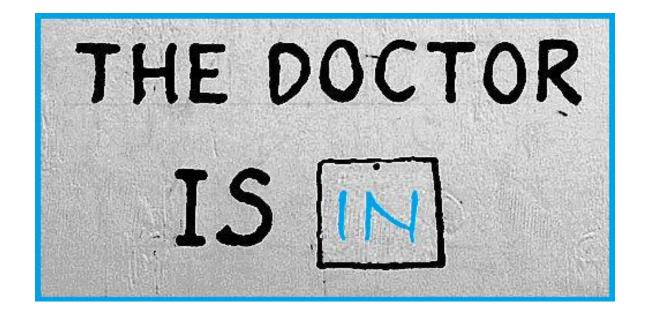
Next Step Office Hours: Biochemistry Review

- Welcome to Office Hours!
- Introduction
- What Do I Need for this Session?
- Biochem Content Review
- What Next?





Office Hours: Biochem Content Review

Introduction to Office Hours

- Thanks for coming to Next Step Office Hours!
- If you haven't been here before, here's how it works...
- These sessions are meant to be:

Interactive

Problem-focused

Specific to your needs (so ask questions!)

- Today's focus: Review of Biochem
- Future sessions: content review, FL review
- This is NOT a lecture! You can benefit most by:

Raising your hand and speaking Commenting in the chat box

Responding to poll questions

Before Getting Started

- 1. If you have a microphone, make sure it is turned on and easily available.
- 2. Locate the hand-raise button on the toolbar on your screen.
- 3. Locate the chat box on the toolbar.
- 4. Let me know if you're having any technical issues!

Think of your question after Office Hours are over?

Post on the forums!
 forum.nextstepmcat.com



Biochem Content Review

• Overall study strategies

Active learning

Big-picture perspective

Test-like thinking

• High-yield topics

Enzymes & enzyme kinetics

Amino acids

Glycolysis

Citric acid cycle

Electron transport chain



Biochem Study Strategies

Recurring theme for biochem:

Don't miss the forest for the trees!

When studying, ask yourself ...

- Why does this matter physiologically?
 - Biomolecules: how does chemical structure connect to biological function?
 - Pathways: what does a pathway DO?
- What are the inputs & outputs of a pathway?
- How is a pathway regulated (big-picture?)
- Does a pathway have any especially important steps?





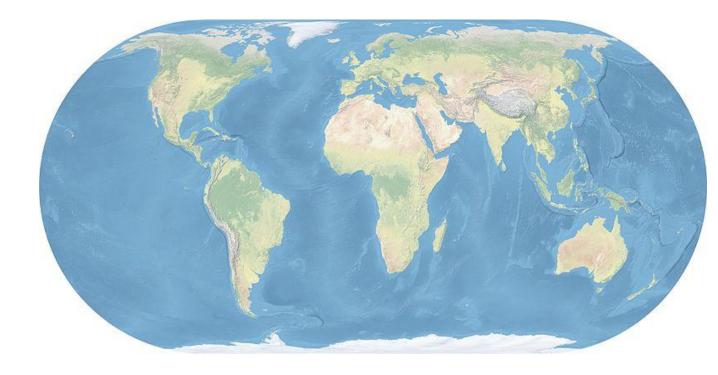
Biochem Study Strategies

A big-picture approach to biochem:

How is biochem tested on the MCAT? How do you get the most bang for your buck in terms of studying?

Focus on:

- Principles
- Physiological function
- Interconnections with other subject matter
 Amino acids & acid-base chemistry
 Carbohydrates & stereochemistry
 Metabolism & physiology



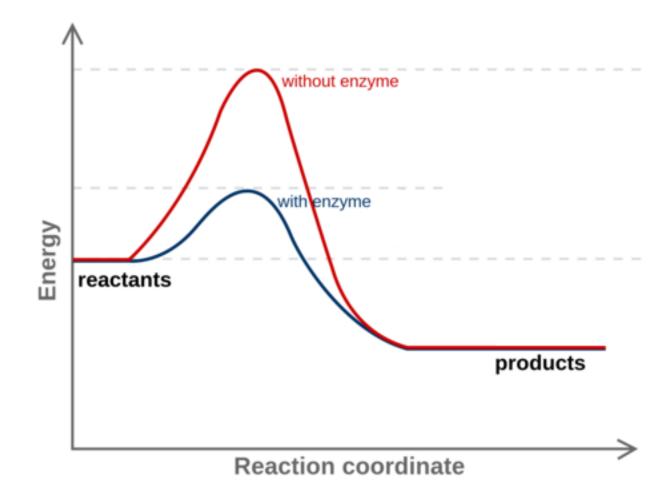
What have your biochem experiences been like? What strategies work for you?



What do enzymes do?

- Enzymes are biological catalysts.
- Enzymes reduce activation energy of rxn.
- Reduced activation energy \rightarrow faster rate
- What do enzymes NOT do?
- Major types of enzymes:

Oxidoreductases	Lyases
Transferases	Isomerases
Hydrolases	Ligases

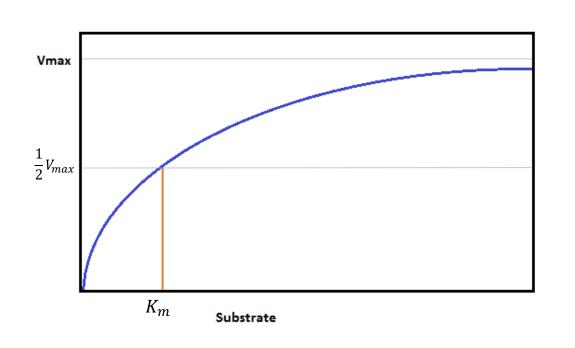




Office Hours: Biochem Content Review

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Michaelis-Menten saturation curve



V_{max}: the maximum rate of the reaction

K_m: the amount of substrate needed for the enzyme to work half as fast as it can

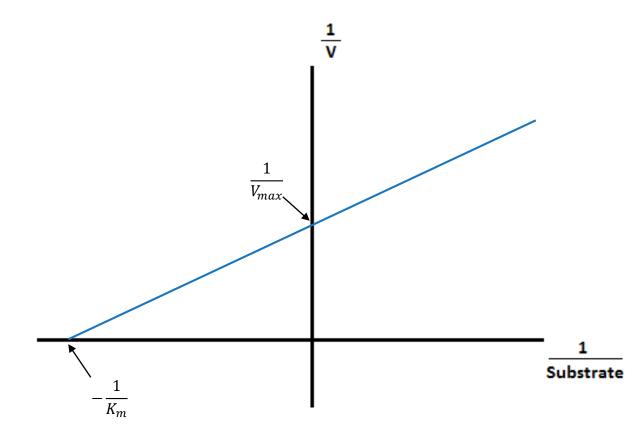
Why is K_m a useful thing to measure?

What are units for K_m ? What about K_{eq} and rate constant k?

What other assumptions do MM curves make? (Hint: [enzyme])



Lineweaver-Burk plot



Why use LB plots?

- *V_{max}* and *K_m* can be more precisely determined.
- Types of inhibition can be visualized more clearly.

Remember, info about [substrate] is still on x-axis, and info about rate is still on y-axis.



Office Hours: Biochem Content Review

Types of inhibition: understand first, memorize K_m and V_{max} effects second!

Competitive inhibition: inhibitor binds at active site V_{max} unchanged & K_m increased: why?

Noncompetitive inhibition: inhibitor binds at allosteric site V_{max} reduced & K_m unchanged: why?

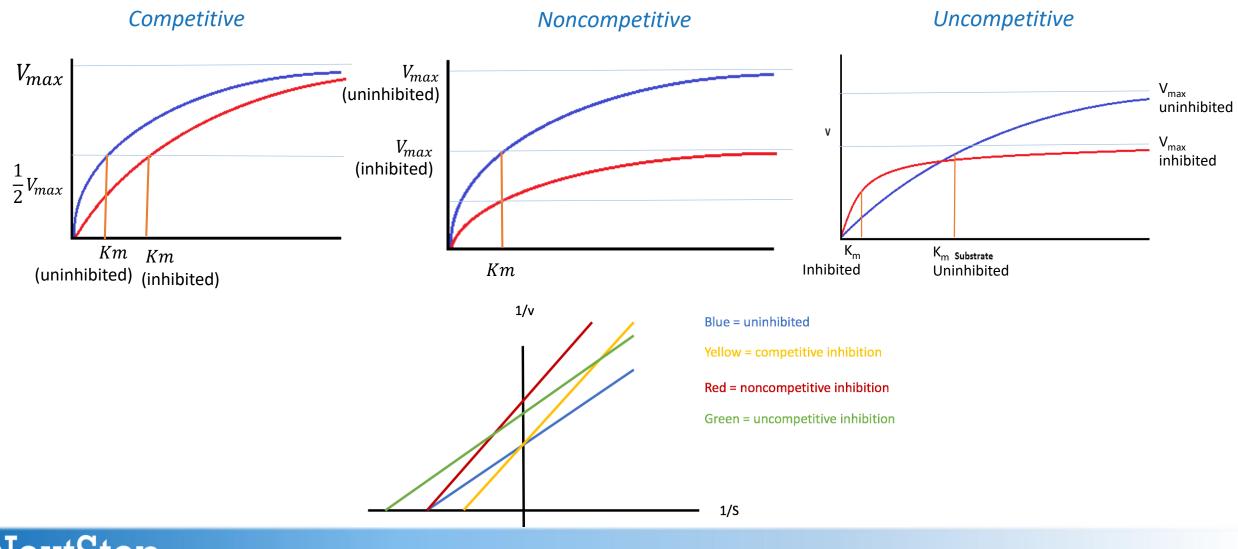
 $\mathbf{\hat{}} \rightarrow \mathbf{\hat{}} \mathbf{\hat{}} \rightarrow \mathbf{\hat{}} \mathbf{\hat{}}$



Uncompetitive inhibition: inhibitor binds E-S complex

V_{max} reduced & K_m reduced: why?





Office Hours: Biochem Content Review

1. A biochemist is investigating a reaction featured in human metabolism. He notes that all necessary reactants are present in his test tube and that the process should proceed spontaneously given the conditions, but observes that no products are being made. The most likely explanation is:

A) the rate of the uncatalyzed reaction is extremely slow.B) a particular enzyme within live cells changes the mechanism of the reaction.

C) catalysts within human cells increase the amount of free energy released during the course of the reaction.D) A and B only.

2. Catalase is an enzyme found in especially high concentrations in the liver. This molecule catalyzes the conversion of the reactive oxidative species hydrogen peroxide into water and oxygen. In the presence of this enzyme:

A) the conversion of hydrogen peroxide to water and oxygen gas is made spontaneous.

B) the rate of conversion of hydrogen peroxide to water and oxygen gas is increased.

C) the rate of conversion of water and oxygen gas to hydrogen peroxide is increased.

D) more than one of the above.



3. Priya is investigating the function of Enzyme D, which has a K_m value of 0.175 mM. She adds a large quantity of competitive inhibitor (Compound G) into her test tube. Which of the following is the apparent K_m value that she subsequently observes?

A) 0.09 mM B) 0.10 mM C) 0.175 mM D) 0.500 mM 4. Which of the following changes may impact the V_{max} or maximal reaction rate?

A) Altering the amount of enzymeB) Altering the amount of a noncompetitive inhibitorC) Altering the amount of a mixed inhibitorD) All of the above



The artificial sweetener aspartame is the methyl ester of the dipeptide of L-phenylalanine and L-aspartic acid (Figure 1). There are two general approaches to prepare aspartame. The chemical approach involves reacting the methyl ester of phenylalanine with an N-protected anhydride of aspartic acid. The protecting group, either a benzyl or formyl group is then removed by mild acid hydrolysis. In addition to the desired product, a beta structural isomer is also formed due to formation of a peptide bond with the wrong carboxylate group, which must be removed since it produces a bitter taste. A second enzymatic synthesis has been developed in which proteases catalyze the selective peptide bond formation and avoids the formation of the beta isomer.

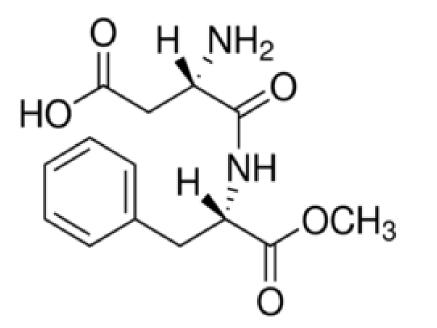


Figure 1 Structure of N-(L- α -Aspartyl)-L-phenylalanine, 1-methyl ester (Note: $pK_{a1} = 3.2$ and $pK_{a2} = 7.7$)

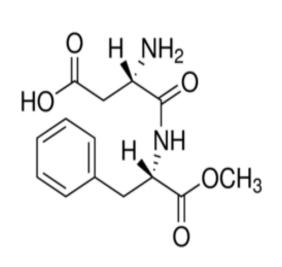


Upon ingestion, aspartame is broken down in the duodenum into its components, aspartic acid, phenylalanine and methanol, with the subsequent formation of metabolites such as formaldehyde and formic acid. Some research has raised concerns that aspartame may lead to the formation of certain cancers as a result of the formation of some of these potentially toxic compounds. A new drug, known as protein AT7 (MW = 5×10^4 amu), has been developed to counter this possibility.



25. According to the passage, the pl of aspartame is most nearly:

A. 3.2 B. 5.5 C. 7.0 D. 7.7



26. How many stereocenters are in aspartame?

- A. 1
- B. 2
- C. 3
- D. 4

Figure 1 Structure of N-(L- α -Aspartyl)-L-phenylalanine, 1methyl ester (Note: $pK_{a1} = 3.2$ and $pK_{a2} = 7.7$)

27. The two amino acids that form the basis for the dipeptide structure of aspartame, aspartic acid and phenylalanine, are most accurately be classified as:

A. hydrophilic and hydrophilic, respectively.B. hydrophobic and hydrophilic, respectively.C. hydrophilic and hydrophobic, respectively.D. hydrophobic and hydrophobic, respectively.

28. Prior to its digestion in the small intestine, aspartame must pass through the stomach. What is net charge on aspartame while in the stomach?

- A. -1 B. 0
- C. +1

D. +2

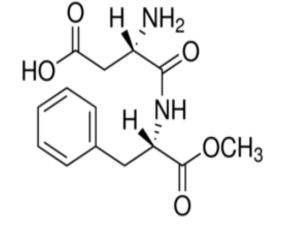


Figure 1 Structure of N-(L- α -Aspartyl)-L-phenylalanine, 1methyl ester (Note: $pK_{a1} = 3.2$ and $pK_{a2} = 7.7$)



Office Hours: Biochem Content and Difficult Passage Review

29. How many amino acid residues are in AT7?

Α.	2
Β.	50
C.	450
D.	900

"A new drug, known as protein AT7 ($MW = 5 \times 10^4$ amu), has been developed to counter this possibility." 30. Peptides are stable in water because:

A) peptide bonds cannot be cleaved by hydrolysis.B) electron sharing between the carbonyl and amino group contributes resonance stabilization across the amide bond.

C) the breakdown of peptides into individual amino acids is entropically unfavorable.

D) peptides hydrogen bond with free-floating proline residues to promote stabilization.



Any Questions?

- Any questions about the content we've reviewed?
- Thoughts about approaches to studying biochem in future? (Study sheets, etc.)



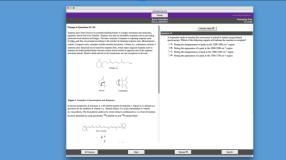
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- Clara: 526 MCAT, 5 years experience
- Phil: 5 years experience; 98% score
- Andrew: U Chicago PhD, **523 MCAT**



Bryan





Phil

Andrew

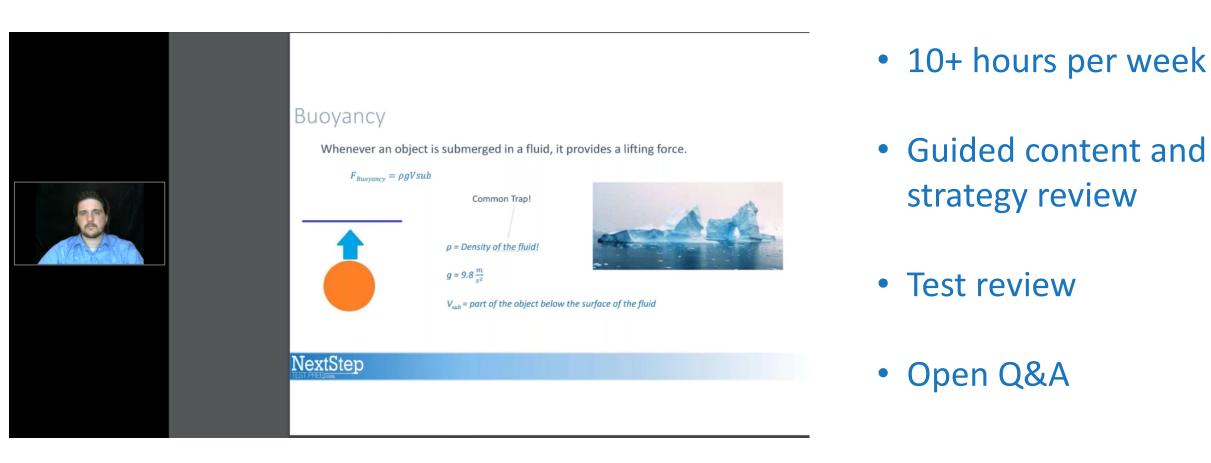
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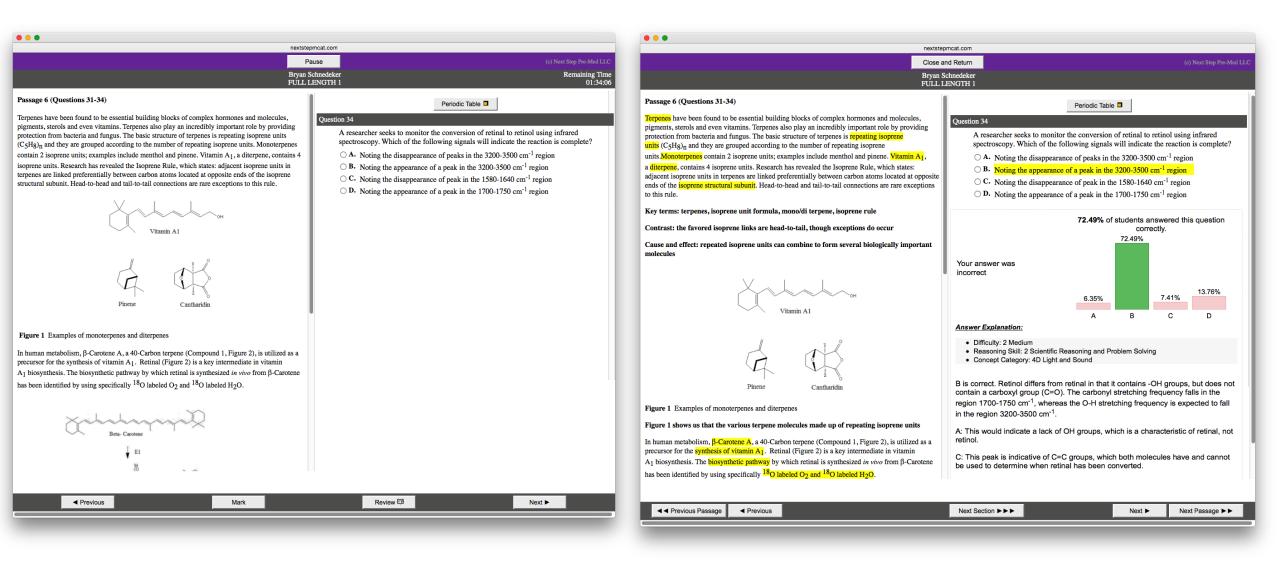
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Section	Scaled Percentile			Chemical Correct:	Results by Cor	tegory	
Chemical an Systems	Results by Reason	ing S	kills	Critical Ar	Concept Category	Total Correct	Correct Percentage
Critical Analy		Total	Correct	Correct:	1A Amino Acids and Proteins	4	57.14% (4/7)
Biological an	Concept Category	Correct	Percentage	Biological	1B Molecular Genetics	9	81.82% (9/11)
Systems	1 Knowledge of Scientific Concepts and 49 71.01 Principles	71.01% (49/69)	Correct:	1C Classical Genetics	9	69.23% (9/13)	
Psychologica Behavior	2 Scientific Reasoning and Problem Solving	39	68.42% (39/57)	Conect.	1D Metabolism	3	37.5% (3/8)
Total	3 Reasoning About the Design and Execution	23	69.7% (23/33)	Psycholog	2A Cell Biology	1	100% (1/1)
	of Research		/	e Correct:	2B Microbiology	4	80% (4/5)
_	4 Data-based and Statistical Reasoning	10	55.56% (10/18)	127 ¹²⁸ 506	2C Reproduction	3	60% (3/5)
	5 Foundations of Comprehension	12	75% (12/16)		3A Nerve and Endocrine	4	50% (4/8)
	6 Reasoning within the Text	14	60.87% (14/23)		3B Organ Systems	1	100% (1/1)
	7 Reasoning Beyond the Text	9	64.29% (9/14)		4A Kinematics and Force	5	50% (5/10)



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We've had to step away for a minute but...



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