

Australian and New Zealand College of Anaesthetists

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PRIMARY EXAMINATION

THURSDAY, 28 MAY 2009

3:15 pm – 4:35 pm

PHARMACOLOGY

MULTIPLE CHOICE QUESTIONS

NOTICE

- **ALL** questions must be answered
- Write the answer to each question on book provided.
- No queries regarding individual questions can be answered.
- There are 75 questions to answer in 75 minutes. You should answer each question in 1 minutes or less.
- All questions are worth equal marks.

1. Drug distribution and clearance:

- A. Volume of distribution of a drug is independent of plasma-protein binding
- B. Highly ionized drugs may have a renal clearance approximating to the glomerular filtration rate**
- C. Phenytoin undergoes zero-order kinetics at low plasma concentrations
- D. A two compartment model can describe the profile of drug concentrations following intramuscular injection
- E. The terminal half-life of a drug provides a useful guide to optimal bolus drug administration

2. Which of the following crosses the blood-brain barrier?

- A. GABA
- B. Propranolol**
- C. Suxamethonium
- D. Edrophonium
- E. Dopamine

3. The following drugs inhibit the cytochrome P450:

- A. Omeprazole
- B. Dexamethasone
- C. Rifampicin
- D. Phenytoin
- E. Fluconazole**

4. Which of the following statements is false:

- A. The volume of distribution (Vd) for water-soluble drugs is smaller in neonates than adults**
- B. In pregnancy the clearance of drugs excreted by the biliary system may be attenuated
- C. Plasma-protein binding is unchanged in obese patients
- D. The clearance of drugs with low hepatic extraction ratios is not limited by hepatic blood flow
- E. Oral propranolol has greater bioavailability in patients with chronic renal failure

5. With isoflurane anaesthesia, MAC awake is:

- A. 0.1% vol
- B. 0.3% vol**
- C. 0.5% vol
- D. 0.8% vol
- E. 1% vol

6. Which of the following statements are true:

- A. The peripheral nerve stimulator generates a square wave electrical pulse**
- B. Double-burst stimulation consists of two bursts of three stimuli at 50 Hz with each triple burst separated by 50 ms
- C. The ratio of twitches with train of four stimulation (T4:T1) in a depolarizing block is less than one
- D. A depolarizing block produces detectable fade in muscle contraction with tetanic stimulation
- E. Post-tetanic count reflects the reversal of neuromuscular blockade

7. Which is NOT correct regarding opioids administered via the epidural space

- A. urinary retention is relatively common
- B. low dose naloxone alleviates pruritis without significantly modifying analgesia
- C. biphasic respiratory depression may occur
- D. lipid solubility affects speed of onset
- E. morphine is commonly used because it has the lowest side effect profile

8. Which of the following statements are true:

- A. At a pH below their pKa, weak acids will be more un-ionized
- B. Phagocytosis is an important mechanism in the transport of insulin
- C. According to Graham's law the rate of diffusion is inversely proportional to the molecular size
- D. Erythromycin decreases the bioavailability of warfarin
- E. A volatile agent with high solubility in blood has rapid onset of effect

9. Uptake of N₂O when breathing 70%:

- A. More than one litre absorbed in the first minute
- B. Equilibrium is achieved in 3mins
- C. Absorb 10 litres in the first 3 mins
- D. At steady state, uptake is 200mls/min
- E. Produces surgical anaesthesia

10. What is NOT correct regarding rramadol:

- A. Is only a weak agonist at the μ -receptor
- B. Inhibits the reuptake of norepinephrine
- C. Increases the levels of serotonin at descending inhibitory neurons
- D. Is more effective for neuropathic pain than morphine
- E. In equianalgesic doses causes greater respiratory depression than morphine

11. Propofol

- A. is a weak organic acid with a pKa of 11
- B. contains Cremophor EL as the commercial solvent
- C. partially antagonises alfentanil
- D. is an effective analgesic
- E. requires a blood concentration of approximately 12-14 μ g/ml for induction of anaesthesia in unpremedicated patients

12. The following is true of opioids and their routes of administration EXCEPT:

- A. Oral oxycodone has a higher bioavailability than oral morphine
- B. Patient's age is a more reliable indicator of opioid dose via the intramuscular route than weight
- C. Pethidine is irritating and painful when administered by subcutaneous injection
- D. The analgesic efficacy of epidural opioids is less than that of parenteral opioids
- E. Transdermal fentanyl patch uses the principle of passive diffusion

13. Phenylephrine:

- A. Metabolised by COMT
- B. used in glaucoma

- C. Safely be used with a monoamine oxidase inhibitor
- D. Effect lasts longer than noradrenaline**
- E. Acts by indirect method only

14. Which of the following DO NOT decrease the lower oesophageal sphincter pressure:

- A. Thiopentone
- B. Atropine
- C. Progesterone
- D. Isoflurane
- E. Suxamethonium**

15. Gelofusine

- A. is derived from human plasma
- B. contains molecules with an average weight of 30 000**
- C. contains calcium
- D. contains preservative
- E. interferes with cross matching

16. The following decreases the intraocular pressure:

- A. Suxamethonium
- B. Acetazolamide**
- C. Hypercarbia
- D. Atropine
- E. Elevated central venous pressure

17. In clinical doses, captopril

- A. may cause hypokalaemia
- B. may cause a moist cough
- C. should be given with food
- D. is given orally once a day
- E. causes an increased level of angiotensin I**

18. Physostigmine

- A. Causes excitatory activity on the EEG**
- B. Doesn't cross the blood brain barrier
- C. Doesn't cause sedation
- D. Only has its effects at nicotinic receptors
- E. Causes retrograde amnesia

19. In a clinical trial, a patient either vomits or not. What type of data is this?

- A. Ordinal
- B. Nominal**
- C. Ratio
- D. Interval
- E. Continuous

20. The following are correct regarding pethidine EXCEPT:

- A Readily crosses the placenta
- B Is a phencyclidine derivative**
- C Norpethidine has proconvulsant properties
- D Pethidine trapping occurs in the fetal compartment
- E. has local anaesthetic properties

21. Low molecular weight heparins

- A. contain unfractionated heparins
- B. is usually given by infusion
- C. should be monitored during use by the activated partial thromboplastin time (APTT)
- D. should be discontinued at least 6 hours before major surgery
- E. may cause thrombocytopenia**

22. The following is correct regarding local anaesthetics EXCEPT:

- A. Cocaine undergoes significant metabolism by ester hydrolysis**
- B. Potency of local anaesthetic agents is quantified as octanol partition coefficient
- C. Acidic environment of an abscess decreases the non- ionized proportion of local anaesthetic
- D. At lower concentrations, the block produced by ropivacaine may be motor sparing
- E. Levobupivacaine is the s-enantiomer of racemic bupivacaine

23. Ketamine

- A. increases cerebral blood flow**
- B. is an N-methyl-D-aspartate (NMDA) receptor agonist
- C. causes a reduction in airway secretions
- D. is a poor analgesic
- E. causes premature labour contractions

24. Regarding sevoflurane:

- A. The vapour pressure is less than enflurane**
- B. The vapour pressure is greater than isoflurane
- C. Cardiovascular side effects are similar to isoflurane
- D. Molecular weight less than isoflurane
- E. Boiling point less than enflurane

25. The following are correct regarding halothane hepatitis EXCEPT:

- A. The incidence of halothane hepatitis is lower in children compared with adults
- B. Patients on liver-enzyme inducing drugs have a lower incidence of halothane hepatitis**
- C. The latency period before presentation with hepatitis is directly proportional to the time between multiple exposures to halothane
- D. The oxidative metabolite of halothane is the trigger for hepatic injury
- E. Enflurane has been reported to cause halothane hepatitis

26. Sulphonylureas:

- A. High incidence of lactic acidosis
- B. Good in patients with depleted insulin stores
- C. Metformin & phenformin are examples

- D. Increased glucose utilisation in the peripheries
- E. Are related to sulphonamides

27. The following drugs cause uterine relaxation EXCEPT:

- A. Nifedipine
- B. Terbutaline
- C. Magnesium sulphate
- D. Atracurium
- E. Atosiban

28. The benzodiazepine with the longest elimination half-life is:

- A. Diazepam
- B. Oxazepam
- C. Temazepam
- D. Midazolam
- E. Lorazepam

29. The following are all amides except:

- A. Bupivacaine
- B. Prilocaine
- C. Etidocaine
- D. Tetracaine
- E. Dibucaine

30. The following statements regarding the drugs used to treat nausea and vomiting are true EXCEPT:

- A Prolonged QT interval is a potentially serious side effect of chronic administration of droperidol
- B Opioid antagonists have ineffective anti-emetic properties
- C Metoclopramide has some 5HT₃ antagonism at higher doses
- D Tropisetron is a 5-HT₃ antagonist, which is metabolized by cytochrome P450 enzyme system
- E Dexamethasone works as an anti-emetic by antagonizing the dopamine receptor

31. With regards to diffusion through a membrane:

- A. Directly proportional to thickness
- B. Inversely proportional to thickness
- C. Inversely proportional to Surface area
- D. Inversely proportional to concentration difference
- E. Directly proportional to square root of the molecular weight

32. Desflurane

- A. has a molecular weight of 185
- B. has a minimum alveolar concentration (MAC) value of 2%
- C. has a blood:gas solubility of 0.42
- D. is flammable in a concentration of 6%
- E. should not be used with soda lime

33. Regarding Neostigmine:

- A. Binds covalently to esteric site on AChEsterase
- B. Binds electrostatically to esteric site on AChEsterase
- C. Binds to anionic site
- D. Forms complex with AChEsterase with a shorter half life than acetylcholine
- E. is the main antidote to Calabar bean poisoning

34. Which is FALSE regarding the mechanism of action of drugs used perioperatively:

- A. Benzodiazepines act as agonists at GABAA receptors by increasing chloride entry to hyperpolarize the synaptic membrane
- B. Ondansetron is a 5-HT3 antagonist that acts both centrally and peripherally
- C. Clonidine is an agonist predominantly at α_2 -adrenoreceptors
- D. Heparin acts by inhibition of antithrombin III
- E. Paracetamol has no effect on the cyclo-oxygenase pathway

35. The statement INCORRECTLY describes how the inotropes increases cardiac contractility:

- A. Adrenaline increases cardiac cAMP
- B. Dobutamine stimulates predominantly β_1 -receptors
- C. Dopexamine acts on β_2 -receptors
- D. Milrinone increases cAMP by stimulating phosphodiesterase
- E. Digoxin increases intracellular calcium by inhibiting the Na-K pump

36. Rocuronium administered in 2 times the ED95 dose:

- A. Rapid onset, short duration
- B. Rapid onset, Intermediate duration
- C. Slow onset, intermediate duration
- D. Slow onset, long duration
- E. Slow onset, short duration

37. Regarding the power of a two-sample t test, all are true EXCEPT:

- A. Increases if the sample sizes are increased
- B. Depends on the difference between the population means which we wish to detect
- C. Depends on the difference between the sample means
- D. Is the probability that the test will detect a given population difference
- E. Cannot be zero

38. The following drugs, in therapeutic dosage have the described approximate percentage plasma protein binding

- A. morphine, 10%
- B. thiopentone 99%
- C. pancuronium 50%
- D. diazepam, 75%
- E. fentanyl, 85%

39. Anti-arrhythmic drugs that DO NOT block the sodium pump include:

- A. Amiodarone

- B. Flecainide
- C. Lidocaine
- D. Propafenone
- E. Verapamil

40. Regarding digoxin toxicity:

- A. Serum level > 2.1 ng/ml is toxic
- B. Calcium acts as an antidote
- C. Causes a long PR interval
- D. Causes xanthopsia
- E. Causes a long QT interval and torsades

41. Drugs WITHOUT α antagonist properties include:

- A. Yohimbine
- B. Doxazosin
- C. Phenoxybenzamine
- D. Phentolamine
- E. Hydralazine

42. With regard to the ionisation of acids and bases

- A. weak acids have a $pK_a > 7.4$
- B. weak acids in solution cannot exist in alkaline pH
- C. the concentration of the ionised form of weak bases increases with an increase in pH
- D. decreasing the H^+ concentration decreases the ionised form of weak acids
- E. strong acids are highly ionised at pH 7.4

43. Adverse effects of epidural opioids include

- A. Lower limb motor blockade
- B. Respiratory depression
- C. Surgical site bleeding
- D. Pulmonary embolism
- E. Interactions with local anaesthetics

44. Which of the following drugs are contraindicated in patients with acute intermittent porphyria

- A. propofol
- B. fentanyl
- C. thiopentone
- D. ketamine
- E. halothane

45. NMDA antagonists include

- A. Ketamine
- B. Propofol
- C. Adrenaline
- D. Ketorolac

E. Dextrometorphane

46. The following drugs cross the placenta in clinically significant amounts when given in conventional dosages during labour

- A. intravenous fentanyl
- B. intravenous neostigmine
- C. intravenous vecuronium
- D. epidural bupivacaine
- E. intravenous ketamine

47. Genetic variability for the clinical efficacy of morphine is suggested to be related to all EXCEPT:

- A. Differences in mu-opioid receptors
- B. Differences in the metabolism of morphine
- C. Differences in renal clearance of morphine
- D. Differences in transport of morphine across the bloodbrain barrier.
- E. Differences in adrenergic modulation of opioid analgesia

48. Guanethidine:

- A. Acts primarily within the CNS
- B. Produces anti-hypertensive effect primarily by presynaptically inhibiting release of noradrenaline
- C. Highly lipid soluble
- D. Mental depression is a troublesome side effect
- E. Orthostatic hypotension is not a prominent side effect

49. Remifentanyl

- A. is a pure μ opioid receptor agonist
- B. is metabolised by plasma cholinesterase
- C. has clinically active metabolites
- D. is equipotent with alfentanil
- E. does not cause nausea and vomiting

50. The blood gas coefficient for the agents are correct EXCEPT:

- A. Isoflurane 1.4
- B. Sevoflurane 0.69
- C. Ether 12
- D. Desflurane 0.42
- E. Halothane 0.8

51. Metabolism of analgesics in neonates is characterised by:

- A. Clearance of paracetamol at the level of older children
- B. Impaired sulphation
- C. Impaired glucuronidation
- D. A morphine clearance of about twice that of older infants
- E. Same pathways of elimination as in adults

52. Droperidol

- A. is a thioxanthine
- B. increases cerebral blood flow
- C. is antiemetic is large doses only
- D. may cause hypotension**
- E. is metabolised by the kidneys

53. Opioid metabolism

- A. Catechol-O-methyltransferase takes part in degradation of morphine
- B. Morphine-6-glucuronide has no analgesic efficacy
- C. Morphine-6-glucuronide is found in higher concentrations than morphine during chronic oral morphine administration**
- D. Less than 5% of Caucasians are poor metabolisers of codeine
- E. Tramadol is metabolized by the same pathway as morphine

54. Which of the following is NOT TRUE regarding aminophylline:

- A. is a mixture of theophylline and ethylenediamine
- B. is metabolized by xanthine oxidase
- C. has an elimination half-life of 12 hours**
- D. causes peripheral vasodilation
- E. is a central nervous system (CNS) stimulant

55. Adenosine

- A. Causes AV block via action at A1 receptors**
- B. Causes bronchoconstriction via A2 receptors
- C. Causes renal vasodilation
- D. Causes profound depression of the SA node
- E. has an exaggerated response in the presence of methyl xanthines

56. Malignant hyperthermia:

- A. Is an inherited hypermetabolic disorder of smooth muscle
- B. Is triggered by non-depolarizing muscle relaxants
- C. Reconstituted dantrolene has a pH of 9 which is an irritant to veins**
- D. Does not occur in children under 2 years of age
- E. Is caused by a single gene defect on chromosome 5p

57. Sugammadex:

- A. inhibits acetylcholinesterase
- B. reverses atracurium
- C. Binds flucloxacillin**
- D. undergoes hepatic metabolism
- E. is a beta-cyclodextrin

58. The following drugs are effective treatments for post-anaesthetic shivering:

- A. Pethidine

- B. Clonidine
- C. Ketanserin
- D. Ondansetron
- E. All are correct

59. The following are correct examples of isomeric forms:

- A. tautomer: isoflurane and enflurane
- B. enantiomer: keto and enol forms of thiopentone
- C. racemic: cisatracurium
- D. structural: S and R forms of ketamine
- E. geometric: cis and trans forms

60. Compared to bupivacaine; ropivacaine:

- A. Has a higher pKa
- B. Is more lipid soluble
- C. Blocks type C nerve fibres faster
- D. Blocks type A nerve fibres more slowly
- E. Is more cardiotoxic

61. Amiloride:

- A. Potassium sparing antidiuretic which blocks the aldosterone receptor
- B. Blocks luminal sodium channels in the collecting tubules
- C. Increases potassium excretion.
- D. Is metabolised by the liver.
- E. Has a short elimination half time.

62. As an in vivo buffer, THAM (Tris-Hydroxymethyl- Amino-Methane) has the following advantages over sodium bicarbonate:

- A. Does not cause pulmonary oedema
- B. It does not increase PaCO₂
- C. Less irritation to peripheral veins
- D. It does not cross the blood brain barrier
- E. Less respiratory depression

63. Regarding the nicotinic acetylcholine receptor, all are true EXCEPT:

- A. Possesses four transmembrane domains.
- B. Each receptor has one binding site for acetylcholine.
- C. Consists of a- and b subunits.
- D. Is a ligand-gated ion channel receptor.
- E. Is a pentamer.

64. Of cardiovascular agents

- A. Dopamine prevents renal failure
- B. Milrinone is antiarrhythmic
- C. Levosimendan is a lusitropic agent
- D. Vasopressin improves inotropy

E. T3 is routinely administered in most ICUs

65. The following drugs can be used safely in a patient on long-term monoamine oxidase inhibitors:

- A. Ephedrine
- B. Pethidine
- C. Ketamine
- D. Fentanyl
- E. Tricyclic antidepressants

66. Regarding voltage-gated sodium channels, all are true EXCEPT:

- A. work on a second or minute timescale
- B. exist in three major conformational states
- C. inactivate very slowly
- D. are the molecular basis for the rapid transient sodium current first described by Hodgkin and Huxley
- E. are crucial for controlling the cellular resting potential of excitable cells

67. The following drugs induce hepatic microsomal enzymes:

- A. Cimetidine
- B. Sodium valproate
- C. Isoniazid
- D. Rifampicin
- E. Phenezine

68. Stomach emptying is stimulated by:

- A. atropine
- B. dopamine
- C. salbutamol
- D. small bowel distention
- E. propranolol

69. Aprotinin

- A. Is a polypeptide
- B. Inhibits plasmin and kallikrein
- C. Is derived from bovine lung
- D. Is metabolized and excreted by the kidney
- E. All of the above

70. The drug with the largest volume of distribution at steady state is:

- A. Propofol
- B. Midazolam
- C. Etomidate
- D. Thiopentone
- E. Methohexitone

71. Lignocaine has a pKa of 7.9 At pH 6.9, the percentage ionised is:

- A. 1%
- B. 10%
- C. 50%
- D. 90%**
- E. 99%

72. Which of the following is the most toxic effect of atropine in children?

- A. Hypotension
- B. Tachycardia
- C. Hyperthermia**
- D. Hypertension
- E. Dry mouth

73. Neuroleptic malignant syndrome:

- A. Occurs only with chronic use
- B. 80% (60%) mortality
- C. dantrolene acts as an antidote
- D. Can be caused by acute withdrawal of L-Dopa therapy**
- E. Is treated with cyproheptadine

74. Drugs which bind preferentially to and stabilize the inactivated conformation of voltage-gated sodium channels include

- A. prilocaine
- B. lidocaine
- C. gabapentin
- D. flecainide
- E. all of the above**

75. In reversing neuromuscular blockade, which of the following combinations is best matched with respect to time of onset?

- A. Atropine & neostigmine
- B. Atropine & glycopyrrolate
- C. Atropine & edrophonium**
- D. Atropine & physostigmine
- E. Glycopyrrolate and edrophonium