

Solution:- Assignment 7

Ans 1. At 400K, $t_{1/2} = \frac{0.693}{k_{400K}}$

$$\Rightarrow \frac{0.693}{10 \times 60} \text{ sec}^{-1} = 1.55 \times 10^{-3} \text{ sec}^{-1}$$

Again, $\frac{k_{420K}}{k_{400K}} = 3$

$$\Rightarrow k_{420K} = 3 \times 1.55 \times 10^{-3} \text{ sec}^{-1} = 4.65 \times 10^{-3} \text{ sec}^{-1}$$

$$\Rightarrow t_{0.5} = \frac{1}{4.65 \times 10^{-3} \text{ sec}^{-1}} \ln \frac{1}{1-0.25}$$

$$= 61.8 \text{ sec.}$$

Ans 2. $r = \frac{dP}{dt} = k[A][B]$

let the initial concentration be $[A]_0 = A_0$, $[B]_0 = B_0$ and $[P]_0 = 0$. Then, when P is formed in concentration x, the concentration of A changes to $A_0 - 2x$ and that of B changes to $B_0 - 3x$. Therefore,

$$\frac{dP}{dt} = \frac{dx}{dt} = k(A_0 - 2x)(B_0 - 3x) \quad \text{with } x=0 \text{ at } t=0$$

$$\int_0^t k dt = \int_0^x \frac{dx}{(A_0 - 2x)(B_0 - 3x)}$$

$$= \int_0^x \frac{6}{(2B_0 - 3A_0)} \times \left(\frac{1}{3(A_0 - 2x)} - \frac{1}{2(B_0 - 3x)} \right) dx$$

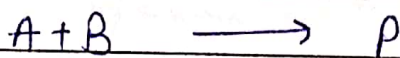
$$= \left(\frac{-1}{2B_0 - 3A_0} \right) \times \left(\int_0^x \frac{dx}{x - (1/2)A_0} - \int_0^x \frac{dx}{x - (1/3)B_0} \right)$$

$$kt = \frac{-1}{(2B_0 - 3A_0)} \times \left[\ln \left(\frac{x - 1/2 A_0}{-1/2 A_0} \right) - \ln \left(\frac{x - 1/3 B_0}{-1/3 B_0} \right) \right]$$

$$= \frac{-1}{(2B_0 - 3A_0)} \ln \left(\frac{(2x - A_0) B_0}{A_0 (3x - B_0)} \right)$$

$$= \left(\frac{1}{(3A_0 - 2B_0)} \right) \ln \left(\frac{(2x - A_0) B_0}{A_0 (3x - B_0)} \right)$$

Ans 3:- for second order-reaction of like:



Integrated rate law is:

$$kt = \frac{1}{[B]_0 - [A]_0} \ln \frac{[A]_0 ([B]_0 - x)}{([A]_0 - x) [B]_0}$$

Here, A represents $\text{CH}_3\text{COOC}_2\text{H}_5$ & B represents NaOH .

(a) For 10s :-

$$(0.21 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}) \times (10\text{s}) = \frac{1}{(0.030 - 0.20) \text{ dm}^3 \text{ mol}^{-1}} \ln \left[\frac{0.20}{0.03} \left(\frac{0.03 - x}{0.20 - x} \right) \right]$$

$$2.1 \text{ dm}^3 \text{ mol}^{-1} = \frac{2.303}{(-0.17 \text{ mol dm}^{-3})} \log \left(6.66 \left(\frac{0.03 - x}{0.20 - x} \right) \right)$$

$$\frac{-2.1 \text{ dm}^3 \text{ mol}^{-1}}{13.54 \text{ dm}^3 \text{ mol}^{-1}} = \log \left(6.66 \left(\frac{0.03 - x}{0.20 - x} \right) \right)$$

$$-0.155 = \log \left(6.66 \left(\frac{0.03 - x}{0.20 - x} \right) \right)$$

$$0.699 = 6.66 \left(\frac{0.03 - x}{0.20 - x} \right)$$

$$0.104 = \frac{0.03 - x}{0.20 - x}$$

$$0.104(0.20 - x) = 0.03 - x$$

$$0.0208 - 0.104x = 0.03 - x$$

$$x - 0.104x = 0.03 - 0.0208$$

$$0.896x = 0.0092$$

$$x = 0.010 \text{ mol dm}^{-3}$$

$$\begin{aligned} \text{Conc. of ester after 10s} &= 0.20 \text{ mol dm}^{-3} - 0.010 \text{ mol dm}^{-3} \\ &= 0.19 \text{ mol dm}^{-3} \end{aligned}$$

(b) For 10 minutes:-

$$(0.21 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}) \times (600 \text{ s}) = -(13.54 \text{ dm}^3 \text{ mol}^{-1}) \log \left(6.66 \left(\frac{0.03-x}{0.2-x} \right) \right)$$

$$-\frac{126 \text{ dm}^3 \text{ mol}^{-1}}{13.54 \text{ dm}^3 \text{ mol}^{-1}} = \log \left(6.66 \left(\frac{0.03-x}{0.2-x} \right) \right)$$

$$-9.30 = \log \left(6.66 \left(\frac{0.03-x}{0.2-x} \right) \right)$$

$$+5.01 \times 10^{-10} = 6.66 \left(\frac{0.03-x}{0.2-x} \right)$$

$$0.75 \times 10^{-10} = \left(\frac{0.03-x}{0.2-x} \right)$$

$$(0.2-x)(0.75 \times 10^{-10}) = (0.03-x)$$

$$0.15 \times 10^{-10} - 0.75 \times 10^{-10} x = 0.03 - x$$

$$0.15 \times 10^{-10} - 0.03 = -0.75 \times 10^{-10} x - x$$

$$x \approx 0.03 \text{ mol dm}^{-3}$$

$$\begin{aligned} \text{Conc of ester after 10 minutes} &= 0.20 - 0.03 \\ &= 0.17 \text{ mol dm}^{-3} \end{aligned}$$

Ans 4:- A reaction nth-order in A has the following rate law:

$$-\frac{d[A]}{dt} = k[A]^n \text{ so } \frac{d[A]}{[A]^n} = -k dt = [A]^{-n} d[A]$$

Integration yields

$$\frac{[A]^{1-n} - [A]_0^{1-n}}{1-n} = -kt$$

let $t_{1/3}$ be the time at which $[A] = [A]_0/3$.

$$\text{So, } -kt_{1/3} = \frac{([A]_0/3)^{1-n} - [A]_0^{1-n}}{1-n}$$

$$= [A]_0^{1-n} \left[\left(\frac{1}{3} \right)^{1-n} - 1 \right]$$

$$\text{And } t_{1/3} = \frac{3^{n-1} - 1}{k(n-1)} [A]_0^{1-n}$$

Ans 5 $\ln \frac{k_2}{k_1} = -\frac{E_a}{R} \left[\frac{1}{T_2} - \frac{1}{T_1} \right]$

$$\frac{-R \ln k_2/k_1}{E_a} = \frac{1}{T_2} - \frac{1}{T_1}$$

$$T_2 = \frac{1}{\frac{1}{T_1} - \frac{R \ln k_2/k_1}{E_a}}$$

$$T_2 = \frac{1}{\frac{E_a - T_1 R \ln k_2/k_1}{E_a T_1}}$$

$$T_2 = \frac{E_a T_1}{E_a - T_1 R \ln k_2/k_1}$$

$$T_2 = \frac{(154000 \text{ J/mol}) \times (290 + 273.15 \text{ K})}{154000 \text{ J/mol} - (290 + 273.15 \text{ K})(8.314 \text{ J/mol K}) \ln \left(\frac{3.20 \times 10^{-4} \text{ s}^{-1}}{9.50 \times 10^{-4} \text{ s}^{-1}} \right)}$$

$$= \frac{86795100 \text{ J/mol} \cdot \text{K}}{152844.19 \text{ J/mol}}$$

$$= 567.40 \text{ K}$$

Ans 6:- The first order half life is related to rate constant by:-

$$t_{1/2} = \ln 2 / k \quad \text{So } k = \frac{\ln 2}{t_{1/2}}$$

$$\Rightarrow k = \frac{\ln 2}{22.14} = 0.0313 \text{ y}^{-1}$$

The Integrated rate law tells us

$$[{}^{90}\text{Sr}] = [{}^{90}\text{Sr}]_0 e^{-kt} \quad \text{So } m = m_0 e^{-kt}$$

Where m is the mass of ${}^{90}\text{Sr}$.

7

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PAGE NO

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(a) After 15 year :-

$$m = (2.0 \mu\text{g}) \times \exp(-0.0313\text{y}^{-1} \times 15\text{y})$$
$$= 1.25 \mu\text{g}$$

(b) After 60 year :-

$$m = (2.0 \mu\text{g}) \times \exp(-0.0313\text{y}^{-1} \times 60\text{y})$$
$$= 0.3057 \mu\text{g}$$