of the Five-Year Degree Course

Maximum Marks: 60

VETERINARY PHARMACOLOGY PAPER-I

| In and Systyemic Veterinan, Di | irks: 60 |
|--|----------------------|
| Section D. | Marks 30 Marks 30 |
| Attempt all questions Answer of all questions is to be written in the space provided at the question in question-booklet. | long with |
| Overwriting is not allowed in the objective type question. | - Mg WALL |
| SECTION - A | |
| General and Systyemic Veterinary Pharmacology: VPT-311 Maximum N | Tarks 30 |
| | 0.5 = 4.5 |
| i) is regarded as a father of chemotherap | |
| ii) Study of absorption, clishibution, metabolism and | excretion |
| of drug is known as pharmacokinetics. | |
| iii) Drugs which cause contraction of non-pregnant uterine muscle is known | own as |
| £ cholic tendence that | |
| iv) Enzymes responsible for metabolism of drug, located in endoplasmi | c reticul |
| of hepatic cell are known as nicrosomal | |
| v) Metaclopramide is an antagonist ofrec | eptors. |
| vi) Potassium citrate acts as urinary Alkalinium | |
| vii) Muscular dystrophy in cattle can be treated usi | ng vit |
| type of diuretics | |
| viii) Mannitol is a Osmohč type of diuretics. | enhance |
| ix) produces local hyperemia and | Cilitano |
| process of inflammation | |
| VPT-I/28/S1/3 rd yr/2017/M F | Page : |

wille across this line

| Q.2 Choose the most suitable answer and write the number of the correct answer $(9x0.5 = 4.5)$ 1 or 2 or 3 or 4 in the space given against each sub question: $(9x0.5 = 4.5)$ | | |
|--|--------|-------|
| table answer and waste each sub question: | (9x0.5 | = 4 5 |
| Chanse the most suitable given against | | 1.0) |
| Q.2 Choose the most of the space given as the space given given as the space given | (| 1 |
| tong obtained from Syllar | | |
| i) Following is a drug of | | |
| | | |
| 2. Quinine | | |
| 3. Iodine | (| 1 |
| 4. Meloxicani | | , |
| 3. Iodine 4. Meloxicam ii) Following is an example of buffered antacid? 1. Alluminum hydroxide 1. Alluminum phosphate | | |
| 1. Jum phosphate | | |
| Alluminum partialicate Magnesium trisilicate | | |
| 3. Magnesium distribution of the second of t | - | - |
| 4. All of above | |) |
| Magnesium under All of above Hypoglycemic convulsion in mice is a bioassay used for? Hypoglycemic convulsion in mice is a bioassay used for? | | |
| | | |
| 2. Immunglobulin | | |
| _ 3. Insulin | | |
| 4. Thyroxine iv) Which of following is an action of bodyon a drug? | (|) |
| iv) Which of following is an account | | |
| 1. Excretion | | |
| 2. Disrtibution | | |
| 3. Metabolism | | |
| All of above Following is a metabolite of chloral hydrate? | (| · · |
| | minima | 1 |
| 1. Arylamine | | |
| 2. Trichloro acetic acid | | |
| Trifluoro acetic acid | | |
| 4. Trichloroethanol | | |
| vi) Liver has tendency to accumulate following drug. | (|) |
| Arsenic and fluoride | | |
| Chloroquine and fluoride | | |
| 3. Chloroquine and paracetamol | | |
| Paracetamol and iodine | | |
| vii) For filtration, drug must have molecular weight: | | |
| 1. Less than 100 dalton | (|) |
| 2. Less than 1000 dalton | | |
| 3. More than 100 dalton | | |
| 4 More than 100 dalton | | |
| 4. More than 10 dalton | | |
| viii) Following is an example of astringent? | | - |
| 1. Citric acid | (| 9 |
| 2. Calcium carbonate | | |
| 3. Acetic acid | | |
| A. Zinc ovid- | | |
| VPT-I/28/S1/3 rd yr/2017/M | | |
| yr/2017/M | | |

| Best available route of drug administration in unconscious animal 1. Intravenous 2. Intramuscular 3. Subcutaneous 4. All of above 3. Attempt any nine out of the following twelve questions. Answ question should be in 2 to 3 lines. What is phase I of drug development? | |
|--|--|
| i) Define generic name of drug. | |
| iii) Explain receptors with examples. | |
| iv) Define efficacy of drug. | |

| v) Write therapeutic indication of diethylsti | lboestrol. |
|---|--------------------------------|
| | |
| vi) Write two examples of physical antagonis | sm? |
| | |
| vii) Define spare receptors. | gards to sense sussess sussess |
| viii) What is an anhydrotics? Give example. | |
| | |
| y) Write use of vincristine. | |
| | |
| | |

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SECTION - B

| WRT 321 Maxin | num Mar | |
|--|-----------|--------|
| Veterinary Neuropharmacology: VPT-321 Maxim | (9x0.5 | Ks 30 |
| Q.6 Fill in the blanks. | | = 4.5) |
| i) Passage of impulse along the axon to untake of acetylcholine | into | - |
| ii) inhibits uptake of decly strong | THO VESIO | cle |
| during synthesis. | | |
| V. Lering is an | | |
| is an indirect acting sympathom | ımetic dr | ug |
| which inhibits functions of monoamine oxidase. | | |
| Stimulation of β ₂ (beta-2) receptor in bronchia causes | | |
| vi) is an example of G protein coup | led choli | Dergio |
| receptor. | | - Sic |
| vii) Butoxamine is an s receptor antag | roniet | |
| | | |
| viii) Glutamate and aspartate aretyp | e of CNS | |
| receptor in action. | | |
| ix) is an alkaloid derived from Rauwolfia s | serpentin | e. |
| Q.7 Choose the most suitable answer and write the number of the c | Orrect a | newar |
| 1 or 2 or 3 or 4 in the space given against each sub question: | (9x0.5 | =4.5 |
| i) Following is an example of selective serotonin reuptake inhibite | | |
| - runopine | or? (|) |
| 2. Citalopam | | |
| 3. Imipramine | | |
| 4. Amitriptyline | | |
| ii) Gallamine is an example of: | , | Α. |
| Natural competitive neuromuscular blocker Natural non-competitive neuromuscular blocker | (| , |
| Natural non-competitive neuromuscular blocker Synthetic competitive neuromuscular blocker | | |
| Synthetic competitive neuromuscular blocker Synthetic non competitive neuromuscular blocker | | |
| Synthetic competitive neuromuscular blocker Synthetic non competitive neuromuscular blocker Strychnine produces CNS stimulation via: | | |
| iii) Strychnine produces CNS stimulation via: | | |
| | (|) |
| 2. Stimulation of GABA receptors 3. Stimulation of GABA | | |
| 4 Inhibition of GABA and Cu | | |
| 3. Stimulation of GABA and Glycene receptors 4. Inhibition of GABA and Glycene receptors | | |
| 4. Inhibition of GABA and Glycene receptors | | |
| | | |

i) Give mechanism of action of methanicon

| ii) Define five ideal properties of general anaesthetics? |
|---|
| |
| |
| iii) Give two examples of reversible acetylcholine esterase inhibitors. |
| |
| |
| iv) Give two examples of topical anaesthetics? |
| I Company and least to the grown and least property |
| The anyoned gala at modificance to VI ego |
| v) Enlist therapeutic applications of atropine. |
| |
| |
| i) Explain stage III of general anaesthesia. |
| |
| |

| | | Please write | Roll No. above th | is line | |
|-----------------|---|---|-------------------|---------------|---------------|
| xii) Discu | ss mechanism of | faction of d | liazepams. | | Mary and |
| | *************************************** | *************************************** | | | |
| | | | | | |
| ************ | | | | | |
| Q.9 Attempt a | any three out of | the follow | ing four qu | estions. Ans | wer of an a |
| question | should be in 5 to | 8 lines. | | | (3x2 |
| i) Discuss | mechanism of a | ection of loc | cal anaesthe | sia. | |
| | | | | | |
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| | | | | rousinii laab | tologydy aver |
| | | | | | |
| ii) Discuss the | erapeutic uses of | phenothia | zine tranoi | ilizers | |
| | | | | zers. | |
| | Area chan 11 | | m) a lugue | rie to real | |
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| | | | ************** | | |

i) Discuss therapeutic uses of sympatholytics with their mechanism of action in

detail.

ii) Discuss intravenous anaesthetics with advantages, disadvantages, classification and mechanism of action.