

Fundamentals of Organic Chemistry CHEM 109

For Students of Health Colleges

Credit hrs.: (2+1)

King Saud University

College of Science, Chemistry Department

Learning Objectives



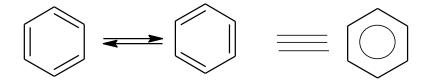
At the end of this chapter, students will able to:

- Understand the hybridization in benzene
- Understand the resonance description of structure of benzene
- Recognize the general properties of aromatic compounds, the criteria of aromaticity and Huckel's rule
- Recognize the methods for naming aromatic compounds, including IUPAC method and common names.
- Know the types of electrophilic aromatic substitution reactions.
- Understand the reactivity of aromatic compounds and orientation of monosubstituted benzene's reactions.
- Know the reactions of alkyl side chains of aromatic compounds.

Aromatic Hydrocarbons



- Originally called aromatic due to fragrant odors, although this definition seems inaccurate as many products posses distinctly non-fragrant smells!
- Currently a compound is said to be aromatic if it has benzene-like in its properties.



Their properties differ markedly from those of aliphatic hydrocarbons.

Aromatic hydrocarbons undergo electrophilic substitution whereas **aliphatic hydrocarbons** undergo electrophilic addition to double and triple bonds and free radical substitution.

The Structure of Benzene Ring



Structure of Benzene

- Molecular formula = C_6H_6 The carbon-to-hydrogen ratio in benzene, suggests a highly unsaturated structure.
- Benzene reacts mainly by substitution.

 It does not undergo the typical addition reactions of alkenes or alkynes.



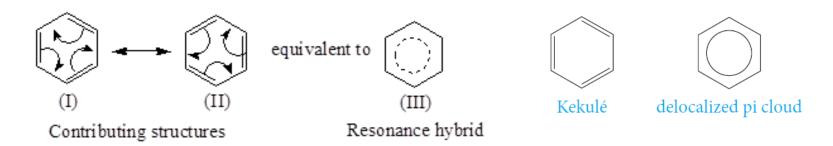
Kekulé structure for benzene.

- ➤ He suggested that six carbon atoms are located at the corners of a regular hexagon, with one hydrogen atom attached to each carbon atom.
- > He suggested that single and double bonds alternate around the ring (conjugated system of double bonds).
- > Kekulé suggested that the single and double bonds exchange positions around the ring so rapidly that the typical reactions of alkenes cannot take place.

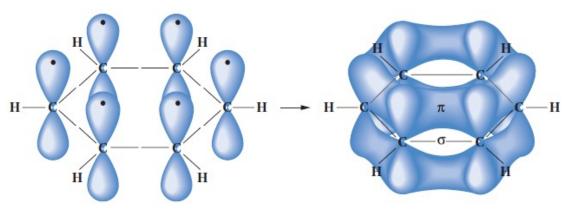
the Kekulé structures for benzene



Resonance Model for Benzene.



- Benzene is planar.
- All of the carbon-carbon bond lengths are identical: 1.39 A°, intermediate between typical single (1.54A°) and double (1.34 A°) carbon-carbon bond lengths.
- Each carbon is therefore sp2-hybridized.
- Bond angles of 120°.



Aromatic Character (Aromaticity)



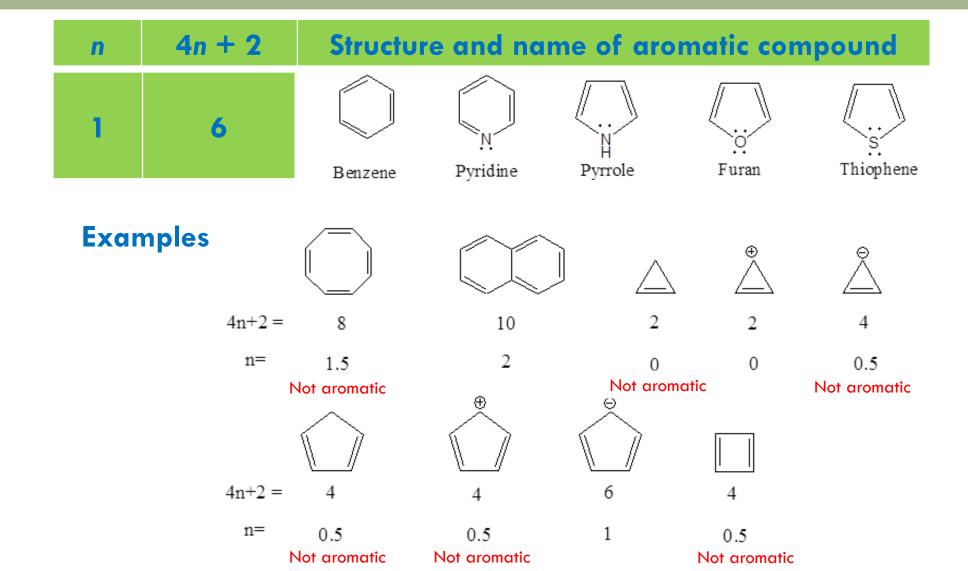
To be classified as aromatic, a compound must have:

- Oyclic structure
- 2 Cyclic structure contains what looks like a continuous system of alternating double and single bonds
- 3 Aromatic compounds must be planar
- 4 Fulfill Huckel rule

The number of \prod electrons in the compound = (4n + 2)

Where (n = 0,1, 2, 3, and so on).

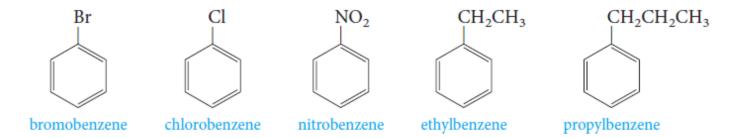




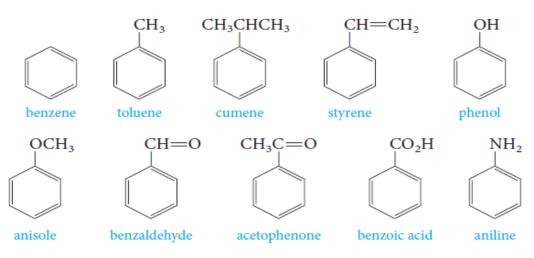
Nomenclature of Aromatic Compounds



 Monosubstituted benzenes that do not have common names accepted by IUPAC are named as derivatives of benzene.



Common names are accepted by IUPAC (parent compounds).





- When two substituents are present, three isomeric structures are possible.
 - They are designated by the prefixes; ortho- (o-), meta- (m-) and para- (p-).
 - If substituent X is attached to carbon 1; o- groups are on carbons 2 and 6, m-groups are on carbons 3 and 5, and p-groups are on carbon 4.

Examples;



■ The prefixes; ortho- (o-), meta- (m-) and para- (p-) are used when the two substituents are not identical.

Br
$$Cl$$
 CH_3 CH_2 CH_3 CH_4 CH_5 CH_5

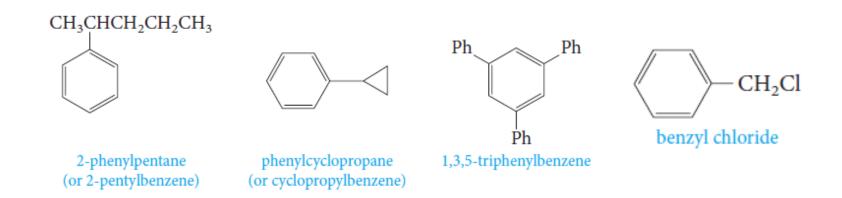
 When more than two substituents are present, their positions are designated by numbering the ring.



 Two groups with special names occur frequently in aromatic compounds; the phenyl group and the benzyl group.

$$C_6H_5-$$
 or $C_6H_5CH_2-$ or $C_6H_5CH_2-$ benzyl group

Examples;



Reactions of Benzene



Electrophilic Aromatic Substitution Reactions

1) Halogenation

$$+ X_2 \xrightarrow{FeX_3} + HX$$

$$X = Cl, Br$$

2) Nitration

$$+ HONO_2^*$$
 $\xrightarrow{H_2SO_4}$ $+ H_2O$

3) Sulfonation

$$+ HOSO_3H \xrightarrow{SO_3} + H_2O$$



4) Alkylation (Friedel-Crafts)

$$+ RCl \xrightarrow{AlCl_3} + HCl$$

$$R = alkyl group$$

5) Acylation (Friedel-Crafts)

We can generalize this two-step mechanism for all the electrophilic aromatic substitutions.

$$+ E^{+} \xrightarrow{\text{step 1}} + E \xrightarrow{\text{step 2}} E + H^{+}$$

The Mechanism of Electrophilic Aromatic Substitution



 We can generalize this two-step mechanism for all the electrophilic aromatic substitutions.

$$+ E^{+} \xrightarrow{\text{step 1}} + E^{+} \xrightarrow{\text{step 2}} E + H^{+}$$

> Halogenation

$$\begin{array}{c} \text{Cl} \\ \text{Cl}$$



Nitration

In aromatic nitration reactions, the sulfuric acid catalyst protonates the nitric acid, which then loses water to generate the nitronium ion (NO_2^+) , which contains a positively charged nitrogen atom.

$$H = \ddot{O} = \ddot{O} + \ddot{O} = \ddot{O} + \ddot{O} + \ddot{O} = \ddot{O} + \ddot{O} + \ddot{O} = \ddot{O} + \ddot{O$$

Sulfonation

We use either concentrated or fuming sulfuric acid, and the electrophile may be sulfur trioxide, SO_3 , or protonated sulfur trioxide, $^+SO_3H$.



> Friedel-Crafts Alkylation

The electrophile is a carbocation, which can be formed either by removing a halide ion from an alkyl halide with a Lewis acid catalyst (for example, AICl₃).

$$\begin{array}{c} \text{Cl} & \text{Cl} \\ | \\ \text{Cl} - \text{Al} + \text{ClCH}_2\text{CH}_3 & \Longrightarrow \text{Cl} - \text{Al}^- - \text{Cl} + \overset{+}{\text{CH}}_2\text{CH}_3 & \overset{\text{H}^+}{\longleftarrow} \text{CH}_2 = \text{CH}_2 \\ | \\ \text{Cl} & \text{cation} \end{array}$$
 (4.20)



Friedel-Crafts Acylation

The electrophile is an acyl cation generated from an acid derivative, usually an acyl halide. The reaction provides a useful general route to aromatic ketones.

$$CH_{3}CCl + AlCl_{3} \Longrightarrow CH_{3}\overset{+}{C} = O + AlCl_{4}^{-}$$

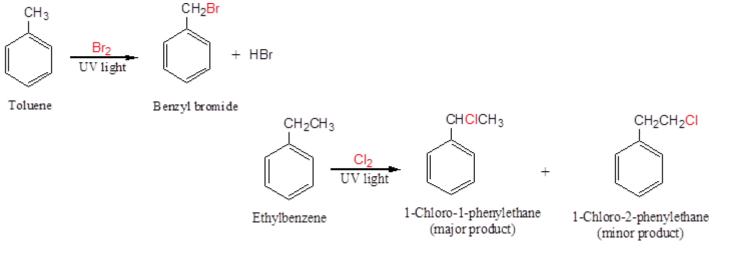
$$acetyl \ choride \qquad acetyl \ cation$$

$$+ CH_{3}\overset{+}{C} = O \Longrightarrow \overset{+}{\leftarrow} \overset{+}{\leftarrow} \overset{-}{\leftarrow} \overset{-}{\leftarrow}$$

Side-Chain Reactions of Benzene-Derivatives



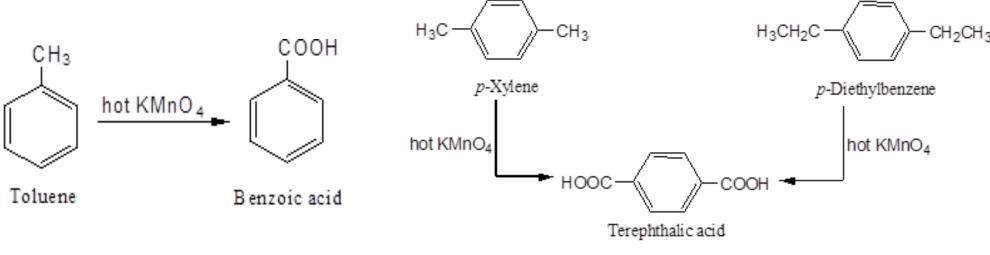
1. Halogenation of an Alkyl Side Chain





2. Oxidation of an Alkyl Side Chain

- Conversion into a carboxyl group, -COOH, by treatment with hot potassium permanganate.
- Regardless the length of the alkyl chain, the product is always the same.



(a raw material used in synthesizing Dacron and other polyesters)



- □ Side-chain oxidation of alkylbenzenes is important in certain metabolic processes.
- One way in which the body rids itself of foreign substances is by oxidation in the liver to compounds more easily excreted in the urine. Toluene, for example, is oxidized to benzoic acid by this process and is eliminated rather readily.
- Benzene, with no alkyl side chain, undergoes a different reaction in the presence of these enzymes, which convert it to a substance capable of inducing mutations in DNA.
 This difference in chemical behavior seems to be responsible for the fact that benzene is carcinogenic but toluene is not.

Disubstituted Benzenes: Orientation & Reactivity



Directing and Activating Effects of Common Functional Groups

- Substituents that release electrons to the ring will activate the ring toward electrophilic substitution.
- Substituents that withdraw electrons from the ring will deactivate the ring toward electrophilic substitution.

	Substituent group	Name of group	
ho	$-NH_2$, $-NHR$, $-NR_2$	amino	
recting	$-\overset{\cdot \cdot \cdot}{O}$ H, $-\overset{\cdot \cdot \cdot}{O}$ CH ₃ , $-\overset{\cdot \cdot \cdot}{O}$ R	hydroxy, alkoxy	Acti
Ortho, Para-Directing	O -NHCR	acylamino	Activating
Ort	$-CH_3$, $-CH_2CH_3$, $-R$	alkyl	
	-F:, -Cl:, -Br:, -I:	halo	
	:0: :0: 	acyl, carboxy	
	:0: :0: -C-NH ₂ -C-OR	carboxamido, carboalkoxy	D
Meta-Directing	:0: -S-OH :0:	sulfonic acid	Deactivating
	-C≡N:	cyano	
	\(\dot{\dot{\dot{\dot{\dot{\dot{\dot{	nitro	



- Substituents already present on an aromatic ring determine the position taken by a new substituent.
- \circ **Example**; nitration of toluene gives mainly a mixture of o- and p-nitrotoluene.

 On the other hand, nitration of nitrobenzene under similar conditions gives mainly the meta isomer.

Uses of Aromatic compound



- Aromatic compounds are important in industry.
- Aromatic compounds are extracted from complex mixtures obtained by the refining of oil or by distillation of coal tar, and are used to produce a range of important chemicals and polymers.
- Styrene is used to make polymers and plastics.
- Benzene used to make some types of rubber, dye, drugs and pesticides.
- Toluene used as a fullerene indicator, and it can be used as an octane booster in gasoline fuels used in internal combustion engines.
- Phenanthrene derivatives are used as pain medications (morphine, codeine, oxycodone).
- Naphthalene acts as a raw material in manufacture of dyes, plasticizers, and insecticides. Also used in production of some pharmaceutical products

First Midterm Exam



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Good Luck!