Chapter 2

Alkanes and Cycloalkanes; Conformational and Geometrical Isomerism

- **Hydrocarbons** are compounds that contain only carbon and hydrogen.

There are three main classes of hydrocarbons, based on the types of carbon–carbon bonds present.

1-**Saturated hydrocarbons** contain only carbon–carbon *single* bonds.

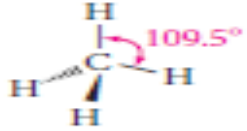
2-**Unsaturated hydrocarbons** contain carbon–carbon *multiple* bonds, double bonds, triple bonds, or both.

3-**Aromatic hydrocarbons** are a special class of cyclic compounds related in structure to benzene.

- Saturated hydrocarbons are known as **alkanes** if they are acyclic, or as
- **cycloalkanes** if they are cyclic.

2.1 The Structure of Alkanes

- Alkanes are hydrocarbons containing only single saturated bonds. General formula: C_nH_{2n+2}.
- The simplest alkane is methane.
- Its tetrahedral three-dimensional structure .
- Alkanes with carbon chains that are unbranched are called **normal alkanes** or *n*-alkanes.



- Each member of this series differs from the next higher and the next lower member by a -CH_2-group (called a **methylene group**).
- A series of compounds in which the members are built up in a regular, repetitive way like this is called a **homologous series**.

Table 2.1 Names and Formulas of the First Ten Unbranched Alkanes

Name	Number of carbons	Molecular formula	Structural formula	Number of structural isomers
methane	1	CH_4	CH_4	1
ethane	2	C_2H_6	CH_3CH_3	1
propane	3	C_3H_8	$CH_3CH_2CH_3$	1
butane	4	C_4H_{10}	$CH_3CH_2CH_2CH_3$	2
pentane	5	C_5H_{12}	$CH_3(CH_2)_3CH_3$	3
hexane	6	C_6H_{14}	$CH_3(CH_2)_4CH_3$	5
heptane	7	C_7H_{16}	$CH_3(CH_2)_5CH_3$	9
octane	8	C_8H_{18}	$CH_3(CH_2)_6CH_3$	18
nonane	9	C_9H_{20}	$CH_3(CH_2)_7CH_3$	35
decane	10	$C_{10}H_{22}$	$CH_3(CH_2)_8CH_3$	75

- Compounds of a **homologous series** differ by *a regular unit* of structure and share *similar properties*.

- **PROBLEM 2.2** Which of the following are alkanes?

a. C_7H_{16} ✓

b. C_7H_{12} ✗ cyclic alkene

c. C_8H_{16} ✗ cyclic alkane / alkene

d. $C_{29}H_{60}$

2.2 Nomenclature of organic compounds

- In the early days of organic chemistry, each new compound was given a name that was usually based on its source or use.

  Examples include
  limonene (from lemons), *α*-pinene (from pine trees), coumarin (from the tonka bean, known to South American natives as *cumaru*)

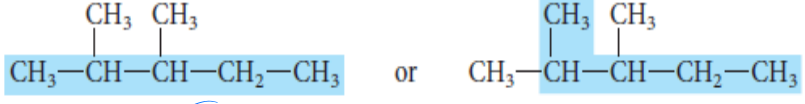
- Internationally recognized systems of nomenclature were devised by a commission of the International Union of Pure and Applied Chemistry; they are known as the IUPAC

2.3 IUPAC rules for naming alkanes

1. The general name for acyclic saturated hydrocarbons is *alkanes*. The -***ane*** ending is used for all saturated hydrocarbons. This is important to remember because later other endings will be used for other functional groups.

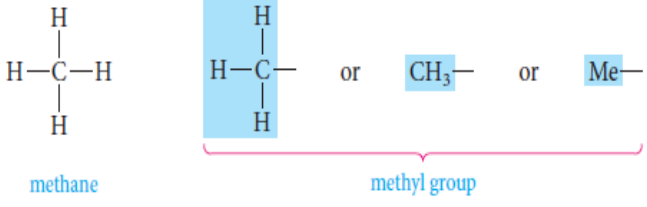
2. Alkanes without branches are named according to the *number of carbon atoms*. These names, up to ten carbons, are given in the first column of Table 2.1.

3. For alkanes with branches, the **root name** is that of the longest continuous chain of carbon atoms. For example, in the structure



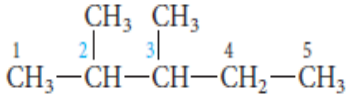
the longest continuous chain (in color) has five carbon atoms. The compound is therefore named as a substituted *pent*ane, even though there are seven carbon atoms altogether.

4. Groups attached to the main chain are called **substituents**. Saturated substituents that contain only carbon and hydrogen are called **alkyl groups**. An alkyl group is named by taking the name of the alkane with the same number of carbon atoms and changing the - *ane* ending to *-yl*.



5. The main chain is numbered in such a way that the first substituent encountered along the chain receives the lowest possible number. Each substituent is then located by its name and by the number of the carbon atom to which it is attached.

- When two or more identical groups are attached to the main chain, prefixes such as *di-, tri-*, and *tetra-* are used. *Every substituent must be named and numbered,* even if two identical substituents are attached to the same carbon of the main chain.



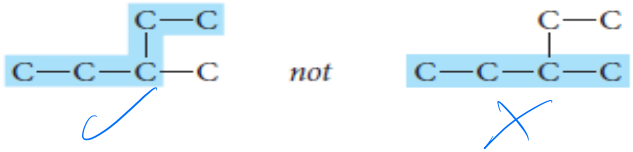
2,3-dimethylpentane

6. If two or more different types of substituents are present, they are listed alphabetically, except that prefixes such as *di-* and *tri-* are not considered when alphabetizing.

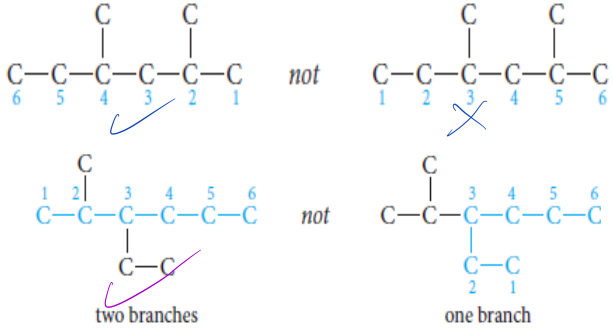
7. Punctuation is important when writing IUPAC names. IUPAC names for hydrocarbons are written as one word. Numbers are separated from each other by commas and are separated from letters by hyphens. There is no space between the last named substituent and the name of the parent alkane that follows it.

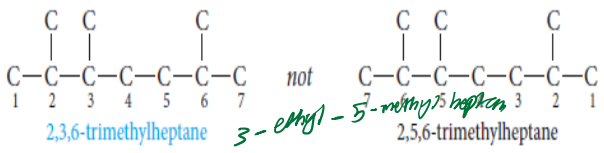
notes

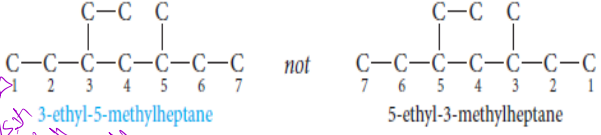
1. Locate the longest continuous carbon chain. This gives the name of the parent hydrocarbon. For example,



2. Number the longest chain beginning at the end nearest the first branch point. For example,

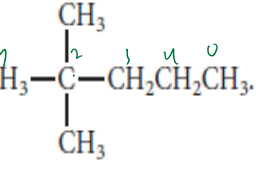






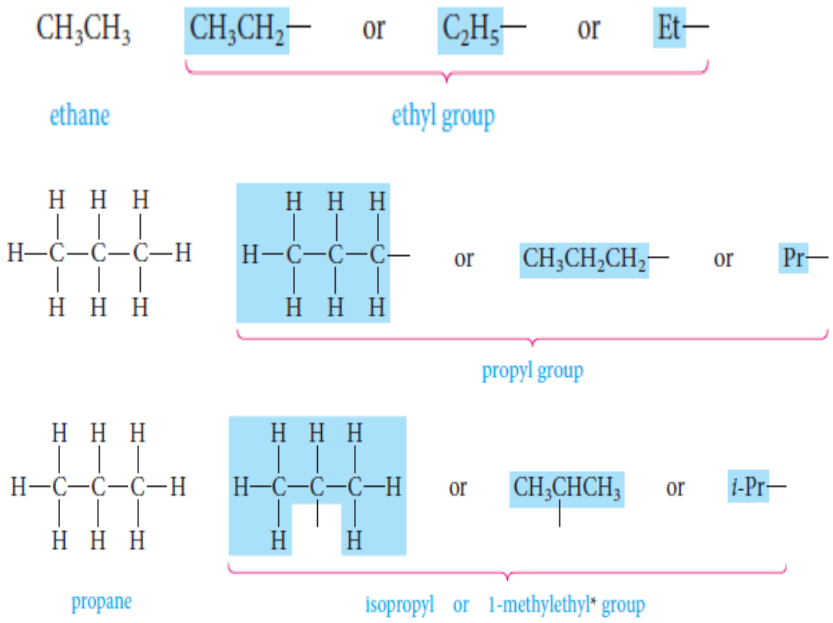
3. Write the name as one word, placing substituents in alphabetic order and using proper punctuation.

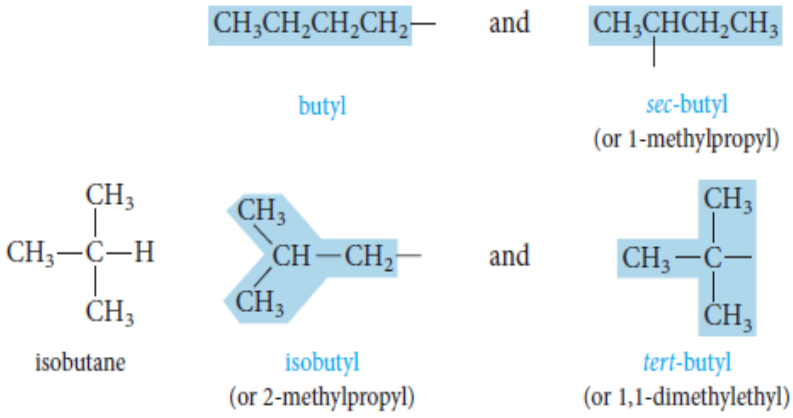
Example 2.2

Give an IUPAC name for  .

2,2-dimethylpentane

2.4 Alkyl and Halogen Substituents





R is the general symbol for an alkyl group.

Halogen substituents are named by changing the *-ine* ending of the element to *-o*.

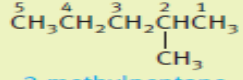
F-	Cl-	Br-	I-
fluoro-	chloro-	bromo-	iodo-

Give the common and IUPAC names for $CH_3CH_2CH_2Br$.

Bromobutane

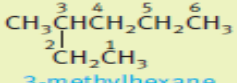
2.5 Use of the IUPAC Rules

Table 2.2 Examples of Use of the IUPAC Rules



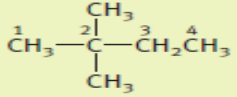
2-methylpentane
(*not* 4-methylpentane)

The ending *-ane* tells us that all the carbon–carbon bonds are single; *pent-* indicates five carbons in the longest chain. We number them from right to left, starting closest to the branch point.



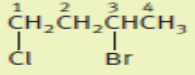
3-methylhexane
(*not* 2-ethylpentane
or 4-methylhexane)

This is a six-carbon saturated chain with a methyl group on the third carbon. We would usually write the structure as $CH_3CH_2CHCH_2CH_2CH_3$ (with CH_3 on the third carbon).



2,2-dimethylbutane
(*not* 2,2-methylbutane
or 2-dimethylbutane)

There must be a number for each substituent, and the prefix *di-* says that there are two methyl substituents.

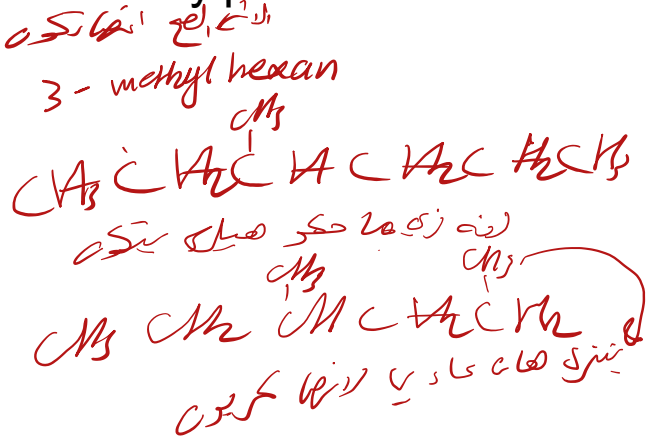


3-bromo-1-chlorobutane
(*not* 1-chloro-3-bromobutane
or 2-bromo-4-chlorobutane)

First, we number the butane chain from the end closest to the first substituent. Then we name the substituents in alphabetical order, regardless of position number.

- **PROBLEM 2.8**

Explain why 1,3-difluorobutane is a correct IUPAC name, but 1,3-dimethylpentane is *not* a correct IUPAC name.

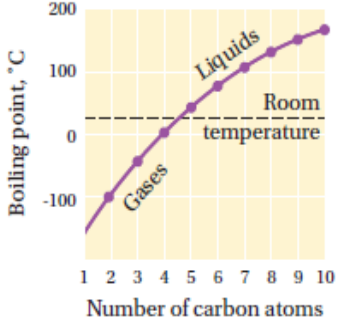


2.7 Physical properties and Intermolecular Interactions

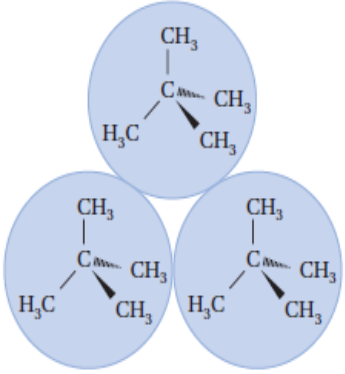
- Alkanes are insoluble in water. That is because water molecules are polar, whereas alkanes are nopolar. (all C-C and C-H bonds are nearly purely covalent.)
- Alkanes have lower boiling points for a given molecular weight than most other organic compounds. The electrons in a nonpolar molecule can become unevenly distributed within the molecule, causing the molecule to have partially positive and partially negative end. The temporarily polarized molecules causes its neighbor molecules polarized as well. Such interaction are called **Van der Waals attraction.**

Properties of alkanes

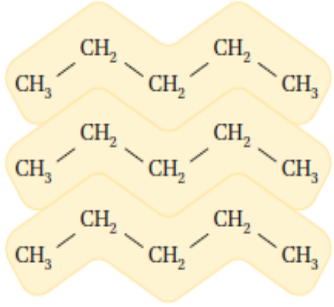
- The boiling points for alkanes rise as the chain length increases and fall as the chains become branched and more nearly spherical in shape.



Name	Formula	Boiling point, °C
pentane	$CH_3CH_2CH_2CH_2CH_3$	36
2-methylbutane (isopentane)	$CH_3CH(CH_3)CH_2CH_3$	28
2,2-dimethyl-propane (neopentane)	$C(CH_3)_4$	10



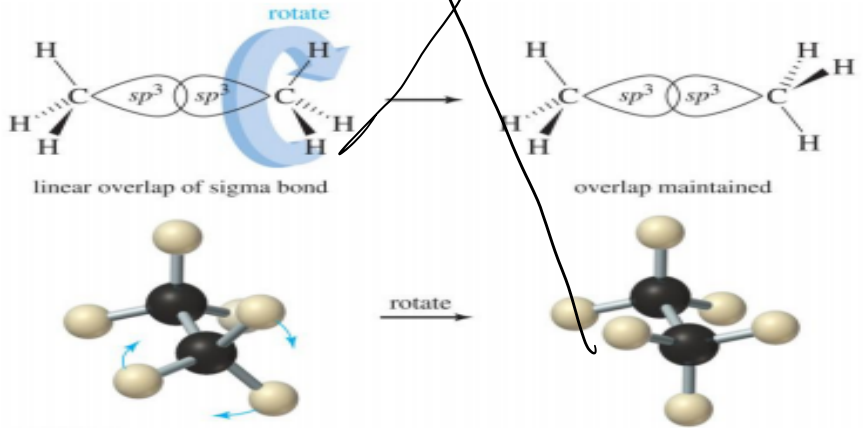
2,2-dimethylpropane
bp 10°C



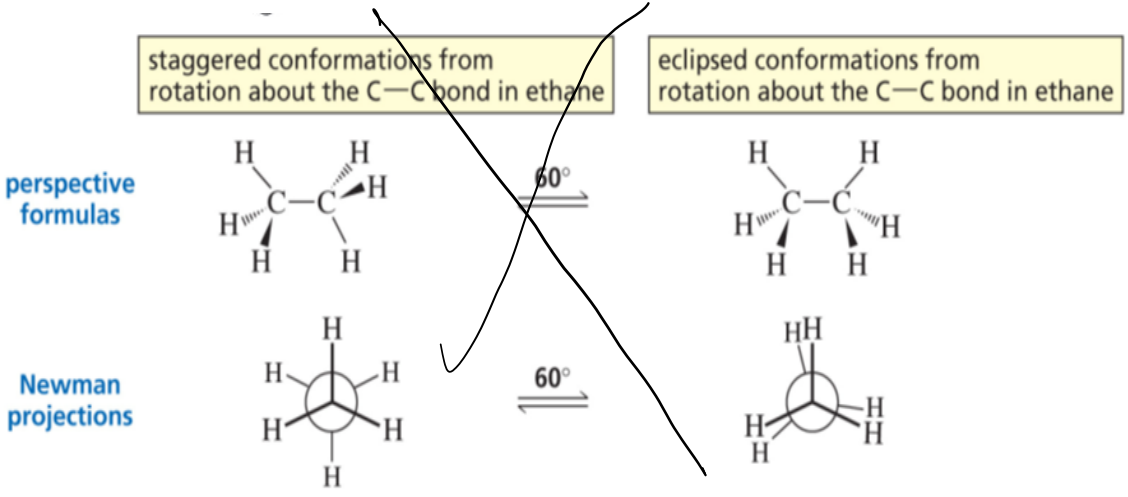
pentane
bp 36°C

2.8 Conformations of Alkanes

- A simple molecule has an infinite number of shapes as a consequence of rotating one single bond. These arrangements are called **conformations** or **conformers**. Conformers are **stereoisomers**, isomers in which the atoms are connected in the same order but are arranged differently in space.

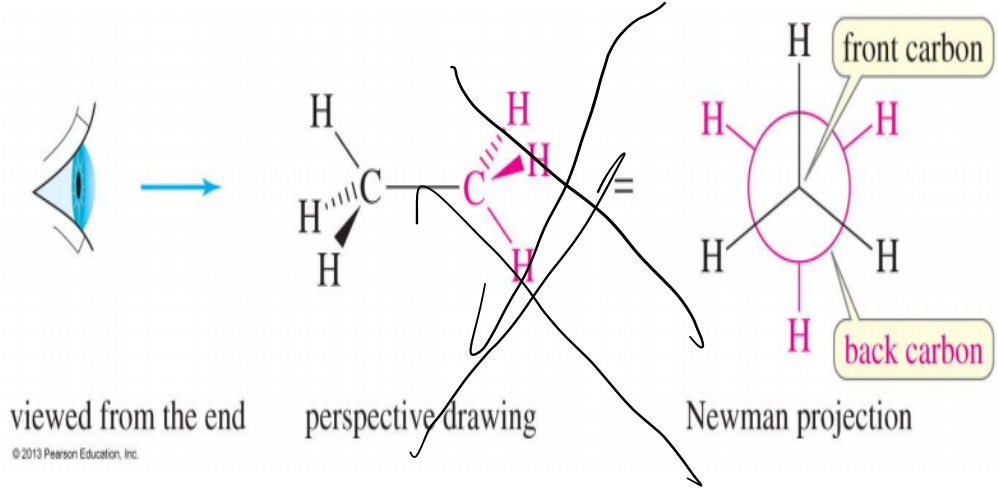


- Two possible conformers for ethane are staggered and eclipsed.



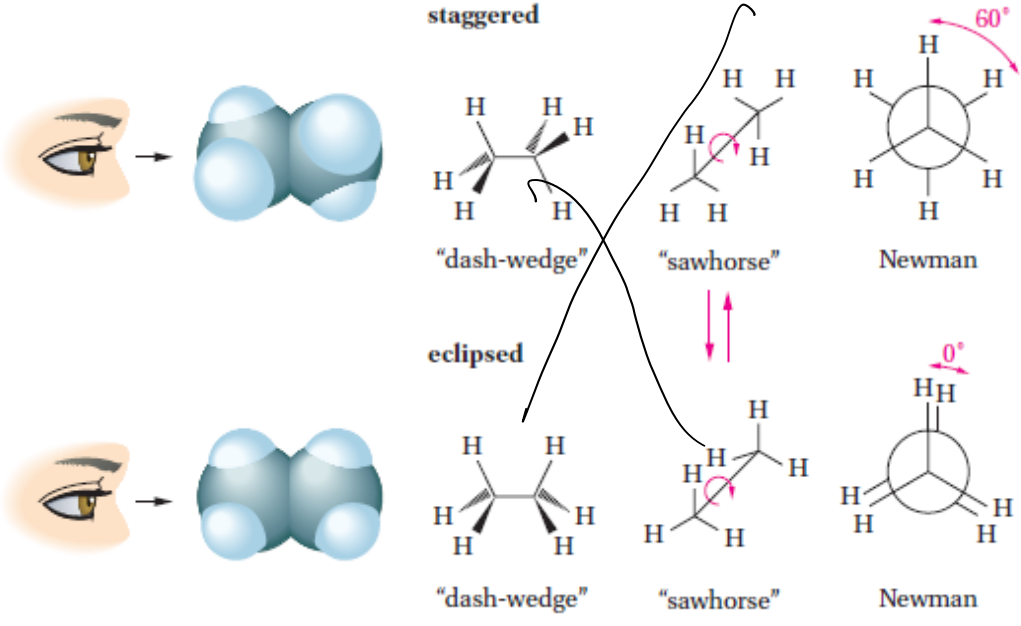
Newman Projections

The molecule is drawn as if it is viewed straight down the C-C bond.



The ***front*** **carbon** is drawn with 3 bonds in a Y shape

The ***back*** **carbon** is drawn as a circle with 3 bonds pointing out from it.



Staggered more stable than eclipsed rotamers

https://youtu.be/oG1aCQvkkD0

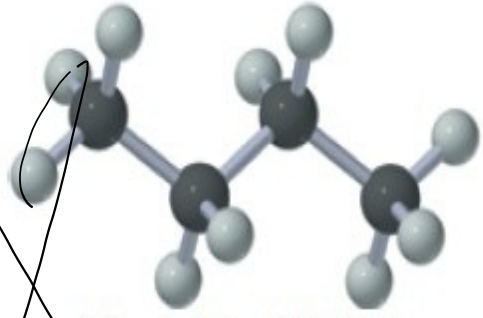
the C-2—C-3 bond

1 2 3 4
$CH_3—CH_2—CH_2—CH_3$

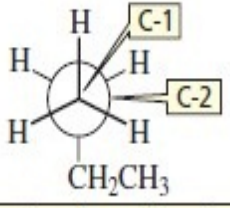
butane

the C-1—C-2 bond

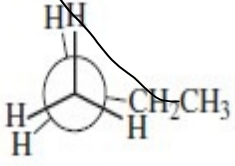
the C-3—C-4 bond



ball-and-stick model of butane



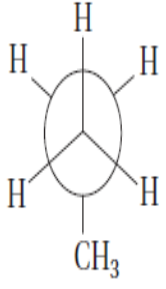
staggered conformation for rotation about the C-1—C-2 bond in butane



eclipsed conformation for rotation about the C-1—C-2 bond in butane

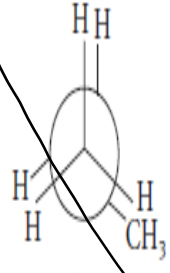
Example 2.4

Draw the Newman projections for the staggered and eclipsed conformations of propane.



staggered

The projection formula is similar to that of ethane, except for the replacement of one hydrogen with methyl.

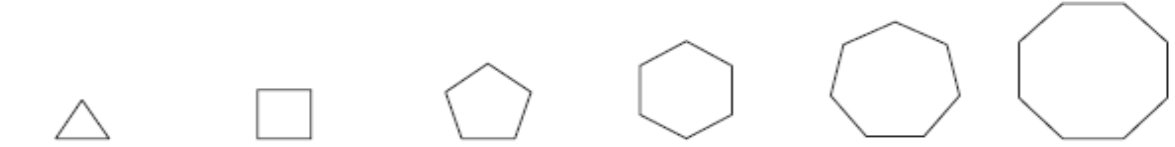


eclipsed

Rotation of the "rear" carbon of the staggered conformation by 60° gives the eclipsed conformation shown.

2.9 Cycloalkanes Nomenclature and Conformation

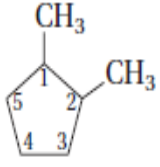
- Cycloalkanes are saturated hydrocarbons that have at least one ring of carbon atoms.
- Cycloalkanes are named by placing the prefix *cyclo*- before the alkane name that corresponds to the number of carbon atoms in the ring.



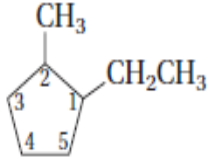
- Nomenclature
- Alkyl or halogen substituents attached to the rings are named in the usual way.
- If only one substituent is present, no number is needed to locate it.

- If there are several substituents , numbers are required. One substituent is always located at ring carbon number 1, and the remaining ring carbons are then numbered consecutively in a way that
  gives the other substituents the lowest possible numbers

methylcyclopentane
(*not* 1-methylcyclopentane)

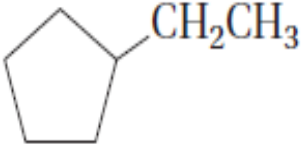


1,2-dimethylcyclopentane
(*not* 1,5-dimethylcyclopentane)

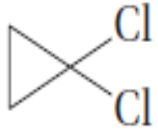


1-ethyl-2-methylcyclopentane
(*not* 2-ethyl-1-methylcyclopentane)

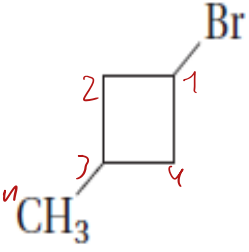
PROBLEM 2.12 Give IUPAC names for

a. 

Ethylcyclopentan

b. 

1,1-dichlorocyclopropan

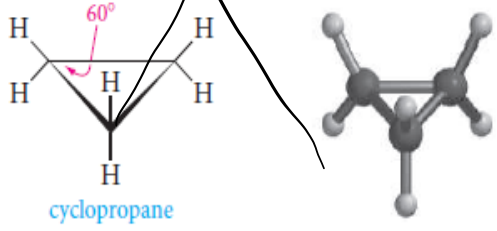
c. 

1-Bromo-3-methylcyclobutan

a. ethylcyclopentane
b. 1,1-dichlorocyclopropane
c. 1-bromo-3-methylcyclobutane

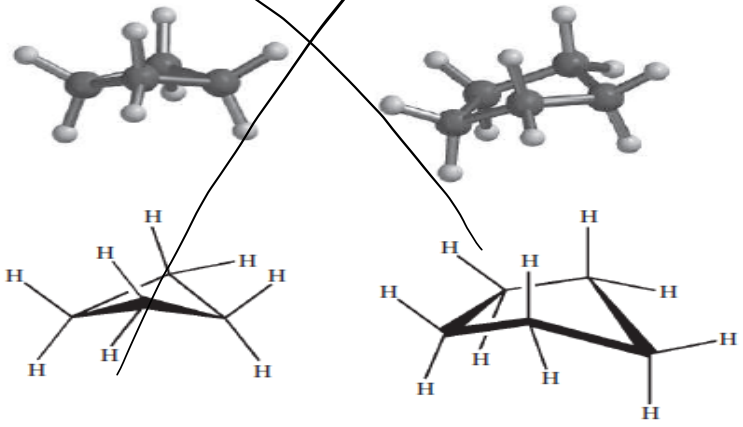
What are the conformations of cycloalkanes?

- Cyclopropane, with only three carbon atoms, is necessarily planar. The C-C-C angle is only 60° (the carbons form an equilateral triangle), much less than the usual sp^3 tetrahedral angle of 109.5°. The hydrogens lie above and below the carbon plane, and hydrogens on adjacent carbons are eclipsed.



cyclopropane

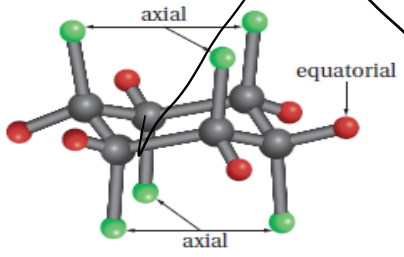
- Cycloalkanes with more than three carbon atoms are nonplanar and have “puckered” conformations. In cyclobutane and cyclopentane, puckering allows the molecule to adopt the most stable conformation (with the least strain energy). Puckering introduces strain by making the C-C-C angles a little smaller than they would be if the molecules were planar; however, less eclipsing of the adjacent hydrogens compensates for this.



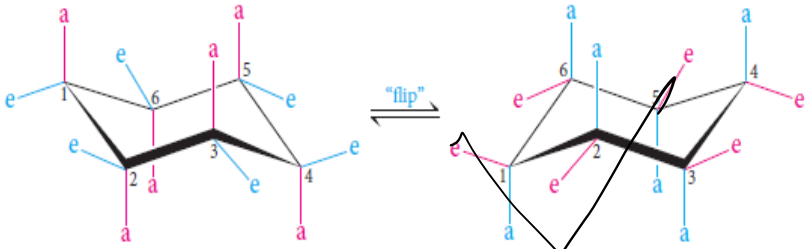
C—C—C angle	cyclobutane	cyclopentane
for planar molecule	90°	108°
observed experimentally	88°	105°

- Six-membered rings are more common and stable. If cyclohexane were planar, the internal C -C-C angles would be those of a regular hexagon, 120°—quite a bit larger than the normal tetrahedral angle (109.5°). The resulting strain prevents cyclohexane from being planar (flat). Its most favored conformation is the **chair conformation**, an arrangement in which all of the C-C-C angles are 109.5° and all of the hydrogens on adjacent carbon atoms are perfectly staggered.

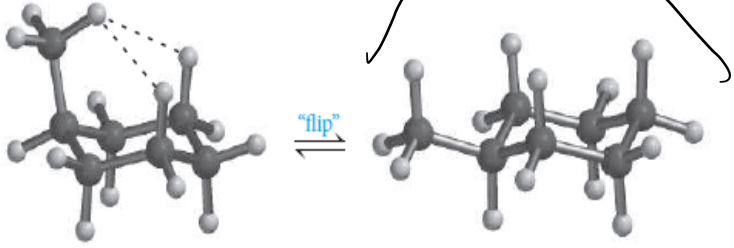
- In the **chair conformation** of cyclohexane, the six **axial** hydrogen atoms lie above and below the mean plane of the ring, while the six **equatorial** hydrogens lie in the plane.



ball-and-stick model



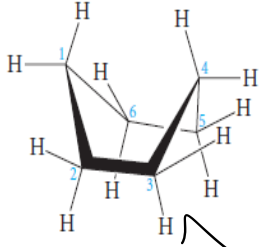
Axial bonds (red) in the left structure become equatorial bonds (red) in the right structure when the ring "flips."



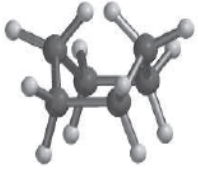
methyl axial
5%

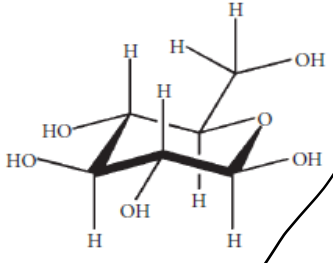
methyl equatorial
95%

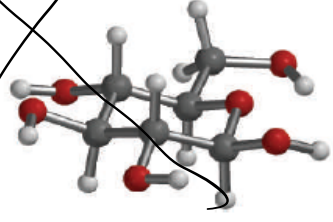
Another puckered conformation for cyclohexane, one in which all C-C-C angles are the normal 109.5°, is the boat conformation.



boat cyclohexane



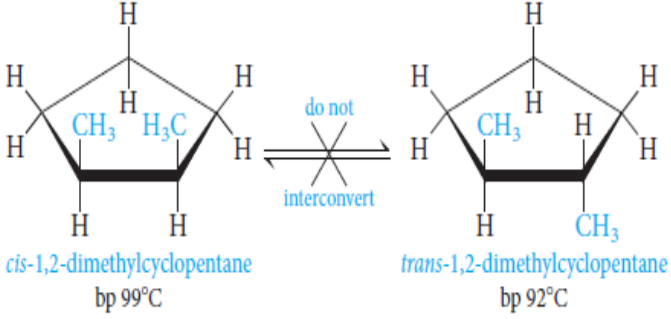




glucose (β-D-glucopyranose)

2.10 *Cis-Trans* Isomerism in Cycloalkanes

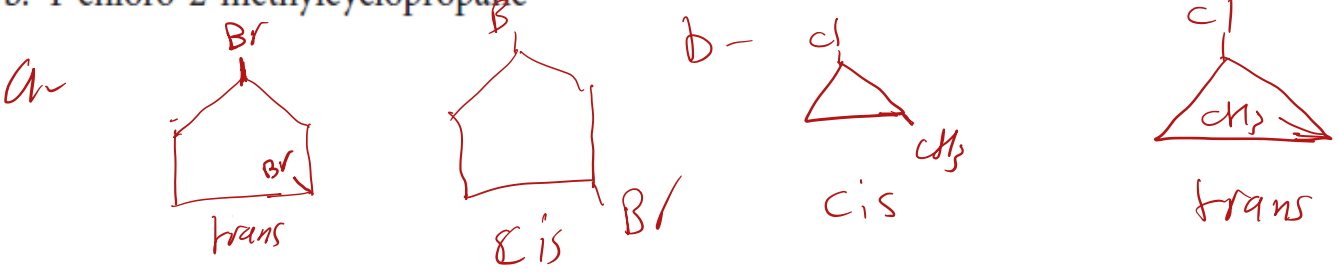
- **Stereoisomerism** deals with molecules that have the same order of attachment of the atoms, but different arrangements of the atoms in space. ***Cis–trans* isomerism** (sometimes called **geometrical isomerism**) is one kind of stereoisomerism, and it is most easily understood with a specific case.



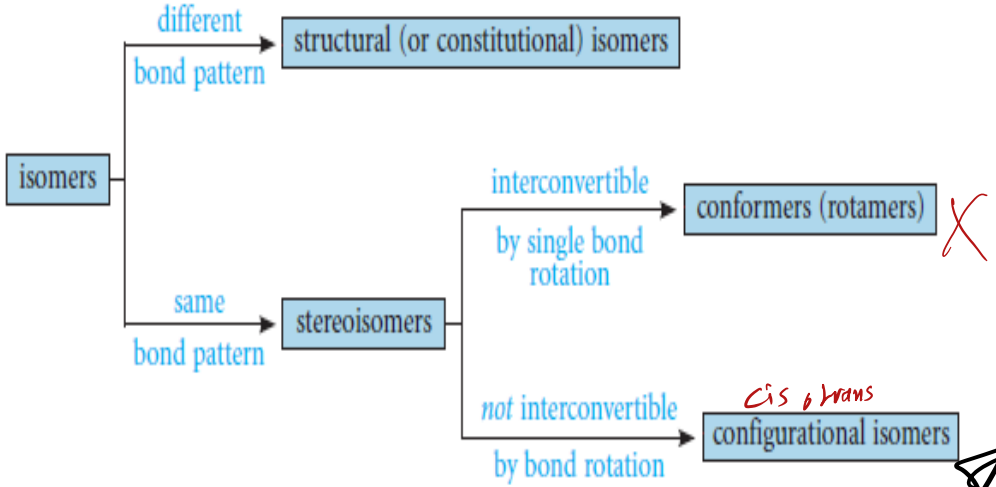
- *Cis*: like groups on same side of ring
- *Trans*: like groups on opposite sides of ring

PROBLEM 2.16 Draw the structure for the *cis* and *trans* isomers of

a. 1,3-dibromocyclopentane
b. 1-chloro-2-methylcyclopropane



2.11 Summery of Isomerism



PROBLEM 2.17 Classify each of the following isomer pairs according to the scheme

- a. *cis*- and *trans*-1,2-dimethylcyclohexane
- b. chair and boat forms of cyclohexane conformational
- c. 1-fluoropropane and 2-fluoropropane structural

2.12 Reactions of Alkanes

- All of the bonds in alkanes are single, covalent, and nonpolar. Hence alkanes are relatively inert.
- Because of this inertness, alkanes can be used as solventsfor extraction or crystallization as well as for carrying out chemical reactions of other substances.

1) Oxidation and Combustion: alkanes as fuels

$$CH_4 + 2\,O_2 \longrightarrow CO_2 + 2\,H_2O + \text{heat}\ (212.8\ \text{kcal/mol})$$

methane

$$C_4H_{10} + \tfrac{13}{2}O_2 \longrightarrow 4\,CO_2 + 5\,H_2O + \text{heat}\ (688.0\ \text{kcal/mol})$$

butane

Partial oxidation of hydrocarbon

$$2\,CH_4 + 3\,O_2 \longrightarrow \underset{\text{carbon monoxide}}{2\,CO} + 4\,H_2O$$

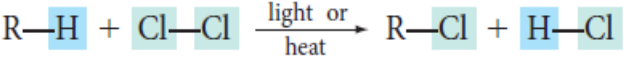
$$CH_4 + O_2 \longrightarrow \underset{\text{carbon}}{C} + 2\,H_2O$$

$$CH_4 + O_2 \longrightarrow \underset{\text{formaldehyde}}{CH_2O} + H_2O$$

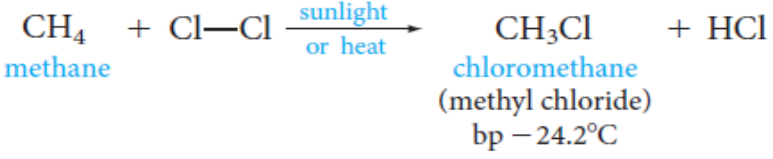
$$2\,C_2H_6 + 3\,O_2 \longrightarrow \underset{\text{acetic acid}}{2\,CH_3CO_2H} + 2\,H_2O$$

2.12 b Halogenation of alkanes.

- General formula

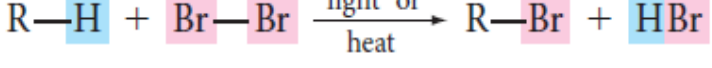


For methane

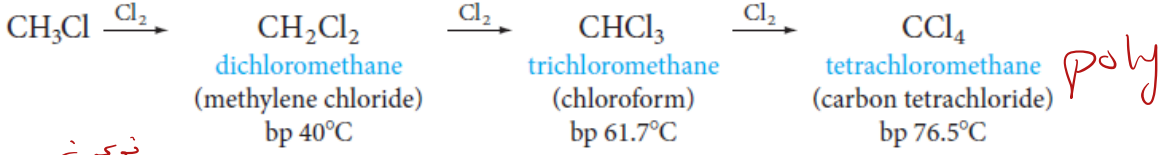


- The reaction is called **chlorination**. This process is a **substitution reaction**, as a chlorine is substituted for a hydrogen . An analogous reaction, called **bromination**, occurs when the halogen source is bromine.

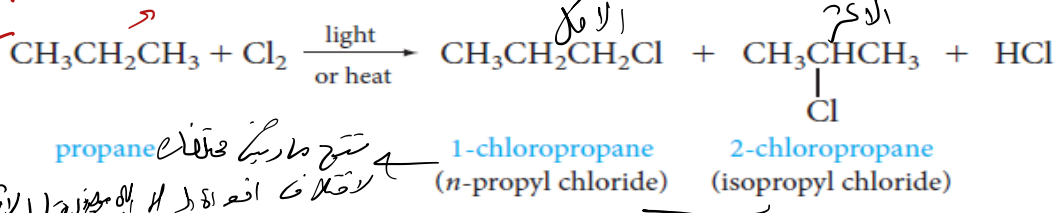
monohalo genation

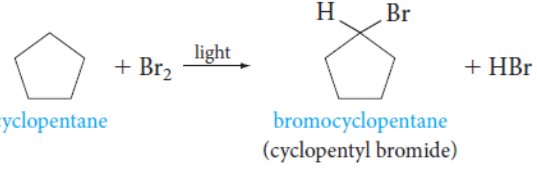


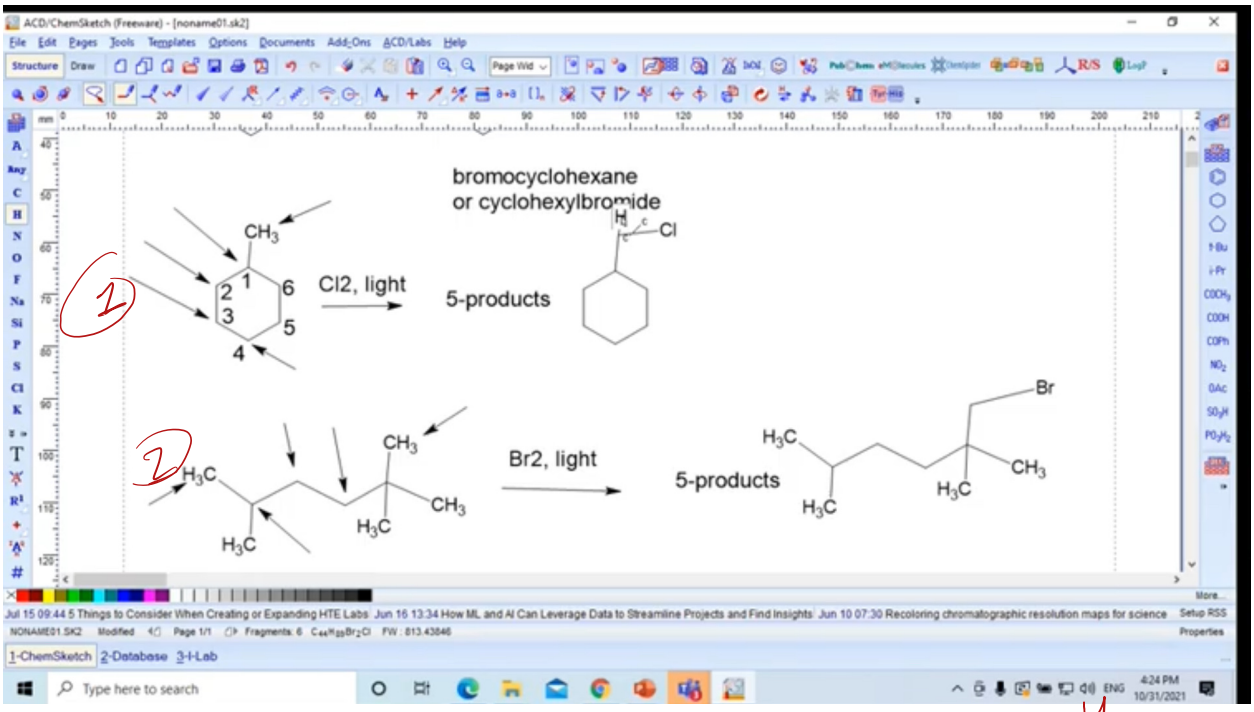
- If excess halogen is present, the reaction can continue further to give polyhalogenated products. Thus, methane and excess chlorine can give products with two, three, or four chlorines



poly

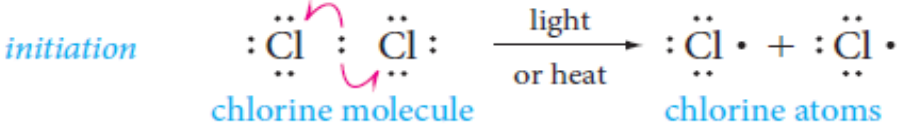




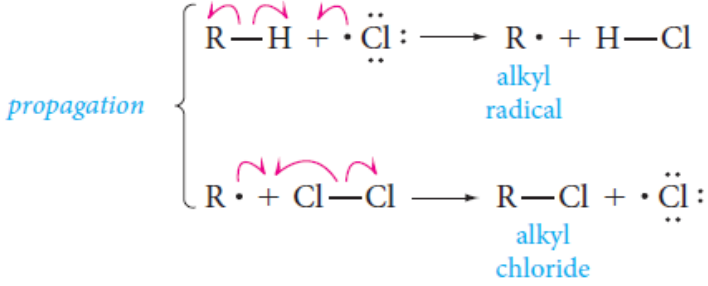

ACD/ChemSketch (Freeware) - [noname01.sk2]
bromocyclohexane
or cyclohexylbromide
Cl2, light
5-products
Br2, light
5-products
1-ChemSketch
2-Database
3-I-Lab

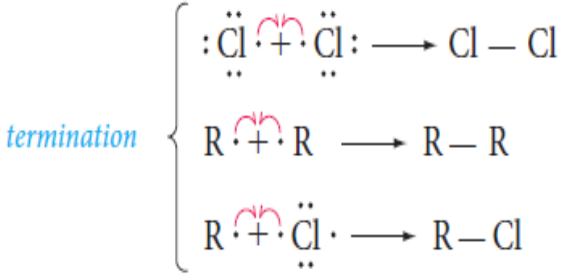
2.13 Free Radical Chain Mechanism of Halogenation

- A **reaction mechanism** is a step-by-step description of the bond-breaking and bond-making processes that occur when reagents react to form products. In the case of halogenation, various experiments show that this reaction occurs in several steps , and not in one magical step. Indeed, halogenation occurs via a **free-radical chain** of reactions.
- The **chain-initiating step** is the breaking of the halogen molecule into two halogen atoms.

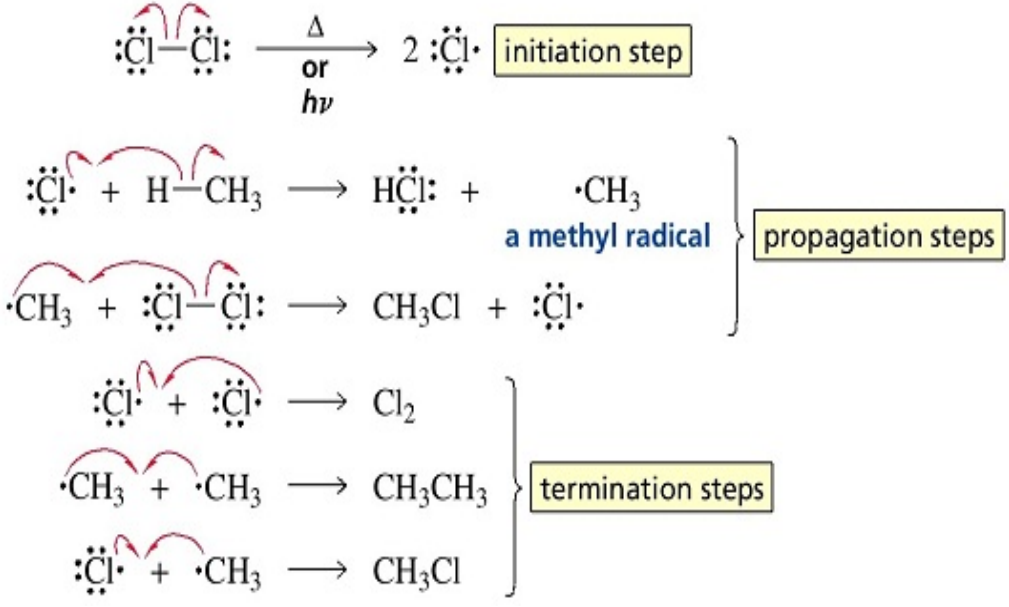


The **chain-propagating steps** are





mechanism for the monochlorination of methane



- Homework 1

26 28 31 35 41 44 45 47 48 50