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Clinical Pharma

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Equations And

Calculations

Calculations

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PHARMACOLOGY  
EQUATIONS for  
USMLE STEP 1

---

259 FA 12 : PHARMA  
COKINETICS  
EQUATIONS WITH  
EXAMPLES

---

Pharmacokinetics:  $V_d$ ,  
Clearance, Half-life:  
Calculation Drug  
Distribution,

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Elimination, Rate<sup>259</sup>

FA 12 : PHARMACO  
KINETICS

EQUATIONS PART 1

One compartment  
model calculations | |

Pharmacokinetics

Calculations -

Bioavailability and

Pharmacokinetics First

~~Order Elimination Rate~~

~~Constant and Half-life |~~

~~A closer look - Lect 11~~

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**PHARMACOKINETICS**

**EQUATIONS PART 3**

Applied Pharmacology

7. Drug dose

calculations John

~~Murphy talks about~~

~~Basic \u0026 Applied~~

~~Pharmacokinetics Self~~

~~Assessment~~

---

Medical Pharmacology:

Pharmacokinetics -

Steady State

Concentration

Pharmacology - PHAR

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MACOKINETICS

(MADE EASY)

Pharmacokinetics:

Volume Of Distribution

animation video

---

Volume of distribution

of drugs

Calculation of

Steady state

concentration on IV

infusion ~~Beer's Law~~

~~Unknown Calculation~~

~~How to Calculate AUC~~

~~Calculation of the area~~

~~under the plasma~~

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~~concentration vs. time~~

curve HOW TO

STUDY

PHARMACOLOGY!

Clearance \u0026amp; Half-

Life - The

Pharmacokinetics Series

Bioavailability And

Intravenous Versus Oral

Administration

Pharmacokinetics

animation: Dosing

Interval Pharmacy

~~Calculations - The~~



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Basics pKa and Drug  
Solubility: Absorption  
and Distribution –  
Pharmacokinetics (PK)

| Lecturio

Pharmacokinetics in  
Patients Requiring  
Renal Replacement  
Therapy Part 1 -

Module 4, Session 1

Pharmacokinetics:

Analyzing

Concentration Data

(Bio) Pharmacokinetics

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~~in Clinical Practice (1.  
Basic Concepts and  
Clinical Relevance)  
Bioequivalence +  
Bioavailability and  
Bioequivalence +  
Biopharmaceutics and  
Pharmacokinetics +  
Clinical Nursing  
Calculations: Teach  
Your Students to  
Medicate Safely Volume  
of Distribution -  
Pharmacology Lect 5~~

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## Clinical Pharmacokinetic Equations And Calculations

Useful Pharmacokinetic  
Equations. Symbols. e.

D = dose = dosing  
interval CL = clearance

Vd = volume of  
distribution ke =

elimination rate

constant ka = absorption  
rate constant F =

fraction absorbed

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(bioavailability)  $K_0 =$   
infusion rate  $T =$   
duration of infusion  $C =$   
plasma concentration.  
General. Elimination  
rate constant.  $k$   $CL$   $V_d$   
 $C$   $C_{tt}$   $CC$ .

## Useful Pharmacokinetic Equations

Clinical  
pharmacokinetic dosage  
calculations are  
conducted using the

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easiest possible equations and methods that produce acceptable results. This is because there are usually only a few (sometimes as little as 1-2) drug serum concentrations on which to base the calculations. Drug serum concentrations are expensive (typically \$35-100 each), and obtaining them can

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cause minor discomfort  
and trauma to the  
patient.  
Equations And  
Calculations

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Pharmacokinetic  
Equations And  
Calculations Clinical  
pharmacokinetic dosage  
calculations are

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conducted using the easiest possible equations and methods that produce acceptable results. This is because there are usually only a few (sometimes as little as 1-2) drug serum concentrations on which to base the calculations.

Clinical  
Pharmacokinetic  
Equations And

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### Calculations

Clinical pharmacokinetic dosage calculations are conducted using the easiest possible equations and methods. This is because there are usually only a few (sometimes as little as 1 – 2) drug serum concentrations on which to base the calculations.



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## Chapter 2. Clinical Pharmacokinetic Equations and Calculations

Clinical  
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Calculations ... Clinical  
pharmacokinetic dosage  
calculations are  
conducted using the  
easiest possible  
equations and methods.  
This is because there are

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usually only a few  
(sometimes as little as  
1 – 2) drug serum  
concentrations on which  
to base the calculations.

## Chapter 2. Clinical Pharmacokinetic Equations and Calculations

### Clinical Pharmacokinetic Equations And Calculations

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clinical pharmacokinetic

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## Clinical Pharmacokinetic Equations And Calculations

- In pharmacokinetic calculations, the term  $e^{-kel(t)}$  represents the fraction of the serum concentration that remains. Thus,  $1 - e^{-kel(t)}$  represents the fraction of the serum concentration that is eliminated.  $t_{1/2}$  or

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Half-life • The time required for the TOTAL amount of remaining drug in the body to decline by 50%.<sup>1</sup>

Pharmacokinetic  
Training Packet for  
Pharmacists  
Clinical  
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Calculations Getting the

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of the options to  
accompany you next

## Clinical Pharmacokinetic Equations And Calculations

Evidence-based clinical  
decision support tools  
and calculators for

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medical professionals.

Includes mobile applications, advanced pharmacokinetic

utilities, and a wealth of evidence-based medicine.

Clinical tools and calculators for medical professionals ...

With a known CL vanco and Vd, an elimination constant ( $K_{el}$ ) can be



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calculated ( $K_{el} = C L_{vanco} / V_d$ ) Once the most likely values of  $K_{el}$  and  $V_d$  have been estimated, one-compartment pharmacokinetic equations are used to identify a dose and its associated peak, trough, and AUC/MIC values.

Vancomycin Calculator  
- ClinCalc.com

*Page 25/38*

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### Basic Pharmacokinetics

**OBJECTIVES** After completing Lesson 2, you should be able to: 1.

Define the concept of apparent volume of distribution and use an appropriate mathematical equation to calculate this parameter. 2. Identify the components of body fluids that make up extracellular and

# Read PDF Clinical Pharma intracellular fluids and know the percentage of each ... Calculations

## Concepts in Clinical Pharmacokinetics, 6th Edition ...

Formula | Volume of  
Distribution = Total  
Dose / Concentration  
Say that a question asks  
you to determine the  
volume of distribution  
(VD) of a drug with a

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total dose of 2,000 mg  
and a concentration...

## How to Simplify Pharmacokinetics Calculations | by ...

Author summary The use of orally inhaled drugs for treating lung diseases is appealing since they have the potential for lung selectivity, i.e. high exposure at the site of

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action — the lung —  
without excessive side  
effects. However, the  
degree of lung selectivity  
depends on a large  
number of factors,  
including  
physiochemical  
properties of drug  
molecules, patient  
disease state, and ...

A mechanistic  
framework for a priori

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pharmacokinetic ...

Pharmacokinetics provides a mathematical basis to assess the time course of drugs and their effects in the body. It enables the following processes to be quantified: Absorption  
Distribution Metabolism  
Excretion These pharmacokinetic processes, often referred to as **ADME**, determine

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the drug concentration  
in the body when  
medicines are  
prescribed. A

Basic pharmacokinetics  
- Pharmaceutical Press

This website currently  
uses the Bauer equation  
which works well. On  
the update this Fall  
2020 I will update the  
CLvanco equation:

CLvanco (L/hr) =

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$0.06 \times (0.70 \times \text{CrCl} + 8)$ .

The equation is based on data from over 1300 SS peak and trough levels. CrCl can be calculated with the Cockcroft-Gault equation, with an adjusted BW used for overweight patients.

Vancomycin

Pharmacokinetics

Review - VancoPK



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Equation (6.1) describes the changes in mass of absorbable drug over time at the site of

administration.  $\frac{dX}{dt} = K_a X - K_e X$  (6.1) where  $\frac{dX}{dt}$  is the decrease in the amount of absorbable drug present at the site of

administration per unit time (e.g., mg h<sup>-1</sup>);  $K_a$  is the first-order absorption rate constant (h<sup>-1</sup>;

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kinetic  
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min); and  $X_a/t$  is the  
mass

Basic Pharmacokinetics

Sample Chapter

Practice problems for  
the calculations required  
when evaluating drug  
bioavailability or  
performing  
pharmacokinetics

LINKS Lecture -  
Pharmacokinetics &  
Bioava...

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## Calculations - Bioavailability and Pharmacokinetics -

### YouTube

proportional. Use the following equation and let 's target 15 mg/L in this case: Eq. 12 New

$$\text{Dose} = (1000 \text{ mg}/24 \text{ hr}) * 15 \text{ mg}/\text{dL} / 11.4$$

$$\text{mg}/\text{dL} = 54.8 \text{ mg}/\text{hr}$$

Generally, if the ratio of C desired / C measured

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is 1.5 or more then decrease the dosing interval ( $\tau$ ) from 12 hours to 8 hours (or 24 hours to 12 hours).

Cases: Gentamicin

& Vancomycin

Pharmacokinetics

Pharmacokinetic models are useful to: A) describe concentration-time data sets. B) predict drug serum concentrations

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after several doses or  
after different routes of  
administration. C)  
calculate  
pharmacokinetic  
constants (clearance,  
volume of distribution,  
half-life). D) a and c: E)  
a, b, and c

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