

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. Describe the mechanisms on how non-steroidal anti-inflammatory drugs exert their therapeutic and adverse effects (75%). Outline the advantages and disadvantages of using COX-2 selective agents (25%).
- 2. Describe the pharmacokinetic and pharmacodynamic properties of ketamine (75%). Outline the advantage of using ketamine as a sedative agent for change of dressing in burned patients (25%).
- 3. What are the differences between addiction, physical dependence and tolerance (60%)? Explain why methadone has a low addictive potential (40%).
- 4. Write short notes on the potential effects of epidural administration with 6ml of 0.25% levobupivacaine and 50mcg of fentanyl at the T6/7 level.
- 5. Describe the factors determining the speed of onset of non-depolarizing neuromuscular blocking drugs.
- 6. Outline the mechanisms of action, side effects, and indications, of 1) ACE inhibitors, 2) beta-blockers and 3) calcium channel blocker (70%). Compare their specific benefits in patients with ischaemic heart disease (30%).
- 7. Define saturated vapor pressure (SVP) and boiling point of a volatile anaesthetic agent (50%). Explain why contemporary variable bypass vaporizers are unsuitable for delivering desflurane (50%).

- 8. Define minimum alveolar concentration (MAC) of an inhaled anaesthetic agent (20%). What are the physiological, pharmacological and pathological factors that may affect MAC (80%)?
- Define randomization in clinical trials (5%). Explain the purpose of randomization in clinical trials (15%). Describe the following types of randomization, and outline their advantages and limitations (80%).
 - A. Simple randomization
 - **B.** Block randomization
 - C. Stratified randomization
- 10. Outline the mechanisms of action of sodium-glucose co-transporter 2 (SGLT-2) receptor inhibitor (40%). Describe the mechanisms where SGLT-2 blockers may induce perioperative euglycemic diabetic ketoacidosis (60%).
- 11. Outline the mechanisms of action and clinical indications of different classes of drugs that are useful in inducing diuresis clinically.
- 12. Define "volume of distribution" (25%).What factors influence the size of the volume of distribution (50%)?How may it be used in the calculation of a loading dose (25%)?

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