## Chemistry and Biochemistry 153A, Winter 2011 Final Exam Answers – Sheet 1

- 1. a. (1) False Yeast cells can perform alcoholic fermentation; Human cells can perform homolactic fermentation
  - b. (1) False Fermentation yields less energy than oxidation
  - c. (1) True
  - d. (1) False *The reduced form is* <u>NADH</u>
  - e. (1) False Coenzyme Q is an electron carrier; Coenzyme A is an acyl carrier
  - f. (1) True
  - g. (1) True
  - h. (1) True
  - i. (1) False Coenzyme Q is found in membranes
  - j. (1) False  $FADH_2$  is not diffusible, it is part of Complex II
- 2. (3) b
- 3. a. (2) 2, 5
  - b. (2) C
  - c. (2) D, E, or F
  - d. (2) A or B
  - e. (4) Concentrations of pathway intermediates are measured and used to calculate  $\Delta G$  for each reaction. Steps with large, negative  $\Delta G$ are likely regulated.
- 4. (5) Any 5 words or phrases related to ATP synthase
- 5. a. (3) True
  - b. (3) False
  - c. (3) True
- 6. (3) Adjacent functional groups in the enzyme (e.g. heme R-groups, aa R-groups) influence the electron affinity
- 7. (4) a, b, d
- 8. a. (2) reductant; oxidized
  - b. (3)  $\Delta E^{\circ} = E^{\circ}_{acceptor} E^{\circ}_{donor}$  $= E'^{\circ}_{NAD^+} - E'^{\circ}_{ethanol} = -0.315 V - (-0.197 V)$ = -0.118 V
    - c. (2) False  $\Delta E^{\prime \circ}$  is negative, so the reaction is favored in the reverse direction

- d. (3)  $\Delta G^{\circ} = -nF\Delta E^{\circ}$ = -2(96.5 kJ/Vmol)(-0.118V) = 22.8 kJ/mol
- e. (5) At equilibrium,  $\Delta G = \Delta G^{\circ} + RT \ln Q = 0$ So  $O = e^{-\Delta G'^{\circ}/RT}$

Also,  $Q = [acet.][NADH]/[EtOH][NAD^+]$ Setting the two equal and rearranging, ratio [EtOH]/[acetaldehyde]

- $= \{ [NADH] / [NAD^+] \} \cdot e^{\Delta G'^{\circ}/RT} \\= (1/700) \cdot e^{(22.8kJ/mol)/(0.00831kJ/molK)(310K)}$
- = 10, at equilibrium

To favor forward reaction, there must be more reactant than at equilibrium, so [EtOH]/[acetaldehyde] > 10

- f. (2) C
- g. (3)  $\Delta G^{\circ}$  is measured at higher [H<sup>+</sup>] than  $\Delta G^{\circ}$  (1M versus 10<sup>-7</sup> M). Since a proton is produced in the rxn, higher  $[H^+]$  favors the reverse direction.
- 9. a. (4) nuc: -OH, hydride; elec: acetaldehyde, NAD<sup>+</sup>
  - b. (2) isozymes
  - c. (2) enz: GAPDH (2) sim: both oxidize an aldehyde and reduce NAD<sup>+</sup> (2) diff: GAPDH couples oxidation with phosphorylation
- 10. a. (2) ligase, synthetase
  - b. (2) succinyl-CoA synthetase
  - c. (6) 1. Phosphoryl transfer (to acetate from ATP), nuc: acetate; elec: ATP 2. Thioester formation. nuc: CoA. elec: acetylphosphate
- 11. a. (3) Although carbons of ethanol enter TCA cycle, they don't contribute to net synthesis of intermediates (not anaplerotic rxns)
  - b. (5) Carbons enter TCA cycle, stay and label oxaloacetate. Oxaloacetate can be converted to glucose via gluconeogenesis.
  - c. (4) carbons 3 & 4 (lower left & far left)

- 12. a. (4) ADH rxn: 1NADH
  ALDH rxn: 1NADH
  Acetate → AcCoA: -1ATP
  TCA cycle: 3NADH, 1FADH<sub>2</sub>, 1GTP
  Net: 5NADH, 1FADH<sub>2</sub>, 1GTP, -1ATP
  - b. (3) 1 NADH  $\rightarrow$  2.5 ATP, so 5NADH  $\rightarrow$ 12.5 ATP; 1FADH<sub>2</sub>  $\rightarrow$  1.5 ATP, 1GTP=1ATP
    - Sum: 12.5 + 1.5 + 1 1 = 14 ATP
  - c. (3) glc oxidation yields 32 ATP, so 34/180 = 0.18 mol ATP/gram glc
  - d. (3) ethanol is more reduced
  - e. (2) True
- 13. a. (1) Lineweaver-Burk or double-reciprocal
  - b. (4) A defined [S] is allowed to react with enzyme, and [P] vs time is measured. The initial slope of this curve, V<sub>o</sub>, is then calculated.
  - c. (2) competitive
  - d. (2) non-competitive
  - e. (3) A, C
  - f. (3) In the NAD<sup>+</sup> binding site, where the nicotinamide binds
  - g. (2) A
  - h. A. (3) 6, B. (2) 3
- 14. a. (2) A
  - b. (3) To create smaller fragments for accurate sequencing and to allow reassembly of sequence fragments (via overlapping peptides)
  - c. (2) Lys is longer but not branched, ionizable, and has a positive charge (versus neutral Gln). Also, Lys is usually not an Hbond acceptor.
  - d. (2) 5
  - e. (4)  $pI = (pKa_1+pKa_2)/2 = (4+4)/2 = 8/2 = 4$ Ionizable groups (from lowest pKa to highest): C-term COOH (~3), Glu-R COOH (~4), Glu-R COOH (~4), N-term NH<sub>3</sub><sup>+</sup> (~8), Lys-R NH<sub>3</sub><sup>+</sup> (~10.5). Below pH 4, the predominant charge states of the groups sum to a net positive charge. Above pH 4, the predominant charge states of the groups sum to a net negative charge.
  - f. (2) B
  - g. (2) No
- 15. a. (5) x-axis: pO<sub>2</sub>; y-axis:  $\theta$  (from 0 to 1.0) <u>Mb:</u> left-most curve, hyperbolic, approaching  $\theta=1$

<u>Mb + NO</u>: right-shifted from Mb curve, hyperbolic, approaching  $\theta < 1$ <u>Hb</u>: sigmoidal curve, approaching  $\theta = 1$ <u>Hb + NO</u>: left-shifted from Hb curve, less sigmoidal than Hb, approaching  $\theta < 1$ 

- b. (1) False *Hb is not an enzyme*
- 16. a. (2) B
  - b. (2) membrane lipid or membrane structure
- 17. a. (3) B
  - b. (5) (a) amylose
    - (b)  $\alpha$ -D-glucopyruanose
    - (c)  $\alpha(1 \rightarrow 4)$  O-glycosidic
    - (d) amylopectin
    - (e)  $\alpha(1 \rightarrow 6)$  O-glycosidic
  - c. (2) hydrolase, glycosidase
  - d. (3) C, G, (F)
  - e. (2) a
  - f. A. (2) Schiff base or imine
    - B. (2) amino acid (or peptide)
    - C. (2) Aldolase
    - D. (3) By acting as an electron sink, stabilizing the carbanion that results from cleavage
  - g. A. (3) The heat induces movements that break the weak interactions that stabilize tertiary structure
    - B. (2) Exposed hydrophobic patches cause protein aggregation into large, insoluble clusters
  - h. (3) O<sub>2</sub> is available, allowing aerobic breakdown of glucose; this is more efficient than fermentation and promotes yeast growth
  - i. (2) Because CO<sub>2</sub> is produced during fermentation
  - j. (2) d