



# SHIKSHA CLASSES

Subject : Chemistry

BOARD ANSWER PAPER

Total Marks : 20

Class : XII

Topic: 6) Chemical Kinetics

## Section (A)

**Q.1 : a) Select and write the most appropriate answer from the following alternatives of each sub question. (05)**

i) The rate constant of reaction .....

Ans : a) decreases with increasing  $E_a$

ii) **The slope of straight line obtained by plotting rate versus conc. of reactant for a first order reaction is**

Ans : d) K

iii) **The half life of the first order reaction having rate constant  $K = 1.7 \times 10^{-5} \text{ s}^{-1}$  is**

Ans : c) 11.1 hour

iv) **The rate constant of zero order reaction has the unit**

Ans : b)  $\text{mol L}^{-1} \text{ s}^{-1}$

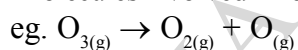
v) **The order of the reaction for which the units of rate constant are  $\text{mol dm}^{-3} \text{ s}^{-1}$  is**

Ans : c) 0

**Q.1 : (b) Very short answer type Question. [2]**

**1) What is molecularity of the reaction ?**

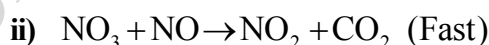
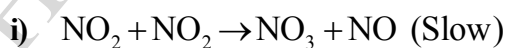
Ans : The molecularity refers to How many reactant molecules involved in reaction



In above reaction there is only one reactant molecule. These are unimolecular reaction or molecularity is one.

**2) What is rate law for a reaction.**

$\text{NO}_{2(\text{g})} + \text{CO}_{(\text{g})} \rightarrow \text{NO}_{(\text{g})} + \text{CO}_{2(\text{g})}$  **If the reaction occurs in the following steps?**



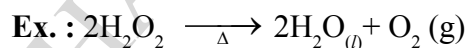
Ans : The rate law for reaction is  $\text{rate} = -K[\text{NO}_2]^2$   
 $\text{NO}_3$  is reaction intermediate.

## Section (B)

**Q.2 : Attempt Any Three of the following question. (06)**

**1) Define rate law. Explain with one example.**

Ans : Rate law is defined as an experimentally determined equation that expresses the rate of a chemical reaction in terms of molar concentrations of the reactants.



The rate law for reaction is

$$\text{rate} = -\frac{1}{2} \frac{d[\text{H}_2\text{O}_2]}{dt} = K[\text{H}_2\text{O}_2]$$

The reaction is first order reaction

**2) Derive the expression for half life of zero order reaction.**

Ans : The integrated rate law for zero order

$$\text{reaction is } K = \frac{[\text{A}]_0 - [\text{A}]_t}{t}$$

where  $[\text{A}]_t$  is concentration of A that remains unreacted at time t.

$$\text{at } t = t_{1/2}, [\text{A}]_t = \frac{[\text{A}]_0}{2}$$

$$K = \frac{[\text{A}]_0 - [\text{A}]_0/2}{t_{1/2}} \quad K = \frac{[\text{A}]_0}{2t_{1/2}} \quad \text{i.e. } t_{1/2} =$$

$$\frac{[\text{A}]_0}{2K}$$

**3) Write Arrhenius equation and explain the terms involved.**

Ans : The Arrhenius equation is -

$$\ln k = \ln A - \frac{E_a}{RT}$$

$$\text{or } \log_{10} K = \log_{10} A - \frac{E_a}{2.303RT}$$

when  $K$  = rate constant

$A$  = frequency factor or pre exponential factor

$E_a$  = activation energy

$T$  = temperature

**4) What is zero order reaction? Derive integrated rate law for zero order reaction.**

**Ans :** The rate of zero order reaction is independent of the reactant concentration

Integrated rate law for zero order reaction  $A \rightarrow P$

The differential law is give by

$$\text{rate} = -\frac{d[A]}{dt} = K[A]^0 = K \quad \dots\dots (i)$$

By rearrangement above equ<sup>n</sup> given  $d[A] = -K dt$

Integration between the limits

$[A] = [A]_0$  at  $t = 0$  and  $[A] = [A]_t$  at time  $t$

$$\int_{[A]_0}^{[A]_t} d[A] = -K \int_0^t dt$$

$$[A]_t - [A]_0 = -Kt$$

$$\text{Hence } Kt = [A]_0 - [A]_t$$

**Section (C)**

**Q.3 : Answer the following question (any one). (03)**

**1) In a first order reaction  $A \rightarrow B$ , 60% of the given sample of compound decomposes in 45 minutes. What is half life of reaction?**

**Ans :** Given,

$$[A]_0 = 100\%$$

$$\chi = 60\%$$

$$\therefore [A]_t = [A]_0 - \chi = 100 - 60 = 40\%$$

$$t = 45 \text{ min}$$

Formula :-

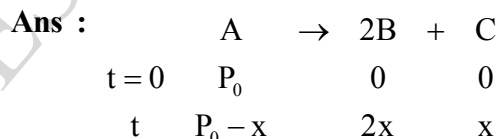
$$\begin{aligned} K &= \frac{2.303}{t} \log_{10} \frac{[A]_0}{[A]_t} \theta \\ &= \frac{2.303}{45} \times \log_{10} \frac{100}{40} \\ &= \frac{2.303}{45} \times \log_{10} 2.5 \\ &= \frac{2.303}{45} \times 0.3979 \\ &= 0.0204 \text{ min}^{-1} \end{aligned}$$

$$\text{Half life} = t_{1/2} = \frac{0.693}{K}$$

$$= \frac{0.693}{0.0204}$$

$$= 34 \text{ min}$$

**ii) For the first order reaction  $A_{(g)} \rightarrow 2B_{(g)} + C_{(g)}$  the initial pressure of A is 90 mm Hg and pressure after 10 min is found to be 180 mm Hg. Find rate constant of reaction.**



$$P_T = P_0 + 2x$$

$$x = \frac{P_T - P_0}{2}$$

$$x = \frac{180 - 90}{2} = 45$$

Pressure of A at time 10 min

$$P_0 - x = 90 - 45 = 45 \text{ mm Hg}$$

$$K = \frac{2.303}{t} \log \frac{[P_0]}{[P_t]}$$

$$K = \frac{2.303}{10} \log \frac{90}{45}$$

$$= 0.2303 \times 0.3010$$

$$= 0.0693 \text{ min}^{-1}$$

**Section (D)**

**Q.4 : Answer the following question. (Any one) (04)**

**1) a) Derive integrated rate law for first order reaction.**

**Ans :** Consider a general first order reaction is.

$A \rightarrow \text{products}$

The differential rate law is  $-\frac{d[A]}{dt} = K[A]$

where  $[A] = [A]_0$  at time  $t = 0$  and

$[A] = [A]_t$  at time  $t = t$

Integrating above equation in limits.

$$\int_{[A]_0}^{[A]_t} \frac{d[A]}{A} = -K \int_0^t dt$$

$$\{\ell n [A]\}_{[A]_0}^{[A]_t} = -K(t)_0^t$$

$$\ell n [A]_t - \ell n [A]_0 = -Kt$$

$$\ell n \frac{[A]_t}{[A]_0} = -Kt$$

$$Kt = \ell n \frac{[A]_0}{[A]_t}$$

$$K = \frac{1}{t} \ell n \frac{[A]_0}{[A]_t}$$

$$K = \frac{1}{t} \ell n \frac{[A]_0}{[A]_t}$$

converting  $\ell n$  to  $\log_{10}$  the integrated rate law becomes.

$$K = \frac{2.303}{t} \log_{10} \frac{[A]_0}{[A]_t}$$

**b) Consider the reaction  $2N_2O_{5(g)} \rightarrow 4NO_{2(g)} + O_{2(g)}$  in liquid bromine. At a particular moment during the reaction  $N_2O_5$  disappears at a rate  $0.02 \text{ M/s}$ . At what rates  $NO_2$  and  $O_2$  are formed? what is rate of reaction.**

**Ans. :**  $-\frac{d[N_2O_5]}{dt} = 0.02 \text{ m/s}$ .

The reaction shows that the rate of formation

of  $NO_2$  is twice the rate of consumption of  $N_2O_5$ .

$$\therefore \frac{d[NO_2]}{dt} = -2 \frac{d[N_2O_5]}{dt}$$

$$= -2 \times (-0.02 \text{ M/s}) = 0.04 \text{ M/s}$$

The rate of consumption of  $N_2O_5$  is twice at the rate of formation of  $O_2$

$$-\frac{d[N_2O_5]}{dt} = 2 \frac{d[O_2]}{dt} \text{ or}$$

$$\frac{d[O_2]}{dt} = -\frac{1}{2} \frac{d[N_2O_5]}{dt}$$

$$\text{Hence } \frac{d[O_2]}{dt} = -\frac{1}{2} \times (-0.02 \text{ M/S})$$

$$= 0.01 \text{ M/s}$$

Rate of reaction =

$$= -\frac{1}{2} \frac{d[N_2O_5]}{dt} = \frac{1}{4} \frac{d[NO_2]}{dt} = \frac{d[O_2]}{dt}$$

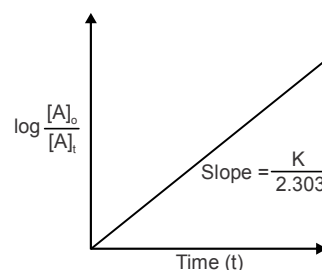
$$= -\frac{1}{2} \times (-0.02 \text{ M/s})$$

$$= -0.01 \text{ M/s}$$

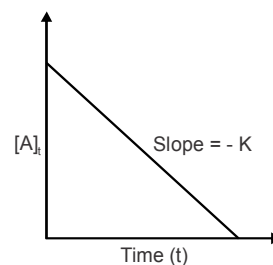
**OR**

**2) a) Draw the following graphs**

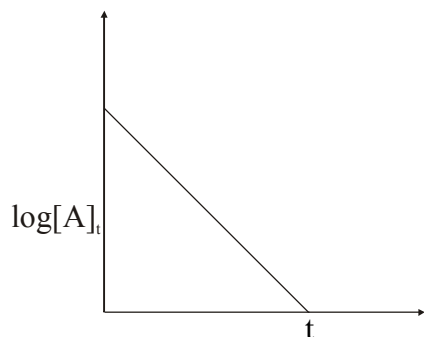
**Ans :** i)  $\log \frac{[A]_0}{[A]_t}$  vs time(t) for first order reaction



ii)  $[A]_t$  vs  $t$  for zero order reaction.



iii)  $\log[A]_t$  vs t for first order



b) The rate of the reaction,  $A + B \rightarrow P$  is  $3.6 \times 10^{-2} \text{ mol dm}^{-3} \text{ s}^{-1}$  where

$[A] = 0.2 \text{ moles dm}^{-3}$  and  $[B] = 0.1 \text{ moles dm}^{-3}$ .

Calculate the rate constant if the reaction is first order in A and second order in B.

Ans : The reaction is first order in 'A' and second order in 'B'. Hence the rate law gives  
rate =  $K [A][B]^2$

$$\text{or } K = \frac{\text{rate}}{[A][B]^2}$$

$$\text{rate} = 3.6 \times 10^{-2} \text{ Ms}^{-1}$$

$[A] = 0.2 \text{ mol dm}^{-3}$  and  $[B] = 0.1 \text{ mol dm}^{-3}$   
all value substituting gives

$$K = \frac{3.6 \times 10^{-2} \text{ mol dm}^{-3} \text{ s}^{-1}}{0.2 \text{ mol dm}^{-3} \times (0.1 \text{ mol dm}^{-3})^2}$$
$$= \frac{3.6}{0.2} = 18$$

$$K = 18 \text{ mol}^{-2} \text{ dm}^6 \text{ s}^{-1}$$

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