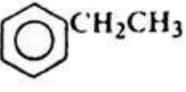


Arenes

Aliphatic-aromatic hydrocarbons >>

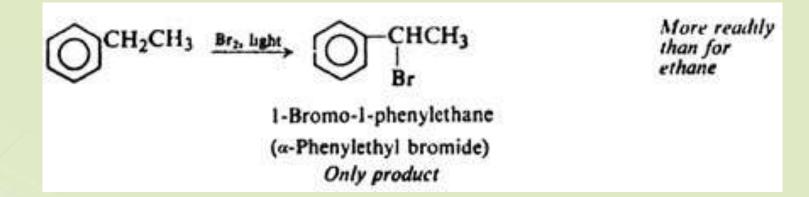
Important compounds are contain both aliphatic and aromatic units; hydrocarbons of this kind are known collectively as arenes for example:



Ethylbenzene

The ring of ethylbenzene should undergo the electrophilic substitution characteristic of benzene, and the side chain should undergo the free radical substitution characteristic of ethane

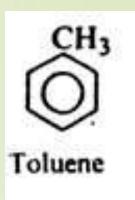
More readily H2CH3 HNO, H2SO4 CH2CH3 CH₂CH₃ than for benzene a-Nitroethylbenzene p-Nitroethylbenzene Chief products



Thus each portion of the molecule affects the reactivity of the other portion and determines the orientation of attack

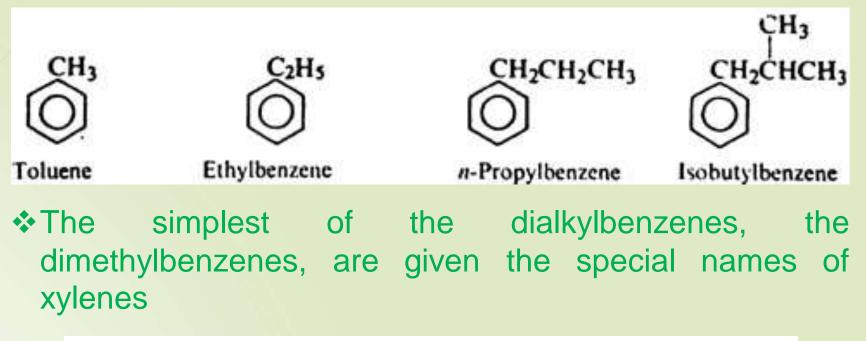
Structure and nomenclature >>

methylbenzene,alkylbenzenes,theof simplestThe <>



is

Compounds containing longer side chains are named by prefixing the name of the alkyl group to the word -______ benzene

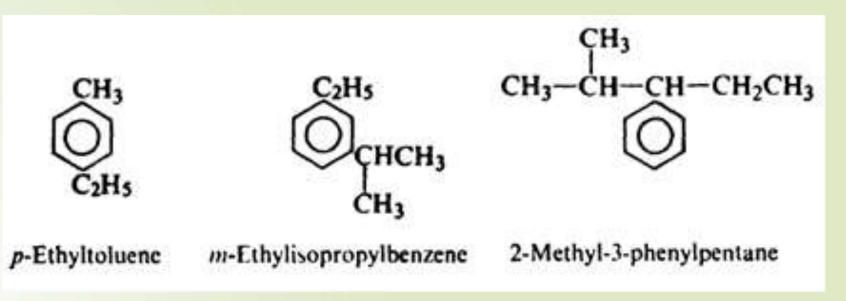




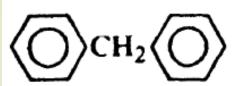
are group methylone containingDialkylbenzenes named as derivatives of toluene

Others are named by prefixing the names of both alkyl groups to the word -benzene

chain side complicated verya containing A compound right might be named as a phenylalkane (C₆H₅ = phenyl)



Compounds containing more than one benzene ring are nearly always named as derivatives of alkanes



Diphenylmethane

CH₂CH₂

1.2-Diphenylethane

simplestThe hasalkenylbenzene namespecialthe Othersstyrene. substitutedas namedgenerallyare substitutedas occasionallyalkenes, benzenes. Alkynylbenzenes are named as substituted alkynes

Preparation of alkylbenzenes ≻

1. Attachment of alkyl group: Friedel-Crafts alkylation.

$$\bigcirc + RX \xrightarrow{\text{Lewis acid}} \bigodot + HX R may rearrange$$

Lewis acid: AlCl₃, BF₃, HF, etc. Ar-X cannot be used in place of R-X

prepareto possibleis it alkanespolyhalogenated compounds containing more than one aromatic ring

$$2C_6H_6 + CH_2Cl_2 \xrightarrow{AlCl_3} C_6H_5CH_2C_6H_5 + 2HCl$$

Diphenylmethane

$$2C_{6}H_{6} + ClCH_{2}CH_{2}Cl \xrightarrow{AlCl_{3}} C_{6}H_{5}CH_{2}CH_{2}C_{6}H_{5} + 2HCl$$

1,2-Diphenylethane

$$3C_{6}H_{6} + CHCl_{3} \xrightarrow{AICl_{3}} C_{6}H_{5} - C_{6}C_{6}H_{5} + 3HCl + HCl_{4}$$

$$3C_{6}H_{6} + CCl_{4} \xrightarrow{AICl_{3}} C_{6}H_{5} - C_{6}C_{6}H_{5} + 3HCl + HCl_{5}$$

$$3C_{6}H_{6} + CCl_{4} \xrightarrow{AICl_{3}} C_{6}H_{5} - C_{6}C_{6}H_{5} + 3HCl + Cl_{6}$$

$$C_{1}$$

$$Triphenylchloromethane$$
the involvesalkylationFriedel-Craftsformechanism
following steps
$$(1) \qquad RCl + AlCl_{3} \rightleftharpoons AlCl_{4} + R \cdot y$$

$$(2) \qquad R \cdot y + C_{6}H_{6} \rightleftharpoons C_{6}H_{5}$$

$$H$$

$$(3) \qquad C_{6}H_{5} + AlCl_{4} - \rightleftharpoons C_{6}H_{5}R + HCl + AlCl_{3}$$

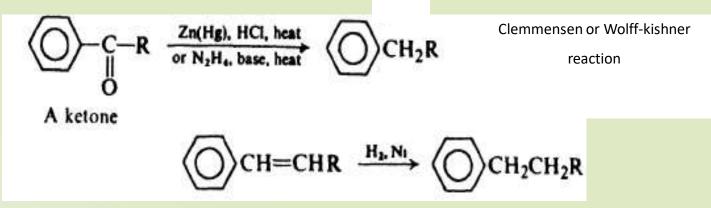
We might expect the benzene ring to be attacked by carbonium ions generated in other ways: by the action of acid on alcohols and on alkenes.

 $ROH + H^+ \rightleftharpoons ROH_2 \odot \rightleftharpoons R \odot + H_2O$ Carbonium ions from alcohols $-c = c - + H^+$, $\rightarrow -c - c - c = c$ and from alkenes H₂SO4 $C_6H_6 + (CH_3)_3COH$ $C_6H_5-C(CH_3)_3$ tert-Butyl alcohol tert-Butylbenzene H_2SO_4

 $C_{6}H_{6} + (CH_{3})_{2}C = CH_{2} \xrightarrow{H_{2}SO_{4}} C_{6}H_{5} - C(CH_{3})_{3}$ Isobutylene *tert*-Butylbenzene We might expect Friedel-Crafts alkylation accompanied by the kind of rearrangement isthat characteristic of carbonium ion reactions

$$C_{6}H_{6} + CH_{3}CH_{2}CH_{2}CI \xrightarrow{AICl_{3}}{-18^{\circ} to 80^{\circ}} C_{6}H_{5}CH_{2}CH_{2}CH_{3} \text{ and } C_{6}H_{5}CHCH_{3} \text{ Isopropylbenzene} 35-31^{\circ}/_{6} \text{ and } C_{6}H_{5}CHCH_{3} \text{ Isopropylbenzene} 65-69^{\circ}/_{6} \text{ and } C_{6}H_{5}CHCH_{2}CH_{3} \text{ and } C_{6}H_{5}CCH_{2}CH_{3} \text{ and } C_{6}H_{5}CHCH_{2}CH_{3} \text{ and } C_{6}H_{5}CHCH_{2}CH_{3} \text{ and } C_{6}H_{5}CHCH_{2}CH_{3} \text{ and } C_{6}H_{5}CCH_{2}CH_{3} \text{ and } C_{6}H_{5}CCH_{2}CH_{5} \text{ and } C_{6}H_{5}CCH_{5}CCH_{5}CH_{5}CH_{5}CH_{5}CH_{5}CH_{5}CH_{5}CH_{5$$

2. Conversion of side chain.

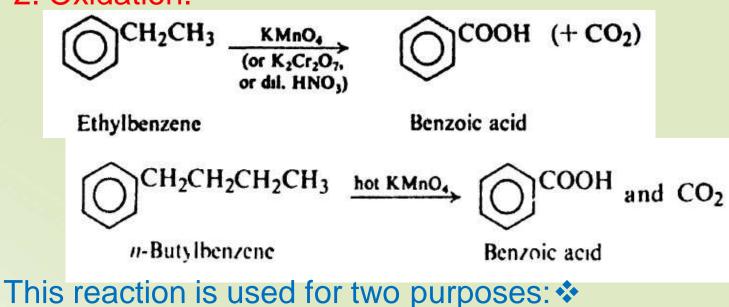


Reactions of alkylbenzenes

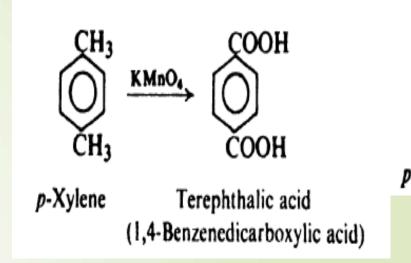
Hydrogenation1.

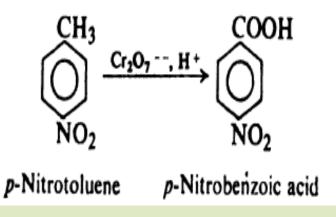
$$\underbrace{\bigcirc^{CH_2CH_3}}_{\text{Ethylbenzene}} + 3H_2 \xrightarrow{N_1, P_1, Pd} \underbrace{\bigcirc^{CH_2CH_3}}_{\text{Ethylcyclohexane}}$$

2. Oxidation.

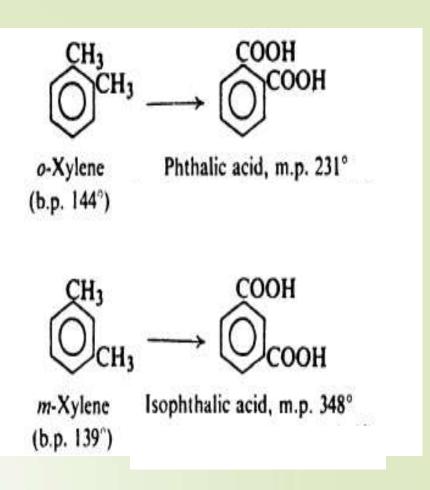


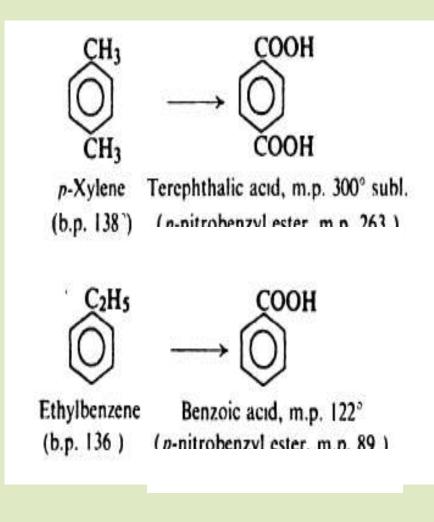
Synthesis of carboxylic acids



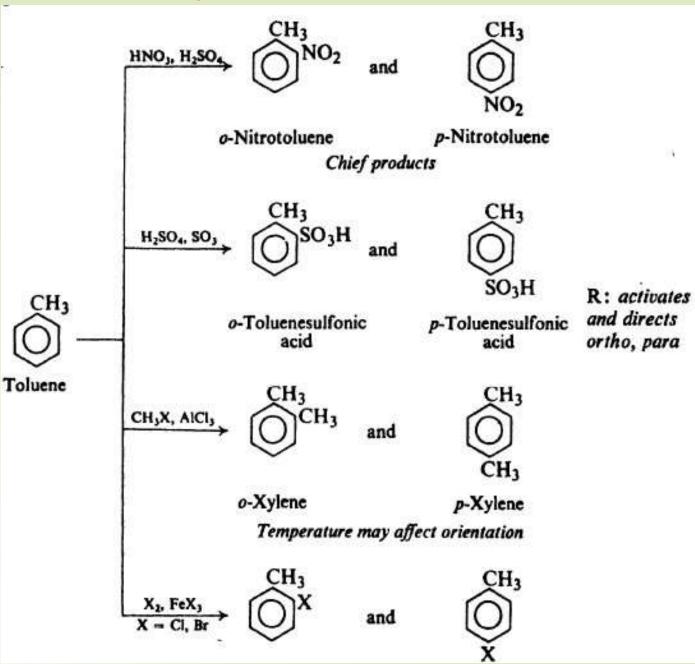


Identification of alkylbenzenes





3. Substitution in the ring. Electrophilic aromatic substitution.



4. Substitution in the side chain. Free-radical halogenation.

The ring and the side chain. We can control the position of attack simply by choosing the proper reaction conditions

