

Clinical Pharmacokinetics Part 1

Rational Dosing and the Time Course of Drug Action

Leslie Goldstein, Pharm.D.

Associate Professor

Department of Clinical Specialties

lgolds01@nyit.edu

**NEW YORK INSTITUTE
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I am available to groups and individuals for pharmacology help and discussions by appointment.

lgolds01@nyit.edu

After completing the preparation materials, students should be able to:

1. Describe clearance, volume of distribution, bioavailability, elimination half-life, steady state, extraction ratio, and area under the curve using pharmacokinetic models.
2. Differentiate first-order and zero-order elimination kinetics and linear and non-linear processes in relation to drug concentration and effect.
3. Graph drug accumulation, half-life, and steady state for drugs with first-order elimination administered by continuous and intermittent dosing.
4. Calculate the loading dose and maintenance dose for an individual patient to achieve a target concentration within the therapeutic window when given specific pharmacokinetic parameters.
5. Identify pharmacokinetic variables that influence drug plasma concentrations and the criteria for therapeutic drug monitoring.

Preparation Materials (links are in the CPG and on the next slide)

Required

- ScholarRx Bricks | Practice Questions

Highly relevant optional materials:

- Dr. Goldstein's Word handout | Lecture Videos | Guided Reading Questions
- Textbook resources with links are listed on the next slide

SUGGESTIONS:

- *Use the resources that work best for you.*
- *You do not need to study all of them.*
- *Work through the GUIDED READING QUESTIONS with pen/pencil and paper.
Try answering them in your OWN WORDS first without looking at the readings, and then again after reviewing.*
- *Practice questions (not graded): Simple Recall and Case Vignettes*

Links

Scholar Rx Bricks: (Required)

General Principles

Pharmacokinetics and Pharmacodynamics > Pharmacokinetics: Drug Concentration and Dosing

<https://exchange.scholarrx.com/brick/drug-concentration-and-dosing>

Enzymes as Drug Targets <https://exchange.scholarrx.com/brick/enzymes-as-drug-targets>

Katzung & Vanderah's Basic & Clinical Pharmacology, 16e, 2024; Chapter 3: Pharmacokinetics

<https://accessmedicine-mhmedical-com.nyit.idm.oclc.org/content.aspx?bookid=3382§ionid=281747069>

THE REVIEW BOOKS HAVE PRACTICE QUESTIONS

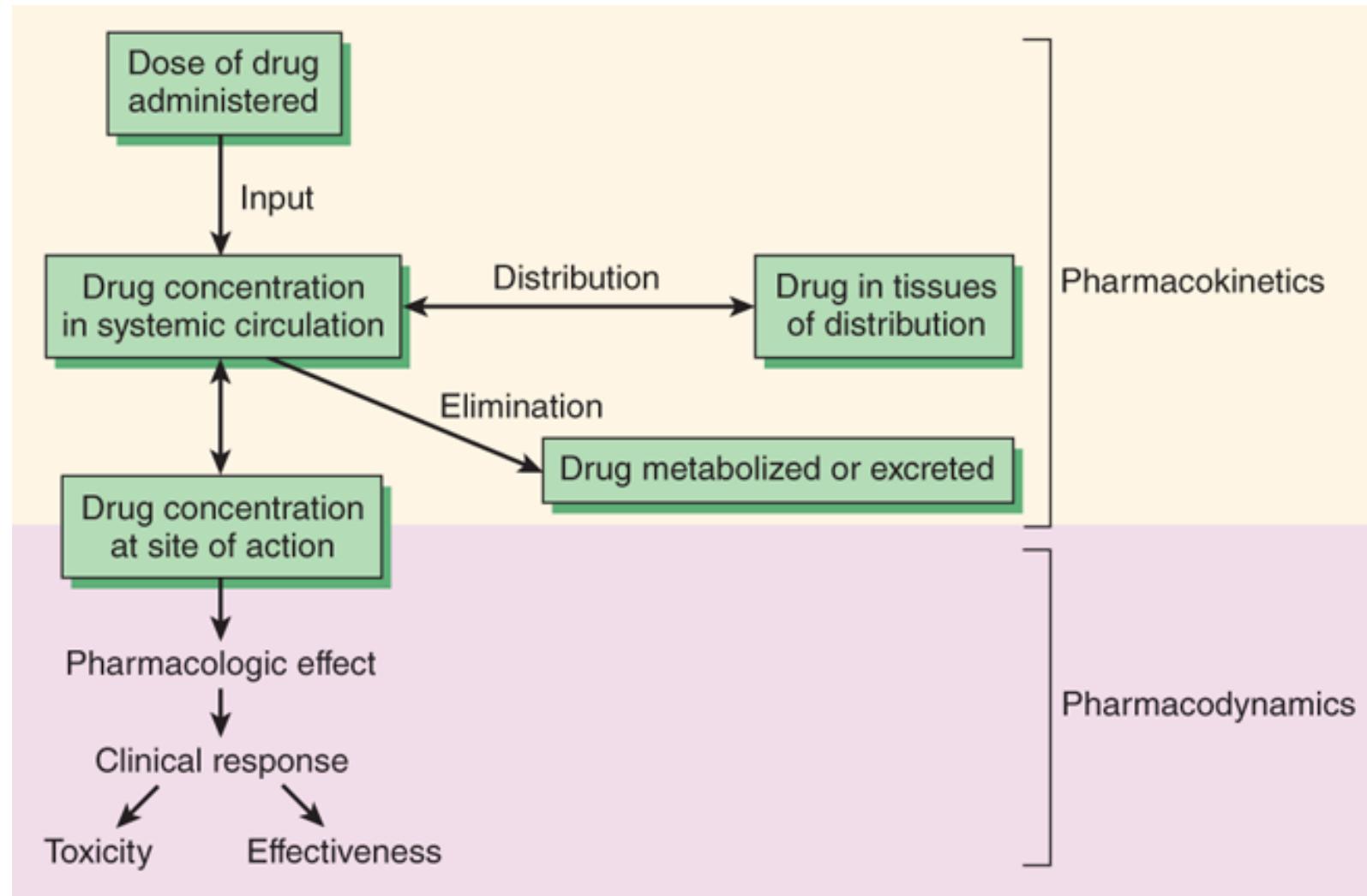
Katzung & Trevor's Pharmacology: Examination and Board Review, 14e, 2024

Chapter 3: Pharmacokinetics

<https://accessmedicine-mhmedical-com.nyit.idm.oclc.org/content.aspx?bookid=3461§ionid=285589076>

LWW Health Library Premium Basic Sciences; Lippincott Illustrated Reviews: Pharmacology, 8e, 2023: Chapter 1: Pharmacokinetics

<https://premiumbasicsciences-lwwhealthlibrary-com.nyit.idm.oclc.org/content.aspx?sectionid=253324942&bookid=3222>



A rational approach to achieving ***a desired beneficial effect with minimal adverse effects*** combines the principles of pharmacokinetics and pharmacodynamics to clarify the dose-effect relationship.

The relationship between dose and effect can be separated into pharmacokinetic (dose-concentration) and pharmacodynamic (concentration-effect) components. Concentration provides the link between pharmacokinetics and pharmacodynamics and is the focus of the target concentration approach to rational dosing. The three primary processes of pharmacokinetics are input, distribution, and elimination.

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Citation: Chapter 3 Pharmacokinetics & Pharmacodynamics: Rational Dosing & the Time Course of Drug Action, Katzung BG, Vanderah TW. *Basic & Clinical Pharmacology*, 15e; 2021, Figure 3-1. Available at: <https://accessmedicine.mhmedical.com/content.aspx?bookid=2988§ionid=250594300> Accessed: August 17, 2021
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Key points: What you need to know and understand

Terms

<p>Clearance (CL)</p> $CL = \text{elimination rate} / C_p$ <p>Thus,</p> $\text{Elimination rate} = CL \times C_p$ <p>C_p: plasma concentration</p>	<p>The volume of blood from which the drug is removed per unit of time: it is the proportionality factor that relates the rate of elimination to drug concentration.</p> <p>Units: volume/time (mL/min or L/h)</p>
<p>Bioavailability (F)</p>	<p>The fraction of the administered dose of drug that reaches the systemic circulation</p> <p>Intravenous (IV) administration: $F = 1$ (100%)</p>
<p>Apparent Volume of distribution (V_d)</p> $V_d = \text{Dose} / C_p$	<p>The proportionality factor that relates the amount of drug in the body to the concentration of drug in the blood or plasma</p> <p>Units: Liters or Liters/kg body weight</p>
<p>Elimination half-life ($t_{1/2}$)</p> $t_{1/2} = (0.693 \times V_d) / CL$ <p>Units: time</p>	<p>The time required for the amount of drug in the body or blood to fall by 50%. For drugs eliminated by first-order kinetics, this number is a constant regardless of the concentration.</p>

What you need to know and understand

Terms to know

<p>Extraction ratio $E_H = CL_H / Q$ and $F = 1 - ER$ H: hepatic; Q: blood flow</p>	<p>A measure of an organ's intrinsic capacity for eliminating a given drug from the systemic circulation over a single pass through the organ (usually liver or kidney)</p>
<p>Area under the concentration-time curve (AUC) $F = AUC_{oral} / AUC_{IV}$ $F = AUC_{formulation\ A} / AUC_{formulation\ B}$ Units: Cp x time (mg x min)/ml</p>	<p>The integration of the variation in plasma drug concentration over time after a single dose or during a single dosing interval AUC reflects <i>the actual (total) body exposure</i> to an administered dose of a drug, irrespective of the rate of absorption.</p>
<p>First-order elimination A constant <i>fraction</i> of drug is eliminated per unit of time</p>	<p>Drug clearance is a constant. That is, the ratio of rate of elimination to plasma concentration is the same over a broad range of plasma concentration. $t_{1/2}$ is constant.</p>
<p>Zero-order elimination A constant <i>amount</i> (not fraction) of drug is eliminated per unit of time</p>	<p>Capacity-limited elimination: the drug metabolizing enzymes eventually will become saturated as the concentration of substrate increases. The rate of elimination ($t_{1/2}$) varies depending on drug concentration.</p>

What you need to know and understand

Terms to know

Multidose kinetics Continuous or intermittent dosing rate	Drug accumulation: The average concentration of drug will increase until a mean steady state (plateau) is reached (first-order elimination only)
Steady state	Dynamic equilibrium (first-order elimination only): the rate of drug elimination = the rate of drug administration It takes 4 half-lives to achieve 94% of steady state.
Therapeutic window	The range of safe and effective drug doses between the minimum therapeutic and minimum toxic concentrations for the drug
Peak concentration	the maximum concentration the drug achieves for a given dose
Trough concentration	the minimum concentration of the drug immediately before the next dose is given
Minimum effective concentration	the minimum concentration required to produce a therapeutic effect

What you need to know and understand: Terms to know

Don't forget to factor in bioavailability, F.	<p>Loading dose $LD = \frac{C_{desired} \times V_d}{F}$</p> <p>Maintenance dose $MD = [(C_p \times CL) / F] \times \tau$ τ: dosing interval</p>	<p>The dose of drug that promptly raises the concentration of drug in plasma to the target concentration. A loading dose may be desirable when the steady state concentration needs to be achieved rapidly, e.g. life-threatening situation or a drug with a very large V_d or long $t\frac{1}{2}$.</p> <p>➤ Switch to maintenance dose after loading dose to reduce toxicity risk.</p>
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Simple models to describe drug distribution and elimination (first-order elimination).
The time course of drug elimination is plotted on a graph.

One-compartment model of distribution and elimination	This model assumes that the body acts like a single, uniform compartment and that a rapid intravenous injection of a dose of drug equilibrates immediately.
Two compartment model of distribution and elimination	Initial rapid changes in plasma concentration are observed because of a distribution phase – the time required for the drug to reach equilibrium distribution between the central compartment (blood) and a second compartment (tissues and fluid) followed by a slower elimination phase.

What you need to know and understand

Terms to know

Equivalence between drug products:

Pharmaceutical equivalence	Drug products that contain the same active ingredients and are identical in strength or concentration, dosage form, and route of administration
Bioequivalence	Two pharmaceutically equivalent drug products are considered to be <i>bioequivalent</i> when the rates and extents of bioavailability of the active ingredient in the two products are not significantly different under suitable test conditions.
Therapeutic equivalence	Bioequivalent drug products having similar safety and efficacy profiles.
Biosimilar	Biological (protein) product highly similar to an FDA-approved biological product and having no clinically meaningful differences in safety and effectiveness from the reference product.

Pharmacokinetic variables influencing plasma concentration and clearance.

Factors affecting drug half-life	Effect on $t_{1/2}$
<i>Effects on volume of distribution (V_d)</i>	
Aging: \downarrow muscle mass $\rightarrow \downarrow V_d$	Decreased $t_{1/2}$
Obesity: \uparrow adipose mass $\rightarrow \uparrow V_d$ of lipophilic drugs	Increased $t_{1/2}$
Pathologic fluid: $\uparrow V_d$ of hydrophilic drugs	Increased $t_{1/2}$
<i>Effects on clearance</i>	
CYP induction: \uparrow rate of metabolism	Decreased $t_{1/2}$
CYP inhibition: \downarrow rate of metabolism	Increased $t_{1/2}$
Cardiac failure: \downarrow clearance	Increased $t_{1/2}$
Hepatic failure: \downarrow clearance	Increased $t_{1/2}$
Renal failure: \downarrow clearance	Increased $t_{1/2}$

Disease states that can alter the pharmacokinetics of absorption, distribution, metabolism, excretion.

- Renal disease
- Liver disease
- Heart failure
- Shock
- Edema, ascites, pleural effusion

Therapeutic drug monitoring for optimizing the dose of a drug *in an individual patient* ⇒ **maximizing effect while minimizing toxicity**

- Only clinically meaningful when the efficacy of a drug treatment can be enhanced by achieving a certain concentration or effect range and/or the toxicity of drug therapy can be reduced by maintaining a certain concentration or effect range:
 - Measuring the concentration of drug in plasma has utility when there is a(n):
 1. Relationship between the concentration of drug in plasma and the clinical effect
 2. Significant inter- and/or intra-patient pharmacokinetic variability
 3. Established target concentration
 4. Narrow therapeutic window
 5. Availability of a reliable, cost-effective drug assay for clinical use

Optimization of drug therapy is based on in-depth understanding of factors that determine an individual's response to drug treatment.

Fundamental principles should guide prescribing of drugs:

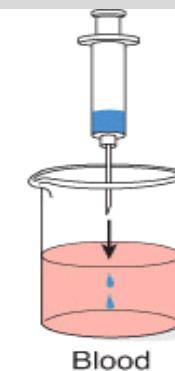
Benefits v. Risk	The benefits of drug therapy, however defined, should always outweigh the risk.
Dose	The smallest dosage necessary to produce the desired effect should be used.
Treatment regimen	The number of medications and doses per day should be minimized.
Changes in drug regimen	Prescribers should be particularly wary when adding or stopping specific drugs that are especially liable to provoke interactions and adverse reactions.
Individual variabilities	Genetics play a role in determining variability in drug response and may become a part of clinical practice
Knowledge	Keeping abreast of the literature and unbiased opinion will provide a foundation for decision-making.
Familiarity	Prescribers should use only a limited number of drugs, with which they are thoroughly familiar.

Parameters Governing Drug Disposition

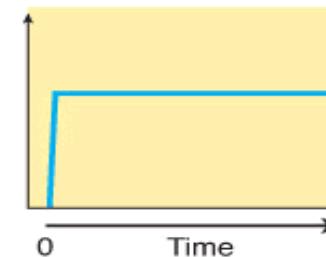
- Clearance
- Bioavailability
- Volume of distribution
- Elimination half-life
- Area under the concentration vs time curve (AUC)
- Steady state dynamic equilibrium with continuous dosing
the rate of elimination = the rate of drug administration

Models of drug distribution and elimination

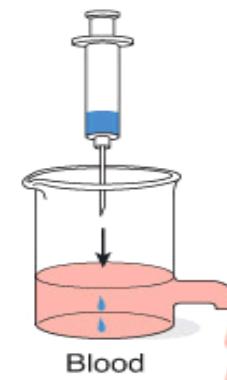
A

Amount
in blood

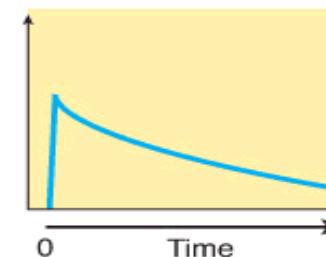
Blood



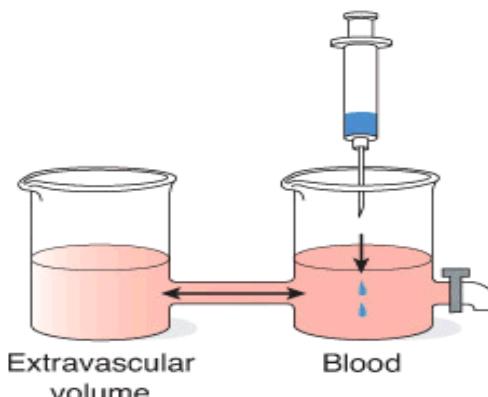
B

Amount
in blood

Blood



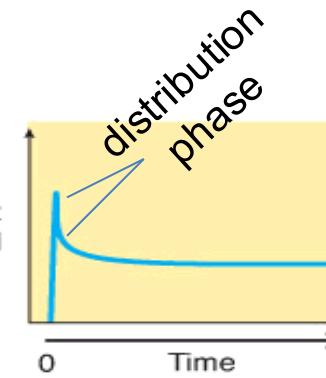
C

Extravascular
volume

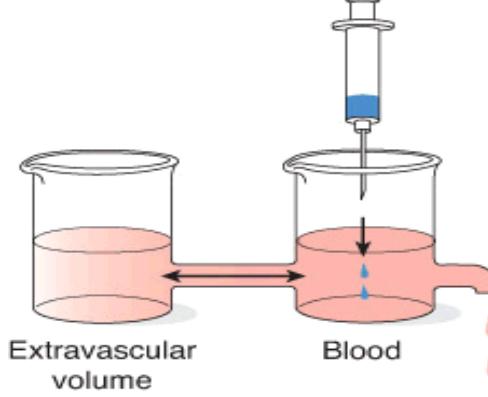
Blood

Amount
in blood

Blood



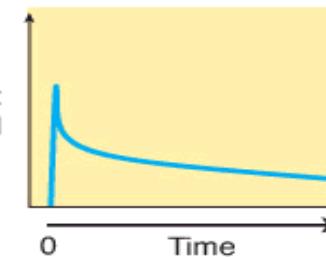
D

Extravascular
volume

Blood

Amount
in blood

Blood



One compartment model:

no movement of drug out of the beaker \Rightarrow steep rise to maximum concentration followed by a plateau

route of elimination is present \Rightarrow a sharp rise to a maximum followed by a slow decay

Two compartment model:

distribution from blood to extracellular fluids and tissues (rapid equilibration)

drug in blood \leftrightarrow drug in extravascular volume

1st compartment

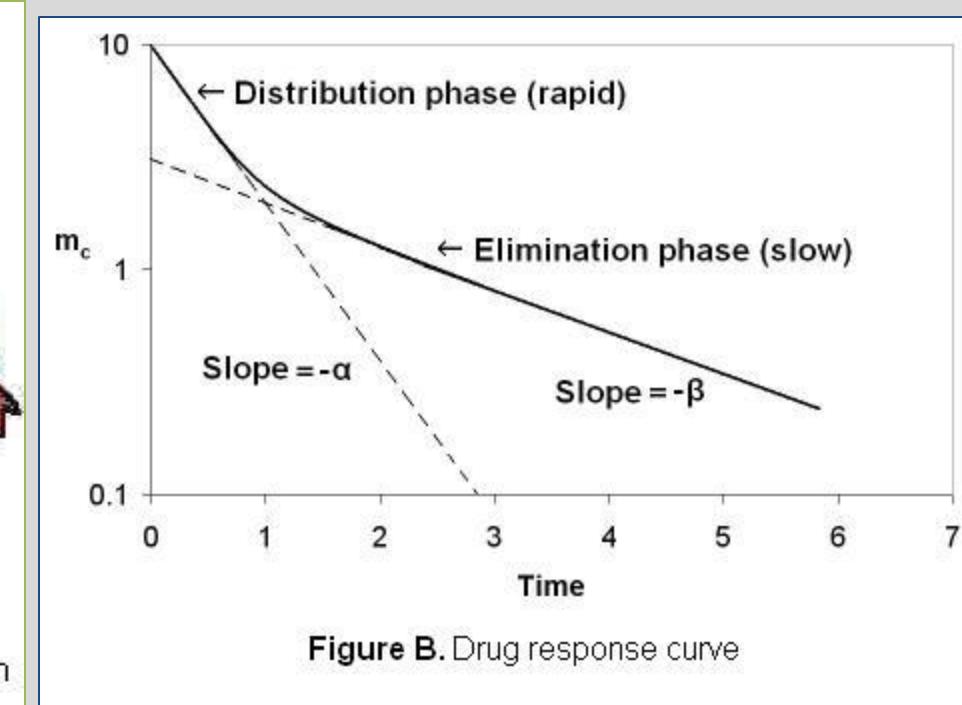
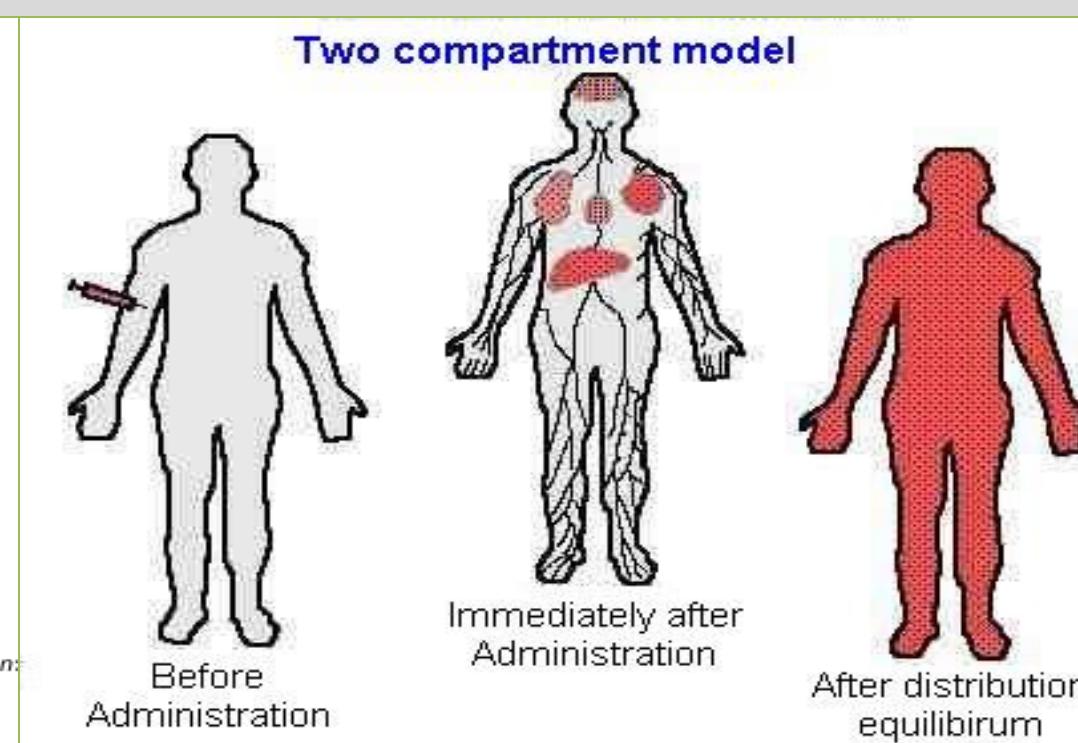
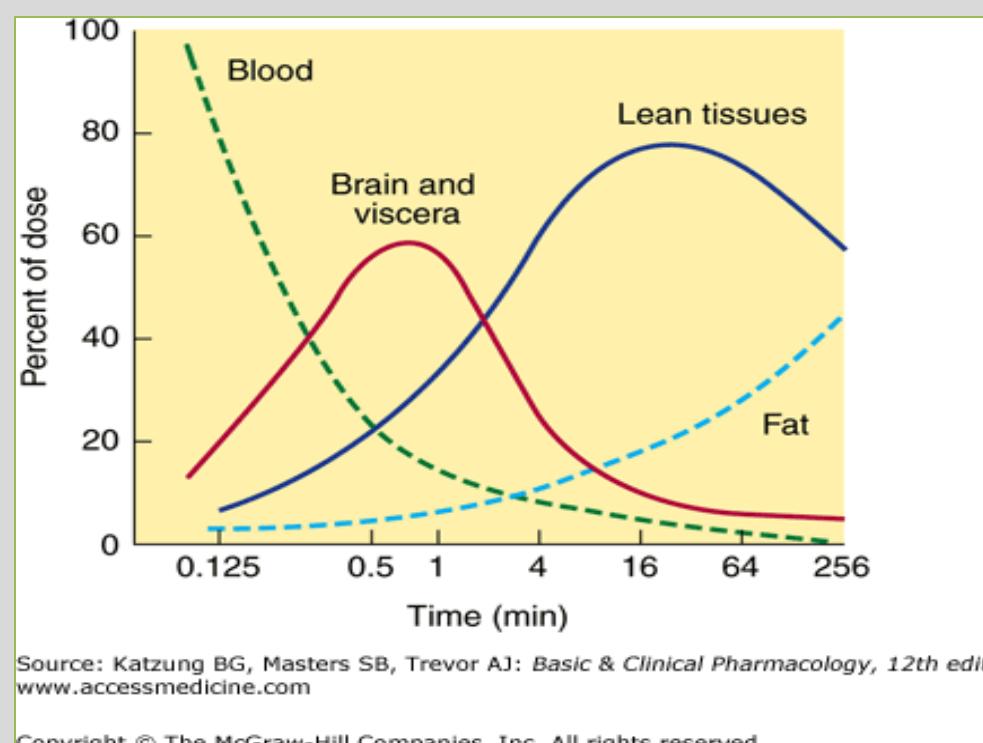
2nd compartment

distribution phase followed by the slower elimination phase

Multicompartment model (first-order kinetics)

- Central compartment: the highly perfused tissues
- Final compartment: the more slowly perfused tissues

central compartment ↔ peripheral compartment



Katzung Figure 25-7

Variation in drug distribution rate

Change in blood flow to various tissues



***Change in rates of
drug distribution to tissues***

Thus, the effect of a drug at various sites of action can vary ***depending on perfusion of these sites.***

Clearance (*CL*)

CLEARANCE

- total systemic clearance:

$$CL = CL_{\text{renal}} + CL_{\text{hepatic}} + CL_{\text{other}}$$

Excretion *Biotransformation to inactive drug* *bile, sweat, saliva, breast milk, lungs (gases)*

Clearance depends on:

- the drug,
- the blood flow, and
- the condition of the organs of elimination in the patient

CLEARANCE

Clearance is the proportionality factor that predicts the rate of elimination in relation to the drug concentration.

- the **rate** of drug elimination is directly proportional to drug **concentration** when clearance is constant

$$CL = \frac{\text{rate of drug elimination (mg/min)}}{\text{drug concentration in measured fluid (mg/mL/kg)}}$$

- Rearranging: Drug elimination rate = $CL \times C_{\text{plasma}}$

At steady state

- Dosing rate_{ss} = Rate of elimination_{ss}
- Dosing rate_{ss} = $CL_{ss} \times C_{ss}$

Extraction ratio_{organ} (E_{organ}),
Bioavailability (F), and
Area under the concentration-time curve (AUC)

Extraction Ratio (E):

a clearance model of an organ's ***intrinsic capacity***

for eliminating a drug in a single pass through the organ at steady state

Extraction ratio compares plasma levels of free drug at steady state immediately before entering and just after exiting the organ:

$$E = \frac{(C_{in} - C_{out})}{C_{in}}$$

$$CL = Q \times \frac{(C_{in} - C_{out})}{C_{in}}$$



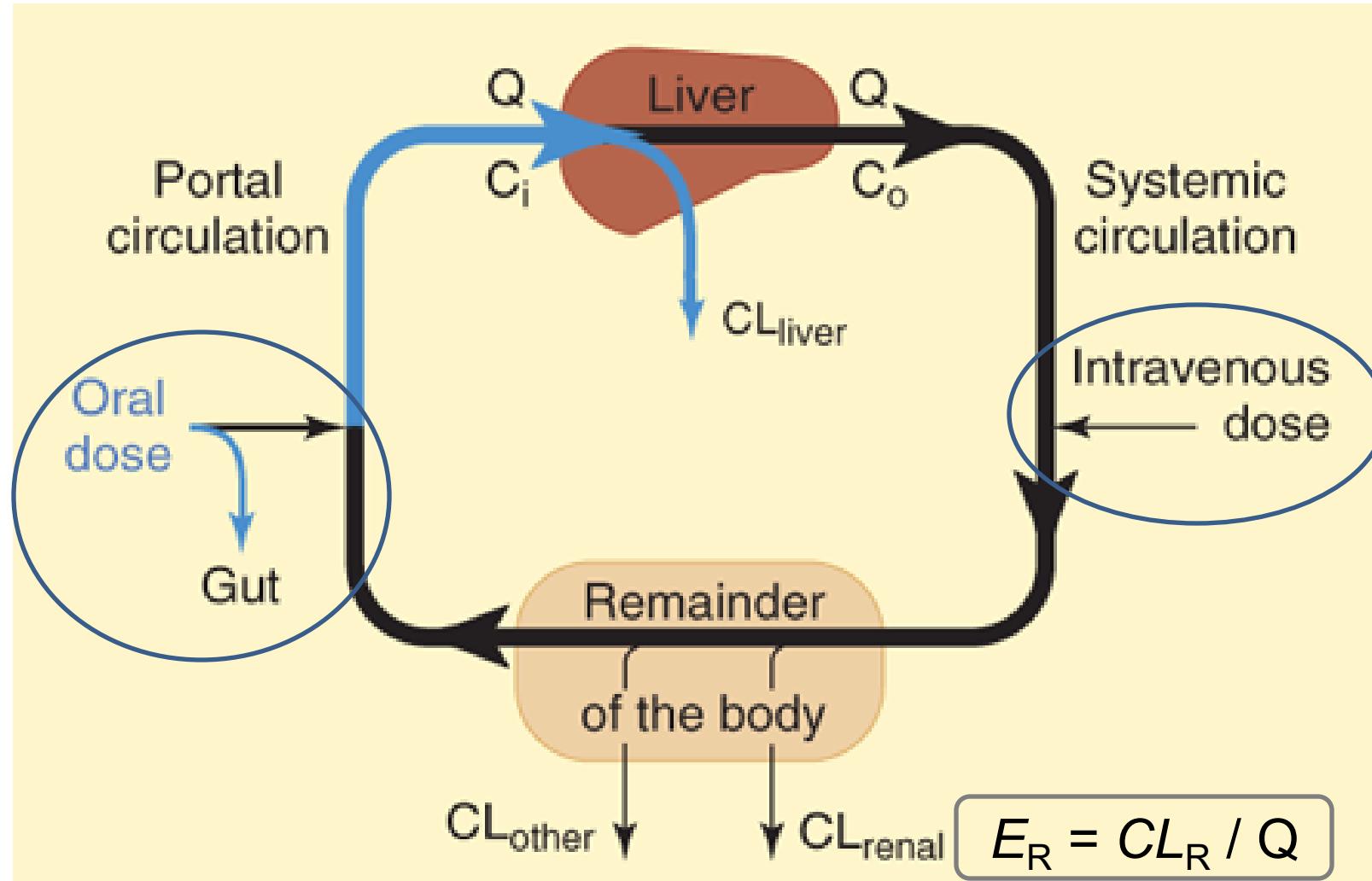
$$CL = Q \times E$$

$$E = CL / Q$$

Drug clearance is determined by:

1. Blood flow (Q) through the eliminating organ
2. Free (unbound) fraction of drug in plasma
3. Intrinsic clearance – the intrinsic ability of clearance mechanisms of the organ (ie, hepatic enzymes to metabolize the particular drug or renal excretory processes)

Illustration of Hepatic and Renal Extraction (E_H and E_R) of a Drug



B. G. Katzung, M. Kruidering-Hall, R. L. Tuan, T. W. Vanderah, A. J. Trevor
Katzung & Trevor's Pharmacology: Examination & Board Review, 13e
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The principles of organ extraction and first-pass effect are illustrated. Part of the administered oral dose (blue) is lost in the gut in the feces or to metabolism, and lost to metabolism in the liver before it enters the systemic circulation: This is the first-pass effect. The extraction of drug from the circulation by the liver is equal to blood flow (Q) times the difference between entering and leaving drug concentration, ie, $Q \times (C_i - C_o)$. CL, clearance. (Modified with permission from Katzung BG: Basic & Clinical Pharmacology, 8th ed. New York, NY: McGraw Hill; 2001.)

$E = (C_{in} - C_{out}) / C_{in}$
C: concentrations of unbound drug
Blood flow determines the rate of presentation of drug to the liver and the rate of exit from the liver.
 $CL = Q \times (C_{in} - C_{out}) / C_{in}$
 $E_H = CL_H / Q$
Bioavailability of oral drugs can be predicted:
 $F = f \times (1 - E)$

f: fraction of dose absorbed

Effect of reduced hepatic blood flow on systemic drug levels

Blood flow-dependent extraction

For drugs with high E_H (>0.7)

- Shunting of blood past the liver will result in substantial increases in drug availability in systemic circulation.**

Hepatic clearance approximates blood flow.

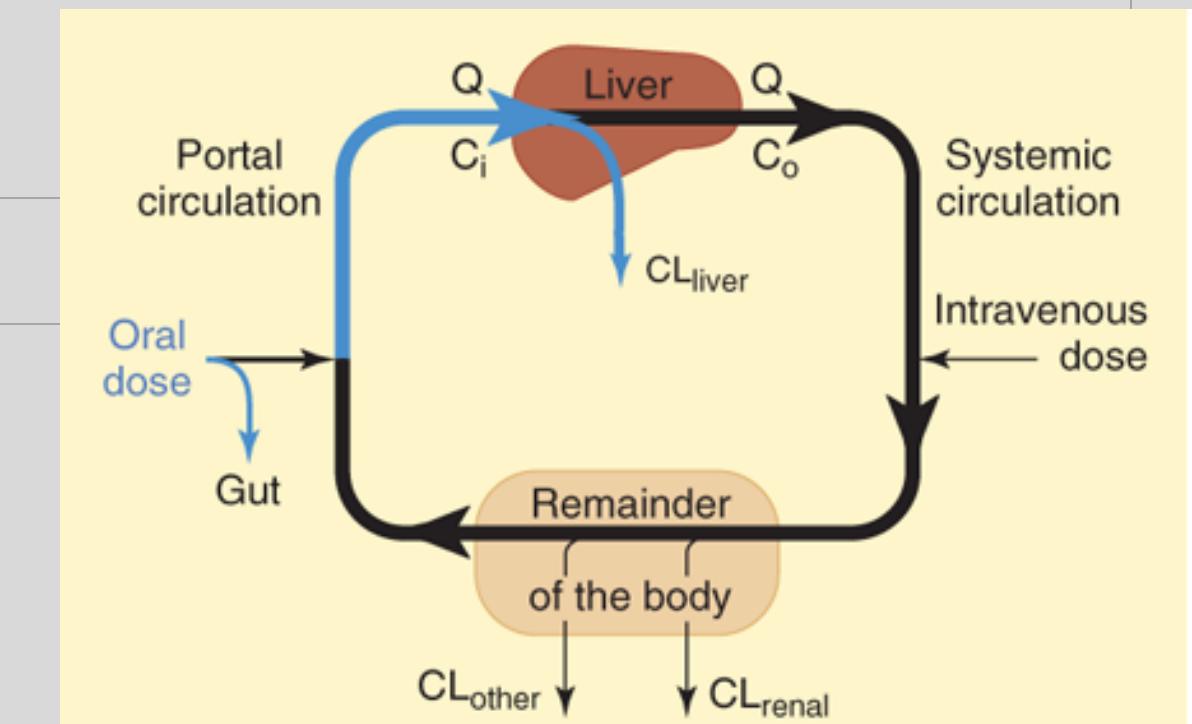
Capacity-limited extraction

For drugs with low E_H (<0.3)

- Shunting of blood past the liver will cause little change in bioavailability.**

Clearance will be proportional to:

the unbound fraction and the drug's intrinsic clearance.



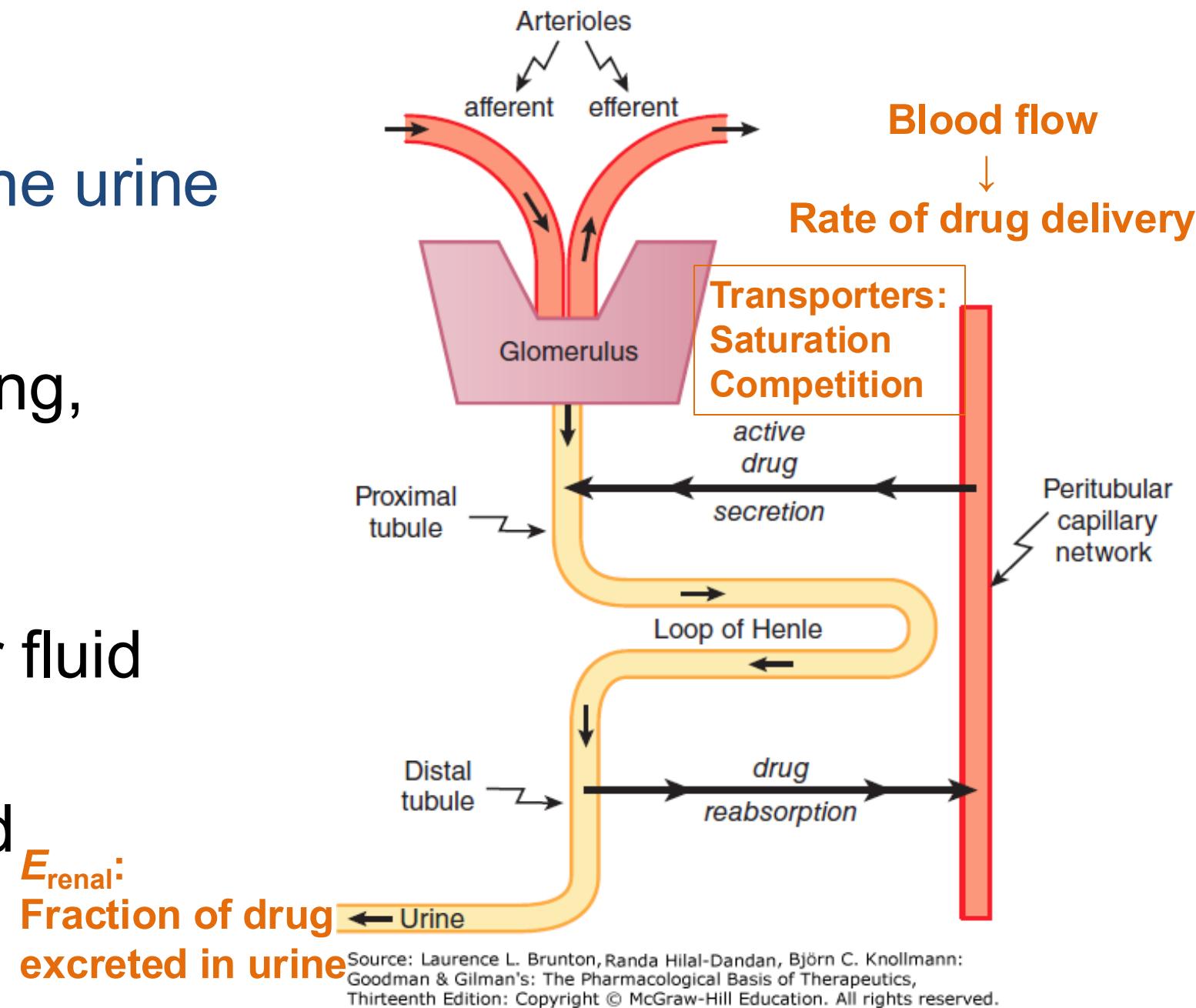
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Renal Extraction:

Fraction of active drug excreted in the urine

Renal clearance is affected by:

1. Renal blood flow, protein binding, function of nephrons
2. Glomerular filtration rate
3. Secretion rate from peritubular fluid into tubular fluid
4. Reabsorption from tubular fluid back into the blood stream



Renal drug handling. Drugs may be filtered from the blood in the renal glomerulus, secreted into the proximal tubule, reabsorbed from the distal tubular fluid back into the systemic circulation, and collected in the urine. Membrane transporters (OAT, OCT, MDR1, and MRP2, among others) mediate secretion into the proximal tubule (see Figures 5–12 and 5–13 for details). Reabsorption of compounds from the distal tubular fluid (generally acidic) is pH sensitive: Ionizable drugs are subject to ion trapping, and altering urinary pH to favor ionization can enhance excretion of charged species (see Figure 2–2).

Bioavailability (*F*)

the fraction of active drug that reaches the systemic circulation

Bioavailability is determined by the:

- 1) **Dose** and
- 2) **Fraction** of the dose

that is absorbed *and* escapes first-pass elimination

$$F = f \times (1 - E)$$

f: fraction of dose absorbed

Drug absorption may be reduced due to:

- incomplete release of drug from dosage formulation
- lesser ability of drug to cross physiologic barriers
- efflux of drug by P-glycoprotein
- metabolism in the intestinal epithelium

Bioavailability is factored into oral dosing calculations.

Example

- Verapamil $E_H \approx 0.67$
- Expected oral bioavailability of verapamil

$$\begin{aligned} F_{\text{verapamil}} &= 1 - 0.67 \\ &= 0.33 \Rightarrow 33\% \end{aligned}$$

Note: Considerable interpatient variability; the actual oral bioavailability of verapamil varies from 20% to 35%

$$\text{dosing rate} = \frac{CL \times C_{ss}}{F}$$

Converting a patient from verapamil I.V. to oral form

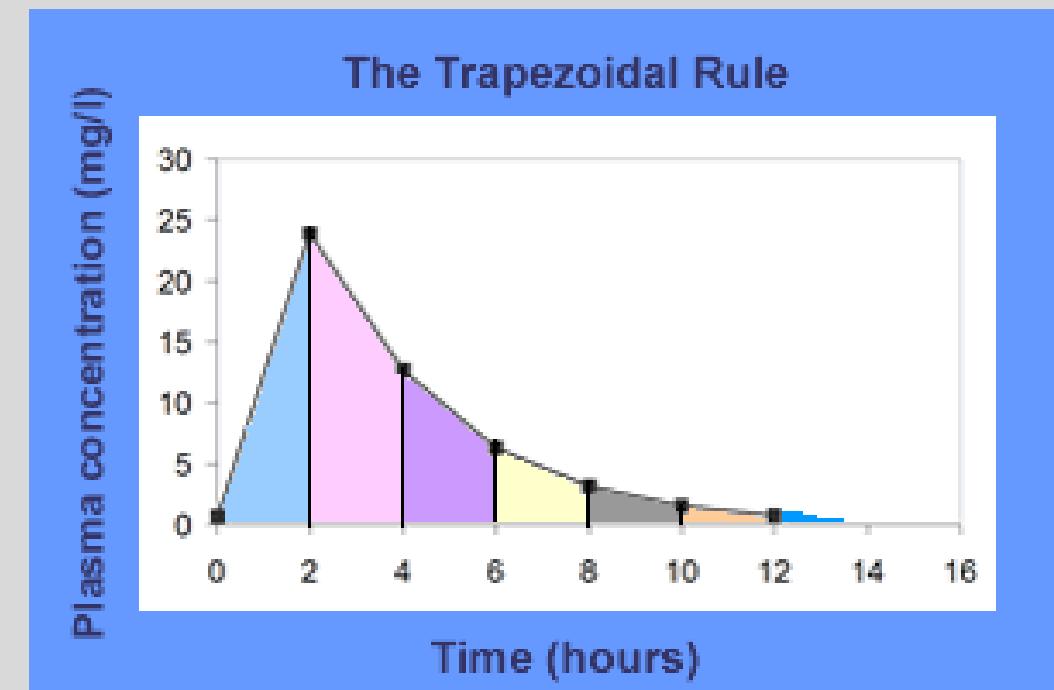
- I.V. dose = 5 mg / hour
= 120 mg / 24 hours
- oral dose = dose / $F_{\text{verapamil}}$
oral dose = 120 mg / 0.33
oral dose = 360 mg / 24 hrs
extended release capsule

AUC

area under the plasma concentration-time curve

AUC assesses the extent of ***the total body exposure*** to a dose of drug

- By integrating the concentration of drug in plasma over time
- For a single dose or during a single dosing interval at steady state.



- Clearance may be estimated from AUC:

$$CL = \text{dose} / C_{ss} \text{ therefore, } CL = \text{dose} / \text{AUC}$$

- The bioavailability of a drug determines its concentration in plasma, which, over time, determines the actual body exposure to the drug.

<http://sepia.unil.ch/pharmacology/index.php?id=66>

Equivalency between Drug Products

Bioavailability can be derived from the area under the curve.

Clinical application:

Comparing bioavailability of different formulations or different manufacturers' products of the same drug

- **Absolute bioavailability**

comparing bioavailability of oral (or other route) to the I.V. formulation of the same drug

$$F = \frac{AUC_{\text{oral}}}{AUC_{\text{IV}}}$$

- **Relative bioavailability**

comparing the bioavailability of:

Different formulations of the same drug(eg, tablet, capsule, solution), or

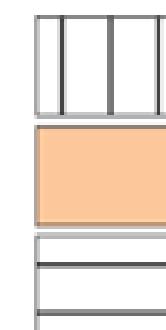
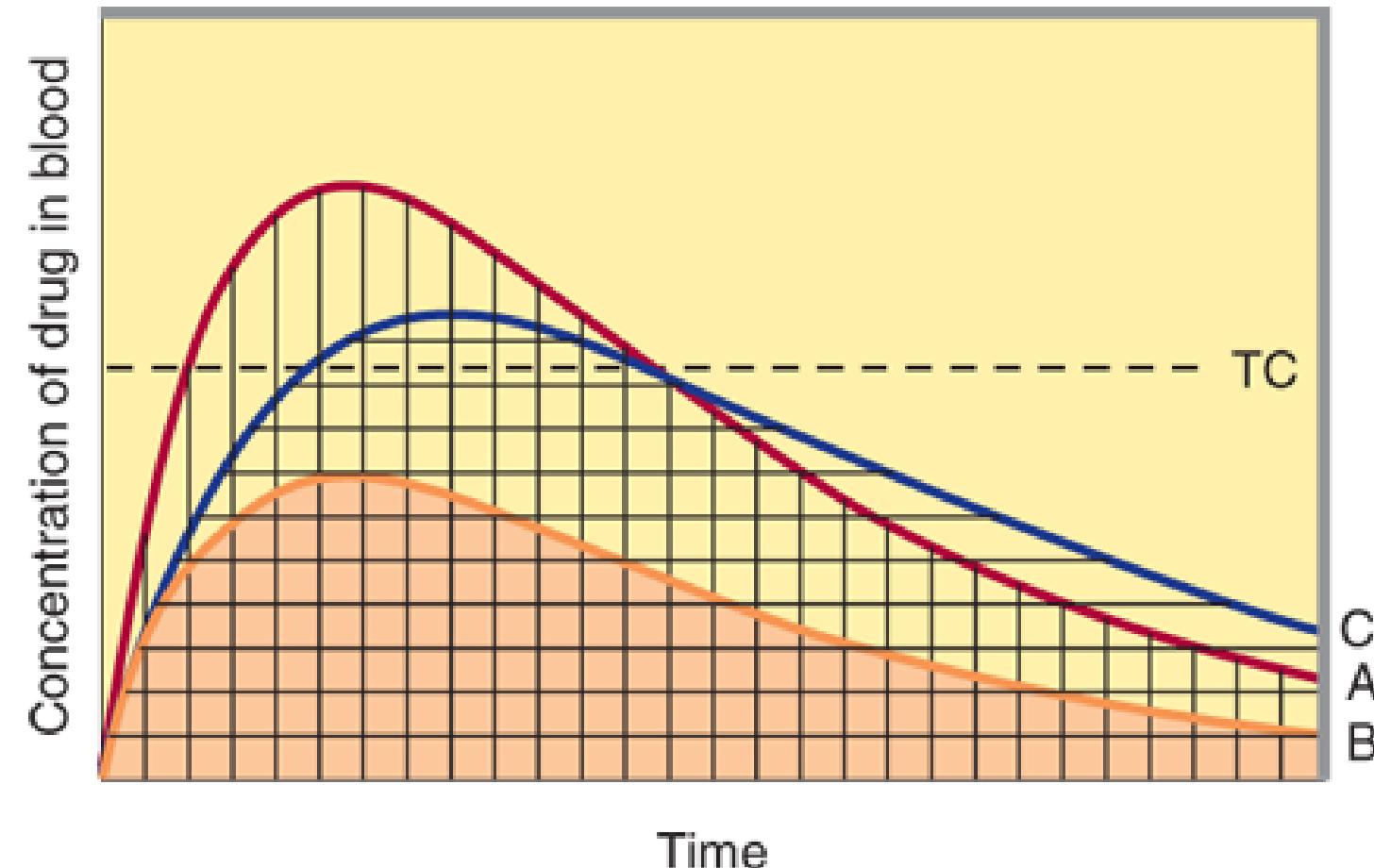
The same formulation of the same drug made by different companies (eg brand and generic tablets)

$$F = \frac{AUC_{\text{version A}}}{AUC_{\text{version B}}}$$

The area under the blood concentration-time curve (AUC) is proportional to the dose and the extent of bioavailability for a drug if its elimination is first-order.

The blood concentration-time curves illustrate how changes in the rate of absorption and extent of bioavailability can influence both the duration of action and the effectiveness of the same total dose of a drug administered in three different formulations.

The dashed line indicates the target concentration (TC) of the drug in the blood.



- A: Drug rapidly and completely available
- B: Only half of availability of A but rate equal to A
- C: Drug completely available but rate only half of A

Source: Bertram G. Katzung, Todd W. Vanderah:
Basic & Clinical Pharmacology, Fifteenth Edition
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Apparent Volume of Distribution (V_d)

Dose

Volume of Distribution:

The proportionality constant that relates the total amount of drug in the body to the plasma concentration at a given time.

$$V_d = \frac{\text{Dose}}{C_{\text{blood or plasma}}}$$

This model considers the body a single homogeneous compartment.

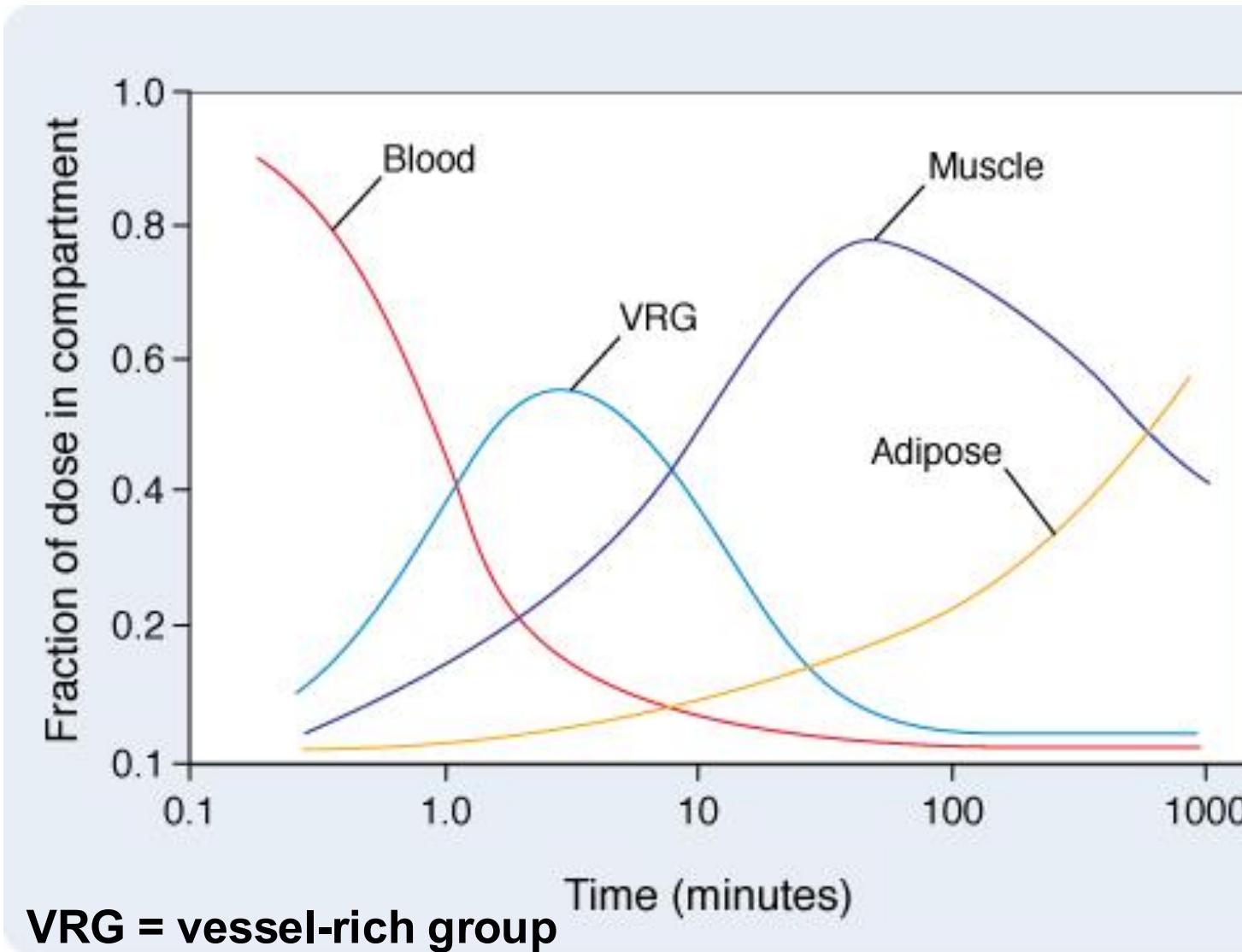
Clearance
 CL

It is an apparent (theoretical) volume that represents a drug's propensity to remain in plasma or distribute to other tissue compartments.
It is a calculated parameter.

From: 3 Pharmacokinetics

Principles of Pharmacology: The Pathophysiologic Basis of Drug Therapy Fourth Edition, 4e, 2017

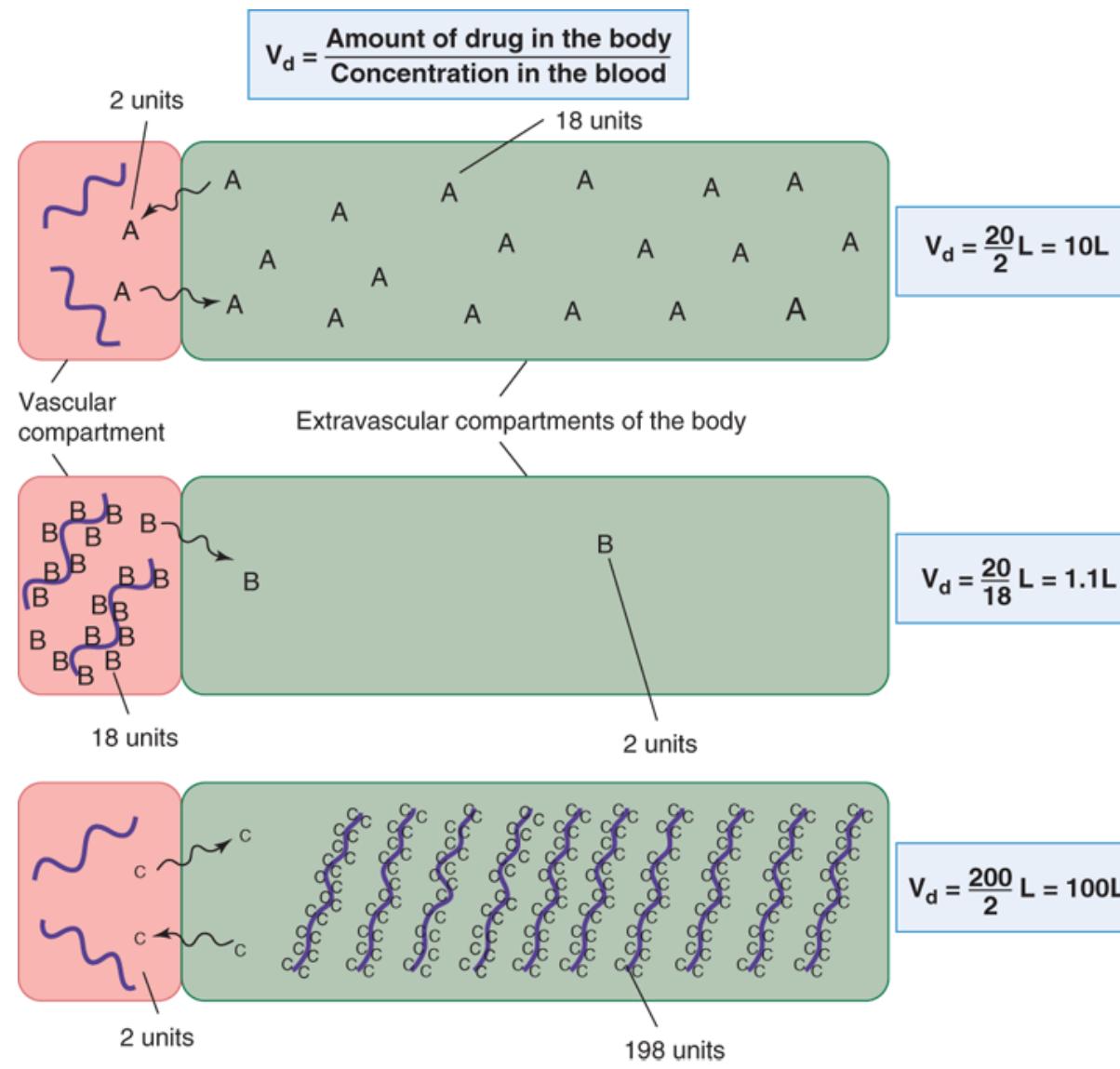
Four-compartment model of drug distribution.



Multi-compartment model:
Drug concentration in a sample varies over time as the drug distributes around the body.

The drug concentration in a sample immediately after IV bolus will produce a different V_d value than a sample taken several hours later.

- 1. Initial distribution phase**
 - Highly-perfused organs – brain, kidney, liver, heart – receive most of the drug
- 2. Second distribution phase**
 - Delivery to muscle, most viscera, skin, and fat is slower
 - May require minutes to several hours before the concentration of drug in tissue is in equilibrium with that in blood
 - Involves far larger fraction of body mass relative to initial phase



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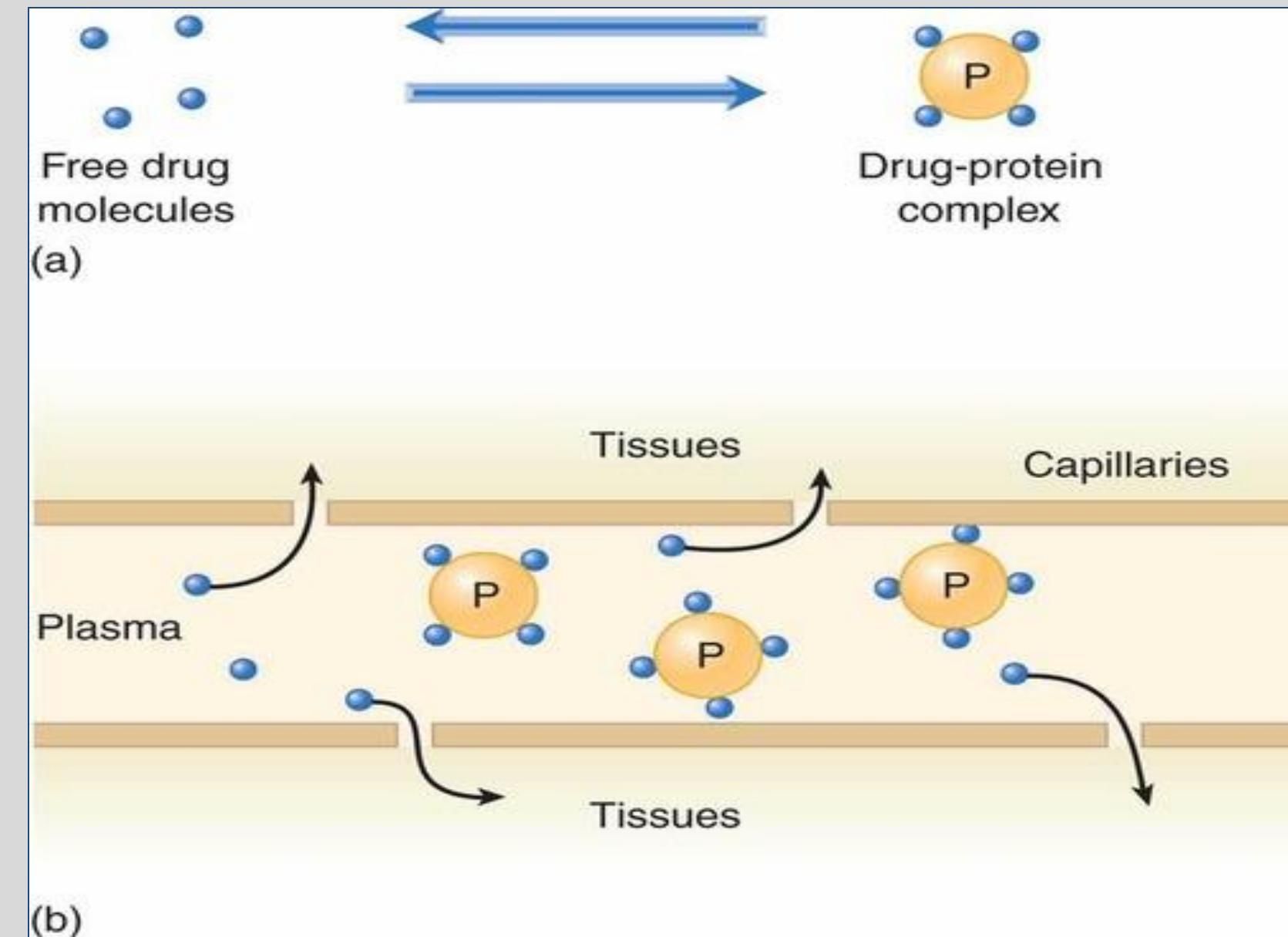
- Drug A diffuses freely between the 2 compartments.
- Drug B binds avidly to plasma proteins (wavy lines).
 The drug is retained in plasma compartment → low V_d
- Drug C binds avidly to molecules in peripheral tissues.
 - Low concentration in blood
 - High V_d
 - Larger total dose is required to achieve measurable plasma concentrations

Effect of drug binding on volume of distribution. Drug A diffuses freely between the 2 compartments and does not bind to macromolecules (heavy wavy lines) in the vascular (volume 1 L) or the extravascular compartments (volume 5 L) of the hypothetical organism in the diagram. With 20 units of the drug in the body, the steady-state distribution leaves a blood concentration of 2 units/L. Drug B, on the other hand, binds avidly to proteins in the blood. At equilibrium, only 2 units of the total are present in the extravascular volume, leaving 18 units still in the blood. In each case, the total amount of drug in the body is the same (20 units), but the apparent volumes of distribution are very different. Drug C is avidly bound to molecules in peripheral tissues, so that a larger total dose (200 units) is required to achieve measurable plasma concentrations. At equilibrium, 198 units are found in the peripheral tissues and only 2 units in the plasma, so that the calculated volume of distribution is greater than the physical volume of the system.

When plasma protein binding is reversible, a chemical equilibrium exists between bound and unbound drug.

Processes that lower the free drug concentration lead to rapid dissociation of drug from the drug-protein complex, thereby

reestablishing equilibrium between bound and free drug.



Anticipated effect of drug displacement from plasma proteins:

Displacement
of drug from
plasma proteins



Transient
increase in
free fraction



**Rapid
equilibration**

The increase in free fraction will increase the apparent clearance of the drug, thus the mean unbound plasma concentration at steady state would not change.

The **rate** of drug metabolism is directly proportional to the free drug concentration and the **fraction** of drug removed is constant.

🔑 Changes in protein binding due to disease states and drug-drug interactions are clinically relevant mainly for a small subset of drugs with a **narrow therapeutic window and potential for significant toxicity**.

Summary: Clinical Pharmacokinetics Part 1

- The plasma concentration is a function of the rate of input of the drug (by absorption) into the plasma, the rate of distribution into other tissues, and the rate of elimination.
- If the rate of input is known, the remaining processes are well described by two primary parameters: **apparent volume of distribution (V_d)** and **clearance (CL)**. These parameters are unique for a particular drug and a particular patient but have average values in large populations that can be used to predict drug concentrations.
- Clearance of a drug is the factor that predicts the rate of elimination in relation to the drug concentration.
- Volume of distribution is the proportionality constant that relates the total amount of drug in the body to the plasma concentration at a given time.

- Change in blood flow to various tissues change in rates of drug distribution to tissues. Thus, the effect of a drug at various sites of action can vary depending on perfusion of these sites.
- Removal of a drug by an organ can be specified as the extraction ratio – the fraction or percentage of the drug removed from the perfusing blood during one passage through the organ.
- The bioavailability of a drug is the fraction (F) of the administered dose that reaches the systemic circulation.
- The area under the concentration vs. time curve (AUC) describes the measured concentration of drug in the systemic circulation as a function of time (from zero to infinity) – the concentration of drug appearing in plasma occurs over time. It can be used to calculate bioavailability.

Matching

Drug products that contain the same active ingredients and are identical in strength or concentration, dosage form, and route of administration

Bioequivalent drug products having similar safety and efficacy profiles.

Biosimilar product and meets additional standards for interchangeability

Biological product highly similar to an FDA-approved biological product and having no clinically meaningful differences in safety and effectiveness from the reference product

Pharmaceutically equivalent drug products show no significant difference in the rate and extent of absorption of the active pharmaceutical ingredient

Bioequivalence

Biosimilar

Interchangeable
biologic product

Pharmaceutical
equivalence

Therapeutic
equivalence

References

- Access Medicine Goodman & Gilman's The Pharmacological Basis of Therapeutics 14e; 2023; Chapter 2: Pharmacokinetics: The Dynamics of Drug Absorption, Distribution, Metabolism, and Elimination
- Access Medicine Harrison's Principles of Internal Medicine, 22e, 2025; Chapter 71: Principles of Clinical Pharmacology
- Access Medicine Katzung's Basic & Clinical Pharmacology 16e, 2024: Chapter 3: Pharmacokinetics & Pharmacodynamics: Rational Dosing & the Time Course of Drug Action, and Chapter 66: Rational Prescribing & Prescription Writing
- Access Medicine Principles and Practice of Hospital Medicine, 2e, 2017; Chapter 9: Principles of Evidence-Based Prescribing

Lecture Feedback Form:

<https://comresearchdata.nyit.edu/redcap/surveys/?s=HRCY448FWYXREL4R>

Clinical Pharmacokinetics Part 2

Rational Dosing and the Time Course of Drug Action

Leslie Goldstein, Pharm.D.

Associate Professor

Department of Clinical Specialties

lgolds01@nyit.edu

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After completing the preparation materials, students should be able to:

1. Describe clearance, volume of distribution, elimination half-life, steady state, bioavailability, extraction ratio, and area under the curve using pharmacokinetic models.
2. Differentiate first-order and zero-order elimination kinetics and linear and non-linear processes in relation to drug concentration and effect.
3. Graph drug accumulation, half-life, and steady state for drugs with first-order elimination administered by continuous and intermittent dosing.
4. Calculate the maintenance and loading doses for an individual patient to achieve a target concentration within the therapeutic window in when given specific pharmacokinetic parameters.
5. Identify pharmacokinetic variables that influence drug plasma concentrations and the criteria for therapeutic drug monitoring.

Preparation Materials (links are in the CPG and on the next slide)

Required

- ScholarRx Bricks | Practice Questions

Highly relevant optional materials:

- Dr. Goldstein's Word handout | Video Lectures | Guided Reading Questions
- Textbook resources with links are listed on the next slide

SUGGESTIONS:

- *Use the resources that work best for you.*
- *You do not need to study all of them.*
- *Work through the GUIDED READING QUESTIONS with pen/pencil and paper.
Try answering them in your OWN WORDS first without looking at the readings, and then again after reviewing.*
- *Practice questions (not graded): Simple Recall and Case Vignettes*

Links

Scholar Rx Bricks: (Required)

General Principles

Pharmacokinetics and Pharmacodynamics > Pharmacokinetics: Drug Concentration and Dosing

<https://exchange.scholarrx.com/brick/drug-concentration-and-dosing>

Enzymes as Drug Targets <https://exchange.scholarrx.com/brick/enzymes-as-drug-targets>

Katzung & Vanderah's Basic & Clinical Pharmacology, 16e, 2024; Chapter 3: Pharmacokinetics

<https://accessmedicine-mhmedical-com.nyit.idm.oclc.org/content.aspx?bookid=3382§ionid=281747069>

THE REVIEW BOOKS HAVE PRACTICE QUESTIONS

Katzung & Trevor's Pharmacology: Examination and Board Review, 14e, 2024

Chapter 3: Pharmacokinetics

<https://accessmedicine-mhmedical-com.nyit.idm.oclc.org/content.aspx?bookid=3461§ionid=285589076>

LWW Health Library Premium Basic Sciences; Lippincott Illustrated Reviews: Pharmacology, 8e, 2023: Chapter 1: Pharmacokinetics

<https://premiumbasicsciences-lwwhealthlibrary-com.nyit.idm.oclc.org/content.aspx?sectionid=253324942&bookid=3222>

For the list of “What you need to know and understand”,
please see the PowerPoint presentation for
Clinical Pharmacokinetics, Part 1

Half-life

Half-life

The time required to reduce the amount of drug in the body by one-half during elimination – that is, with each half-life the drug concentration decreases by 50% (first-order kinetics).

 $t_{1/2}$ changes as a function of both clearance and volume of distribution

$$t_{1/2} = \frac{0.693 \times V_d}{CL}$$

- **$t_{1/2}$ is an indicator of:**
 1. Time required to reach steady state
 2. Time required for drug to be removed from the body
 3. A way to estimate dosing interval

Half-life depends on both the rate of elimination and the volume of distribution.

- $t_{1/2}$ is prolonged in proportion to either an increase in V_d or decrease in CL .

Half-life is dependent on exponential elimination kinetics. 0.693 is the logarithm of 2 ($\ln 2$).

First-order elimination

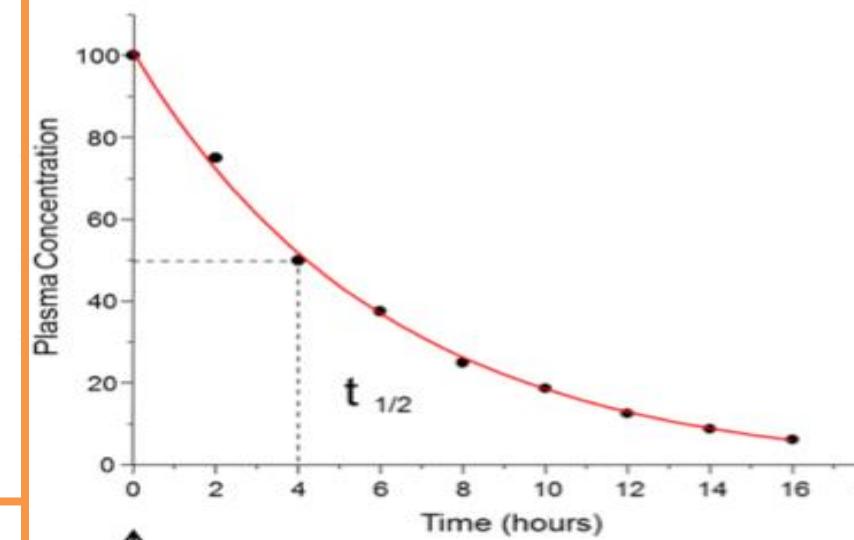
Zero-order elimination (capacity-limited elimination)

First-Order Elimination Kinetics

A constant proportion (fraction) of drug is eliminated per unit of time (linear elimination).

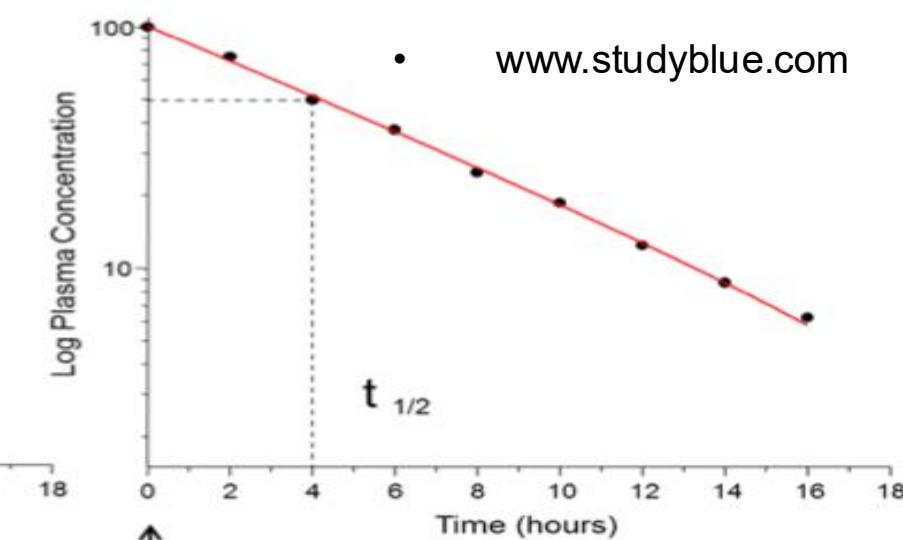
The **rate** of drug metabolism is directly proportional to the free drug concentration and the **fraction** of drug removed is constant.

The vast majority of drugs follow first-order elimination kinetics at therapeutic doses.



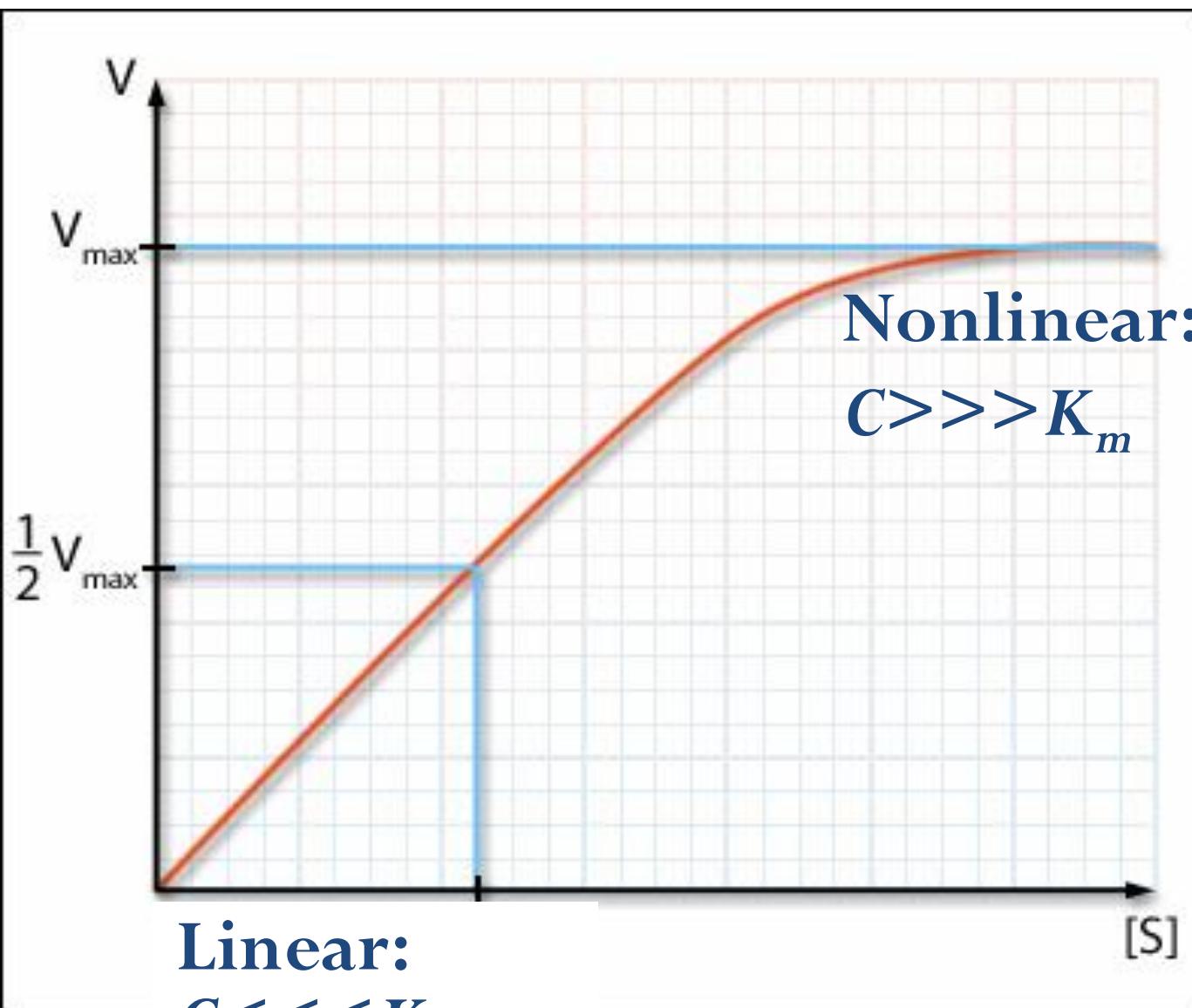
↑ Serum level curve observed from a drug eliminated by a first-order process.

$$t_{1/2} = 4 \text{ hours in this illustration}$$



↑ A plot of the same data using a log scale on the y-axis results in a straight line.

Zero-order kinetics: Clearance mechanisms become saturated – the rate of metabolism remains constant – a constant amount (not fraction) of drug is metabolized per unit of time. Dangerously elevated drug concentration can result.



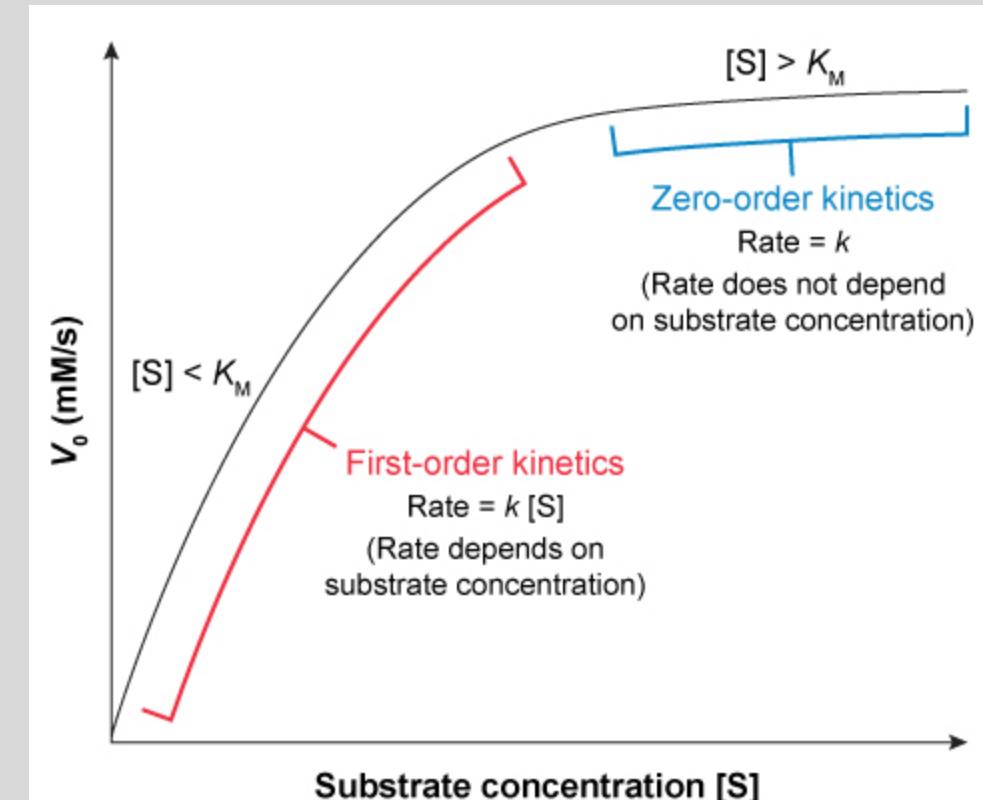
$$\text{Rate of Elimination} = \frac{V_{max} \times C}{K_m + C}$$

$$C_{ss} = \frac{\text{Dosing rate} \times K_m}{V_{max} - \text{Dosing rate}}$$

As dosing rate approaches V_{max} , $V_{max} - \text{dosing rate}$ approaches zero → disproportionate increase in concentration.
At $C \gg K_m$ elimination rate is almost independent of concentration.

K_m is the concentration of substrate at which enzyme activity is at half maximal – $\frac{1}{2} V_{max}$.

Enzyme Kinetics Curve comparing zero-order and first-order kinetics.



Linear Plot of Cp Versus Time Showing High Cp and Low Cp Zero Order and First Order Elimination

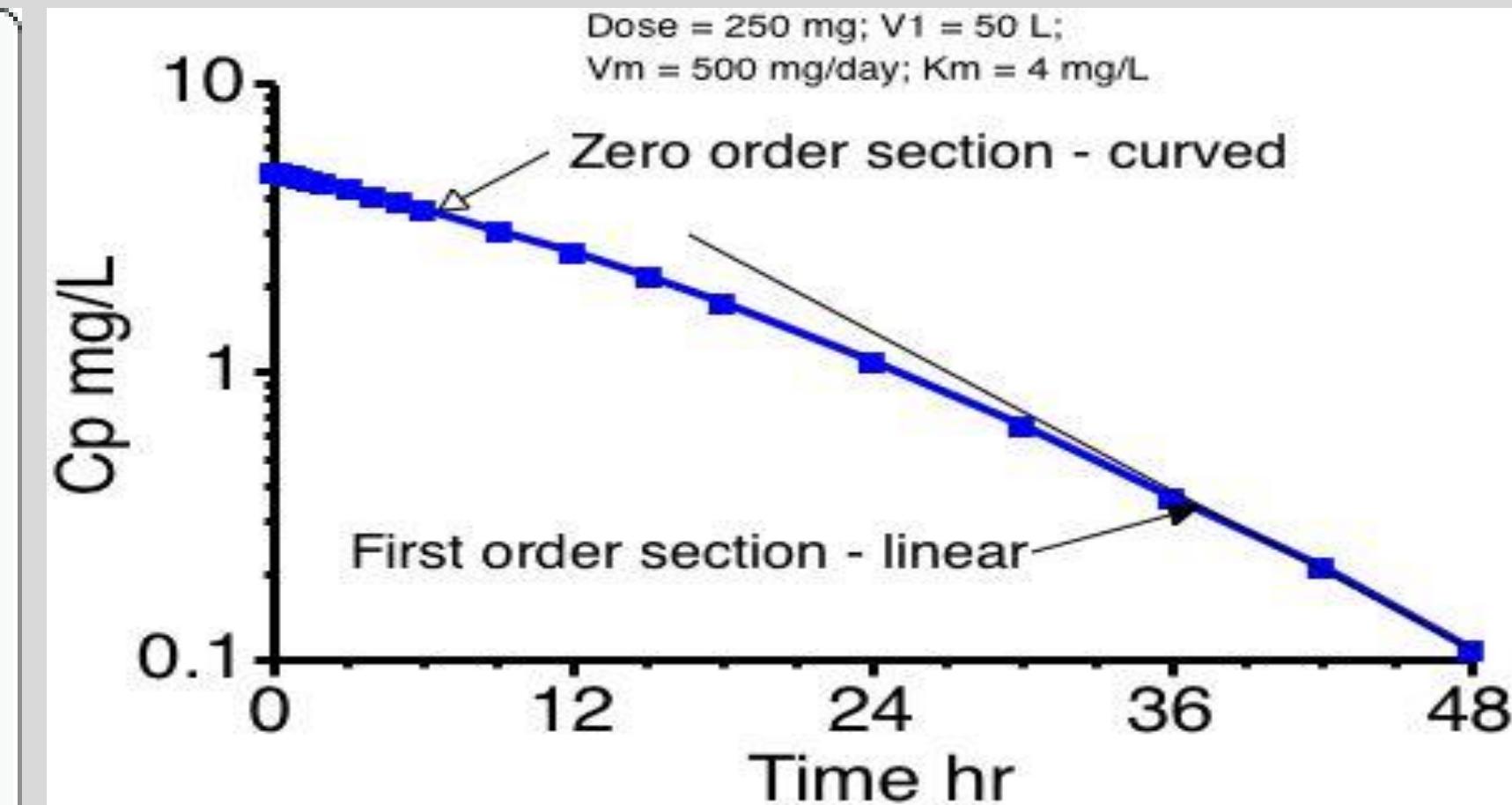
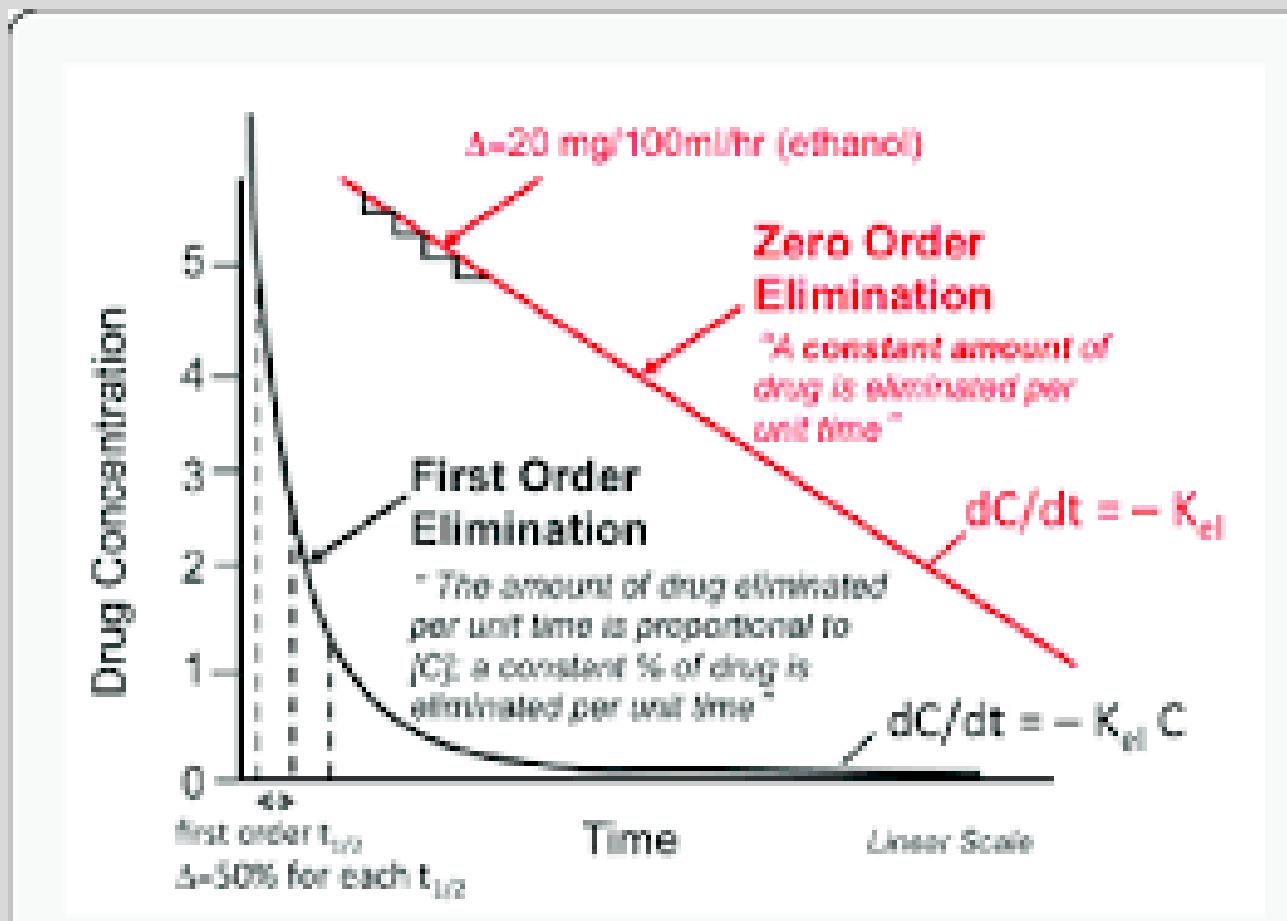


Figure 21.3.2 <http://www.boomer.org/c/p4/c21/c2103.html>

If dosing rate exceeds elimination capacity, steady state cannot be achieved ...the concentration will keep rising as long as dosing continues.

Changing the dosing rate for a drug with nonlinear kinetics is difficult and unpredictable.

Three drugs exhibit zero-order kinetics at therapeutic concentrations:
Ethanol, aspirin (anti-inflammatory doses), and phenytoin.

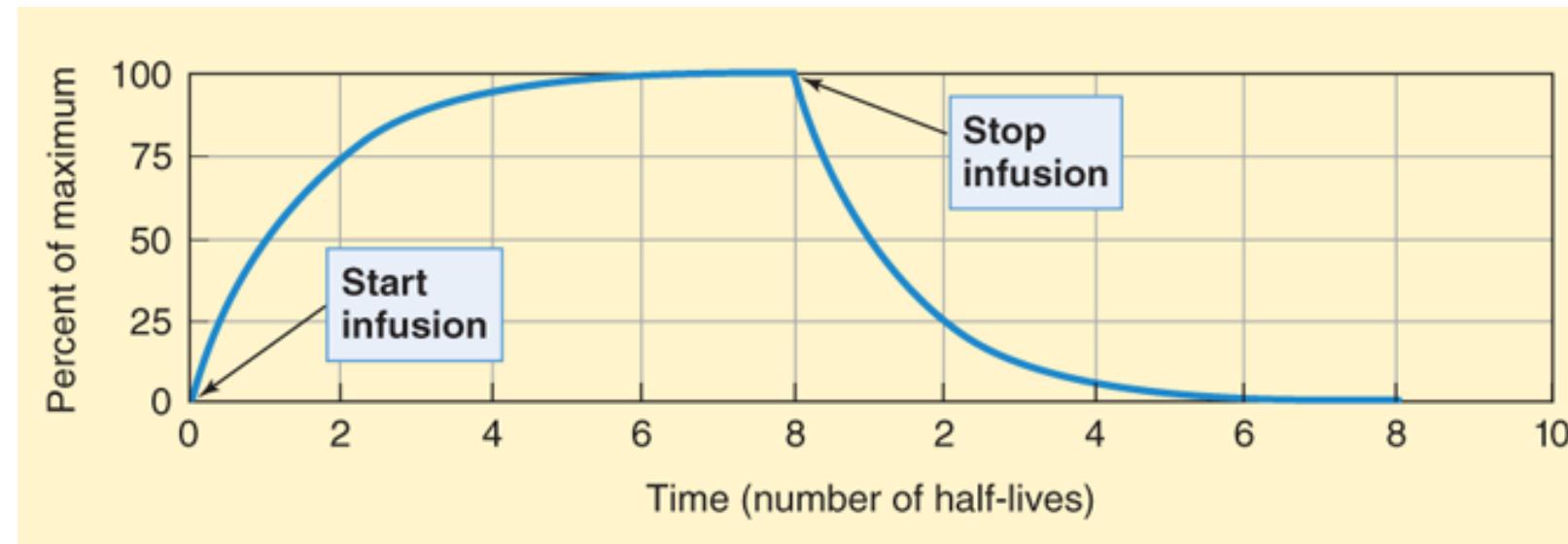
Most drug elimination pathways will become saturated if the dose is high enough→...→ toxicity.

Multidose kinetics: Drug Accumulation and Steady State for drugs that exhibit first-order elimination

Continuous administration results in drug accumulation until the rate of drug going in = rate going out.

After that point, no additional accumulation occurs.

Dynamic equilibrium – steady state – is reached.



B. G. Katzung, M. Kruidering-Hall, R. L. Tuan, T. W. Vanderah, A. J. Trevor
Katzung & Trevor's Pharmacology: Examination & Board Review, 13e
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4 half-lives is the time required ...

- to approach steady state and full effects to be seen, and
- for drug to be removed from the body by elimination

1 $t_{1/2}$ → 50%
2 $t_{1/2}$ → 75%
3 $t_{1/2}$ → 87.5%
4 $t_{1/2}$ → ~94%

Plasma concentration (plotted as percentage of maximum) of a drug given by constant intravenous infusion for 8 half-lives and then stopped. The concentration rises smoothly with time and always reaches 50% of steady state after 1 half-life, 75% after 2 half-lives, 87.5% after 3 half-lives, and so on. The decline in concentration after stopping drug administration follows the same type of curve: 50% is left after 1 half-life, 25% after 2 half-lives, and so on. The asymptotic approach to steady state on both increasing and decreasing limbs of the curve is characteristic of drugs that have first-order kinetics.

Drug accumulation with a continuous or intermittent dosing rate is based on the fundamental principle of clearance:

- Clearance = the volume of blood from which a drug is removed per unit of time (in mL/min or L/h)

- Clearance relates the rate of elimination to the plasma concentration (in $\mu\text{g/mL}$ or mg/L)

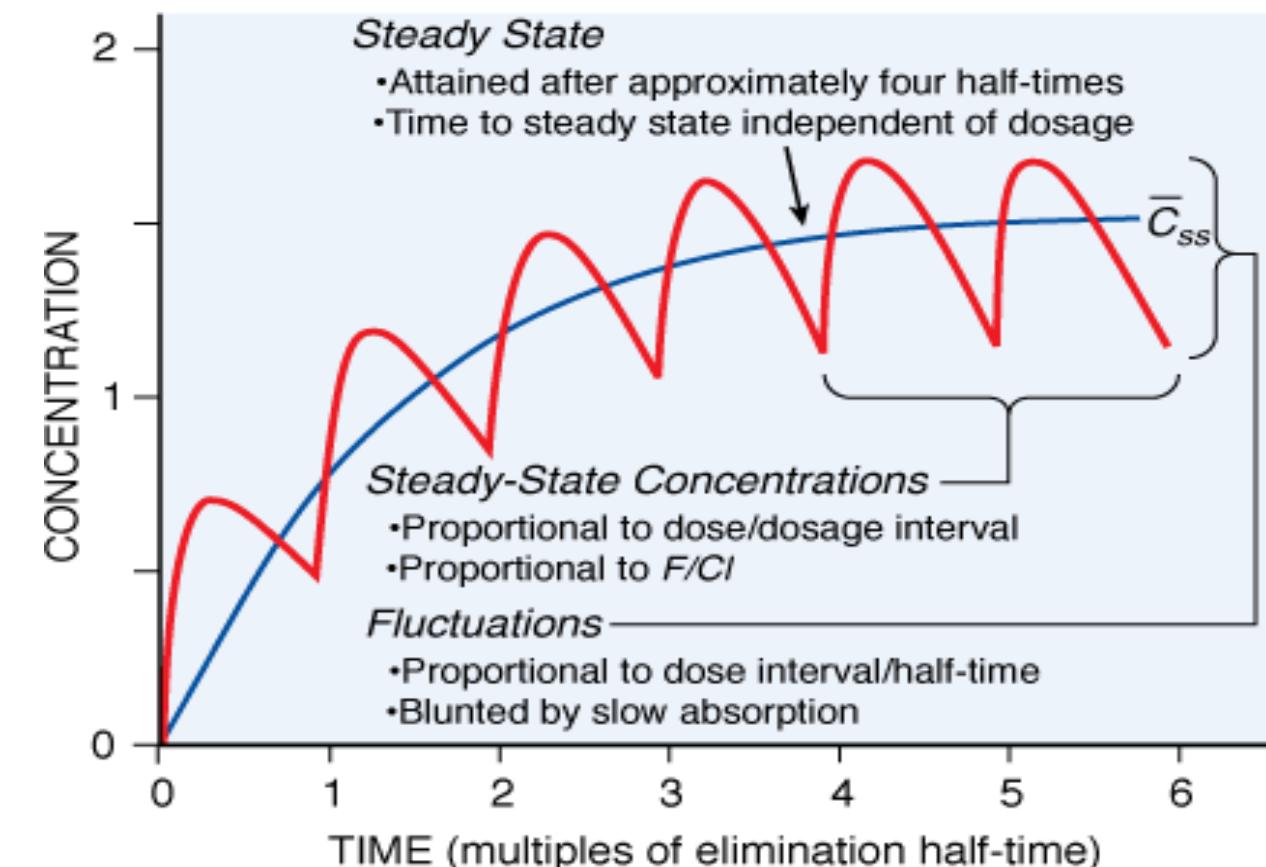
$$\text{dosing rate} \times F = CL \times C_p$$

- Clearance of a particular drug remains constant although the actual amount of drug in the clearance volume varies with the plasma drug concentration.

Steady state:

Rate of drug elimination = Rate of drug administration

when drug is administered at a constant rate.



Source: Brunton LL, Chabner BA, Knollmann BC: *Goodman & Gilman's The Pharmacological Basis of Therapeutics*, 12th Edition:
www.accessmedicine.com
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The legend for this figure is in the notes section of the PowerPoint slide.