

Chapter 02: The Drug Approval Process Test Bank

MULTIPLE CHOICE

1. The nurse is preparing to administer a schedule II injectable drug and is drawing up half of the contents of a single-use vial. Which nursing action is correct?
- Ask another nurse to observe and cosign wasting the remaining drug from the vial.
 - Keep the remaining amount in the patient's drawer to give at the next dose.
 - Record the amount unused in the patient's medication record.
 - Dispose of the vial with the remaining drug into a locked collection box.

ANS: A

Schedule II drugs are controlled substances, and all must be accounted for. When wasting a portion of a drug, another nurse should observe and cosign that a drug was wasted.

DIF: COGNITIVE LEVEL: Applying (Application)

REF: Page 16

TOP: NURSING PROCESS: Nursing Intervention

MSC: NCLEX: Physiological Integrity: Pharmacological and Parenteral Therapies

2. A patient is prescribed a medication and asks the nurse if the drug is available in a generic form. The nurse understands that a generic medication will have a name that
- is a registered trademark.
 - is always capitalized.
 - describes the drug's chemical structure.
 - is non-proprietary.

ANS: D

The generic name is the official, non-proprietary name for a drug. The brand name is the trademark name and is always capitalized. The chemical name describes the chemical structure of the drug.

DIF: COGNITIVE LEVEL: Understanding (Comprehension)

REF: Page 18

TOP: NURSING PROCESS: N/A

MSC: NCLEX: Physiological Integrity: Pharmacological and Parenteral Therapies

3. A patient receives a prescription on which the provider has noted that a generic medication may be given. The patient asks the nurse what this means. What will the nurse tell the patient about generic drugs?
- They contain the same inert ingredients as brand-name drugs.
 - They have chemical structures that are identical to proprietary drugs.
 - They tend to be less expensive than brand-name drugs.
 - They undergo extensive testing before they are marketed.

ANS: C

Generic drugs are approved by the FDA if they are proved to be bioequivalent to the brand-name drug. They tend to be less expensive because manufacturers of these drugs do not have to do the extensive testing required of brand-name drugs before marketing. They are not identical to brand-name drugs and often have different inert ingredients.

DIF: COGNITIVE LEVEL: Applying (Application)

REF: Page 18

Kee Pharmacology Chapter 1test Bank

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