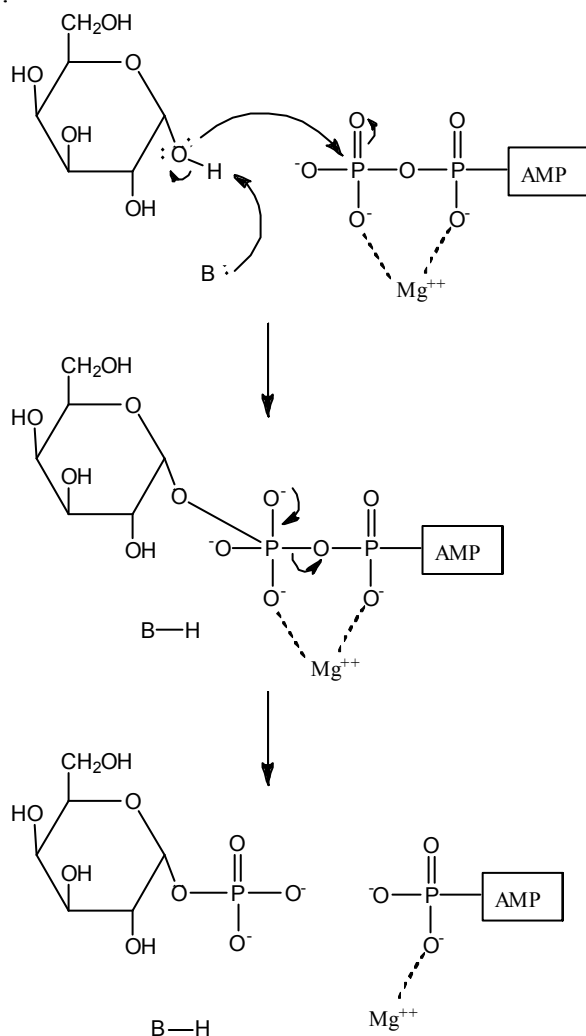


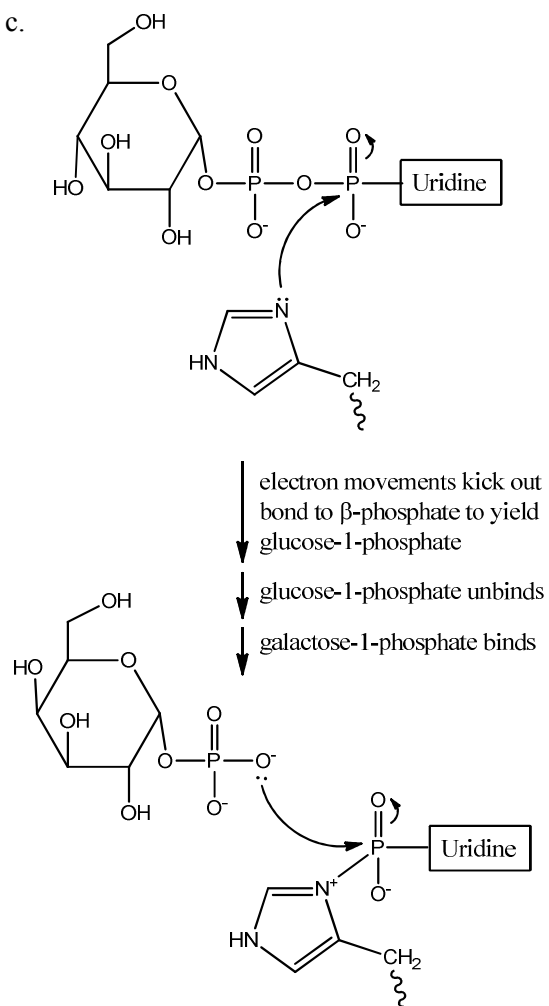
1. No. (All are formed within each subunit.)
2. 22 (10 in α subunit, 12 in β)
3. d (Asn is necessary since there is an N-linked glycosylation)
4. b
5. a. Heating would separate and denature the subunits because it breaks the weak interactions that stabilize 2°, 3°, and 4° structure.
b. D
6. a. C
b. B
7. a. B
b. The protein hormone, in its normal conformation, with its carbohydrates attached
8. a
9. The lack of sequence similarity suggests convergent evolution (in which 2 sequences start out different). However, the structures are so similar that it's possible the subunits started as the same sequence and diverged to the point where they no longer show any sequence similarity.
10. d
11. True
12. a
13. hydrophobic (van der Waals)
14. a. $\text{glucose} + \text{ATP} \rightarrow \text{glucose-6-phosphate} + \text{ADP} + \text{H}^+$
b. PIPES. The reaction produces H^+ , so a buffer that has more molecules in the deprotonated state ($\text{pK}_a < \text{pH}$) will be better able to resist the change in pH.
- c. $\text{pH} = \text{pK}_a + \log \frac{[\text{A}^-]}{[\text{HA}]}$
 $\frac{[\text{A}^-]}{[\text{HA}]} = 10^{\text{pH}-\text{pK}_a} = 10^{7.2-6.8} = 10^{0.4} = 2.51$
 $[\text{A}^-] + [\text{HA}] = 100 \text{ mM}; [\text{HA}] = 100 \text{ mM} - [\text{A}^-]$
 $\frac{[\text{A}^-]}{[\text{HA}]} = \frac{[\text{A}^-]}{(100 \text{ mM} - [\text{A}^-])} = 2.51$
 $[\text{A}^-] = 251 \text{ mM} - 2.51[\text{A}^-]$
 $3.51[\text{A}^-] = 251 \text{ mM}$
 $[\text{A}^-] = 71.5 \text{ mM}$
 $[\text{HA}] = 100 \text{ mM} - 71.5 \text{ mM} = 28.5 \text{ mM}$
- d. Mg^{2+}
- e. A, D, G, H
- f. $\Delta G'^{\circ} = -RT \ln K'_{eq} = -RT \ln \left(\frac{[\text{G6P}][\text{ADP}]}{[\text{Glc}][\text{ATP}]} \right)$
 $[\text{ADP}] = [\text{G6P}]$ and $[\text{ATP}] = [\text{Glc}]$, so $K'_{eq} = \frac{[\text{ADP}][\text{ADP}]}{([\text{ADP}][\text{ATP}]^2)} = e^{-\Delta G'^{\circ}/RT} = e^{16.7/(0.00831)(298)} = 848.7$
 $[\text{ADP}]/[\text{ATP}] = (848.7)^{0.5} = 29.1$
- g. $[\text{ADP}] + [\text{ATP}] = 10 \text{ mM}$, so
 $[\text{ATP}] = 10 \text{ mM} - [\text{ADP}]$
 $[\text{ADP}]/[\text{ATP}] = [\text{ADP}]/(10 \text{ mM} - [\text{ADP}]) = 29.1$
 $[\text{ADP}] = 291 \text{ mM} - 29.1[\text{ADP}]$
 $30.1[\text{ADP}] = 291 \text{ mM}; [\text{ADP}] = 9.67 \text{ mM}$
- h. $[\text{ADP}] = [\text{H}^+]$ produced = 9.67 mM
so, 9.67 mM deprot. buffer \rightarrow prot.
 $\text{pH} = \text{pK}_a + \log \left(\frac{[\text{A}^-]}{[\text{HA}]} \right)$
 $= 6.8 + \log \left\{ \frac{(71.5 - 9.67)}{(28.5 + 9.67)} \right\}$
 $= 6.8 + 0.2 = 7.0$
15. a. Production of glycogen in sheep relies on the activity of (non-glucokinase) hexokinase.
b. double-reciprocal or Lineweaver-Burk
c. Sheep hexokinase has a lower K_m than rat hexokinase.
d. Yes. Since the [glucose] in sheep is lower, you would expect the K_m of sheep hexokinase to be lower. (Sheep hexokinase has higher binding affinity for glucose.)
e. The higher purity sample has less of a non-competitive inhibitor.
f. The authors cannot be sure that all inhibitors have been removed.
g. True

16. a. transferase, kinase

b.



c.



d. A

e. isomerase, mutase

f. Phosphoglycerate mutase (PGM)

g. Enzyme 1. This step couples ATP hydrolysis to galactose phosphorylation, so it's likely to be highly spontaneous (like the hexokinase reaction).

h. ATP, NADH

i. galactose, ADP, AMP, P_i , NAD^+

j. D

17. a. Triacylglycerols or triglycerides
- b. 1-myristoyl-2-myristoyl-3-palmitoyl-glycerol
- c. Reactant: ATP, products: ADP + H⁺; enzyme 1 is a transferase (class) and kinase (subclass)
- d. Either glyceraldehyde-3-phosphate or dihydroxyacetonephosphate is produced by enzyme 2. Enzyme 2 is an oxidoreductase (class) and dehydrogenase (subclass)
- e. enz 1: -1 ATP
enz 2: +1 NADH
GAPDH: +1 NADH
PGK: +1 ATP
PK: +1 ATP
PDH complex: +1 NADH
isocitrate DH: +1 NADH
 α -KGDH complex: +1 NADH
sucCoA synthetase: +1 GTP
succinate DH: +1 FADH₂
malate DH: +1 NADH
Total: ATP+GTP: 2; NADH: 6; FADH₂: 1
- f. 3: oxidoreductase, dehydrogenase; 4: lyase, (hydratase); 5: oxidoreductase, dehydrogenase; 6 transferase, acyltransferase
- g. Because the β -carbon of the fatty acid is oxidized.
- h. 2x myristic acid (14 carbons) \rightarrow 2 \times 7 acetyl-CoA
1x palmitic acid (16 carbons) \rightarrow 8 acetyl-CoA
14 + 8 = 22 acetyl-CoA
- i. transport across mb: -3 ATP
 β -oxidation: produces 1 FADH₂ and 1 NADH per round
6 rounds for myristic acid; 7 for palmitic
2 \times 6 + 7 = 19 rounds, so 19 FADH₂ & 19 NADH
TCA: 3 NADH, 1 FADH₂ & 1 GTP per round
for 22 acetyl-CoA: 66 NADH, 22 FADH₂, 22 GTP
Total: ATP+GTP: 19; NADH: 85; FADH₂: 41
- j. 2+19 directly from ATP+GTP = 21
(6+85 NADH) \times 2.5 ATP/NADH = 227.5
(1+41 FADH₂) \times 1.5 ATP/FADH₂ = 63
total ATP equivalents = 311.5
- k. 1 glucose yields 32 ATP
$$\frac{311.5 \text{ ATP/lipid}}{750 \text{ Da}} \div \frac{32 \text{ ATP/glucose}}{180 \text{ Da}} = 2.3x$$