# 8.1 Explaining the Properties of Acids \& Bases 



SCH4U - Chemistry, Gr. 12, University Prep

## Equilibrium \& Acids \& Bases

$\square$ So far, we have looked at equilibrium of general chemical systems:
$\square$ We learned about the equilibrium constant K , reversibility, factors that affect or shift equilibrium, using ICE charts and calculating $\mathrm{K}_{\mathrm{c}}$
$\square$ Now we are going to look at equilibrium, applying the concept to Acids \& Bases
${ }^{\square}$ Calculating $\mathrm{K}_{\mathrm{a}}$ and $\mathrm{K}_{\mathrm{b}}$
$\square$ Before we begin, we better review Acids \& Bases from Grade 11!!!

## Common Properties of Acids \& Bases

| Property | Acid | Sour |
| :---: | :---: | :---: |
| Taste | Sase |  |

## Arrhenius Theory of Acids \& Bases

$\square$ Based on the ions produced when they are in water
$\square$ ACID: dissociates in water to form $\mathbf{H}^{+}{ }_{(a q)}$
$-\mathrm{HCl}_{(\mathrm{aq})} \rightarrow \mathrm{H}^{+}{ }_{(\mathrm{aq})}+\mathrm{Cl}^{-}{ }_{(\mathrm{aq})}$
$\square$ BASE: dissociates in water to form $\mathbf{O H}^{-}{ }_{(\mathrm{aq})}$
$-\mathrm{NaOH}_{(\mathrm{aq})} \rightarrow \mathrm{Na}^{+}{ }_{(\mathrm{aq})}+\mathrm{OH}^{-}{ }_{(\mathrm{aq})}$
$\square$ LIMITATIONS:
$\square$ does not explain some bases (i.e. ammonia, salt solutions)
$\square$ does not explain acid-base reactions without water (i.e. gas)

## Brønsted-Lowry Theory

$\square$ Defines acids and bases regarding protons $\left(\mathrm{H}^{+}\right)$

$$
\text { proton }=\text { nucleus of a hydrogen atom }\left(\mathrm{H}^{+} \text {ion }\right)
$$

$\square$ ACID: substance from which a proton can be removed

- "proton-donor"
$\square$ BASE: substance that can accept a proton
- "proton-acceptor"

$$
\underset{\text { acid }}{\mathrm{HCl}_{(a q)}+\mathrm{H}_{2} \mathrm{O}_{(l)} \leftrightarrow} \underset{\text { base }}{\leftrightarrow} \underset{\substack{\text { conjugate } \\ \text { acid }}}{\mathrm{H}_{3} \mathrm{O}_{(a q)}^{+}}+\underset{\substack{\text { conjugate } \\ \text { base }}}{\mathrm{Cl}_{(a q)}^{-}}
$$

## Conjugate Acid-Base Pairs

$\mathrm{NH}_{3(a q)}+\mathrm{H}_{2} \mathrm{O}_{(l)} \leftrightarrow \mathrm{NH}_{4(a q)}^{+}+\mathrm{OH}_{(a q)}^{-}$ base acid conjugate conjugate acid base
dissociation is an equilibrium reaction because it proceeds in both directions
$\square \mathrm{H}_{2} \mathrm{O}$ donates a proton in forward $r \times n \rightarrow$ acid
$\square \mathrm{NH}_{3}$ accepts a proton from $\mathrm{H}_{2} \mathrm{O} \rightarrow$ base
If a substance acts as a proton donor and a proton accepter,
it is termed "amphoteric" (i.e. water)


Fgyrin. Canjugate acid-bese pairs in the diasecintion of acetic acid in water

## Conjugate Acid-Base Pairs

Conjugate Base - The species remaining after an acid has transferred its proton.

Conjugate Acid - The species produced after base has accepted a proton.

| donation of |  |
| :---: | :---: |
| $\begin{gathered} \text { HX } \\ \text { acid } \end{gathered}$ | $\mathrm{X}^{-}$ conjugate base |
| HCl | $\mathrm{Cl}^{-}$ |
| $\mathrm{H}_{2} \mathrm{SO}_{4}$ | $\mathrm{HSO}_{4}^{-}$ |
| $\mathrm{HNO}_{3}$ | $\mathrm{NO}_{3}{ }^{-}$ |
| $\mathrm{H}_{3} \mathrm{O}^{+}$ | $\mathrm{H}_{2} \mathrm{O}$ |
| $\mathrm{HSO}_{4}^{-}$ | $\mathrm{SO}_{4}{ }^{2-}$ |
| $\mathrm{CH}_{3} \mathrm{COOH}$ | $\mathrm{CH}_{3} \mathrm{COO}^{-}$ |
| $\mathrm{NH}_{4}{ }^{+}$ | $\mathrm{NH}_{3}$ |
| $\mathrm{H}_{2} \mathrm{O}$ | $\mathrm{OH}^{-}$ |

## Amphoteric behaviour

Note: Water can act as acid or base (Amphoteric)

| Acid | Base |  | Conjugate Acid | Conjugate Base |
| :---: | :---: | :---: | :---: | :---: |
| HCl | $+\mathrm{H}_{2} \mathrm{O}$ | $\leftrightarrow$ | $\mathrm{H}_{3} \mathrm{O}^{+}+$ | $\mathrm{Cl}^{-}$ |
| $\mathrm{H}_{2} \mathrm{PO}_{4}{ }^{-}$ | $+\mathrm{H}_{2} \mathrm{O}$ | $\leftrightarrow$ | $\mathrm{H}_{3} \mathrm{O}^{+}+$ | $\mathrm{HPO}_{4}{ }^{2-}$ |
| $\mathrm{NH}_{4}{ }^{+}$ | $+\mathrm{H}_{2} \mathrm{O}$ | $\leftrightarrow$ | $\mathrm{H}_{3} \mathrm{O}^{+}+$ | $\mathrm{NH}_{3}$ |
| Base | Acid |  | Conjugate Acid | Conjugate Base |
| : $\mathrm{NH}_{3}$ | $+\mathrm{H}_{2} \mathrm{O}$ | $\leftrightarrow$ | $\mathrm{NH}_{4}{ }^{+}+$ | $\mathrm{OH}^{-}$ |
| $\mathrm{PO}_{4}{ }^{3-}$ | $+\mathrm{H}_{2} \mathrm{O}$ |  | $\mathrm{HPO}_{4}{ }^{2-}+$ | $\mathrm{OH}^{-}$ |

Identify the conjugate acid-base pairs. Connect as shown.

$\mathrm{NH}_{2}{ }^{-}+\mathrm{H}_{2} \mathrm{O} \quad \leftrightarrow \quad \mathrm{NH}_{3}+\mathrm{OH}^{-}$
$\mathrm{CN}-+\mathrm{H}_{2} \mathrm{O} \leftrightarrow \mathrm{HCN}+\mathrm{OH}^{-}$
$\mathrm{HClO}_{4}+\mathrm{CH}_{3} \mathrm{COOH} \leftrightarrow \mathrm{ClO}_{4}^{-}+\mathrm{CH}_{3} \mathrm{COOH}_{2}{ }^{+}$
$\mathrm{HCN}+\mathrm{H}_{2} \mathrm{O} \leftrightarrow \mathrm{H}_{3} \mathrm{O}^{+}+\mathrm{CN}^{-}$
$\mathrm{HSO}_{4}^{-}+\mathrm{HCl} \leftrightarrow \quad \mathrm{H}_{2} \mathrm{SO}_{4}+\quad \mathrm{Cl}-$
$\mathrm{SO}_{4}{ }^{2-}+\mathrm{HNO}_{3} \leftrightarrow \mathrm{HSO}_{4}-\quad+\mathrm{NO}_{3}-$

## Lewis Acids \& Bases

Lewis defined them as:
Acid - an electron pair acceptor
Base - an electron pair donor


Base


## Strong Acids

$\square$ Examples of Strong acids:
$\square$ Binary acids: $\mathrm{HCl}, \mathrm{HBr}, \mathrm{HI}$
$\square$ Oxoacids (contain polyatomics): $\mathrm{HNO}_{3}, \mathrm{H}_{2} \mathrm{SO}_{4}, \mathrm{HClO}_{3}, \mathrm{H}_{3} \mathrm{PO}_{4}$

Strong acids completely dissociate in water
(equilibrium favours products, lies to the right)

This means $100 \%$ of the acid will turn into ions. So if you have 0.2 M HCl , it will turn into $0.2 \mathrm{M} \mathrm{H}_{3} \mathrm{O}^{+}$
ie) $\mathrm{HCl}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{H}_{3} \mathrm{O}^{+}+\mathrm{Cl}^{-}$


$$
\begin{aligned}
& \text { isoressing acid strength } \\
& \text { isoressing acid strength }
\end{aligned}
$$

FIFIC:C The relative atrength of axsacids increases with the number of axpgen stoms.

## Strong Acids

$\square$ Strong acids dissociate 100\%
$\square \mathrm{HA}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{A}^{-}+\mathrm{H}_{3} \mathrm{O}^{+}$

Before dissociation


HA

HA

After dissociation

$\mathrm{H}_{3} \mathrm{O}^{+}$
(any strong acid) (any strong acid)

## Strong Bases

$\square$ Examples of strong bases:
$\square$ Oxides \& Hydroxides of alkali metals (Group 1) and of of alkali earth metals (Group 2) below beryllium [e.g. NaOH sodium hydroxide, MgO magnesium oxide, $\mathrm{Na}_{2} \mathrm{O}$ sodium oxide]

Strong bases completely dissociate in water (equilibrium favours products, lies to the right)

- This means $100 \%$ of the base will turn into ions. So if you have 0.2 M NaOH , it will turn into $0.2 \mathrm{M} \mathrm{OH}^{-}$
ie) $\mathrm{NaOH}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{Na}^{+}+\mathrm{OH}^{-}$
$\square$ Strong bases dissociate 100\%
$\square \mathrm{B}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{OH}^{-}+\mathrm{BH}^{+}$

Before dissociation


B
B

After dissociation

(any strong base) (any strong base)

## Calculations that involve strong acids \& bases

$\square$ Strong acids/bases (and strong electrolytes) completely dissociates into ions in water
$\square\left[\mathrm{H}_{3} \mathrm{O}\right]^{+}{ }_{(a q)}$ is equal to the [strong acid]
$\square\left[\mathrm{OH}^{-}{ }_{(\text {aq) }}\right.$ is equal to the [strong base]
$\square$ You cannot determine the concentrations of ions of weak acids/bases/electrolytes this way because they do not completely dissociate in solution (which means we will have use the concept of EQUILIBRIUM!!!)

## Example

$\square$ Find the concentration of hydronium ions in $4.5 \mathrm{~mol} / \mathrm{L} \mathrm{HCl}$

- Since HCl is a strong acid, it will completely dissociate into ions
$-\mathrm{HCl}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{H}_{3} \mathrm{O}^{+}+\mathrm{Cl}^{-}$
$-4.5 \mathrm{M} \quad 4.5 \mathrm{M}$
- So, the $\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]$is 4.5 M


## Example

$\square$ Is this solution acidic or basic: 31.9 mL of 2.75 M HCl added to 125 mL of $0.05 \mathrm{M} \mathrm{Mg}(\mathrm{OH})_{2}$ ?
$\square$ Find the moles of $\mathrm{H}_{3} \mathrm{O}^{+}$(since HCl is strong)
$\square \mathrm{HCl}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{H}_{3} \mathrm{O}^{+}+\mathrm{Cl}^{-}$
(1:1 ratio)
$\square \mathrm{n}=\mathrm{cv}=(2.75 \mathrm{M})(0.0319 \mathrm{~L})=0.0877 \mathrm{~mol}$
$\square$ Find the moles of $\mathrm{OH}^{-}\left(\mathrm{Mg}(\mathrm{OH})_{2}\right.$ is strong)
$\square \mathrm{Mg}(\mathrm{OH})_{2}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{Mg}^{2+}+2 \mathrm{OH}^{-}$(1:2 ratio)
$\square \mathrm{n}=\mathrm{cv}=(0.05 \mathrm{M})(0.125 \mathrm{~L})=0.00625 \mathrm{~mol} \times 2=0.0125$ mol of hydroxide ion
$\square$ Combine the two! $0.0877 \mathrm{~mol} \mathrm{H}_{3} \mathrm{O}^{+}-0.0125 \mathrm{~mol} \mathrm{OH}^{-}$ leaves us with $0.0752 \mathrm{~mol} \mathrm{H}_{3} \mathrm{O}^{+}$
$\square \mathrm{c}=\mathrm{n} / \mathrm{v}=0.0752 / 0.1569 \mathrm{~L}=0.479 \mathrm{M}\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]$

## Homework

p. 24 in workbook

