

**Final Exam Sample Questions for
Biopharmaceutics and Pharmacokinetics
PT606 Pharm D Students**

1. If an IV bolus of a drug declines to 30% of the dose after 8 hours. What is the elimination half-life of the drug assuming first-order kinetics?

- a. 2.3 hours
- c. 4.6 hours.**
- e. 1.6 hours
- b. 3.7 hours.
- d. 6.3 hours.

2. Assumptions for a one compartment model are:

- a. Body is homogeneous
- b. Distribution instantaneous
- c. Drug concentration proportional to dose
- d. Rate of elimination proportional to how much is there.
- e. All the above**

Questions 3-5 apply to the following:

A drug was given as an intravenous bolus to a 70 Kg patient in a dose of 80 mg. The elimination follows first order kinetics. Two hours after drug administration the concentration was found to be 1.48 µg/mL. Four hours after administration drug concentration was 0.74 µg/mL.

3- What is the initial plasma drug concentration directly after the injection?

- a) 1.9 µg/mL
- b) 2.9 µg/mL.**
- c) 3.9 µg/mL
- d) 4.9 µg/mL
- e) none of the above

4- What is the elimination half-life of the drug?

- a) 1 hour
- b) 2 hour**
- c) 3 hour
- d) 4 hour
- e) 5 hour

5- What is the amount of the drug in the body after 4 hours of the injection?

- a) 19.9 mg**
- b) 19.9 µg
- c) 19.9 mg/L
- d) 19.9 µg/L
- e) 19.9 L

Questions 6-7 apply to the following

A drug was given in an intravenous dose of 2.29 mg/Kg to a 70 Kg patient. The drug follows two compartment model with an equation describing its kinetics as follows:

$$C_p = 8.2 e^{-3.45t} + 2.5 e^{-0.049t}$$

6. What is the plasma level after 5 hours of the IV dose?

- a) 2.368 µg/mL
- b) 1.968 µg/mL.**
- c) 4.689 µg/mL
- d) 5.142 µg/mL
- e) 10.254 µg/mL

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Barbara Lacher



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Pharmacokinetics will help you to Understand the basic concepts in biopharmaceutics and pharmacokinetics Use raw data and derive the pharmacokinetic models and parameters that best describe the process of drug absorption distribution and elimination Critically evaluate biopharmaceutic studies involving drug product equivalency and unequivalency Design and evaluate dosage regimens of drugs using pharmacokinetic and biopharmaceutic parameters Detect potential clinical pharmacokinetic problems and apply basic pharmacokinetic principles to solve them Practical problems and clinical examples with discussions are included in each chapter to help you apply these principles to patient care and drug consultation situations Chapter Objectives Chapter Summaries and Frequently Asked Questions along with additional application questions appear within each chapter to identify and focus on key concepts Most of the chapters have been revised to reflect our current understanding of drug product performance bioavailability bioequivalence pharmacokinetics pharmacodynamics and drug therapy **Applied Biopharmaceutics & Pharmacokinetics, Fifth Edition** Leon Shargel, Susanna Wu-Pong, Andrew Yu, 2004-08-19 The most comprehensive text on the practical applications of biopharmaceutics and pharmacokinetics 4 STAR DOODY S REVIEW The updated edition provides the reader with a solid foundation in the basic principles of pharmacokinetics and biopharmaceutics Students will be able to apply the information to their clinical practice and researchers will find this to be a valuable reference This modestly priced book should be the gold standard for student use Doody s Review Service The primary emphasis of this book is on the application and understanding of concepts Basic theoretical discussions of the principles of biopharmaceutics and pharmacokinetics are provided along with illustrative examples and practice problems and solutions to help the student gain skill in practical problem solving

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Candidate's Review Guide to the Pharmacist Licensure Examination California. State Board of Pharmacy,1996

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