CHEM 261 October 13, 2022

RECALL:

Identification of Chiral and Achiral (not chiral) compounds

Example: Diaminopimelic acid

- The above molecule is achiral even though there are stereogenic center (s), because there is symmetry within the molecule

- These kinds of molecules are called **meso compounds**, which are compounds that contain stereocenters yet because of their symmetry, have mirror images that can be superimposed.

- All achiral molecules, including meso compounds do not rotate polarized light (i.e. $[\alpha]_D = 0$)

- Diaminopimelic acid - a component of bacterial cell wall and biosynthetic precursor to the amino acid known as lysine

- this R,S diaminopimelic acid (above) is a diastereomer of the enantiomers (S,S or R,R diaminopimelic acid) below:

A racemic mixture (racemate) of two enantiomers in a 1:1 ratio also has an $[\alpha]_D = 0$

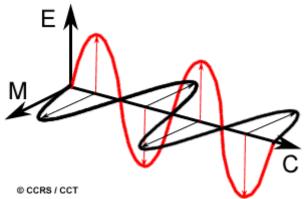
Physical Properties of Enantiomers

- Same physical properties with achiral agents or procedures
 - o Melting point, boiling point, solubility in achiral solvents
- Separation of enantiomers (resolution) require a chiral agent
- Diastereomers have different physical properties (m.p, b.p, density, solubility)

Optical Activity/Rotation

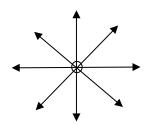
- Rotation of plane of polarized light
- α = measured rotation (°)
- $[\alpha]_D$ = Absolute rotation
 - o (+) (clockwise)
 - o (-) (counter-clockwise)
- Absolute rotation is 0 ° for achiral molecules

Pure enantiomers show equal but opposite rotation

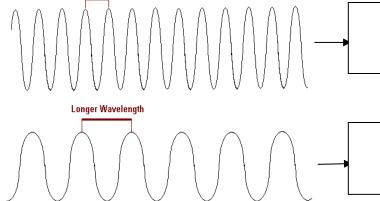


Light has an oscillating electric field (red) intersecting with a magnetic field (black)

Shorter Wavelength



End on view of vector components of normal light. It is possible to polarize light and remove some of those vectors.



- Shorter Wavelength
- Higher Energy
- Higher Frequency
- ➤ Longer Wavelength
- ➤ Low Energy
 - Low Frequency

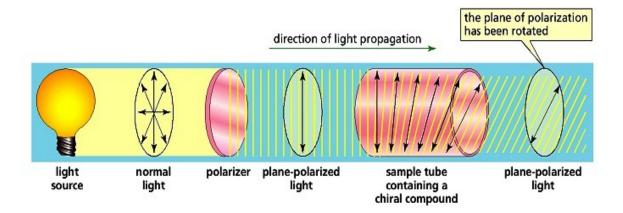
$$E = h\nu = \frac{hc}{\lambda}$$

E = energy

H = Planck's Constant (6.6 x 10⁻³⁴ Joules•sec

v = frequency

 λ = wavelength



Optical Rotation

Factors affecting optical rotation:

- Concentration of compound, g/cm³ (c)
- Path length that light travels through the solution, cm (1)

 $[\alpha]_D$ = Absolute rotation at the D line of sodium (589 nm or 5890 Å)

$$\left[\alpha\right]_D = \frac{\alpha}{c \cdot l}$$

 α = measured rotation (°) c = concentration (g/cm³) l = path length (cm)

D = D-line of sodium light $[\alpha]$ = absolute rotation

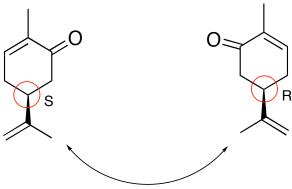
Degrees (°) = +: Clockwise (dextrorotatory/D)

- : Anticlockwise (levorotatory/L)

Note: not correlated with R/S configuration

Example: Carvone

$$[\alpha]_D = -60^{\circ}$$
 [$\alpha]_D = +60^{\circ}$ (made up values, not real rotation)



Enantiomers

S-carvone (caraway) R-carvone (spearmint)

Note: enantiomers will always have equal but opposite rotation, as such, they would cancel the rotation of the other when present as mixtures in solution.

Calculating Optical Purity

For this example for S-carvone, assume the pure S enantiomer has -60° rotation

<u>R</u>	<u>S</u>	Rotation (°)	Optical Purity (%)
100 %	0 %	+ 60 °	100 %
75 %	25 %	+ 30 °	50 %
50 %	50 %	0 о	0 %
25 %	75 %	−30 °	50 %
0 %	100	-60 °	100 %

If a solution is an equivalent mix of 1:1 R + S enantiomers, α measured = 0 ° A 50:50 mixture of enantiomers is called a *racemic mixture* (or racemate)

Optical Purity (measured experimentally) = Enantiomeric Excess (ee) (Theoretical value)

Resolution of Enantiomers

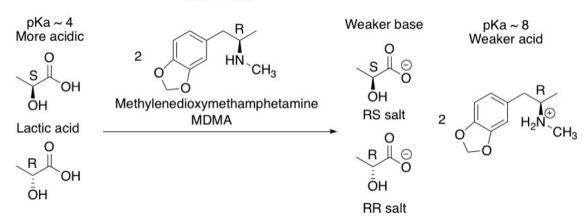
Definition: separation of two enantiomers

- Requires a chiral reagent to convert enantiomers to diastereomeric salts

Racemic mixture: 1:1 ratio of enantiomers in a mixture

Example: Lactic Acid

More basic



The starting material lactic acids are enantiomers of each other.

By reacting enantiomers to make a salt with an enantiomer of MDMA (another chiral molecule that is optically pure), also known as ecstasy, one can obtain salts which are now diastereomers of each other (RS and RR). The resulting diastereomers have different melting points, boiling points, solubilities, and can be separated by crystallization.

More examples of amines:

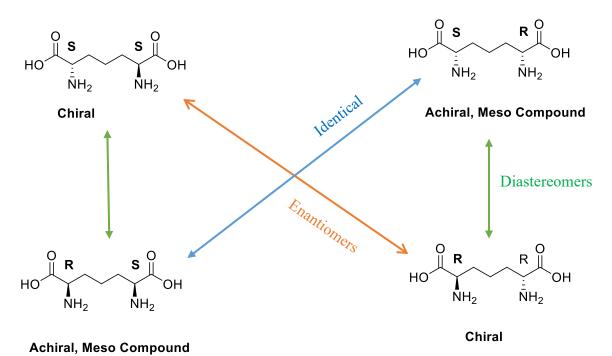
Amphetamine

Methamphetamine

$$NH_2$$

Cannot be used for resolution of enantiomers. Need a stereogenic centre

Review of concepts:



Substitution Reactions

Remember: Radical Substitution

Note: Although radicals are sp³ hybridized, rapid inversion around the central C results in a loss of stereochemistry. Hence, the resulting product would be a racemic mixture.

Ionic Substitution:

Nucleophile is a substance that seeks positive charge

Types of Nucleophilic Substitution (S_N)

S_N1 - rate depends on 1 concentration

 $S_{\rm N}2$ - The rate is dependent on the concentration of the nucleophile and the nucleophile (2 concentrations)

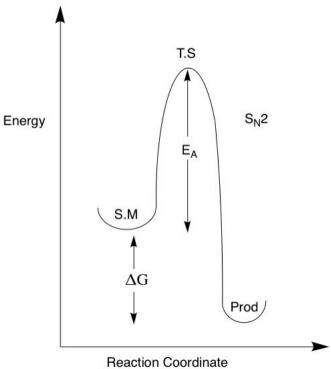
Sn2 Mechanism

Reverse reaction will not occur.

- Inversion of configuration (Walden inversion)
- **Concerted**: The bonds of the starting material break at the same time that the product bonds form.

- **Stereospecific**: stereochemistry of the starting material determines the stereochemistry of the product.
- Works for CH₃-X, R-CH₂X, R₁R₂CHX.

Energy Diagram of S_N2 Reaction



ricaction coordinate

There is no intermediate in the $S_{\rm N}2$ reaction, as it is concerted

Good Leaving Groups	Bad Leaving Groups
I ⁻ , Br ⁻ , Cl ⁻ , <<< F ⁻ (due to solvation)	H-, R- (alkyl), -NR ₂ , -OR, -OH
RSO ₃ -	

As such, the reaction below is not reversible:

More examples of reactions that does not proceed via $S_N 2$

$$\begin{array}{ccc}
& S_{N2} \\
& \times \\
& \times$$

Leaving group activation:

1) Protonation

OH NaCl No Rxn

NaCl
$$H_2SO_4$$
 H_2SO_4 H_2O

Na+Cl H_2O

Na+Cl H_2O

Highlights:

- H₂SO₄ is a proton (H⁺) donor.
- OH is transformed into a better leaving group via protonation of the O atom.
- Cl⁻ can then attack via S_N2, kicking off H₂O in the process.