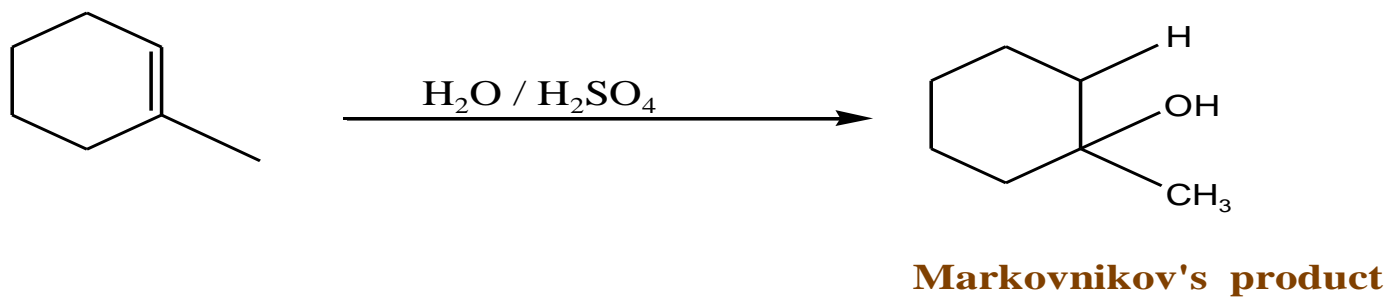
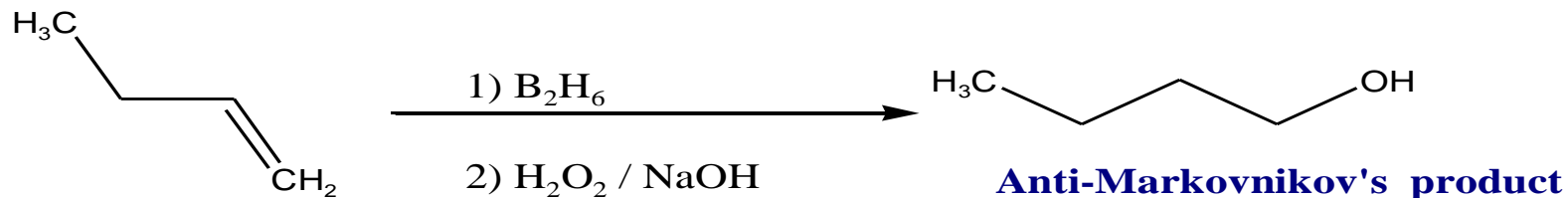
(b) *o*-Iodophenol or *p*-Iodophenol

(c) *o*-Creasol or *o*-Nitrophenol

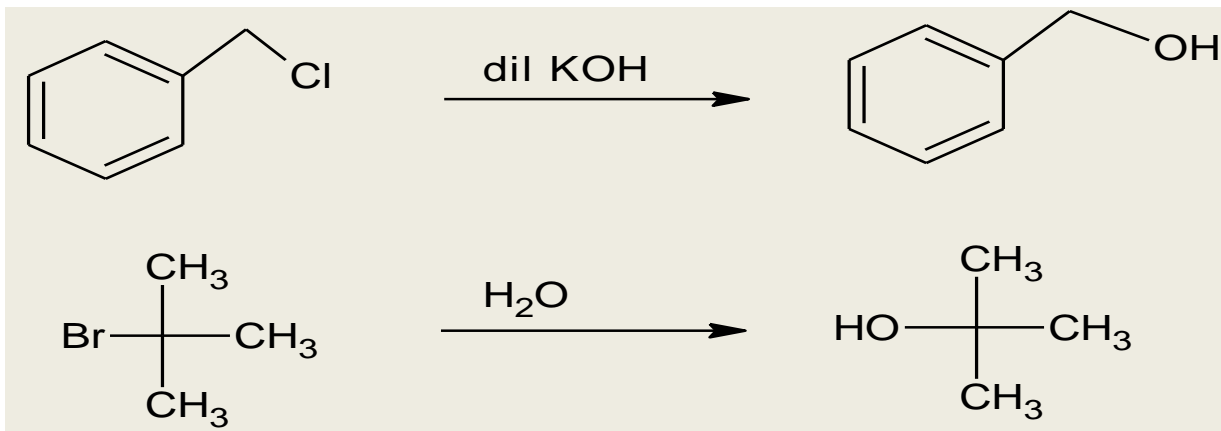
(d) *o*-Nitrophenol or *m*-Nitrophenol

A. Preparation of alcohols:

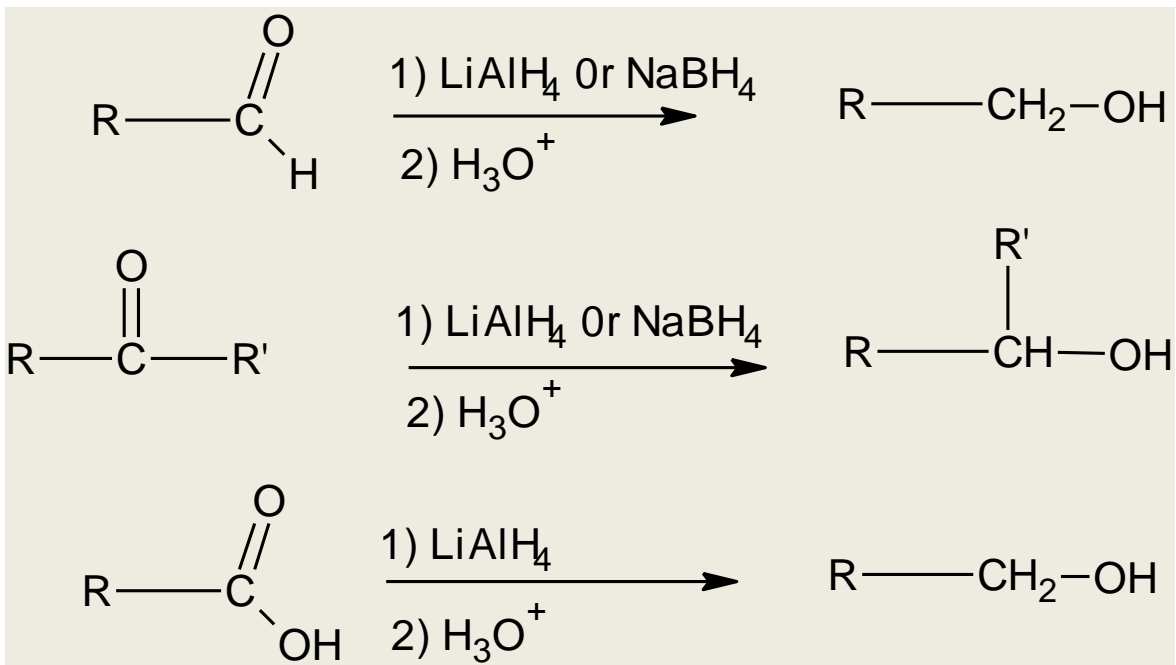
1- From alkenes



2- From alkyl halide

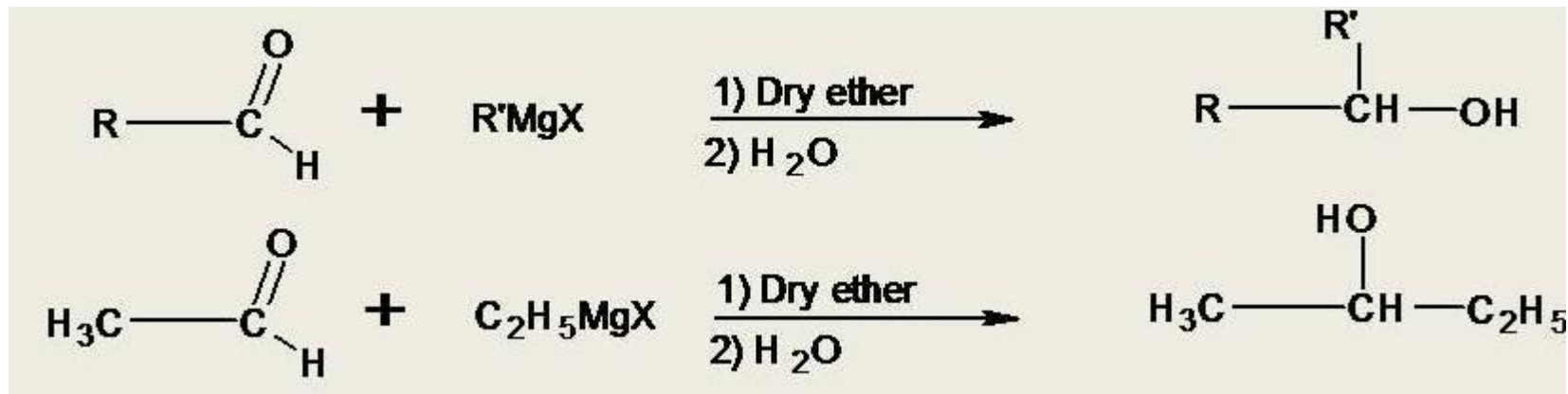


3- From aldehyde, ketone and carboxylic acid (Reduction reaction)

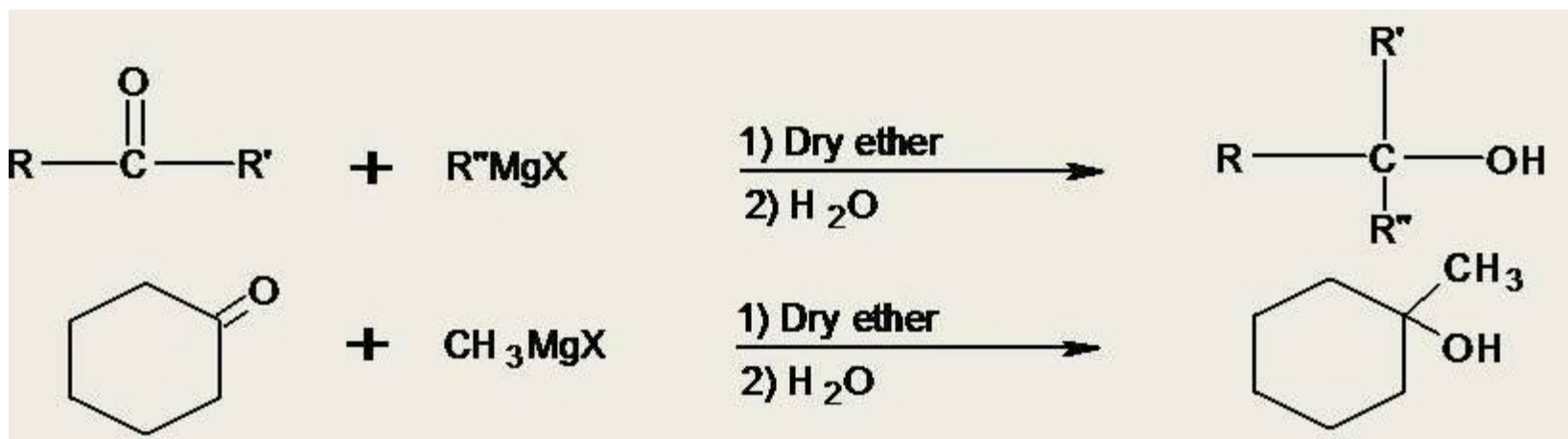


4- From Grignard reagent

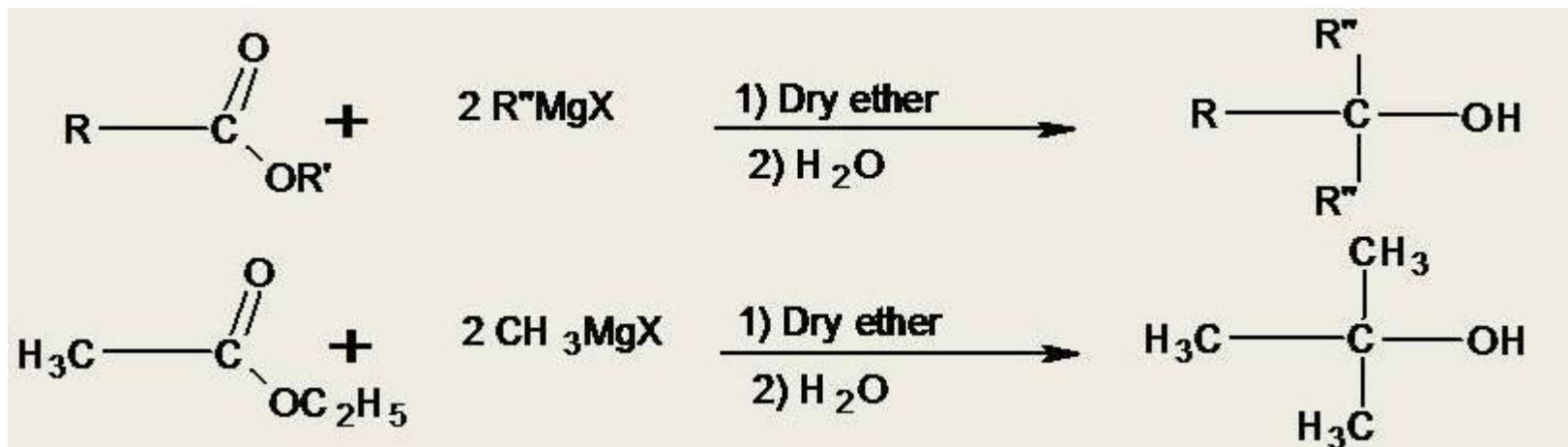
Reaction with Aldehyde



Reaction with ketone

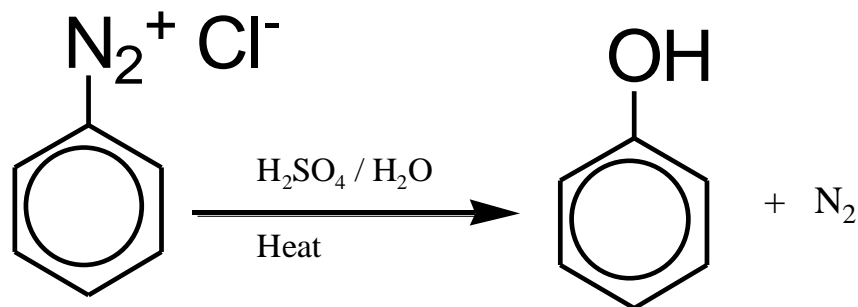


Reaction with Ester

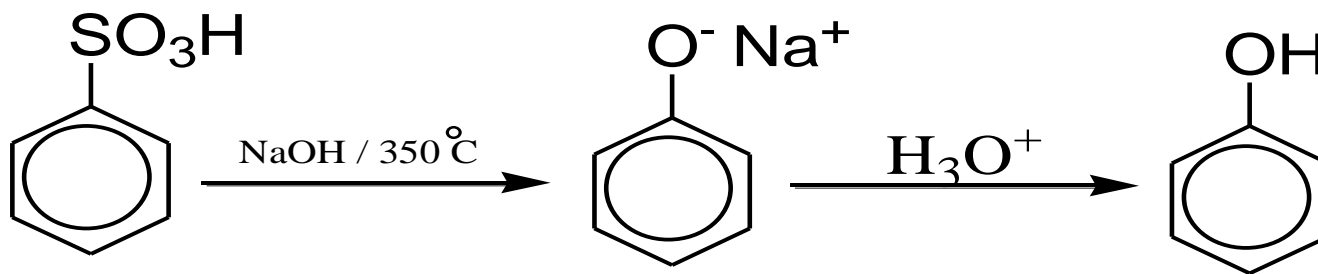


Preparation of Phenols

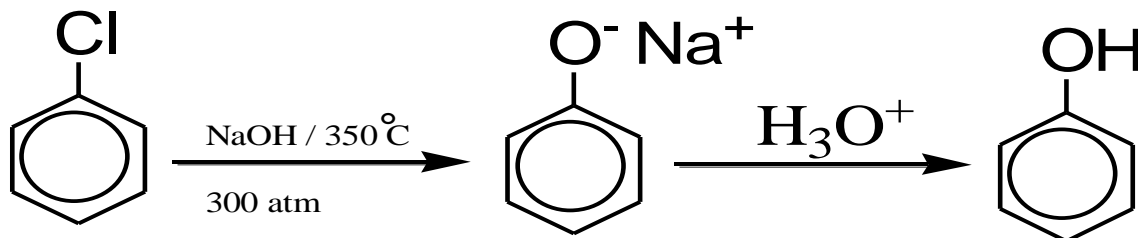
1- Hydrolysis of Diazonium salts



2- Fusion of sodium with benzene-sulfonates:

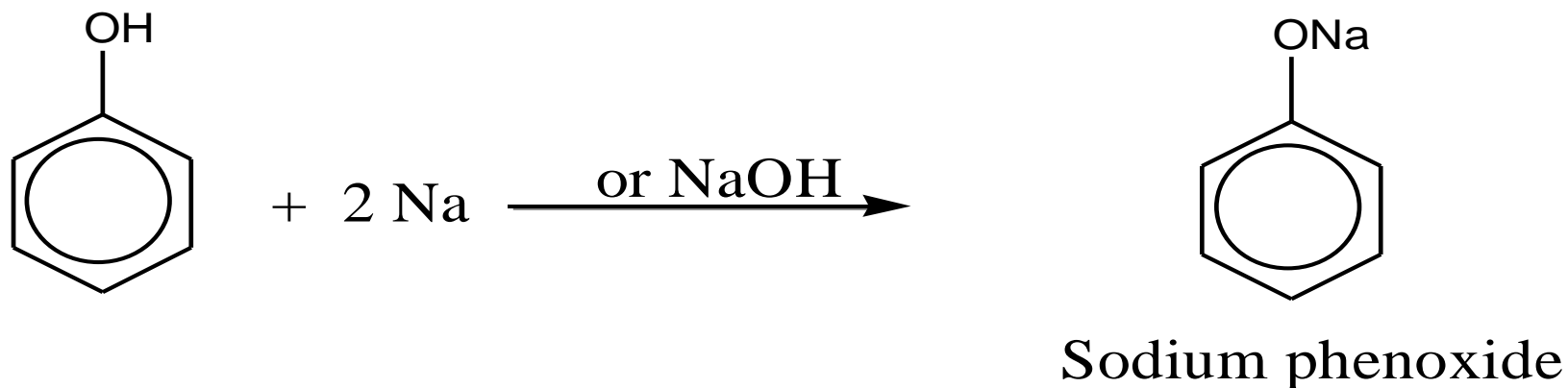


3- From alkyl halide:



Reaction of Alcohols and Phenols

1) Salt Formation By Reaction With Active Metals



2) Elimination Of Water (Dehydration)

Reagent/catalyst

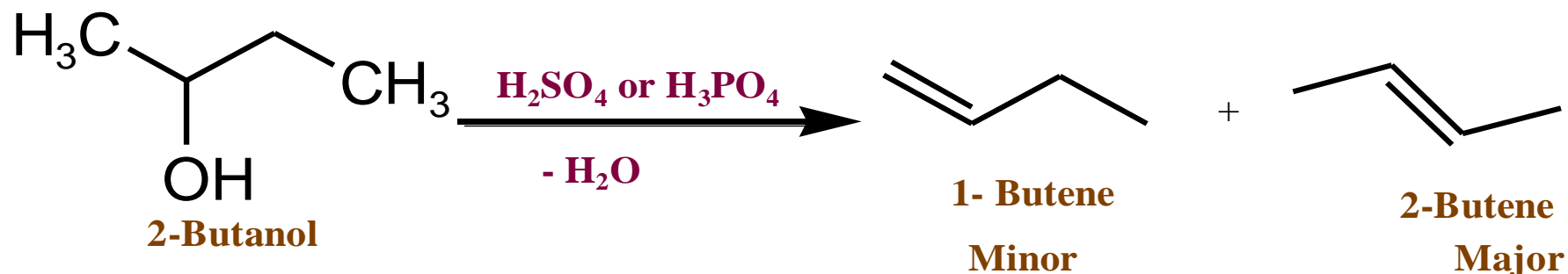
Conditions

Product

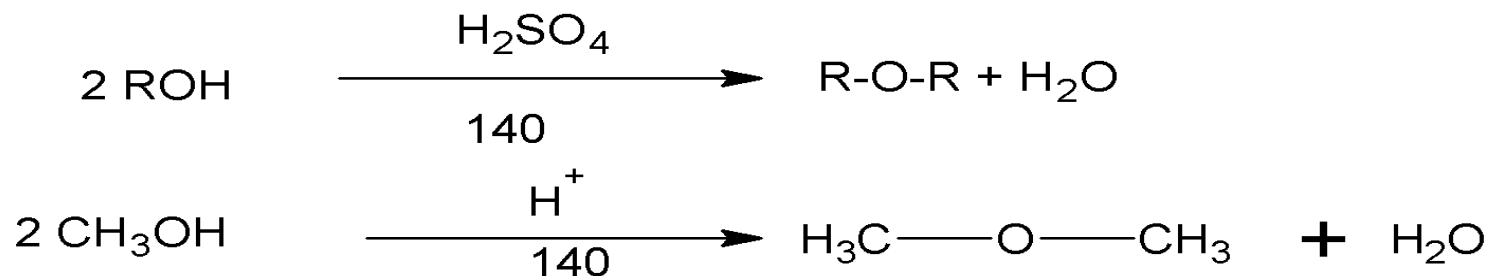
conc. sulphuric acid (H_2SO_4)

reflux at 180°C

alkene

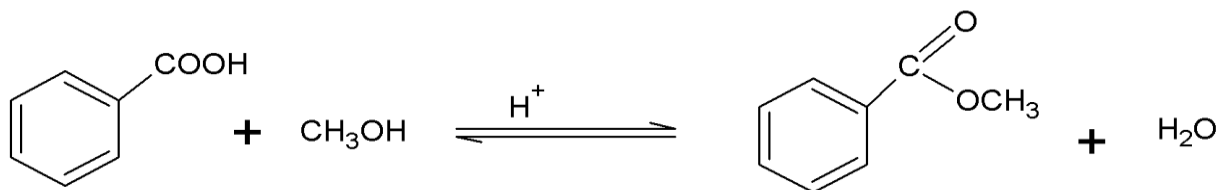
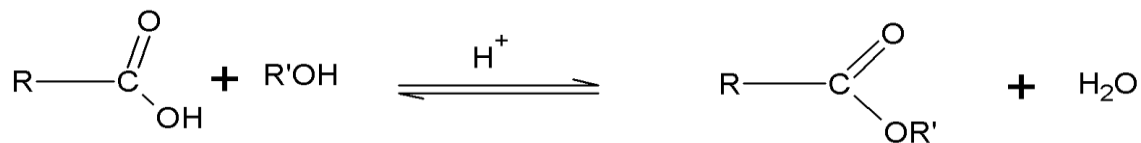


While dehydration of alcohols at lower temperature will give ethers



3) Ester Formation

- Carboxylic acid + alcohol in presence of strong acid catalyst (e.g. conc. H_2SO_4) produces esters



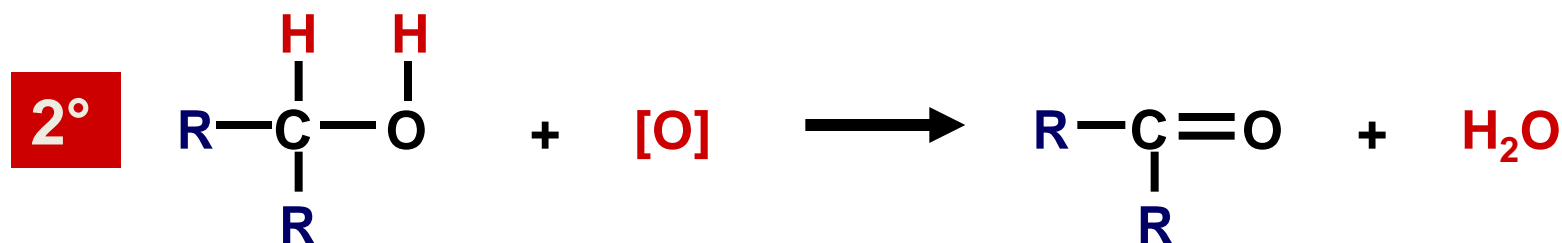
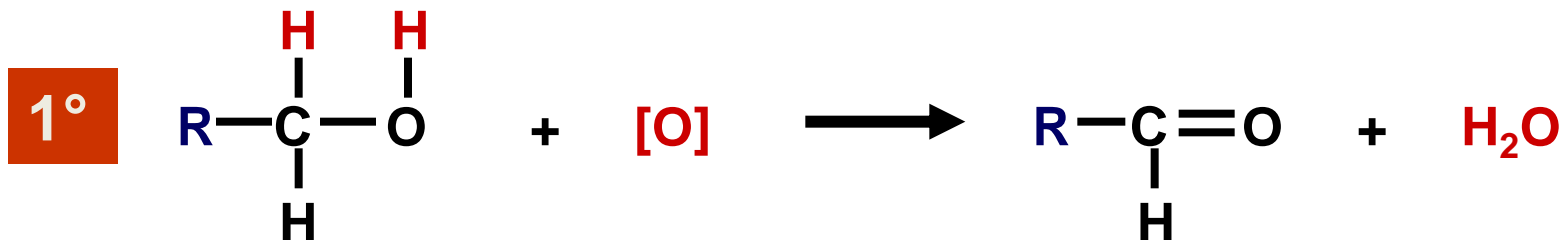
4) Alkyl Halides Formation



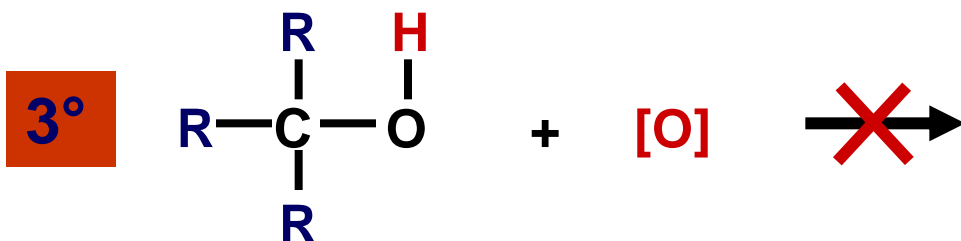
5) Oxidation Of Alcohols

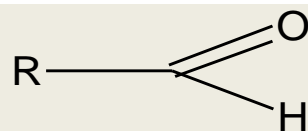
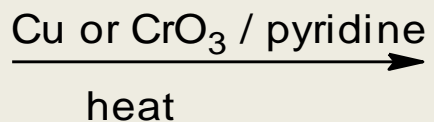
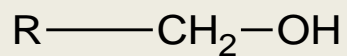
Alcohols can be oxidised depending on their class

➤ For oxidation to take place easily you must have two hydrogen atoms on adjacent C and O atoms.



This is possible in **1°** and **2°** alcohols but not in **3°** alcohols.

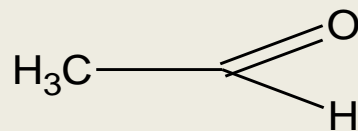
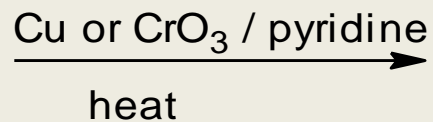
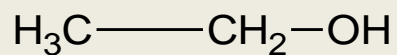




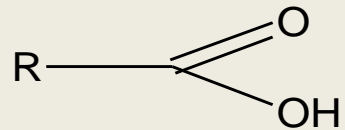
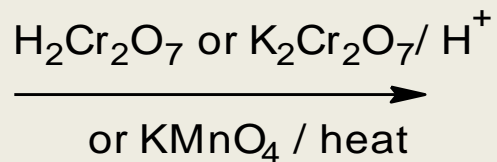
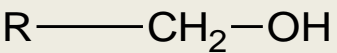
weak oxidizing reagent

Primary alcohol

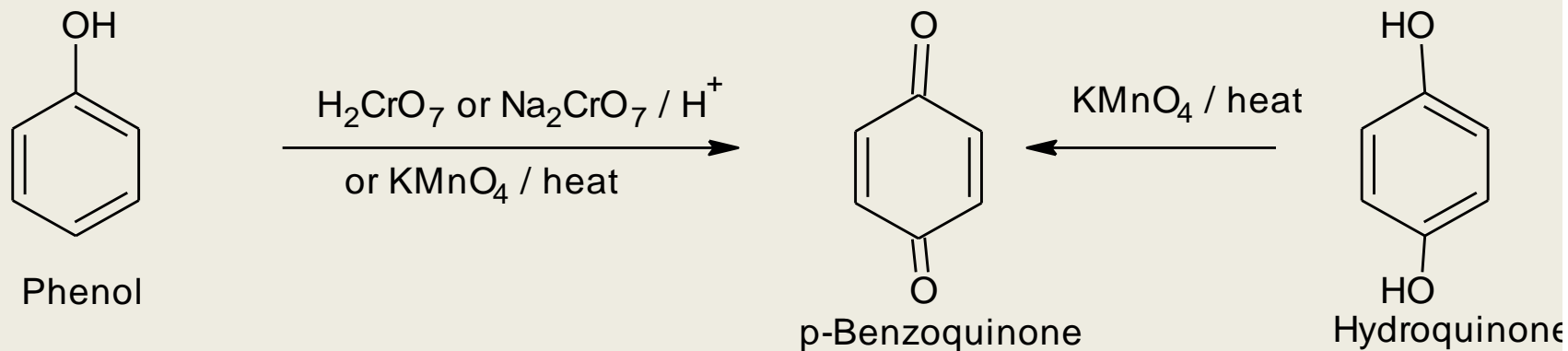
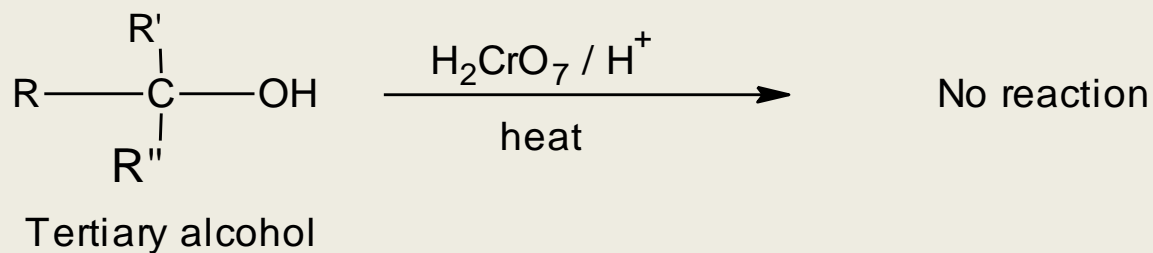
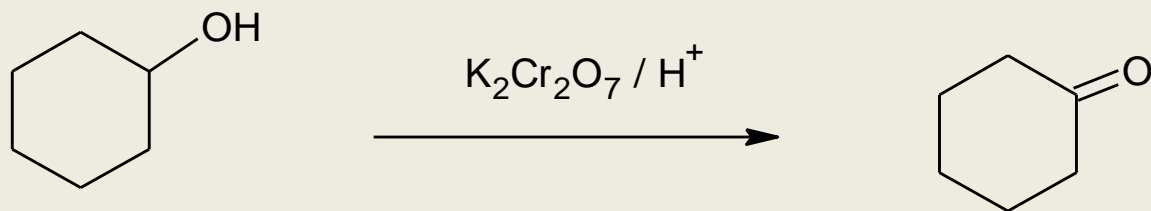
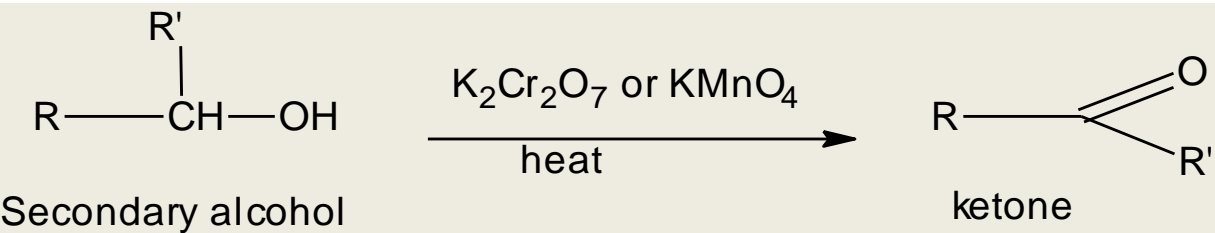
aldehyde



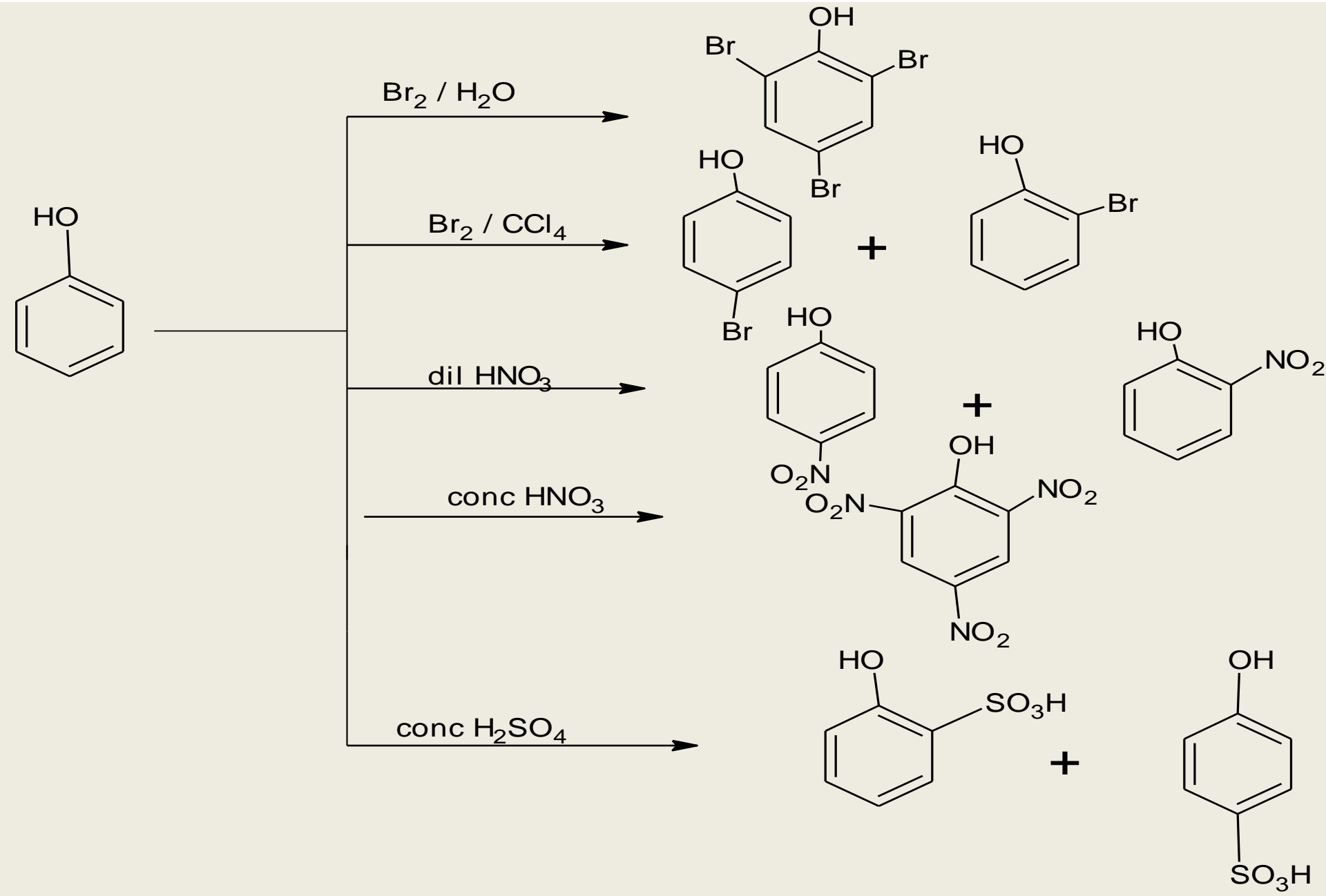
Primary alcohol



carboxylic acid




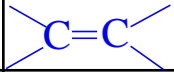

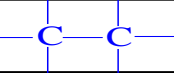
6- Reaction of aromatic ring of phenols



Functional Group Precedence in IUPAC nomenclature

When **two or more functional groups** are present in a molecule, that group highest in the table is used as the parent (suffix) in the IUPAC name and all others are cited as substituents (prefixes).

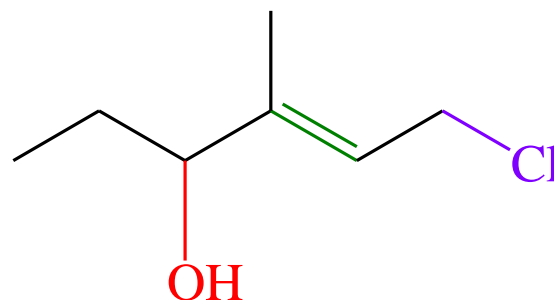


	Function Group Name	Formula	Suffix	Prefix
1-	Carboxyl	-COOH	carboxy-	-oic acid
2-	Ester	-COOR	R-oxycarbonyl-	-R-oate
3-	Aldehyde	-CHO	Formyl-	-al
4-	Ketone	$>C=O$	oxo-	-one
5-	Alcohol	-OH	hydroxy-	-ol
6-	Amine	-NH ₂	Amino-	-Amine
7-	Alkene		-----	-ene
8-	Alkyne		-----	-yne
9-	Alkane		Alkyl-	-----
10-	Ethers	R-O-	Alkoxy-	-----
11-	Halides	F, Cl, Br, I-	Halo-	-----

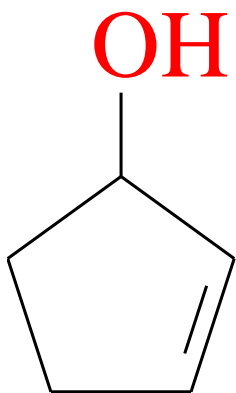
Examples



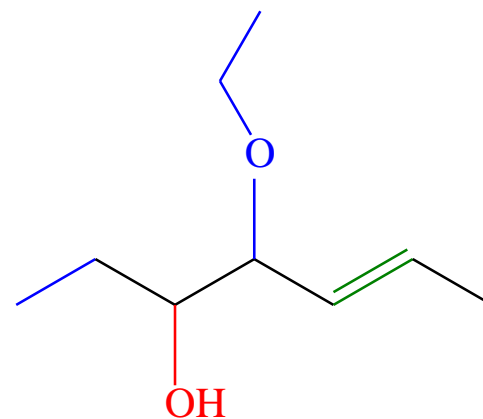
3-penten-1-yne



6-chloro-4-methyl-hexa-4-en-3-ol



2-cyclopent-1-ol



4-Ethoxyhept-5-en-3-ol

H.W(6)

Page: 187; problems 7.12

Page: 194; problem 7.18

