First Year B. Pharm. (Practice): SUMMER - 2019 SUBJECT: APPLIED PHARMACEUTICS

| Day Date | | Friday 28/06/2019 S- 2019-5372 | Time: 10 Max. Mark | 0.00 AM TO 01.00 PM s:60 |
|---------------|-------------------------|---|---|-----------------------------|
| N.B. 1) 2) 3) | que Fig | No. 1 and Q.No. 5 are COMPULSORY. Out of remaining attentions from Section – I and ANY TWO questions from Section qures to the RIGHT indicate FULL marks. Sections should be written in SEPARATE ans | n – II. | WO |
| | | SECTION – I | | |
| Q.1 | A) | Attempt any seven objective questions (MCQs / fill in th following. | e blanks) of | the (07) |
| | i) ii) | Primary reason to formulate paediatric oral liquid is a) to improve palatability b) for ease of admic c) to improve bioavailability d) to improve stability Water is choice of solvent for oral liquid formulations; most for it is a) widely available b) low cost c) physiologically compatible and d) good solvent processors. | lity important rea | |
| | iii) | safe drug molecules The main reason to formulate dry syrup is a) ease of handling b) to improve bioact c) to improve taste d) to protect from respectively. | vailability | |
| | v) vi) vii) viii) | Eye drops and injection are same in requirements, excepta) both are sterile | n free un. | by |
| Q.1 | B) | Give principle of GMP, QA and QC. | | (03) |
| Q.2 | | uss merit and demerits of formulating dosage form and wulating various types of tables. | vrite concept | of (10) |
| Q.3 | a) b) | Discuss properties of active pharmaceutical ingredient considered while designing its dosage form. Discuss mechanism of transdermal penetration of drug. | that should | be (05) (05) |
| Q.4 | Ansv i) ii) iii) iv) v) | wer ANY FIVE of the following. Write with reason two examples of drugs and its proprietary proprietary formulated into enteric coated tablets. Write two examples of drugs that need to store in cold place a Write merit and demerit of insulin syringe over conventional. Compare the concept of dispersible tablet and mouth dissolving Explain method to administer pessary. | and dry place. insulin injecti ng tablet. | (10) are |
| | vi) | Write concept of formulation of lotions with two example products. | P.T. | |

SECTION – II

| Q.5 | A) | Attempt any seven objective questions (MCQs / fill in the blanks) of the following. | (07) | | |
|-------------|--|---|------|--|--|
| | i) | A drug having Low solubility and low Permeability is related to | | | |
| | , | a) BCS Class I b) BCS Class II | | | |
| | | c) BCS Class III d) BCS Class IV | | | |
| | ii) | Name of antibiotics which causes staining of infant teeth | | | |
| | , | a) Penicillin b) erythromycin | | | |
| | | c) Tetracycline d) All the above | | | |
| | iii) | One of the following reaction does not take place in Phase I metabolic | | | |
| | | a) Oxidation b) Reduction | | | |
| | | c) Hydrolysis d) Conjugation | | | |
| | iv) | For absorption of drug from gastrointestinal tract, preferentially it should | | | |
| | , | have log P valve | | | |
| | | a) Less than 1 b) 1–2 | | | |
| | | c) 4–5 d) Above 5 | | | |
| | v) | The drug passage from stomach to the small intestine is called . | | | |
| | vi) molecular size of drug pass from blood to extracellular fluid. vii) BCS Class II drug shows Solubility and Permeability. | | | | |
| | | | | | |
| | viii) | | | | |
| Q.5 | B) | Explain the mechanism response for loss dicoumarol activity due to (03) Phenobarbital co-administration? | | | |
| Q.6 | Expla | ain in details various barriers for drug distribution. | (10) | | |
| Q. 7 | a) b) | Define Bioequivalence and discuss importance of BA/BE Studies. Explain patient related factors affecting absorption of drugs. | | | |
| Q.8 | Answer ANY FIVE of the following. (10 | | | | |
| | i) | Define phagocytosis and pinocytosis. | | | |
| | ii) | Why sodium and potassium salt of diclofenac available in market. | | | |
| | iii) Write note on pro-drug approach of levodopa in Parkinsonism patient | | | | |
| | iv) | | | | |
| | , | examples? | | | |
| | v) | What are various sites of drug metabolism in the body? Why is liver | | | |
| | • | considered the major site for such a process? | | | |
| | vi) | Write significance of biological half-life (t 1/2) of drug. | | | |
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