## **BIOS 452/CHEM 452**

Second Exam

Fall, 2010

12:00-12:55pm, Wednesday, November 3, 2010

UIN:	

Circle Discussion Section:

Mon 8

Tue 9:30

Wed 9

Thu 9:30

Fri 9

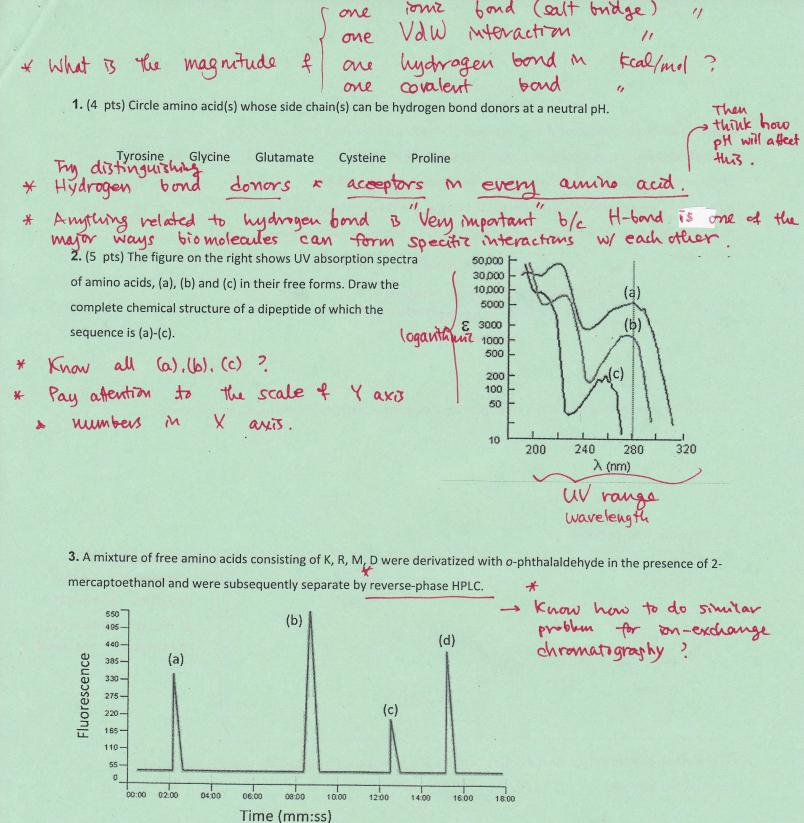
Fri 11

## General Instruction

- \* Do not turn the page until you are told to do so.
- \* You may take the exam with you only after 12:35pm.
- \* No calculators allowed. For calculations and graphing, show all your work!!!
- \* The exam is total 9 pages (including cover page), 31 questions grouped into 13 for 55 minutes.
- \* Read the question carefully to the end.

Pay attention to units.

Remember that X-ray structures do not show hydrogen atoms.

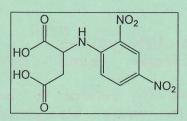


(1) (4 pts) Match the peaks with the corresponding amino acid: (a) \_\_\_\_\_; (b) \_\_\_\_\_; (c) \_\_\_\_\_; (d) \_\_\_\_\_.

(2) (2 pts) To elute the bound derivatives from the column as above, one should (increase, decrease – *Circle one*) the ratio of acetonitrile (CH<sub>3</sub>CN) to water in the mobile phase.

Page 3 of 9 How do you clute the bound molecules?

**4.** (6 pts) When a single polypeptide Z was treated with Sanger's reagent followed by acid hydrolysis, it gave an  $\alpha$ -amino derivative shown on the right. When peptide Z was treated with trypsin, it yielded two fragments. When subsequently treated with dithiothreitol (DTT), two fragments were further split into three. The sequences of the resulting three fragments are determined and they are



as follows:

EQCNK DCSR GVF

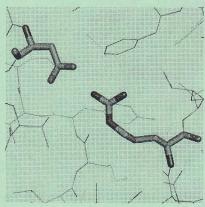
Describe the primary structure of the intact peptide Z including any covalent bond involving side chains.

## 5. Choose one from below.

(1) (3 pts) You want to determine the sequence of the first 5 residues of a protein or a peptide. Compound \_\_\_\_\_\_\_best suits this purpose.

(2) (3 pts) Compound \_\_\_\_\_ can be used to selectively determine the C-terminal residue of a protein.

6. The figure shows a portion of a protein structure determined by X-ray at pH 8. Two residues that interact with each other inside the protein core are highlighted in a stick representation. Oxygen is in red, nitrogen in blue and carbon in green.



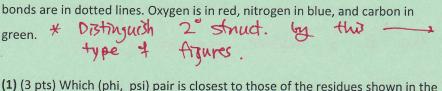
(1) (2 pts) The interaction between the two highlighted residues is a (van der Waals interaction, hydrogen bond, salt bond, metal-ligand bond - Circle one).

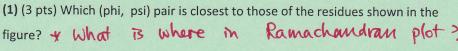
Why is this a saft band, not a H-bond?

(2) (2 pts) The strength of this interaction would (decrease, increase, remain what is this? how does it the same – Circle one) on the surface of the protein.

why the catalytic triad is partially protein hydrophobic core (in dymotrypsin, PDB: 7GCH. - Look into its catalytic site

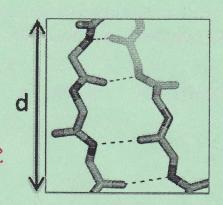
7. This figure shows a segment of a protein structure determined by X-ray crystallography. Only backbone atoms are shown in a stick model. Hydrogen 2° struct. Argures.





(molecular

(-90, -90)(-90, 90)(90, -90)(90, 90)



(2) (3 pts) Which of the following is most enriched with the type of protein scaffold shown in the figure?

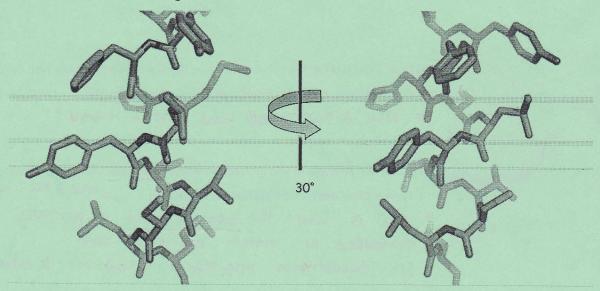
sitk fibrain wool kenation proteins farming these Tendon Name the Hair Silk Red blood Wool Hemoglobin Collagen «-Kevatin (3) (3 pts) Which of the following is closest to the distance indicated as "d" in the figure?

1x10<sup>-12</sup> m  $1x10^{-11} \text{ m}$   $1x10^{-10} \text{ m}$ 1x10<sup>-9</sup> m  $1x10^{-8}$  m  $1x10^{-7}$  m 1x10<sup>-6</sup> m confertable with

Common

(C-C) covalen

8. This is a portion of a protein structure determined by X-ray in a stick representation. Oxygen is in red, nitrogen in blue, sulfur in yellow and carbon in green.



(1) (3 pts) What is the amino acid sequence of this protein segment? ((a), (b) - Circle one)

amino auds by its 3-0 structures Recognize

(a) Y F H M L Y L V C L M

Distinguish N-ter r c-ter & a peptide regment in

(b) M L C V L **Y** L M H F Y

(2) (4 pts) On your answer for (1), circle the TWO residues that form hydrogen bonds with the backbone atoms of the underlined residue, Y.

rules that - makes up the 2° structures H-band Know

(3) (4 pts) On one of the figures, draw lines between atoms indicating the TWO hydrogen bonds that the backbone of the underlined residue Y makes.

on 3D structure Korow

> (4) (3 pts) When you transfer this protein segment from a non-polar to a polar environment, it will results in an (decrease, increase – Circle one) in  $\Delta G$  and an (decrease, increase – Circle one) in  $\Delta S$ .

Thus, the segment is likely to (exposed outside of, embedded inside of - Circle one) of a protein due to this the d-help that spans across lipid bilayer membran

a transmembrane helix Page 6 of 9

an amphiphatic helix? What Joes "amphiphatic" helix mean?

- 9. (4 pts) Which of the following statement(s) is/are NOT true?
- (a) The tertiary structure of a protein can be predicted from a known 3D structure of a protein with a similar primary structure.
- (b) The secondary structures of a protein cannot be predicted without a known 3D structure of a homologous protein.
- (c) Collagen structure is stabilized by hydrogen bonds, van der Waals interactions and covalent crosslinks between side chains.
- (d) When curly hair is converted into straight hair using  $\beta$ -mercaptoethanol during a "permanent wave (perm)" process, the constituent protein changes its secondary structure to a more extended form.

\* can you explain why " why not of I (a) - (h)?

- 10. (1) (4 pts) Hemoglobin is a conjugated protein. It contains a complex of a protoporphyrin and Fe<sup>2+</sup> which is called a
   Such non-amino acid component of a conjugated protein is called a
- (2) (4 pts) CO<sub>2</sub> is a (Substrate, Allosteric effector, Competitive inhibitor *Circle one*) for hemoglobin. And CO is a (Substrate, Allosteric effector, Competitive inhibitor – *Circle one*) for hemoglobin.
- (3) (3 pts) Myoglobin has a molecular weight of 17,000 g/mole. Estimate the molecular weight of hemoglobin in kilodalton (kDa). Show your reasoning.

  What toes it mean to say you can't to anything else.

  Myoglobin & homologous to hemoglobin?

  Da=9/mole "FDa=1000 Da".
  - 11. (5 pts) The monosaccharide drawn in a Fisher representation on the right is a
  - (D, L -Circle one)- (aldose, ketose Circle one) and also is a (triose, tetrose, pentose Circle one).

one).

Distinguish D & L M Carts protein & Lipids?

(Spart if Muder and)

Freely convert different ways to describe same structures

(2) (Bonus 3 pts) Choose the appropriate chemical structure for the monosaccharide.

(a) (b) (c) (d) OH OH

OH

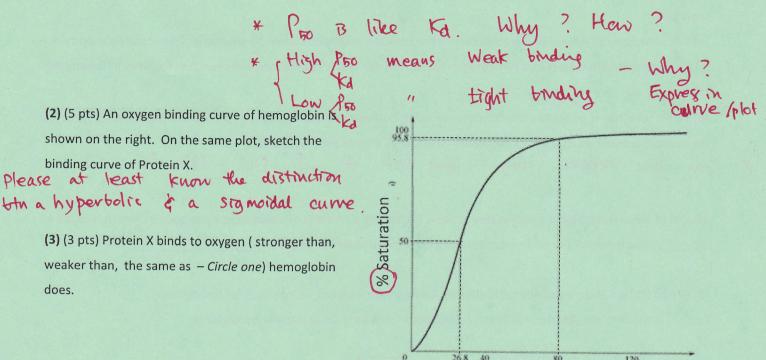
ŌН

CH<sub>2</sub>OH

\* This type of question , can be applied to any protein binding to anything.

12. Consider an oxygen binding protein X with  $P_{50}$  value of 100 mmHg and a Hill coefficient of 1.

(1) (3 pts) Calculate the ratio of the concentration of oxygen-bound X to the concentration of total X at 25 mmHg.



You can determine  $P_{50}$  looking at a hyperbolic  $O_2$  partial pressure (pO<sub>2</sub>, mmHg) curve f %. Saturation vs binding molecule's concentration.

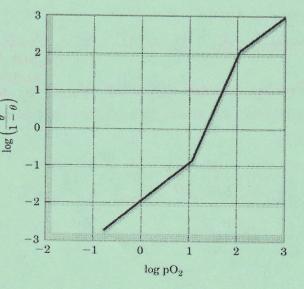
This is similar to how you can determine Km in Vo us [S] plot You see that?

(4) (3 pts) A Hill plot of hemoglobin is shown on the right. What is the Hill coefficient of hemoglobin?

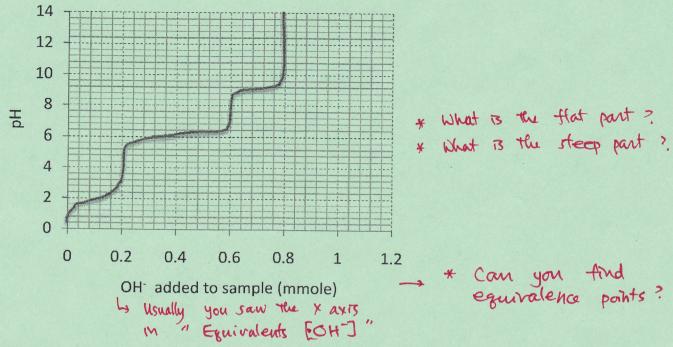
What is a Hill coefficient?

\* What is a cooperative binding? Not cooperative binding

(5) (Bonus 3 pts) On the same plot, sketch the Hill plot of Protein X.



13. Following is a titration curve of a peptide sample. The molecular weight of the peptide is 1.1 kDa.



(1) (3 pts) This peptide consists of approximately (1, 2, 10, 20, 100, 200 – Circle one) residues.

\* M.W. of an amino acid ~ 110 Da = 0.11 kDa 2 ON \* mw. of a nucleotide pair ~ 660 Pa = 0.66 FDa ) average \* (2) (3 pts) Circle amino acid residue(s) that the peptide MUST contain? Pa = 0.33 KDa

WYGSH

(3) (6 pts) What is the pl of this peptide? Show all your work.

\* Please practice w other peptide w different composition For certain, there will be a suestion of a pI.

(4) (Bonus 3 pts) Given the total volume of the sample during titration was 100 mL, what is the concentration of the peptide in this sample in molarity (M)?

\* Why figuring out pI is important/useful?

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E.g. How will you use this info for ion-exchange chromato- graphy?