

31.(A) For 1st order reaction, half life is independent of initial concentration $t_{1/2} = \frac{\ln 2}{k}$

32.(C) $k = pZ.e^{-E/RT}$, in this equation k is inversely related to E. so decrease in E will result in increase of k.

33.(B) $\frac{k_{T+10}}{k_{10}} \approx 2 \text{ to } 3$

34.(C) Learn as a fact.

35.(C) Value of rate constant at two different temperature will give value of activation energy.

$$\ln \left(\frac{k_2}{k_1} \right) = \frac{E_a}{R} \left(\frac{T_2 - T_1}{T_1 T_2} \right)$$

36.(A) $k = Ae^{-E_a/RT}$, temperature dependent.

37.(D) For first order reaction, $t_{1/2}$ is constant and to complete 99.9% reaction, it requires approximately 10 times of half-life time.

$$kt = \ln \frac{100}{0.1} \quad \frac{0.693}{t_{1/2}}, t = 2.303 \log 1000$$

$$t = \frac{2.303 \times 3 \times t_{1/2}}{0.693} \approx 10t_{1/2}$$

$$\text{i.e. } \approx 4 \times 10 = 40 \text{ min.}$$

38.(C) Hydrolysis reactions are pseudo first order reactions.

39.(C) $r = k [A]^2 [B]$

40.(B) Intercept = $\ln A$

41.(C) Enzyme works as catalyst for biochemical reaction. So without enzyme, reaction has different activation energy.

42.(C) $kt = \ln \frac{100}{5} \quad t = \frac{2.303 \times t_{1/2} \times \log 20}{0.693} = 4.324 \times t_{1/2} = 17.3 \text{ days.}$

43.(C) 300 K 290 K 280 K
 $k_1 = 4k_2$ $2k_2$ k_2
Hence $k_2 = 0.25 k_1$

44.(C) Bimolecular reaction are explained by collision theory.

45.(C) $kt = \ln \left(\frac{100}{10} \right) = \ln 10$

$$t = \frac{\ln 10}{k} = \frac{\ln 10}{\ln 2} \times t_{1/2}$$

$$t = \frac{2.303}{0.693} t_{1/2} \approx 3.3 t_{1/2}$$