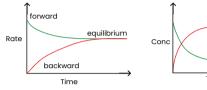
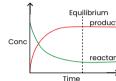
EQUILIBRIUM

- The chemical reactions which takes place in both directions are called reversible reactions
- Equilibrium is the end state of a reversible reaction.
- Gaseous Equilibrium is established only in a closed container.
- At equilibrium, the rate of forward and backward reactions are equal.
- At equilibrium, the concentration of reactants & products becomes constant.

GRAPHICAL REPRESENTATIONS





Q. For the equilibrium $A \longrightarrow B$, the variation of the rate of the forward (a) & reverse (b) reaction with time is given by



Applications of K & Q

- Value of K depends only on temperature.
- If K for the reaction $aA + bB \implies cC+dD$ is K, then K for the reaction cC+dD \iff aA+ bB will be $\frac{1}{2}$
- If K for the reaction $aA + bB \iff cC+dD$ is K, then K for the reaction $naA + nbB \implies ncC + ndD$ will be $(K)^n$
- During the addition of two reactions having equilibrium constants $K_1 \& K_2$, then the net Constant $K = K_1 \times K_2$
- \bullet During the subtraction of a reaction having constant K_2 from a reaction having constant K_1 , then the net constant $K = K_1/K_2$
- If Q<K, the reaction will proceed in forward direction
- If Q>K, the reaction will proceed in backward direction
- If Q=K, the system is in equilibrium.
- If $K > 10^3$, the reaction is almost complete in forward direction.
- If $K < 10^3$, the reaction is in backward direction.
- If K is in $b/w 10^3 \& 10^{-3}$ almost same reaction takes place in both forward and backward direction
- QAt a given temperature, the equilibrium constants for the reactions, $NO(g) + \frac{1}{2}O_2(g) \longrightarrow NO_2(g) \& 2NO_2(g) \longrightarrow 2NO(g) + O_2(g)$

are K₁ and K₂ respectively. If K₁ is 4×10^{-3} . then K₂ will be

(A)
$$8x10^{-3}$$
 (B) $16x10^{-3}$ (C) $6.25x10^{4}$ (D) $6.25x10^{6}$

PHYSICAL EQUILIBRIUM

- Such equilibrium is established in physical reactions.
- It is dynamic in nature.

LIQUID — VAPOUR EQUILIBRIUM

Here vapour pressure is constant at a constant temp for given.

SOLID - LIQUID EQUILIBRIUM

- Established only at a constant temparature
- Ice-water equilibrium established at 0°C (at1 atm)

SOLID IN LIQUID EQUILIBRIUM

- Established only in a Saturated solution
- eg: Saturated sugar solution. Sugar (dissolved) ______ Sugar (undissolved)

GAS IN LIQUID EQUILIBRIUM

- Here solubility depends upon pressure (Henry's law)
- eg: Soda water
- CO_2 (dissolved) \longleftarrow CO_2 (undissolved)

Q. Which of the given statements does not elucidate the equilibrium state precisely?

- (A) The equilibrium can be approached from either direction.
- (B) The equilibrium can be attained only if the system is an isolated system.
- (C) The free energy change at constant pressure and temperature is zero.
- (D) It is dynamic in nature.

CHEMICAL EQUILIBRIUM

HOMOGENEOUS EQUILIBRIUM

- If they are in solid or liquid phase(or aqueous), K can be represented as Kc.
- If they are in gaseous phase, K can be represented as Kp
- For a general reaction

$$aA + bB \longrightarrow cC + dD$$

$$K_{c} = \frac{[C]^{c}[D]^{d}}{[A]^{a}[B]^{b}} & \& K_{p} = \frac{P_{c}^{c}P_{D}^{d}}{P_{A}^{a}P_{B}^{b}}$$

 $K_p = K_c (RT)^{\Delta n_g} \Delta n_g = n_g - n_g$

UNIT OF EQUILIBRIUM CONSTANT

- Unit of $K_a = (\text{mol/L})^{\Delta n_g}$
- Unit of $K_p = (atm)^{\Delta n_g}$
- If △n = 0, equilibrium constant has no unit

HETEROGENEOUS EQUILIBRIUM

- In heterogeneous equilibrium, concentration of pure solid &

Q For the reaction,

$$\mathrm{Fe_2}\ \mathrm{N}_{(s)} + rac{3}{2}\mathrm{H}_{2(g)}
ightleftharpoons 2\mathrm{Fe}_{(s)} + \mathrm{NH}_{3(g)}$$

(a)
$$K_c = K_p(RT)$$

(a)
$$K_c = K_p(RT)$$
 (b) $K_c = K_p(RT)^{-1/2}$ (c) $K_c = K_p(RT)^{1/2}$ (d) $K_c = K_p(RT)^{3/2}$

(c)
$$K_c = K_P(RT)^1$$

d)
$$K_c = K_P (RT)^{3/2}$$

CHEMICAL EQUILIBRIUM

- Chemical equilibrium approaches from both forward & backward direction
- Chemical equilibrium is dynamic in nature.

EQUILIBRIUM CONSTANT[K]

For a general reversible reaction $aA + bB \iff cC + dD$

$$K_c = \frac{[C]^c [D]^d}{[A]^\alpha [B]^b}$$

REACTION QUOTIENT [Q]

- At any time during the reaction aA+bB ← cc+dD the ratio $[C]^c[D]^d$ is known as concentration quotient, Q_c . [A]a [B]b
- At equilibrium Q_c = K_c

CHARACTERISTICS OF K [eab. const]

- Value of K does not depends upon initial concentration of reactants and products.
- 2. Value of K does not depends upon the direction from which
- \bigcirc In the given reaction: A+2B \Longrightarrow 2C, 2 moles each of A & B present in 10 L of solution combine to form 1 mole of C. Calculate K, for the reaction.

(A) 1.5 (B) 6.67 (C) 0.15 (D) 2.3

LE CHATELIER'S PRINCIPLE

According to Le-chatelier's principle, if a system at equilibrium is subjected to a change in concentration, temperature or pressure, the equilibrium will shifts automatically in one direction which will nullify the effect of the change

- 1. Conc. of reactant increases —shift towards forward reaction.
- 2. Conc. of product decreases→shift towards forward reaction.
- 3. Conc. of reactant decreases—shift towards backward reaction.
- 4 Conc. of product increases → shift towards backward reaction. 5. Pressure increases \rightarrow shift towards lesser number of gaseous moles
- 6. Pressure decreases → shift towards higher number of gaseous moles
- 7 No. of gaseous moles of reactants & products are equal, pressure has no effect
- If temperature increases → shift towards endothermic
- g If temperature decreases -- shift towards exothermic
- 10. Catalyst helps to attain eqm state easily.
- 11 Addition of inert gas at constant volume, no effect.
- 12. Addition of inert gas at constant pressure → shift towards higher number of gaseous moles
- Q Which one of the following conditions will favour maximum formation of the product in the reaction $A_2(g) + B_2(g) \longrightarrow X_2(g)$. $\Delta rH = -XkJ/mol$?
- (A) Low temperature and high pressure (B) High temperature and high pressure

(C) Low temperature and low pressure (D) High temperature and low pressure