

## Chapter 01

### Multiple Choice

*Identify the choice that best completes the statement or answers the question.*

- \_\_\_\_ 1. The use of specific drugs to prevent, treat, or diagnose disease is known as
  - a. toxicology
  - b. pharmacokinetics
  - c. pharmacotherapeutics
  - d. toxicokinetics
  - e. dynamic equilibrium
  
- \_\_\_\_ 2. The study of how the body deals with a drug in terms of the way the drug is absorbed, distributed, and eliminated is known as
  - a. toxicology
  - b. pharmacodynamics
  - c. pharmacy
  - d. pharmacokinetics
  - e. biodynamics
  
- \_\_\_\_ 3. Toxicology is
  - a. the study of the harmful effects of chemicals
  - b. the study of a drug's beneficial effects
  - c. the analysis of drug absorption, distribution, and metabolism
  - d. the preparation and dispensing of therapeutic medications
  - e. the analysis of a drug's molecular structure
  
- \_\_\_\_ 4. The generic name of a drug
  - a. is also known as the "official" or "nonproprietary" name
  - b. is often derived from the chemical name
  - c. tends to be somewhat shorter than the drug's chemical name
  - d. all of the above are true
  
- \_\_\_\_ 5. If there is no existing patent for a drug, or if the patent has expired, the same drug may be marketed by separate drug companies under different
  - a. chemical names
  - b. generic names
  - c. trade names
  - d. nonproprietary names
  - e. organic names
  
- \_\_\_\_ 6. The generic form of a drug is considered to be as safe and effective as the original, brand-name product if the generic form \_\_\_\_\_ as the brand-name drug.
  - a. has the same type and amount of the active ingredient(s)
  - b. uses the same administration route
  - c. has the same pharmacokinetic profile (drug absorption, plasma levels, and so forth)
  - d. produces the same therapeutic effects
  - e. all the above are true

- \_\_\_\_ 7. During drug testing and approval, the drug is usually tested in a relatively small sample (200 to 300 people) with a specific disease or pathologic condition during
- preclinical trials
  - phase 1 clinical trials
  - phase 2 clinical trials
  - postmarketing surveillance
- \_\_\_\_ 8. Drugs that can be purchased directly by the consumer without a prescription are also known as
- nonproprietary medications
  - Schedule I controlled substances
  - Schedule II controlled substances
  - Schedule III controlled substances
  - over-the-counter drugs
- \_\_\_\_ 9. The point at which there is no further increase in the response to a drug even if the dosage continues to be increased is known as the
- ceiling effect
  - maximal efficacy
  - potency
  - all of the above
  - A and B only
- \_\_\_\_ 10. When two drugs are compared, the drug that requires a lower dosage to produce the same effect as a higher dose of the second drug is said to
- be more potent
  - be less potent
  - have a great maximal efficacy
  - have a greater therapeutic index
  - be pharmacosuperior
- \_\_\_\_ 11. When evaluating drug safety, the dosage that causes 50 percent of subjects to exhibit a specific adverse effect is known as the
- median therapeutic dose
  - median toxic dose
  - therapeutic index
  - ceiling effect
  - threshold dose
- \_\_\_\_ 12. In general, the greater the value of the \_\_\_\_\_, the safer the drug is considered to be.
- median effective dose
  - threshold dose
  - therapeutic index (TI)
  - ceiling effect
  - potency index
- \_\_\_\_ 13. Regarding drug development and approval, an “orphan drug” is a drug that is
- prescribed for conditions other than those approved by the FDA
  - given special funding for development because it is used in a small patient population with a relatively rare disease
  - available directly to consumers without a prescription
  - only available in countries outside the United States

- \_\_\_\_\_ 14. Prescription use of a drug to treat conditions other than those that the drug was originally approved to treat (off label prescribing)
- a. is illegal and punishable by revoking a physician's license
  - b. is legal only if the Center for Disease Control provides written permission to the physician
  - c. is legal only if the drug is not a controlled substance
  - d. is legal and quite common in the United States
  - e. is legal only after the patent for a drug has expired

## **Chapter 01**

### **Answer Section**

#### **MULTIPLE CHOICE**

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|------------|--------|
| 1. ANS: C  | PTS: 1 |
| 2. ANS: D  | PTS: 1 |
| 3. ANS: A  | PTS: 1 |
| 4. ANS: D  | PTS: 1 |
| 5. ANS: C  | PTS: 1 |
| 6. ANS: E  | PTS: 1 |
| 7. ANS: C  | PTS: 1 |
| 8. ANS: E  | PTS: 1 |
| 9. ANS: E  | PTS: 1 |
| 10. ANS: A | PTS: 1 |
| 11. ANS: B | PTS: 1 |
| 12. ANS: C | PTS: 1 |
| 13. ANS: B | PTS: 1 |
| 14. ANS: D | PTS: 1 |