

Chemical Kinetics

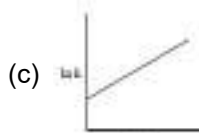
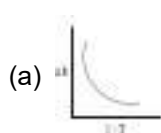
12th Standard

Chemistry

Exam Time : 02:05:00 Hrs

Total Marks : 125

- For a first order reaction $A \rightarrow B$ the rate constant is $x \text{ min}^{-1}$. If the initial concentration of A is $0.01M$, the concentration of A after one hour is given by the expression.
 (a) $0.01 \cdot e^{-x}$ (b) $1 \times 10^{-2}(1 - e^{-60x})$ (c) $(1 \times 10^{-2})e^{-60x}$ (d) none of these
- A zero order reaction $X \rightarrow \text{XProduct}$, with an initial concentration $0.02M$ has a half life of 10 min. if one starts with concentration $0.04M$, then the half life is
 (a) 10 s (b) 5 min (c) 20 min (d) cannot be predicted using the given information
- Among the following graphs showing variation of rate constant with temperature (T) for a reaction, the one that exhibits Arrhenius behavior over the entire temperature range is



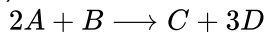
(d) both (b) and (c)

- For a first order reaction $A \rightarrow \text{product}$ with initial concentration $x \text{ mol L}^{-1}$, has a half life period of 2.5 hours. For the same reaction with initial concentration $\left(\frac{x}{2}\right) \text{ mol L}^{-1}$ the half life is
 (a) $(2.5 \times)$ (b) (c) 2.5 (d) Without knowing the rate constant, $t_{1/2}$ cannot be determined from the given data
- For the reaction, $2NH_3 \rightarrow N_2 + 3H_2$, if $-\frac{d[NH_3]}{dt} = k_1[NH_3]$, $\frac{d[N_2]}{dt} = k_2[NH_3]$, $\frac{d[H_2]}{dt} = k_3[NH_3]$ then the relation between k_1 , k_2 and k_3 is
 (a) $k_1 = k_2 = k_3$ (b) $k_1 = 3k_2 = 2k_3$ (c) $1.5k_1 = 3k_2 = k_3$ (d) $2k_1 = k_2 = 3k_3$
- The decomposition of phosphine (PH_3) on tungsten at low pressure is a first order reaction. It is because the (NEET)
 (a) rate is proportional to the surface coverage (b) rate is inversely proportional to the surface coverage (c) rate is independent of the surface coverage (d) rate of decomposition is slow
- For a reaction $\text{Rate} = k[\text{acetone}]^{3/2}$ then unit of rate constant and rate of reaction respectively is
 (a) $(\text{mol L}^{-1} \text{ s}^{-1})$, $(\text{mol}^{1/2} \text{ L}^{1/2} \text{ s}^{-1})$ (b) $(\text{mol}^{-1/2} \text{ L}^{1/2} \text{ s}^{-1})$, $(\text{mol L}^{-1} \text{ s}^{-1})$ (c) $(\text{mol}^{1/2} \text{ L}^{1/2} \text{ s}^{-1})$, $(\text{mol L}^{-1} \text{ s}^{-1})$ (d) (mol L s^{-1}) , $(\text{mol}^{1/2} \text{ L}^{1/2} \text{ s}^{-1})$
- The addition of a catalyst during a chemical reaction alters which of the following quantities? (NEET)
 (a) Enthalpy (b) Activation energy (c) Entropy (d) Internal energy
- Consider the following statements :
 (i) increase in concentration of the reactant increases the rate of a zero order reaction.
 (ii) rate constant k is equal to collision frequency A if $E_a = 0$
 (iii) rate constant k is equal to collision frequency A if $E_a = 0$
 (iv) a plot of $\ln(k)$ vs T is a straight line.
 (v) a plot of $\ln(k)$ vs $\left(\frac{1}{T}\right)$ is a straight line with a positive slope.
 Correct statements are
 (a) (ii) only (b) (ii) and (iv) (c) (ii) and (v) (d) (i), (ii) and (v)
- In a reversible reaction, the enthalpy change and the activation energy in the forward direction are respectively $-x \text{ kJ mol}^{-1}$ and $y \text{ kJ mol}^{-1}$. Therefore, the energy of activation in the backward direction is
 (a) $(y-x) \text{ kJ mol}^{-1}$ (b) $(x+y) \text{ kJ mol}^{-1}$ (c) $(x-y) \text{ kJ mol}^{-1}$ (d) $(x+y) \times 10^3 \text{ J mol}^{-1}$
- What is the activation energy for a reaction if its rate doubles when the temperature is raised from $200K$ to $400K$? ($R = 8.314 \text{ JK}^{-1} \text{ mol}^{-1}$)
 (a) $234.65 \text{ kJ mol}^{-1} \text{ K}^{-1}$ (b) $434.65 \text{ kJ mol}^{-1} \text{ K}^{-1}$ (c) $434.65 \text{ J mol}^{-1} \text{ K}^{-1}$ (d) $334.65 \text{ J mol}^{-1} \text{ K}^{-1}$
- For a first order reaction, the rate constant is 6.909 min^{-1} . the time taken for 75% conversion in minutes is
 (a) $\left(\frac{3}{2}\right) \log 2$ (b) $\left(\frac{2}{3}\right) \log 2$ (c) $\left(\frac{3}{2}\right) \log \left(\frac{3}{4}\right)$ (d) $\left(\frac{2}{3}\right) \log \left(\frac{4}{3}\right)$

- 13) In a first order reaction $x \rightarrow y$ if k is the rate constant and the initial concentration of the reactant x is 0.1M, then, the half life is

(a) $\left(\frac{\log 2}{k}\right)$ (b) $\left(\frac{0.693}{(0.1)k}\right)$ (c) $\left(\frac{\ln 2}{k}\right)$ (d) none of these

- 14) Predict the rate law of the following reaction based on the data given below



Reaction number	[A] (min)	[B] (min)	Initial rate (M s ⁻¹)
1	0.1	0.1	x
2	0.2	0.1	2x
3	0.1	0.2	4x
4	0.2	0.2	8x

- (a) rate = $k[A]^2[B]$ (b) rate = $k[A][B]^2$ (c) rate = $k[A][B]$ (d) rate = $k[A]^{1/2}[B]^{1/2}$
- 15) **Assertion:** rate of reaction doubles when the concentration of the reactant is doubles if it is a first order reaction.

Reason: rate constant also doubles

- (a) Both assertion and reason are true and reason is the correct explanation of assertion. (b) Both assertion and reason are true but reason is not the correct explanation of assertion. (c) Assertion is true but reason is false. (d) Both assertion and reason are false.

- 16) The rate constant of a reaction is 5.8×10^{-2} . The order of the reaction is

- (a) First order (b) zero order (c) Second order (d) Third order

- 17) For the reaction $N_2O_5(g) \rightarrow 2NO_2(g) + \frac{1}{2}O_2(g)$ value of rate of

disappearance of N_2O_5 is given as $6.5 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1}$ The rate of formation of NO_2 and O_2 is given respectively as

- (a) $(3.25 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1})$ (b) $(1.3 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1})$ (c) $(1.3 \times 10^{-1} \text{ mol L}^{-1} \text{ s}^{-1})$ (d) and $(1.3 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1})$ and $(3.25 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1})$ and $(3.25 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1})$ None of these

- 18) During the decomposition of H_2O_2 to give dioxygen, 48 g O_2 is formed per minute at certain point of time. The rate of formation of water at this point is

- (a) $0.75 \text{ mol min}^{-1}$ (b) 1.5 mol min^{-1} (c) $2.25 \text{ mol min}^{-1}$ (d) 3.0 mol min^{-1}

- 19) If the initial concentration of the reactant is doubled, the time for half reaction is also doubled. Then the order of the reaction is

- (a) Zero (b) one (c) Fraction (d) none

- 20) In a homogeneous reaction $A \rightarrow B + C + D$ the initial pressure was P_0 and after time t it was P . expression for rate constant in terms of P_0 , P and t will be

(a) $k = \left(\frac{2.303}{t}\right) \log \left(\frac{2P_0}{3P_0 - P}\right)$ (b) $k = \left(\frac{2.303}{t}\right) \log \left(\frac{2P_0}{P_0 - P}\right)$

(c) $k = \left(\frac{2.303}{t}\right)$

- 21) If 75% of a first order reaction was completed in 60 minutes, 50% of the same reaction under the same conditions would be completed in

- (a) 20 minutes (b) 30 minutes (c) 35 minutes (d) 75 minutes

- 22) The half life period of a radioactive element is 140 days. After 560 days, 1 g of element will be reduced to

(a) $\left(\frac{1}{2}\right) g$ (b) $\left(\frac{1}{4}\right) g$ (c) $\left(\frac{1}{8}\right) g$ (d) $\left(\frac{1}{16}\right) g$


- 23) The correct difference between first and second order reactions is that (NEET)

- (a) A first order reaction first order reaction can be catalysed; a second order reaction cannot be catalysed. (b) The half life of a second order reaction does not depend on $[A_0]$. (c) The rate of a first order reaction does not depend on reactant concentrations; the rate of a second order reaction does depend on reactant concentrations. (d) The rate of a first order reaction does depend on reactant concentrations; the rate of a second order reaction does not depend on reactant concentrations.

- 24) After 2 hours, a radioactive substance becomes $\left(\frac{1}{16}\right)^{th}$ of original amount Then

the half life (in min) is

- (a) 60 minutes (b) 120 minutes (c) 30 minutes (d) 15 minutes

- 25)  This reaction follows first order kinetics. The rate constant at particular temperature is $2.303 \times 10^{-2} \text{ hour}^{-1}$. The initial concentration of cyclopropane is 0.25 M. What will be the concentration of cyclopropane after 1806 minutes? ($\log 2 = 0.3010$)

(a) 0.125 M (b) 0.215 M (c) 0.25×2.303 M (d) 0.05 M

- 26) $2\text{N}_2\text{O}_5 \rightarrow \text{NO}_2 + \text{O}_2$, $\frac{d[\text{N}_2\text{O}_5]}{dt} = k_1[\text{N}_2\text{O}_5]$, $\frac{d[\text{NO}_2]}{dt} = k_2[\text{N}_2\text{O}_5]$ and $\frac{d[\text{O}_2]}{dt} = k_3[\text{N}_2\text{O}_5]$,

the relation between k_1 , k_2 and k_3 is

(a) $2k_1 = 4k_2 = k_3$ (b) $k_1 = k_2 = k_3$ (c) $2k_1 = k_2 = 4k_3$ (d) $2k_1 = k_2 = k_3$

- 27) What would be the rate of disappearance of oxygen, if the rate of formation of nitric oxide (NO) is $3.6 \times 10^{-3} \text{ mol L}^{-1} \text{ s}^{-1}$?

(a) $4 \times 10^{-3} \text{ mol L}^{-1} \text{ s}^{-1}$ (b) $4 \times 10^{-3} \text{ mol}^{-1} \text{ L}^{-1} \text{ s}^{-1}$ (c) $4.5 \times 10^{-3} \text{ mol L}^{-1} \text{ s}^{-1}$ (d) $4.5 \times 10^{-3} \text{ mol}^{-1} \text{ L}^{-1} \text{ s}^{-1}$

- 28) For a reaction, $2\text{A} + \text{B} \rightarrow 3\text{C}$, The rate of appearance of C at time 't' is $1.2 \times 10^{-4} \text{ mol L}^{-1} \text{ s}^{-1}$. Identify the rate of reaction.

(a) $4 \times 10^{-5} \text{ mol L}^{-1} \text{ s}^{-1}$ (b) $4.5 \times 10^{-1} \text{ mol L}^{-1} \text{ s}^{-1}$ (c) $3.6 \times 10^{-4} \text{ mol L}^{-1} \text{ s}^{-1}$ (d) None of these

- 29) For the reaction, $2\text{N}_2\text{O}_5 \rightarrow 4\text{NO}_2 + \text{O}_2$, select the correct statement.

(a) Rate of formation of O_2 is same as rate of formation of NO_2 (b) Rate of disappearance of N_2O_5 is two times the rate of formation of NO_2 (c) Rate of formation of O_2 is 0.5 times the rate of disappearance of N_2O_5 (d) Rate of formation of NO_2 is equal to rate of disappearance of N_2O_5

- 30) In pseudo-order reactions

(a) The actual order of reaction is different from that expected using rate law expression (b) The concentration of at least one reactant is taken in large excess. (c) The concentration of reactant taken in excess may be taken as constant of these (d) All

- 31) The depletion of ozone involves the following steps:

Step 1: $\text{O}_2 + \text{O} \xrightleftharpoons[k_2]{k_1} \text{O}_3$ (fast)

Step 2: $\text{O}_3 + \text{O} \xrightarrow{k} 2\text{O}_2$ (slow)

The predicted order of the reaction will be

(a) I (b) II (c) III (d) Zero

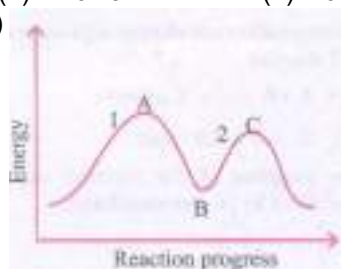
- 32) What would be the activation energy of a reaction when the temperature is increased from 27°C to 37°C ?

(a) 534 kJ mol^{-1} (b) 53.4 kJ mol^{-1} (c) 5.34 kJ mol^{-1} (d) None of these

- 33) $\text{A} + \text{B} \rightarrow \text{C}$; $\Delta H = 60 \text{ kJ mol}^{-1}$ $E_a = 150 \text{ kJ}$. What is the activation energy of the backward reaction?

(a) 210 kJ (b) 105 kJ (c) 90 kJ (d) 145 kJ

- 34)



(a) It is a 2 steps reaction, step 1 is slower than step 2 (b) It is a 2 steps reaction, step 2 is slower than step 1. (c) Single step reaction where B is activated complex (d) Single step reaction in which B is a reaction intermediate.

- 35) The Unit of rate constant and rate of reaction are same for

(a) First order (b) second order (c) Third order (d) zero order

- 36) Which of the following does not affect the rate of reaction?

(a) Amount of the reactant taken (b) Physical state of the reactant (c) ΔH of reaction (d) Size of vessel

- 37) Which of the following statement is not correct?

(a) Molecularity of a reaction cannot be fractional (b) Molecularity of a reaction cannot be more than three (c) Molecularity of a reaction can be zero (d) Molecularity is assigned for each elementary step of mechanism.

- 38) Which order reaction obeys the expression $t_{1/2} \propto \frac{1}{[A]}$?

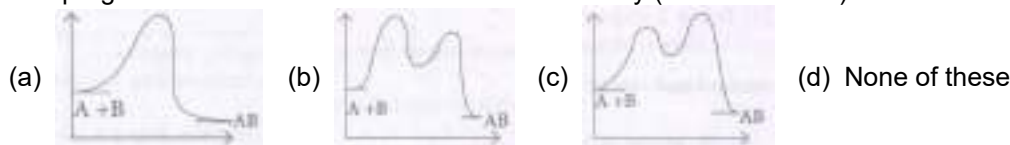
(a) First (b) Second (c) Third (d) Zero

- 39) For an exothermic chemical process occurring in 2 steps as

(i) $\text{A} + \text{B} \rightarrow \text{X}$ (slow) ;

(ii) $X \rightarrow AB$ (fast)

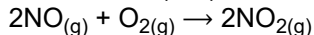
The progress of the reaction can be best described by (x- intermediate).



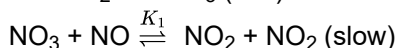
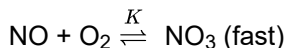
40) The graph between the $\log K$ versus $\frac{1}{T}$ is a straight line. The slope of the line is

- (a) $\frac{-2.303R}{E_a}$ (b) $\frac{E_a}{2.303R}$ (c) $\frac{2.303R}{E_a}$ (d) $\frac{E_a}{2.303R}$

41) Nitric oxide (NO) reacts with oxygen to produce nitrogen dioxide



If the mechanism of reaction is



- (a) Rate = $K' [\text{NO}] [\text{O}_2]$ (b) Rate = $K' [\text{NO}] [\text{O}_2]^2$ (c) Rate = $K' [\text{NO}]^2 [\text{O}_2]$ (d) Rate = $K' [\text{NO}]^3 [\text{O}_2]$

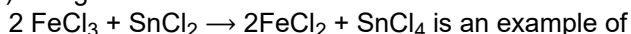
42) The addition of a catalyst during a chemical reaction alters which of the following quantities?

- (a) Activation energy (b) Entropy (c) Internal energy (d) Enthalpy

43) A reaction having equal activation energies for forward and reverse reactions has

- (a) $\Delta G = 0$ (b) $\Delta H = 0$ (c) $\Delta H = \Delta G = \Delta S = 0$ (d) $\Delta S = 0$

44) The given reaction



- (a) I order (b) II order (c) III order (d) None of these

45) Enzymes increase the rate of reactions.

- (a) By lowering activation energy (b) By increasing activation energy (c) By changing equilibrium constant (d) By forming enzyme substrate complex

46) Activation energy of a chemical reaction can be determined by

- (a) Evaluating rate constants at two different temperatures (b) Evaluating velocities of reaction at two different temperatures (c) Evaluating rate constant at standard concentration of reactants (d) Changing temperature

47) $2A \rightarrow B + C$. It would be a zero order reaction when

- (a) The rate of reaction is proportional to square of conc. of A (b) The rate of reaction remains same at any conc. of A (c) The rate of reaction remains unchanged at any conc. of B and C (d) The rate of reaction doubles if conc. of B is increased to double.

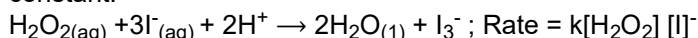
48) What will be the rate constant of a I order reaction if its half life is given to be 20 min?

- (a) 13.86 min^{-1} (b) 28.86 min^{-1} (c) $3.47 \times 10^{-2} \text{ min}^{-1}$ (d) None of these

49) How much time will be taken for 20 gm to reduce 5 g? [$R = 2 \times 10^{-3} \text{ s}^{-1}$ (I order reaction)]

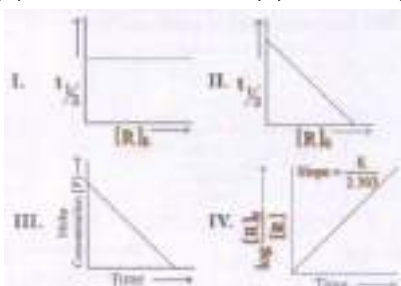
- (a) 693.1 s (b) 693.1 s^{-1} (c) 6.931 s (d) 6.931 s^{-1}

50) For the following reaction, identify the order of reaction and dimension of the rate constant.



- (a) $2, \text{L mol}^{-1} \text{ s}^{-1}$ (b) $1, \text{s}^{-1}$ (c) $\frac{3}{2}, \text{L}^{1/2} \text{ mol}^{-1/2} \text{ s}^{-1}$ (d) None of these

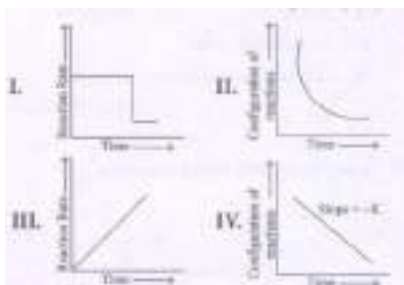
51)



Which of the above graphs is correct for first order reactions?

- (a) I, II (b) II, III (c) I, IV (d) I, III

52)



Which of the above graphs is correct for zero order reactions?

- (a) I, II (b) I, III (c) I, IV (d) II, III

53) During decomposition of an activated complex

- (a) energy is always released (b) energy is always absorbed (c) energy does not change (d) products may be formed.

54) A first order reaction is 50% completed in 1.26×10^{14} . How much time would it take for 100% completion?

- (a) 1.26×10^{15} s (b) 2.52×10^{14} s (c) 2.52×10^{28} s (d) Infinite

55) Concentration is expressed in?

- (a) $\frac{\text{number of moles/litre}}{\text{time in sec}}$ (b) $\frac{\text{time in sec}}{\text{number of moles/litre}}$ (c) $\frac{\text{number of moles/litre}}{\text{volume}}$ (d) $\frac{\text{volume}}{\text{number of moles/litre}}$

56) During a chemical reaction, the concentration of reaction

- (a) increases (b) decreases (c) remains constant (d) first increases and then decreases

57) For the second order reaction $t_{\frac{1}{2}} \propto$

- (a) $\frac{1}{a^2}$ (b) $\frac{1}{a}$ (c) Constant (d) a

58) The unit of zero order rate constant is

- (a) $\text{litre mol}^{-1} \text{sec}^{-1}$ (b) $\text{mol litre}^{-1} \text{sec}^{-1}$ (c) sec^{-1} (d) $\text{litre}^2 \text{sec}^{-1}$

59) The excess energy which a molecule must possess to become active is known as

- (a) kinetic energy (b) threshold energy (c) potential energy (d) activation energy

60) Arrhenius equation is

- (a) $k = Ae^{-1/RT}$ (b) $k = Ae^{RT/E_a}$ (c) $k = Ae^{-E_a/RT}$ (d) $k = Ae^{E_a/RT}$

61) The term A in Arrhenius equation is called as

- (a) Probability factor (b) Activation energy (c) Collision factor (d) Frequency factor

62) The half life period of a first order reaction is 10 minutes. Then its rate constant is

- (a) $6.93 \times 10^{-2} \text{ min}^{-1}$ (b) $0.693 \times 10^{-2} \text{ min}^{-1}$ (c) $6.932 \times 10^{-2} \text{ min}^{-1}$ (d) $69.3 \times 10^{-1} \text{ min}^{-1}$

63) For a reaction: $aA \rightarrow bB$, the rate of reaction is doubled when the concentration of A is increased by four times. The rate of reaction is equal to

- (a) $k[A]^a$ (b) $k[A]^{\frac{1}{2}}$ (c) $k[A]^{\frac{1}{a}}$ (d) $K[A]$

64) Compound A reacts by first order kinetics. At 25°C , the rate constant of the reaction is 0.60 sec^{-1} . What is the half life of A?

- (a) 1.15 sec (b) 0.4158 sec (c) 0.093 sec (d) 1.29 sec

65) Pick out the odd one.

- (a) ${}_{92}\text{U}^{238} \rightarrow {}_{90}\text{Th}^{234} + {}_2\text{He}^4$ (b) $\text{SO}_2 + \text{Cl}_2 \xrightarrow{\text{H}^+} \text{H}_2\text{SO}_3$ (c) $\text{CH}_3\text{COOCH}_3 + \text{H}_2\text{O} \xrightarrow{\text{H}^+} \text{CH}_3\text{COOH} + \text{CH}_3\text{OH}$ (d) $\text{C}_{12}\text{H}_{22}\text{O}_{11} + \text{H}_2\text{O} \xrightarrow{\text{H}^+} \text{C}_6\text{H}_{12}\text{O}_6 + \text{C}_6\text{H}_{12}\text{O}_6$

66) If the initial concentration of the reactants is doubled, then rate

- (a) remains constant (b) doubled its value (c) is reduced to half of initial rate (d) is increased by four times of initial rate

67) The rate of a reaction is expressed as $k[A]^2[B]^1$, the reaction is said to be

- (a) second order with respect to A (b) first order with respect to B (c) overall order of the reaction is third order (d) all the above

68) Which one of the following is an example of pseudo first order reaction?

- (a) Acid hydrolysis of ester (b) Decomposition of HI (c) Synthesis of NH_3 (d) All radioactive transformations

69) The rate constant of a first order reaction is $2.0 \times 10^{-6} \text{ s}^{-1}$. The initial concentration of the reactant is 0.10 mol dm^{-3} . What is the value of the initial rate?

- (a) $2.0 \times 10^{-7} \text{ mol dm}^{-3} \text{ s}^{-1}$ (b) $2.0 \times 10^{-6} \text{ mol dm}^{-3} \text{ s}^{-1}$ (c) $2.0 \times 10^{-5} \text{ mol dm}^{-3} \text{ s}^{-1}$ (d) $2.0 \times 10^7 \text{ mol dm}^{-3} \text{ s}^{-1}$

70) If the activation energy is high then the rate of the reaction is

- (a) high (b) moderate (c) low (d) cannot be predicted

71) Pick out the effect of catalyst on activation energy?

- (a) Catalyst lowers the (b) Catalyst provides alternate (c) Both (a) (d) None of

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- activation energy path to the reaction and (b) these
- 72) In Arrhenius equation, if EA is positive and $T_2 > T_1$, then _____
 (a) $k_1 = k_2$ (b) $k_2 < k_1$ (c) $k_1 = \frac{1}{k_2}$ (d) $k_2 > k_1$
- 73) For the reaction $A \rightarrow C$, it is found that the rate of the reaction quadruples when the concentration of A is doubled. The rate for the reaction is $\text{Rate} = [A]^n$. The value of n is _____
 (a) 1 (b) 2 (c) 0 (d) 3
- 74) In the graph showing Maxwell, Boltzmann distribution of energy _____
 (a) area under the curve must not change with increase in temperature. (b) area under the curve increases in temperature. (c) area under the curve decreases with increase in temperature. (d) None of these above
- 75) For a complex reaction _____
 (a) Order of overall reaction is same as molecularity of the slowest step. (b) Order of overall reaction is less than the molecularity of the slowest step. (c) Order of overall reaction is greater than the molecularity of the slowest step. (d) None of these above.
- 76) In any unimolecular reaction _____
 (a) Only two reacting species is involved in the rate determining step. (b) The order and the molecularity of slowest step are equal to one. (c) The molecularity of the reaction is one and order is zero. (d) Both molecularity and order of the reaction are one.
- 77) Rate law cannot be determined from balanced chemical equation if _____
 (a) Reverse reactions is not involved (b) It is an elementary reaction (c) It is a sequence of elementary reactions (d) All of the reactants is in excess. Rate law can be determined from balanced chemical equation if it is an elementary reaction.
- 78) The value of rate constant of a pseudo first order reaction _____
 (a) Independent on the concentration of reactants present in small amount (b) Depends on the concentration of reactants present in excess (c) Independent of the concentration of reactants (d) Depends only on temperature.
- 79) $\text{CH}_3\text{COOCH}_3 + \text{H}_2\text{O} \xrightarrow{\text{H}^+} \text{CH}_3\text{COOH} + \text{CH}_3\text{OH}$ is an example of _____ order reaction.
 (a) first (b) zero (c) third (d) pseudo
- 80) When the E_a of a reaction zero then the rate constant of the reaction is equal to _____
 (a) 2.303 K (b) $\frac{K}{2.303}$ (c) $\left(t_{\frac{1}{2}}\right)$ (d) A
- 81) By the order of reaction we mean _____
 (a) the sum of powers to which the concentration terms are raised in the rate equation (b) the number of reactants take part in the reaction (c) the number of concentration terms in the velocity equation for the reaction (d) the least number of product molecule needed for the reaction
- 82) The oxidation of potassium iodide by potassium persulphate as per the rate law, rate $k[\text{K}_2\text{S}_2\text{O}_8][\text{KI}]$. The order with respect to potassium iodide is, _____
 (a) two (b) one (c) three (d) four
- 83) If the rate law for a reaction $A + B \rightarrow$ is $C \frac{d[A]}{dt} \times \frac{1}{2} = k[A]^n[B]^m$ then the order of a reaction is _____
 (a) n (b) m (c) n + m (d) m - n
- 84) For a reaction $2A + B \rightarrow 3C$, express the rate of reaction in terms of formation of the product _____
 (a) $\frac{1}{2} \frac{d[A]}{dt}$ (b) $-\frac{1}{3} \frac{d[C]}{dt}$ (c) $\frac{1}{3} \frac{d[C]}{dt}$ (d) $-\frac{1}{2} \frac{d[B]}{dt}$
- 85) Energy of activation of a reactant is reduced by _____
 (a) reduced pressure (b) increased pressure (c) reduced temperature (d) increased temperature
- 86) Activation energy is equal to _____
 (a) Threshold energy + Energy of colliding molecules (b) Threshold energy (c) Threshold energy x Energy of colliding molecules (d) Threshold energy - Energy of colliding molecules
- 87) For the reaction $\text{N}_{2(g)} + 3\text{H}_{2(g)} \rightarrow 2\text{NH}_{3(g)}$ the rate of the reaction in terms of ammonia is _____
 (a) $+\frac{1}{2} \frac{-d[\text{NH}_3]}{dt}$ (b) $-\frac{1}{2} \frac{d[\text{NH}_3]}{dt}$ (c) $-\frac{d[\text{NH}_3]}{dt}$ (d) $+\frac{d[\text{NH}_3]}{dt}$
- 88) The magnitude of order of a reaction may be _____

(a) fractional (b) zero (c) integral values (d) all of these
89) For a reaction, $E_a = 0$ and $k = 4.2 \times 10^5 \text{ sec}^{-1}$ at 300 K, the value of k at 310 K will be

(a) $4.2 \times 10^5 \text{ sec}^{-1}$ (b) $8.4 \times 10^5 \text{ sec}^{-1}$ (c) $8.4 \times 10^5 \text{ sec}^{-1}$ (d) unpredictable
90) For an elementary reaction, its order is never _____ since it is a _____ step process.

(a) integral, single (b) fractional, single (c) zero, one (d) fractional, zero
91) In the acid hydrolysis of an ester, what is the time taken for complete hydrolysis?
(a) 8 hours (b) 45 minutes (c) both (a) and (b) (d) None

92) $\text{CH}_3\text{COOCH}_3 + \text{H}_2\text{O} \xrightarrow{\text{H}^+} \text{CH}_3\text{COOH} + \text{CH}_3\text{OH}$ is an example of _____ order reaction.
(a) first (b) zero (c) third (d) pseudo

93) The minimum energy that all colliding molecules must possess so as to make the collisions more effective and successful is _____.
(a) activation energy (b) colliding energy (c) threshold energy (d) kinetic energy

94) For a general reaction $a\text{A} + b\text{B} \rightarrow \text{Products}$, the rate of the reaction is equal to _____.
(a) $k[\text{A}]^p [\text{B}]^q$ (b) $k [\text{A}] [\text{B}]$ (c) k (d) $\frac{1}{k}$

95) The time required for 50% completion of the reaction is known as _____.
(a) Average life period (b) Half-life period (c) Rate (d) None of these

96) In a second order reaction, if one of the concentration is excess, the order of the reaction is _____ order reaction.
(a) first (b) pseudo first (c) third (d) zero

97) The half-life period of a first order reaction is 69.3 seconds. Its rate constant is _____.
(a) 10^{-2} s^{-1} (b) 10^{-4} s^{-1} (c) 10 s^{-1} (d) 10^2 s^{-1}

98) If $[\text{A}]$ is the concentration of A at any time t and $[\text{A}_0]$ is the concentration at $t = 0$, then for the first order reaction, the rate equation can be written as _____.
(a) $k = \frac{2.303}{t} \log \left[\frac{\text{A}}{\text{A}_0} \right]$ (b) $k_t = 2.303 \log \left[\frac{\text{A}_0}{[\text{A}]} \right]$ (c) $k = 2.303 \log \left[\frac{\text{A}_0}{[\text{A}_0] - [\text{A}]} \right]$ (d) $k = \frac{2.303}{t} \log$

99) All radioactive transformations follow _____ order kinetics.
(a) zero (b) first (c) second (d) third

100) Activation energy of a reactant is reduced by _____.
(a) increased temperature (b) reduced temperature (c) increased pressure (d) reduced pressure

Match the following

101) Arrhenius (1) pseudo first order
102) $\text{N}_2\text{O}_{5(\text{g})} \rightarrow 2\text{NO}_{2(\text{g})} + \frac{1}{2} \text{O}_{2(\text{g})}$ (2) 1
103) $\text{CH}_3\text{COCH}_3 + \text{I}_2 \xrightarrow{\text{H}} \text{ICH}_2\text{COCH}_3 + \text{HI}$ (3) $k = Ae^{-\left(\frac{E_a}{RT}\right)}$
104) $2\text{HI} \rightarrow \text{H}_2\text{I}_2$ (4) 0
105) $\text{CH}_3\text{COOCH}_3(\text{aq}) + \text{H}_2\text{O} \xrightarrow{\text{H}} \text{CH}_3\text{COOH}(\text{aq}) + \text{CH}_3\text{OH}(\text{aq})$ (5) 2
106) Pseudo first order reaction (6) involves at least two reactants

Assertion and reason

107) Assertion: If temperature does not affect the rate of reaction, $E_a = 0$.

Reason: Lesser the activation energy, slower will be the reaction

- a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
c) Assertion is true but reason is false.
d) Both assertion and reason are false.

108) Assertion: A positive catalyst increases the rate of reaction.

Reason: A positive catalyst alters reaction mechanism and decreases activation energy.

- a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
c) Assertion is true but reason is false.
d) Both assertion and reason are false.

- 109) Assertion: Order and molecularity of a reaction are always equal.
Reason: Complex reactions takes place in steps and fastest step determine the molecularity of reaction
a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
c) Assertion is true but reason is false.
d) Both assertion and reason are false.
- 110) Assertion: Hydrolysis of ester in acidic medium follows pseudo first order kinetics.
Reason: Hydrolysis of ester is independent of the concentration of acid used
a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
c) Assertion is true but reason is false.
d) Both assertion and reason are false.
- 111) Assertion: In presence of positive catalyst, activation energy & threshold energy decreases.
Reason: Minimum energy required to permit a reaction is known as threshold energy.
a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
c) Assertion is true but reason is false.
d) Both assertion and reason are false.
- 112) Assertion: Rate constant determined from Arrhenius equation are fairly, accurate for simple as well as complex molecule.
Reason: Reactant molecules undergo chemical change irrespective of their orientation during collision.
a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
c) Assertion is true but reason is false.
d) Both assertion and reason are false.
- 113) Assertion: All collision of reactant molecule lead to product formation.
Reason: Product formation is independent of orientation of the reactant molecules.
a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
c) Assertion is true but reason is false.
d) Both assertion and reason are false.
- 114) Assertion: The enthalpy of reaction remains constant in the presence of a catalyst.
Reason: A catalyst participating in the reaction forms a different activated complex and lowers down the activation energy of reactant and product remains the same.
a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
c) Assertion is true but reason is false.
d) Both assertion and reason are false.
- 115) Assertion: Order and molecularity are same.
Reason: Order is determined experimentally and molecularity is the sum of the stoichiometric coefficient of rate determining elementary step.
a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
c) Assertion is true but reason is false.
d) Both assertion and reason are false.

116) Assertion: Order of the reaction zero or fractional.

Reason: We cannot determine order from balanced chemical equation.

- a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
- b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
- c) Assertion is true but reason is false.
- d) Both assertion and reason are false.

117) Assertion: Rate constant of zero order reaction has same units as the rate of reaction.

Reason: Rate constant of zero order reaction does not depend upon unit of concentration.

- a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
- b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
- c) Assertion is true but reason is false.
- d) Both assertion and reason are false.

118) Assertion: The order of reaction can have fractional value.

Reason: Order cannot be determined from a stoichiometrically balanced equation.

- a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
- b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
- c) Assertion is true but reason is false.
- d) Both assertion and reason are false.

119) Assertion: A catalyst is a substance which alters the rate of a reaction. Reason: In the presence of catalyst the energy of activation is increased.

- a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
- b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
- c) Assertion is true but reason is false.
- d) Both assertion and reason are false.

120) Assertion: Hydrolysis of methyl acetate by HCl is a pseudo first order reaction.

Reason: HCl is used as catalyst in the above reaction.

- a) Both assertion and reason are true and the reason is the correct explanation of the assertion.
- b) Both assertion and reason are true but the reason is not the correct explanation of the assertion.
- c) Assertion is true but reason is false.
- d) Both assertion and reason are false.

Find the wrong statement

121) Mark the incorrect statements.

- a) Catalyst provides an alternative pathway, to reaction mechanism.
- b) Catalyst raises the activation energy.
- c) Catalyst lowers the activation energy.
- d) Catalyst not alters enthalpy change of the reaction.

122) Consider the following statements and identify the incorrect statement(s).

- (i) Decomposition of H_2O_2 is an II order reaction
 - (ii) $t_{1/2}$ is independent of initial concentration of a reaction.
 - (iii) Fractional order reactions are observed depending on their rates.
 - (iv) $\text{Rate} = k [\text{A}]^p [\text{B}]^q$; $p + q = \text{order}$.
- a) only (ii)
 - b) both (ii) and (iii)
 - c) only (i)
 - d) None of these

123) a) The rate of a reaction increase with increasing temperature.

- b) The rate of a reaction increases with the increase in the concentration of the reactants
- c) Gas phase reactions are slower than reactions involving solid reactants
- d) The rate of a reaction is affected by the surface area of the reactant.

124) About "Collision theory"

- a) Collision theory used to predict the rates of chemical reactions for gases.



- b) This theory is based on the kinetic theory of gases
 c) Chemical reactions occur as a result of collision between the reacting molecules
 d) Decrease in concentration of the reactant brings about more collisions.
- 125) a) Rate of a reaction depends on the initial concentration of reactants
 b) Rate constant of a reaction does not depend on the initial concentration of reactants
 c) Concentration and surface area decreases the number of collision
 d) The molecularity of a reaction is the no. of molecules or ions that participate in the rate determining step.

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