# (Chapter 4)(Chemical Kinetics) XII

## **Intext Questions**

#### Question 4.1:

For the reaction  $R \rightarrow P$ , the concentration of a reactant changes from 0.03 M to 0.02 M in 25 minutes. Calculate the average rate of reaction using units of time both in minutes and seconds.

Answer

$$=-\frac{\Delta[R]}{\Delta t}$$

$$=-\frac{\left[\mathbf{R}\right]_{2}-\left[\mathbf{R}\right]_{1}}{t_{2}-t_{1}}$$

Average rate of reaction
$$= -\frac{\Delta[R]}{\Delta t}$$

$$= -\frac{[R]_2 - [R]_1}{t_2 - t_1}$$

$$= -\frac{0.02 - 0.03}{25} \text{ M min}^{-1}$$

$$= -\frac{-0.01}{25} \text{ M min}^{-1}$$

$$= 4 \times 10^{-4} \text{ M min}^{-1}$$

$$= \frac{4 \times 10^{-4}}{60} \text{ M s}^{-1}$$

$$= 6.67 \times 10^{-6} \text{ M s}^{-1}$$

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#### Question 4.2:

In a reaction,  $2A \rightarrow Products$ , the concentration of A decreases from 0.5 mol  $L^{-1}$  to 0.4  $mol \ L^{-1}$  in 10 minutes. Calculate the rate during this interval?

Answer

$$\text{Average rate} = -\frac{1}{2} \frac{\Delta \left[ \mathbf{A} \right]}{\Delta t}$$

$$= -\frac{1}{2} \frac{\left[A\right]_2 - \left[A\right]_1}{t_2 - t_1}$$

$$= -\frac{1}{2} \frac{0.4 - 0.5}{10}$$

$$= -\frac{1}{2} \frac{-0.1}{10}$$

 $= 0.005 \text{ mol } L^{-1} \min^{-1}$ 

 $= 5 \times 10^{-3} \text{ M min}^{-1}$ 

#### Question 4.3:

For a reaction, A + B  $\rightarrow$  Product; the rate law is given by,  $r = k[A]^{1/2}[B]^2$ . What is the order of the reaction?

The order of the reaction  $= \frac{1}{2} + 2$   $= 2\frac{1}{2}$ 

$$=2\frac{1}{2}$$

= 2.5

#### Question 4.4:

The conversion of molecules X to Y follows second order kinetics. If concentration of X is increased to three times how will it affect the rate of formation of Y?

Answer

The reaction  $X \rightarrow Y$  follows second order kinetics.

Therefore, the rate equation for this reaction will be:

 $Rate = k[X]^2 (1)$ 

Let  $[X] = a \mod L^{-1}$ , then equation (1) can be written as:

Rate<sub>1</sub> =  $k . (a)^2$ 

$$= ka^2$$

If the concentration of X is increased to three times, then  $[X] = 3a \text{ mol } L^{-1}$ 

Now, the rate equation will be:

Rate =  $k (3a)^2$ 

 $= 9(ka^2)$ 

Hence, the rate of formation will increase by 9 times.

#### Question 4.5:

A first order reaction has a rate constant  $1.15 \ 10^{-3} \ s^{-1}$ . How long will 5 g of this reactant take to reduce to 3 g?

Answer

From the question, we can write down the following information:

Initial amount = 5 g

Final concentration = 3 q

Rate constant =  $1.15 \ 10^{-3} \ s^{-1}$ 

We know that for a 1st order reaction,

$$t = \frac{2.303}{k} \log \frac{[R]_0}{[R]}$$

$$=\frac{2.303}{1.15\times10^{-3}}\log\frac{5}{3}$$

$$=\frac{2.303}{1.15\times10^{-3}}\times0.2219$$

= 444.38 s

= 444 s (approx)

#### Question 4.6:

Time required to decompose  $SO_2Cl_2$  to half of its initial amount is 60 minutes. If the decomposition is a first order reaction, calculate the rate constant of the reaction.

Answer

We know that for a 1st order reaction,

$$t_{1/2} = \frac{0.693}{k}$$

It is given that  $t_{1/2} = 60$  min

$$\therefore k = \frac{0.693}{t_{1/2}}$$

$$=\frac{0.693}{60}$$

Or 
$$k = 1.925 \times 10^{-4} \,\mathrm{s}^{-1}$$

### Question 4.7:

What will be the effect of temperature on rate constant?

Answer

The rate constant of a reaction is nearly doubled with a 10° rise in temperature. However, the exact dependence of the rate of a chemical reaction on temperature is given by Arrhenius equation,

$$k = Ae^{-Ea/RT}$$

Where,

A is the Arrhenius factor or the frequency factor

*T* is the temperature

R is the gas constant

 $E_a$  is the activation energy

Question 4.8:

The rate of the chemical reaction doubles for an increase of 10 K in absolute temperature from 298 K. Calculate  $E_a$ .

Answer

It is given that  $T_1 = 298 \text{ K}$ 

$$..T_2 = (298 + 10) \text{ K}$$

= 308 K

We also know that the rate of the reaction doubles when temperature is increased by 10°.

Therefore, let us take the value of  $k_1 = k$  and that of  $k_2 = 2k$ 

Also, 
$$R = 8.314 \text{ J K}^{-1} \text{ mol}^{-1}$$

Now, substituting these values in the equation:

$$\log \frac{k_2}{k_1} = \frac{E_a}{2.303 R} \left[ \frac{T_2 - T_1}{T_1 T_2} \right]$$

We get:

$$\log \frac{2k}{k} = \frac{E_{\rm a}}{2.303 \times 8.314} \left[ \frac{10}{298 \times 308} \right]$$

$$\Rightarrow \log 2 = \frac{E_{\text{a}}}{2.303 \times 8.314} \left[ \frac{10}{298 \times 308} \right]$$

$$\Rightarrow E_{u} = \frac{2.303 \times 8.314 \times 298 \times 308 \times \log 2}{10}$$

 $= 52897.78 \text{ J mol}^{-1}$ 

 $= 52.9 \text{ kJ mol}^{-1}$ 

Note: There is a slight variation in this answer and the one given in the NCERT textbook.

#### Question 4.9:

The activation energy for the reaction

$$2HI(g) \rightarrow H_2 + I_{2(g)}$$

is 209.5 kJ mol<sup>-1</sup> at 581K. Calculate the fraction of molecules of reactants having energy equal to or greater than activation energy?

Answer

In the given case:

 $E_{\rm a} = 209.5 \; {\rm kJ} \; {\rm mol}^{-1} = 209500 \; {\rm J} \; {\rm mol}^{-1}$ 

T = 581 K

 $R = 8.314 \text{ JK}^{-1} \text{ mol}^{-1}$ 

Now, the fraction of molecules of reactants having energy equal to or greater than activation energy is given as:

$$x = e^{-Ea/RT}$$

$$\Rightarrow \ln x = -E_a / RT$$

$$\Rightarrow \log x = -\frac{E_a}{2.303 \ RT}$$

$$\Rightarrow \log x = \frac{209500 \,\mathrm{J \, mol^{-1}}}{2.303 \times 8.314 \,\mathrm{JK^{-1} \, mol^{-1}} \times 581} = 18.8323$$

Now, x = Anti log (18.8323)

= Anti  $\log \overline{19}.1677$ =  $1.471 \times 10^{-19}$