

Chapter 1: Drug Definitions, Standards, and Information Sources

Test Bank

MULTIPLE CHOICE

1. What is the name under which a drug is listed by the U.S. Food and Drug Administration (FDA)?

- a. Brand
- b. Nonproprietary
- c. Official
- d. Trademark

ANS: C

The official name is the name under which a drug is listed by the FDA. The brand name, or trademark, is the name given to a drug by its manufacturer. The nonproprietary, or generic, name is provided by the U.S. Adopted Names Council.

DIF: Cognitive Level: Knowledge REF: p. 1 OBJ: 2

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Safe, Effective Care Environment

2. Which source contains information specific to nutritional supplements?

- a. *USP Dictionary of USAN & International Drug Names*
- b. *Natural Medicines Comprehensive Database*
- c. *United States Pharmacopoeia/National Formulary (USP NF)*
- d. *Drug Interaction Facts*

ANS: C

United States Pharmacopoeia/National Formulary contains information specific to nutritional supplements. *USP Dictionary of USAN & International Drug Names* is a compilation of drug names, pronunciation guide, and possible future FDA approved drugs; it does not include nutritional supplements. *Natural Medicines Comprehensive Database* contains evidence based information on herbal medicines and herbal combination products; it does not include information specific to nutritional supplements. *Drug Interaction Facts* contains comprehensive information on drug interaction facts; it does not include nutritional supplements.

DIF: Cognitive Level: Knowledge REF: p. 2 OBJ: 4

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Physiological Integrity

3. What is the most comprehensive reference available to research a drug interaction?

- a. *Drug Facts and Comparisons*
- b. *Drug Interaction Facts*
- c. *Handbook on Injectable Drugs*
- d. *Martindale—The Complete Drug Reference*

ANS: B

First published in 1983, *Drug Interaction Facts* is the most comprehensive book available on drug interactions. In addition to monographs listing various aspects of drug interactions, this information is reviewed and updated by an internationally renowned group of physicians and pharmacists with clinical and scientific expertise.

DIF: Cognitive Level: Comprehension REF: p. 3 OBJ: 3

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Physiological Integrity

4. The physician has written an order for a drug with which the nurse is unfamiliar. Which section of the *Physicians' Desk Reference (PDR)* is most helpful to get information about this drug?

- a. Manufacturer's section
- b. Brand and Generic Name section
- c. Product Category section
- d. Product Information section

ANS: B

A physician's order would include the brand and/or generic name of the drug. The alphabetic index in the *PDR* would make this section the most user friendly. Based on a physician's order, manufacturer's information and classification information would not be known. The Manufacturer's section is a roster of manufacturers. The Product Category section lists products subdivided by therapeutic classes, such as analgesics, laxatives, oxytocics, and antibiotics. The Product Information section contains reprints of the package inserts for the major products of manufacturers.

DIF: Cognitive Level: Comprehension REF: p. 3 OBJ: 4

TOP: Nursing Process Step: Planning

MSC: NCLEX Client Needs Category: Physiological Integrity

5. Which online drug reference makes available to health care providers and the public a standard, comprehensive, up to date look up and downloadable resource about medicines?

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- a. *American Drug Index*
 - b. *American Hospital Formulary*
 - c. DailyMed
 - d. *Physicians' Desk Reference (PDR)*
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ANS: C

DailyMed makes available to health care providers and the public a standard, comprehensive, up to date look up and downloadable resource about medicines. The *American Drug Index* is not appropriate for patient use. The *American Hospital Formulary* is not appropriate for patient use. The *PDR* is not appropriate for patient use.

DIF: Cognitive Level: Knowledge REF: p. 4 OBJ: 5

TOP: Nursing Process Step: Implementation

MSC: NCLEX Client Needs Category: Physiological Integrity

6. Which legislation authorizes the FDA to determine the safety of a drug before its marketing?

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- a. Federal Food, Drug, and Cosmetic Act (1938)
 - b. Durham Humphrey Amendment (1952)
 - c. Controlled Substances Act (1970)
 - d. Kefauver Harris Drug Amendment (1962)
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ANS: A

The Federal Food, Drug, and Cosmetic Act of 1938 authorized the FDA to determine the safety of all drugs before marketing. Later amendments and acts helped tighten FDA control and ensure drug safety. The Durham Humphrey Amendment defines the kinds of drugs that cannot be used safely without medical supervision and restricts their sale to prescription by a licensed practitioner. The Controlled Substances Act addresses only controlled substances and their categorization. The Kefauver Harris Drug Amendment ensures drug efficacy and greater drug safety. Drug manufacturers are required to prove to the FDA the effectiveness of their products before marketing them.

DIF: Cognitive Level: Knowledge REF: p. 4 OBJ: 8

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Physiological Integrity

7. Meperidine (Demerol) is a narcotic with a high potential for physical and psychological dependency. Under which classification does this drug fall?

-
- a. I
 - b. II
 - c. III
 - d. IV
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ANS: B

Meperidine (Demerol) is a Schedule II drug; it has a high potential for abuse and may lead to severe psychological and physical dependence. Schedule I drugs have high potential for abuse and no recognized medical use. Schedule III drugs have some potential for abuse. Use may lead to low to moderate physical dependence or high psychological dependence. Schedule IV drugs have low potential for abuse. Use may lead to limited physical or psychological dependence.

DIF: Cognitive Level: Comprehension REF: p. 4 | p. 5 OBJ: 7

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Safe, Effective Care Environment

8. What would the FDA do to expedite drug development and approval for an outbreak of smallpox, for which there is no known treatment?

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- a. List smallpox as a health orphan disease.
 - b. Omit the preclinical research phase.
 - c. Extend the clinical research phase.
 - d. Fast track the investigational drug.
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ANS: D

Once the Investigational New Drug Application has been approved, the drug can receive highest priority within the agency, which is called fast tracking. A smallpox outbreak would become a priority concern in the world. Orphan diseases are not researched in a priority manner. Preclinical research is not omitted. Extending any phase of the research would mean a longer time to develop a vaccine. The FDA must ensure that all phases of the preclinical and clinical research phase have been completed in a safe manner.

DIF: Cognitive Level: Knowledge REF: p. 7 OBJ: 8

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Safe, Effective Care Environment

9. Which statement is true about over the counter (OTC) drugs?

- a. They are not listed in the *USP NF*.
- b. A prescription from a health care provider is needed.
- c. They are sold without a prescription.
- d. They are known only by their brand names.

ANS: C

OTC medications do not require a prescription. A variety of names, both generic and trade, can be used for individual drugs sold OTC. OTC drugs are listed in the *USP NF*. Prescription drugs require an order by a health professional who is licensed to prescribe, such as a physician, nurse practitioner, physician assistant, or dentist.

DIF: Cognitive Level: Comprehension REF: p. 2 OBJ: 2

TOP: Nursing Process Step: Planning

MSC: NCLEX Client Needs Category: Physiological Integrity

10. Which is the most authoritative reference for medications that are injected?

- a. *Physician's Desk Reference*
- b. *Handbook on Injectable Drugs*
- c. DailyMed
- d. *Handbook of Nonprescription Drugs*

ANS: B

The *Handbook on Injectable Drugs* is the most comprehensive reference available on the topic of compatibility of injectable drugs. It is a collection of monographs for more than 300 injectable drugs that are listed alphabetically by generic name.

DIF: Cognitive Level: Comprehension REF: p. 3 OBJ: 4

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Physiological Integrity

11. The nurse is administering Lomotil, a Schedule V drug. Which statement is true about this drug's classification?

- a. Abuse potential for this drug is low.
- b. Psychological dependency is likely.
- c. There is a high potential for abuse.
- d. This drug is not a controlled substance.

ANS: A

Lomotil, a Schedule V drug, has an abuse potential of limited physical or psychological dependence liability compared with drugs in Schedule IV. Because abuse potential is low with a Schedule V drug, a prescription may not be required. Psychological dependency is not likely with a Schedule V drug. Schedule V drugs are classified as controlled substances.

DIF: Cognitive Level: Knowledge REF: p. 5 OBJ: 7

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Physiological Integrity

12. The nurse is transcribing new orders written for a patient with a substance abuse history. Choose the medication ordered that has the greatest risk for abuse.

- a. Lomotil
- b. Diazepam
- c. Phenobarbital
- d. Lortab

ANS: D

Lortab is a Schedule III drug with a high potential for abuse but less so than drugs in Schedules I and II. Lomotil is a Schedule V drug with a low potential for abuse compared with those in Schedule V. Diazepam is a Schedule IV drug with a low potential for abuse compared with those in schedule III. Phenobarbital is a Schedule IV drug with a low potential for abuse compared with those in Schedule III.

DIF: Cognitive Level: Application REF: pp. 4-5 OBJ: 7

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Physiological Integrity

MULTIPLE RESPONSE

13. The nurse is caring for a patient newly diagnosed with type 1 diabetes mellitus. Which approach(es) to therapeutic methods would be considered in this patient's treatment? (*Select all that apply.*)

- a. Therapeutic drugs
- b. Concentrated carbohydrate diet
- c. Family centered care
- d. Regular daily exercise and activity
- e. Daily baths

ANS: A, B, D

Therapeutic methods include drug therapy, diet therapy, physiotherapy, and psychological therapy. Therapeutic methods do not include family centered care or daily baths.

DIF: Cognitive Level: Application REF: p. 1 OBJ: 1

TOP: Nursing Process Step: Planning

MSC: NCLEX Client Needs Category: Physiological Integrity

14. An older adult experiencing shortness of breath is brought to the hospital by her daughter. While obtaining the medication history from the patient and her daughter, the nurse discovers that neither has a list of the patient's current medications or prescriptions. All the patient has is a weekly pill dispenser that contains four different pills. The prescriptions are filled through the local pharmacy. Which resource(s) would be appropriate to use in determining the medication names and doses? (*Select all that apply.*)

- a. *Martindale—The Complete Drug Reference*
- b. *Physicians' Desk Reference*, Section 4
- c. Senior citizens' center
- d. Patient's home pharmacy

ANS: B, D

The *Physicians' Desk Reference*, Section 4, has full color images of commonly dispensed tablets and capsules. The patient's pharmacy would have an accurate account of all the medications the client is currently taking. *Martindale—The Complete Drug Reference* has written information on medications and would not be an appropriate resource. The senior citizens' center is not likely to have specific patient medication information.

DIF: Cognitive Level: Application REF: p. 3 OBJ: 3 | 4

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Physiological Integrity

15. The nurse planning patient teaching regarding drug names would include which statement(s)?(Select all that apply.)

- a. Most drug companies place their products on the market under generic names.
- b. The official name is the name under which the drug is listed by the U.S. Food and Drug Administration (FDA).
- c. Brand names are easier to pronounce, spell, and remember.
- d. The first letter of the generic name is not capitalized.
- e. The chemical name is most meaningful to the patient.

ANS: B, C, D

The official name is the name under which the drug is listed by the FDA. Brand names are easier to pronounce, spell, and remember. The first letter of the generic name is not capitalized. Most drug companies place their products on the market under brand names instead of generic names. The chemical name is most meaningful to the chemist.

DIF: Cognitive Level: Application REF: p. 1 OBJ: 2

TOP: Nursing Process Step: Planning

MSC: NCLEX Client Needs Category: Physiological Integrity

16. When categorizing, the nurse is aware that which drug(s) would be considered Schedule II?(Select all that apply.)

- a. Marijuana
- b. Percodan
- c. Amphetamines
- d. Fiorinal
- e. Flurazepam

ANS: B, C

Schedule II drugs have a high potential for abuse, they are currently accepted in the United States, and use may lead to severe psychological or physical dependence. Percodan and amphetamines are considered Schedule II drugs. Marijuana is a Schedule I drug. Fiorinal is a Schedule III drug. Flurazepam is a Schedule IV drug.

DIF: Cognitive Level: Analysis REF: pp. 4-5 OBJ: 7

TOP: Nursing Process Step: Assessment

MSC: NCLEX Client Needs Category: Physiological Integrity