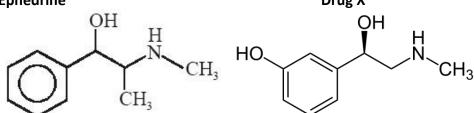


Instructions:

- a. There are twelve pre-labelled answer books. Please make sure you answer the questions in the respective answer book.
- b. Write your candidate number on the cover of each answer book.
- c. Use ink or ball-point pen.
- d. Answer ALL questions. They are worth equal marks and you should spend approximately **ten minutes** for each question. For questions with multiple parts, allocation of marks is indicated in the brackets.
- 1. The structures for ephedrine (left) and drug X (right) are given below: Ephedrine Drug X



- (a) Explain the term structure activity relationship (10%).
- (b) Based on the structure-activity relationships, describe the similarities and differences in pharmacokinetic and pharmacodynamic properties of drug X compared with ephedrine (90%).
- 2. Describe the pharmacology of suxamethonium under the following headings:
 - (a) mechanism of action, including a description of the appropriate receptor (40%);
 - (b) pharmacokinetic properties (30%); and
 - (c) adverse effects (30%).
- 3. Empagliflozin and rosiglitazone are drugs used for the treatment of diabetes mellitus. For each drug, describe the mechanism(s) of action (40%), and important adverse effects (40%). Explain the potential mechanism for adverse cardiovascular effect for rosiglitazone (20%).
- 4. Compare and contrast the pharmacodynamic properties of dexmedetomidine and propofol.
- 5. Classify, with example, the drugs that affect uterine tone (40%). Outline the adverse effect of these agents (60%).

- 6. Describe the dose and mechanisms leading to paracetamol toxicity (75%). Outline the conditions that predispose to paracetamol toxicity (25%). (Note: treatment for paracetamol toxicity is not required)
- 7. (a) Outline the mechanism of action of corticosteroid in the management of asthma (40%).
 - (b) List four side effects associated with long term use of corticosteroid and outline their underlying mechanisms (60%).
- 8. (a) Define drug tolerance (20%).
 - (b) Draw a dose-response curve to illustrate the phenomenon of drug tolerance (20%).
 - (c) Describe THREE possible pharmacodynamic mechanisms by which drug tolerance may develop (60%).
- 9. Describe and use examples to illustrate the clinical uses of opioids.
- 10. Outline the chemistry of soda lime and its use in anaesthesia (50%). Describe the interaction of soda lime with sevoflurane and desflurane (50%).
- **11.** A new "Aspiration Risk Test" is used to predict the risk of aspiration with a sensitivity of 86% and specificity of 70%.
 - (a) Define the terms "sensitivity", "specificity", "positive predictive value" and "negative predictive value". Explain how they can be obtained in a 2x2 table (50%).
 - (b) How will the incidence of aspiration affect performance of this test in terms of predicting the aspiration risk of an individual (50%)?
- 12. Describe the factors that may influence the plasma concentration of local anaesthetics after a single bolus of local anaesthetics for a peripheral nerve block (50%). How would term pregnancy affect the plasma concentration (50%)?

- End -