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DENTAL PHARMACOLOGY EXAMINATION # 1

February 3, 2005

You have ONE (1) hour and 15 minutes to complete this examination. The examination contains 50 questions, each worth 2 points.

Answer all questions on the computer sheet provided; use a soft lead pencil. Be sure that you have correctly identified your answer sheet by PRINTING your name and social security number and correctly filling in the grid spaces. You may keep your exam booklet. Answers and grades will be posted on Blackboard.

This examination is being administered under the Honor Code of Temple University Dental School.

Quest	tions 1 - 50: Select the single, most appropriate answer.	
X.	Which of the following local anesthetics has a prolonged half-life in patients with liver disease compared to patients with normal hepatic function?	
* * *	A. Procaine B. Tetracaine C. Benzocaine D. Cocaine Mepivacaine A. A	
2.)	effects of norepinephrine?	Agenist .
· (?)	A. Metoprolot - B. Blocker Longs B. Acetylcholine - ACL aganist C. Amphetamine D. Phenylaphrine - A. Aganist E. Proprandol - P. P. Blocker	NO (NE \(\beta \)
13.	Which of the following drugs would be most useful for the relief of bronchoconstriction? By Agonist	1 H.R. 1 B.P.
	A. Terazosin B. Guanethidine C. Timolol D. Terbutaline E. Naloxone Albuterol, Ten	В, r.
JA!	Which of the following drugs <u>activates</u> alpha-2 adrenoceptors located presynaptically in the central nervous system?	
	A. Prazosin B. Albuterol C. Phentolamine D. Metoprolol E. Clonidine Blocker	↓ H.R
<i>J</i> 8.	All of the following statements apply to propranolol EXCEPT:	1 Bronchodilatio
, * * * .	Reduces elevated blood pressure Possesses low lipid solubility, therefore it does not enter the CNS Is a competitive antagonist of beta-1 and beta-2 receptors Is used for the prophylactic management of angina pectoris Is contraindicated in patients with bronchial asthma	Broncho Ernst

Frankhe-

Anti-Psychotic

All of the following statements apply to chlorpromazine EXCEPT:

Stimulates histamine receptors A.

Causes orthostatic hypotension T

Blocks muscarinic receptors

Increases the rate of secretion of prolactin D.

Is a member of the phenothiazine class of antipsychotic agents E.

All of the following drugs have a latency period of at least one week before the onset of their therapeutic action EXCEPT:

Tranyleypromine MAOT - ALTI-dep. A.

Imipramine - Anti - Pep B.

Lithium - Mania C.

Fluoxetine - Anti- Dep D.

Diazepam - Benzo -E

Which of the following drugs is an antidepressant that selectively inhibits serotonin re-uptake with minimal effect on norepinephrine re-uptake?

Desipramine A.

Diazepam B.

Protriptyline C.

Fluoxetine.

Mirtazepine

>5.5, R,I.

Which of the following drugs is most useful to control the manic phase of bipolar depression?

- Fluoxetine A.
- Doxepine B.
- Protriptyline C.
- Paroxetine D.
- Lithium

All of the following statements concerning buspirone are true EXCEPT:

Acts as a partial agonist at the serotonin (5HT_{1A}) receptor イ A.

Lacks muscle relaxant activity T B.

Is used in the management of generalized anxiety disorder T C.

Lacks sedative activity TD.

Is used in the management of panic attacks and psychosis (E.)

Non- Benzodiazepine

		the state of the s
· V.	Which of the following drugs would be the best choice for controlling seizures associated with status epilepticus?	
	seizures associated with status epitepheda.	
	A. Gabapentin	
	B. Topiramate	
.!	O Lorazepam	
	D. Haloperidol	4. 11. 11. 11. 11. 11. 11. 11. 11. 11. 11.
	E. Ethosuximide	
(12)	Which of the following antiepileptic drugs is most likely to cause hepat	ic failure?
17		
	A. \ Clonazepam	7.20
-12)	B. Valproic acid	Control of the second
	C. Whenytoin	
	(D) Carbamazepine	Contract Contract
	E. / Ethosuximide	
	Pilocarpine is most likely to produce which of the following effects?	
/3.		
	(A) Increase in salivary secretions	14 10
elle.	B. Increase in intraocular pressure	all Holes
2.00	Constipation	
	D. Tachycardia	
	E. Seizures	May ago was in
	Treatment of organophosphate intoxication should include which of t	he following
<i>y</i>	drugs?	S. Charles
	drugs:	14
	A. Physostigmine	
	B. Bethanechol	12-1-15 mm
	© Pralidoxime	A CALL SOL
	D. Pilocarpine	
	E. Propranolol	
	/ and the complete and the second	A Parket State Co.
J	8. Atropine is LEAST likely to produce	En alex
/	-A Mydrasis will dilation	100
	B. Tachycardia Will	
	-C. Dry mouth Will	
	(D) Bronchoconstriction	
	E. Fever	
THE REAL PROPERTY.	E. Fever won't Vasodilator	
~	Vasodilator	

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The major site of drug metabolism by the cytochrome P450 enzyme system is Heroprotein

- Plasma
- Lungs
- Liver
- Gastrointestinal tract
- Kidney

A 45-year-old man enters the emergency room with an overdose of a weak acid (pKa=6). The drug is excreted by the kidney. Approximately what percent of drug is passively reabsorbed when the pH of the tubular Log NI = pka-ph I+ 1-10% filtrate and urine is 5?

- 1%
- 10%
- 50% C.
- 90%
- 100%

All of the following statements about drug excretion are true EXCEPT:

- An inhalational drug (e.g. nitrous oxide) is excreted through
- Lipid soluble drugs can enter breast milk, causing effects in a nursing infant
- Drugs with a molecular weight greater than 450 can be excreted in the bile
- Both free and protein-bound drugs in the plasma are susceptible to glomerular filtration
- Water soluble drugs and metabolites are susceptible to renal excretion

Drugs A and B have similar pharmacokinetic parameters, except that Drug A is 95% bound to plasma proteins (Drug B is not bound). Which of the following statements is most likely true?

- Drug W is more slowly absorbed
 - Drug A is more rapidly distributed to its site of action
 - Drug A has a smaller volume of distribution F.C.
- Dring A is more rapidly metabolized CD.
- Drug A is more rapidly excreted F.E.

All of the following responses are a result of activation of cholinergic muscarinic receptors EXCEPT: Decrease of heart rate T A. Constriction of the pupil (miosis) T B. Increased gastrointestinal tract motility and tone T Relaxation of bronchial smooth muscle Dilation of erectile tissue E. Immediately following intravenous injection of a drug, the plasma concentration was measured as 300 ng/ml. Four hours later, the plasma concentration was determined to be 75 ng/ml. Assuming first-order kinetics, which of the following values represents the half-life of the drug? 30 minutes A.-60 minutes B. 2 hours 4 hours 6 hours E. A drug with an elimination half-life of 2 hours is given by intravenous infusion of 5 μ g/min. The minimum time needed for the drug to achieve 95% of its steady-state value is which of the following? 2 hours A. 4.3 hours B. 8.6 hours 18 hours Halothane - causes E. 24 hours Identify the drug most useful to treat malignant hyperthermia. Dantrolene Atracurium C. Diazepam Succinylcholine D. E. Thiopental Selegiline is used to treat Parkinsonism. It acts by Acting as an agonist to the D₁ dopamine receptor Increasing dopamine activity by inhibiting MAOb activity Decreasing acetylcholine activity by blocking muscarinic receptors Inhibiting dopamine decarboxylase activity in the periphery Stimulating dopamine release E.

All of the following effects occur with acute or chronic alcohol ingestion Korsakoff's psychosis Poe! -Decrease in high density lipoproteins 7 Cutaneous vasodilation Doe's Increased fat production in the liver Doc 5 Inhibition of antidiuretic hormone Does Which benzodiazepine is preferred as a hypnotic because of its relative short duration of action? Chlordiazepoxide A. Flurazepam B. Diazepam C. ' Nitrazepam Triazolam All of the following drugs achieve muscular relaxation via action in the central nervous system EXCEPT: Dantrolene Mephenesin A. Diazepam B. Baclofen C. ' Dantrolene Chlordiazepoxide Sinemet, a combination of levodopa and carbidopa, is used to treat parkinsonism. Carbidopa is important in this preparation because it Releases stores of endogenous dopamine Inhibits the breakdown of levodopa to dopamine in the periphery Enhances the breakdown of levodopa to dopamine in the central nervous Is a dopamine receptor agonist D. Is a dopamine receptor antagonist acting in the area postrema to E. prevent nausea Phase II block at the neuromuscular junction occurs when succinylcholine Releases excessive amounts of histamine Is used in combination with d-tubocurarine Is used in a patient who has recently engaged in excessive museular activity Is used in a patient who has had excessive bed rest Is used for a prolonged period and in excessive amounts

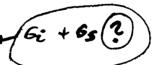
For questions 30 and 31:

Agent	Blood/Gas P.C.	MAC
71gont		•
Desflurane	0.42	6
Sevoflurane	0.69	2.0
Enflurane	1.80	1.7
Isoflurane	1.40	1.4
Halothane	2.30	0.75

- Which of the following drugs has the most rapid rate of induction?
 - Desflurane
 - Sevoflurane
 - Enflurane
 - Isoflurane D.
 - Halothane E.
- Which of the following drugs is the LEAST potent?
 - Desflurane
 - Sevoflurane
 - Enflurane C.
 - Isoflurane D.
 - Halothane



- Which of the following enzymes is directly regulated by the guanine nucleotide binding protein Gq?
 - Protein kinase A A.
 - Protein kinase C B.
 - Phospholipase C
 - Adenylyl cyclase
 - Guanylyl cyclase E.





A drug that binds to the receptor for epinephrine, without activating the receptor is called

- A physiologic antagonist
- A competitive antagonist
- A partial agonist C.
- A reverse agonist D.
- A chemical antagonist E.

A live of the state of interest and interest and interest and protection	
34. An ultra-short acting barbiturate administered intravenously for induction	and the same of th
of anesthedia:	aka Mada wasan
	N. O. W. Ange
A. Nitrous oxide	
B. Thopental	
C Midazolam	
D. Enflurane	
E. Propofol	
35. A benzodiazepine that enhances GABA activity, produces sedation, skele	tal
muscle relaxation and amnesia:	
	The second second
A. Nitrous oxide	
B. Thiopental - Barb	
© Midazolam	
	Market
D. Enflurane General Anes E. Desflurane	
All of the following statements about local anesthetics are true EXCEPT	
20. Am of the logic was	The state of
TA. Benzocaine can only be used topically	
T B. Local anesthetics block sodium channels	A CONTRACTOR OF THE SECOND
C. The charged form of lidocaine is important for penetrating neurona	1
cell membranes, while the uncharged form is the most active at its	A STATE OF THE STA
binding site	A Long When A Vision
Epinephrine is included in lidocaine preparations to decrease the	and the second
absorption of lidocaine	
E. Seizures, induced by lidocaine, can be treated with diazepam	
E. Scizures, induced by indocame, can be weated with diazopain	• 4
37. All of the following statements apply to phenylephrine EXCEPT:	∠, agon's+
All of the following statements apply to phenylephine EXCELT:	
Reduces secretions by inhibiting parasympathetic stimulation	> Mydriasis
B. Is a direct-acting alpha adrenergic receptor agonist	11 +
	WITHOUT
TC. Reverses hypotension during anesthesia	without Cycloplegi
D. Is added to local anesthetic solutions to prolong anesthesia	
TE. Constricts small vessels in the nasal mucosa	1.00
(a) All Cil Cil :	DT.
(38) All of the following statements are true concerning amphetamine EXCE	P1:
T A. Tolerance develops to the appetite suppressant effects	
→ B. Is used to reverse the sleepiness of narcolepsy	
7 -> C. Is a Schedule III controlled substance	
D.) Prevents the release of dopamine from nerve endings	
in the central nervous system	
T E. Is used in the management of attention-deficit hyperactivity disord	er

note and	*. *	Benz
<i>36</i> .	Diaz	repam is useful for all of the following indications EXCEPT:
Ager	Δ.	Musculo-skeletal disorders Obsessive compulsive disorders
Ab 1. 143	B	Obsessive-compulsive disorders
Te	C.	Generalized anxiety disorder
Se Se	D.	Acute alcohol withdrawal
	Ε.	Oral sedation in dentistry
AN	Flor	nazenil reverses the behavioral depressant effects of which one of
Benzogonist		following drugs?"
		Diamon
W. of the Ban	(A)	Diazepam Manufication of the control
	В.	Morphine Newstart at the least of the least
7.4	C.	Phenobarbital
A STATE OF THE STA	. D.	Haloperidol
PL	E.	Thioridazine
<i>y</i> 11.	pres	ich of the following drugs is most likely to increase heart rate and blood sure, increase gastric acid secretion, and cause seizures at high doses?
	A.	Bethanechol
	$^{\circ}$	Nicotine
	C.	Propranolol
	D.	Edrophonium '
	E.	Atropine
) <u>4</u>	. Dru	gs that block muscarinic receptors are most useful in the treatment of
	YAA	Gastrointestinal spasm
•	NB.	Hypertension
	N C.	Atony of the bladder
	۷D.	Xerostomia
	۸E.	Myasthenia gravis
(43)) All	of the following statements are true about routes of administration EXCEPT:
11 -	T A	A drug administered intravenously has a bioavailability of 100%
W 60-	→ R	Sublingual administration leads to delay in onset of action,
1	, U.,	compared to oral administration
	TC.	Inhalation can lead to central nervous system effects
{	7 D.	Transdermal administration is useful for producing a systemic effect
	1 E	Inhalation is useful for a local effect in the lungs

Increased cardiac contractility and conduction, lipolysis, glycogenolysis, and relaxation of bronchial smooth muscle are due to activation of which of the following classes of receptors? Cholinergic muscarinic receptors Cholinergic nicotinic receptors Alpha adrenergic receptors Beta adrenergic receptors Serotonin receptors When alcohol and disulfiram are present in the body, disulfiram inhibits resulting in nausea and vomiting. Alcohol dehydrogenase Acetaldehyde Mixed function oxidase Aldehyde oxidase Alcohol oxidase Aldehyde dehydrogenase Which of the following compounds blocks acetylcholine release? Botulinum toxin Chlordiazepoxide C. Baclofen Black widow spider toxin D. Ediophondrum E. Diazepam Which of the following drugs reverses the action of the curare-type drugs (non-depolarizing neuromuscular blockers)? Isoflurane Diazepam - 13 en Z

Neostigmine ACL E In

Propranolol Bus B

Halothane

For question 48:

Drug Dose	Percent Responding to Drug X	Percent responding to Drug Y	
0.1 mg	1	10	
0.3 mg	5 .	20	
1.0 mg	10 *	50	
3.0 mg	50∗	68	
10.0 mg	70	92	
30 mg	90	100	
100 mg	100	100'	

- 48. Which of the following statements can be concluded from the table above?
 - A. Drug X is safer than Drug Y N
 - B. Drug Y is more effective than Drug X No
 - C. The two drugs act on the same receptors N Drug X is less potent than Drug Y
 - E. The therapeutic index of Drug Y is 10 ∧
- Which of the following best describes the signaling involved in steroid hormone action?
 - A. Action on a tyrosine kinase-linked receptor
 - B. Diffusion across the cell membrane followed by action on soluble guanylyl cyclase
 - —C. Activation of a G-protein that inhibits adenylyl cyclase
 - —D. Opening of a transmembrane ion channel
 - E. Diffusion into the cell and activating a receptor to allow binding to DNA
- Most rapid acting inhalational anesthetic, but low potency; useful in many dental procedures:
 - A.) Nitrous oxide
 - B. Thiopental
 - C. Midazolam
 - D. Enflurane
 - E. Propofol

