-	I 4053 FINAL I 23, 2001	EXAM NAMI	Ξ	
Answer question 1 or 2 (16 points)		Page	Points	
1.	Asp-52 and Glu-35 have been shown to be est the enzyme lysozyme. Asp-52 has a pK of 3. hydrophobic environment, has a pK of 6.3.	5	is by 1 2 3 4	
	<ul><li>(a) What <b>fraction</b> of each of these residues is</li><li>(b) Describe the substrate for lysozyme, and (structure or word description).</li></ul>	* <u>1</u>	alyzes 5 Total	

A heptapeptide gave 2,4 dinitrophenylleucine on N-terminal analysis, and free Met as the first 2. product with carboxypeptidase. Trypsin treatment gave a tripeptide (composition: Lys, Leu, Tyr) and a tetrapeptide (composition: Trp, Met, Ala, Glu). Chymotrypsin teatment of the tetrapeptide gave two dipeptides, one migrating to the positive electrode and one migrating to the negative electrode upon electrophoresis at pH 4.5 The dipeptide migrating to the positive electrode showed U.V. absorption at 280 nm (indicating an aromatic residue). What is the sequence of the heptapeptide?

## Answer question 3 or 4 (16 points)

3. Aldolase catalyses the following reaction of glycolysis:

fructose-1,6-bisphosphate — dihydroxyacetone phosphate + glyceraldehyde-3-phosphate

- (a) Give the structures of the reactant and products of this reaction.
- (b)  $\ddot{A}G^{o'}$  for this reaction is +23.9 kJ mol<sup> $\bar{I}$ </sup>. Calculate K', the equilibrium constant. (Assume T = 37 °C or 310 K)
- (c) Calculate  $\ddot{A}G'$  for the reaction when fructose-1,6-diphosphate is  $1.0 \times 10^{-4}$  M, dihydroxyacetone phosphate is  $4.0 \times 10^{-5}$  M, and glyceraldehyde-3-phosphate is  $2.5 \times 10^{-6}$  M.
- 4. Following are three models for reversible inhibition:

(1) (2) (3)  $E + S \rightleftharpoons ES \rightarrow E + P$   $E + S \rightleftharpoons ES \rightarrow E + P$   $E + S \oiint ES \rightarrow E + P$   $E + I \rightleftharpoons EI$   $ES + I \rightleftarrows ESI$   $E + I \rightleftarrows EI$ where  $K_I = \frac{[E][I]}{[EI]}$  and  $K'_I = \frac{[ES][I]}{[ESI]}$   $ES + I \rightleftarrows ESI$ 

- (a) Identify each model by the **name** of the inhibition.
- (b) Give the kinetic equation in the reciprocal Lineweaver-Burk form that corresponds to each model.
- (c) Draw a Lineweaver Burk plot for each model, showing one line for the uninhibited reaction and a second line for a reaction containing inhibitor.

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## Answer question 5 or 6 (16 points)

- 5. You have prepared phospholipid vesicles containing  $K^+$  at a concentration of 10 mM and you place them in a solution that is 50 mM in  $K^+$ . Adding either gramicidin or valinomycin to the solution will allow more  $K^+$  to enter the cell, creating an electrical potential across the membrane (inside positive). (T = 25°C, F = 96.5 kJ-V<sup>-1</sup>-mol<sup>-1</sup>; R = 8.314 J-K<sup>-1</sup>-mol<sup>-1</sup>)
  - (a) What will be the magnitude of the potential  $(\Delta \Psi)$  when the system comes to equilibrium?
  - (b) You measure the rate of transport of K<sup>+</sup> with each antibiotic and find that transport with gramicidin is about the same at 25°C and 15°C, while the transport with valinomycin is much slower at 15°C than at 25°C. Explain this result in terms of the structure of the membrane and the mode of action of the two antibiotics.
- 6. Give the stepwise mechanism catalyzed by the enzyme **pyruvate dehydrogenase**, showing the partial structure of all the enzyme-bound intermediates. Identify the protein components catalyzing each step and the coenzymes involved as either cosubstrates or prosthetic groups.

#### Answer 7 or 8 (16 points)

- 7. In the conversion of glucose to pyruvate, three enzymes of glycolysis operate at concentrations of substrate and product removed from equilibrium, and these steps are sites of regulation. For each of the three steps, give:
  - 1. The **structure** of the reactants and products (names of coenzymes okay)
  - 2. The **name** of the enzyme.
  - 3. Metabolites or intermediates which **activate** and those which **inhibit** the enzyme.
- 8. In the oxidation of acetyl-CoA by the TCA cycle, three enzymes operate at concentrations of substrate and product removed from equilibrium, and these steps are sites of regulation. For each of the three steps, give:
  - 1. The structure of the reactants and products (names of coenzymes okay)
  - 2. The **name** of the enzyme.
  - 3. Metabolites or intermediates which **activate** and those which **inhibit** the enzyme.

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# 9. (10 points) Put in the blanks the names or abbreviations for the amino acid or acids that:

	contain sulfur.	have an amide in the side chain.		
	contain an aromatic alcohol.	contain an aliphatic alcohol		
	migrate to the negative electrode at pH 8.5.	contain a second chiral carbon.		
	migrate to the positive electrode at pH 4.0.	form a peptide bond that is cleaved by trypsin.		
	form a peptide bond that is cleaved by cyanogen bromide.	contain an imidazole ring.		
10.	(6 points) Tell whether each of the following would <b>increase</b> or <b>decrease</b> the fraction of hemoglobin sites containing oxygen.			
	a decrease in pO <sub>2</sub> .	an increase in [bis-phosphoglycerate].		
	dissociation into subunits.	an increase in pH.		
	an increase in [CO <sub>2</sub> ]	an increase in [H <sup>+</sup> ].		
11.	(10 points) Identify by name the following carbohydrates or lipids:			
	the 2-epimer of glucose	a glycerophospholipid containing three glycerol molecules		
	an 18-carbon omega-6 fatty acid.	the disaccharide repeating unit of cellulose		
	the disaccharide repeating unit of amylose	the monosaccharide repeating unit of chitin		
	the 2-epimer of D-erythrose	the lipid formed when Phosphoryl- choline is attached to ceramide		
	a non-reducing disaccharide	the characteristic sugar which defines a ganglioside		

12. (6 points) Draw the structures of guanine, cytosine, and thymine.

guanine cytosine thymine

13. (4 points) What is meant by Chargaff's rules?