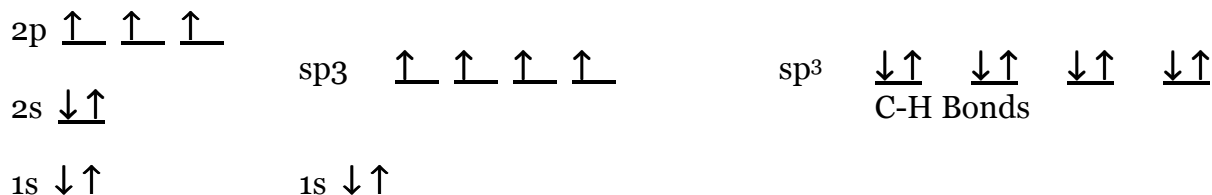


These Notes are to SUPPLEMENT the Text, They do NOT Replace reading the Text Book Material. Additional material that is in the Text Book will be on your tests! To get the most information, READ THE CHAPTER prior to the Lecture, bring in these lecture notes and make comments on these notes. These notes alone are NOT enough to pass any test!

NOTE THESE ARE DRAFT LECTURE NOTES!

CARBON: 4 electrons in the outer shell and sp^3 hybridized



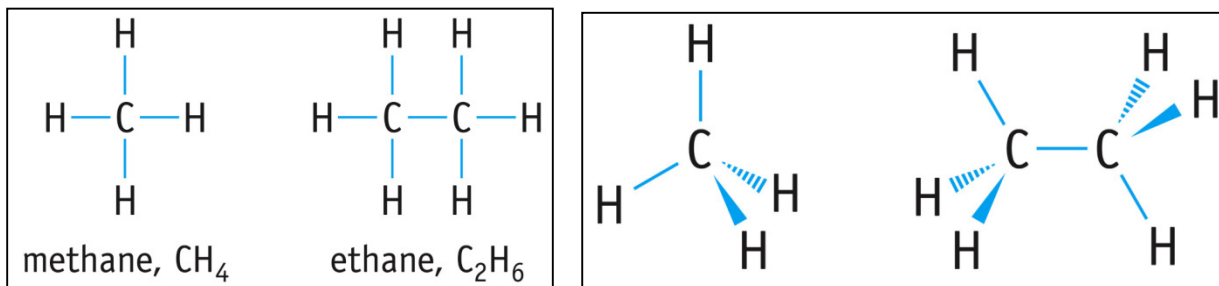
With 4 covalent bonds to many different molecules, carbon forms a large number of bonds to create more complex structures than with In-Organic compounds. Carbon can form the following types of bonds:

- | | | | | |
|---------------------------------------|---------|--------------------|--------|-----|
| 1. Single Bonds | C-C | C-O | C-N | C-X |
| 2. One Double Bond & Two Single Bonds | H-C=C-H | H ₂ C=O | HR-C=N | |
| 3. Two Double Bonds | O=C=O | | | |
| 4. Triple Bond | H-C≡C-H | | | |

Remember the geometry of these:

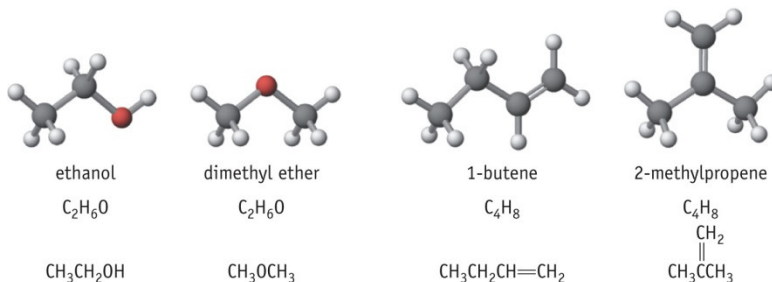
- | | | | |
|------------------------------------|---------------|-----------------|---------------|
| 1. Single Bonds | SP_3 Hybrid | Tetrahedral | 109.5° |
| 2. Double Bond | SP_2 Hybrid | Trigonal Planar | 120° |
| 3. Two Double Bonds or Triple Bond | SP Hybrid | Linear | 180° |

Picture the molecules in 3-D and not flat on a sheet of paper:



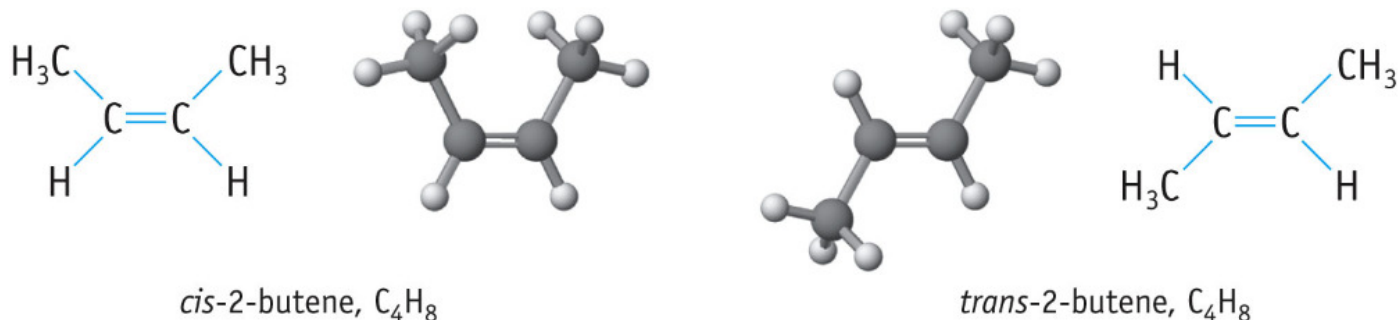
Isomers

Structural: Compounds having the same composition but different molecular geometry:



Stereoisomers: compounds with the same formulae and same attachments but have different orientations in space.

Geometric Isomers: Different orientations in space (e.g. cis / trans):



Optical Isomers: Molecules with structures that are mirror images of each other

Chiral: Non-superimposable mirror images

Enantiomers: Pairs of non-superimposable, mirror image molecules.

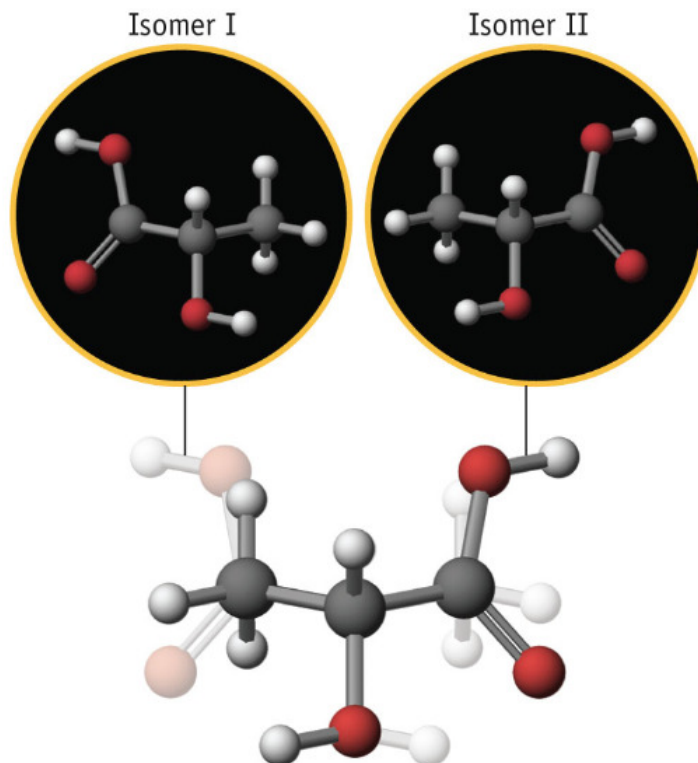
These compounds have the same physical properties: MP, BP, Density, Solubility, IR, NMR

But: in solution, pure enantiomers will rotate plane-polarized light – **Optical Isomerism**

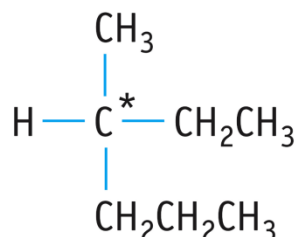
Your Left Hand is an Enantiomer for your Right Hand

They are mirror images and non-superimposable on each other.

Carbon with 4 different functional groups attached is usually Optically Active – forms L & R Isomers



(a) Lactic acid enantiomers are nonsuperimposable.



Stability of Carbon Compounds:

Organic compounds do not decompose thermally under normal conditions

10.2 Hydrocarbons: Compounds made of only Carbon and Hydrogen

Alkanes: only single bonds

CycleAlkanes: Single bonds, cyclic structure

Alkenes: at least one double bond

Alkynes: at least one triple bond

Aromatic: 6 member unsaturated ring

Alkanes: General formulae $\text{C}_n\text{H}_{2n-2}$

Alkanes have Carbon - Carbon single bonds and they are called **Saturated Hydrocarbons**

Remember, there is free rotation about Carbon - Carbon single bonds

<u>n</u>	<u>Structure</u>	<u>Name</u>	
1	CH_4	Methane	
2	$\text{H}_3\text{C}-\text{CH}_3$	Ethane	
3	$\text{H}_3\text{C}-\text{CH}_2-\text{CH}_3$	Propane	
4	$\text{H}_3\text{C}-\text{CH}_2-\text{CH}_2-\text{CH}_3$	n-Butane	n = normal
	$\begin{array}{c} \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{H}_3\text{C}- \end{array}$	iso-Butane	

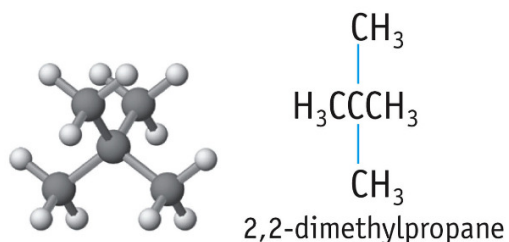
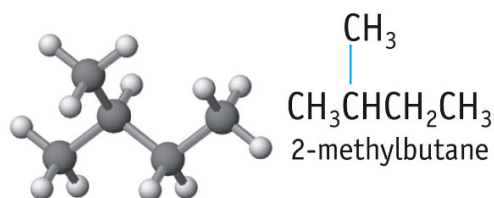
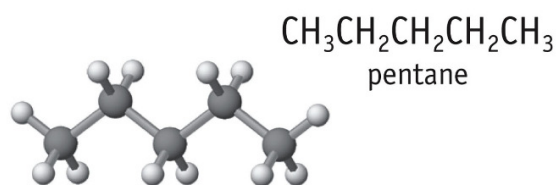
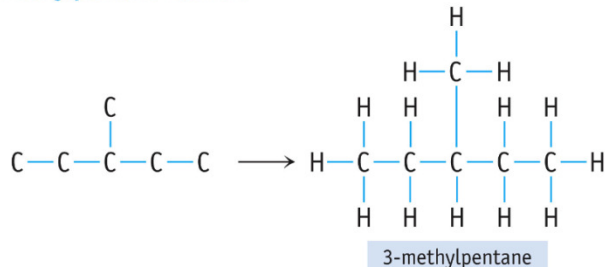
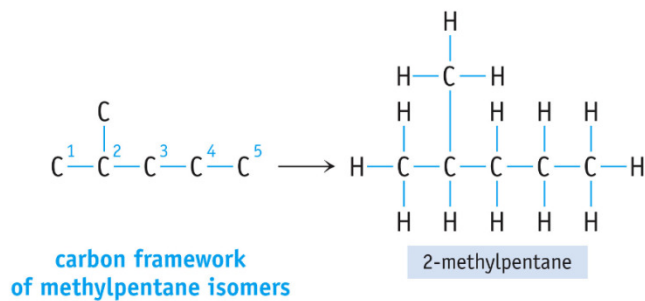


Table 10.2 Selected Hydrocarbons of the Alkane Family, $\text{C}_n\text{H}_{2n+2}$ *

Name	Molecular Formula	State at Room Temperature
Methane	CH_4	
Ethane	C_2H_6	
Propane	C_3H_8	Gas
Butane	C_4H_{10}	
Pentane	C_5H_{12} (pent- = 5)	
Hexane	C_6H_{14} (hex- = 6)	
Heptane	C_7H_{16} (hept- = 7)	
Octane	C_8H_{18} (oct- = 8)	Liquid
Nonane	C_9H_{20} (non- = 9)	
Decane	$\text{C}_{10}\text{H}_{22}$ (dec- = 10)	
Octadecane	$\text{C}_{18}\text{H}_{38}$ (octadec- = 18)	
Eicosane	$\text{C}_{20}\text{H}_{42}$ (eicos- = 20)	Solid

Structural Isomers: Isomers that have the same molecular formulae, but different orientations in space. See above: Pentane, 2-methylbutane, 2,2-dimethylpropane



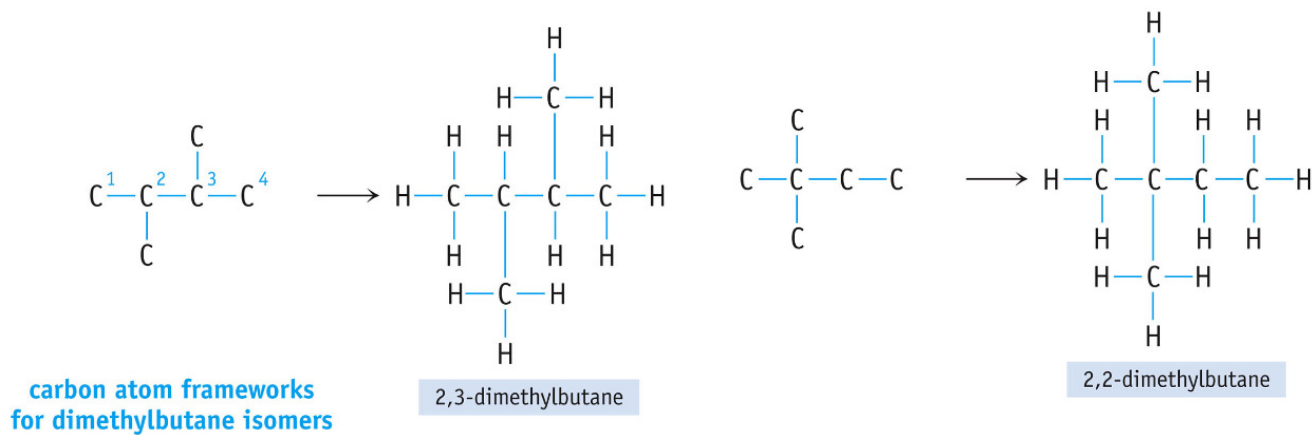
Naming:

1. Draw the structural isomer of the alkane with the longest possible C atom chain

This is the Root Structure

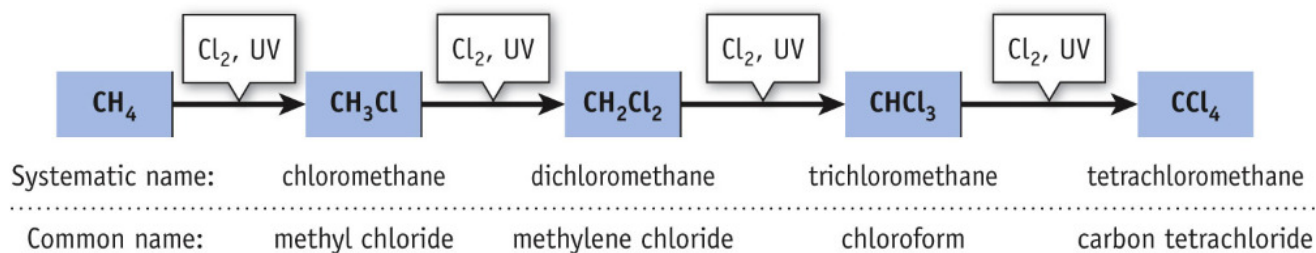
2. Substituent / Additional groups on the longest hydrocarbon chain are identified by a name and the position on the carbon chain. Numbering begins on the chain end that allows the 1st additional group to have the lowest possible number. Names of these Alkane Groups are called Alkyl Groups

3. If more than one group is attached to a particular long chain carbon atom, then use prefixes: di, tri and tetra. (Why is there no Mono or Hepta?).



Properties of Alkanes

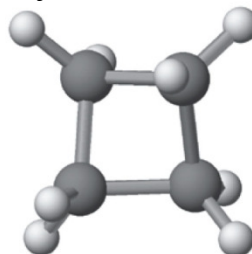
- Methane, Ethane, Propane and Butane are gases at RT and pressure
- Higher Mw compounds are liquids or solids
- Increase MP or BP with increase Mw
- Alkanes are colorless and insoluble in water - they are non-polar
- Alkanes burn in air to give CO₂ and H₂O
- Alkanes are relatively inert – they do not undergo chemical reactivity
- Alkanes do react with Chlorine:



Cycloalkanes: C_nH_{2n} Formed from tetrahedral carbon joined to form a ring

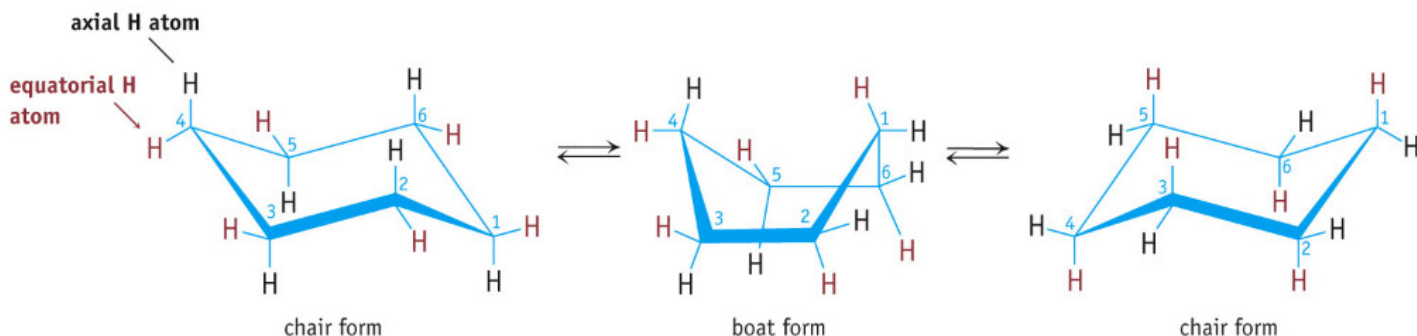


cyclopropane, C_3H_6



cyclobutane, C_4H_8

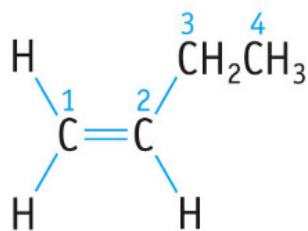
Cyclohexane, C_6H_{12} forms a 6 member ring that exists in a chair and boat 3D configuration. If the carbon atoms were flat (as in the plane of this piece of paper) they would be strained having 120° bond angles. But in the chair/boat configuration they have the normal 109.5° bond angle.



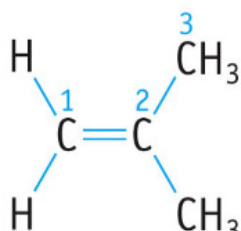
Alkenes and Alkynes

C_nH_{2n} Alkenes are hydrocarbons with one or more $\text{C}=\text{C}$ (carbon-carbon double) bonds
 $\text{C}_n\text{H}_{2n-2}$ Alkynes are hydrocarbons with one or more $\text{C}\equiv\text{C}$ (carbon-carbon triple) bonds.

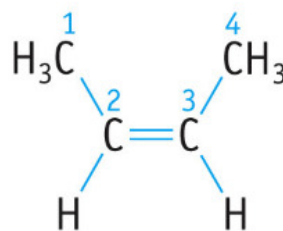
Alkenes have the base Alkane name, but end in -ene. The root name is determined by the longest carbon chain that contains the double bond. It may include the prefix: cis or trans



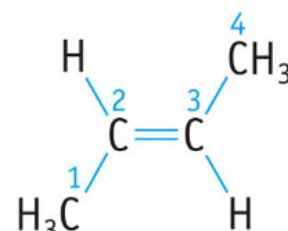
1-butene



2-methylpropene



cis-2-butene



trans-2-butene

Alkynes have the carbon-carbon triple bond. $\text{H}-\text{C}\equiv\text{C}-\text{H}$ is acetylene, a welding gas

Properties of Alkenes and Alkynes

They are colorless

Low Mw compounds are gases

Are oxidized, react with air, to form CO₂ and H₂O and very exothermic

They are called **Unsaturated Compounds**

Alkenes are Trigonal Planer with bond angles of 120°

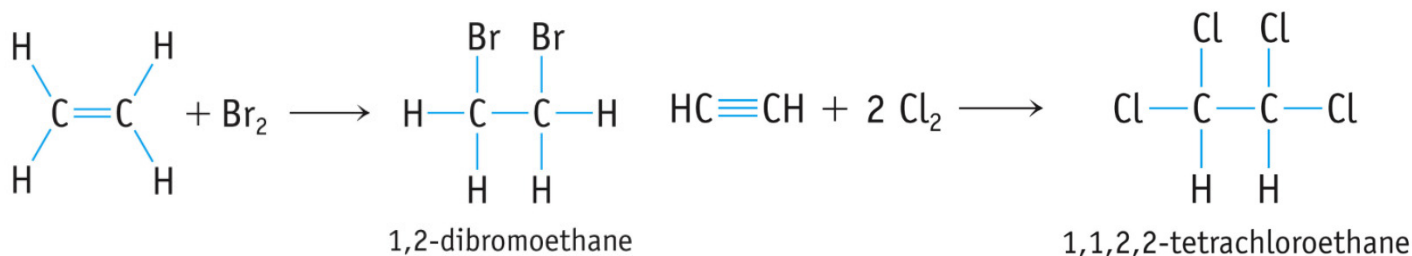
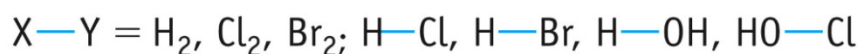
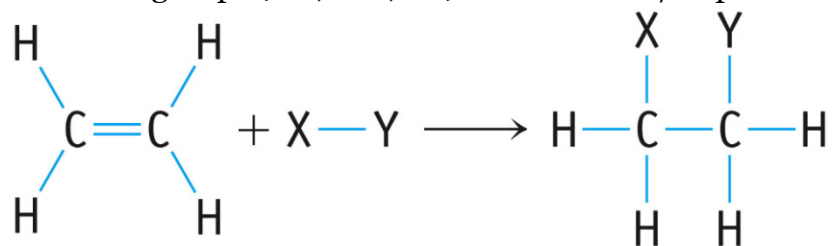
Alkynes are Linear with bond angles of 180°

Table 10.4 Some Simple Alkynes C_nH_{2n-2}

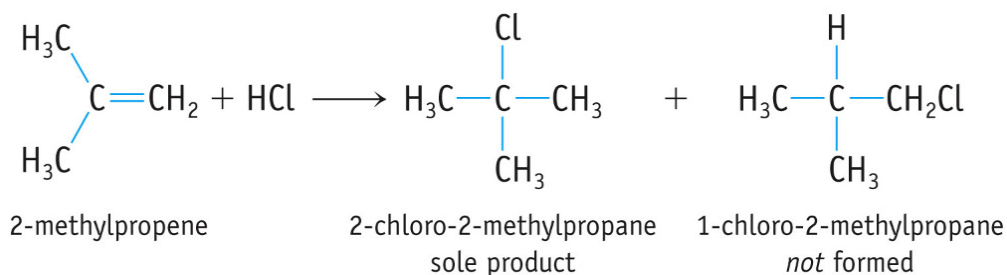
Structure	Systematic Name	Common Name	BP (°C)
HC≡CH	ethyne	acetylene	-85
CH ₃ C≡CH	propyne	methylacetylene	-23
CH ₃ CH ₂ C≡CH	1-butyne	ethylacetylene	9
CH ₃ C≡CCH ₃	2-butyne	dimethylacetylene	27

Reactions:

Addition Reactions: Add groups (Cl₂, HCl, etc) to the double / triple bond

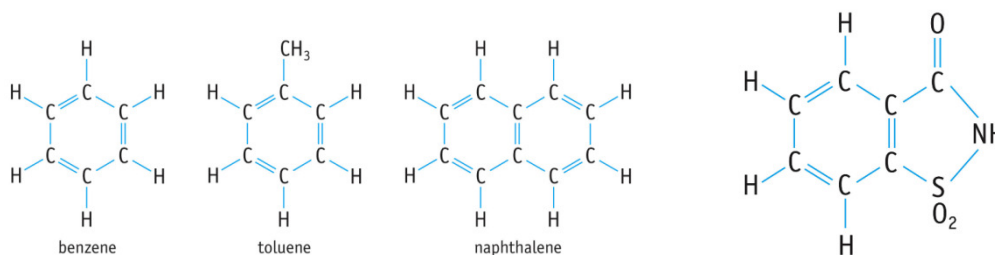


Markovnikov's Rule: when HX is added to an unsymmetrical alkene, the hydrogen is attached to the carbon that already has the largest number of hydrogens:



Hydrogenation: The H-X reagent is H₂. $\text{CH}_2=\text{CH}_2 + \text{H}_2 \rightarrow \text{CH}_3-\text{CH}_3$

Aromatic Compounds: C₆H₆ is a class of compounds with a 6 membered unsaturated ring

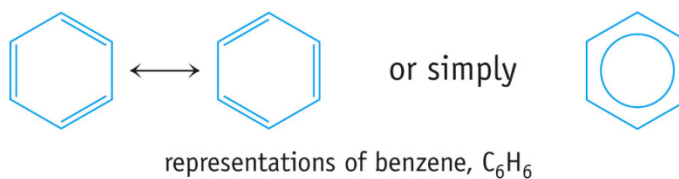


Properties:

- Usually obtained from coal
- Starting point for making many different compounds and drugs
- Do not undergo the normal alkene addition reactions
- Benzene is a colorless liquid
- They are insoluble in water
- Oxidized with oxygen to produce O₂, H₂O and energy (exothermic)
- Very stable due to its Resonance Stabilization
- Undergoes Substitution reactions in place of addition

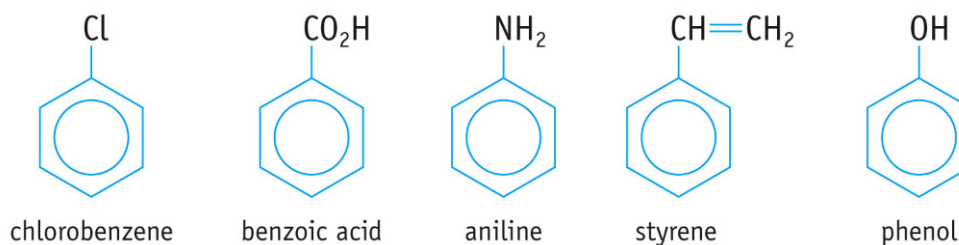
Benzene: common industrial solvent and reaction starting reactant
Toluene: from tolu balsam, pleasant smelling gum (don't smell it!)
Naphthalene: moth balls

The Benzene bonds are **Resonance Formulae** :

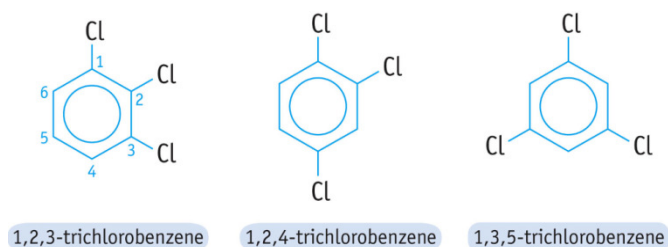


Benzene Derivatives:

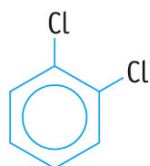
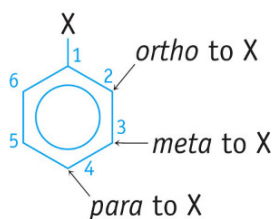
Mono Substituted:



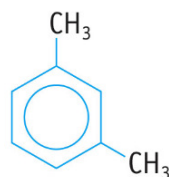
Tri Substituted:



Di-Substituted:



Systematic name: 1,2-dichlorobenzene
Common name: *o*-dichlorobenzene

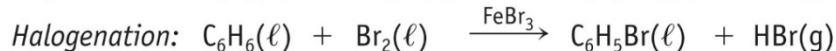
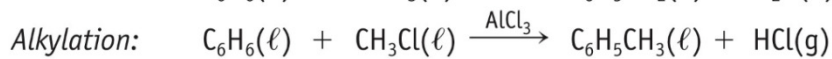
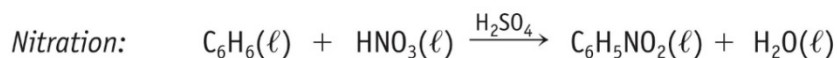


Systematic name: 1,3-dimethylbenzene
Common name: *m*-xylene



Systematic name: 1,4-dinitrobenzene
Common name: *p*-dinitrobenzene

Substitution Reactions: require a strong Bronsted Acid



10.3 Alcohols, Ethers and Amines

Functional Group (X): an atom or group of atoms attached to a carbon atom in the hydrocarbon. R-X
-X = Alcohol -OH, Amine -NH₂, Ether C-O-C, Halogen, Acid COOH.

Table 10.6 Common Functional Groups and Derivatives of Alkanes

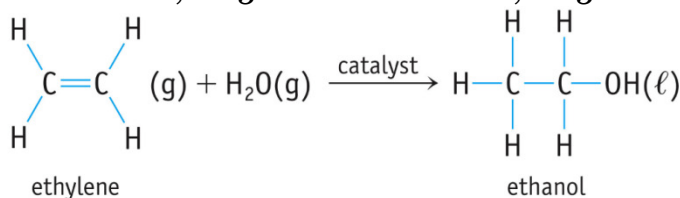
Functional Group*	General Formula*	Class of Compound	Examples
F, Cl, Br, I	RF, RCl, RBr, RI	Haloalkane	CH ₃ CH ₂ Cl, chloroethane
OH	ROH	Alcohol	CH ₃ CH ₂ OH, ethanol
OR'	ROR'	Ether	(CH ₃ CH ₂) ₂ O, diethyl ether
NH ₂ †	RNH ₂	(Primary) Amine	CH ₃ CH ₂ NH ₂ , ethylamine
$\begin{array}{c} \text{O} \\ \parallel \\ -\text{CH} \end{array}$	RCHO	Aldehyde	CH ₃ CHO, ethanal (acetaldehyde)
$\begin{array}{c} \text{O} \\ \parallel \\ -\text{C}-\text{R}' \end{array}$	RCOR'	Ketone	CH ₃ COCH ₃ , propanone (acetone)
$\begin{array}{c} \text{O} \\ \parallel \\ -\text{C}-\text{OH} \end{array}$	RCO ₂ H	Carboxylic acid	CH ₃ CO ₂ H, ethanoic acid (acetic acid)
$\begin{array}{c} \text{O} \\ \parallel \\ -\text{C}-\text{OR}' \end{array}$	RCO ₂ R'	Ester	CH ₃ CO ₂ CH ₃ , methyl acetate
$\begin{array}{c} \text{O} \\ \parallel \\ -\text{C}-\text{NH}_2 \end{array}$	RCONH ₂	Amide	CH ₃ CONH ₂ , acetamide

* R and R' can be the same or different hydrocarbon groups.

† Secondary amines (R₂NH) and tertiary amines (R₃N) are also possible, see discussion in the text.

Alcohols and Ethers

Alcohol: R-OH CH₃-OH Methanol, CH₃-CH₂-OH Ethanol, CH₃-CH₂CH₂-OH n-Propanol



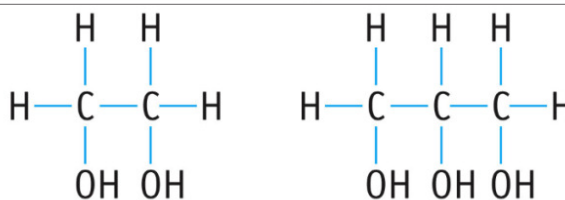
Usually prepared by Fermentation.

C3 and above alcohols can have structural isomers:

Ethanol is used in "Beverages" IsoPropyl Alcohol is rubbing alcohol

Table 10.7 Some Important Alcohols

Condensed Formula	BP (°C)	Systematic Name	Common Name	Use
CH ₃ OH	65.0	Methanol	Methyl alcohol	Fuel, gasoline additive, making formaldehyde
CH ₃ CH ₂ OH	78.5	Ethanol	Ethyl alcohol	Beverages, gasoline additive, solvent
CH ₃ CH ₂ CH ₂ OH	97.4	1-Propanol	Propyl alcohol	Industrial solvent
CH ₃ CH(OH)CH ₃	82.4	2-Propanol	Isopropyl alcohol	Rubbing alcohol
HOCH ₂ CH ₂ OH	198	1,2-Ethandiol	Ethylene glycol	Antifreeze
HOCH ₂ CH(OH)CH ₂ OH	290	1,2,3-Propanetriol	Glycerol (glycerin)	Moisturizer in consumer products



Systematic name: 1,2-ethanediol
Common name: ethylene glycol

Systematic name: 1,2,3-propanetriol
Common name: glycerol or glycerin

Ethylene Glycol
Glycerol

Car antifreeze
Backbone for soap and living organic matter

Ether: R-O-R' H₃C-CH₂-O-CH₂-CH₃ Diethyl Ether
Lower BP than the corresponding alcohol – Why??
Lower Mw is slightly soluble in water

Properties of Alcohols

Methane CH₄ a gas, Methanol CH₃OH a liquid soluble in water

Alcohols are polar, like water, hydrogen bond and are soluble in polar solvents

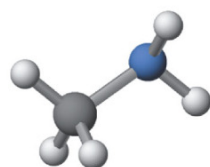
As R in R-OH increases, the BP increases and solubility in water decreases (Why ??)

Amines

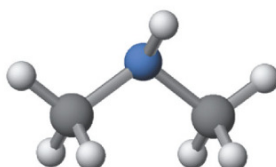
R-NH₂ = **Primary**

RR'-NH = **Secondary**

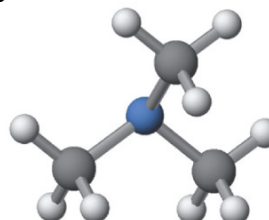
RR'R''-N **Tertiary Amine**



CH₃NH₂
primary amine
methylamine



(CH₃)₂NH
secondary amine
dimethylamine

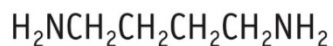


(CH₃)₃N
tertiary amine
trimethylamine

Remember: Nitrogen is Trigonal Pyramidal with a lone pair of electrons sticking out!

Properties of Amines:

Offensive Odors:



putrescine
1,4-butanediamine

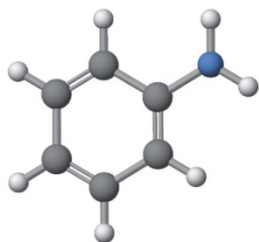


cadaverine
1,5-pentanediamine

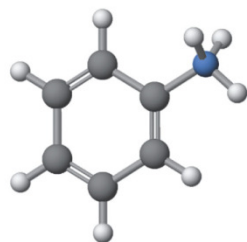
Low Mw amines are soluble in water

All amines are bases and react with acids to give a salt (H^+ reacts with N^- : Nitrogen lone pair)

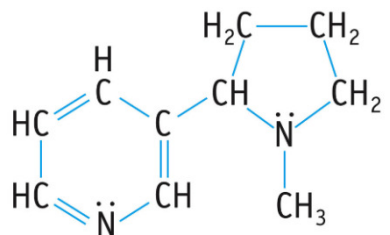
Aromatic Amines:



aniline

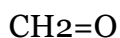


anilinium ion



nicotine

Aldehydes:



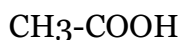
Formaldehyde

Ketones:



Acetone

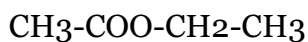
Carboxylic Acids



Acetic Acid

Have a sour taste

Ester

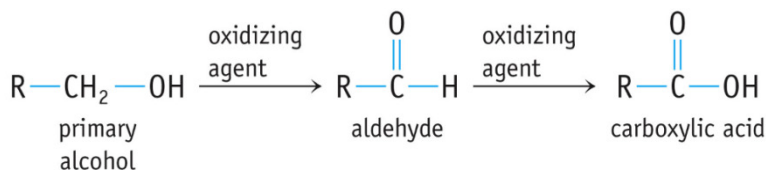


Ethyl Acetate (Nail Polish Remover)

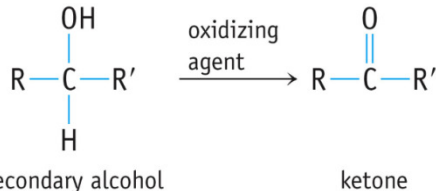
Amide



Oxidation of a primary alcohol:

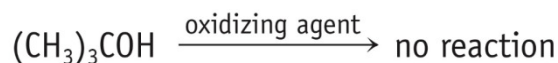


Oxidation of a secondary alcohol:



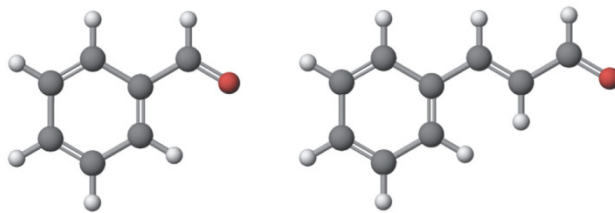
(—R and $\text{—R}'$ are organic groups. They may be the same or different.)

Oxidation of a tertiary alcohol:



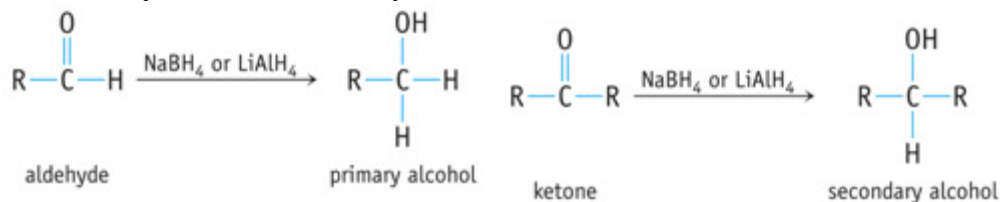
Aldehydes and Ketones

Have a pleasant odor



benzaldehyde, C_6H_5CHO *trans*-cinnamaldehyde, $C_6H_5CH=CHCHO$

Aldehydes and ketones are the oxidation products of primary and secondary alcohols
Reduction of Aldehydes and ketones yield the alcohol



Carboxylic Acids Produced by the oxidation of an alcohol or aldehyde (Why not ketone ?)

Sour taste: Acetic Acid, Tartaric Acid, Citric Acid, Formic Acid, Butyric Acid

Named by adding "oic" to the alkane name

Low Mw acids dissolve in water

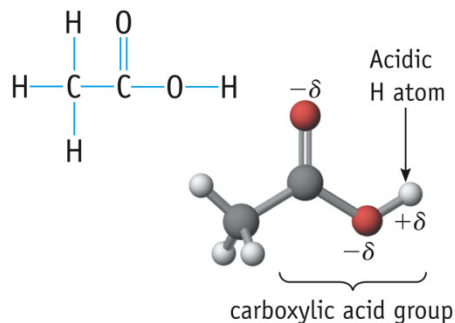


Table 10.9 Some Naturally Occurring Carboxylic Acids

Name	Structure	Natural Source
Benzoic acid		Berries
Citric acid	$\text{HO}_2\text{C}-\text{CH}_2-\text{C}(\text{OH})(\text{CO}_2\text{H})-\text{CH}_2-\text{CO}_2\text{H}$	Citrus fruits
Lactic acid	$\text{H}_3\text{C}-\text{CH}(\text{OH})-\text{CO}_2\text{H}$	Sour milk
Malic acid	$\text{HO}_2\text{C}-\text{CH}_2-\text{CH}(\text{OH})-\text{CO}_2\text{H}$	Apples
Oleic acid	$\text{CH}_3(\text{CH}_2)_7-\text{CH}=\text{CH}-(\text{CH}_2)_7-\text{CO}_2\text{H}$	Vegetable oils
Oxalic acid	$\text{HO}_2\text{C}-\text{CO}_2\text{H}$	Rhubarb, spinach, cabbage, tomatoes
Stearic acid	$\text{CH}_3(\text{CH}_2)_{16}-\text{CO}_2\text{H}$	Animal fats
Tartaric acid	$\text{HO}_2\text{C}-\text{CH}(\text{OH})-\text{CH}(\text{OH})-\text{CO}_2\text{H}$	Grape juice, wine

Table 10.10 Some Simple Carboxylic Acids

Structure	Common Name	Systematic Name	BP (°C)
$\begin{array}{c} \text{O} \\ \parallel \\ \text{HCOH} \end{array}$	Formic acid	Methanoic acid	101
$\begin{array}{c} \text{O} \\ \parallel \\ \text{CH}_3\text{COH} \end{array}$	Acetic acid	Ethanoic acid	118
$\begin{array}{c} \text{O} \\ \parallel \\ \text{CH}_3\text{CH}_2\text{COH} \end{array}$	Propionic acid	Propanoic acid	141
$\begin{array}{c} \text{O} \\ \parallel \\ \text{CH}_3(\text{CH}_2)_2\text{COH} \end{array}$	Butyric acid	Butanoic acid	163
$\begin{array}{c} \text{O} \\ \parallel \\ \text{CH}_3(\text{CH}_2)_3\text{COH} \end{array}$	Valeric acid	Pentanoic acid	187

Esters: RCOOR'

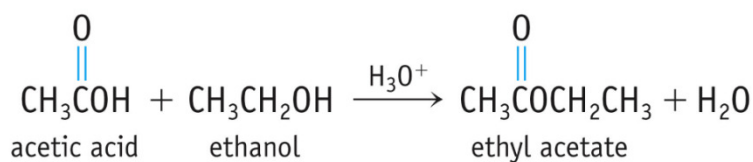
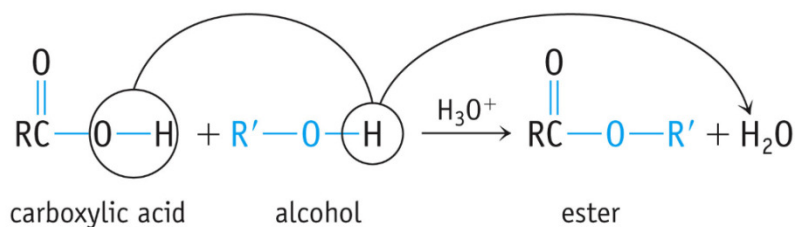

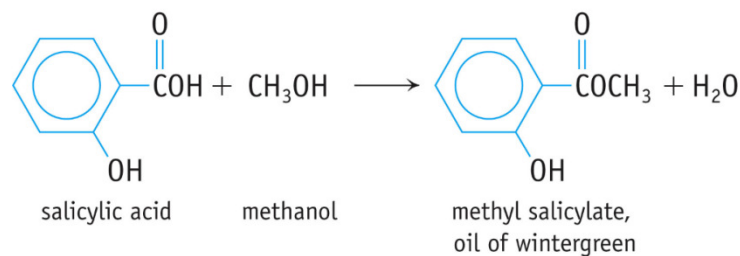
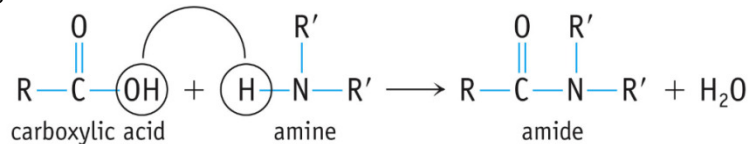


Table 10.11 Some Acids, Alcohols, and Their Esters

Acid	Alcohol	Ester	Odor of Ester
$\text{CH}_3\text{CO}_2\text{H}$ acetic acid	$\begin{array}{c} \text{CH}_3 \\ \\ \text{CH}_3\text{CHCH}_2\text{CH}_2\text{OH} \end{array}$ 3-methyl-1-butanol	$\begin{array}{c} \text{O} \quad \text{CH}_3 \\ \parallel \quad \\ \text{CH}_3\text{COCH}_2\text{CH}_2\text{CHCH}_3 \end{array}$ 3-methylbutyl acetate	Banana
$\text{CH}_3\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$ butanoic acid	$\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{OH}$ 1-butanol	$\begin{array}{c} \text{O} \\ \parallel \\ \text{CH}_3\text{CH}_2\text{CH}_2\text{COCH}_2\text{CH}_2\text{CH}_2\text{CH}_3 \end{array}$ butyl butanoate	Pineapple
$\text{CH}_3\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$ butanoic acid	 benzyl alcohol	$\begin{array}{c} \text{O} \\ \parallel \\ \text{CH}_3\text{CH}_2\text{CH}_2\text{COCH}_2-\text{C}_6\text{H}_5 \end{array}$ benzyl butanoate	Rose



Amides: From a organic acid and an amine



10.5 Polymers From many parts, or a giant molecule made from small molecules or monomers

Thermoplastics Polyethylene soften and flow when heated and harden when cooled

Thermosetting Formica initially soft, but set hard when heated and cannot be resoftened

Addition Polymers made directly by adding monomer units

Condensation Polymers made by combing monomer units and splitting out a small molecule such as water

We will not go into

Addition Polymers

Polyethylene and other Polyolefins

Natural and Synthetic Rubber

Condensation Polymers

Polyesters

Polyamids