

### Assignments for Week-12

Q1. Human serum albumin has binding sites for

- a. Diazepam and fatty acids
- b. Fatty acid and warfarin
- c. Diazepam and warfarin
- d. Fatty acids and cholesterol

Q2. Ketoprofen is an

- a. Anti-bacterial drug
- b. Anti-depressant drug
- c. Anti-coagulatory drug
- d. Anti-inflammatory drug

Q3. Enthalpy-entropy compensation can be correlated to

- a. A higher negative value of enthalpy corresponding to a higher positive value of entropy
- b. A higher positive value of enthalpy corresponding to higher positive value of entropy
- c. A higher positive value of enthalpy corresponding to higher negative value of entropy
- d. A higher negative value of enthalpy corresponding to higher negative value of entropy
- e.

Q4. Thermal unfolding of proteins can be studied by

- a. Fluorescence spectroscopy
- b. Isothermal titration calorimetry
- c. High performance liquid chromatography
- d. UV spectroscopy

Q5. The heat flow ( $dQ/dt$ ) is directly proportional to the

- a. Rate of product formation ( $d[P]/dt$ )
- b. Enthalpy ( $\Delta H$ ) of the reaction
- c. Volume of the reactants used
- d. Volume of the products formed

Q6. The rate of velocity of a reaction given by Michaelis-Menten relationship is:

a. 
$$v = \frac{d[P]}{dt} = \frac{v_0[S]}{K_M + [S]}$$

$$\text{b. } v = \frac{d[P]}{dt} = \frac{v_{\max}[S]}{K_a + [S]}$$

$$\text{c. } v = \frac{d[P]}{dt} = \frac{v_{\max}[E]}{K_M + [E]}$$

$$\text{d. } v = \frac{d[P]}{dt} = \frac{v_{\max}[S]}{K_M + [S]}$$

Q7. Enthalpy-entropy compensation can be observed due to the following reasons:

- a. Hydrogen bond dissociation
- b. Formation of new bonds
- c. Solvent reorganization
- d. Increase in Gibbs free energy

Q8. The common assumptions made in estimation of binding constant are

- a. Binding is a two-state process
- b. Activity coefficients are less than unity
- c. Ligands bind to the folded form of the protein
- d. All  $\Delta C_p$  values are temperature-independent

Q9. Rational drug design can benefit from

- a. Identification of nature of drug-protein interactions
- b. Identification of functional groups responsible for binding
- c. Understanding toxicity of the drug molecules
- d. Understanding ability of the drug to destabilize protein

Q10. Enthalpy of binding of a ligand to protein can be determined by using

- a. Differential Scanning Calorimetry
- b. Isothermal titration Calorimetry
- c. Fluorescence spectroscopy
- d. pH measurements