- 1. False
- 2. True
- 3. True
- 4. False
- 5. True
- 6. a, c
- 7. b
- 8. Coordinates heme iron (proximal his) H-bonds to oxygen (distal his) Reduces binding affinity of CO (distal his) Stabilizes BPG binding to T-state (through salt bridges with His2 and His143 of β-chains) Forms (stabilizing) salt bridges in T-state (His146 of β-chains) (Additional answers may also be correct)
- 9. Myoglobin facilitates diffusion of O<sub>2</sub> through muscle
- 10. Mb p50 is *lower than* Hb p50. Myoglobin must be able to pick up (bind)  $O_2$  at partial pressures where hemoglobin is releasing  $O_2$ , so it must have a higher  $O_2$  affinity (and lower p50).
- 11. False
- 12. True
- 13. True
- 14. True
- 15. True
- 16. False
- 17. a, (d), g
- 18. a.



- 18. b. The catalytic efficiency would be greater for hydrolysis of the peptide, because the peptide would bind the enzyme more tightly (in the specificity pocket and elsewhere – *p*-nitrophenylacetate doesn't bind in the specificity pocket). This would lower Km and thus increase catalytic efficiency.
- 19. a
- 20.  $V = k_3[A] k_4[B][C]$
- 21. M<sup>-1</sup>s<sup>-1</sup>
- 22. Lyase
- 23. Here, Q = [B][C]/[A] = 0.1M[C]/[A]For a spontaneous reaction,  $Q < K_{eq}$ , or 0.1M[C]/[A] < 100M, or [C]/[A] < 1000.
- 24. Yes. In the derivation we assume that [product]=0 at early time points, so any expressions that contain [B] or [C] would drop out (just as expressions with [P] drop out in the usual derivation).
- 25.  $k_{.1}[EA] + k_2[EA] = k_1[E][A] + k_{.2}[E][B][C]$ (Rate of formation of E = rate of breakdown of E)
- 26. True
- 27. 6. Oxidoreductase
  - 7. Transferase
  - 8. Isomerase
  - 9. Lyase
- Lys229: Covalent; Preferential binding of TS Glu187: General base
  Lys107: Electrostatic; Proximity & orientation effects
  Asp33: General base; General acid
- 29. The Schiff base is more electrophilic than the carbonyl of the substrate, and so it is better at accepting electrons from the broken bond. (It is better at promoting formation of a carbanion transition state at C3.)
- 30. Greater  $[H^+]$  favors the protonation of groups ( $\alpha$ -chain N-termini and  $\beta$ -chain His sidechains) on hemoglobin that must be protonated to form T-state salt bridges. So increased  $[H^+]$  favors salt bridge formation, which stabilizes the T-state, which is less likely to re-bind O<sub>2</sub> released in the capillaries.