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3.1

Q3.1.1 Short Answer: What is psychopharmacology, and why do we study it?

Answer: The scientific study of the effects of drugs on the nervous system and behavior; to learn about psychotherapeutic drugs and gain a better understanding of how these things work.

- Q3.1.2 Multiple Choice: Which of the following is NOT a function of psychopharmacology?
 - (A) Study the effects of drugs on the nervous system
 - (B) Study the effects of drugs on behavior
 - (C) Study the effects of drugs on the immune system
 - (D) Study the effects of drugs on neurotransmitter systems

Answer: C.

Q3.1.3 Fill in the Blank: The location at which a drug interacts with the body to produce its effects is called the _____.

Answer: Site of action.

Q3.1.4 Short Answer: What is the difference between an agonist and an antagonist?

Answer: An agonist mimics or enhances the effects of a neurotransmitter, while an antagonist blocks or inhibits the effects of a neurotransmitter.

Q3.1.5 True or False: Drugs directly create effects in the body.

Answer: False. They modulate ongoing cellular activity.

- Q3.1.6 Multiple Choice: Which of the following is an example of a drug that acts as an agonist?
 - (A) Naloxone (B) Morphine (C) Curare (D) Atropine

Answer: B.

Q3.1.7 Short Answer: What is selective action?

Answer: The ability of a drug to affect only certain types of receptors or neurotransmitter systems, minimizing side effects.

Q3.1.8 Short Answer: What is an example of how an agonistic effect can become antagonistic?

Answer: If a drug increases the release of a neurotransmitter and also blocks its reuptake, it can lead to an excess of the neurotransmitter, which may inhibit further release.

Q3.1.9	3.1.9 Multiple Choice: What is a precursor?				
	 (A) A substance that inhibits neurotransmitter release (B) A substance that enhances neurotransmitter release (C) A substance from which another substance is formed (D) A substance that blocks neurotransmitter receptors Answer: C.				
Q3.1.10	Fill in the Blank: The process of creating a called Answer: Synthesis.	neurotransmitter from its precursors is			
Q3.1.11	Prill in the Blanks: A(n) as neurotransmitter and its effer binds to a different site on the receptor and transmitter. Answer: Direct, mimics, indirect, enhances.	cts, while a(n) agonist			
Q3.1.12	2 Fill in the Blanks: A(n) are receptor and the effects of the antagonist binds to the same receptor as the its effects. Answer: Indirect, blocks, direct, inhibits.	neurotransmitter, while a(n)			
Q3.1.13	3 Multiple Choice: Drugs that cause the action are called:	n potential to stay in a depolarized state			
	(C) Antagonists (D)	Depolarizing agents Inverse agonists			
Q3.1.14	Answer: B 4 Fill in the Blanks: In the following diagram, and identify the outcome:	label the specific enzyme for each arrow,			

Q3.1.15 Long(-ish) Answer: Describe the difference between a neurotransmitter and a neuromodulator.

Answer: A neurotransmitter is a chemical messenger that transmits signals across a synapse from one neuron to another, while a neuromodulator is a substance that modulates the activity of neurotransmitters, often affecting a larger area of the brain and influencing the overall tone of neural activity.

Q3.1.16 Matching: Match the following examples with them either being an antagonist or an agonist. (Some of these may be direct or indirect. Specify each one.)

Choices
(a) Curare
(b) Atropine
(c) Morphine
(d) Naloxone
(e) Botulinum Toxin
(f) Interfering with docking proteins
(g) Blocking the reuptake of a neurotransmitter
(h) Sarin
(i) Interfering with vesicles
(j) Blocking receptors
(k) Black widow spider venom
(l) Cobra and Krait Venom
(m) Parathion
(n) DFP
(o) Physostigmine
(1) Direct antagonist
(2) Indirect antagonist
(3) Direct agonist
(4) Indirect agonist
(5) Antagonist
(6) Agonist

3.2

Q3.2.1 Multiple Choice: Which of the following neurochemicals does NOT transmit information (according to our notes)?

	(A) Dopamine	(B) Glutamate
	(C) GABA	(D) Glycine
	Answer: A.	
Q3.2.2	Fill in the Blank: Peptides are short chanswer: Amino acids.	ains of
Q3.2.3	Fill in the Blanks: The difference between and opiates are	een opioids and opiates are that opioids are
Q3.2.4	Short Answer: What is the pain pathw down? (Generally speaking.)	ay for the face? What about from the neck
	Answer: From the face (specifically the transfer conducting, C-fibers (unmyelinated) s	rigeminal nerve); A-delta fibers (mylinated) slow conducting fibers
Q3.2.5	Fill in the Blanks: The three types of opi and Answer: Mu, Kappa, Delta.	oid receptors are
Q3.2.6	neurochemicals bind to each the most? Answer: Mu: Analgesia and euphoria,	ee opioid receptors responsible for, and what Endorphins; Delta: Analgesia, Enkephalins; mines, learning and memory, emotional con- rphins.
Q3.2.7	Multiple Choice: Prostaglandins become	e active during
	(A) Resting-and-Digesting	(B) Crying
	(C) Daydreaming	(D) Bleeding
	Answer: D.	
Q3.2.8	True or False: Celecoxib (Celebrex), market because it causes heart attacks an Answer: False; Rofecoxib (Vioxx) is the a	
Q3.2.9	Fill in the Blank: Cylooxygenase (COX) to its active state. Answer: Prostaglandins	is an enzyme that converts inactive
Q3.2.10	pain pathway.	or the direct pain pathway and the indirect zed pain, immediate and brief, mechanical

and dull pain, takes longer, but lingers, chemical (inflammatory) pain,

(strong) and thermal (extreme temperature); Indirect: Slow pain, throbbing, aching

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Q3.2.11	Fill in the Blanks: Pain arrives at the, then travels to the, then travels to the Once there, it is processed by several brain regions. First, the
	contributes to arousal. Then, the, particularly the
	anterior cingulate cortex (ACC), processes the emotional aspects of pain. When the
	pain is overwhelming, the activates and releases endogenous opioids
	to reduce the sensation—this allows a person, for example, to escape danger despite a
	severe injury. Finally, the and other areas help interpret and associate the pain with context.
	Answer: Brain stem reticular formation (BSRF); Thalamus; Thalamus; Limbic system
	(ACC); Periaqueductal gray region; Frontal lobes
Q3.2.12	Matching: Match the following drugs with their respective NSAID class.
	Choices
	(a) Ibuprofen
	(b) Aspirin
	(c) Diflunisal
	(d) Naproxen
	(e) Salsalate
	(f) Ketoprofen
	(1) Proprionic Acid Derivatives
	(2) Salicylates
	(2) Suite, 18005
	Answer: (1): a, d, f; (2): b, c, e

Q3.2.13 Long Answer: Some studies show that both the placebo effect and acupuncture can be blocked by Naloxone, an opioid antagonist. What does this suggest about the mechanism of acupuncture's pain-relieving effects? Does this prove that acupuncture is not entirely a placebo?

Answer: This suggests that acupuncture's pain-relieving effects may be mediated by the body's endogenous opioid system, similar to the placebo effect. However, this does not prove that acupuncture is entirely a placebo; it only indicates that placebo-like mechanisms (such as expectation-induced opioid release) may contribute to its effects. Other mechanisms may also be involved.

Q3.2.14 Short Answer: What are some of the functions of opioids? (List the main effects and the side effects.)

Answer: Prevents diarrhea, gives euphoria, analgesia, changes the stress response, body temperature, emotion, feeding motivation, sexual behavior, learning, drowsiness, and promotes pro-social behavior in some cases.

Q3.2.15	True or False: The term <i>colocalized</i> means two or more neurotransmitters are released from two separate neurons at the same time. Answer: False; from the same neuron.		
Q3.2.16	Short Answer: What is the definition of pain? (DO NOT say this exam!!!!!!!) Answer: Unpleasant sensory and emotional experience associated with actual or potential tissue damage.		
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3.3			
Q3.3.1	<b>True or False:</b> The amines (monoamines) are derived from amino acids. $Answer:$ True.		
Q3.3.2	$\begin{tabular}{ll} \textbf{Short Answer:} & \textbf{Name three neurotransmitters that fall under the amino acid category.} \\ & \textbf{Answer:} & \textbf{Glutamate, GABA, Glycine.} \end{tabular}$		
Q3.3.3	Fill in the Blank: The two indolamines are and  Answer: Serotonin (5-HT) and Melatonin.		
Q3.3.4	Multiple Choice: What amino acid are indolamines derived from?		
	(A) Tryptophan (B) Tyrosine (C) Glutamate (D) Glycine		
	Answer: A.		
Q3.3.5	Fill in the Blank: The precursor to glutamate is, and the enzyme that synthesizes glutamate from it is  Answer: The precursor is glutamine; the enzyme is glutaminase.		
Q3.3.6	<b>Short Answer:</b> What receptor does ketamine bind to, and what is its effect? <i>Answer:</i> Ketamine binds to the NMDA receptor and acts as a dissociative anesthetic, which is being studied as a treatment for depression.		
Q3.3.7	Fill in the Blank: The enzyme deactivates anandamide.  Answer: Fatty acid amide hydrolase (FAAH)		
Q3.3.8	<b>True or False:</b> The most common excitatory neurotransmitter in the brain is GABA. <i>Answer:</i> False; it is glutamate.		
Q3.3.9	Fill in the Blank: The drug is a direct antagonist of the NMDA receptor and can cause hallucinations and dissociation.		

Q3.3.10 Short Answer: What transporters are responsible for glutamate reuptake, and why is this process important?

Answer: Excitatory amino acid transporters (EAATs); it prevents excitotoxicity, which can lead to brain damage (e.g., in stroke or ALS).

Answer: Phencyclidine (PCP).

Q3.3.11	Multiple Choice: Which receptor is clost tant for synaptic plasticity and memory for	•	te and is impor-
	(A) GABA receptor	(B) NMDA receptor	
	(C) Serotonin receptor	(D) Dopamine receptor	
	Answer: B.		
Q3.3.12	Short Answer: What enzyme converts neurotransmitter is GABA?  Answer: Glutamic acid decarboxylase (Gneurotransmitter in the brain.		Ų <u>-</u>
Q3.3.13	Fill in the Blanks: The three catecholan		·
Q3.3.14	Multiple Choice: What do all catecholar derived from?	nines contain, and what ami	no acid are they
	<ul><li>(A) Catechol and are derived from tryptop</li><li>(B) Catechol and are derived from tyrosine</li><li>(C) Indole and are derived from tryptophs</li><li>(D) Indole and are derived from tyrosine</li></ul>	9	
	Answer: B.		
Q3.3.15	Fill in the Blank: The enzyme	converts tyrosine in	nto L-DOPA.
Q3.3.16	Short Answer: Explain how botox interaction and thus inhibiting factors.	at the neuromuscular junc	tion, preventing
Q3.3.17	Fill in the Blank: The orbicularis oculi muscle influences  Answer: Happiness.		
Q3.3.18	3 True or False: Tyrosine is the precursor for serotonin.  Answer: False; it is the precursor for catecholamines.		
Q3.3.19	<b>Short Answer:</b> What are the names of the and epinephrine? <i>Answer:</i> Dopaminergic		
Q3.3.20	Fill in the Blanks: Melatonin is synthed in regulating  Answer: Serotonin; sleep-wake cycles (circular)		and is involved

Q3.3.21 Short Answer: What is another name for peptides in the context of neurotransmitters, and give an example. Answer: Neuropeptides; example: Endogenous opioids Q3.3.22 Multiple Choice: What is the name of the endogenous cannabinoid neurotransmitter whose name means "bliss" in Sanskrit? (A) Anandamide (B) Cannabidiol (C) Tetrahydrocannabinol (THC) (D) 2-Arachidonoylglycerol (2-AG) Answer: A. Q3.3.23 Short Answer: How are lipid-based neurotransmitters synthesized and stored? Answer: They are synthesized on demand and not stored in synaptic vesicles. Q3.3.24 Fill in the Blank: The gaseous neurotransmitter that is required for an erection is Answer: Nitric Oxide (NO). Q3.3.25 Long Answer: Describe the study that addressed the question, "Does Botox decrease emotional experience?" Describe the population, the method, and the results Answer: The study involved participants receiving Botox injections and then received either botox or restylane (dermal filler) injected. Everyone then was shown emotion evoking movies. The results are that the Botox group reported less emotional experience than the restylane group. Q3.3.26 Short Answer: Name one neurotransmitter that is a nucleoside. What is its function? Answer: Adenosine; it is involved in sleep regulation and has inhibitory effects on neurotransmission. Q3.3.27 Fill in the Blanks: Fill in the following spaces that describe the process of dopamine metabolism: DA is broken down by _____ into _____. Then, _____ converts it into . Answer: Monoamine oxidase (MAO); Dihydroxyphenylacetric acid (DOPAC); Catechol-

Q3.3.28 Short Answer: What are the results of the study into depression that asks "Can Botox be used as a good thing?" Answer

O-methyltransferase (COMT); Homovanillic acid (HVA).

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Q3.4.1	<b>Short Answer:</b> What does cholinergic in <i>Answer:</i> Cholinergic refers to the neuro ceptors.		ACh) and its re-
Q3.4.2	Multiple Choice: Who first discovered	acetylcholine in 1921?	
	<ul><li>(A) Otto von Loewy</li><li>(C) Neal Miller</li></ul>	<ul><li>(B) James Olds</li><li>(D) Peter Milner</li></ul>	
	Answer: A.		
Q3.4.3	<b>Short Answer:</b> What experiment led to <i>Answer:</i> Otto von Loewy took a frog hear pathetic part of the vagus nerve which slanother frog heart, it also slowed down, slanother frog heart from the frog heart from the frog heart from the f	ert, put it in saline, stimula owed the heart. When he	ted the parasym- put the saline in
Q3.4.4	Fill in the Blank: The original name	given to acetylcholine by i	ts discoverer was
	Answer: Vagusstoff.		
Q3.4.5	Short Answer: What are the two types Answer: Nicotinic and muscarinic recept		
Q3.4.6	<b>True or False:</b> Acetylcholine is the only no branch of the autonomic nervous system. <i>Answer:</i> True.	eurotransmitter used in the	parasympathetic
Q3.4.7	Fill in the Blanks: In the sympathetic new while NE is used at the  Answer: Preganglionic synapse (or paraw with smooth muscles and glands.		
Q3.4.8	Multiple Choice: Which of the following	g is NOT a function of AC	h in the CNS?
	(A) Learning and alertness	(B) Memory	
	(C) REM sleep generation	(D) Pain modulation	
	Answer: D.		
Q3.4.9	Short Answer: Describe the synthesis of Answer: Acetylcoenzyme-A (Acetyl-CoA from acetic acid. Then, choline acetyltra the acetyl-CoA to choline, forming acetylcoenzyme-A (Acetyl-CoA)	) attaches to an acetate ion nsferase (ChAT) transfers	
Q3.4.10	Fill in the Blanks: The precursor to acet that synthesizes acetylcholine is	·	_ and the enzyme

Q3.4.11 Short Answer: Explain how acetylcholine is metabolized.

Answer: ACh is broken down by the enzyme acetylcholinesterase (AChE) into acetate and choline. The choline is taken back up by active transport and reused, while acetate is broken down and eliminated.
Q3.4.12 Multiple Choice: Which type of ACh receptor is ionotropic?

(A) Nicotinic receptors

(B) Muscarinic receptors

(C) Both nicotinic and muscarinic receptors

Answer: A.

(D) Neither nicotinic nor muscarinic receptors

Q3.4.13 Short Answer: Explain what the sympathetic chain is, and where it is located.

Answer: The sympathetic chain is a series of ganglia located along the spinal cord that connects the sympathetic nervous system to the spinal nerves. It allows for the rapid transmission of signals throughout the body.

Q3.4.14 Fill in the Blank: The drug ______ is a direct antagonist of nicotinic receptors, causing paralysis.

Answer: Curare.

Q3.4.15 True or False: Atropine blocks muscarinic receptors and is derived from the plant known as belladonna alkaloids (deadly nightshade).

Answer: True.

Q3.4.16 Short Answer: How does Botulinum Toxin interfere with acetylcholine function? *Answer:* It interferes with Ca²⁺ influx channels, preventing the release of ACh.

Q3.4.17 Fill in the Blanks: Black Widow Spider venom causes ______ of ACh, while Cobra venom _____ ACh receptors.

Answer: Continual release; blocks.

Q3.4.18 Multiple Choice: Which of the following is a reversible AChE blocker used to treat myasthenia gravis?

(A) Sarin (B) Parathion

(C) Neostigmine (Prostigmin) (D) DFP (Diisopropylfluorophosphate)

Answer: C.

Q3.4.19 True or False: Donepezil (Aricept) crosses the blood-brain barrier and is used to treat the cognitive symptoms of Alzheimer's disease.

Answer: True.

Q3.4.20	True or False: Nicotinic recptors are antagonists at low does, but agonists at high doses.		
	Answer: False; they are agonists at low de	oses and antagonists at high doses.	
Q3.4.21	Multiple Choice: In the PNS, where are (A) Brain and spinal cord (C) Autonomic ganglia	e nicotinic receptors predominantly located? (B) Neuromuscular junctions (D) All of the above	
		(B) III of the above	
	Answer: B.		
Q3.4.22	Long Answer: Define the neuromusclar Answer:	junction and the paravertebral ganglion.	
O2 4 22	Multiple Choice. In the sympathetic new	vrous system, which nouvetransmitter is used	
Q3.4.23	at the neuromuscular junction with smoot	vous system, which neurotransmitter is used the muscles and glands?	
	(A) Acetylcholine	(B) Norepinephrine	
	(C) Dopamine	(D) Serotonin	
	Answer: B.		
Q3.4.24	<b>True or False:</b> In the sympathetic nerve mitter used at the neuromuscular junction <i>Answer:</i> True.	ous system, acetylcholine is the neurotransa with sweat glands.	
Q3.4.25	Fill in the Blanks: The the spinal cord. This is why when you gresponds at once.  Answer: Sympathetic chain; ALL	_ is a chain of ganglia that runs parallel to get anxious, of your body	
Q3.4.26	<b>Short Answer:</b> Compare the neurotranss system versus the sympathetic nervous sy <i>Answer:</i>	mitters used in the parasympathetic nervous stem.	
Q3.4.27	Multiple Choice: Which of the following tonomic nervous system is FALSE?	g statements about acetylcholine in the au-	
	(A) ACh is the only neurotransmitter in t	he parasympathetic branch	
	(B) ACh is used at preganglionic synapses branches	s in both sympathetic and parasympathetic	
	(C) ACh is used at postganglionic synapse	s to sweat glands in the sympathetic branch	
	(D) ACh is the primary neurotransmitter	at the neuromuscular junction with smooth	

muscles in the sympathetic branch

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Answer: D.

Q3.4.28 Fill in the Blank: In the somatic nervous system, ACh ______ the neuromuscular junction.

Answer: Excites.

Q3.4.29 Short Answer: Explain the role of acetylcholine in the somatic nervous system.

Answer: ACh excites the neuromuscular junction in the somatic nervous system, making it crucial for transmitting motor messages from the nervous system to skeletal muscles, resulting in voluntary movement.

Q3.4.30 Matching: Match each brain structure with its projection target.

Choices

- (a) Nucleus Basalis
- (b) Medial Septal Nucleus and Nucleus of Diagonal Band
- (c) Pedunculopontine nucleus (PPT) and Laterodorsal Tegmental Nucleus (LDT)
- Q3.4.31 Multiple Choice: Which structure in the basal forebrain that uses ACh is primarily responsible for activating the cortex and facilitating learning?
  - (A) Nucleus Basalis

- (B) Medial Septal Nucleus
- (C) Nucleus of Diagonal Band
- (D) Pedunculopontine nucleus

Answer: A.

Q3.4.32 True or False: The Medial Septal Nucleus, which uses ACh, primarily modulates the amygdala.

Answer: False. It primarily modulates the hippocampus.

Q3.4.33 Long Answer: Explain the function of acetylcholine in REM sleep generation, including the specific brain structures involved.

Answer: Acetylcholine is important for REM sleep generation through the actions of the Pedunculopontine nucleus (PPT) and Laterodorsal Tegmental Nucleus (LDT). These cholinergic structures project to the pons and thalamus, activating brain regions during REM sleep that are responsible for the vivid dreaming and rapid eye movements characteristic of this sleep phase. ACh levels are highest during REM sleep, facilitating the cortical activation seen in this paradoxical sleep state.

(A) Tyrosine (B) L-DOPA (C) Tryptophan (D) Choline

Answer: A.

Q3.5.3 Fill in the Blank: The rate-limiting enzyme in the synthesis of catecholamines is

Answer: Tyrosine Hydroxylase.

Q3.5.4 Short Answer: Describe the pathway of dopamine synthesis from its amino acid precursor.

Answer: Tyrosine is converted to L-DOPA by tyrosine hydroxylase, then L-DOPA is converted to dopamine by DOPA decarboxylase.

Q3.5.5	<b>True or False:</b> The word "tyrosine" is derived from a word meaning "tire." <i>Answer:</i> False; it is derived from "cheese"				
Q3.5.6	Multiple Choice	: Which pathway is	involved in movemen	nt and motor control?	
	<ul><li>(A) Nigrostriatal sy</li><li>(C) Mesolimbic sys</li></ul>		<ul><li>(B) Mesocortical</li><li>(D) Tuberoinfunction</li></ul>		
	Answer: A.				
Q3.5.7	Fill in the Blanks: The nigrostriatal pathway starts in the and ends in the  Answer: Substantia nigra; striatum (caudate nucleus and putamen).				
Q3.5.8	Short Answer: List four symptoms of Parkinson's disease.  Answer: Any four of: weakness, tremor at rest, muscle rigidity, problems with balance, abnormal gait, trouble learning.				
Q3.5.9	Multiple Choice Parkinson's disease		ed to the developme	ent of an animal model for	
	(A) MPTP	(B) MPPP	(C) MPP+	(D) MAO	
	Answer: A.				
Q3.5.10		alled		ains of people with Parkin	
Q3.5.11	to the Substantia I	Nigra. Huntington's Chore	,	GABA from the Striatus GABA from the Striatus	
Q3.5.12	Long Answer: E standing of Parkins Answer:		ΓP incident in 1982	contributed to our unde	
00 7 10			(D:4 1: ) :		
Q3.5.13	and	<b>k:</b> Methylphenidate in the brain. (DA); Norepinephri		levels of	
Q3.5.14	Multiple Choice ment?	: Which system is p	orimarily responsible	e for reward and reinforc	

	(A) Nigrostriatal system	(B) Mesocortical system	
	(C) Mesolimbic system	(D) Tuberoinfundibular system	
	Answer: C.		
Q3.5.15	<b>Short Answer:</b> What neuropeptide, also called orexin, is involved in the regulation of sleep and wakefulness?  Answer: Hypocretin.		
Q3.5.16	Fill in the Blank: The drug to treat insomnia.  Answer: Suvorexant (Belsomra).	is an orexin receptor antagonist used	
Q3.5.17	<b>True or False:</b> The mesocortical system is involved in short-term memory, planning and problem-solving. $Answer:$ True.		
Q3.5.18	Multiple Choice: Which researchers discovered that electrical stimulation of certain brain areas could be rewarding rather than aversive?		
	(A) Otto von Loewy and Vagusstoff	(B) James Olds and Peter Milner	
	(C) Neal Miller and Delgado	(D) Lateral hypothalamus researchers	
	Answer: B.		
Q3.5.19	Short Answer: What structure within the limbic system is considered the "pleasure center" of the brain?  Answer: Nucleus accumbens.		
Q3.5.20	Fill in the Blanks: The following is a paragraph the describes dopamine synthesis: Tyrosine is converted to by the enzyme This converted form is then used to create dopamine by the enzyme  Answer: L-DOPA; Tyrosine Hydroxylase; DOPA Decarboxylase.		
Q3.5.21	Multiple Choice: Which of the following is NOT a function of dopamine in the CNS?		
	(A) Movement and motor control	(B) Reward and reinforcement	
	(C) Learning and memory	(D) Sleep-wake cycles and REM sleep	
	Answer: D.		
Q3.5.22	<b>Short Answer:</b> Describe the metabolism of dopamine.  Answer: Dopamine is metabolized by the enzyme monoamine oxidase (MAO) into Dihydroxyphenylacetic acid (DOPAC), which is further metabolized by catechol-Omethyltransferase (COMT) into Homovanillic acid (HVA).		

 ${\bf Q3.5.23~Short~Answer:}$  Define choreoathetotic movements.

Answer: Choreoathetotic movements refer to excessive movement disorders.

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Q3.5.24		refers to slow, continuous writhing the Greek word for "dance") refers to rapid,	
Q3.5.25	<b>True or False:</b> Both athetosis and choreic movements are characterized by too little movement.  Answer: False. Both are characterized by too much movement.		
Q3.5.26	<b>Short Answer:</b> Where in the brain is hypocretin produced? <i>Answer:</i> In the lateral hypothalamus.		
Q3.5.27	Multiple Choice: Which drug increases both dopamine and norepinephrine in the brain and can be used to treat narcolepsy?		
	(A) Suvorexant (Belsomra)	(B) Methylphenidate (Ritalin)	
	(C) TAK-994	(D) Hypocretin	
	Answer: B.		
Q3.5.28	<b>Short Answer:</b> Explain the difference between athetosis and choreic movements. <i>Answer:</i> Athetosis involves slow, continuous writhing movements, while choreic movements are rapid, purposeless, involuntary movements (like dancing).		
Q3.5.29	Fill in the Blank: is a neuropeptide involved in the regulation of sleep and wakefulness that is also known as orexin.  Answer: Hypocretin		
Q3.5.30	<b>Short Answer:</b> What is the role of adenosine in the body? <i>Answer:</i> Adenosine is a nucleoside that plays a role in sleep regulation and has inhibitory effects on neurotransmission. It accumulates in the brain during wakefulness and promotes sleepiness.		