

THE pH SCALE

Acidic strength refers to an acid's inclination to yield H_3O^+ or H^+ ions when dissolved in water. Therefore, the greater the propensity to release H^+ ions, the stronger the acidity of the substance. On the other hand, basic strength pertains to a base's ability to provide OH^- ions when mixed with water.

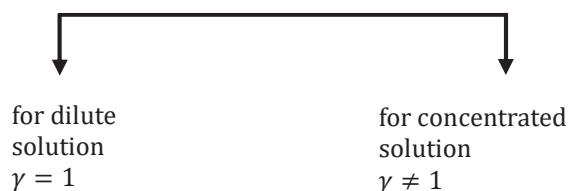
So, greater the tendency to give OH^- ions, more will be basic strength of the substance. The

concentration of H^+ ions is written in a simplified form introduced by Sorenson known as pH scale.

pH is defined as negative logarithm of activity of H^+ ions. activity (a) = $\gamma \cdot \frac{C}{C_0}$

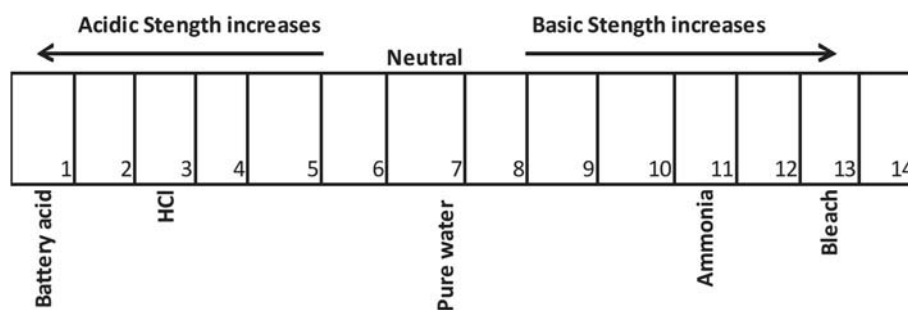
$C \rightarrow$ unitless, $C_0 \rightarrow$ standard concentration

$$a = \gamma \cdot C$$



$$\therefore a = C = [\text{H}^+]$$

$$\therefore \text{pH} = -\log (\text{where is the activity of } \text{H}^+ \text{ ions})$$



The activity of H^+ ions represent the concentration of unbound H^+ or H_3O^+ ions within a solution. The pH scale, established from 0 to 14, designates a midpoint at 7 when using water as the solvent at 25 °C. It's essential to recognize that alterations in temperature and solvent will result in a modification of the pH scale's range.

For example,

$$0 - 14 \quad \text{at } 25^\circ\text{C} (K_w = 10^{-14}) \quad \text{Neutral point, pH} = 7$$

$$0 - 13 \quad \text{at } 80^\circ\text{C} (K_w = 10^{-13}) \quad \text{Neutral point, pH} = 6.5$$

pH can also be negative or > 14

Relation Between pH and pOH

The pH (potential of hydrogen) is determined by the negative logarithm (base 10) of the concentration of hydronium ions (H_3O^+), expressed as $\text{pH} = -\log [\text{H}_3\text{O}^+]$. Similarly, the pOH (potential of hydroxide) is calculated as the negative logarithm (base 10) of the concentration of hydroxide ions (OH^-), denoted as $\text{pOH} = -\log [\text{OH}^-]$.

The product of the concentrations of hydronium and hydroxide ions in a solution, represented as $[\text{H}_3\text{O}^+][\text{OH}^-]$, is equal to the ion product constant (K_w), which is 10^{-14} at 25°C . Mathematically, this is expressed as $[\text{H}_3\text{O}^+][\text{OH}^-] = K_w = 10^{-14}$.

Taking the logarithm of both sides of the equation, we get $[\text{H}_3\text{O}^+] + \log[\text{OH}^-] = \log K_w = \log 10^{-14} = -14$. This equation signifies the relationship between the concentrations of hydronium and hydroxide ions and their connection to the ion product constant.

Furthermore, the sum of the pH and pOH in a solution is equal to the negative logarithm (base 10) of the ion product constant, denoted as $\text{p}K_w$, which is 14 at 298 K. In other words, $\text{pH} + \text{pOH} = \text{p}K_w = 14$ represents the equilibrium between the acidic and basic components in a solution at a given temperature.

Common Ion Effect

In the case of pure water, the concentrations of hydronium (H_3O^+) and hydroxide (OH^-) ions are both equal to 10^{-7} M. When an acid is introduced to pure water, the concentration of hydronium ions ($[\text{H}_3\text{O}^+]$) becomes greater than 10^{-7} , disrupting the equilibrium.

Conversely, when a base is added to pure water, the concentration of hydroxide ions ($[\text{OH}^-]$) exceeds 10^{-7} , while the concentration of hydronium ions ($[\text{H}_3\text{O}^+]$) is determined by K_w divided by the concentration of hydroxide ions, represented as $[\text{H}_3\text{O}^+] = K_w/[\text{OH}^-]$.

The concept of the common ion effect comes into play when a weak electrolyte's ionization is hindered in the presence of another electrolyte, particularly a strong electrolyte that shares a common ion with the weak electrolyte. This effect occurs because the addition of a common ion reduces the extent to which the weak electrolyte dissociates, impacting its ionization process.