# Wisdom Education Academy 

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## Physical and Chemical Processes

Physical processes involve such changes, which only affects the physical properties of the substance undergoing changes but have no effect on the chemical composition and properties.

Chemical processes involve changes in chemical composition and properties. Whenever a chemical change occurs, we can say that a chemical reaction has taken place.

## Types of Chemical Reactions

1. Combination Reactions

In such reactions two or more substances combine to form a single compound.
e.g.,
$2 \mathrm{Mg}+\mathrm{O}_{2} \rightarrow 2 \mathrm{MgO}$

## 2. Decomposition Reactions

In these reactions. a compound decomposes to produce two or more different substances.
e.g., $\mathrm{PCI}_{5} \Leftrightarrow \mathrm{PCI}_{3}+\mathrm{CI}_{2}$

Digestion of food is also a decomposition reaction.
[Decomposition by heat IS called thermal decomposition and decomposition by sunlight is called photo decomposition.]

## 3. Displacement Reactions

These reactions involve displacement of one element or group by another. These are infact, redox reactions, e.g.,

## $\mathrm{Zn}(\mathrm{s})+\mathrm{H}_{2} \mathrm{SO}_{4}$

## 4. Double Displacement or Metathesis Reactions

In these. reactions two compounds react to form two new compounds and no change in oxidation state take place, e.g., precipitation reactions, neutralisation, reactions.
$\mathrm{AgNO}_{3}(\mathrm{aq})+\mathrm{NaCI}(\mathrm{aq}) \rightarrow \rightarrow \mathrm{AgCl}(\mathrm{s})+\mathrm{NaNO}_{3}(\mathrm{aq})$
5. Reversible and Irreversible Reactions

| Irreversiplesreactions |  |
| :--- | :--- |
| 1. | Chemical reactions which always <br> proceed to completion in only forward <br> direction, e.g., <br> $A_{g N O}+\mathrm{NaCl}$ |
| 2. | These reactions never attain equilibrium. |

Reactions which do not proceed to completion in forward direction and also proceed in the backward direction under suitable conditions, e.g. .

$$
\mathrm{N}_{2}(g)+3 \mathrm{H}_{2}(g) \rightleftharpoons 2 \mathrm{NH}_{3}
$$

These reactions attain equilibrium.

## Equilibrium State

Under given set of conditions if a reversible process or chemical reaction is carried out in a closed container, a constancy in some observable properties like colour intensity, pressure, density, is observed. Such a state is referred to as an equilibrium state.

Equilibrium may be classified as :

## Physical Equilibrium

Equilibrium set up in physical processes like evaporation of water, melting of solids, dissolution of solutes, etc., is called physical equilibrium, e.g., Ice $\Leftrightarrow$ Water

At equilibrium,
Rate of melting of ice $=$ Rate of freezing of water

## Chemical Equilibrium

If a reversible reaction is carried out in a closed vessel, a stage is attained where the speed of the forward reaction equals the speed of the backward reaction. It corresponds to chemical equilibrium. At equilibrium,

Rate of forward reaction = Rate of backward reaction


## Characteristics of Chemical Equilibrium

1. Equilibrium can be attained from either side.
2. Equilibrium is dynamic in nature, i.e., at equilibrium reaction does not stop.
3. At equilibrium, there is no change in the concentration of various species.
4. The equilibrium state remains unaffected by the presence of catalyst. Catalyst helps to attain the equilibrium state rapidly.
5. The observable physical properties of the process become constant.

## Law of Mass Action

Guldberg and Waage states that the rate of a chemical reaction is directly proportional to the product of the active masses of the reacting substances. For a general reaction,

$$
a A+b B \rightleftharpoons c C+d D
$$

## Rate of forward reaction $\propto[A]^{a}[B]^{b}=k_{f}[A]^{a}[B]^{b}$

## Rate of backward reaction $\propto[C]^{c}[D]^{d}=k_{b}[C]^{c}[D]^{d}$

where, $\mathrm{k}_{\mathrm{f}}$ and $\mathrm{k}_{\mathrm{b}}$ are rate constants.
In heterogeneous equilibrium, the active mass of pure solids and liquids are taken as
At equilibrium,
Rate of forward reaction = Rate of backward reaction

$$
\begin{aligned}
k_{f}[A]^{a}[B]^{b} & =k_{b}[C]^{c}[D]^{d} \\
\frac{k_{f}}{k_{b}} & =K_{c}=\frac{[C]^{c}[D]^{d}}{[A]^{a}[B]^{b}}
\end{aligned}
$$

$\mathbf{K}_{\mathrm{c}}$ is called the equilibrium constant.

## Use of Partial Pressures Instead of Concentration

For gaseous reactions, partial pressures are conveniently used since at any fixed temperature partial pressure is directly proportional to concentration. For a general gaseous reaction,

$$
\begin{aligned}
& a A+b B \rightleftharpoons c C+d D \\
& K_{p}=\frac{p_{C}^{c} \times p_{D}^{d}}{p_{A}^{a} \times p_{B}^{b}}
\end{aligned}
$$

## Relation between $K_{c}$ and $K_{c}$

where, $\Delta \mathrm{n}_{\mathrm{s}}=$ moles of products - moles of reactants (gaseous only)
Relation between $K_{c}$ and $K_{p}$ for different types of reactions
(i) When $\Delta \mathrm{n}_{\mathrm{g}}=0, \mathrm{~K}_{\mathrm{p}}=\mathrm{K}_{\mathrm{c}}$
(ii) When $\Delta \mathrm{n}_{\mathrm{g}}=+\mathrm{ve}, \mathrm{K}_{\mathrm{p}}>\mathrm{K}_{\mathrm{c}}$
(iii) When $\Delta \mathrm{n}_{\mathrm{g}}=-\mathrm{ve}, \mathrm{K}_{\mathrm{p}}<\mathrm{K}_{\mathrm{c}}$

Units of $K_{p}$ and $K_{\text {c }}$
(i) Unit of $K_{p}=(a t m)^{a^{n}}$
(ii) Unit of $\mathrm{K}_{\mathrm{c}}=\left(\mathrm{mol} \mathrm{L}^{-1}\right)^{\Delta n}$,

Characteristics of Equilibrium Constant $K_{p}$ or $K_{c}$

1. It has definite value for every chemical reaction at a particular temperature.
2. The more is the value of $K_{c}$ or $K_{p}$, the more is the extent of completion of reaction, i.e., $K_{c}<1$ indicates lesser concentration of products than reactants.
$\mathrm{K} \geq 10^{3}$ shows completion of reaction and $\mathrm{K} \leq 10^{-3}$ shows that the reaction does not proceed at all.
3. When the reaction can be expressed as sum of two other reactions, the $\mathrm{K}_{\mathrm{c}}$ of overall reaction is equal to the product of equilibrium constants of individual reactions.
4. The equilibrium constant is independent of initial concentrations of reactants.
5. Equilibrium constant is independent of presence of catalyst.
6. $\mathrm{K}_{\mathrm{c}}$ for backward reaction is inverse of $\mathrm{K}_{\mathrm{c}}$ for forward reaction.
7. If an equation is multiplied by $n$, the $K$ becomes $K^{n}$, and if it is divided by m, the $k$ becomes ${ }^{m} \sqrt{ } k$.
8. In equilibrium constant expression if activities are used in places of molar concentration, h becomes dimensionless.

## Types of Equilibrium <br> Homogeneous Equilibrium

In homogeneous equilibrium, the reactants and products arc present in the same phase or physical suite (gaseous or liquid).
$2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \Leftrightarrow 2 \mathrm{SO}_{3}(\mathrm{~g})$

## Heterogeneous Equilibrium

In heterogeneous equilibrium the reactants and products are present in two or more physical states or phases.
$3 \mathrm{Fe}(\mathrm{s})+4 \mathrm{H}_{2} \mathrm{O}(\mathrm{g}) \Leftrightarrow \mathrm{Fe}_{3} \mathrm{O}_{4}(\mathrm{~s})+4 \mathrm{H}_{2}(\mathrm{~g})$

## Reaction Quotient

For any reversible reaction at any stage other than equilibrium, the ratio of the molar concentrations of the products to that of the reactants. where each concentration term is raised to the power equal to the stoichiometric coefficient to the substance concerned, is called the reaction quotient, $\mathrm{Q}_{c}$.
For a general reaction
$\mathrm{aA}+\mathrm{bB} \Leftrightarrow \mathrm{cC}+\mathrm{dD}$
which is not at equilibrium,
$\mathrm{Q}_{\mathrm{c}}=[\mathrm{C}]^{\mathrm{c}}+[\mathrm{D}]^{\mathrm{d}} /[\mathrm{A}]^{\mathrm{a}}[\mathrm{B}]^{\mathrm{b}}$
If
(i) $\mathrm{Q}_{c}>\mathrm{K}_{c}$, the value of $\mathrm{Q}_{\mathrm{c}}$ will tend to decrease to reach the value of $\mathrm{K}_{\mathrm{c}}$ (towards equilibrium) and the reaction will proceed in the reverse direction.
(ii) $\mathrm{Q}_{\mathrm{c}}<\mathrm{K}_{\mathrm{c}}$ it will lend to increase and the reaction will proceed in the forward direction.
(ii) $\mathrm{Q}_{\mathrm{c}}=K_{c}$, the reaction is at equilibrium.

Le - Chatelier's Principle
There are three main factors which affect the state of equilibrium.
They are

## concentration

temperature
pressure.
Le - Chatelier's principle states that if a system at equilibrium is subjected to a change in concentration.
pressure or temperature. the equilibrium equilibrium change.

## Effect of Change of Concentration

If at equilibrium the concentration of one of the reactants is increased. the equilibrium will shift in the forward direction and vice-versa.

## Effect of Change in Pressure

No effect of pressure on equilibria having same moles of reactants and products. e.g., $\mathrm{N}_{2}+\mathrm{O}_{2} \Leftrightarrow 2 \mathrm{NO}$. When there is change in the number of moles, the equilibrium will shift in the direction having smaller number of moles when the pressure is increased and vice-versa, e.g.,
$\mathrm{N}_{2}+3 \mathrm{H}_{2} \Leftrightarrow 2 \mathrm{NH}_{3}$ [High p. high yield of $\mathrm{NH}_{3}$ ]
Effect of Temperature
When process is exothermic, low temperature favours the forward reaction. When process is endothermic. high temperature favours the formation of products.

## Effect of Addition of Inert Gas

(i) Addition of inert gas at constant pressure At constant pressure. if an inert gas is added. it will increase the volume of the system. Therefore. the equilibrium will shift in a direction in which there is an increase in the number of moles of gases.
(ii) Addition of inert gas at constant volume If keeping volume of the system constant, an inert gas is added. the relative molar concentration of the substance will not change. Hence. the equilibrium position of the reaction remains unaffected.

## Effect of Catalyst

The presence of catalyst does not change the position of equilibrium. It simply fastens the attainment of equilibrium.

## Le-Chatelier's Principle Applicable to Physical Equilibrium

(i) Effect of pressure on solubility The increased pressure, will increase the solubility of gas and vice-versa.
(ii) Effect of temperature on solubility Some substances dissolve with the absorption of heat. Solubility of such substances will increase with increase of temperature and vice-versa, e.g., dissolution of $\mathrm{NH}_{4} \mathrm{CI}, \mathrm{KCI}$, $\mathrm{KNO}_{3}$, etc. The dissolution of calcium acetate and calcium hydroxide is exothermic, so their solubility is lowered at higher temperature.
(iii) Effect of pressure on the melting point of ice

Ice $\Leftrightarrow$ liquid water

The ice occupy the more volume than liquid water, so increased pressure will result in melting of ice according to Le-Chatelier principle.

Favourable conditions for some chemical equilibria to get higher yield of product.

| Chemical equilibria | Favourable condition for forward direction |
| :---: | :---: |
| $\mathrm{N}_{2}+3 \mathrm{H}_{2} \rightleftharpoons 2 \mathrm{NH}_{3}$ | High pressure, low temperature and isolation of $\mathrm{NH}_{3}$ by liquefaction. |
| $\mathrm{N}_{2}+\mathrm{O}_{2} \rightleftharpoons 2 \mathrm{NO}$ | High temperature and isolation of NO. |
| $\mathrm{PCl}_{5} \rightleftharpoons \mathrm{PCl}_{3}+\mathrm{Cl}_{2}$ | Low pressure, high temperature and isolation of $\mathrm{PCl}_{3}$ and $\mathrm{Cl}_{2}$ |

## Calculation of the Degree of Dissociation ( $\alpha$ ) from Density Measurement

 $\alpha=\mathrm{D}-\mathrm{d} / \mathrm{d}$where, $\mathrm{D}=$ theoretical vapour density
$\mathrm{d}=$ observed vapour density
Now, molecular mass $=2 * \mathrm{VD}$
$\therefore \alpha=\mathrm{M}_{\mathrm{c}}-\mathrm{M}_{\mathrm{o}} / \mathrm{M}_{\text {。 }}$
where, $\mathrm{M}_{\mathrm{c}}=$ calculated molecular weight
$\mathrm{M}_{\mathrm{o}}=$ observed molecular weight

## Thank You

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