Semester-IV

Sub Name-medicinal chemistry-I (sub code-BP-402T)

Objective

Drugs acting on Central Nervous System Sedatives and Hypnotics

- · Benzodiazepines: SAR of Benzodiazepines, Chlordiazepoxide, Diazepam*, Oxazepam, Chlorazepate, Lorazepam, Alprazolam, Zolpidem
- · Barbiturtes: SAR of barbiturates, Barbital*, Phenobarbital, Mephobarbital, Amobarbital, Butabarbital, Pentobarbital, Secobarbital
 Miscelleneous
- · Amides and imides: Glutethmide.
- · Alcohol and their carbamate derivatives: Meprobomate, Ethchlorvynol. Aldehyde and their derivatives: Triclofos sodium, Paraldehyde. Antipsychotics
- · Phenothiazeines: SAR of Phenothiazeines Promazine hydrochloride, Chlorpromazine hydrochloride*, Triflupromazine, Thioridazine hydrochloride, Piperacetazine hydrochloride, Prochlorperazine maleate, Trifluoperazine hydrochloride.
- · Ring Analogues of Phenothiazeines: Chlorprothixene, Thiothixene, Loxapine succinate, Clozapine.
- · Fluro buterophenones: Haloperidol, Droperidol, Risperidone.
- · Beta amino ketones: Molindone hydrochloride. Benzamides: Sulpieride.

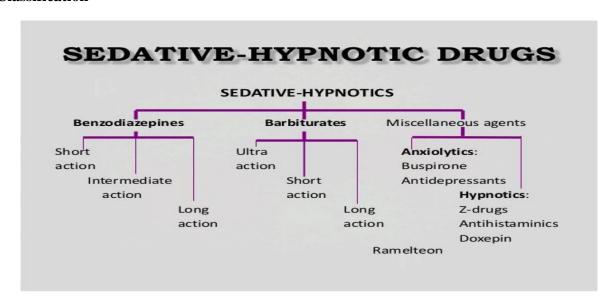
Anticonvulsants: SAR of Anticonvulsants, mechanism of anticonvulsant action

- · Barbiturates: Phenobarbitone, Methabarbital.
- · Hydantoins: Phenytoin*, Mephenytoin, Ethotoin.
- · Oxazolidine diones: Trimethadione, Paramethadione.
- · Succinimides: Phensuximide, Methsuximide, Ethosuximide*.
- · Urea and monoacylureas: Phenacemide, Carbamazepine*.
- · Benzodiazepines: Clonazepam.
- · Miscellaneous: Primidone, Valproic acid, Gabapentin, Felbamate.

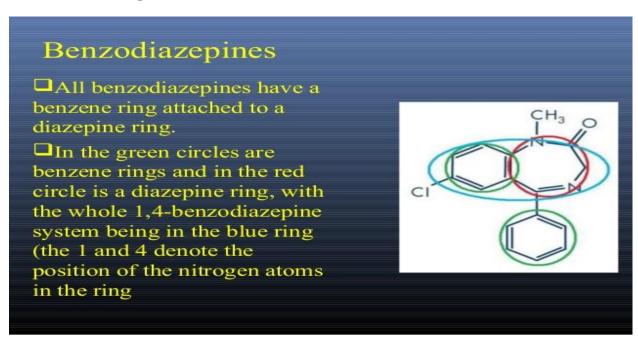
Sedative and Hypnotics

Sedative-hypnotics are a class of drugs that cause a dose-dependent depression of the CNS function, inducing sedation, sleep, and unconsciousness with increasing dose. Agents in this class of drugs include benzodiazepines and Z-drugs, barbiturates, and melatonin agonists.

Classification



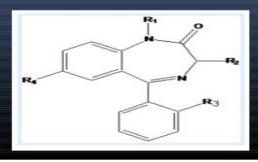
SAR of Benzodiazepines

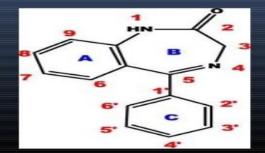


Benzodiazepines

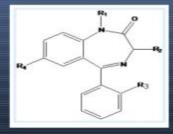
Different benzodiazepines have been developed through chemical substitutions at two major positions on the benzodiazepine structure

Therefore, all benzodiazpines are simply variations on the same core chemical structure.



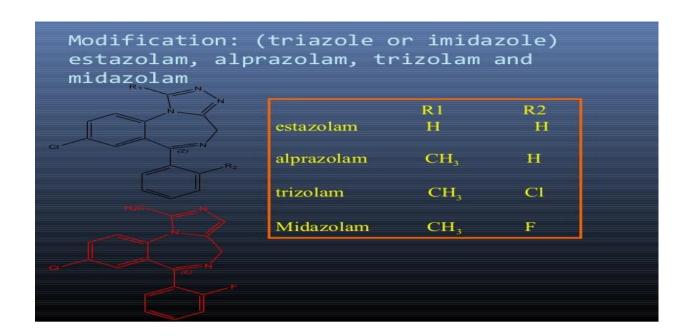


Benzodiazepine



	R1	R2	R3	R4
a	СНЗ	H	H	C1
b	CH3	Н	F	C1
c	H	H	H	NO2
d	H	H	Cl	NO2

- a. diazepam b.flutoprazepam c. nitrazepam
- d. clonazepam



1. Chlordiazepoxide

2. Diazepam

3. Oxazepam

4. Chlorazepate

5. Lorazepam

6. Alprazolam

7. Zolpidem

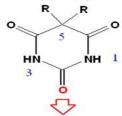
$$H_3C$$
 N
 CH_3
 H_3C

Barbiturtes:

SAR of Barbiturtes



5,5,-disubstituted & 1,5,5trisubstituted are <u>active</u>



All other substitution ▶ inactive

1,3-disubstituted or 1,3,5,5tetrasubstituted are <u>inactive</u> or produce <u>convulsions</u>

if replaced with S give thiobarbiturates

- * Replacement of C-2 O by S → ↑ lipid solubility. Thiopental used as IV anesthetics due to rapid onset & quick brain levels achieved.
- * Introduction of more sulfur atoms (2,4-dithio derivatives) destroys potency, due to decreased hydrophilic character beyond required limits.

4

1. Barbital

2. Phenobarbital

3. Mephobarbital

4. Amobarbital

$$H_3C$$
 CH_3

5. Butabarbital

6. Pentobarbital

$$\begin{array}{c} & & & \\ & & \\ & & \\ \text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3 \\ & & \\ & & \\ \text{CH}_3 \end{array} \begin{array}{c} & & \\ & & \\ & & \\ & & \\ \end{array} \begin{array}{c} & \\ & \\ \end{array} \begin{array}{c} & \\ & \\ & \\ \end{array} \begin{array}{c} & \\ \\ \end{array} \begin{array}{c} & \\ & \\ \end{array} \begin{array}$$

7. Secobarbital

Miscelleneous

1. Glutethmide

2. Meprobomate

3. Ethchlorvynol

4. Triclofos sodium

eBiochemicals

5. Paraldehyde

$$H_3C$$
 O CH_3 CH_3

Sedative-Hypnotics: Clinical Uses

- For relief of anxiety
- For insomnia
- For sedation and amnesia before and during medical and surgical procedures
- For treatment of epilepsy and seizure states
- As a component of balanced anesthesia (intravenous administration)
- For control of ethanol or other sedative-hypnotic withdrawal states
- For muscle relaxation in specific neuromuscular disorders
- As diagnostic aids or for treatment in psychiatry

Adverse Effects

- · Drowsiness and confusion: Most common AE
- Ataxia occurs at high doses
- Cognitive impairment (decreased long-term recall and retention of new knowledge) can occur with use of benzodiazepines.
- Benzodiazepines should be used cautiously in patients with liver disease.
- Alcohol and other CNS depressants enhance the sedative—hypnotic effects of the benzodiazepines.
- Administration in third trimester can result in "floppy-infant syndrome"



Antipsychotics

Antipsychotics, also known as narcoleptics or major tranquilizers, are a class of medication primarily used to manage psychosis (including delusions, hallucinations, paranoia or disordered thought), principally in schizophrenia and bipolar disorder. Antipsychotics are usually effective in relieving symptoms of psychosis in the short term.

The long-term use of antipsychotics is associated with adverse effects such as involuntary movement disorders, gynecomastia, impotence, weight gain and metabolic syndrome.

CLASSIFICATION OF ANTIPSYCHOTIC DRUGS

Typical antipsychotics

- Phenothiazines
 - e.g. chlorpromazine, fluphenazine, thioridazine
- Butyrophenones
 - e.g. haloperidol, droperidol
- Thioxanthines
 - e.g. chlorprotixen, thiothixene

Atypical antipsychotics

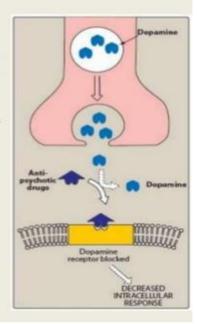
- Clozapine
- Risperidone
- · Sulpiride
- Sertindole
- · Seroquel
- Olanzapine
- · Quetiapine.

Mechanism of action

antipsychotic agents currently >A11 typical block employed clinically post synaptic dopaminergic D2 receptors in the mesolimbic and cortex regions of the brain and act competetive antagonist of dopamine. The blokade of D₂ receptors is thought to be responsible for decreasing the positive symptoms schizophrenia.

ightharpoonupCentral dopamine receptors are subdivided into D_1 , D_2 and according to some sources, D_3 receptors. These receptors have a high affinity for dopamine, but they differ in selectivity to neuroleptics of various chemical classes.

Adenylyl cyclase is involved which is responsible for the conversion of adenosine triphosphate to cyclic adenosine monophosphate (cAMP).



SAR of Phenothiazeines

Structure activity relationship of Phenothiazine...

THE NEUROLEPTIC PROPERTIES OF PHENOTHIAZINE MAY BE AFFECTED BY FOLLOWING:

- A) NATURE OF THE CHAIN IN POSITION 10
- B) NATURE OF THE AMINO GROUP
- C) R2 SUBSTITUTION
- ✓ Replacement of the h in position 2
- ✓ Substitution at position 3
- ✓ Substitution at position 1
- ✓ Three carbon chain
- ✓ Branching with larger groups
- ✓ Replacement of terminal alkyl amino group

1. Promazine hydrochloride

2. Chlorpromazine hydrochloride

3. Triflupromazine

4. Thioridazine hydrochloride

5. Piperacetazine hydrochloride

6. Prochlorperazine maleate

7. Trifluoperazine hydrochloride

8. Chlorprothixene

9. Thiothixene

10. Loxapine succinate

11. Clozapine

12. Haloperidol

13. Droperidol

14. Risperidone

15. Molindone hydrochloride

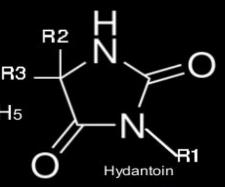
Anticonvulsants:

Anticonvulsants are a diverse group of pharmacological agents used in the treatment of epileptic seizures. Anticonvulsants are also increasingly being used in the treatment of bipolar disorder and borderline personality disorder, since many seem to act as mood stabilizers, and for the treatment of neuropathic pain.

SAR of Anticonvulsants

1. Hydantoins

- Phenylethylhydation
 R₁ = H R₂ = C₂H₅ R₃ = C₆H₅
- Phenytoin
 R1 = H R2 = R3 = C6H5
- Mephenytoin
 R1 = CH3 R2 = C2H5 R3 = C6H5
- Ethotoin
 R₁ = C₂H₅
 R₅ = H
 R₅ = C₆H₅





Contd..

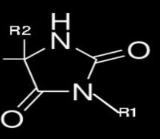
- A phenyl or other aromatic substituents at C₅ is essential for the activity.
- Alkyl substituents at position 5 may contribute to sedation, a property absent in phenytoin. Hydantoin

Hydantoin



Contd..

- Among other hydantoins, like spirohydantoins, thiohydantoins, dithiohydantoins, and 1, 3disubstituted hydantoins, some exhibit activity against chemically induced convulsions.
- While remaining are ineffective against electroshock induced convulsions.



Hydantoin



Mechanism of Action

Anticonvulsants suppress the excessive rapid firing of neurons during seizures. Anticonvulsants also prevent the spread of the seizure within the brain. Conventional antiepileptic drugs may block sodium channels or enhance γ -aminobutyric acid (GABA) function.

1. Phenobarbitone

2. Methabarbital

3. Phenytoin

4. Mephenytoin

5. Ethotoin

$$N-C_2H_5$$

6. Trimethadione

7. Paramethadione

8. Phensuximide

9. Methsuximide

10. Ethosuximide

$$O = \bigvee_{N = O}^{CH_3} CH_3$$

11. Phenacemide

12. Carbamazepine

13. Clonazepam

14. Primidone

15. Valproic acid

16. Gabapentin

17. Felbamate

$$H_2N$$
 O O NH_2

Uses of Anticonvulsants

- Anticonvulsants work by calming hyperactivity in the brain in various ways.
- For this reason, some of these drugs are used to treat epilepsy, prevent migraines, and treat other brain disorders.
- They are often prescribed for people who have rapid cycling -- four or more episodes of mania and depression in a year.

Adverse effects of Anticonvulsants

- Dizziness.
- Drowsiness.
- Fatigue.
- Nausea.
- Tremor.
- Rash.
- Weight gain

Learning outcomes

➤ Know the structural activity relationship (SAR) of different class of drugs.