

MODULE-2



Rapid Revision Notes



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PHARMACEUTICAL CHEMISTRY

- **MEDICINAL CHEMISTRY**
- **ORGANIC CHEMISTRY**
- **PHYSICAL CHEMISTRY**
- **INORGANIC CHEMISTRY**
- **BIOCHEMISTRY**

Features

- * According To Latest Syllabus
- * All Topics Covered In Concise Forms
- * Designed By Pharmacy India Experts
- * Important For All Pharma Competitive Exam
- * Subject Wise Previous Year Questions



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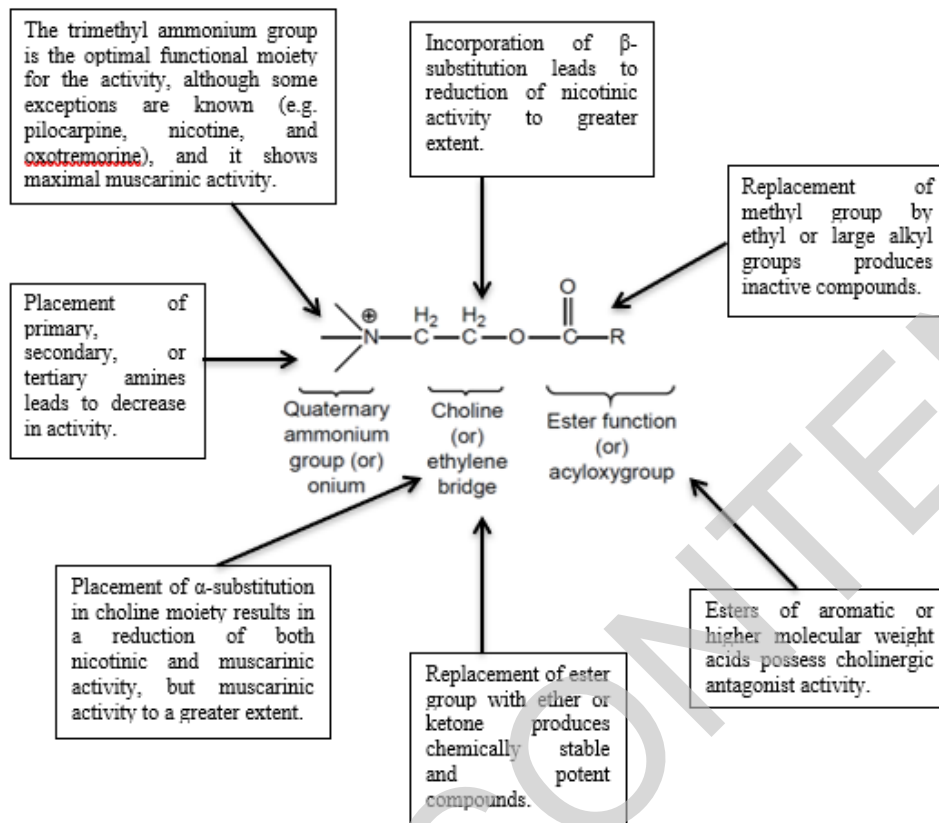
SECTION

1

Medicinal Chemistry

- ➔ **Drugs acting on autonomic nervous system (ANS)**
- ➔ **Drugs acting on central nervous system (CNS)**
- ➔ **Drugs acting on cardiovascular system (CVS)**
- ➔ **Drugs acting on inflammation/ allergy**
- ➔ **Drugs acting on endocrine system**
- ➔ **Drugs acting on respiratory system**
- ➔ **Drugs acting on urinary system**
- ➔ **Drugs acting on blood and blood forming organs**
- ➔ **Chemotherapeutic agents**

Structure activity relationship



Anticholinergic Drugs

Introduction

- The parasympatholytics or cholinergic blocking agents include atropine and related alkaloids obtained from plants, such as *Atropa belladonna*, *A. accuminata*, *Hyosyamus niger*, *Scopolia carniolica*, *Datura stramonium*, and synthetic or semisynthetic atropine substitutes.
- The antinicotinic drugs consist of ganglion blockers and neuromuscular junction blockers.
- Muscarinic antagonists are often called as parasympatholytics.

Classification

1. Solanaceous alkaloids and analogues

Drug name	Structure
Atropine	
Scopolamine	
Homatropine	

Introduction

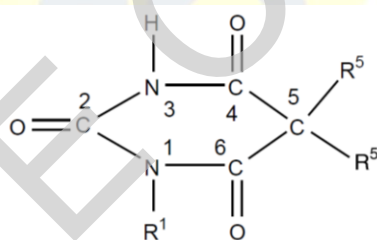
- These drugs can selectively relieve pain, reduce fever, suppress disordered movement, induce sleep or arousal, and reduce the desire to eat or ally the tendency to vomit.

Sedative and Hypnotics**Introduction**

- Sedatives are central nervous system (CNS) depressant drugs that reduce excitement, tension, and produce relaxation.
- Hypnotics are drugs that depress the CNS and produce sleep similar to that of natural sleep.
- Both sedative and hypnotic action may reside in the same drug.

Classification

1. **Barbiturates** → Barbiturates are further classified on the basis of duration of their action.

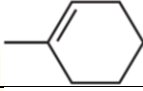
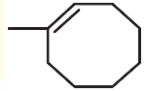


- a. **Long-acting barbiturates (6 h or more than 6 h)**

Drug name	R ¹	R ⁵	R ^{5'}
Barbital	-H	-C ₂ H ₅	-C ₂ H ₅
Phenobarbital	-H	-C ₂ H ₅	-C ₆ H ₅
Mephobarbital	-CH ₃	-C ₂ H ₅	-C ₆ H ₅
Metharbital	-CH ₃	-C ₂ H ₅	-C ₂ H ₅

- b. **Intermediate-acting barbiturates (3–6 h)**

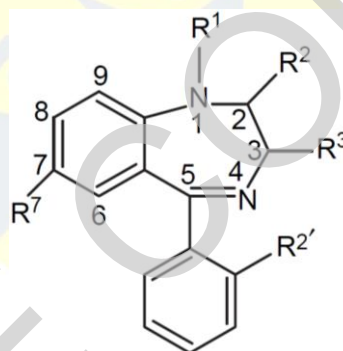
Drug name	R ¹	R ⁵	R ^{5'}
Amobarbital	-H	-C ₂ H ₅	$\text{—CH}_2\text{CH}_2\text{CH} \begin{cases} \text{CH}_3 \\ \text{CH}_3 \end{cases}$
Butobarbital	-H	-C ₂ H ₅	$\begin{array}{c} \text{—CH—CH}_2\text{—CH}_3 \\ \\ \text{CH}_3 \end{array}$
Aprobarbital	-H	CH ₂ =CH—CH ₂ —	(CH ₃) ₂ CH—
Talbutal	-H	CH ₂ =CH—CH ₂ —	CH ₃ CH ₂ CH(CH ₃)—
Butalbital	-H	CH ₂ =CH—CH ₂ —	(CH ₃) ₂ CHCH ₂ —
Hexobarbital	-CH ₃	-CH ₃	

Pentobarbital	-H	-C ₂ H ₅	$\begin{array}{c} \text{CH}_3\text{CH}_2\text{CH}_2\text{CH} - \\ \\ \text{CH}_3 \end{array}$
Secobarbital	-H	CH ₂ =CH-CH ₂ -	$\begin{array}{c} \text{CH}_3(\text{CH}_2)_2\text{CH} - \\ \\ \text{CH}_3 \end{array}$
Cyclobarbital	-H	-C ₂ H ₅	
Heptabarbital	-H	-C ₂ H ₅	

c. Short-acting barbiturates (less than 3 h)

Drug name	R ¹	R ⁵	R ^{5'}
Thiopentone	-H	-C ₂ H ₅	$\begin{array}{c} \text{CH}_3(\text{CH}_2)_2\text{CH} - \\ \\ \text{CH}_3 \end{array}$

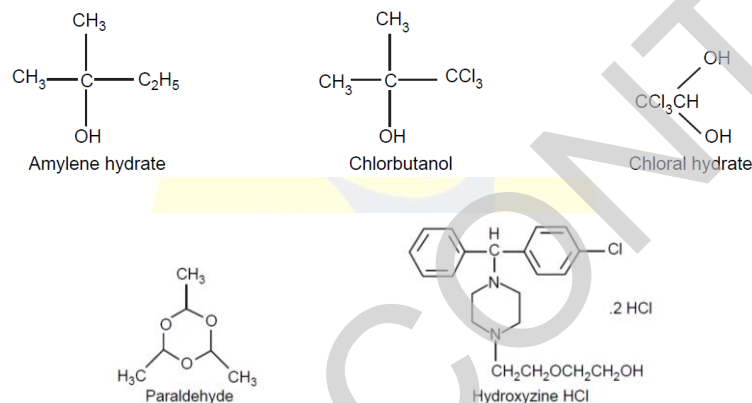
2. Benzodiazepines



Drug name	R ¹	R ²	R ³	R ⁷	R ^{2'}
Diazepam	-CH ₃	= O	-H	-Cl	-H
Oxazepam	-H	= O	-OH	-Cl	-H
Chlordesmethyl diazepam	-H	= O	-H	-Cl	-Cl
Fosazepam	$\begin{array}{c} \text{O} \\ \uparrow \\ \text{---}(\text{CH}_2)\text{P}(\text{CH}_3)_2 \end{array}$	= O	-H	-Cl	-H
Prazepam	$\text{---CH}_2 \text{---} \triangle$	= O	-H	-Cl	-H
Nitrazepam	-H	= O	-H	-NO ₂	-H
Nordiazepam	-H	= O	-H	-Cl	-H
Nimetazepam	-CH ₃	= O	-H	-NO ₂	-H
Flunitrazepam	-CH ₃	= O	-H	-NO ₂	-F
Flurazepam	-(CH ₂) ₂ N (C ₂ H ₅) ₂	= O	-H	-Cl	-F
Quazepam	-CH ₂ CF ₃	= S	-H	-Cl	-F
Halozepam	-CH ₂ CF ₃	= O	-H	-Cl	-H
Temazepam	-CH ₃	= O	-OH	-Cl	-H
Lorazepam	-H	= O	-OH	-Cl	-Cl
Clonazepam	-H	= O	-H	-NO ₂	-Cl
Doxefazepam	-CH ₂ OH	= O	-OH	-Cl	-F

<ul style="list-style-type: none"> Glutethemide 	
Quinazolines	
<ul style="list-style-type: none"> Methaqualone 	
<ul style="list-style-type: none"> Mecloqualone 	

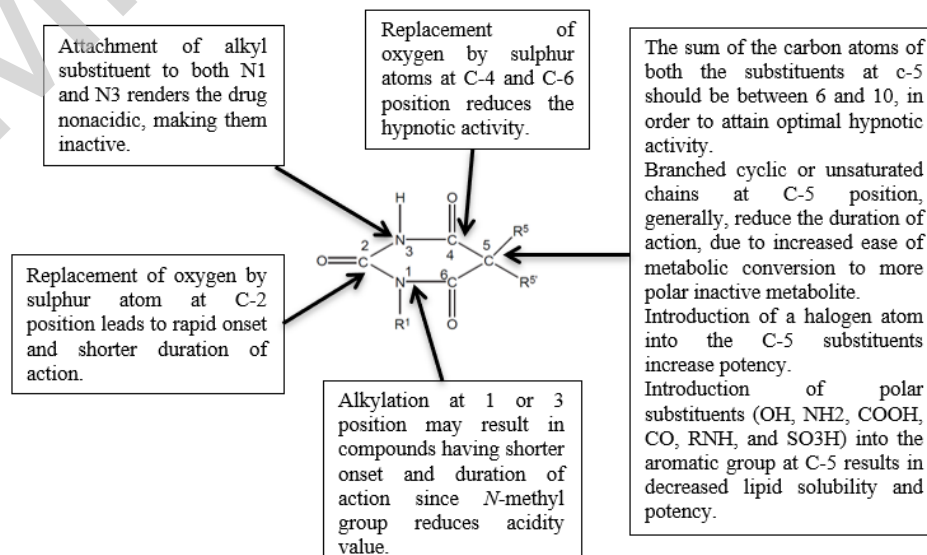
5. Alcohols and aldehydes



6. Miscellaneous

- Inorganic salts → KBr, magnesium sulphate
 - Acids and esters → Tryptophan, 5-hydroxy tryptophan, Etomidate
 - Antihistamines and anticholinergics → Doxylamine, Diphenhydramine, Pyrilamine
 - Sulphones → Toxic-not used
 - Plant extracts → *Radix valerianae*, *Rauwolfia serpentine*, *Avana sativa*, *Glandula lupuli*
 - Endogenous substances: peptides
7. Newer agents → Zaleplon, Zopiclone, Zolpidem

SAR of Barbiturates



Antidepressants

Introduction

- Antidepressants are drugs, which enhance alertness and may result in an increased output of behaviour.
- They are used for the relief of symptoms of moderate and severe depressive disorder.

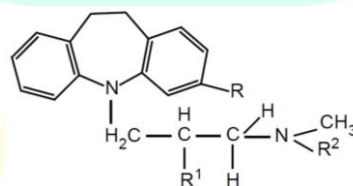
Classification

1. Monoamine oxidase inhibitors (MAOIs)

Drug name	Structure
Isocarboxazide	
Iproniazid	
Phenelzine	
Pheniprazine	
Tranlycypromine	
Pargyline HCl	
Clorgyline	
Deprenyl	

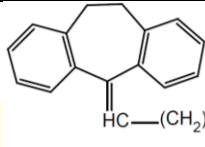
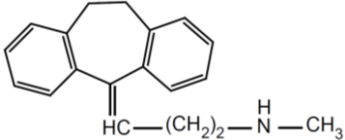
2. Tricyclic antidepressants (TCAs)

a. Imino dibenzyl derivatives

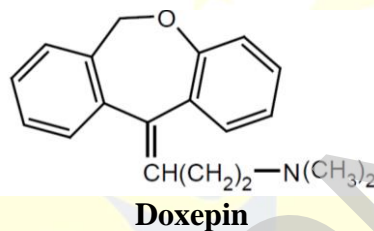


Drug name	R	R ¹	R ²
Imipramine	-H	-H	-CH ₃
Desipramine	-H	-H	-H
Trimipramine	-H	-CH ₃	-CH ₃
Chlorimipramine	-Cl	-H	-CH ₃

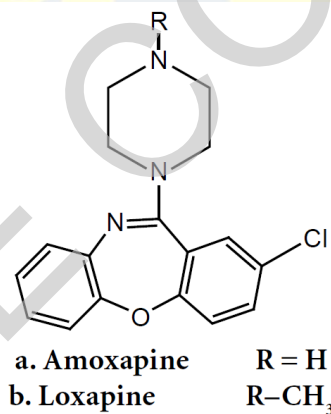
b. Dibenzo cycloheptane derivatives

Drug name	Structure
Amitriptyline	
Nortriptyline	

c. Dibenzoxepine derivatives



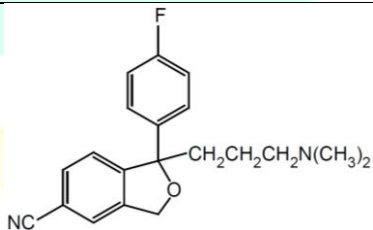
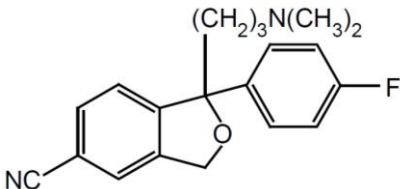
d. Dibenzoxazepine derivatives



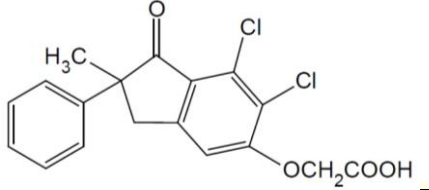
3. Second-generation antidepressants

- a. Bicyclic: Viloxamine, Zimeldine
- b. Tricyclic: Dibenzepine, Amoxapine, Imipramine
- c. Tetracyclic: Mianserin, Mianserin

4. Selective serotonin reuptake inhibitors (SSRIs)

Drug name	Structures
Citalopram(±) Isomer	
Escitalopram (+) Isomer	

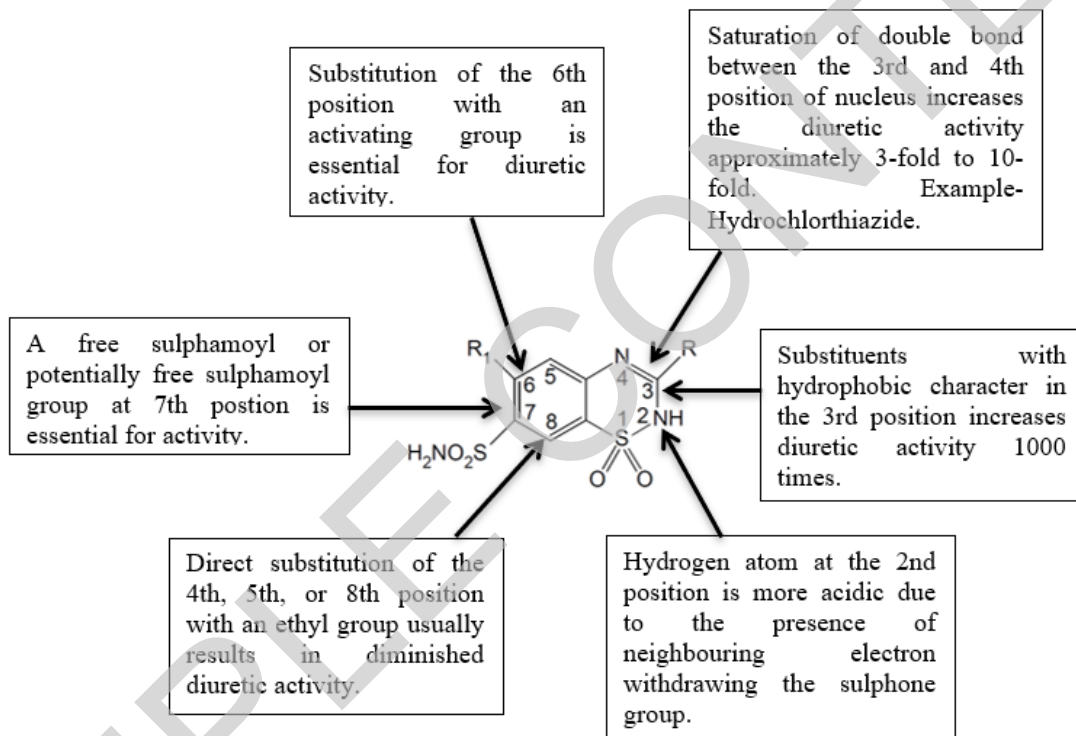
k. Uricosuric diuretics

Drug name	Structure
Indacrinone	

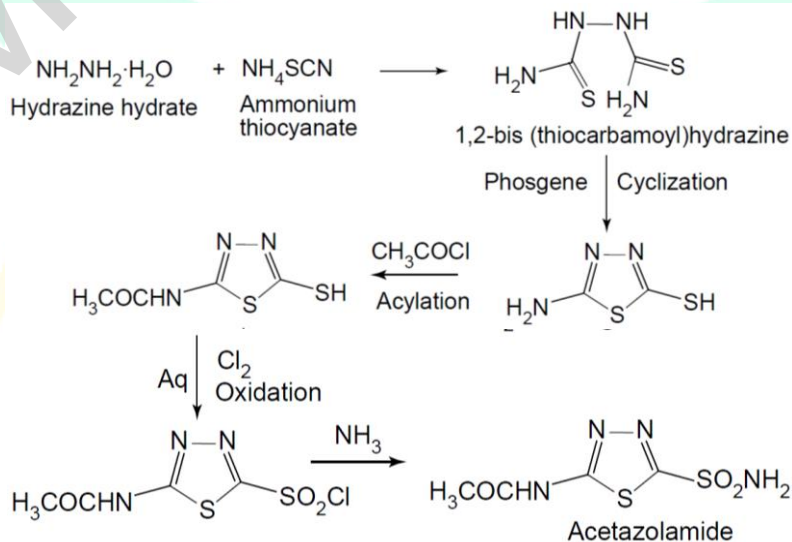
l. Acidic diuretics → Ammonium chloride

m. Miscellaneous → Muzolimine

SAR of Thiazide Diuretics



1. Acetazolamide [N-(Sulphonamido-1,3,4-thiadiazol-2-yl)acetamide]



SECTION

2

Organic Chemistry












- ➔ **Heterocyclic Compounds**
- ➔ **Aromaticity Compounds**
- ➔ **General Principles**
- ➔ **Important Name Reactions**
- ➔ **Isomerization**

Introduction












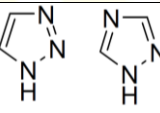
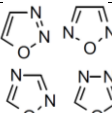
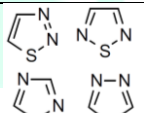

- Heterocyclic compounds are organic compounds that contain a ring structure containing atoms in addition to carbon, such as sulfur, oxygen or nitrogen, as the heteroatom.
- Carbocyclic compounds are organic compounds that contain ring system made up entirely of carbon atoms.

Classification


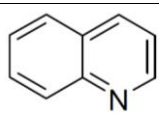
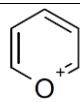
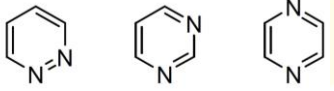
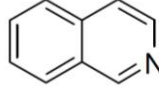
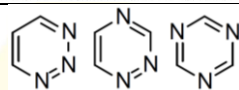
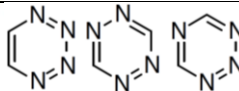
1. Three and four membered heterocycles

Ring size	Heteroatoms			
	N	O	S	Other
3	 Aziridine	 Oxirane	 Thiirane	
	 Diaziridine	 Dioxirane		 Oxaziridine
4	 Oxetane	 Thietane	 Thietane	 Selenetane
				 Phosphetane

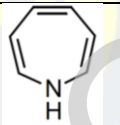
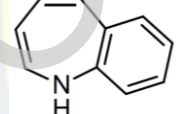
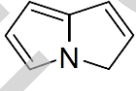
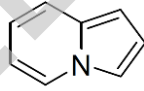
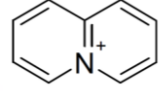
2. Five membered heterocycles

Ring size	Heteroatoms				
	N	Benzo	O	Benzo	S
5	 Pyrrole	 Indole	 Furan	 Benzofuran	 Thiophene
	 Pyrazole		 Isoxazole		 Isothiazole
5	 Imidazole		 Oxazole		 Thiazole
	 Triazoles		 Oxadiazoles		 Thiadiazoles
	 Tetrazole				

3. Six membered heterocycles

Ring size	Heteroatoms		
	N	Benzo	O
6	 Pyridine	 Quinoline	 Pyrilium
	 Diazines Pyridazine Pyrimidine Pyrazine	 Isoquinoline	
	 Triazines		
	 Tetrazines		

4. Other simple heterocycles

Ring size	Heteroatoms	
	N	Benzo
7	 Azepine	 Benzoazepine
5-5	 Pyrroline	
5-6	 Indolizine	
6-6	 Quinolizinium	

Nomenclature of Heterocyclic Compounds

1. Prefix for Hetero Atoms

Hetero atoms	Valence	Prefix
O	2	Oxa
N	3	Aza
S	2	Thia
Se	2	Selena
Te	2	Tellura
P	3	Phospha
As	3	Arsa
Si	4	Sila
Ge	4	Germa

Hybridization of central atom	% of s, p, and d characters	No. of hybrid orbitals	Shape and bond angle of hybrid orbital	Examples
sp	50% s and p character	2	Linear/ 180°	BeF ₂ , BeH ₂ , BeCl ₂ , C ₂ H ₂
sp ²	33.33% s and 66.66% p character	3	Trigonal planar/ 120°	BeF ₃ , BH ₃ , C ₂ H ₄
sp ³	25% s and 75% p character	4	Tetrahedron/ 109.28°	C ₂ H ₆ , CH ₄
sp ³ d	-	5	trigonal bipyramidal / 120° and 90°	PCl ₅ , PF ₅
sp ³ d ²	-	6	Octahedron/ 90°	SF ₆

Relative size of sp, sp² and sp³ -orbitals

- s-orbital, character, in the three hybrid orbitals varies in the following manner:
 $sp > sp^2 > sp^3$
- Since, s-orbitals are closer to the nucleus than p-orbitals, it is, thus, expected that **greater the s-character of the hybrid orbital the smaller is its size**. Therefore, the order of the size of the three hybrid orbitals is:

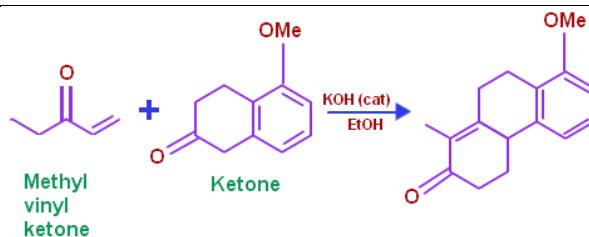
$$sp^3 > sp^2 > sp$$

- On the basis of the sizes, sp-orbital forms shortest and sp³-orbital longest bonds with other atoms.

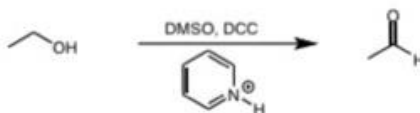
Bond Length

- The average distance between the centres of the nuclei of the two bonded atoms in a molecule is called the **bond length**.
- It also depends on the type of hybridization and the resonance.
- The bond length is:
Single bond > Double bond > Triple bond
- The bond length of the sigma bond in, different types of hybridization follows the following order:
σ-bond (sp³) > σ-bond (sp²) > σ-bond (sp)
- The bond length is expressed in angstrom units (Å) or picometre (pm), (1Å 10⁻¹⁰ m, 1 pm 10⁻¹² m).
- The bond lengths are measured by X-ray crystallography and by microwave spectroscopy.

Bond	Bond length (Å)
C-C	1.54
C=C	1.34
C≡C	1.20
C-H(sp ³ -s)	1.112
C-H(sp ² -s)	1.103
C-H(sp-s)	1.08
C-O	1.40



60. Pfitzner –Moffatt oxidation → It is a chemical reaction for the oxidation of primary and secondary alcohols to aldehydes and ketones, respectively. The oxidant is a combination of dimethyl sulfoxide (DMSO) and dicyclohexylcarbodiimide (DCC).

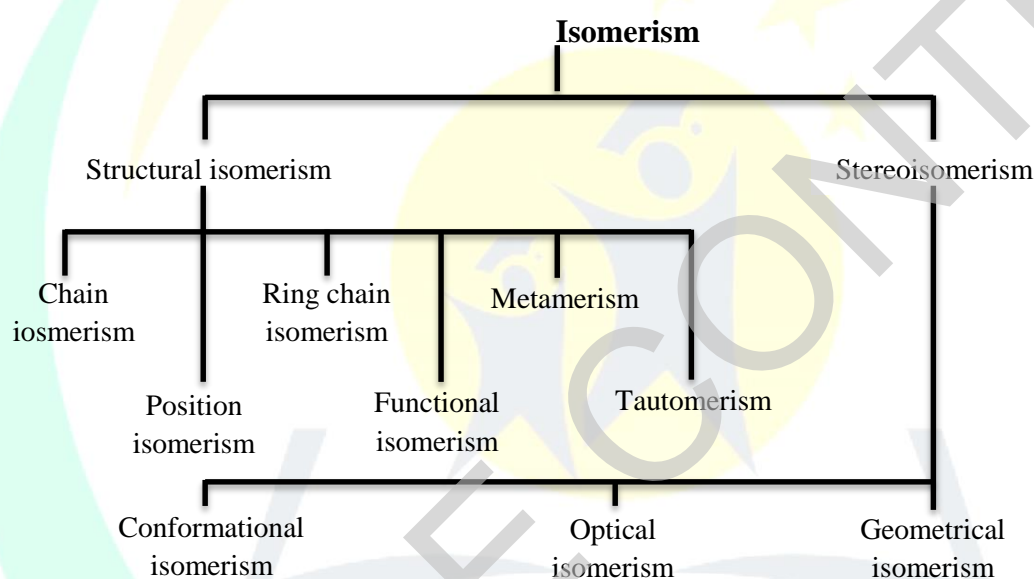


NAME REACTIONS

Name reactions	Starting materials	Catalyst	End product
Wolff Kishner reduction/ Hung-milnon reaction	Aldehydes and ketones	H_2NNH_2 ; $\text{C}_2\text{H}_5\text{ONa}$; Glycol	Methylene group
Meerweili-Ponndorf-Verley (MPV) reduction	Ketones	$[(\text{CH}_3)_2\text{CHO}]_3\text{Al}$; $(\text{CH}_3)_2\text{CHOH}$	2° alcohol
Pinacol-pinacolone rearrangement	Ketones	Mg/ Hg; mineral acid	2,3-dimethyl butane-2,3-diol(Pinacol); 3,3-dimethyl butan-2-one (Pinacolone)
Beckmann rearrangement	Ketoximes	conc. H_2SO_4 , PCl_5 , H_3PO_4 , SOCl_2 or $\text{C}_6\text{H}_5\text{SO}_2\text{Cl}$	N-methyl acetamide
Aldol condensation	carbonyl compounds	NaOH , $\text{Ba}(\text{OH})_2$ or K_2CO_3	Aldol
Baeyer Villiger oxidation	Aliphatic ketones	(per monosulphuric acid, H_2SO_4) or per benzoic acid ($\text{C}_6\text{H}_5\text{CO}_3\text{H}$) or m-chloro perbenzoic acid or per acetic acid ($\text{CH}_3\text{CO}_3\text{H}$) or $\text{CF}_3\text{CO}_3\text{H}$	Esters
Cannizzaro's reactions	Aldehydes (with no α -hydrogen)	Conc. Alkali	Carboxylic acids
Tischenko's reaction	Aldehydes (with or without α -hydrogen)	$(\text{C}_2\text{H}_5\text{O})_3\text{Al}$	Ester
Claisen-Schmidt reaction	aliphatic aldehyde or ketone (with α -hydrogen)	Dil. NaOH	α,β -unsaturated compound
Schmidt reaction	Carbonyl compounds	conc. H_2SO_4	Alkyl cyanide and N-alkyl formamide
Lederer Manase's reaction	Phenol	dilute acid or alkali	o- and p-hydroxy benzyl alcohol

Introduction

- Two or more than two organic compounds having the **same molecular formula and molecular weight but different physical and chemical properties** are called **isomers** and the phenomenon is called **isomerism**.



Structural isomerism:

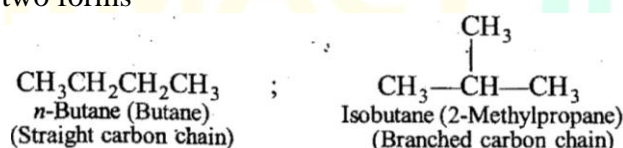
- It is due to the difference in the manner in which the constituent atoms or groups are linked to one another within the molecule, without any reference to space.
- Structural isomers are compounds having same molecular 'formula' but different structural formulae.

1. Chain or nuclear isomerism

- This type of isomerism is due to difference in the arrangement of carbon atoms constituting the chain, *i. e.*, straight or branched chain of carbon atoms. It is also known as, nuclear or skeletal isomerism. The isomers showing chain isomerism belong to same homologous series.

- Examples-**

C_4H_{10} (butane) exists in two forms-



Pentane (C_5H_{12}) exist in three isomers-

SECTION

3

**Physical
Chemistry**

→ **Complete Topic**

PHYSICAL CHEMISTRY

STATES OF MATTER

PARAMETER	SOLID	LIQUID	GAS
Shape	Definite shape	Indefinite shape	Indefinite shape
Volume	Definite Volume	Indefinite Volume	Indefinite Volume
Inter particular Forces	Strong Inter particular Forces	Comparatively weaker Inter particular Forces	Inter particular forces are negligible
Inter particular Space	Negligible inter particular space	Comparatively large inter particular space	Very large Inter particular space
Packing of Particles	Particles are very Closely packed	Particles are loosely packed	Particles are loosely packed
Density	Very High Density	Low Density	Very low density

INTRAMOLECULAR FORCES OF ATTRACTION

Type of Bond	Definition/ Description
Ionic bond	This bond is formed by the complete transfer of valence electron(s) between atoms.
Covalent bond	This bond is formed between atoms that have similar electronegativities.
Nonpolar covalent bond	A nonpolar covalent bond is formed between same atoms or atoms with very similar electronegativities.
Polar covalent bond	A polar covalent bond is formed when atoms of slightly different electronegativities share electrons.
Metallic bonding	This type of covalent bonding specifically occurs between atoms of metals, in which the valence electrons are free to move through the lattice

INTERMOLECULAR FORCES OF ATTRACTION

Force/ Interaction	Description
Dipole-dipole interactions	These forces occur when the partially positively charged part of a molecule interacts with the partially negatively charged part of the neighboring molecule.
Ion-Dipole Interactions	These interactions are similar to dipole-dipole interactions except for the fact that they arise between ions and polar molecules.
Ion Induced Dipole Interactions	In this type of interaction, a non-polar molecule is polarized by an ion placed near it. The non-polar molecules, upon obtaining a charge, behave as induced dipoles
Hydrogen bonding	This is a special kind of dipole-dipole interaction that occurs specifically between a hydrogen atom bonded to either an oxygen, nitrogen, or fluorine atom.
London dispersion forces	These are the weakest of the intermolecular forces and exist between all types of molecules, whether ionic or covalent—polar or nonpolar.

Isobaric process	In this process the pressure remains constant throughout the change i.e., $dP = 0$.
Isochoric process	In this process volume remains constant throughout the change, i.e., $dV = 0$.
Cyclic process	When a system undergoes a number of different processes and finally return to its initial state, it is termed cyclic process. For a cyclic process $dE = 0$ and $dH = 0$.
Reversible process	A process which occurs infinitesimally slowly, i.e. opposing force is infinitesimally smaller than driving force and when infinitesimal increase in the opposing force can reverse the process, it is said to be reversible process.
Irreversible process	When the process occurs from initial to final state in single step in finite time and cannot be reversed, it is termed an irreversible process.

LAWS OF THERMODYNAMICS

Law	Description
Zero law of thermodynamics	"If a system A is in thermal equilibrium with a system C and if B is also in thermal equilibrium with system C, then A and B are in thermal equilibrium with each other whatever the composition of the system."
First law of thermodynamics	"Energy can neither be created nor destroyed although it can be converted from one form into another." $E_2 - E_1 = \Delta E = q + w$ (Change in internal energy) = (Heat added to the system) + (Work done on the system)
Second law of thermodynamics	Kelvin statement: "It is impossible to derive a continuous supply of work by cooling a body to a temperature lower than that of the coldest of its surroundings." Clausius statement: "It is impossible for a self-acting machine, unaided by any external agency, to transfer heat from one body to another at a higher temperature or Heat cannot itself pass from a colder body to a hotter body, but tends invariably towards a lower thermal level."
Third law of thermodynamics	"The entropy of all perfectly crystalline solids is zero at the absolute zero temperature. Since entropy is a measure of disorder, it can be interpreted that at absolute zero, a perfectly crystalline solid has a perfect order of its constituent particles."

TERMS USED IN THERMODYNAMICS

Terms	Description
Enthalpy	Heat content of a system at constant pressure is called enthalpy denoted by 'H'.
Enthalpy change	Heat change at constant pressure can be given as $\Delta q = \Delta E + P\Delta V$ Difference between ΔH and ΔE is significant when gases are involved in chemical reaction.

INORGANIC CHEMISTRY

INDIAN PHARMACOPOEIA

EDITION	YEAR & VOLUME	CHAIRMANSHIP	SUPPLEMENT	ADDENDUM
1 st	1955	Dr. B. N. Ghosh	1960	-
2 nd	1966	Dr. B. Mukherjee	1975	-
3 rd	1985 (2 volumes)	Dr. Nityanand		1989 & 1991
4 th	1996 (2 volumes)	Dr. Nityanand		31 st December 2000 & 30th June 2003
5 th	2007 (3 volume)	Dr. Nityanand		2008
6 th	2010 (3 volume)	Shir K. Chandramouli	2012	-
7 th	2014 (4 volume)	Nabi Azad (Health Minister)	2015	2016
8 th	2018 (4 volume)	Dr. C.K. Mishra	2019	2021

LIMIT TEST

TEST	DESCRIPTION
Chloride	Limit test of chloride is based on the reaction of soluble chloride with silver nitrate in presence of dilute nitric acid to form silver chloride, which appears as solid particles (Opalescence) in the solution.
Sulphate	Limit test of sulphate is based on the reaction of soluble sulphate with barium chloride in presence of dilute hydrochloric acid to form barium sulphate which appears as solid particles (turbidity) in the solution.
Iron	Limit test of Iron is based on the reaction of iron in ammonical solution with thioglycollic acid in presence of citric acid to form iron thioglycolate which is pale pink to deep reddish purple in color.
Lead	Limit test of lead is based on the reaction of lead and diphenyl thiocarbazon (dithizone) in alkaline solution to form lead dithizone complex which is read in color. Dithizone is green in color in chloroform and lead-dithizone complex is violet in color, so the resulting color at the end of process is red.
Arsenic	Limit test of Arsenic is based on the reaction of arsenic gas with hydrogen ion to form yellow stain on mercuric chloride paper in presence of reducing agents like potassium iodide. It is also called as Gutzeit test and requires Gutzeit apparatus .
Heavy Metals	Limit test of heavy metals is based on the reaction of metallic impurities with hydrogen sulfide in acidic medium to form brownish colour solution.

ACID BASE & BUFFER

Acids	Bases
<ul style="list-style-type: none"> ○ Which converts blue litmus paper to red ○ Having the PH <7 ○ Sour taste ○ React with bases to form salts and water. Example :- Hydrochloric acid (HCl)	<ul style="list-style-type: none"> ○ Which converts red litmus paper to blue ○ Having the PH >7 ○ Bitter taste ○ React with Acids to form salts and water. Example: Sodium Hydroxide

THEORY OF ACID AND BASE

Theory	Acids	Base
Arrhenius theory	An Acid is a substance that can release hydrogen ion (H ⁺) when dissolved in water. Example: Hydrochloric acid. HCl → H⁺ + Cl⁻	A Base is a substance that can release a Hydroxyl ion (OH ⁻) when dissolved in water. Example: Sodium Hydroxide. NaOH → Na⁺ + OH⁻
Bronsted Lowry theory	Acid is the substance which donate proton. Conjugate acid: species that forms after a base accepts an H ⁺ .	Base is the substance which accept proton. Conjugate base: species that remains after an acid donates its H ⁺ .
Lewis theory	Acid: Acid is the molecule or ion that accept the lone pair of electrons. Example: H ⁺ , NH ₄ ⁺ , Na ⁺ , Cu ⁺⁺ , Al ⁺⁺⁺	Base: Base is the molecule or ion that donate the lone pair of electrons. Example: OH ⁻ , Cl ⁻ , CN ⁻

BUFFERS

Acidic buffers	Basic buffers	Neutral buffers
It is combination of weak acid and its corresponding salt.	It is combination of weak base and its corresponding salt.	It is single substance showing properties of buffers.
Example: Mixture of acetic acid, sodium acetate	Example: Mixture of Ammonium hydroxide and ammonium chloride	Example : Ammonium acetate

SODIUM DEFICIENCY

Condition	Hyponatremia	Hypertnatremia
Reason	Low level of Na ⁺	High level of Na ⁺
Symptoms	Extreme urine loss, diarrhea, kidney damage, vomiting, excessive sweating	Dehydration. High sodium intact. Intense thirst, fatigue

CHLORINE DEFICIENCY

Condition	Hypochloremia	Hyperchloremia
Reason	Decrease calcium level in body	Increase calcium level in body
Symptoms	Metabolic acidosis, Vomiting, Lack of reabsorption	Excess loss of bicarbonate ions and dehydration Symptoms

			Iron metabolism
Vitamin B₁ (Thiamine)	Pyrimidine ring	Beri-beri	Carbohydrate metabolism
Vitamin B₂ (Riboflavin)	Dimethyl isoalloxazine ring	Glossitis, Cheilosis, Angular stomatitis	Flavin adenosine dinucleotide (FAD) and Flavin mononucleotide(FMN)
Vitamin B₃ (Niacin)	Pyridine	Pellagra	Nicotinamide adenine dinucleotide (NAD ⁺) and Nicotinamide adenine dinucleotide phosphate (NADP ⁺).
Vitamin B₅ (Pantothenic acid)	Alanine and D-pantoic acid	Gopalan's Burning Foot Syndrome (Paresthesia)	Coenzyme A or CoA (A for acetylation)
Vitamin B₆ (Pyrridoxine)	Pyridine	Neurological symptoms, hypochromic microcytic anaemia	transamination, decarboxylation, deamination, transsulfuration, condensation
Vitamin B₇ (Biotin)	Imidazole and thiophene rings	Anemia, Dermatitis, Loss of appetite	Serves as a carrier of CO ₂ in carboxylation reactions.
Vitamin B₉ (Folic acid)	Pteridine ring	Macrocytic anemia	Serves as an acceptor or donor of one carbon units
Vitamin B₁₂ (Cyanocobalamin)	Corrin ring	Pernicious anemia	Synthesis of methionine from homocysteine, Isomerization of methymalonyl CoA to rrccinyl CoA

CLASSIFICATION OF ENZYMES

S.No.	Enzyme class	Examples
1.	Oxidoreductases: Transfer of hydrogen or addition of oxygen	Lactate dehydrogenase (NAD); Glucose-6-phosphate dehydrogenase (NADP); Succinate dehydrogenase (FAD); di-oxygenases
2.	Transferases: Transfer of groups other	Aminotransferase.

	than hydrogen	(Subclass: Kinase, transfer of phosphoryl group from ATP; e.g. Hexokinase)
3.	Hydrolases: Cleave bond and add water.	Acetyl choline esterase; Trypsin
4.	Lyases: Cleave without adding water.	Aldolase; HMG CoA lyase; ATP Citrate lyase. (Subclass: Hydratase; add water to a double bond)
5.	Isomerases: Intramolecular transfers. They include racemases and epimerases.	Triose phosphate isomerase
6.	Ligases: ATP dependent condensation of two molecules.	Acetyl CoA carboxylase; Glutamine synthetase; PRPP synthetase

ENZYMES WITH THEIR RESPECTIVE SUBSTRATE AND COMPETITIVE INHIBITORS

S. No.	ENZYME	SUBSTRATE	INHIBITOR	SIGNIFICANCE OF INHIBITION
1.	Acetylcholine Esterase	Acetylcholine	Succinyl choline	Used in surgery for muscle relaxation in anaesthetized patient
2.	Dihydrofolate reductase	Dihydrofolic acid (DHF)	Aminopterin Amethopterin Methotrexate	Employed in the treatment of leukemias and other cancers
3.	Dihydropteroate synthase	Para-amino benzoic acid (PABA)	Sulphonamides	Prevent bacterial synthesis of folic acid
4.	HMG CoA Reductase	HMG CoA	Lovastatin	Inhibit cholesterol biosynthesis
5.	Monoamine oxidase	Catecholamines (epinephrine, norepinephrine)	Ephedrine Amphetamine	Useful in elevating catecholamine levels
6.	Vitamin K epoxide reductase	Vitamin K	Dicumarol	Act as coagulant
7.	Xanthine oxidase	Hypoxanthine	Allopurinol	Used in control of gout to reduce excess production of uric acid from hypoxanthine

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