CHEMICAL KINETICS

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Content

- Rate of Reaction
- Order of Reaction
- Arrhenius Equation

Introduction

- In chemical Kinetic, Reaction occurs in single step are Simple or Elementary reaction & those which take two or more steps are complex reaction.
- For elementary reaction order of reaction & molecularity are *generally* same.

Rate of Reaction

> Change in concentration of Reactant or Product per unit time called rate of Reacⁿ.

$$\rightarrow$$
 t=t a-x x

$$\rightarrow$$
 => $r = k[a-x]$

Rate Constant

$$Rate = \frac{dx}{dt}$$

$$rate = \frac{a - x - a}{dt} = \frac{-x}{t} = \frac{dx}{dt} (for A)$$

$$rate = \frac{x - 0}{dt} (for P)$$

Rate of Reaction

$$A + B ----> C + D$$

Rate of Reaction
$$(r) = \frac{-d(A)}{dt} = \frac{-d(B)}{dt} = \frac{d(C)}{dt} = \frac{d(D)}{dt} = \frac{dx}{dt}$$

- > Rate of disappearance of reactant = Rate of appearance of Product
- \rightarrow eg. 2A + 3B ---> C + 4D

 \triangleright

$$rate = \frac{-1}{2} \frac{d[A]}{dt} = \frac{-1}{3} \frac{d[B]}{dt} = \frac{d[C]}{dt} = \frac{1}{4} \frac{d[D]}{dt}$$

 \rightarrow Concⁿ = P^r at Temp^r constant

Order of Reaction

- No of molecules of reactant whose concⁿ determines the rate of reaction.
- > It is experimental value. It may be zero or fraction, -ve also.
- > It is derive from rate expression.

$$A \to P$$

$$R = K[A]^x$$

x = order of Reaction

1st order of Reaction

$$R = K[A]^x$$

$$kt = 2.303 \log \frac{a}{a - x}$$

a= initial concⁿ; a-x=left concⁿ; k=time⁻¹

 $t(1/2) = \frac{0.693}{K}$

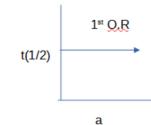
: Independent of initial concentration

$$t=t/2$$

$$a=a; a-x=a/2$$

$$k = \frac{2.303}{t(1/2)} \log \frac{a}{a/2} = \frac{0.693}{t(1/2)}$$

$$t(1/2) = \frac{0.693}{K}$$



x = 1

 $A \rightarrow P$

$$r = k[a - x]$$

$$\frac{dx}{dt} = k [a - x]$$

$$\frac{dx}{a-x} = kdt$$

$$kt = 2.303 \log \frac{a}{a - x}$$

Zero order of Reaction

$$\rightarrow$$
 t=t a-x x

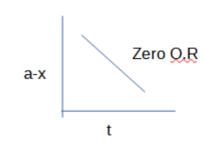
$$R = K[A]^x \quad \text{at } x = 0$$

•
$$x = kt$$
 ; $k = mole L^{-1} time^{-1}$

$$\Rightarrow$$
 If t=t/2 => x=a/2

>
$$t(1/2) = \frac{a}{2k} : t_{1/2} \propto a$$

Rate const. => depends only Temp^r



$$r = k \left[a - x \right]^0$$

$$r = k$$

$$\frac{dx}{dt} = k$$

$$dx = k \cdot dt$$

$$x = kt$$

Arrhenius Eqn

Relation between rate constant & Temperature

$$\rightarrow$$
 K = Ae^{-Ea/RT}

$$\log K = \log A - \frac{Ea}{2.303 RT}$$

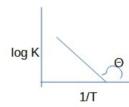
- > A= Arrhenius Constant or Pre-exponential factor
- Ea = activation energy; R= gas const.; T= temp. In Kelvin

$$K = Ae^{-(Ea/RT)}$$

$$\ln K = \ln A - \frac{Ea}{RT}$$

$$2.303 \log K = 2.303 \log A - \frac{Ea}{RT}$$

$$\log K = \log A - \frac{Ea}{2.303 RT}$$



$$slope = tan\Theta = \frac{-Ea}{2.303 R}$$
$$y - intercept = log A$$

Conclusion