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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. (This answer needs work. IF you have a better one, please let me know and I'll update this document.)

- 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 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 from which a neurotransmitter is broken down Answer: C. Q3.1.9 Short Answer: What is an example of how an agonistic effect can become antago-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 Q3.1.10 Fill in the Blank: The process of creating a neurotransmitter from its precursors is Answer: Synthesis. Q3.1.11 Fill in the Blanks: A(n) _____ agonist binds to the same receptor as the neurotransmitter and _____ its effects, while a(n) ____ agonist binds to a different site on the receptor and ______ the effects of the neurotransmitter. Answer: Direct, mimics, indirect, enhances. Q3.1.12 Fill in the Blanks: A(n) _____ antagonist binds to a different site on the receptor and _____ the effects of the neurotransmitter, while a(n) _____ antagonist binds to the same receptor as the neurotransmitter and its effects. Answer: Indirect, blocks, direct, inhibits. Q3.1.13 Multiple Choice: Drugs that cause the action potential to stay in a depolarized state are called: (A) Agonists (B) Depolarizing agents (C) Antagonists (D) Inverse agonists Answer: B Q3.1.14 Fill in the Blanks: In the following diagram, label the specific enzyme for each arrow,

Answer: Synthetic; Metabolic; Inactive Metabolite

and identify the outcome:

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.

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- (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	
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- (2) Indirect antagonist_______
- (3) Direct agonist_______
- (4) Indirect agonist________

Answer: (1): a, b, d, l; (2): e, f, i, j; (3): c, k; (4): g, h, m, n, o.

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: acids.	Peptides are short	chains of	Answer: Amino
Q3.2.3		d opiates are		ates are that opioids are
Q3.2.4	down? (Generally spaces) Answer: From the f	peaking.)	crigeminal nerve); A-	nat about from the neck delta fibers (mylinated)
Q3.2.5	Fill in the Blanks: and	_•	ioid receptors are	,
Q3.2.6	neurochemicals bind Answer: Mu: Anal Kappa: Colocalized	to each the most? gesia and euphoria,	Endorphins; Delta: amines, learning and	esponsible for, and what Analgesia, Enkephalins; memory, emotional con-
Q3.2.7	Multiple Choice:	Prostaglandins becon	ne active during	
	(A) Resting-and-Dig	esting	(B) Crying	
	(C) Daydreaming		(D) Bleeding	
	Answer: D.			
Q3.2.8	market because it ca	elecoxib (Celebrex), auses heart attacks ar coxib (Vioxx) is the a	nd stroke.	was removed from the cor.
Q3.2.9	Fill in the Blank: to its active state. Answer: Prostagland	$Cylooxygenase\ (COX)$ dins) is an enzyme that co	nverts inactive
Q3.2.10	Long Answer: Lis pain pathway.	t the characteristics	for the direct pain pa	athway and the indirect

Answer: Direct: Sharp and well localized pain, immediate and brief, mechanical (strong) and thermal (extreme temperature); Indirect: Slow pain, throbbing, aching and dull pain, takes longer, but lingers, chemical (inflammatory) pain,

Q3.2.11	Fill in the Blanks: Pain arrives at 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
	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.

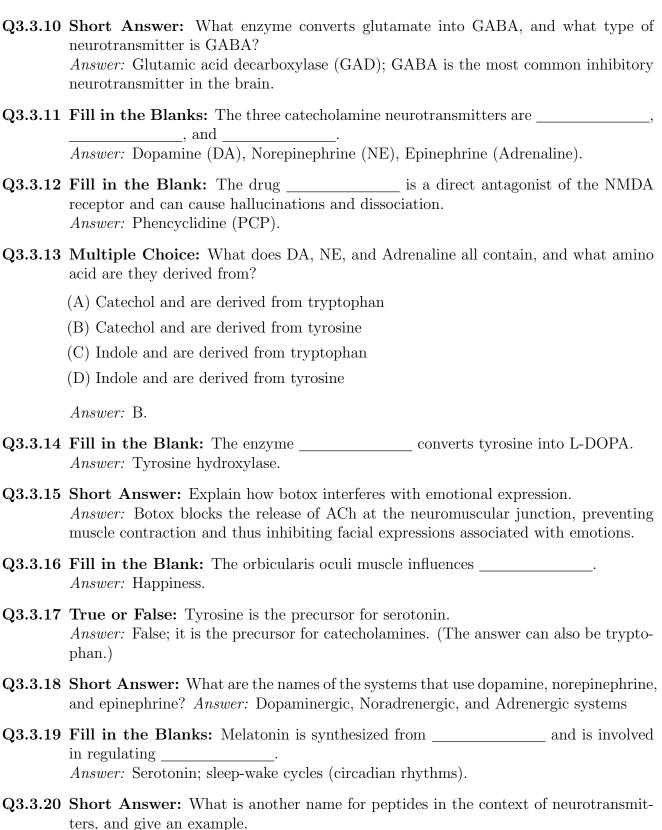
Q3.2.14 Short Answer: What are some of the functions of opioids? (List the main effects and the side effects.)

> 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	2.16 Short Answer: What is the definition of pain? (DO NOT say this exam!!!!!!!!) Answer: Unpleasant sensory and emotional experience associated with actual o tential tissue damage.		
3.3			
Q3.3.1	True or False: The amines (monoamine Answer: True.	s) are derived from amino acids.	
Q3.3.2	Fill in the Blank: The two indolamines Answer: Serotonin and Melatonin.	are and	
Q3.3.3	Multiple Choice: What amino acid are	indolamines derived from?	
	(A) Tryptophan (B) Tyrosine	(C) Thymine (D) Phenylalanine	
	Answer: A.		
Q3.3.4	Fill in the Blank: The precursor to gluthat synthesizes glutamate from it is		
Q3.3.5	Short Answer: What receptor does ketamine bind to, and what is its effect? Answer: Ketamine binds to the NMDA receptor and acts as a dissociative anesthetic, which is being studied as a treatment for depression.		
Q3.3.6	Fill in the Blank: The enzyme	deactivates anandamide.	
Q3.3.7	True or False: The most common excita <i>Answer:</i> False; it is glutamate.	tory neurotransmitter in the brain is GABA.	
Q3.3.8	is this process important?	esponsible for glutamate reuptake, and why rs (EAATs); it prevents excitotoxicity, which or ALS).	
Q3.3.9	Multiple Choice: Which receptor is closely associated with glutamate and is important for synaptic plasticity and memory formation?		
	(A) GABA receptor(C) Serotonin receptor	(B) NMDA receptor(D) Dopamine receptor	

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Answer: B.



Answer: Neuropeptides; example: Endogenous opioids

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Q3.3.21	Multiple Choice: What is the name of the endogenous cannabinoid neurotransmitter whose name means "bliss" in Sanskrit?				
	(A) Anandamide	(B) Cannal	bidiol		
	(C) Tetrahydrocannabinol (THC)	(D) 2-Arac	hidonoylglycerol (2-AG)		
	Answer: A.				
Q3.3.22	2 Short Answer: How are lipid-based neurotransmitters synthesized and stored? Answer: They are synthesized on demand and not stored in synaptic vesicles.				
Q3.3.23	Fill in the Blank: The gaseous ne	urotransmitter tl	hat is required for an erection is		
	Answer: Nitric Oxide (NO).				
Q3.3.24	4 Long Answer: Describe the study that addressed the question, "Does Botox decrease emotional experience?" Describe the sample, 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.				
Q3.3.25	5 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.				
Q3.3.26	Fill in the Blanks: Fill in the following spaces that describe the process of dopamine metabolism:				
	DA is broken down by converts it into Answer: Monoamine oxidase (MAO); O-methyltransferase (COMT); Homo	; Dihydroxypheny	vlacetric acid (DOPAC); Catechol-		
Q3.3.27	Short Answer: What were the rest Botox be used as a good thing?" Answer: 15% of placebo had a decrease in depression.	v	-		
~~~	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	······	·····		
3.4	•				
Q3.4.1	<b>Short Answer:</b> What does choliner <i>Answer:</i> Cholinergic refers to the new tors.	~	cetylcholine (ACh) and its recep-		

Q3.4.2	2 Short Answer: What are the four main functions of acetylcholine in the central			
	nervous system? $Answer: \mbox{ Learning and alertness, memory, REM sleep generation, and reward system.}$			
Q3.4.3	Multiple Choice: Who first	discovered acetylcholine in 1921?		
	(A) Otto von Loewy	(B) James Olds		
	(C) Neal Miller	(D) Peter Milner		
	Answer: A.			
Q3.4.4	covered acetylcholine.	owing describes the experiment of the scientist that dis-		
	part of the vagus nerve, whi	, put it in, and stimulated the ch slowed it. When he put the solution into another ed down, showing a chemical (ACh) was released. ution; parasympathetic; frog heart.		
Q3.4.5		runs parallel to the spinal cord. This is of your body responds at once.		
Q3.4.6	Fill in the Blank: The original	ginal name given to acetylcholine by its discoverer was		
	Answer: Vagusstoff.			
Q3.4.7	Short Answer: What are the two types of ACh receptors?  Answer: Nicotinic and muscarinic receptors.			
Q3.4.8	True or False: Acetylcholin pathetic branch of the autono Answer: True.	e is the primary neurotransmitter used in the parasymmic nervous system.		
Q3.4.9	Multiple Choice: Which of	the following is NOT a function of ACh in the CNS?		
	(A) Learning and alertness	(B) Memory		
	(C) REM sleep generation	(D) Pain modulation		
	Answer: D.			
Q3.4.10	Fill in the Blanks The following describes the synthesis and metabolism process of acetylcholine.			
	attaches to an acetate ion, of which is derived from  Then, transfers the acetate from the first chemical to choline,			
	which forms acetylcholine. Wl	hen it is time to be broken down, ACh is broken down by and The acetate is then broken down		
	and eliminated, while the lat	ter chemical is taken back up by and		
	· · · · · · · · · · · · · · · · · · ·	zyme-A (Acetyl-CoA); acetic acid; choline acetyltransterase (AChE); choline; active transport.		

- Q3.4.11 Multiple Choice: Which type of ACh receptor is ionotropic? (A) Nicotinic receptors (B) Muscarinic receptors (C) Both nicotinic and muscarinic receptors (D) Neither nicotinic nor muscarinic receptors Answer: A. Q3.4.12 Short Answer: Explain what the sympathetic chain is, and identify where it is located. Answer: Its 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. Q3.4.13 Fill in the Blank: The drug ______ is a direct antagonist of nicotinic receptors, causing paralysis. Answer: Curare. Q3.4.14 True or False: Atropine blocks nicotinic receptors and is derived from the plant known as belladonna alkaloids (deadly nightshade). Answer: False; it blocks muscarinic receptors. Q3.4.15 Short Answer: How does Botulinum Toxin interfere with acetylcholine function? Answer: It interferes with Ca²⁺ influx channels, preventing the release of ACh. Q3.4.16 Fill in the Blanks: Black widow spider venom causes of ACh, while (cobra and) krait venom _____ ACh receptors. Answer: Continual release; blocks. Q3.4.17 Multiple Choice: Which of the following is a reversible AChE blocker used to treat myasthenia gravis? (A) Deprenyl (Eldepryl) (B) Tetrabenazine (Xenazine) (C) Physostigmine (Antilirium) (D) Neostigmine (Prostigmin) Answer: D. Q3.4.18 True or False: Xanomelne (Cobenfy) crosses the blood-brain barrier and is used to treat the cognitive symptoms of Alzheimer's disease. Answer: False; Donepezil (Aricept)
- Q3.4.20 Multiple Choice: In the PNS, where are nicotinic receptors predominantly located?

Answer: False; they are agonists at low doses and antagonists at high doses.

Q3.4.19 True or False: Nicotinic receptors are antagonists at low does, but agonists at high

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	(A) Brain and spinal cord	(B) Neuromuscular junctions
	(C) Autonomic ganglia	(D) None of the above
	Answer: B.	
Q3.4.21	and a muscle fiber, where ACh is part of the sy cord, where preganglionic neurons Answer: The neuromuscular june muscle fiber, where ACh is released	is the synapse between a motor neuron is released to stimulate muscle contraction. The empathetic nervous system, located near the spinal synapse with postganglionic neurons. It is the synapse between a motor neuron and a distributed to stimulate muscle contraction. The paravertebral contraction is system, located near the spinal cord, where the postganglionic neurons.
Q3.4.22	Multiple Choice: In the sympat at the neuromuscular junction with	hetic nervous system, which neurotransmitter is used th smooth muscles and glands?
	(A) Acetylcholine	(B) Norepinephrine
	(C) Dopamine	(D) Serotonin

Answer: B.

Q3.4.23 Short Answer: Compare the neurotransmitters used in the parasympathetic nervous system versus the sympathetic nervous system.

Answer: In the parasympathetic nervous system, ACh is used at both the preganglionic

and postganglionic synapses. In the sympathetic nervous system, ACn is used at both the preganglionic and postganglionic synapses. In the sympathetic nervous system, ACh is used at the preganglionic synapse, while NE is used at the postganglionic synapse (except for sweat glands, which use ACh).

- Q3.4.24 Multiple Choice: Which of the following statements about acetylcholine in the autonomic nervous system is FALSE?
  - (A) ACh is the primary neurotransmitter in the parasympathetic branch
  - (B) ACh is used at preganglionic synapses in both sympathetic and parasympathetic branches
  - (C) ACh is used at postganglionic synapses to sweat glands in the sympathetic branch
  - (D) ACh is the primary neurotransmitter at the neuromuscular junction with smooth muscles in the sympathetic branch

Answer: D.

Q3.4.25 Fill in the Blank: In the somatic nervous system, ACh ______ the neuro-muscular junction.

Answer: Excites.

Q3.4.26	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.				
Q3.4.27	True or False: The Medial Septal Nucleu amygdala.  Answer: False. It primarily modulates the	es, which uses ACh, primarily modulates the hippocampus.			
Q3.4.28		anctions in the CNS, acetylcholine facilitates on through the actions of the			
	activating brain regions for this time period	od. ucleus (PPT); Laterodorsal Tegmental Nu-			
Q3.4.29		and are structures that ampus through the fornix. This is important of Diagonal Band.			
Q3.4.30	30 Multiple Choice: Which of the following correctly describes the neurotransmitter pathway in the parasympathetic nervous system?				
	(A) ACh at preganglionic synapse, ACh at postganglionic synapse				
	(B) ACh at preganglionic synapse, NE at postganglionic synapse				
	(C) NE at preganglionic synapse, ACh at	postganglionic synapse			
	(D) NE at preganglionic synapse, NE at postganglionic synapse				
	Answer: A.				
Q3.4.31	muscaria mushrooms. [NOT REQUIRED]	ryaks engage with ACh? o make a hallucinogenic drink from Amanita FOR FULL CREDIT:] The drink was used in similar to those of ACh (e.g., hallucinations,			
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3.5	•			
Q3.5.1	and	·	olamines are	
Q3.5.2	Answer: It is a sel	- ,	itor that increases dop	elegiline (Jumex)) work camine levels in the brain
Q3.5.3	Multiple Choice	: What is the precu	rsor for dopamine?	
	(A) Tyrosine	(B) L-DOPA	(C) Tryptophan	(D) Choline
	Answer: A.			
Q3.5.4	·		g enzyme in the synth	nesis of catecholamines i
	Answer: Tyrosine	Hydroxylase.		
Q3.5.5	precursor. Answer: Tyrosine	_	OPA by tyrosine hydr	esis from its amino acidoxylase, then L-DOPA i
Q3.5.6	word "tire" (spelle	ed "tyre") for its circ		ed British variation of the
Q3.5.7	Multiple Choice	: Which pathway is	involved in movement	and motor control?
	(A) Nigrostriatal s	ystem	(B) Mesocortical s	system
	(C) Mesolimbic sys	stem	(D) Tuberoinfundi	bular system
	Answer: A.			
Q3.5.8	We start at the _	, wh	nich then sends an inh	
	signal to the	, who ser	nds a reciprocal inhibit	sorysig

Answer: Striatium; GABA; substantia nigra; DA; GABA; globus pallidus; thalamus; primary motor cortex.
 Q3.5.9 Short Answer: List four symptoms of Parkinson's disease.
 Answer: Any four of: weakness, tremor at rest, muscle rigidity, problems with balance,

the _____, which causes voluntary movement.

abnormal gait, trouble learning.

nal back. Then, the first system sends an inhibitory ______ signal to the _____, who then excites

Q3.5.10	Short Answer: What is a drug that was used to lower blood pressure, but gave Parkinson's-like symptoms as a side effect? Answer: Reserpine (Raudixin)				
Q3.5.11	Fill in the Blank: The misfolded proteins found in the brains of people with Parkinson's disease are called Answer: Lewy Bodies.				
Q3.5.12	True or False: In Huntington's Chorea, there is too much GABA from the Striatum to the Substantia Nigra. Answer: False. In Huntington's Chorea, there is too little GABA from the Striatum to the Substantia Nigra.				
Q3.5.13	Long Answer: Explain how the MPTP incident in 1982 contributed to our understanding of Parkinson's disease. <i>Answer:</i> In 1982, young California heroin users thought they were using synthetic heroin (MPPP), but were actually exposed to MPTP. They instantly developed Parkinson's-like symptoms. MPTP is converted to MPP+ by the enzyme MAO, which damaged dopaminergic cells in the substantia nigra. This led to the development of animal models for Parkinson's research and potential treatment approaches, including MAO inhibitors like deprenyl (selegiline).				
Q3.5.14	Fill in the Blank: Methylphenidate (Ritalin) increases levels of and in the brain. Answer: Dopamine (DA); Norepinephrine (NE).				
Q3.5.15	5.15 Multiple Choice: Which system is primarily responsible for reward and reiment?				
	(A) Nigrostriatal sys	tem	(B) Mesocortical syst	em	
	(C) Mesolimbic syste	em	(D) Tuberoinfundibu	lar system	
	Answer: C.				
Q3.5.16	Short Answer: What neuropeptide, also called orexin, is involved in the regulation of sleep and wakefulness? Answer: Hypocretin.				
Q3.5.17	17 Multiple Choice: What neurotoxin led to the development of an animal model for Parkinson's disease?				
	(A) MPTP	(B) MPPP	(C) MPP+	(D) MAO	
	Answer: A.				
Q3.5.18	Fill in the Blank: to treat insomnia. Answer: Suvorexant		is an orexin rec	ceptor antagonist used	

Q3.5.19	True or False: The mesocortical system is involved in short-term memory, planning, and problem-solving. Answer: True.				
Q3.5.20	Multiple Choice: Which researchers disbrain areas could be rewarding rather that	covered that electrical stimulation of certain n aversive?			
	(A) Otto von Loewy	(B) James Olds and Peter Milner			
	(C) Neal Miller and Delgado	(D) Lateral hypothalamus researchers			
	Answer: B.				
Q3.5.21	Short Answer: What structure within the limbic system is considered the "pleasure center" of the brain? <i>Answer:</i> Nucleus accumbens.				
Q3.5.22	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.				
Q3.5.23	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.				
Q3.5.24	Short Answer: 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).				
Q3.5.25	Short Answer: Define choreoathetotic movements. Answer: Choreoathetotic movements refer to excessive movement disorders.				
Q3.5.26	Fill in the Blanks: The term refers to slow, continuous writhing movements, while (from the Greek word for "dance") refers to rapid, purposeless, involuntary movements. Answer: Athetosis; Choreic				
Q3.5.27	True or False: Both athetosis and choreic movements are characterized by too little movement. Answer: False. Both are characterized by too much movement.				
Q3.5.28	Short Answer: Where in the brain is hypocretin produced? <i>Answer:</i> In the lateral hypothalamus.				

Q3.5.29	Multiple Choice: Which drug is an orexin agonist and can be used to treat nar-colepsy?				
	(A) Suvorexant (Belsomra)	(B) Methylphenidate (Ritalin)			
	(C) TAK-994	(D) Hypocretin			
	Answer: C.				
Q3.5.30	Short Answer: 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).				
Q3.5.31	Fill in the Blank: is a neuropeptide involved in the regulation of sleep and wakefulness that is also known as orexin. Answer: Hypocretin				
Q3.5.32	Short Answer: 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.				
Q3.5.33	True or False: Nucleosides and neuropeptides are the same thing. <i>Answer:</i> False. Nucleosides are not the same as neuropeptides.				
Q3.5.34	Multiple Choice Spinal nerves leave the spinal cord and synapses in the paravertebra ganglion. This action is part of the system.				
	(A) Sympathetic	(B) Parasympathetic			
	(C) Somatic	(D) Central Nervous System (CNS)			
	Answer: A.				
Q3.5.35	Fill in the Blank What type of neurotra Answer: Acetylcholine (ACh).	ansmitter was used in the previous problem?			