SOLVED QUESTION BANK

Objective / Very Short Answer Type Questions [1 Mark]

- 1. The half life period of first order reaction is 1386 seconds. The specific rate constant of the reaction is
 - (a) $0.5 \times 10^{-2} \text{ s}^{-1}$
- (6) $0.5 \times 10^{-3} \, s^{-1}$
- (c) $5.0 \times 10^{-2} \text{ s}^{-1}$
- (4) 5.0 × 10⁻³ s⁻¹
- $I_{1/2} = \frac{0.693}{}$ Ann. (b) $k = \frac{0.693}{1386} = 0.5 \times 10^{-3} s^{-1}$
 - 2. The rate constant of a reaction A --- B is 0.6 × 10⁻³ mole per second. If the concentration of [A] is 5 M, then what will be concentration of [B] after 20 minutes?
 - (a) 0.36 M

(A) 0.72 NO

(c) 1.08 M

- 6d) 3,60 M
- $k = \frac{[R]_0 [R]}{r}$ for zero order. Ams. (b) \Rightarrow 0.6 × 10⁻³ × 20 × 60 - [R]₀ - [R] == 0.72 M Amount of [B] produced = $[R]_0 - [R] = 0.72 \text{ M}$
 - 3. Which of the following is affected by catalyst?
 - (a) AH

(b) AS

(c) AG

- (4) E.
- Ans. (d) Catalyst affects activation energy. It does not affect. AH, AS, AG.
 - 4. A first order reaction has specific reaction rate 10-2s-1. How much time it will take for 20g of reactant to reduce to 5g?
 - (a) 138.6 s

(b) 346.5 s

(c) 693.0 s

- (d) 238.6 s
- Ans. (a) $I_{3/4} = \frac{1.386}{1.002} = 138.6 \text{ s}$
 - 5. The activation energy of a reaction can be determined from the slope of which of the following graph:
 - (a) lnk Vs -

(b) $\frac{T}{\log k} Vs \frac{1}{T}$

(c) lnk Vs T

- (a) $\frac{\ln k}{\pi}$ Vs T
- Ans. (a) When we plot lnk Vs $\frac{1}{T}$, slope of line = $-\frac{E_d}{R}$.
 - Mechanism of a hypothetical reaction X₂+Y₂→→2 XY is given below:
 - (a) X2 = X + X (fast)
 - (b) $X + Y_2 \longrightarrow XY + Y$ (slow)
 - (c) X + Y → XY (fast)

The overall order of reaction is

- (a) 2
 - (6) 0

- Ans. (c) rate = $k[X][Y_2]$. $K = \frac{[X]^2}{X_2} \implies [X] = k^{12}|_{X_2}$ $\mathsf{rate} = k \times K^{1/2} \{ \mathbf{X}_2 \}^{1/2} \, \{ \mathbf{Y}_2 \} = k^1 \, \{ \mathbf{X}_2 \}^{1/2} \, \{ \mathbf{Y}_3 \}$ Order = $\frac{1}{3} + 1 = \frac{3}{3} = 1.5$
 - 7. The rate of first order reaction is 0.04 $\max_{\mathbf{L}^{-1} \in \mathbb{R}^{d}}$ 10 sec. and 0.03 mol L. at 20 seconds after initials, of the reaction. $I_{1/2}$ of reaction is

$$\lceil \log 4 - \log 3 = 0.12n \rceil$$

- (a) 44.1 s (b) 54.1 s (c) 24.1 s (d) 34.1.
- Ans. (c) $k = \frac{2.303}{20-10} \log \frac{0.04}{6.03} = \frac{2.303}{10} \log \frac{4}{3}$ $=\frac{2.303}{10}(\log 4 - \log 3)$

$$I_{1/2} = \frac{2.303}{10} (0.6021 - 0.4771) = \frac{2.303}{10} \times 0.136$$

$$I_{1/2} = \frac{0.693}{k} = \frac{0.693 \times 10}{2.303 \times 0.1249} = \frac{3.010}{0.1249} = 24.16$$

- 8. If the initial concentration of reactant is doubled $t_{1/2}$ is also doubled, the order of reaction is (a) zero (A) I
- Ans. (a) In zero order of reaction $t_{1/2} \propto [R]_0$
 - For a reaction taking place in three steps, the event rate constant $k = \frac{k_1 k_2}{k_2}$. If E_{a_1} , E_{a_2} and E_{a_1} are \pm
 - 40, 50 and 60 kJ mol-1, the overall E, is | HOIS (a) 30 (c) 60(d).50
- k Ac-EJRT Ans. (a)

$$k = \frac{k_1 k_2}{k_3}$$

$$\Rightarrow Ae^{-E_{g}/RT} = \frac{Ae^{-E_{g}/RT} \times Ae^{-E_{g}/RT}}{Ae^{-E_{g}/RT}}$$

$$e^{-E_{0}/RT} = \frac{Ae^{-E_{0}/RT}}{e^{-E_{0}/RT} \times e^{-E_{0}/RT}}$$

Taking In on both side we get

(4) 2

(a) I

$$\frac{E_{\sigma}}{RT} = \frac{-E_{\sigma_1} - E_{\sigma_2} + E_{\sigma_3}}{RT}$$

$$E_{\sigma} = E_{\sigma_1} + E_{\sigma_2} - E_{\sigma_3}$$

$$E_{\sigma} = 40 + 50 - 60 = 30 \text{ kJ mol}^{-1}$$

(c) 3

10. If conc. of reactant 'A' is increased 10 times and rate of reaction becomes 100 times. What is order with respect to 'A'? (40) 4

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Peter R. Bergethon, Kevin Hallock

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Table of Contents Chemical Kinetics Questions And Answers

- 1. Understanding the eBook Chemical Kinetics Questions And Answers
 - The Rise of Digital Reading Chemical Kinetics Questions And Answers
 - Advantages of eBooks Over Traditional Books
- 2. Identifying Chemical Kinetics Questions And Answers
 - Exploring Different Genres
 - o Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
- 3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Chemical Kinetics Questions And Answers
 - User-Friendly Interface
- 4. Exploring eBook Recommendations from Chemical Kinetics Questions And Answers
 - Personalized Recommendations
 - Chemical Kinetics Questions And Answers User Reviews and Ratings
 - Chemical Kinetics Questions And Answers and Bestseller Lists

- 5. Accessing Chemical Kinetics Questions And Answers Free and Paid eBooks
 - Chemical Kinetics Questions And Answers Public Domain eBooks
 - Chemical Kinetics Questions And Answers eBook Subscription Services
 - Chemical Kinetics Questions And Answers Budget-Friendly Options
- 6. Navigating Chemical Kinetics Questions And Answers eBook Formats
 - ∘ ePub, PDF, MOBI, and More
 - Chemical Kinetics Questions And Answers Compatibility with Devices
 - Chemical Kinetics Ouestions And Answers Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Chemical Kinetics Questions And Answers
 - Highlighting and Note-Taking Chemical Kinetics Questions And Answers
 - Interactive Elements Chemical Kinetics Questions And Answers
- 8. Staying Engaged with Chemical Kinetics Questions And Answers
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Chemical Kinetics Questions And Answers
- 9. Balancing eBooks and Physical Books Chemical Kinetics Questions And Answers
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Chemical Kinetics Questions And Answers
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Chemical Kinetics Questions And Answers
 - Setting Reading Goals Chemical Kinetics Questions And Answers
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Chemical Kinetics Questions And Answers
 - Fact-Checking eBook Content of Chemical Kinetics Questions And Answers
 - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning

- Utilizing eBooks for Skill Development
- Exploring Educational eBooks
- 14. Embracing eBook Trends
 - Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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