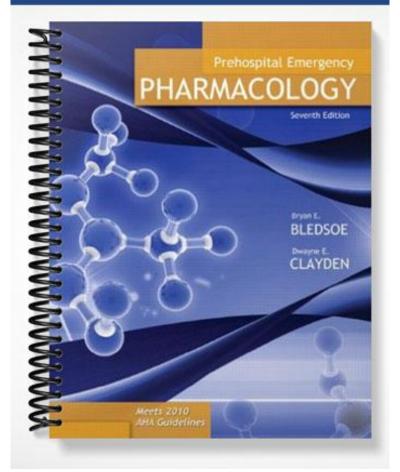
TEST BANK



Bledsoe T	est Bank
Chapter 2	Commented [IP1]: This chapter has 20 questions
1. Two major divisions of pharmacology are:	
a. Pharmacokinetics and pharmacodynamics	
b. Pharmacognosy and biotransformation	
c. Active transport and diffusion	
d. Biotransformation and elimination	
Answer: <u>Aa</u>	
Rationale: a. pharmacokinetics-Pharmacokinetics and pharmacodynamics are the	2- <u>two</u>
major divisions of pharmacology.	
b. pharmacognosy Pharmacognosy refers to the broad study of natural and synthe	tic
drugs and biotransformation is a function of pharmacokinetics.	
c. active Active transport and diffusion explain how a drugmedication moves.	
d. biotransformation Biotransformation and elimination is a function of	
pharmacokinetics.	
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2. Which of the following factors is not- <u>NOT</u> a component of pharmacokine	ics?
a. Absorption	
b. Distribution	
c. Biotransformation	
d. Binding	
Answer: d	
Rationale: a. explains-Explains the movement of a drug medication into the system	m. Commented [IP2]: Change to medication?
b. distribution Distribution is how the drug medication travels to target tissues or	
c. biotransformation Biotransformation is how a drug medication is broken down	

system<u>.</u>

d. binding Binding is a component of pharmacodynamics.

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3. Drug A requires the use of adenosine triphosphate (ATP) in order to move into

the cellular membrane. -This type of diffusion is known as:

- a. Elimination
- b. Active transport
- c. Facilitated
- d. Osmosis

Answer: b

Rationale: a. elimination Elimination is the removal of a drug from the body.
b. active Active transport involves the use of energy, such as ATP to move a substance through a membrane that otherwise would not be able to penetrate.
c. facilitated Facilitated diffusion requires the use of a helper protein (such as insulin with glucose) to cause a change in the cellular membrane allowing entry of the substance.
d. osmosis Osmosis is a passive process involving the movement of water.

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- Glucose in the bloodstream is a large molecule that cannot readily enter a cell unless it binds with insulin. -This <u>best-BEST</u> describes:
 - a. Osmosis
 - b. Diffusion
 - c. Carrier-mediated diffusion
 - d. Absorption

Answer: c

Rationale: a. <u>osmosis Osmosis</u> involves the movement of solvent, normally water.
b. <u>diffusion Diffusion</u> is a general term related to the movement of substances and involves both active and passive forms, of which carrier-mediated diffusion is included.
c. <u>carrierCarrier</u>-mediated diffusion most correctly explains the process described.
d. <u>absorption Absorption</u> encompasses a medications progress from its pharmaceutical dosage form to a biologically available substance that can then pass through or across tissues.

Commented [IP5]: Change to medication? NO

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 The term ______ describes the movement of molecules across a membrane down a pressure gradient from an area of higher pressure to and area of lower pressure normally resulting from hydrostatic forces.

- a. Filtration
- b. Carrier-mediated diffusion
- c. Active transport
- d. Osmosis

Answer: a

Rationale: a. filtration Filtration is correct.

b. <u>carrierCarrier</u>-mediated diffusion occurs when a protein is involved in binding and causing change in membrane shape to allow diffusion.

c. active Active transport involves the use of energy to cause diffusion.

d. osmosis-Osmosis is the movement of water from an area of low solute concentration to an area of high solute concentration.

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- 6. ______ describes the movement of a medication from the site of application into the body and into the extracellular compartment.
 - a. Diffusion
 - b. Osmosis
 - c. Absorption
 - d. Elimination

Answer: c

Rationale: a. diffusion <u>Diffusion</u> is the movement of a substance across a membrane.
b. osmosis <u>Osmosis</u> is the movement of water from an area of low solute concentration to an area of high solute concentration.

c. <u>absorption Absorption</u> encompasses a medications progress from its pharmaceutical dosage form to a biologically available substance that can then pass through or across tissues.

Bledsoe Test Bank d. elimination Elimination refers to removal of a substance from the body. pagePage 27 7. A medication that is acidic in nature when ingested orally will: a. Bypass the stomach and be absorbed slowly in the small intestine b. Be neutralized quickly upon entering the stomach c. Not enter the bloodstream, but will be eliminated in the gastrointestinal tract d. Be rapidly absorbed across the stomach membrane Answer: d Rationale: a. acidic Acidic medications, such as Aspirinaspirin, tend to be rapidly Commented [IP6]: generic name should be lowercase. absorbed in the stomach because of the acidic environment. -The small intestine environment is alkaline and will neutralize excessive hydrogen ions. b. see See rationale for a. c. see <u>See</u> rationale for a. d. correct Correct answer. Commented [IP7]: delete word answer to be consistent pagePage 28 8. Which of the following forms of absorption is generally the slowest due to the decreased vascular supply? a. Subcutaneous b. Intramuscular c. Intravenous d. Sub-lingual Commented [IP8]: delete hyphen? Answer: a

Rationale: a. the <u>The</u> subcutaneous layer of the skin is composed of dense fatty tissue that has poor vascular supply and slow absorption of medications.

b. <u>muscles_Muscles_generally</u> have very good blood supply promoting absorption.

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c. intravenous Intravenous injection provides the most rapid (and common) route of absorption.
d. the The sublingual area is highly vascular resulting in rapid absorption.

Commented [IP9]: hyphen?

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9. ______best_BEST_describes the process in which a medication is transported from the site of absorption to the site of action.

- a. Active transport
- b. Diffusion
- c. Distribution
- d. Elimination

Answer: c

Rationale: a. <u>active_Active_transport</u> involves the movement of a substance across the cellular membrane utilizing an energy source.

b. <u>diffusion_Diffusion</u> describes movement across the cellular membrane and can be an active or passive process.

c. <u>distribution Distribution</u> is the process whereby a medication is transported from the site of absorption to the site of action.

d. elimination Elimination is removal of a substance from the body.

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- 10. You are treating a patient who suffers from renal disease and decreased perfusion of the kidneys. -What effect will this have on the distribution of a drug that normally acts on kidneys, such as the diuretic medication Lasix?
 - a. The drug will be present in increased concentrations at the kidneys resulting in toxicity.
 - b. The drug will be poorly delivered to the kidneys due to a decreased perfusion and will not produce the desired results.
 - c. As the kidneys are a primary organ of elimination, the effects of the drug will be unchanged.

d. The drug will most likely be metabolized by the liver resulting in the same effects systemically.

Answer: b

Rationale: a. <u>decreased Decreased</u> renal blood flow results in poor delivery of the medication to the kidney.

b. correct.

c. the <u>The</u> decreased blood flow to the kidney will prevent the drug from producing the desired results.

d. <u>if-If</u> a drug is meant for a specific organ of tissue, it must reach the site in order to be effective.

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- 11. While researching the drug Amiodarone amiodarone, you read that the drug is considered to be highly protein bound upon entering the body. -Knowing this, you can deduce that:
 - a. The drug is largely ineffective when administered by intravenous route
 - b. The drugs half-life will be relatively short due to the rapid elimination of the free, or unbound, percentage
 - c. Much of the drug remains active in the bloodstream and only a small percentage is bound in tissues
 - d. The drug remains bound to proteins but will slowly release the active component when necessary resulting in a long half-life

Answer: d

Rationale: a. <u>when_When</u> administered in therapeutic dosages, intravenous routes provides the most rapid, effective dose.

b. the <u>The</u> drug will have a long <u>half half-</u>life as the bound component will slowly release as the free or unbound active component is metabolized.

c. <u>when When</u> the drug is highly protein bound, much of the drug will be stored in protein reservoirs. -Only a small amount remains in the bloodstream as an active metabolite.

Commented [IP10]: use of word drug, instead of medication throughout question ok.

Commented [IP11]: generic name lowercase

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d. correctCorrect.

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12. The blood_-brain barrier is a preventive barrier that prevents most_MOST

_____ medications from reaching the brain.

- a. Lipid-soluble
- b. Water-soluble
- c. Non-ionized, unbound
- d. Barbiturate

Answer: b

Rationale: a. lipid-Lipid-soluble medications readily pass the blood_-brain barrier.

b. waterWater-soluble medications cannot permeate the blood_brain barrier.

c. <u>a A</u> non-ionized, unbound medication can pass the blood_brain barrier_

d. <u>a-A</u> barbiturate is an example of a non-ionized, unbound drug.

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13. The process by which a drug medication passing through the liver may be	 Commented [IP12]: change to medication?
partially or completely inactivate many medications is known as:	
a. First-pass effect	Commented [IP13]: hyphen? Correct as is

- b. Proo-drug effect
- c. Protein reservoir binding
- d. Plasma binding effect

Answer: a

Rationale: a. correctCorrect.

b. a-A pro-ddrug is an inactive pre-cursor of a drug prior to conversion to an active metabolite. Commented [IP15]: hyphen

c. protein Protein reservoir binding occurs when a portion of a drug is bound to proteins

but may be released for us at a later time.

d. plasma-Plasma binding occurs with proteins as described above in answer c.

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- 14. A medications half-life refers to the:
 - a. Concentration of a drug
 - b. Dose required to be therapeutic
 - c. Time required for the total dose of a medication to be decreased by onehalf
 - d. Removal of a medication from the body

Answer: c

Rationale: a. concentration <u>Concentration</u> refers to the amount of drug available.

b. <u>the The</u> therapeutic index is the difference between minimal therapeutic dose and toxic level of dose.

c. correctCorrect.

d. this This defines the process of elimination.

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15. The ______ is the length of time that a medication concentration

is sufficient in the blood to produce a therapeutic response.

- a. Half-life
- b. Duration of action
- c. Efficacy
- d. Affinity

Answer: b

Rationale: a. half-Half-life is the time required for one-half of the dose of a medication to be removed from the body.

b. correctCorrect.

c. efficacy Efficacy is the ability of a drug to produce the expected response.

d. affinity Affinity is the force of attraction between a drug and its receptor site.

Commented [IP16]: change to medication? no

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16. Which of the following is the <u>most_MOST</u> common way that medications exert their actions?

- a. Receptor site binding.
- b. Changing the physical properties of a cell.
- c. Chemically combining with another chemical.
- d. Altering the normal metabolic pathway.

Answer: a

Rationale. a. confecteoneet.	Rationale: a.	correctCorrect.
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b. while While all answers listed are ways in which a drug may exert its actions, receptor

site binding is the most common.

c. while While all answers listed are ways in which a drug may exert its actions, receptor

site binding is the most common.

d. <u>while While</u> all answers listed are ways in which a drug may exert its actions, receptor site binding is the most common.

Commented [IP17]: Same sentence

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17. A medication such as nalbuphine binds with a specific receptor site causing an expected effect, but blocks another medication from triggering the receptor. -This

is known as a/an:

- a. Agonist
- b. Antagonist
- c. Agonist-antagonist
- d. Beta blocker

Answer: c

Rationale: a. an <u>An</u> agonist binds to a specific receptor causing a desired response.

b. an-An antagonist blocks a receptor site, but does not produce the expected result.

- c. correctCorrect.
- d. <u>a-A</u>beta blocker is an example of an antagonistic effect.

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18. ______ is the relative amount of a medication required to produce a desired response.

conca response.

- a. Antagonism
- b. Efficacy
- c. Therapeutic threshold
- d. Potency

Answer: d

Rationale: a. an-An antagonism produces a blocking effect at a receptor site.

b. efficacy Efficacy is the power of a medication to produce a therapeutic effect.

c. therapeutic Therapeutic threshold is the minimum concentration of a medication

required to cause a desired response.

d. correct.

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19. If a drug exceeds its therapeutic index, the drug will:

- a. Cause the desired response
- b. Be toxic to the patient
- c. Not have any effect on the patient
- d. Be present in the minimal level and may not cause the desired response

Answer: b

Rationale: a. for For a drug to cause a desired response, it must remain within the therapeutic index.

b. the <u>The</u> difference between the minimum effective concentration and the toxic level is the therapeutic index.

c. if If a drug exceeds the therapeutic index, it will be toxic to the patient.

d. the The minimal dose required is known as the minimum effective concentration.

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- 20. Which of the following statements is <u>true-TRUE</u> regarding medication administration to the children and elderly?
 - a. Liver and kidney functions in both populations is Liver and kidney functions in both populations are altered leaving both susceptible to altered medication responses.
 - b. Doses need to be significantly higher in both populations to produce the desired effect due to decreased liver functions.
 - Both populations should be treated the same as all patients as the therapeutic index does not change.
 - d. Pediatric patients have well-developed livers, while elderly patients suffer from decreased liver function.

Answer: a

Rationale: a. correctCorrect.

b. doses-<u>Doses</u> need to be altered accordingly to the appropriate population being treated.
c. while-<u>While</u> the therapeutic index may the change, the patient's ability to metabolize or excrete the medication will affect how much drug may be available.

d. the <u>The</u> infant population does not have well developed renal and hepatic function.

Commented [IP18]: Hyphen? yes

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