Al-Mustaqbal University College Department of Pharmacy 4th stage Pharmacology II Lecture: 2



# **CNS Stimulants**

## **Overview**

## • CNS stimulants include:

- 1. The **psychomotor** stimulants cause:
  - Excitement and euphoria
  - Decrease feelings of fatigue
  - Increase motor activity
- 2. The hallucinogens produce
  - Changes in thought and mood



• CNS stimulants have diverse clinical uses and are important as drugs of abuse

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## **PSYCHOMOTOR STIMULANTS**



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- They are a **purine-derived** group of pharmacologic agents.
- They are clinically used as bronchodilatory and stimulatory agents.
- The methylxanthines include:
- 1. Theophylline which is found in tea
- 2. Theobromine found in cocoa
- **3. Caffeine**, the most widely consumed stimulant in the world, is found in the highest concentration in certain **coffee** products, **tea**, **cola** drinks, **energy drinks**, **chocolate candy**, and **cocoa**.







## **Mechanisms of action:**

- 1. Translocation of extracellular calcium
- 2. Increase in cyclic AMP and cyclic GMP caused by inhibition of phosphodiesterase
- 3. Blockade of adenosine receptors.
- The latter most likely accounts for the actions achieved by the usual consumption of caffeine-containing beverages.



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## Actions:

## 1. CNS effects:

- The **caffeine** contained in **1-2 cups** of coffee (**100 -200 mg**) causes a <u>decrease in</u> <u>fatigue and increased mental alertness</u>.
- Consumption of 1.5 g of caffeine (12-15 cups of coffee) produces <u>anxiety and</u> tremors.
- The **spinal cord** is stimulated only by **very high doses (2 5 g)** of caffeine.
- Tolerance can rapidly develop to the stimulating properties of caffeine, and withdrawal consists of feelings of fatigue and sedation.

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## **Actions:**

## 2. Cardiovascular system:

- A high dose of caffeine has **positive inotropic** and **chronotropic effects** on the heart.
- In others, an accelerated heart rate can trigger premature ventricular contractions.

## 3. Diuretic action:

• Caffeine has a **mild diuretic** action that increases the urinary output of sodium, chloride, and potassium.

## 4. Gastric mucosa:

 Because methylxanthines stimulate the secretion of gastric acid, individuals with peptic ulcers should avoid foods and beverages containing methylxanthines.

## **Therapeutic uses:**

- Caffeine and its derivatives **relax** the smooth muscles of the **bronchioles**.
- Previously the mainstay of asthma therapy, theophylline has been largely replaced by other agents, such as β2 agonists and corticosteroids
- Caffeine is also used in combination with the analgesics like acetaminophen and aspirin for the management of headaches in both prescription and over-the-counter products.





## **Pharmacokinetics:**

- The methylxanthines are well absorbed **orally**.
- Caffeine **distributes** throughout the body, including the **brain**.
- These drugs cross the placenta to the fetus and are secreted into the breast milk.
- All **methylxanthines** are metabolized in the **liver**, generally by the CYP1A2 pathway, and the metabolites are excreted in the **urine**.

## **Adverse effects:**

- Moderate doses of caffeine cause insomnia, anxiety, and agitation.
- High dosage results in toxicity, which is manifested by emesis and convulsions.
- Lethal dose is 10 g of caffeine (about 100 cups of coffee), which induces cardiac arrhythmias.
- Lethargy, irritability, and headache occur in users who routinely consume more than 600 mg of caffeine per day (roughly 6 cups of coffee per day) and then suddenly stop.

- Nicotine is the active ingredient in **tobacco**.
- It is **not** currently used **therapeutically** (except in **smoking cessation** therapy).
- It is **second** only to caffeine as the most widely used CNS **stimulant**, and it is **second** only to alcohol as the most **abused** drug.
- In combination with the <u>tars and carbon monoxide</u> found in cigarette smoke, nicotine represents a serious risk factor for <u>lung and cardiovascular</u> disease, various cancers, and other illnesses.
- **Dependency** on the drug is not easily overcome.



#### **Mechanism of action:**

- In low doses, nicotine causes ganglionic stimulation by depolarization.
- At high doses, nicotine causes ganglionic blockade.
- Nicotine receptors exist in the CNS, which participate in the stimulant effect.
  Pharmacokinetics:
- It is highly lipid soluble, absorption readily occurs via the <u>oral mucosa</u>, <u>lungs</u>, <u>GI mucosa</u>, and <u>skin</u>.
- Nicotine crosses the **placental** membrane and is secreted in **breast milk**.
- By inhaling tobacco smoke, the average smoker takes in 1 to 2 mg of nicotine per cigarette and the acute **lethal** dose is **60 mg**.
- More than **90%** of the nicotine **inhaled** in smoke is **absorbed**.
- Clearance of nicotine involves metabolism in the lung and the liver and urinary excretion.
- **Tolerance** to the effects of nicotine develops rapidly, often within days.

## **Actions:**

## **CNS:**

- Nicotine is highly lipid soluble and readily crosses the blood-brain barrier.
- Cigarette smoking or administration of **low doses** of nicotine produces some degree of **euphoria** and **arousal**, as well as **relaxation**.
- It improves attention, learning, problem-solving, and reaction time.
- High doses of nicotine result in central respiratory paralysis and severe hypotension caused by medullary paralysis.
- Nicotine is also an **appetite suppressant**.

## **Actions:**

## **Peripheral effects:**

- Stimulation of the **sympathetic ganglia** as well as of the **adrenal medulla increases** blood pressure and heart rate.
- In addition, nicotine-induced **vasoconstriction** can decrease coronary blood flow, adversely affecting a patient with **angina**.
- Stimulation of the parasympathetic ganglia also increases motor activity of the bowel.

## **Adverse effects:**

- The CNS effects of nicotine include irritability and tremors.
- Nicotine may also cause intestinal cramps, diarrhea, and increased heart rate and blood pressure.
- In addition, cigarette smoking **increases** the rate of **metabolism** for a number of drugs.

## Withdrawal syndrome:

- Nicotine is an **addictive** substance, and **physical dependence** develops rapidly and can be severe.
- Withdrawal is characterized by
- ✓ Irritability
- ✓Anxiety
- ✓ Restlessness
- ✓ Difficulty concentrating, headaches, and insomnia
- ✓ Appetite is affected, and GI upset often occurs

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## Varenicline

- Varenicline is a **partial agonist** at neuronal **nicotinic** acetylcholine receptors in the CNS.
- It produces **less euphoric effects** than nicotine (nicotine is a full agonist at these receptors).
- Thus, it is useful as an adjunct in the management of smoking cessation in patients with nicotine withdrawal symptoms.
- Additionally, varenicline tends to attenuate the **rewarding effects** of nicotine if a person relapses and uses tobacco.
- Patients taking varenicline should be monitored for suicidal thoughts, vivid nightmares, and mood changes.



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## Cocaine

- Cocaine is a widely available and highly **addictive drug**.
- Cocaine is classified as a **Schedule II drug** of controlled substances.
- The mechanism of action of cocaine is the **blockade of the reuptake of the monoamines** (norepinephrine, serotonin, and dopamine) into the presynaptic terminals.
- This **potentiates** and **prolongs** the **CNS** and **peripheral** actions of these monoamines.
- In particular, the **prolongation** of **dopaminergic** effects in the brain (limbic system) produces intense **euphoria**.
- Chronic intake of cocaine depletes dopamine triggering the vicious cycle of craving for cocaine which temporarily relieves severe depression.

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## Cocaine

Schedule	Definitions	Examples
Schedule I	No accepted medical use with a lack of	Heroin, peyote, ecstasy,
	accepted safety and high abuse potential;	lysergic acid diethylamide
	medications within this schedule may not be	(LSD)
	prescribed, dispensed, or administered for	
	medical use	
Schedule II	High abuse potential with severe	Morphine, codeine,
	psychological or physical dependence;	hydromorphone, methadone,
	however, these medications have an accepted	oxycodone, fentanyl,
	medical use and may be prescribed,	methylphenidate,
	dispensed, or administered	amphetamine, pentobarbital,
		combination products with <
		15 mg of hydrocodone per
		dosage unit (eg, Vicodin)
Schedule III	Abuse potential less than Schedules I or II but	Products with < 90 mg of
	more than Schedule IV medications; abuse	codeine per dosage unit (eg,
	may lead to moderate or low physical	Tylenol with codeine),
	dependence or high psychological dependence	dronabinol, anabolic steroids,
		ketamine
Schedule IV	Abuse potential less than Schedule III but	Propoxyphene, various
	more than Schedule V medications	benzodiazepines, sibutramine
Schedule V	Medications with the least potential for abuse	Robitussin AC, Phenergan
	among the controlled substances	with codeine, pregabalin

#### Table I. Schedules of Controlled Substances within the CSA.<sup>3,4</sup>

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- Amphetamine is a sympathetic amine that shows neurologic and clinical effects similar to cocaine.
- **Dextroamphetamine** is a major member of this class of compounds.
- Methamphetamine is a derivative of amphetamine available for prescription use.
- **3,4-Methylenedioxymethamphetamine** (also known as MDMA, or Ecstasy) is a synthetic derivative of methamphetamine with both stimulant and hallucinogenic properties.

#### **Mechanism of action:**

- Amphetamine has an **indirect** effect via an **elevation** of the level of **catecholamines** in synaptic spaces by **releasing intracellular stores.**
- Also, it inhibits MAO, and it is a weak reuptake inhibitor.
- Despite different mechanisms of action, the **behavioral** effects of amphetamine and its derivatives are **similar** to those of **cocaine**.



## **Actions:**

- CNS:
- The **major behavioural effects** of amphetamine result from a combination of its dopamine and norepinephrine release-enhancing properties.
- Amphetamine <u>increased alertness</u>, <u>decreased fatigue</u>, <u>depressed appetite</u>, and <u>insomnia</u>.
- At high doses, psychosis and convulsions can ensue.

## Sympathetic nervous system:

• In addition to its CNS action, amphetamine acts on the adrenergic system, indirectly stimulating the receptors through norepinephrine release.

## **Therapeutic uses:**

## **Attention deficit hyperactivity disorder (ADHD):**

- Some young children are **hyperkinetic** and **lack** the ability to be involved in any one activity for **longer** than a few minutes.
- Dextroamphetamine, methamphetamine, and methylphenidate can help **improve** attention **span** and **alleviate** many of the **behavioral** problems associated with ADHD and **reduce** hyperkinesia.
- Lisdexamfetamine is a prodrug that is converted to the active component dextroamphetamine after GI absorption and metabolism.



## **Therapeutic uses:**

## **Attention deficit hyperactivity disorder (ADHD):**

- Atomoxetine is a non-stimulant drug approved for ADHD in children and adults.
- Unlike methylphenidate, which blocks dopamine reuptake more than norepinephrine reuptake, atomoxetine is more selective for inhibition of norepinephrine reuptake.
- Therefore, it is **not** considered **habit forming** and is **not** a **controlled substance**.



#### **Therapeutic uses:**

#### Narcolepsy:

- Narcolepsy is a relatively **rare** sleep disorder that is characterized by **uncontrollable bouts** of sleepiness during the **day**.
- It is sometimes accompanied by **catalepsy**, a loss in muscle control, and even paralysis brought on by strong emotions such as laughter.
- The sleepiness can be treated with drugs, such as mixed amphetamine salts or methylphenidate.
- Modafinil and its R-enantiomer derivative, Armodafinil, are considered first-line agents for the treatment of narcolepsy.
- Modafinil is effective **orally**, it is well distributed throughout the body and undergoes extensive **hepatic** metabolism, and the metabolites are excreted in the **urine**.
- Headaches, nausea, and nervousness are the primary adverse effects, they may have some potential for abuse and physical dependence.



## **Therapeutic uses:**

## **Appetite suppression:**

- **Phentermine** and **diethylpropion** are sympathomimetic amines that are related structurally to amphetamine.
- These agents are used for their **appetite-suppressant** effects in the management of obesity.





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## Pharmacokinetics:

- Amphetamine is completely **absorbed** from the GI tract, **metabolized** by the liver, and **excreted** in the urine.
- Note: Administration of urinary alkalinizing agents such as sodium bicarbonate will increase the <u>nonionized species</u> of the drug and <u>enhance</u> the reabsorption of dextroamphetamine from the renal tubules into the bloodstream.
- Amphetamine abusers often administer the drugs by IV injection and/or by smoking.
- The **euphoria** caused by amphetamine lasts 4 to 6 hours, or four- to eightfold longer than the effects of **cocaine**.

## **Adverse effects:**

## **CNS effects:**

- Adverse effects of amphetamine usage include insomnia, irritability, weakness, dizziness, tremor, and hyperactive reflexes.
- Amphetamine can also cause <u>confusion</u>, <u>delirium</u>, <u>panic</u> states, and <u>suicidal</u> tendencies, <u>especially in mentally ill patients</u>.
- Chronic amphetamine use produces a state of "amphetamine psychosis" that resembles the psychotic episodes associated with schizophrenia.



## **Adverse effects:**

## **Cardiovascular effects:**

 Amphetamine causes <u>palpitations</u>, <u>cardiac arrhythmias</u>, <u>hypertension</u>, <u>anginal</u> <u>pain</u>, <u>and circulatory collapse</u>.

## **GI system effects:**

• Amphetamine acts on the GI system, causing <u>anorexia</u>, <u>nausea</u>, <u>vomiting</u>, <u>abdominal cramps</u>, <u>and diarrhea</u>.

### **Contraindications:**

- Patients with hypertension, cardiovascular disease, hyperthyroidism, glaucoma.
- History of drug abuse or those taking MAO inhibitors.

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## Methylphenidate

- Methylphenidate has **CNS-stimulant** properties similar to those of amphetamine and may also lead to **abuse**, although its **addictive** potential is controversial.
- It is a **Schedule II** drug.
- Methylphenidate is presently one of the most prescribed medications in **children**.
- It is estimated that **4 to 6 million** children in the United States take methylphenidate daily for **ADHD**.
- Its active isomer, **dexmethylphenidate**, is also a **Schedule II** drug used for the treatment of **ADHD**.



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## **Mechanism of action:**

- Children with ADHD may produce weak dopamine signals, which suggests that onceinteresting activities provide fewer rewards to these children.
- Methylphenidate is a dopamine and norepinephrine transport inhibitor and may act by increasing both dopamine and norepinephrine in the synaptic space.
- Methylphenidate may have **less potential for abuse** than cocaine, because it enters the brain much **more slowly** than cocaine and, thus, does not increase dopamine levels as **rapidly**.



## Methylphenidate

## **Therapeutic uses:**

- Methylphenidate has been used for the treatment of **ADHD**.
- Methylphenidate is also effective in the treatment of **narcolepsy**.
- Unlike, methylphenidate, **dexmethylphenidate** is **not** indicated in the treatment of **narcolepsy**.

## **Pharmacokinetics:**

- Both methylphenidate and dexmethylphenidate are readily **absorbed** after oral administration.
- Methylphenidate is available in **extended-release** oral formulations and as a **transdermal patch** for once-daily application.
- The de-esterified product, **ritalinic acid**, is excreted in urine.

## Methylphenidate

## **Adverse effects:**

- GI adverse effects are the most common and include abdominal pain and nausea.
- Other reactions include <u>anorexia</u>, insomnia, nervousness, and fever.
- In **seizure** patients, methylphenidate may **increase** seizure **frequency**, especially if the patient is taking **antidepressants**.
- It is **contraindicated** in patients with <u>glaucoma</u>.
- Methylphenidate can **inhibit the metabolism** of warfarin, phenytoin, phenobarbital, primidone, and tricyclic antidepressants.

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## HALLUCINOGENS

- A few agents have, as their primary action, the ability to induce **altered** <u>perceptual states reminiscent of dreams</u>.
- Many of these altered states are accompanied by visions of bright, colourful changes in the environment and by the plasticity of constantly changing shapes and color.
- The individual under the influence of these agents is **incapable** of **normal decision making** because the drug interferes with **rational thought**.
- These compounds are known as hallucinogens, and lysergic acid diethylamide (LSD) and tetrahydrocannabinol (from marijuana) are examples of agents in this class.



## THANK YOU FOR YOUR ATTENTION

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