ACIDS, BASES \& BUFFERS

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## Acid \& Base Theories

## Arrhenius Water Theory

Acids are substances which produce hydrogen ions in solution.
Alkalis are substances which produce hydroxide ions in solution.
NOTE: Alkalis are soluble BASES

$$
\mathrm{HA}_{(\mathrm{aq})}+\mathrm{BOH}_{(\mathrm{aq})} \longrightarrow \mathrm{BA}_{(\mathrm{aq})}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})}
$$

Neutralisation happens because hydrogen ions and hydroxide ions react to produce water.

## acid + base


salt + water
Dissociate to produce $\mathrm{OH}^{-}$
Accept protons
Donate electron pairs
$\mathrm{pH}>7$
WATER $\left(\mathrm{H}_{2} \mathrm{O}\right)$

$$
\mathrm{H}^{+}+\mathrm{OH}^{-} \longrightarrow \mathrm{H}_{2} \mathrm{O}
$$

Neutralisation
$\mathrm{pH}=7$

## Acids \& Bases

## Common Acids

| Chemical Name | Formula | Uses | Strength |
| :---: | :---: | :---: | :---: |
| Nitric Acid | $\mathrm{HNO}_{3}$ | Explosives, fertilizer, dyes, <br> glue | Strong |
| Sulfuric Acid | $\mathrm{H}_{2} \mathrm{SO}_{4}$ | Explosives, fertilizer, dyes, <br> glue, batteries | Strong |
| Hydrochloric Acid | HCl | Metal cleaning, food <br> preparation, ore refining, <br> stomach acid | Strong |
| Phosphoric Acid | $\mathrm{H}_{3} \mathrm{PO}_{4}$ | Fertilizer, plastics and <br> rubber, food preservation | Moderate |
| Acetic Acid | $\mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$ | Plastics and rubber, food <br> preservation, Vinegar | Weak |
| Hydrofluoric Acid | HF | Metal cleaning, <br> glass etching | Weak |
| Carbonic Acid | $\mathrm{H}_{2} \mathrm{CO}_{3}$ | Soda water | Weak |
| Boric Acid | $\mathrm{H}_{3} \mathrm{BO}_{3}$ | Eye wash | Weak |

## Common Bases

| Chemical <br> Name | Formula | Common <br> Name | Uses | Strength |
| :---: | :---: | :---: | :---: | :---: |
| Sodium <br> hydroxide | NaOH | Lye, <br> caustic soda | Soap, plastic, <br> petrol refining | Strong |
| Potassium <br> hydroxide | KOH | Caustic <br> potash | Soap, cotton, <br> electroplating | Strong |
| Calcium <br> hydroxide | $\mathrm{Ca}(\mathrm{OH})_{2}$ | Slaked lime | Cement | Strong |
| Sodium <br> bicarbonate | NaHCO | Baking soda | Cooking, antacid | Weak |
| Magnesium <br> hydroxide | $\mathrm{Mg}(\mathrm{OH})_{2}$ | Milk of <br> magnesia | Antacid | Weak |
| Ammonium <br> hydroxide | $\mathrm{NH}_{4} \mathrm{OH}$, | Ammonia |  |  |
| $\left\{\mathrm{NH}_{3}(\mathrm{aq})\right\}$ | water | Detergent, <br> fertilizer, <br> explosives, <br> fibers | Weak |  |

## Acids \& Base Theories

## Lewis Acid-Base Theory

Lewis Acid: a species that accepts an electron pair (i.e., an electrophile) and will have vacant orbitals.

Lewis Base: a species that donates an electron pair (i.e., a nucleophile) and will have lone-pair electrons.


Lewis Acids are Electrophilic ie. electron attracting. Various species can act as Lewis acids.

- All cations are Lewis acids since they are able to accept electrons. (e.g., $\mathrm{Cu}^{2+}$, $\mathrm{Fe}^{2+}, \mathrm{Fe}^{3+}$ )
- An atom, ion, or molecule with an incomplete octet of electrons can act as an Lewis acid (e.g., $\mathrm{BF}_{3}, \mathrm{AlF}_{3}$ ).
- Molecules where the central atom can have more than 8 valence shell electrons can be electron acceptors, and thus are classified as Lewis acids (e.g., $\mathrm{SiBr}_{4}, \mathrm{SiF}_{4}$ ).

Lewis Bases are Nucleophilic meaning that they "attack" a positive charge with their lone pair.

- Atoms, ions, or molecules with a lone-pair of electrons can thus be a Lewis base e.g., $\mathrm{OH}^{-}, \mathrm{CN}^{-}, \mathrm{CH}_{3} \mathrm{COO}^{-},: \mathrm{NH}_{3}, \mathrm{H}_{2} \mathrm{O}:, \mathrm{CO}:$


## Acids \& Base Theories

## Bronsted-Lowry Acid-Base Theory

An acid is a proton (hydrogen ion) donor.
A base is a proton (hydrogen ion) acceptor.


## Properties of Acids

Acids change the colour of litmus from blue to red.
Some Acids are highly corrosive
Acids react with active metals to evolve hydrogen gas

## Acids lose their acidity when mixed with a base.

When equal amounts of acid and base are combined the process of neutralisation occurs and salt and water is formed,

## Acids have a pH value of less than 7

Acids are sour in taste.
Acids react with carbonates and hydrogen carbonates to form a salt, water, and carbon dioxide gas.

Acids react with sulphites and bisulphites to form a salt, water and sulfur dioxide

Acids and metal sulphides react to form salt and hydrogen sulphide.

Acids are classified on the basis of their sources, strength, concentration, the presence of oxygen and its basicity.

The different types of acids are organic acids, mineral strong acids, weak acids, concentrated acids, dilute acids, Oxy-acids, Hydracids, monobasic acids, dibasic acids, and tribasic acids.

## Properties of Bases

Bases change the colour of litmus from red to blue.
Bases have a bitter taste.
Bases lose their basicity when mixed with acids.
Bases react with acids to form salt and water.
Bases feel slippery or soapy.
Strong bases are highly corrosive in nature whereas other bases are mildly corrosive.

The $\mathbf{p H}$ value of bases ranges from 8-14.
Bases and ammonium salts produce ammonia.
Bases are classified on the basis of strength, concentration and acidity.

The different kinds of acids are strong base acid, weak base acid, concentrated base, dilute base, monoacidic base, diacidic base and triacidic base.

## Reactions of Acids \& Bases

Acid + Base $\longrightarrow$ Salt + Water (Neutralisation) eg. $\mathrm{HCl}_{(\mathrm{aq})}+\mathrm{NaOH}_{(\mathrm{aq})} \longrightarrow \mathrm{NaCl}_{(\mathrm{aq)}}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})}$
$\mathrm{H}^{+}{ }_{(\mathrm{aq})}+\mathrm{OH}^{-}{ }_{(\mathrm{aq})} \longrightarrow \mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})}$

Acid + Reactive Metal $\longrightarrow$ Salt + Water
eg. $\mathrm{H}_{2} \mathrm{SO}_{4(\mathrm{aq})}+\mathrm{Cu}_{(\mathrm{s})} \longrightarrow \mathrm{CuSO}_{4(\mathrm{aq})}+\mathrm{H}_{2(\mathrm{~g})}$

Acid + Carbonate $\longrightarrow$ Salt + Carbon Dioxide + Water
eg. $\mathrm{HNO}_{3(a q)}+\mathrm{CaCO}_{3(\mathrm{aq})} \longrightarrow \mathrm{CaCO}_{3(\mathrm{aq})}+\mathrm{CO}_{2(\mathrm{~g})}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})}$

Acid + Sulfite $\longrightarrow$ Salt + Sulfur Dioxide + Water
eg. $2 \mathrm{HCl}_{(\mathrm{aq})}+\mathrm{Na}_{2} \mathrm{SO}_{3(\mathrm{aq})} \longrightarrow 2 \mathrm{NaCl}_{(\mathrm{aq})}+\mathrm{SO}_{2(\mathrm{~g})}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})}$

## Acids \& Bases

$\mathbf{p H}=-\log \left[\mathbf{H}^{+}\right]$

| $\mathbf{p H}$ | $\left[\mathbf{H}^{+}\right] \mathbf{~ M}$ | $\left[\mathbf{H}^{+}\right] \mathbf{~ M}$ | Examples |
| :--- | :--- | :--- | :--- |
| 0 | 1 | $10^{0}$ | Battery acid |
| 1 | 0.1 | $10^{-1}$ | Stomach acid |
| 2 | 0.01 | $10^{-2}$ | Lemon Juice |
| 3 | 0.001 | $10^{-3}$ | Vinegar |
| 4 | 0.0001 | $10^{-4}$ | Tomato juice |
| 5 | 0.00001 | $10^{-5}$ | Black coffee |
| 6 | 0.000001 | $10^{-6}$ | Saliva |
| 7 | 0.0000001 | $10^{-7}$ | Pure Water |
| 8 | 0.00000001 | $10^{-8}$ | Sea water |
| 9 | 0.000000001 | $10^{-9}$ | Baking soda |
| 10 | 0.0000000001 | $10^{-10}$ | Bar of soap |
| 11 | 0.00000000001 | $10^{-11}$ | Ammonia |
| 12 | 0.000000000001 | $10^{-12}$ | Household bleach |
| 13 | 0.0000000000001 | $10^{-13}$ | Oven cleaner |
| 14 | 0.00000000000001 | $10^{-14}$ | Sodium hydroxide |

## Strong Acids \& Bases

Strong acids (HA) and bases (BOH) completely dissociate into their ions in water.

$$
\begin{aligned}
& \mathrm{HA} \longrightarrow \mathrm{H}^{+}+\mathrm{A}^{-} \\
& \mathrm{BOH} \longrightarrow \mathrm{~B}++\mathrm{OH}^{-}
\end{aligned}
$$

## Strong Acids

HCl : hydrochloric acid
$\mathrm{HNO}_{3}$ : nitric acid
$\mathrm{H}_{2} \mathrm{SO}_{4}$ : sulfuric acid
HBr : hydrobromic acid
HI: hydroiodic acid
$\mathrm{HClO}_{4}$ : perchloric acid

## Strong Bases

LiOH: lithium hydroxide NaOH : sodium hydroxide KOH : potassium hydroxide RbOH: rubidium hydroxide CsOH : cesium hydroxide $\mathrm{Ca}(\mathrm{OH})_{2}$ : calcium hydroxide $\mathrm{Sr}(\mathrm{OH})_{2}$ : strontium hydroxide $\mathrm{Ba}(\mathrm{OH})_{2}$ : barium hydroxide

## Weak Acids \& Bases

Weak acids and bases partially dissociate in water.

$$
\begin{aligned}
& \mathrm{HA} \leftrightharpoons \mathrm{H}^{+}+\mathrm{A}^{-} \\
& \mathrm{BOH} \leftrightharpoons \mathrm{~B}^{+}+\mathrm{OH}^{-}
\end{aligned}
$$

The extent to which weak acids and bases dissociate is given by their dissociation constant value.

$$
\begin{aligned}
& \text { Acid dissociation constant, } \mathrm{K}_{\mathrm{a}}=\frac{\left[\mathrm{H}^{+}\left[\mathrm{A}^{-}\right]\right.}{[\mathrm{HA}]} \\
& \text { Base dissociation constant, } \mathrm{K}_{\mathrm{b}}=\frac{\left[\mathrm{B}^{+}\left[\mathrm{OH}^{-}\right]\right.}{[\mathrm{BOH}]}
\end{aligned}
$$

The higher the value if $\mathrm{K}_{\mathrm{a}}, \mathrm{K}_{\mathrm{b}}$, the stronger the weak acid or base.
Weak Acids

| Acid | Formula |
| :---: | :---: |
| Formic | $\mathrm{HCOOH}^{2}$ |
| Acetic | $\mathrm{CH}_{3} \mathrm{COOH}$ |
| Trichloroacetic | $\mathrm{CCl}_{3} \mathrm{COOH}$ |
| Hydrofluoric | HF |
| Hydrocyanic | HCN |
| Hydrogen sulfide | $\mathrm{H}_{2} \mathrm{~S}$ |
| Water | $\mathrm{H}_{2} \mathrm{O}$ |

Weak Bases

| Acid | Formula |
| :---: | :---: |
| Ammonia | $\mathrm{NH}_{3}$ |
| Pyridine | $\mathrm{C}_{5} \mathrm{~N}_{5} \mathrm{~N}$ |
| Methylamine | $\mathrm{CH}_{3} \mathrm{NH}_{2}$ |
| Water | $\mathrm{H}_{2} \mathrm{O}$ |

## Water Dissociation Constant ( $\mathrm{K}_{\mathrm{w}}$ )

Water partially dissociates (autoionises):

$$
\mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})} \leftrightharpoons \mathrm{H}^{+}{ }_{(\mathrm{aq})}+\mathrm{OH}^{-}{ }_{(\mathrm{aq})}
$$

$\mathrm{H}^{+}$is a proton and combines with a molecule of water to form the hydroxonium ion, $\mathrm{H}_{3} \mathrm{O}^{+}$

$$
2 \mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})} \leftrightharpoons \mathrm{H}_{3} \mathrm{O}^{+}{ }_{(\mathrm{aq})}+\mathrm{OH}^{-}{ }_{(\mathrm{aq})}
$$

## Water behaves as both a weak acid and a weak base.

The Water Dissociation Constant (KW) is just another acid dissociation constant.

This gives you:

$$
\mathrm{K}_{\mathrm{w}}=\left[\mathrm{H}^{+}\right]\left[\mathrm{OH}^{-}\right]
$$

Kw always equals $\mathbf{~ m o l}^{\mathbf{2}} \mathbf{d m}^{-6}$

In pure water, $\left[\mathrm{H}^{+}\right]=\left[\mathrm{OH}^{-}\right]$. Under normal temperature and pressure, $\left[\mathrm{H}^{+}\right]=1 \times 10^{-7} \mathrm{M}$.

$$
\mathrm{pH} \text { of pure water }=-\log _{10}\left[1 \times 10^{-7}\right]=7
$$

## Buffers

A buffer solution is one which resists changes in pH when small quantities of an acid or an alkali are added to it.

Acid Buffer: mixtures of a weak acid (HA) and its conjugate base (A-)

## HA/NaA

The presence of $\mathrm{A}^{-}$from the dissolution of NaA suppresses the dissociation of HA.

Buffer mixture contains 'reservoirs of HA and $\mathrm{A}^{-}$

$$
\begin{aligned}
& \mathrm{HA} \leftrightharpoons \mathrm{H}^{+}+\mathrm{A}^{-} \\
& \mathrm{NaA} \leftrightharpoons \mathrm{Na}++\mathrm{A}-
\end{aligned}
$$



Buffering capacity determined by concentrations of HA and A-

```
pH of buffer influenced by [A-]/[HA] ratio
```


## Buffers

Basic Buffer: mixtures of a weak base $(\mathrm{BOH})$ and its conjugate base $(\mathrm{B}+)$

## BOH/BA

The presence of $B+$ from the dissolution of $B A$ suppresses the dissociation of BOH.

Buffer mixture contains 'reservoirs of BOH and $\mathrm{B}^{+}$

Weak acid:
$\mathrm{BOH} \leftrightharpoons \mathrm{B}^{+}+\mathrm{OH}^{-}$

Salt of weak acid:
$B A \longrightarrow B^{+}+A^{-}$
Buffer destroyed once added $\left[\mathrm{H}^{+}\right]$exceeds [BOH]

## Addition of ACID $\left(\mathrm{H}^{+}\right)$

$$
\mathrm{BOH}+\mathrm{H}^{+} \longrightarrow \mathrm{B}^{+}+\mathrm{H}_{2} \mathrm{O}
$$

## BOH



Buffering capacity determined by concentrations of BOH and $\mathrm{B}^{+}$
pH of buffer influenced by $\left[\mathrm{B}^{+}\right] /[\mathrm{BOH}]$ ratio

## pH of Buffers

## Henderson-Hasselbalch equation

## Acidic Buffers

$$
\mathrm{pH}=\mathrm{pK}_{\mathrm{a}}+\frac{\log \left[\mathrm{A}^{-}\right]}{[\mathrm{HA}]}
$$

## Basic Buffers

$$
\mathrm{pOH}=\mathrm{pK}_{\mathrm{b}}+\frac{\log [\mathrm{B}]}{[\mathrm{BOH}]}
$$

$$
\begin{aligned}
\mathrm{pK}_{\mathrm{a}} & =-\log \mathrm{K}_{\mathrm{a}} \\
\mathrm{~K}_{\mathrm{a}} & =10^{-\mathrm{pKa}}
\end{aligned}
$$

$$
\begin{aligned}
p K_{b} & =-\log K_{b} \\
K_{a} & =10^{-\mathrm{pKb}}
\end{aligned}
$$

