

CAREER ENDEAVOUR ACADEMY PRIVATE LIMITEE Best Institute for NET-JRF, GATE & IIT-JAM Exams

CHEMICAL SCIENCES (NET-JRF/GATE)

Unit Test: Chemical Kinetics

Time : 00: 50 Hour

Date : 06-02-2014 M.M. : 60

Instructions:

- 1. Question Paper contains two parts: Part-A and Part-B. Part-A contains 10 objective type questions, each question carry 2 marks. Part-B contains 10 objective type questions, each question carry 4 marks.
- 2. There is nagative marking, 0.5 mark will be deducted for each wrong answer in Part-A and 1 marks for Part-B.
- 3. Attempt all the questions, use of calculator is not allowed.

| PART-A | | | | | | | | | |
|--------|---|----------|-----------------------|-------------------|---------|---|--|--|--|
| | | | | | | | | | |
| 1. | The concentration of R in the reaction $R \rightarrow P$ was measured as a function of time and the following da | | | | | | function of time and the following data is | | |
| | obtained | | | | | | | | |
| | [R] (molar) | 1.0 | 0.75 (| 0.40 | 0.10 | | | | |
| | t (min) (| 0.0 | 0.05 (| 0.12 | 0.18 | | | | |
| | The order of reaction is: | | | | | | | | |
| | (a) Zero | | (b) One | | | (c) Two | (d) Three | | |
| | | | | | | 1 | | | |
| 2. | For the reaction | n aA — | \longrightarrow Pro | ducts t | he plot | of $\overline{[A]^2}$ versus tin | ne gives a straight line. The order of the | | |
| | reaction is: | | | | | | | | |
| | (a) 0 | | (b) 1 = 1 | | | (c) | (d) 3 | | |
| | (a) 0 | | | RE | ER | EDEAV | | | |
| 3. | For a reaction w | ith rate | equation | l | | | | | |
| | dC dC | | | | | | | | |
| | $-\frac{1}{dt} = kC^2$ | | | | | | | | |
| | C_a and C are the concentration at time t = 0 and 't' respectively. If 10 min were required for C _a to become C _a / | | | | | | | | |
| | 2, the time required for C_0 to become $C_0/4$ is: | | | | | | | | |
| | (a) 10 min | | (b) 20 m | in | | (c) 30 min | (d) 40 min | | |
| 4. | The molar extind | ction c | oefficient | oface | mplexi | s 12000 L mol ⁻¹ cm ⁻ | ¹ and the minimum detectable absorbance | | |
| | is 0.01. Calculate the minimum concentration of the complex that can be detected in a Lambert-Beer law cell | | | | | | | | |
| | of path length 1. | 00 cm. | | | | - | | | |
| | (a) 8 33×10 ⁻⁷ M | 1 | (b) 1 20 | ×10 ⁻⁵ | М | (c) 1 20×10 ⁻⁴ M | (d) 1 20×10 ⁻³ M | | |
| | (u) 0.55/(10 10 | 1 | (0) 1.20 | | | (0) 1.20/10 101 | (a) 1.20/(10/10) | | |
| 5. | For a reaction $2A \rightarrow$ Product the initial concentration of A is 0.1 M and the rate consta | | | | | | A is 0.1 M and the rate constant is | | |
| | $2 \times 10^{-3} \text{ dm}^3 \text{mol}^{-1} \text{s}^{-1}$. The half life period of the reaction is: | | | | | | | | |
| | (a) 5 sec | | (b) 50 se | ec | | (c) 500 sec | (d) 5000 sec | | |
| | (, | | (3) 2 3 5 | | | (-) 0 0 0 000 | | | |



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The decomposition of ethane follows the rate law $-\frac{d}{dt} [CH_3CH_3] = k [CH_3CH_3]$ with $k = \left(\frac{k_1k_3k_4}{k_5}\right)^{\frac{1}{2}}$. 6.

Where k_1, k_3, k_4 and k_5 are the rate constants of various elementary steps.

The values of E_as of elemantary reactions were found to be

$$E_{a_1} = 350 \text{ kJ mole}^{-1}$$

 $E_{a_3} = 160 \text{ kJ mole}^{-1}$
 $E_{a_4} = 30 \text{ kJ mole}^{-1}$

and

 $E_{a_5} = 10 \text{ kJ mole}^{-1}$

t-O

The activation energy of the overall reaction is (b) 265 kJ mole⁻¹ (a) 165 kJ mole^{-1} (c) 275 kJ mole^{-1} (d) 375 kJ mole^{-1}

7. Consider the elementary reaction $A + A + A \longrightarrow$ Product

(a)
$$\frac{15}{2a^2k}$$
 (b) $\frac{15}{6a^2k}$ (c) $\frac{13}{6a^2k}$ (d) $\frac{7}{18a^2k}$

The reaction $2H_2 + 2NO \rightarrow N_2 + 2H_2O$ is assumed to proceed by the following mechanism: 8.

$$2NO \Longrightarrow N_2O_2 \qquad \text{fast}$$

$$N_2O_2 + H_2 \Longrightarrow N_2O + H_2O \qquad \text{slow}$$

$$N_2O + H_2 \Longrightarrow N_2 + H_2O \qquad \text{fast}$$

which is the rate law for this reaction?

| (a) Rate = K[NO][H ₂] (b) Rate = K[NO] ² [H ₂] (c) Rate = K[NO][H ₂] ² (d) Rate = K[NO] ² [H ₂] |
|--|
|--|

Column-II

| 9. Column-I | |
|-------------|--|
|-------------|--|

| (i) Collisional quenching | | | | | (P) $S_1 \longrightarrow S_0 + hv$ | | |
|--|----------|----------|-----|-------|--|--|--|
| (ii) Fh | lorescei | nce | | | (Q) $T_1 \longrightarrow S_0 + hv'$ | | |
| (iii) Photosensitization | | | | | (R) $S^* + M \longrightarrow S + M$ + heat | | |
| (iv) Pl | nosphor | rescence | | | (S) $S^* + A_2(g) \longrightarrow S + 2A(g)$ | | |
| Correct match for (P), (Q), (R), (S) in column (I) is: | | | | | | | |
| | (P) | (Q) | (R) | (S) | | | |
| (a) | (ii) | (iv) | (i) | (iii) | | | |

| (a) | (ii) | (iv) | (i) | (iii) |
|-----|------|------|-------|-------|
| (b) | (ii) | (iv) | (iii) | (i) |
| (c) | (iv) | (ii) | (i) | (iii) |
| (d) | (iv) | (ii) | (iii) | (i) |
| | | | | |



 $A + B \xrightarrow{k}$ Products (k = rate constant for the reaction). Exp. NO. $[A] (mol L^{-1})$ $[B] (mol L^{-1})$ Initial rate (mol $L^{-1} s^{-1}$) 1×10⁻³. 1. 0.024 0.035 2. 0.070 8×10⁻³. 0.012 3. 0.024 0.070 8×10⁻³. (a) $-\frac{d[A]}{dt} = k[A][B]^2$ (b) $-\frac{d[A]}{dt} = k[A]^2[B]$ (c) $-\frac{d[A]}{dt} = k[A][B]^3$ (d) $-\frac{d[A]}{dt} = k[B]^3$ Part-B The rate constant of decomposition of a compound is represented by

The value of
$$E_A$$
 (in Kcal mol⁻¹) for this compound at 300 K is:
(a) 24 (b) 12 (c) 240 (d) 120

 $lnK = 5.0 - \frac{12000}{T}$

$$Br_{2} + M \xleftarrow{k_{1}}{k_{-1}} 2Br + M (fast), \left(K = \frac{k_{1}}{k_{-1}}\right)$$

$$Br + H_{2} \xrightarrow{k_{2}} HBr + H (slow)$$

$$H + Br_{2} \xrightarrow{k_{3}} HBr + Br (fast)$$
(a) Rate = $k_{2} (K)^{\frac{1}{2}} [H_{2}] [Br_{2}]^{\frac{1}{2}}$
(b) Rate = $k_{2} [H_{2}] [Br_{2}]^{\frac{1}{2}}$
(c) Rate = $k_{2} k_{1}^{\frac{1}{2}} [H_{2}] [Br_{2}]^{\frac{1}{2}}$
(d) Rate = $k_{2} [H_{2}] [Br_{2}]^{\frac{1}{2}}$

13. For the reaction $2A + B \rightleftharpoons X^{\#} \longrightarrow P$, $E_a = 25.0 \text{ kJ mol}^{-1}$ at 300 K. The enthalpy change for the formation of activated complex from the reaction in kJ/ol is: (a) 20 kJ/mole (b) 17.5 kJ/mole (c) 27.5 kJ/mole (d) 15.0 kJ/mole

14.
$$\begin{array}{c} k_1 \\ A(g) \\ 3k_1 \\ C(g) \end{array}$$

11.

At t = 0, pressure due to A only = 8 atm.

Partial pressure due to B(g), C(g), C(g) and total pressure at the end of reaction is:

(a)
$$P_B = 2$$
 atm, $P_C = 6$ atm, $P_{\text{total}} = 8$ atm (b) $P_B = 6$ atm, $P_C = 2$ atm, $P_{\text{total}} = 8$ atm

(c)
$$P_B = 2 \text{ atm}, P_C = 6 \text{ atm}, P_{\text{total}} = 32 \text{ atm}$$
 (d) $P_B = \frac{8}{3} \text{ atm}, P_C = 2 \text{ atm}, P_{\text{total}} = \frac{14}{3} \text{ atm}$

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| 15. | For the sequential read | etion. A $\frac{k_1 = (0.602) \sec^{-1}}{k_1 = (0.602) \sec^{-1}}$ | $\rightarrow B \xrightarrow{k_2 = (0.301) \text{sec}^{-1}} C$ | - | | | |
|-----|--|--|---|---|--|--|--|
| | Time when concentration of 'B' will reach maximum is: | | | | | | |
| | (a) 2303 sec | (b) 1 sec | (c) 4 606 sec | (d) 2 sec | | | |
| 16 | When light is incident | on any chemical substa | nce then 20% of light is | absorbed and rest is transmitted | | | |
| 10. | So transmittance and absorbance values are respectively. | | | | | | |
| | | | | | | | |
| | (a) 80%, $\log \frac{8}{10}$ | (b) 80%, $\log \frac{10}{8}$ | (c) 20%, log 5 | (d) 20%, log 2. | | | |
| 17. | Consider the following | reactions: | | | | | |
| | (i) $A(g) \longrightarrow \operatorname{Produ}$ | cts, | (Unimolecular reaction following Lindemann mechanism) | | | | |
| | (ii) $A(g) \xrightarrow{M(s)} Production$ | oducts | (Unimolecular hetrogeneous catalysis reation) | | | | |
| | Choose the correct sta Low Pressure | tements. | High Pressure | | | | |
| | (a) Reaction (i) – Ist o | rder | Reaction (i) – 2nd or | der | | | |
| | Reaction (ii) $-2nd$ | lorder | Reaction (ii) – Ist ord | ler | | | |
| | (b) Reaction (i) $-2nd$ Reaction (ii) $-1st$ | order | Reaction (i) -1 st order Reaction (ii) -2 nd order | | | | |
| | (c) Both reaction (i) at | nd (ii) – 2nd order | Both reaction (i) and | (ii) – 1st order | | | |
| | (d) Reaction (i) $-2nd$ | order | Reaction (i) -1 st ord | ler | | | |
| | Reaction (ii) – 1st | order | Reaction (ii) – zeroth | order | | | |
| 18. | For the reaction A | \rightarrow B, the rate constant | at $k_1 = A_1 e^{-E_{a_1}/RT}$ and | for the reaction $k_2 = A_2 e^{-E_{a_2}/RT}$. If | | | |
| | $A_1 = 10^8, A_2 = 10^{10}, A_3$ 2 cal/k mol) | $E_{a_1} = 600 \text{ cal / mol, } E_a$ | $_{2} = 1800 \text{ cal} / \text{m}$, then t | he temperature 'a' which $k_1 = k_2 (R =$ | | | |
| | (a) 1200 K | (b) 1200×4.606 | (c) $\frac{1200}{4.606}$ K | (d) $\frac{600}{4.606}$ K | | | |
| 19. | 19. The following mechanism has been suggested for thermal decomposition of NO | | | | | | |
| | Overall reaction: $2NO_2 = 2NO + O_2$ | | | | | | |
| | Mechanism: | $NO_2 + NO_2 \xrightarrow{k_1} NO_2$ | $NO + NO + O_2; NO_2 +$ | $NO_2 \xrightarrow{k_2} NO_3 + NO_3$ | | | |
| | | $NO_3 + NO \xrightarrow{k_3} 2N$ | $NO_2; NO_3 + NO_2 - \frac{k_4}{k_4}$ | \rightarrow NO + O ₂ + NO ₂ | | | |
| | The above mechanism leads to the following rate law: $-\frac{d[NO_2]}{dt} = 2 k_d [NO_2]^2$. The value of $k_d = ?$ | | | | | | |
| | (a) $k_1 + \frac{k_2 k_4 [No]}{k_1 [No] + k_2}$ | 0] [NO.] | (b) $k_1 + \frac{k_2 k_4 [NO_2]}{k [NO] + k [NO]}$ | | | | |
| | | L-·~2] | ~3[110] + K | 4 L ^{- · · ·} 2 J | | | |
| | (c) $1 + \frac{k_2 k_4 [NO_2]}{k_3 [NO] + k_4 [2]}$ | $\frac{1}{NO_2}$ | (d) None of these | | | | |
| | | | | | | | |
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20. The incorrect statement is/are

(I) For KCl $a = \sqrt{(\gamma \pm m)^4}$ (II) Ionic strength of 0.25 molal K₂SO₄ solution is 0.50. (III) Salt effect is +ve, for the reaction $A^{2+} + B^+ \longrightarrow P$

(IV) For $2A^+ + B^- + C^{--} \longrightarrow P$ $\log\left(\frac{k}{k_0}\right) = +6A\sqrt{I}$ (a) II, III (b) IV only

(c) III and IV

(d) None of these





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