

Chemistry, The Central Science, 10th edition
Theodore L. Brown; H. Eugene LeMay, Jr.;
and Bruce E. Bursten

Chapter 25

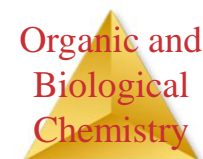
Organic and Biological Chemistry

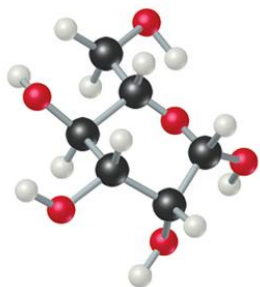
John D. Bookstaver

St. Charles Community College

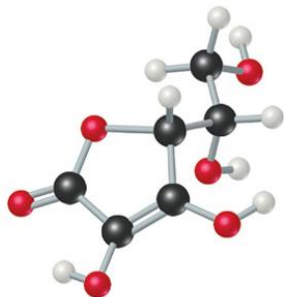
St. Peters, MO

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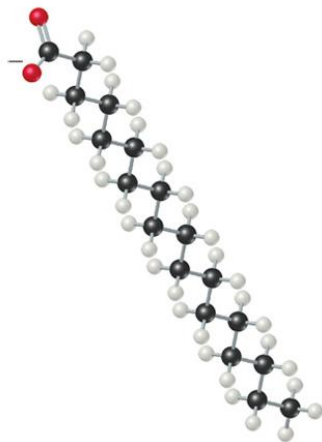




Glucose ($C_6H_{12}O_6$)



Ascorbic acid ($C_6H_7O_6$)



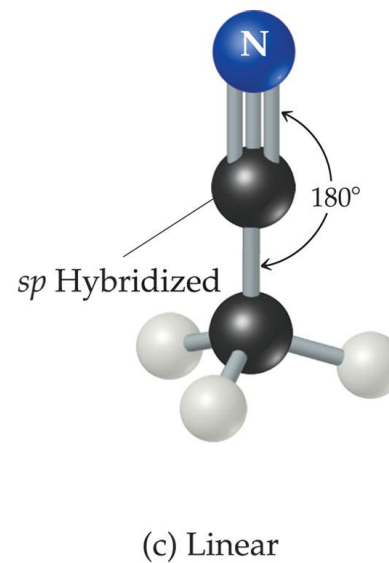
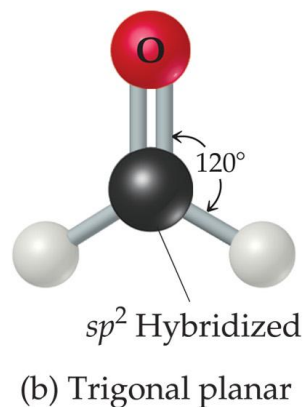
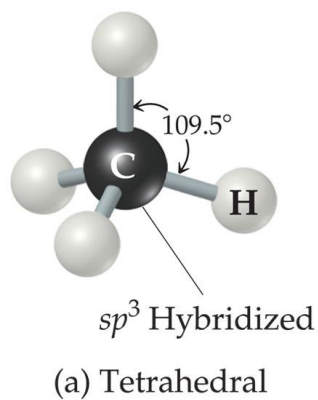
Surfactant ($C_{17}H_{35}COO^-$)

Organic Chemistry

- The chemistry of carbon compounds.
- Carbon has the ability to form long chains.
- Without this property, large biomolecules such as proteins, lipids, carbohydrates, and nucleic acids could not form.

Structure of Carbon Compounds

- There are three hybridization states and geometries found in organic compounds:
 - sp^3 Tetrahedral
 - sp^2 Trigonal planar
 - sp Linear



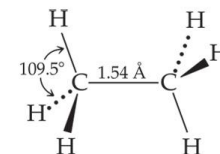
Hydrocarbons

- Four basic types:
 - Alkanes
 - Alkenes
 - Alkynes
 - Aromatic hydrocarbons

ALKANE
Ethane CH_3CH_3



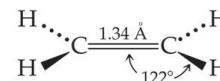
(a)



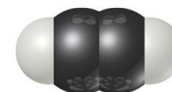
ALKENE
Ethylene $\text{CH}_2=\text{CH}_2$



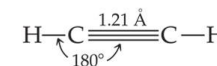
(b)



ALKYNE
Acetylene $\text{CH}\equiv\text{CH}$



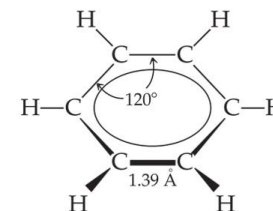
(c)



AROMATIC
Benzene C_6H_6

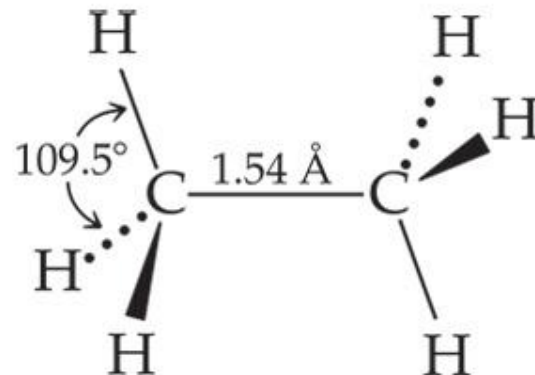


(d)



Alkanes

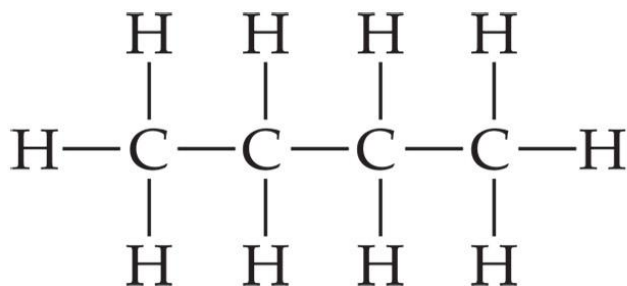
ALKANE
Ethane



- Only single bonds.
- Saturated hydrocarbons.
 - “Saturated” with hydrogens.

Formulas

- Lewis structures of alkanes look like this.
- Also called structural formulas.
- Often not convenient, though...

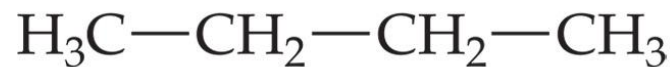
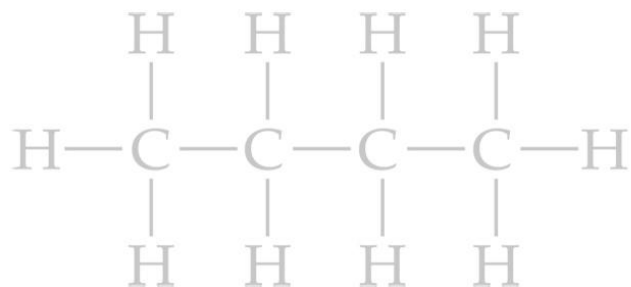


or



Formulas

...so more often condensed formulas are used.



or

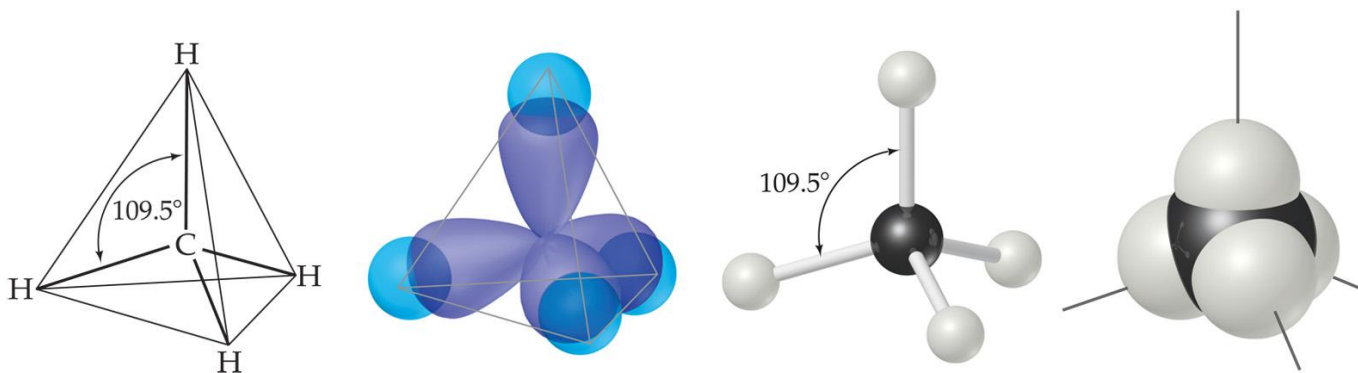


Properties of Alkanes

Molecular Formula	Condensed Structural Formula	Name	Boiling Point (°C)
CH ₄	CH ₄	Methane	-161
C ₂ H ₆	CH ₃ CH ₃	Ethane	-89
C ₃ H ₈	CH ₃ CH ₂ CH ₃	Propane	-44
C ₄ H ₁₀	CH ₃ CH ₂ CH ₂ CH ₃	Butane	-0.5
C ₅ H ₁₂	CH ₃ CH ₂ CH ₂ CH ₂ CH ₃	Pentane	36
C ₆ H ₁₄	CH ₃ CH ₂ CH ₂ CH ₂ CH ₂ CH ₃	Hexane	68
C ₇ H ₁₆	CH ₃ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₃	Heptane	98
C ₈ H ₁₈	CH ₃ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₃	Octane	125
C ₉ H ₂₀	CH ₃ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₃	Nonane	151
C ₁₀ H ₂₂	CH ₃ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₃	Decane	174

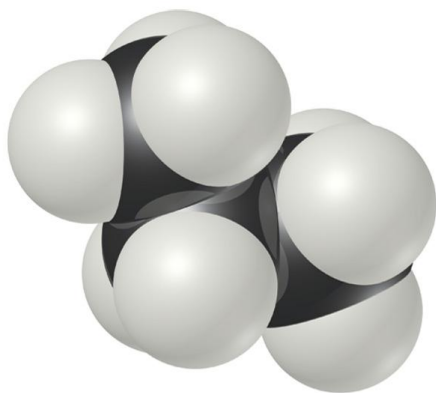
- Only van der Waals force: London force.
- Boiling point increases with length of chain.

Structure of Alkanes

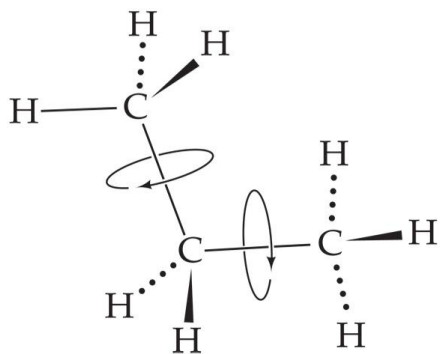


- Carbons in alkanes sp^3 hybrids.
- Tetrahedral geometry.
- 109.5° bond angles.

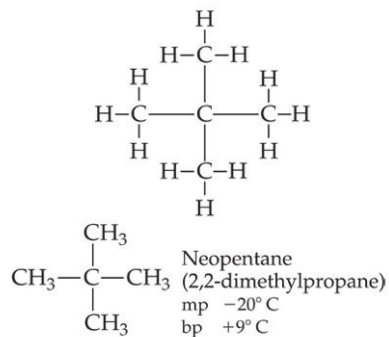
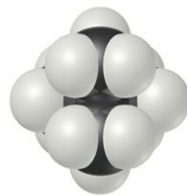
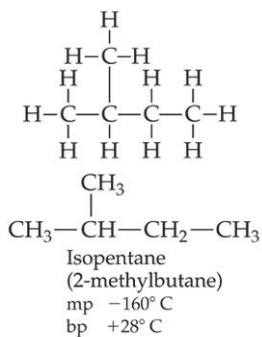
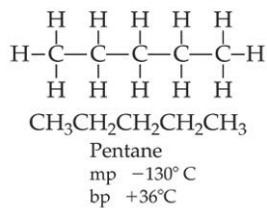
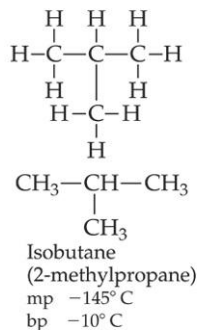
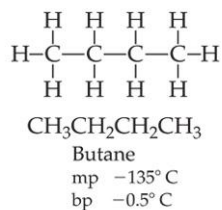
Structure of Alkanes



- Only σ -bonds in alkanes
- Free rotation about C—C bonds.



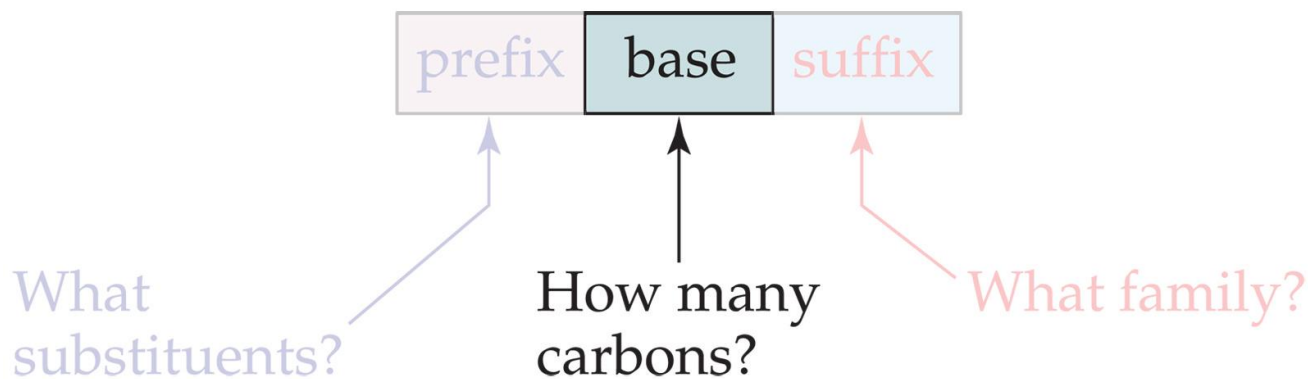
Isomers



Have same molecular formulas, but atoms are bonded in different order.

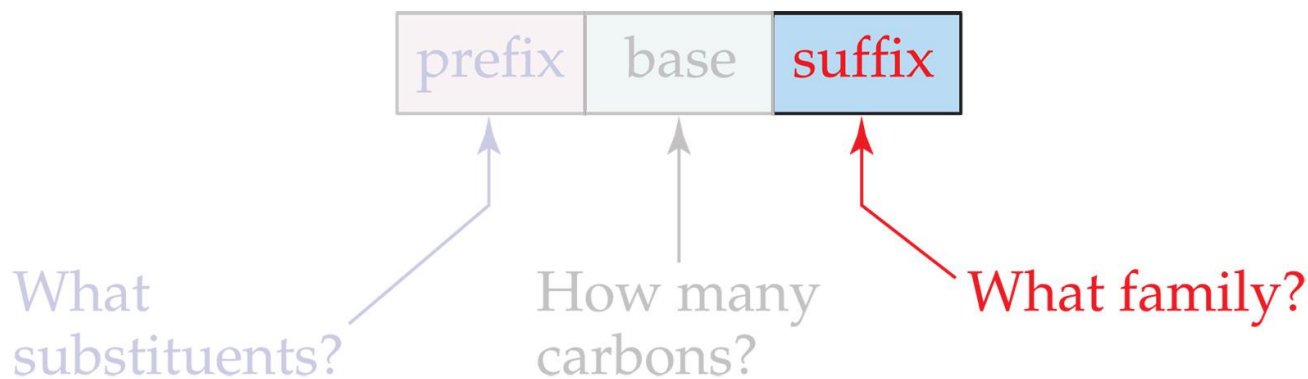
Organic Nomenclature

- Three parts to a compound name:
 - *Base*: Tells how many carbons are in the longest continuous chain.



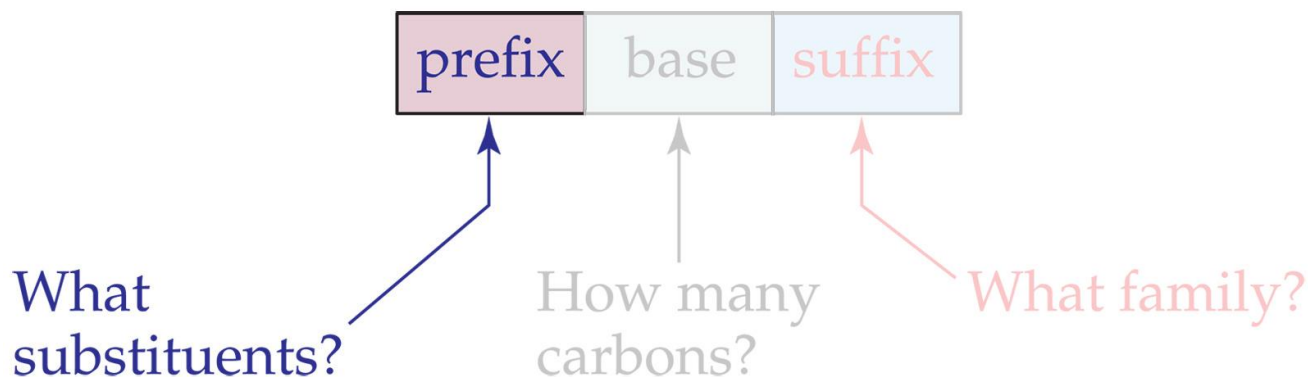
Organic Nomenclature

- Three parts to a compound name:
 - Base: Tells how many carbons are in the longest continuous chain.
 - Suffix: Tells what type of compound it is.

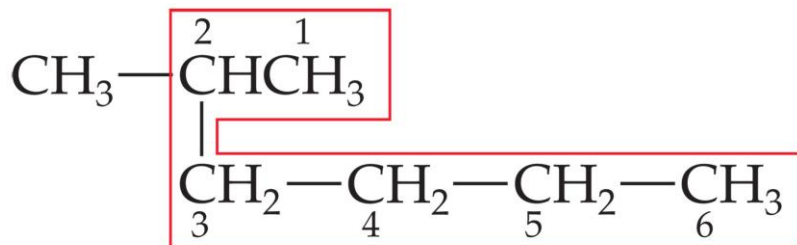


Organic Nomenclature

- Three parts to a compound name:
 - Base: Tells how many carbons are in the longest continuous chain.
 - Suffix: Tells what type of compound it is.
 - Prefix: Tells what groups are attached to chain.



To Name a Compound...

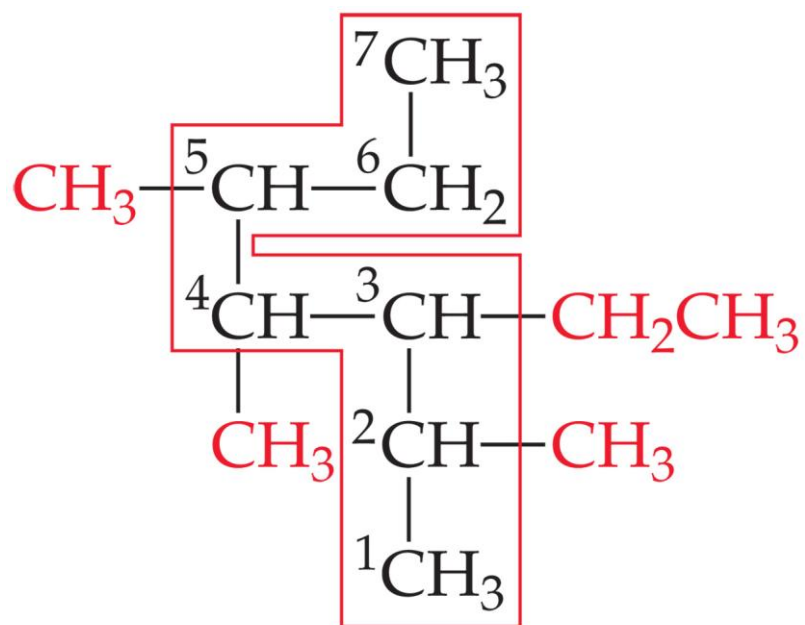


2-Methylhexane

1. Find the longest chain in the molecule.
2. Number the chain from the end nearest the first substituent encountered.
3. List the substituents as a prefix along with the number(s) of the carbon(s) to which they are attached.

Group	Name
CH ₃ —	Methyl
CH ₃ CH ₂ —	Ethyl
CH ₃ CH ₂ CH ₂ —	Propyl
CH ₃ CH ₂ CH ₂ CH ₂ —	Butyl
$\begin{array}{c} \text{CH}_3 \\ \\ \text{HC}— \\ \\ \text{CH}_3 \end{array}$	Isopropyl
$\begin{array}{c} \text{CH}_3 \\ \\ \text{CH}_3—\text{C}— \\ \\ \text{CH}_3 \end{array}$	<i>tert</i> -Butyl

To Name a Compound...

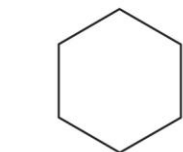
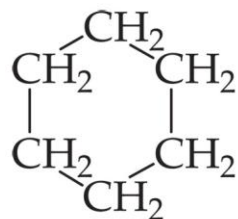


3-Ethyl-2,4,5-trimethylheptane

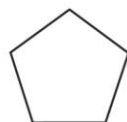
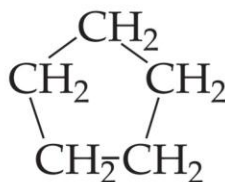
If there is more than one type of substituent in the molecule, list them alphabetically.

Cycloalkanes

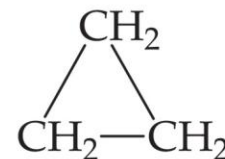
- Carbon can also form ringed structures.
- Five- and six-membered rings are most stable.
 - Can take on conformation in which angles are very close to tetrahedral angle.
 - Smaller rings are quite strained.



Cyclohexane



Cyclopentane



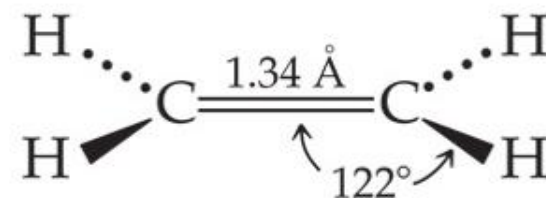
Cyclopropane

Reactions of Alkanes

- Rather unreactive due to presence of only C—C and C—H σ -bonds.
- Therefore, great nonpolar solvents.

Alkenes

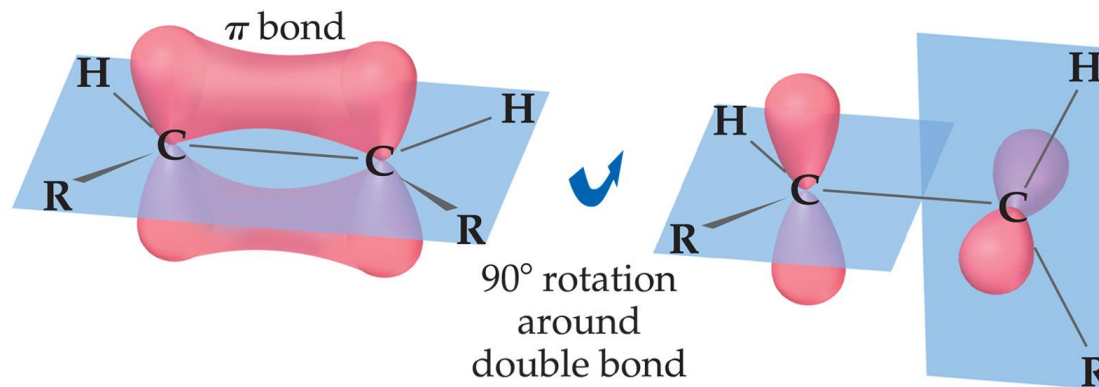
ALKENE
Ethylene



- Contain at least one carbon-carbon double bond.
- Unsaturated.
 - Have fewer than maximum number of hydrogens.

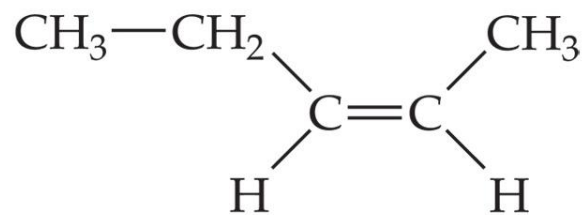
Structure of Alkenes

- Unlike alkanes, alkenes cannot rotate freely about the double bond.
 - Side-to-side overlap makes this impossible without breaking π -bond.

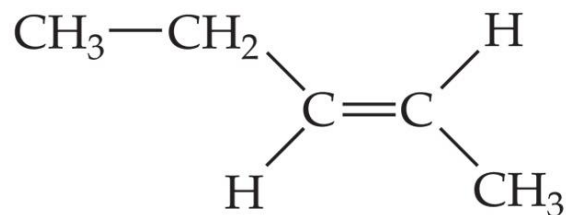


Structure of Alkenes

This creates
geometric isomers,
which differ from
each other in the
spatial arrangement
of groups about the
double bond.

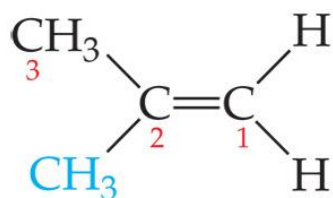


cis-2-Pentene

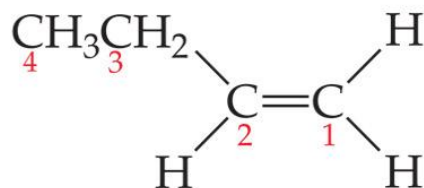


trans-2-Pentene

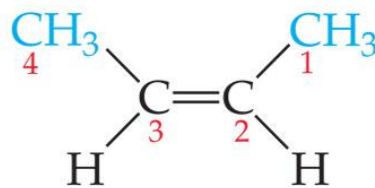
Properties of Alkenes



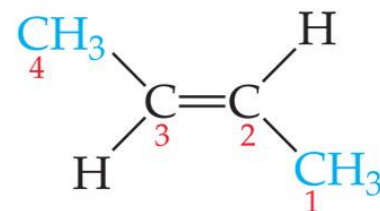
Methylpropene
bp -7°C



1-Butene
bp -6°C



cis-2-Butene
bp $+4^{\circ}\text{C}$

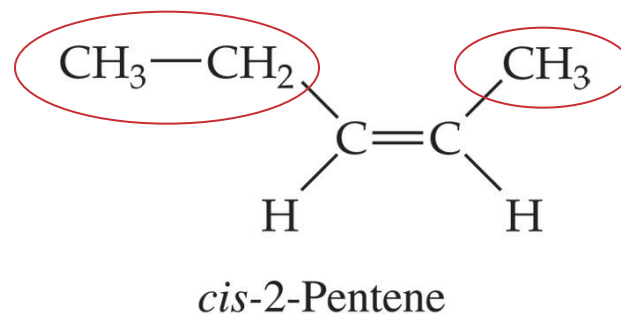
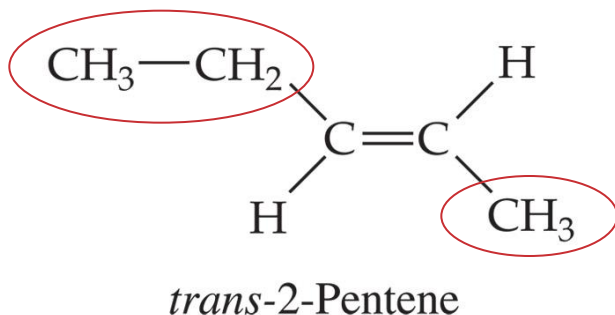
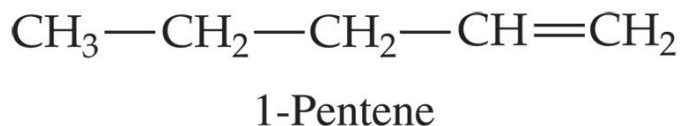


trans-2-Butene
bp $+1^{\circ}\text{C}$

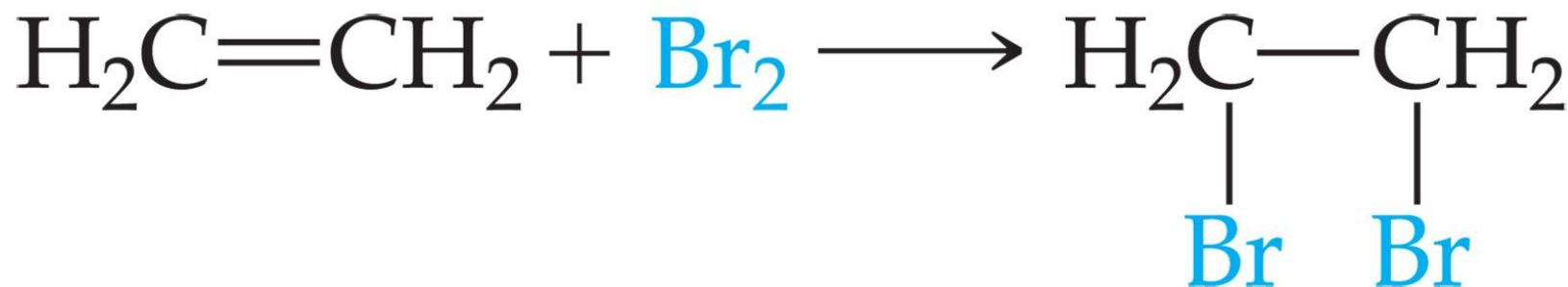
Structure also affects physical properties of alkenes.

Nomenclature of Alkenes

- Chain numbered so double bond gets smallest possible number.
- *cis*- alkenes have carbons in chain on same side of molecule.
- *trans*- alkenes have carbons in chain on opposite side of molecule.



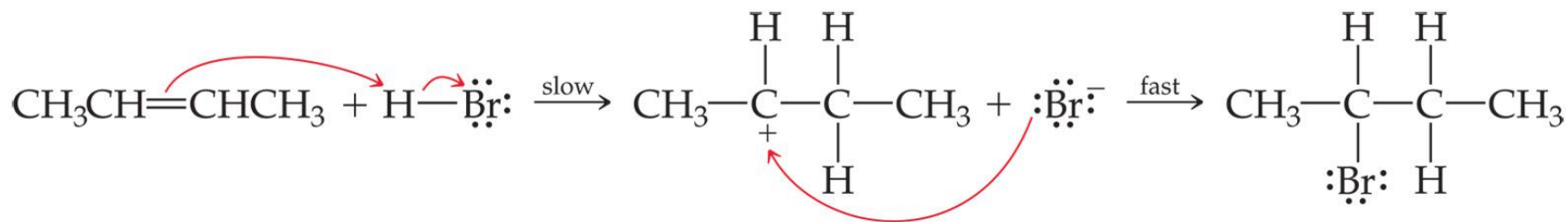
Reactions of Alkenes



- Addition Reactions

- Two atoms (e.g., bromine) add across the double bond.
- One π -bond and one σ -bond are replaced by two σ -bonds; therefore, ΔH is negative.

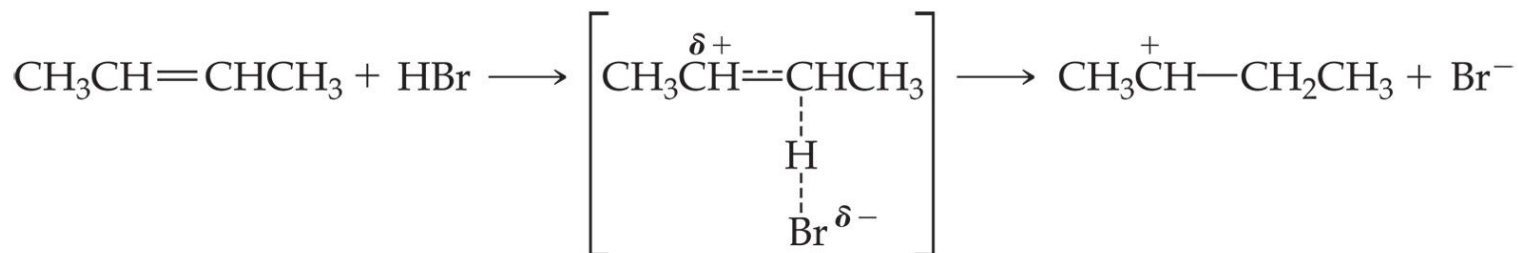
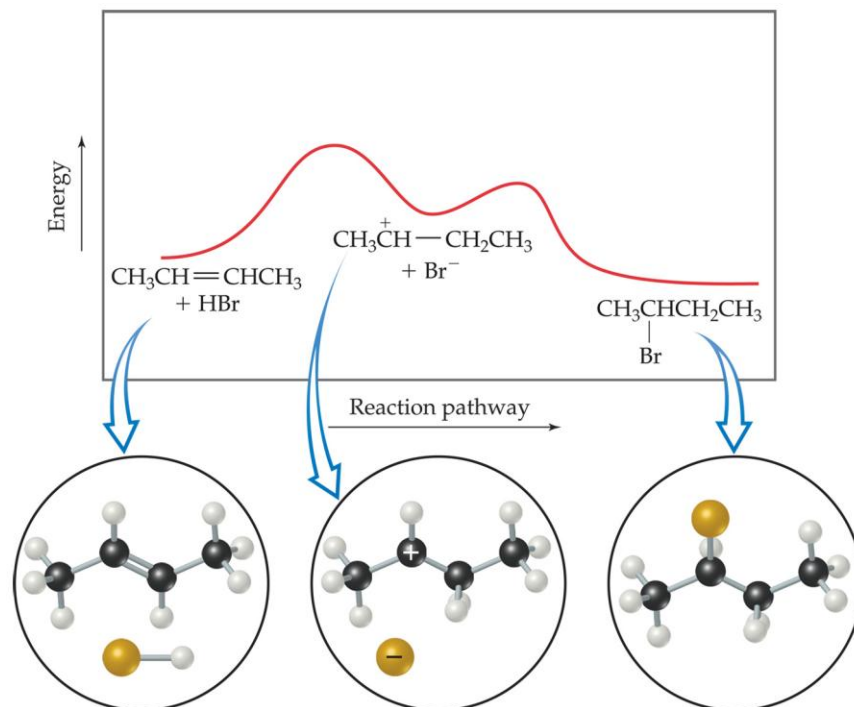
Mechanism of Addition Reactions



- Two-step mechanism:
 - First step is slow, rate-determining step.
 - Second step is fast.

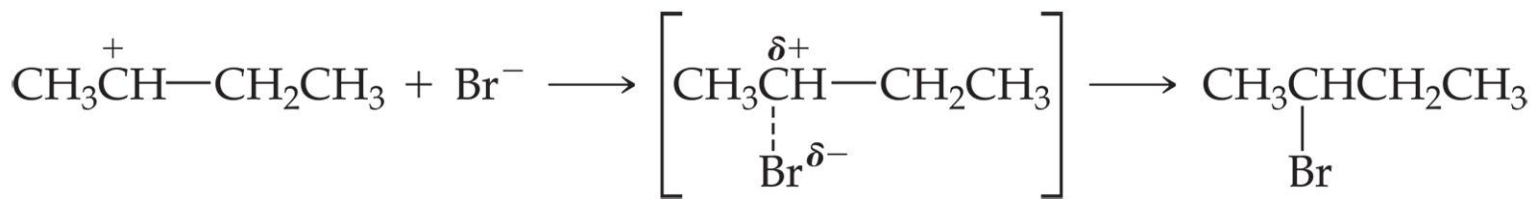
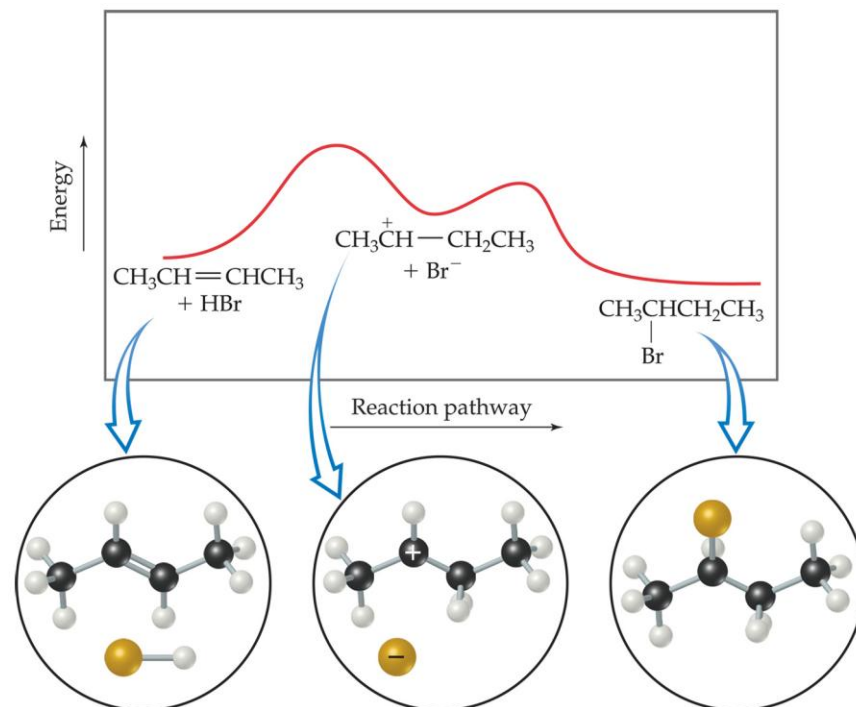
Mechanism of Addition Reactions

In first step, π -bond breaks and new C—H bond and cation form.



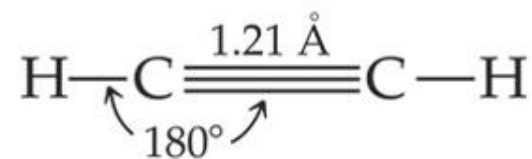
Mechanism of Addition Reactions

In second step, new bond forms between negative bromide ion and positive carbon.



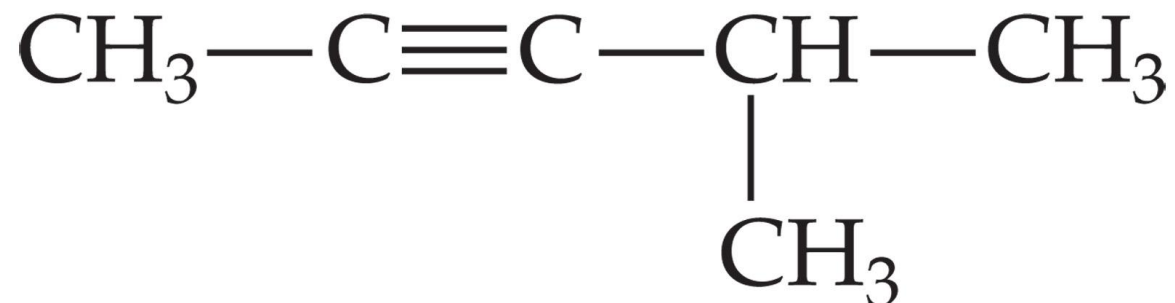
Alkynes

ALKYNE
Acetylene $\text{CH}\equiv\text{CH}$



- Contain at least one carbon–carbon triple bond.
- Carbons in triple bond *sp*-hybridized and have linear geometry.
- Also unsaturated.

Nomenclature of Alkynes

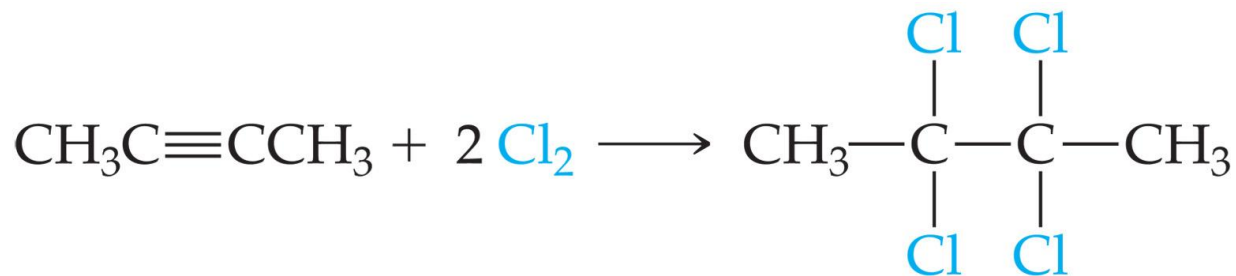


4-methyl-2-pentyne

- Analogous to naming of alkenes.
- Suffix is *-yne* rather than *-ene*.

Reactions of Alkynes

- Undergo many of the same reactions alkenes do.
- As with alkenes, impetus for reaction is replacement of π -bonds with σ -bonds.

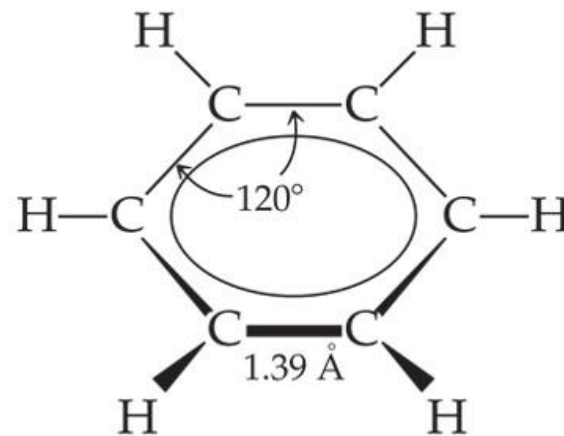
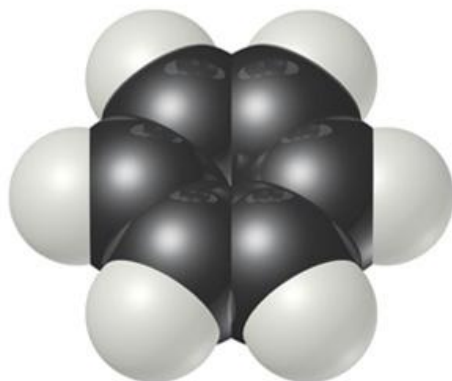


2-Butyne

2,2,3,3-Tetrachlorobutane

Aromatic Hydrocarbons

AROMATIC
Benzene C_6H_6



- Cyclic hydrocarbons.
- *p*-Orbital on each atom.
 - Molecule is planar.
- Odd number of electron pairs in π -system.

Aromatic Nomenclature

Many aromatic hydrocarbons are known by their common names.



Benzene



Naphthalene

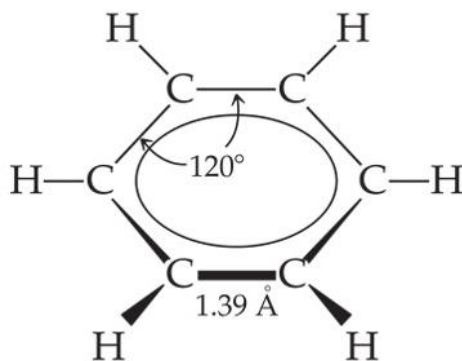
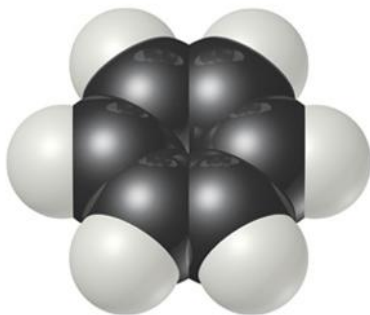


Anthracene



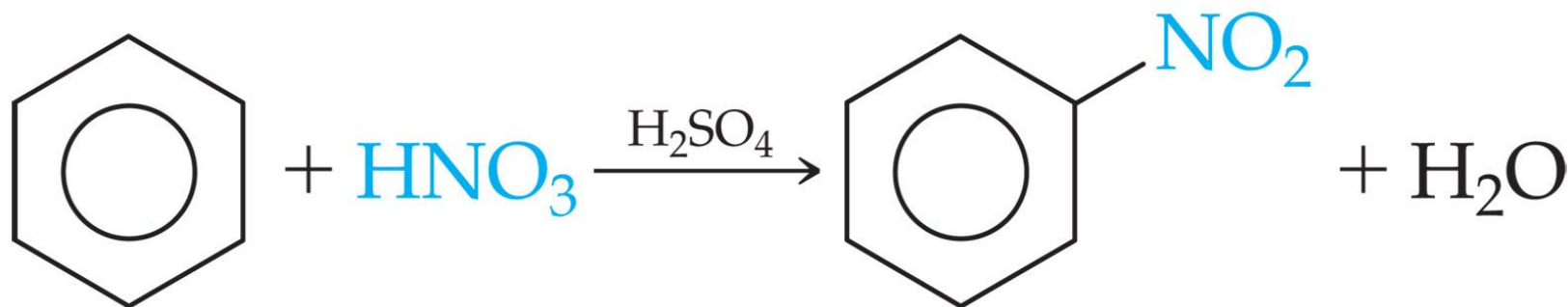
Toluene
Methylbenzene

Reactions of Aromatic Compounds



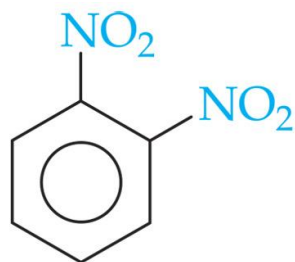
- Unlike in alkenes and alkynes, π -electrons do not sit between two atoms.
- Electrons are delocalized; this stabilizes aromatic compounds.

Reactions of Aromatic Compounds

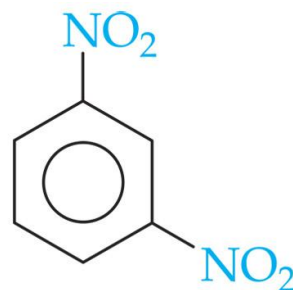


- Due to stabilization, aromatic compounds do not undergo addition reactions; they undergo substitution.
- Hydrogen is replaced by substituent.

Structure of Aromatic Compounds



ortho-Dinitrobenzene
mp 118°C



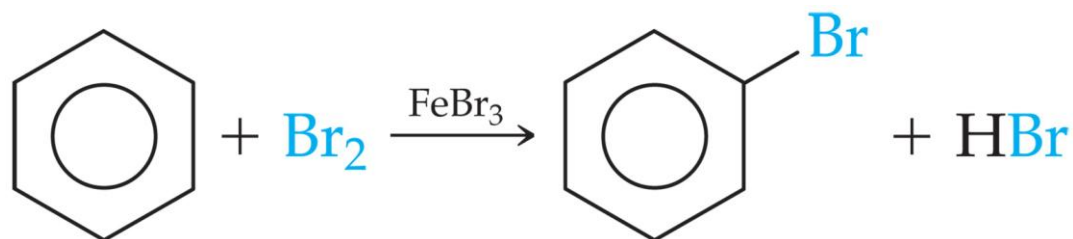
meta-Dinitrobenzene
mp 90°C



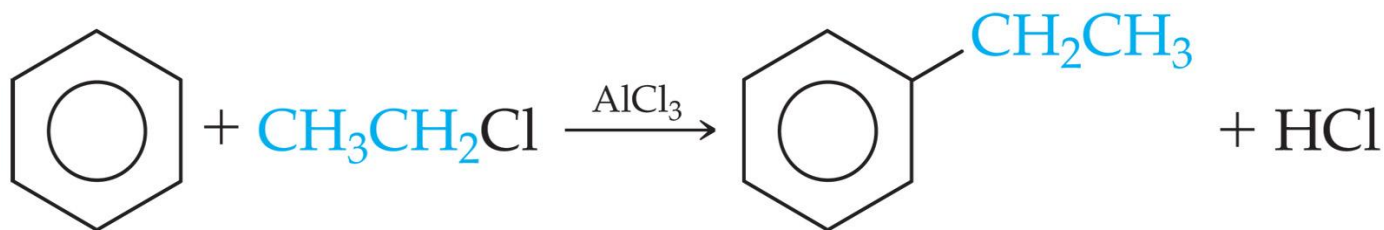
para-Dinitrobenzene
mp 174°C

- Two substituents on a benzene ring could have three possible relationships
 - *ortho*-: On adjacent carbons.
 - *meta*-: One carbon between them.
 - *para*-: On opposite sides of ring.

Reactions of Aromatic Compounds



Halogenation



Friedel-Crafts Reaction

Reactions of aromatic compounds often require a catalyst.

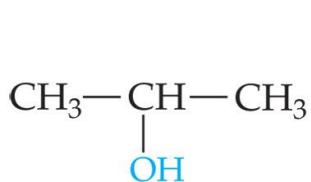
Functional Group	Type of Compound	Suffix or Prefix	Example	Systematic Name (common name)
	Alkene	-ene		Ethene (Ethylene)
	Alkyne	-yne		Ethyne (Acetylene)
	Alcohol	-ol		Methanol (Methyl alcohol)
	Ether	ether		Dimethyl ether
	Haloalkane	halo-		Chloromethane (Methyl chloride)
(X = halogen)				
	Amine	-amine		Ethylamine
	Aldehyde	-al		Ethanal (Acetaldehyde)
	Ketone	-one		Propanone (Acetone)
	Carboxylic acid	-oic acid		Ethanoic acid (Acetic acid)
	Ester	-oate		Methyl ethanoate (Methyl acetate)
	Amide	-amide		Ethanamide (Acetamide)

Functional Groups

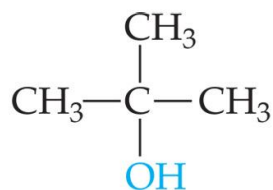
Term used to refer to parts of organic molecules where reactions tend to occur.

Alcohols

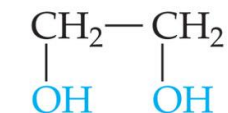
- Contain one or more hydroxyl groups, —OH
 - Named from parent hydrocarbon; suffix changed to *-ol* and number designates carbon to which hydroxyl is attached.



2-Propanol
Isopropyl alcohol;
rubbing alcohol



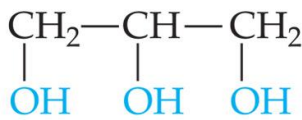
2-Methyl-2-propanol
t-Butyl alcohol



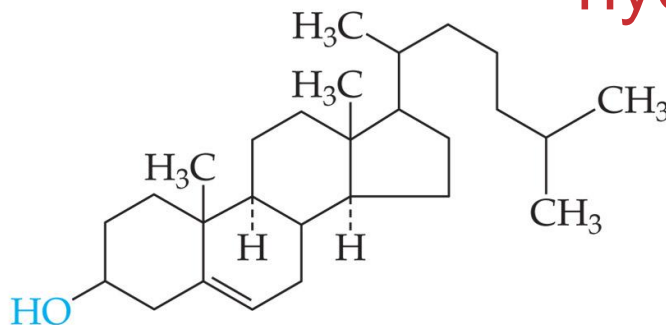
1,2-Ethandiol
Ethylene glycol



Phenol



1,2,3-Propanetriol
Glycerol; glycerin



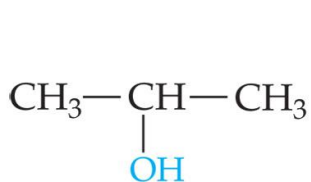
Cholesterol

Alcohols

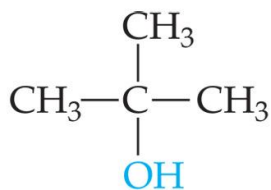
- Much more acidic than hydrocarbons.

➤ $pK_a \sim 15$ for most alcohols.

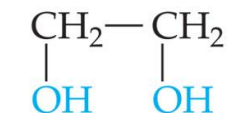
➤ Aromatic alcohols have $pK_a \sim 10$.



2-Propanol
Isopropyl alcohol;
rubbing alcohol



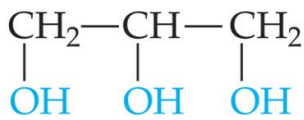
2-Methyl-2-propanol
t-Butyl alcohol



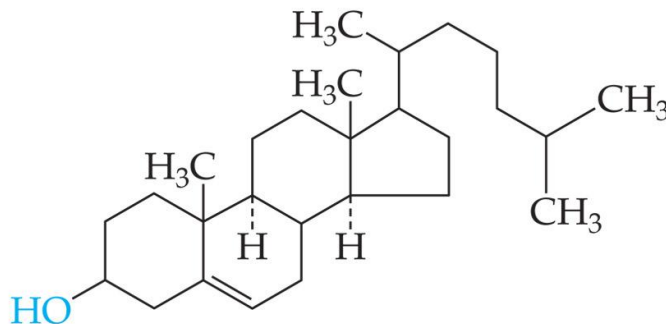
1,2-Ethanediol
Ethylene glycol



Phenol

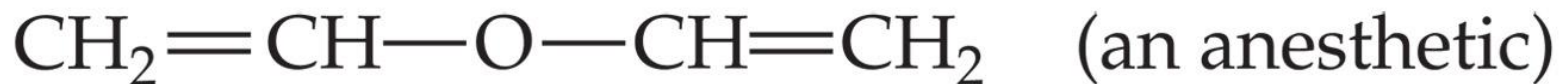


1,2,3-Propanetriol
Glycerol; glycerin



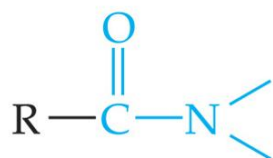
Cholesterol

Ethers



- Tend to be quite unreactive.
- Therefore, they are good polar solvents.

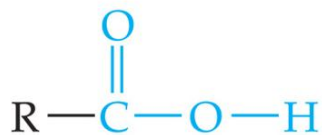
Carbonyl Compounds



Amide



Aldehyde



Carboxylic
acid



Ester

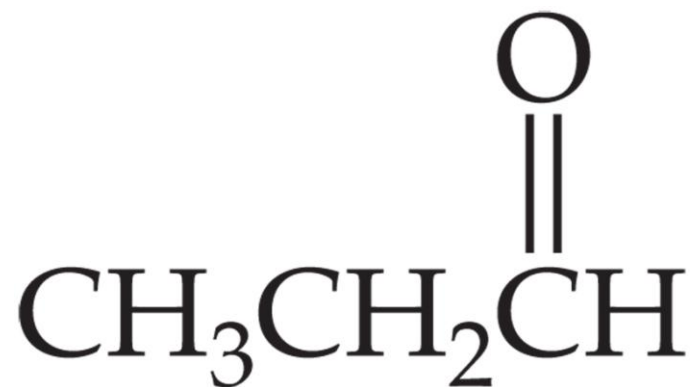


Ketone

- Contain C=O double bond.
- Include many classes of compounds.

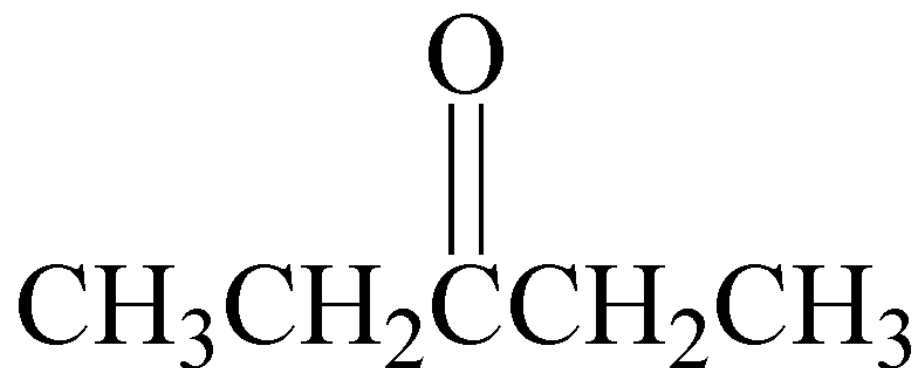
Aldehydes

At least one
hydrogen attached
to carbonyl carbon.



Ketones

Two carbons
bonded to
carbonyl carbon.

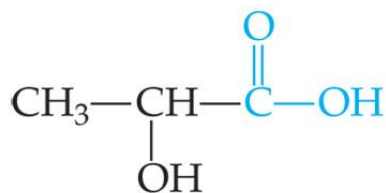


Carboxylic Acids

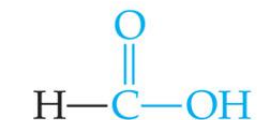
- Have hydroxyl group bonded to carbonyl group.
- Tart tasting.
- Carboxylic acids are weak acids□□.



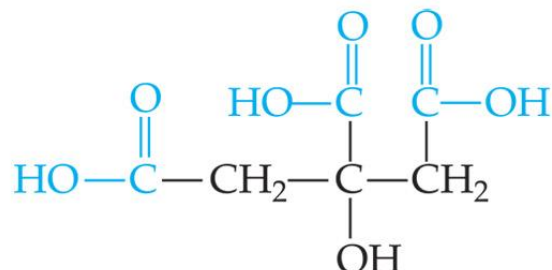
Carboxylic Acids



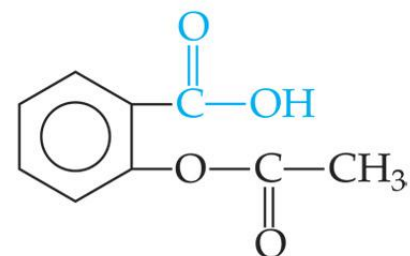
Lactic acid



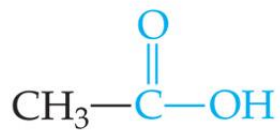
Methanoic acid
Formic acid



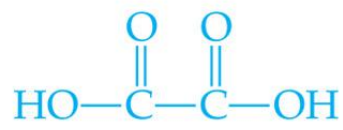
Citric acid



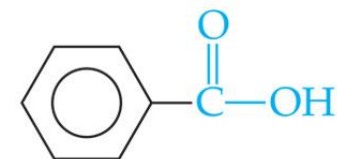
Acetylsalicylic acid
Aspirin



Ethanoic acid
Acetic acid

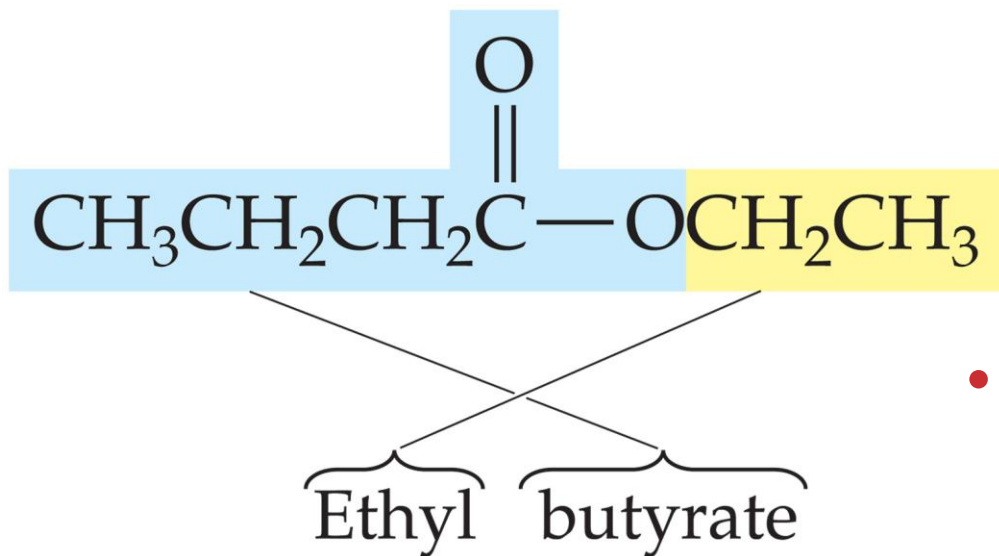


Oxalic acid



Phenyl methanoic acid
Benzoic acid

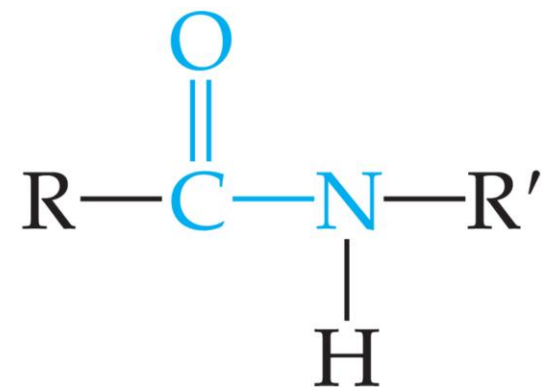
Esters



- Products of reaction between carboxylic acids and alcohols.
- Found in many fruits and perfumes.

Amides

Formed by reaction of carboxylic acids with amines.



Amines

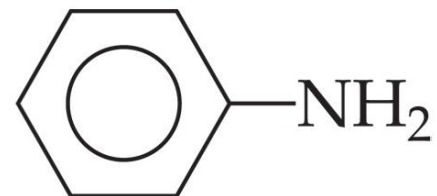
- Organic bases.
- Generally have strong, unpleasant odors.



Ethylamine



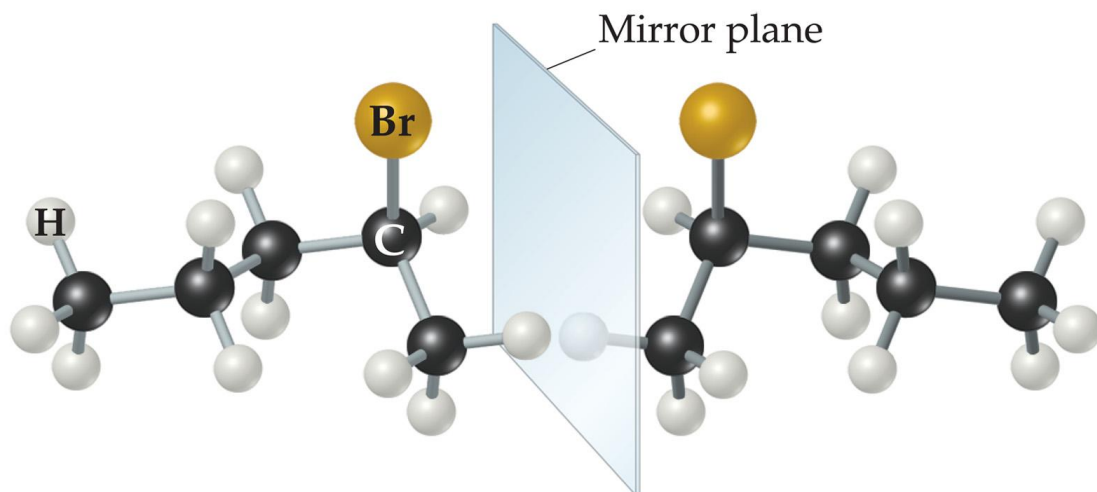
Trimethylamine



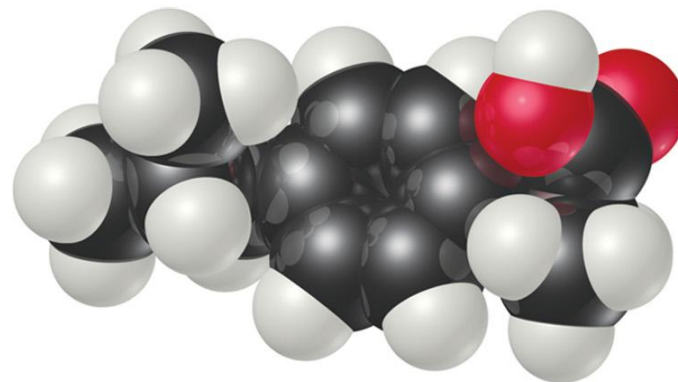
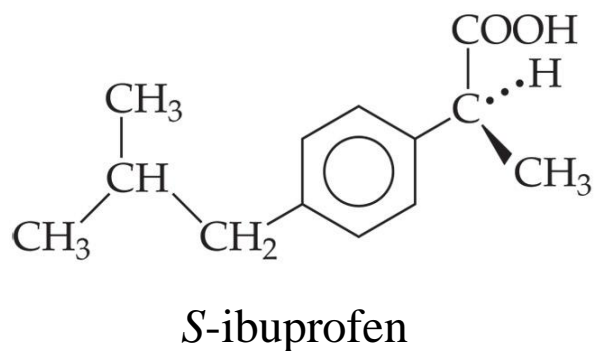
Phenylamine
Aniline

Chirality

- Carbons with four different groups attached to them are handed, or **chiral**.
- Optical isomers or **stereoisomers**
- If one stereoisomer is “right-handed,” its **enantiomer** is “left-handed.”



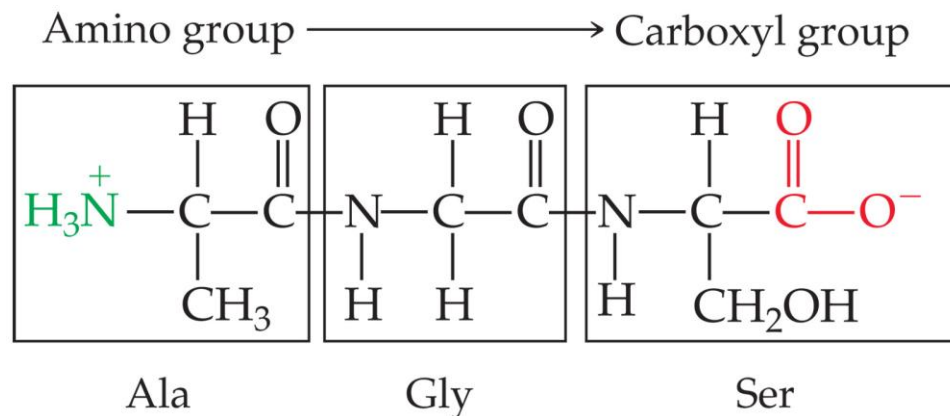
Chirality



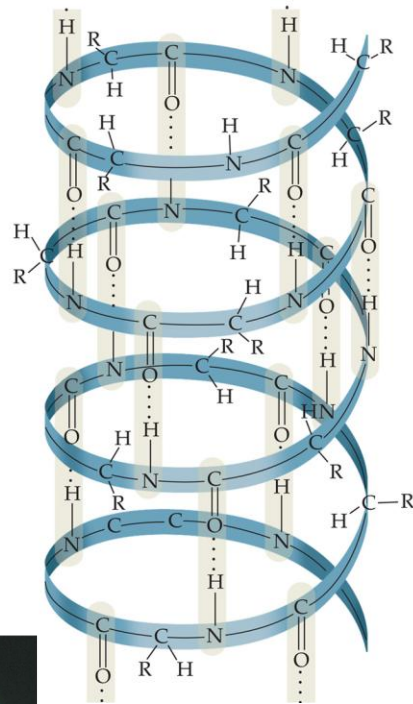
- Many pharmaceuticals are chiral.
- Often only one enantiomer is clinically active.

Amino Acids and Proteins

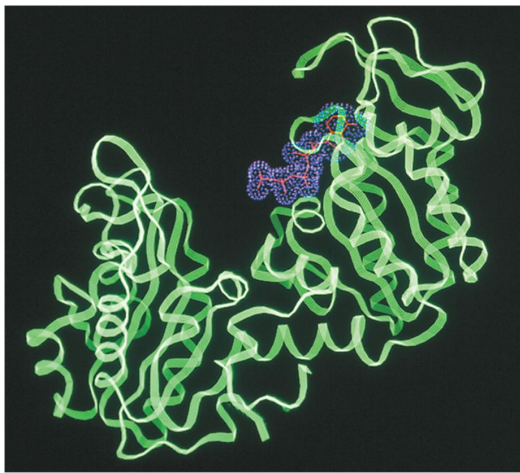
- Proteins are polymers of α -amino acids.
- A condensation reaction between the amine end of one amino acid and the acid end of another produces a peptide bond.



Amino Acids and Proteins

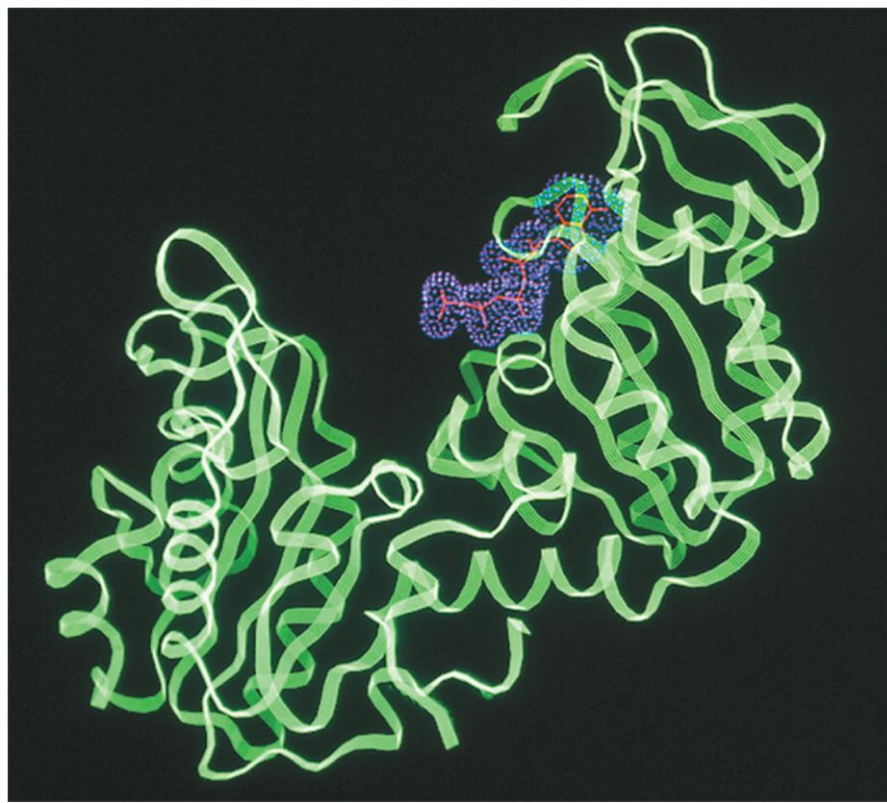


- Hydrogen bonding in peptide chains causes coils and helices in the chain.
- Kinking and folding of the coiled chain gives proteins a characteristic shape.



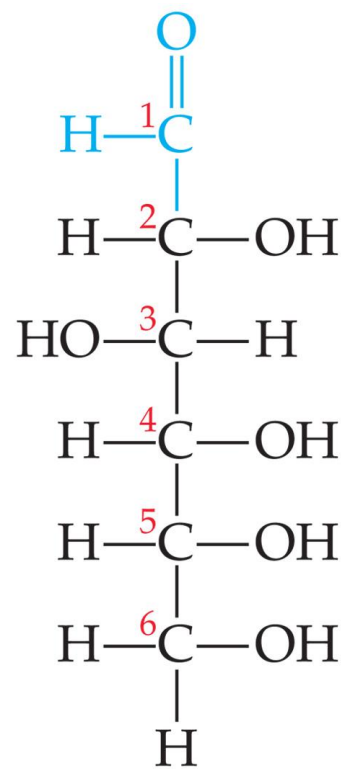
Amino Acids and Proteins

- Most enzymes are proteins.
- The shape of the active site complements the shape of the substrate on which the enzyme acts—hence, the “lock-and-key” model.

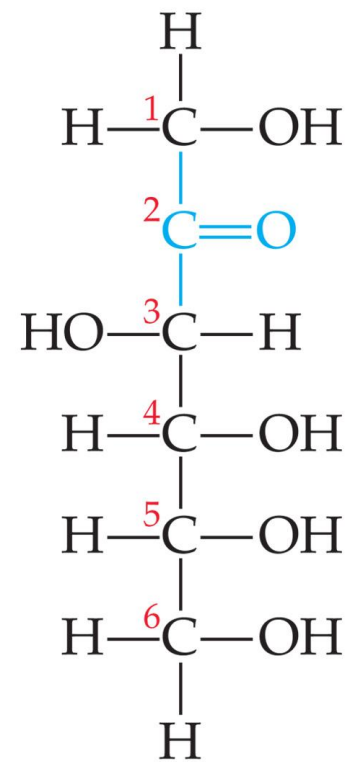


Carbohydrates

Simple sugars are polyhydroxy aldehydes or ketones.

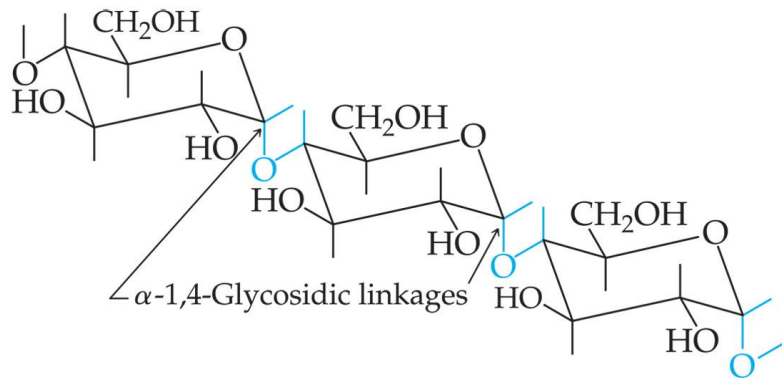
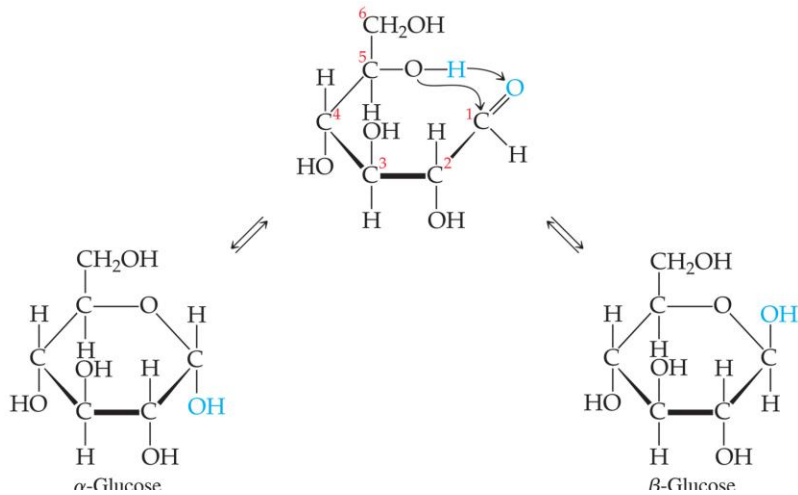


Glucose



Fructose

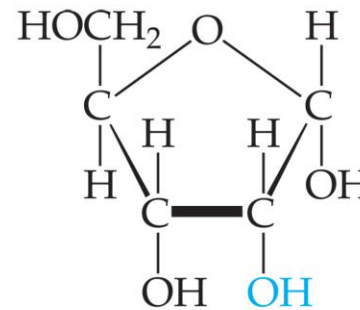
Carbohydrates



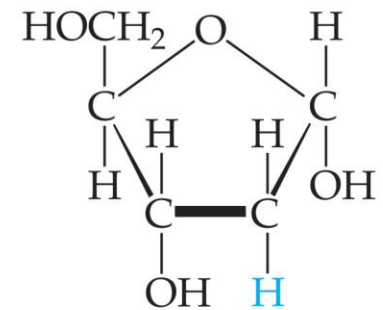
- In solution they form cyclic structures.
- These can form chains of sugars that form structural molecules such as starch and cellulose.

Nucleic Acids

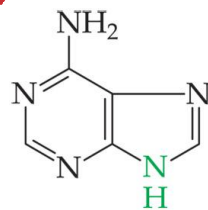
Two of the building blocks of RNA and DNA are sugars (ribose or deoxyribose) and cyclic bases (adenine, guanine, cytosine, and thymine or uracil).



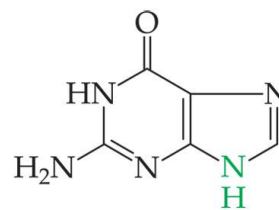
Ribose



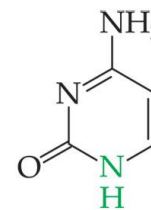
Deoxyribose



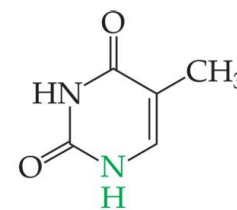
Adenine (A)
DNA
RNA



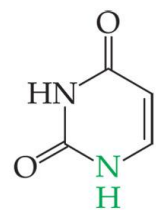
Guanine (G)
DNA
RNA



Cytosine (C)
DNA
RNA



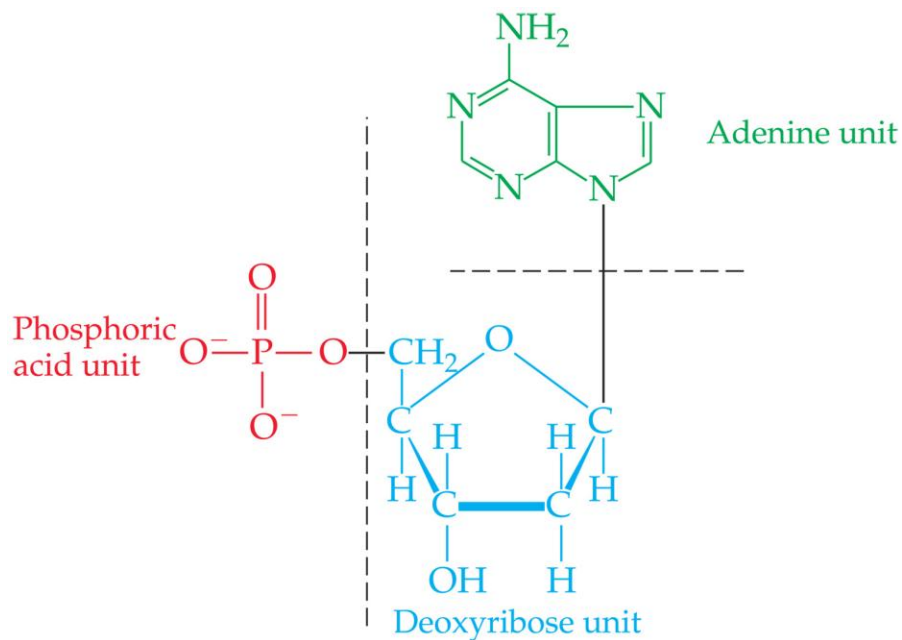
Thymine (T)
DNA



Uracil (U)
RNA

Nucleic Acids

These combine with a phosphate to form a nucleotide.



Nucleic Acids

Nucleotides combine to form the familiar double-helix form of the nucleic acids.

