## Veterinary Pharmacology and Toxicology

Q.	No. 1	The ratio of total concentration (unionized+ionized) of a weak acidic drug
		(pK <sub>a</sub> =4.4) between plasma (pH=7.4) and gastric juice (pH=1.4) at equilibrium
		will be:
		a 10:1
		b 100:1
		c 1000:1
		d 10000:1
Q.	No. 2	If the elimination rate constant $(\beta)$ of a drug is 0.173, its half-life will be:
		a 12 hour
		b 8 hour
		c 1 hour
		d 4 hour
Q.	No. 3	Ocuserts are the drug delivery devices placed in
	5	a Conjuctival sac
	15	b Vaginal cavity
	1- 1	c Uterus
	10 M	d Nasal cavity
Q.	No. 4	Majority of drugs cross biological membranes primarily by:
-1	0	a Passive diffusion
- [	5	b Facilitated diffusion
9	שו	c Pinocytosis
		d Active transport
Q.	No. 5	A drug having half-life of 4h will be eliminated more than 99% in hrs if
	0	it follows first order kinetics.
	6	a 07
	0	b 14
	1	c 21
		d 28
Q.	No. 6	If the total amount of a drug present in the body at a given moment is 2.0 g and
		its plasma concentration is 25 μg/ml, then its volume of distribution will be:
		a 100 L
		b 80 L
		c 60 L
		d 50 L
Q.	No. 7	Cardiac glycoside ouabain is mainly obtained from the leaves of
_		a Digitalis purpurea
		b Digitalis lanata
		c Strophanthus kombe
		d Strophanthus gratus

		a	Mixed order kinetics
		b	First order kinetics
		c	Zero order kinetics
		d	Dose dependent kinetics
Q.	No. 9	In the	e resting state, G protein in a GPCR exists as an
		a	$\alpha$ - $\beta$ - $\gamma$ trimer with GTP occupying the site on $\alpha$ -subunit
		b	$\alpha$ - $\beta$ - $\gamma$ trimer with GDP occupying the site on $\alpha$ -subunit
		c	$\alpha$ - $\beta$ dimer with GDP occupying the site on $\alpha$ -subunit
		d	$\beta$ - $\gamma$ dimer with GDP occupying the site on $\beta$ -subunit
Q.	No. 10	In an	anaesthetized dog, repeated intravenous injection of ephedrine at same
		dose a	and interval shows the phenomenon of:
		a	Anaphylaxis
	3	b	Tachyphylaxis
	5	c	Idiosyncrasy
	100	d	Drug resistance
Q.	No.11	Drug	s producin <mark>g allergic reactions generally</mark> act as:
	1º MA	a	Complete antigens
- /	2	b	Haptens
Ĺ	0	c	Antibodies
I	5	d	Mediators  wing are ionotropic receptors except:  GABA <sub>A</sub> receptor
Q.	No. 12	Follo	wing are ionotropic rec <mark>ep</mark> tors except:
	0	a	GABA <sub>A</sub> receptor
4	F I	b	NMDA-receptor
	0	c	5-HT <sub>2</sub> receptor
	6	d	Glycine receptor
Q.	No. 13	Exam	ple of Ig E mediated all <mark>er</mark> gic reactions is
	1/2	a	Anaphylais
		b	Contact dermatitis
		c	Acute rheumatic fever
		d	Serum sickness
Q.	No. 14	A dru	ig usually attains steady state concentration after its repeated
		admi	nistration for
		a	4.3 half lives
		b	2.3 half lives
		c	6.3 half lives
		d	8.3 half lives

Constant or fixed amount of drug is eliminated per unit of time in

a cGMP b cAMP c DAG d IP3 and DAG  Q. No. 16 Therapeutically undesired but unavoidable pharmacodynamics effect of a drug is called a Idiosyncrasy b Toxic effect c Side effect d Intolerance  Q. No. 17 Identify the incorrect pair of agonist-antagonist acting on the same receptor: a Detomidine: Yohimbine b Dobutamine: Atenolol c Oxotrimorine: Trimethophan d Hexamethonium: Epibatidine  Q. No. 18 The most commonly occurring conjugation reaction for drugs or their metabolites is: a Glucuronidation b Acetylation c Methylation d Glutathione conjugation Q. No. 19 The ratio between LD1 and ED22 i.e., LD1/ED22 is called: a Therapeutic index c Protective index d Certain safety factor  Q. No. 20 Which of the following Phase I enzymes has major share in drugs metabolism? a CYP2D6 c CYP3A4 d CYP2D6 Q. No. 21 The elimination of alcohol follows the a Zero order kinetic b First order kinetic c Second order kinetic	Q.	No. 15	Stimulation of M <sub>1</sub> type of muscarinic receptors results in formation of
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<ul><li>b First order kinetic</li><li>c Second order kinetic</li></ul>	ζ,		
c Second order kinetic			
d Third order kinetic			d Third order kinetic

Q.	No. 22	Drug of choice for treatment of anaphylactic shock is:
		a Salbutamol
		b Noradrenaline
		c Adrenaline
		d Phenylephrine
Q.	No. 23	Drugs producing allergic reactions generally act as:
		a Complete antigens
		b Haptens
		c Antibodies
		d Mediators
Q.	No. 24	Proton pump inhibitors inactivates Na+- K+ ATPase by binding with its amino
		acid:
		a Methionine
	3	b Cysteine
	5	c Proline
	13	d Glycine
Q.	No. 25	Metoclopramide produces its action on GIT by following mechanisms except.
	C M	a D <sub>2</sub> - receptor antagonist
F	7	b 5-HT <sub>4</sub> - receptor antagonist
D	5	c 5-HT <sub>3</sub> - receptor antagonist
I	2	d Cholinomimetic effect
Q.	No. 26	cAMP is the second messenger in following receptor types except:
		a β <sub>1-</sub> adrenergic receptor
4	5	b β <sub>2</sub> -adrenergic receptor
	0	c α <sub>1-</sub> adrenergic receptor
	6	d α <sub>2-</sub> adrenergic receptor
Q.	No. 27	Nitric oxide mediates its action by activation of
	1/4	a Adenylyl cyclase
		b Phospholipase A
		c Phospholipase C
		d Guanylyl cyclase
Q.	No. 28	Glutathion is comprised of the following amino acids except:
		a Glycine
		b Glutamate
		c Cysteine
		d Methionine

Q. No. 29	Monitoring plasma drug concentration is useful while using:
	a Antihypertensive drugs
	b Levodopa
	c Lithium carbonate
	d MAO inhibitors
Q. No. 30	Ketamine exerts majority of its CNS action by
	a Inhibiting NMDA receptor
	b Activating NMDA receptor
	c Inhibition of μ-opioid receptor
	d Inhibition of δ-opioid receptor
Q. No. 31	The receptor transduction mechanism with the fastest time-course of response
	effectuation is:
	a Adenylyl cyclase-cyclic AMP pathway
2	b Phospholipase C-IP3/ DAG pathway
1	c Intrinsic ion channel operation
15	d Protein synthesis modulation
Q. No. 32	Which of the following drugs suppress the level of aldosterone?
(C)	a Captopril
5	b Furosemide
10	
E	d Imipramine
Q. No. 33	After I.V. drug administration, elimination of a drug depends on
	a Lipid solubility
6	b Volume of distribution
The same	c Clearance
8	d Drug concentration
Q. No. 34	When two different chemicals act on two different receptors and their response
Q. 110.01	is opposite to each other on the same cell, this is called as:
	a Non-competitive antagonism
	b Competitive antagonism
	c Chemical antagonism
	d Physiological antagonism
Q. No. 35	Acetyl Co A required for biosynthesis of acetylcholine is derived mainly from
Q. 110.05	a Choline
	b Pyruvate
	c α-Lipoic acid
	d Glutathione
	u GiutatiiiOiic

Q. No. 36	Acetylcholine storage vesicle in cholinergic nerve terminal has diameter of
	about
	a $3-6 A^0$
	b $30-60 \text{ A}^0$
	c $300-600 \text{ A}^0$
	d $3000-6000 \text{ A}^0$
Q. No. 37	An increase in insulin secretion from pancreatic $\beta$ cells occurs by
	a $\alpha_{2-}$ adrenergic receptor stimulation
	b $\alpha_{1-}$ adrenergic receptor stimulation
	c β <sub>1-</sub> adrenergic receptor stimulation
	d β <sub>2-</sub> adrenergic receptor stimulation
Q. No. 38	Precursor for synthesis of endogenous catecholamines is
	a Proline
6	b Serine
5	c Alanine
100	d Tyrosine
Q. No. 39	Identify the wrong pair
10	a Ethacrynic acid: Na <sup>+</sup> -Cl symport inhibitor
R	b Dichlorphenamide: Carbonic anhydrase inhibitor
10	b Dichlorphenamide: Carbonic anhydrase inhibitor c Amiloride: Na <sup>+</sup> channel inhibitor d Canrenone: Mineralocorticoid Receptor antagonist Cholinomimetics are not used in
15	d Canrenone: Mineralocorticoid Receptor antagonist
Q. No. 40	Cholinomimetics are not used in
0	a Glaucoma
- B	b Myasthenia gravis
0	c Post operative atony
6	d Partial heart block
Q. No. 41	The following is a competitive antagonist of GABA but a noncompetitive
1	antagonist of diazepam:
	a Picrotoxin
	b Muscimol
	c Flumazenil
	d Bicuculline
Q. No. 42	Botulin toxin acts by
	a Increasing secretion of Ach
	b Increasing synthesis of Ach
	c Inhibiting Ach release
	d Decreasing uptake of ACh

Q. No. 43	Bromocriptine is
	a An α- adrenergic receptor agonist
	b A dopaminergic receptor agonist
	c A β <sub>1</sub> - adrenergic receptor agonist
	d A $\beta_2$ - adrenergic receptor agonist
Q. No. 44	Identify correct order of potency against β-adrenergic receptor:
	a Epinephrine > Norepinephrine > Isoprenaline
	b Norepinephrine>Epinephrine > Isoprenaline
	c Epinephrine > Isoprenaline > Norepinephrine
	d Isoprenaline > Epinephrine > Norepinephrine
Q. No. 45	Which of the following inhibits the uptake and storage of norepinephrine by the
	storage vesicle?
	a Thioridazine
6	b α- methyl p-tyrosine
5	c Ondansetron
15	d Desipramine
Q. No. 46	Trientine has a potent chelating action against
10 M	a Iron
P	b Copper
10	c Lead
15	d Arsenic  A centrally acting antihypertensive drug is  a Atenolol
Q. No. 47	A centrally acting antihypertensive drug is
0	a Atenolol
· b	b Prazosin
0	c Clonidine
6	d Propranolol W. J. W. W. J. W. W. J. W. W. J. W
Q. No. 48	Which of the following drugs is useful to dissolve gall bladder stone
Y	a Clofibrate
	b Chenodeoxycholic acid
	c Lactulose
	d Lithocholic acid
Q. No. 49	Topiramate is used clinically as
	a Antiepiletic
	b Antiemtic
	c Tocolytic
	d Diuretic

Q.	No. 50	Chemically, kaolin is
		a Hydrated calcium carbonate
		b Purified carbohydrate
		c Activated wood charcoal
		d Hydrated aluminium silicate
Q.	No. 51	Epinephrine produces all of the following effects except:
		a Bronchodilation
		b Hyperglycemia
		c Mydriasis
		d Decrease in oxygen consumption
Q.	No. 52	Indicate the sympathomimetic drug, which is used in a hypotensive emergency:
		a Xylometazoline
		b Ephedrine
		c Terbutaline
	5	d Phenylephrine
Q.	No. 53	Which of the following drugs is a reversible nonselective α, β antagonist?
	15- 11	a Phentolamine
	10 M	b Labetalol
1	2	c Metoprolol
	9	d Propranolol
Q.	No. 54	Carbon monoxide is used as euthanizing agent at concentration of%.
4	0	a 0.5-1
	0	b 2-3
4	B I	c 4-6
	0.	d 8-10
Q.	No. 55	Which of the following cholinomimetics is indirect-acting
	0	a Lobeline
	L	b Carbachol
		c Edrophonium
		d Pilocarpine
Q.	No. 56	Which of the following drugs is both a muscarinic and nicotinic blocker?
		a Atropine
		b Hexamethonium
		c Benztropine
		d Succinylcholine
Q.	No. 57	Montelukast produce its anti-allergic and bronchodilatory action by
		a Inhibiting H <sub>1</sub> receptor
		b Inhibiting HT <sub>1</sub> receptor
		c Inhibiting TXA <sub>2</sub> receptor
		d Inhibiting CysLT <sub>1</sub> receptor

		a Gallamine
		b Physostigmine
		c d-Tubocurarine
		d Pralidoxime
Q.	No. 59	CNS stimulation action of Nikethamide has target cells mainly in
		a Pons
		b Medulla
		c Mid brain
		d Cerebral cortex
Q.	No. 60	b Medulla c Mid brain d Cerebral cortex  Antiarrhythmic drug is a Rosiglitazone b Misoprostol c Acetyl cysteine d Lidocaine
		a Rosiglitazone
		b Misoprostol
	3	c Acetyl cysteine
	5	d Lidocaine
Q.	No. 61	Acetylcholine is not used therapeutically because
	15- 11	a Orally ineffective
	1º M	b Rapidly excreted
- /	2	C High plasma bound
I	0	d Rapidly degraded Rapidly degraded
Q.	No. 62	Drug that acts as selective 5-HT uptake inhibitor is a Fluoxetine
4	0	a Fluoxetine
		b Reserpine A Section 1997
	F I	c Methysergide
	0	d Cimetidine
Q.	No. 63	Chemical mediators in the nociceptive pathway are all of the following except:
	0	a Kinins
	1/4	b Enkephalins
		c Prostaglandins
		d Substance P
Q.	No. 64	Following are neurokinin1 (NK1) recptor antagonists except:
		a Talnetant
		b Vestipitant
		c Morapitant
		d Aprepitant

Atropine poisoning can be best antagonized by

Q. N	No. 65	Which of the following cholinomimetics activates both muscarinic and nicotini
		receptors? a Lobeline
		a Lobeline b Pilocarpine
		271
O N	T- ((	
Q. N	No. 66	Penicillins inhibit the cross linkage betweenamino acids of two
		adjacent glycopeptides to inhibit bacterial cell wall synthesis.
		a Glycine and D-Alanine
		b Glutamate and D-Alanine
		c Glycine and Proline
		d Glycine and L-Lysine
Q. N	lo. 67	Identify the correct order of duration of action of local anaesthetics:
	,cc	a Bupivacaine > Lidocaine > Chlorprocaine > Benoxinate
	D	b Bupivacaine > Chlorprocaine > Lidocaine > Benoxinate
	130	c Bupivacaine > Benoxinate > Lidocaine > Chlorprocaine
//	- 1	d Benoxinate > Chlorprocaine > Lidocaine > Bupivacaine
Q. N	lo. 68	Ionizable g <mark>roup of local anesthetics is respon</mark> sible for:
100		a The potency of local ane <mark>st</mark> hetics
10		b The duration of action local anesthetics
1	. [1]	c The ability of local anesthetics to diffuse to the site of action d The toxicity of local anesthetics
40		d The toxicity of local anesthetics
Q. N	lo. 69	The δ-opioid rece <mark>ptors have greatest affinity for :</mark>
-1	5	a Endorphins
	0.	b Enkephalins
	5	c Etorphine
	0	d Morphine
Q. N	lo. 70	Identify the wrong statement on SAR of barbiturates:
		a Both H-atoms at C-5 be replaced with alkyl or aryl group for CNS depressan
		activity
		b Addition of more than 9 carbon chain at C-5 leads to more depressant activity
		c Unsaturable carbon chain results in short duration
		d Replacement of O-atom at C-2 by sulphur results high potency but short
		duration
Q. N	No. 71	Identify the correct order of MAC values of inhalant anaesthetics:
		a Chloroform <halothane< isoflurane<="" methoxyflurane<="" td=""></halothane<>
		b Chloroform <halothane <="" isoflurane="" methoxyflurane<="" td=""></halothane>

Methoxy flurane < Chloroform < Halothane < Isoflurane

Methoxyflurane < Halothane < Chloroform < Isoflurane

c

d

Q.	No. 72	Indicate the drug belonging to M1-cholinoreceptor blockers
		a Cimetidine
		b Omeprazole
		c Pirenzepine
		d Ranitidine
Q.	No. 73	The drug used as aquaretic is
		a Triamterene
		b Bendroflumethiazide
		c Spironolactone
		d Conivaptan
Q.	No. 74	One Grey (Gy) unit of radiation is equivalent to
		a 1 Rad
		a 1 Rad b 10 Rads c 100 Rads d 1000 Pads
	ුණි	c 100 Rads
	5	d 1000 Rads
Q.	No. 75	Which of the following cholinomimetics is most widely used for paralytic ileus
	15- 11	and atony of th <mark>e urinary bladder?</mark>
		a Lobeline
1	2	b Neostigmine
-		c Pilocarpine
4		d Echothiophate
Q.	No. 76	c Pilocarpine d Echothiophate Indicate the agent, which interferes with GABA binding
	0	a Flurazepam
4	P	b Bicuculline
	0	c Thiopental
	D.	d Zolp <mark>i</mark> dem W Zolpidem
Q.	No. 77	PGF <sub>2a</sub> mediates is action by activation of:
	1	a PLC-IP3/DAG pathway
		b DNA-mRNA-Protein pathway
		c AC-cAMP pathway
		d GC-cGMP pathway
Q.	No. 78	Aglycon of cardiac glycosides is chemically resembles with the structure of
		a Steroidal hormone
		b Pyridine
		c Pyrimidine
		d B-lactam ring

Q.	No. 79	Domperidone produces antiemetic and prokinetic action by stimulating:
		a D1 Receptors in small intestine
		b D2 receptors in gastric mucosa
		c D2 and D3 receptors in CTZ
		d D2 receptors in gastric mucosa and D2 and D3 receptors in CTZ
Q.	No. 80	Which of the following antibiotics is not readily destroyed by penicillinase
		enzymes?
		a Phenoxymethylpenicillin
		b Ticarcillin
		c Flucloxacillin
		d Ampicillin
Q.	No. 81	Which of the following drugs is a COX inhibitor belonging to pyrazolone?
		a Celecoxib
		b Rofecoxib
	1	c Nimesulide
	100	d Phenylbutazone
Q.	No. 82	The long term administration of a thiazide diuretic may also require the
-	C 11	administration of
ŀ	7	a Potassium
D	5	b Sodium
	S I	c Calcium
4	5	d Bicarbonate
Q.	No. 83	Tranquilizer chlorpromazine produce violent incoordination and excitement in:
4	8	a Camel
	-	b Horse
	8	c Cat
	Y (2)	d Dog
Q.	No. 84	Following antidepressants are selective 5-HT reuptake inhibitor except:
		a Doxepine
		b Trazodon
		c Fluoxetine
		d Amoxapine
0.	No. 85	Identify the incorrect pair:
χ.	1,000	a Arsenite : Aquaglyceroporin channels
		b Lead : Voltage gated Ca <sup>2+</sup> channels
		c Thallous ions : Na <sup>+</sup> & K <sup>+</sup> channels
		d α- Amantin : Na-dependent bile acid transport
		a a ramanini . Tra-dependent one acid transport

Q.	No. 86	is a membrane stabilizing agents:							
		a Zileutin							
		b Zafirlucast							
		c Sodium cromoglycate							
		d Montelucast							
Q.	No. 87	Which of the following drugs is used for systemic and deep mycotic infection							
		treatment:							
		a Co-trimoxazol							
		b Griseofulvin							
		c Amphotericin B							
		d Nitrofungin							
Q.	No. 88	5-HT causes platelet aggregation by activatingreceptors.							
		a $5HT_{2A}$							
	5	b 5HT <sub>2B</sub>							
	8	c 5HT <sub>1A</sub>							
	100	d 5HT <sub>1B</sub>							
Q.	No. 89	Identify the antimycobacterial drug belonging to class of antibiotics:							
	C 11	a Isonia <mark>zid</mark>							
	7	b PAS							
Î	U	c Ethambutol							
	5	d Rifampin							
Q.	No. 90	Fluoroquinolones are derived by addition of F-atom atposition of							
7.0		structure of quinolone.							
	8	a 3rd							
	-	b 5th							
	5	c 6th							
	Y (0)	d 7th							
O.	No. 91	Identify the incorrect pair:							
•		a Eukaryote : Topoisomerase IV							
		b Cephalosporins : Thrombocytopenia							
		c Fialuridine : Synthesis of mitochondrial DNA							
		d Fluoroacetate : TCA cycle.							
O.	No. 92	Which of the following is a thiamine antagonist anticoccidial agent::							
٧.	100,7=	a Monensin							
		b Halofuginone							
		c Clopidol							
		d Amprolium							
		4 / Impronum							

Q.	No. 93	Identify the antiviral drug inhibiting viral reverse transcriptase for its action:
		a Zidovudine
		b Vidarabine
		c Rimantadine
		d Gancyclovir
Q.	No. 94	Drug used for toxoplasmosis treatment is:
		a Chloroquine
		b Tetracycline
		c Suramin
		d Pyrimethamine
Q.	No. 95	c Suramin d Pyrimethamine  Colchicine is used to treat a Hyperlipidemia b Jaundice c Goitre
		a Hyperlipidemia
		b Jaundice
	. 3	c Goitre
	5	d Gouty arthritis
Q.	No. 96	Following antimicrobials bind with bacterial 50 S ribosome for their action
	15- 11	except:
	F 11	a Thiamphenicol
- 1	2	b Tetracycline
	U I	c Erythromycin
1		d Lincomycin  The amino acid not required for synthesis of Park neucleotide:
Q.	No. 97	The amino acid not required for synthesis of Park neucleotide:
	0	a Alanine
	P	b Glutamate
	0	c Serine
	DE	d Lysine W. L. Lysine
Q.	No. 98	Nucleated RBCs in very large numbers are found in the toxicity of
	1/4	a Iron
		b Lead
		c copper
		d Cadmium
Q.	No. 99	Ideal copper and molybdenum ration in the diet of animals
		a 6: 1
		b 4:1
		c 3.1
		d 2:1

Q. No. 100	Following are anti-androgens except:
	a Cyproterone
	b Flutamide
	c Finasteride
	d Clomiphene
Q. No. 101	Following anthelmintics are the uncouplers of electron transport except:
	a Rafoxanide
	b Albendazole
	c Niclosamide
	d Hexachlorophene
Q. No. 102	A blood lead concentration reported as 1g / dl is the same as:
	a 1ppm
	b 10 ppm
á	c 100 ppm
5	d 10000 ppm
Q. No. 103	Toxicity ofresembles with thiamine deficiency in equines.
15- 11	a Lantan <mark>a camara</mark>
10	b Strychnus nuxvomica
R	c Atropa belladona
10	d Pteridium aquilinum
Q. No. 104	Identify the wrong pair :
40	a Arsenic - Dimercaprol
0	b Cyanide - Sod. thiosulphate
· P	c Lead - Ca EDTA
0	d Carbaryl - 2-PAM-
Q. No. 105	Saxitoxin causes blockade of inward current of:
0	a Na <sup>+</sup> channels
L	b Ca <sup>2+</sup> channels
	c K <sup>+</sup> channels
	d Cl <sup>-</sup> channels
Q. No. 106	Identify wrong statement regarding nitrate toxicity.
	a Ferric form of Hb is converted ferrous form of Hb
	b MetHb is not able to transport oxygen
	c Methylene blue is antidote for nitrate toxicity
	d Methylene blue is a potent oxidizing agent
Q. No. 107	Which of the following is not an ingredient of universal antidote
	a Activated vegetable charcoal
	b Magnesium oxide
	c Tannic acid
	d Acetyl cysteine

Q.	No. 108	"Turkey X Disease" occurred due intoxication of
		a Aflatoxin
		b Ochratoxin
		c Ergotoxin
		d T-2 toxin
Q.	No. 109	Chemical mediators in the nociceptive pathway are all of the following except
		a Enkephalins
		b Kinins
		c Prostaglandins
		d Substance P
Q.	No. 110	c Prostaglandins d Substance P  Identify the incorrect pair: a Selenium: Blind stagger b Molybdenum: Spectacle disease c Copper: Wilson disease d Mercury: Peat scours
		a Selenium : Blind stagger
		b Molybdenum: Spectacle disease
	. 3	c Copper: Wilson disease
	5	d Mercury: Peat scours
Q.	No. 111	Mechanism of toxicity and treatment of chlorate poisoning resembles with
	15- 11	a Cyanide toxicity
		b Nitrate toxicity
ŀ	2	c Urea toxicity
D	9	d Fluoride toxicity
Q.	No. 112	The correct order of toxicity potential of selenium is
4	0	a Elemental Selenium > Natural organic Selenium > Selenite > Selenide
	0	b Natural organic Selenium > elemental Selenium > Selenite > Selenide
4	P I	c Natural organic Selenium Selenide Selenite elemental Selenium
	0	d Natural organic Selenium Selenite Selenide elemental Selenium
Q.	No. 113	Oonopsis is an example of
	0	a Obligate selenium accumulator plant
	L	b Facultative selenium accumulator plant
		c Non accumulator plant
		d Non accumulator weed
Q.	No. 114	If CS- syndrome is evident then it indicates the poisoning of
		a Type I synthetic pyrethroid insecticides
		b Type II synthetic pyrethroid insecticides
		c Organophosphate insecticides
		d Organochlorine insecticides
Q.	No. 115	Species most resistant to belladona toxicity is
		a Cow
		b Horse
		c Pig
		d Rabbit

- Q. No. 116 Porcine vaginitis is evident following mycotoxicosis due to
  a Citrinin
  b AFG1
  c T-2 Toxin
- Q. No. 117 Knuckling of fetlock joint and roaring sound is characteristic feature of
  - a Lead
  - b Arsenic
  - c Cobalt
  - d Molybdenum

F-2 Toxin

- Q. No. 118 The condition called as "Animal drowns in its own fluid" occurs in poisoning of:
  - a Belladona
  - b HCN
  - c ANTU
  - d Mercury
- Q. No. 119 Rational therapy for acute organophosphate poisoning is:
  - a AChE reactivator alone
  - b Atropine alone
  - c Atropine plus AChE reactivator with supportive therapy
  - d Atropine plus AChE reactivator plus Nicotinic antagonist with supportive therapy
- Q. No. 120 Level of sulfan sulphur (S°) is enhanced to neutralize –CN radicals by the action of the enzyme:
  - a Rhodanese
  - b Aminotransferases
  - c Acetyl transferases
  - d Plasma eserases



## **Key: Veterinary Pharmacology and Toxicology**

Q. No.	Ans	swer	Q. No.	Ans	swer
Q. No. 1	С	1000:1	Q. No. 61	d	Rapidly degraded
Q. No. 2	d	4 hour	Q. No. 62	c	Methysergide
Q. No. 3	a	Conjuctival sac	Q. No. 63	b	Enkephalins
Q. No. 4	a	Passive diffusion	Q. No. 64	a	Talnetant
Q. No. 5	d	28	Q. No. 65	d	Bethanechol
Q. No. 6	Ъ	80 L	Q. No. 66	a	Glycine and D-Alanine
Q. No. 7	d	Strophanthus gratus	Q. No. 67	a	Bupivacaine > Lidocaine >
		- श्रिक्सा वि	STIPA		Chlorprocaine > Benoxinate
Q. No. 8	С	Zero order kinetics	Q. No. 68	c	The ability of local anesthetics
		(2)	4.1	9	to diffuse to the site of action
Q. No. 9	b	α-β-γ trimer with GDP	Q. No. 69	b	Enkephalins
2	80	occupying the site on α-subunit			9
Q. No. 10	b	Tachyphylaxis	Q. No. 70	b	Addition of more than 9 carbon
5	1	M M			chain at C-5 leads to more
100	47.97				depressant activity
Q. No.11	b	Haptens	Q. No. 71	С	Methoxyflurane < Chloroform
15 11	7				< Halothane < Isoflurane
Q. No. 12	С	5-HT <sub>2</sub> receptor	Q. No. 72	С	Pirenzepine
Q. No. 13	a	Anaphylais	Q. No. 73	d	Conivaptan
Q. No. 14	a	4.3 half lives	Q. No. 74	С	100 Rads
Q. No. 15	d	IP <sub>3</sub> and DAG	Q. No. 75	b	Neostigmine
Q. No. 16	С	Side effect	Q. No. 76	b	Bicuculline
Q. No. 17	c	Oxotrimorine: Trimethophan	Q. No. 77	a	PLC-IP3/DAG pathway
Q. No. 18	a	Glucuronidation	Q. No. 78	a	Steroidal hormone
Q. No. 19	d	Certain safety factor	Q. No. 79	d	D2 receptors in gastric mucosa
92		H() (1)	4 3		and D2 and D3 receptors in
7	10			11	CTZ
Q. No. 20	С	CYP3A4	Q. No. 80	С	Flucloxacillin
Q. No. 21	a	Zero order kinetic	Q. No. 81	d	Phenylbutazone
Q. No. 22	С	Adrenaline	Q. No. 82	a	Potassium
Q. No. 23	b	Haptens	Q. No. 83	b	Horse
Q. No. 24	b	Cysteine	Q. No. 84	d	Amoxapine
Q. No. 25	b	5-HT <sub>4</sub> - receptor antagonist	Q. No. 85	С	Thallous ions: Na <sup>+</sup> & K <sup>+</sup>
Q. 1.5. 25		c 1114 1000ptot uniungemen	Q. 1.0. 00		channels
Q. No. 26	С	α <sub>1-</sub> adrenergic receptor	Q. No. 86	С	Sodium cromoglycate
Q. No. 27	d	Guanylyl cyclase	Q. No. 87	С	Amphotericin B
Q. No. 28	d	Methionine	Q. No. 88	a	5HT <sub>2A</sub>
Q. No. 29	С	Lithium carbonate	Q. No. 89	d	Rifampin
Q. No. 30	a	Inhibiting NMDA receptor	Q. No. 90	С	6th
Q. No. 31	С	Intrinsic ion channel operation	Q. No. 91	a	Eukaryote: Topoisomerase IV
Q. No. 32	a	Captopril	Q. No. 92	d	Amprolium
Q. No. 33		Lipid solubility	Q. No. 93		Zidovudine
Q. 110.33	a	Lipiu soiuoiiity	Q. 110.93	a	Zidovadilic

Q. No. 34	d	Physiological antagonism	Q. No. 94	d	Pyrimethamine
Q. No. 35	b	Pyruvate	Q. No. 95	d	Gouty arthritis
Q. No. 36	С	300-600 A <sup>0</sup>	Q. No. 96	b	Tetracycline
Q. No. 37	С	β <sub>1-</sub> adrenergic receptor stimulation	Q. No. 97	c	Serine
Q. No. 38	d	Tyrosine	Q. No. 98	b	Lead
Q. No. 39	a	Ethacrynic acid : Na <sup>+</sup> -Cl symport inhibitor	Q. No. 99	a	6: 1
Q. No. 40	d	Partial heart block	Q. No. 100	d	Clomiphene
Q. No. 41	d	Bicuculline	Q. No. 101	С	Niclosamide
Q. No. 42	c	Inhibiting Ach release	Q. No. 102	d	10000 ppm
Q. No. 43	b	A dopaminergic receptor agonist	Q. No. 103	d	Pteridium aquilinum
Q. No. 44	d	Isoprenaline > Epinephrine > Norepinephrine	Q. No. 104	d	Carbaryl - 2-PAM-
Q. No. 45	d	Desipramine	Q. No. 105	a	Na <sup>+</sup> channels
Q. No. 46	b	Copper	Q. No. 106	a	Ferric form of Hb is converted ferrous form of Hb
Q. No. 47	c	Clonidine	Q. No. 107	d	Acetyl cysteine
Q. No. 48	b	Chenodeoxycholic acid	Q. No. 108	a	Aflatoxin
Q. No. 49	a	Antiepiletic	Q. No. 109	a	Enkephalins
Q. No. 50	d	Hydrated aluminium silicate	Q. No. 110	d	Mercury: Peat scours
Q. No. 51	d	Decrease in oxygen consumption	Q. No. 111	b	Nitrate toxicity
Q. No. 52	d	Phenylephrine	Q. No. 112	d	Natural organic Selenium> Selenite>Selenide> elemental Selenium
Q. No. 53	b	Labetalol	Q. No. 113	a	Obligate selenium accumulator plant
Q. No. 54	С	4-6	Q. No. 114	b	Type II synthetic pyrethroid insecticides
Q. No. 55	c	Edrophonium	Q. No. 115	d	Rabbit
Q. No. 56	c	Benztropine	Q. No. 116	d	F-2 Toxin
Q. No. 57	d	Inhibiting CysLT <sub>1</sub> receptor	Q. No. 117	a	Lead
Q. No. 58	b	Physostigmine	Q. No. 118	c	ANTU
Q. No. 59	b	Medulla	Q. No. 119	c	Atropine plus AChE reactivator with supportive therapy
Q. No. 60	d	Lidocaine	Q. No. 120	a	Rhodanese