

## Acids and Bases

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- Properties
- Defining Acids and Bases
- pH and pOH

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## Arrhenius Definition

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Arrhenius (1884) said that acids and bases release specific ions in water:

Acids - dissociate to produce  $H^+$  ions in water

Bases - dissociate to produce  $OH^-$  ions in water

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## Bronsted-Lowry Definition

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- Bronsted and Lowry independently (1923) said that acids and bases can be thought of  $H^+$  donors and acceptors:

- Acids donate  $H^+$  ions
- Bases accept  $H^+$  ions

Water can either accept or donate a  $H^+$  ions. When water accepts a  $H^+$  ion ( $H_3O^+$ ), it is called hydronium.

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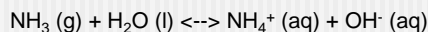
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## Conjugate Acid-Base Pairs

In acid-base equilibria, both the forward and reverse reactions involve proton transfers. In the reaction:



Because it is a reversible reaction,  $\text{NH}_4^+$  is involved in a reverse proton transfer, in which it gives up a proton.

In the reverse reactions, the products are called conjugate acid and conjugate base to identify them as reverse reactants.

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## Categories for acids-bases

- 1) Strong acids completely transfer their protons to water, the conjugate bases do not accept (or negligibly accept) protons
- 2) Weak acids partially dissociate or donate protons to solution. The weak conjugate base also partially accepts protons.
- 3) Substances with negligible acidity that contain hydrogen have strong conjugate bases.

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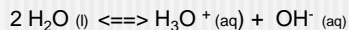
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## Self-ionization of Water

Water can self ionize, which means that if conditions are right, two molecules of water can produce a hydronium ion and a hydroxide ion:



When this happens, we can write the equilibrium constant expression, which is given a special symbol:  $K_w$

All aqueous solutions have a  $K_w = 1.0 \times 10^{-14}$

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## pH

The pH scale, designed by Sorensen, was a proposal that expresses acidity and basicity in a more compact form.

Since the molar concentration of hydronium is different in different substances, we use a scale to show this concentration.

Formula for pH:

$$\text{pH} = -\log [\text{H}_3\text{O}^+]$$

A pH of 0 is very acidic. A pH of 14 is very basic. A pH of 7 is neutral.

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## pOH

Similar to pH, except pOH is a scale to show the concentration of OH<sup>-</sup> ions in solution.

Formula for pOH:

$$\text{pOH} = -\log [\text{OH}^-]$$

Would a substance with a pOH of 6 be an acid or base? How about a pOH of 10?

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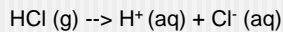
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## Strong Acids & Bases

Strong acids and bases dissociate completely in water. Therefore, the molarity of the [H<sup>+</sup>] will always be equal to the molarity of the monoprotic acid:



If the molarity of the HCl was 0.1 M, then the [H<sup>+</sup>] will also be 0.1M

What if the acid or base is a polyprotic, like H<sub>2</sub>SO<sub>4</sub>?

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## Weak Acids

Because weak acids only partially dissociate in water, the dissociation is in equilibrium, and we can write an equilibrium expression ( $K_a$ )

Ex:  $\text{HCHO}_2(\text{aq}) \rightleftharpoons \text{H}^+(\text{aq}) + \text{CHO}_2^-(\text{aq})$

$$K_a = \frac{[\text{H}^+][\text{CHO}_2^-]}{[\text{HCHO}_2]}$$

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## Weak Acids & pH

From the pH of a given concentration solution, it is possible to determine the  $K_a$  of the acid.

Ex. A prepared solution of 0.10 M formic acid,  $\text{HCHO}_2$ , has a measured pH of 2.38. What is the  $K_a$  for the acid?

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## Weak Acids & pH

Using the  $K_a$  for an acid, it is possible to determine the pH of a solution.

Ex. What is the pH of a 0.30 M acetic acid solution ( $K_a = 1.8 \times 10^{-5}$ )

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## Relative Strengths of Acids & Bases

The more readily a substance gives up a proton, the less readily the conjugate base accepts a proton.  
Or, the stronger the acid, the weaker the conjugate base.

ACID		BASE	
100 percent ionized in H <sub>2</sub> O	Strong	HCl	Cl <sup>-</sup>
		H <sub>2</sub> SO <sub>4</sub>	HSO <sub>4</sub> <sup>-</sup>
		HNO <sub>3</sub>	NO <sub>3</sub> <sup>-</sup>
		H <sup>+</sup> (aq)	H <sub>2</sub> O
		Weak	Weak
		H <sub>2</sub> SO <sub>3</sub>	HSO <sub>3</sub> <sup>-</sup>
		H <sub>3</sub> PO <sub>4</sub>	H <sub>2</sub> PO <sub>4</sub> <sup>-</sup>
		HF	F <sup>-</sup>
		HC <sub>2</sub> H <sub>3</sub> O <sub>2</sub>	C <sub>2</sub> H <sub>3</sub> O <sub>2</sub> <sup>-</sup>
		H <sub>2</sub> CO <sub>3</sub>	HCO <sub>3</sub> <sup>-</sup>
		H <sub>2</sub> S	HS <sup>-</sup>
		H <sub>3</sub> PO <sub>3</sub>	H <sub>2</sub> PO <sub>3</sub> <sup>-</sup>
		NH <sub>4</sub> <sup>+</sup>	NH <sub>3</sub>
		HCO <sub>3</sub> <sup>-</sup>	CO <sub>3</sub> <sup>2-</sup>
		HPO <sub>4</sub> <sup>2-</sup>	PO <sub>4</sub> <sup>3-</sup>
		H <sub>2</sub> O	OH <sup>-</sup>
		Weak	Weak
		HS <sup>-</sup>	S <sup>2-</sup>
		OH <sup>-</sup>	O <sup>2-</sup>
		H <sub>2</sub>	H <sup>-</sup>
		Strong	Strong
		100 percent protonated in H <sub>2</sub> O	

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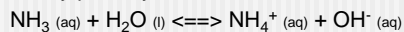
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## Weak Bases

Similar to weak acids, a weak base is a base that only partially dissociates in water, like NH<sub>3</sub>.



The base dissociation constant (K<sub>b</sub>) is calculated in a similar way to K<sub>a</sub>.

$$K_b = \frac{[\text{NH}_4^+][\text{OH}^-]}{[\text{NH}_3]}$$

From the K<sub>b</sub>, it is possible to calculate the pOH.

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## K<sub>b</sub> problems

- 1) A 0.5 M solution of methylamine, CH<sub>3</sub>NH<sub>2</sub>, has a pH of 12.2. What is the K<sub>b</sub> of the base?
- 2) A 0.5 M solution of NH<sub>3</sub> is created. The K<sub>b</sub> for ammonia is 1.8 x 10<sup>-5</sup>.
  - a) What is the OH<sup>-</sup> concentration for the solution at equilibrium?
  - b) What is the pH of the solution?

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## Weak Acid Lab Ka's

Weak Acid Formula	Ka
KHSO <sub>3</sub>	6.4 x 10 <sup>-8</sup>
HC <sub>2</sub> H <sub>3</sub> O <sub>3</sub>	1.6 x 10 <sup>-4</sup>
KHC <sub>8</sub> H <sub>4</sub> O <sub>4</sub>	3.9 x 10 <sup>-6</sup>
CH <sub>3</sub> CO <sub>2</sub> C <sub>8</sub> H <sub>8</sub> COOH	3.2 x 10 <sup>-4</sup>
KHSO <sub>4</sub>	1.0 x 10 <sup>-2</sup>
KH <sub>2</sub> PO <sub>4</sub>	6.2 x 10 <sup>-8</sup>
KHC <sub>4</sub> H <sub>4</sub> O <sub>6</sub>	4.6 x 10 <sup>-5</sup>

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## Relationship of K<sub>a</sub> and K<sub>b</sub>

When two reactions are added together to give a third reaction, the equilibrium constant of the third reaction is the product of the first two.

With acid-conjugate base pairs:

$$K_a \cdot K_b = K_w$$
$$pK_a + pK_b = pK_w$$

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## Salt solutions

Salts that react with water in solution to create H<sup>+</sup> or OH<sup>-</sup> ions undergo hydrolysis and create solutions with pHs that may be different than neutral.

The pH of a salt solution can be predicted by looking at the strength of the acids and bases that made it.

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## Rules for pH of salt solutions

- 1) Salts derived from a strong acid and a strong base is neutral.
- 2) Salts derived from a strong base and a weak acid will have  $\text{pH} > 7$ .
- 3) Salts derived from a weak base and a strong acid will have a  $\text{pH} < 7$ .
- 4) Salts derived from a weak acid and a weak base will have a pH dependent on which is greater between the  $K_a$  of the conjugate acid and the  $K_b$  of the conjugate base.

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## pH of a salt

- A solution of 0.15 M NaF is made. What is the pH of that solution?

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## Titration

Titration is a way to identify unknown concentrations of acids or bases.

In titration reactions, you neutralize an unknown acid (base) with a known concentration of base (acid). By knowing the amount of moles of base (acid) added, you can determine the moles of acid (base) neutralized.

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## Titration Curves

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Titration curves are designed to graphically represent and determine the equivalence point during a titration.

Equivalence point - point at which the  $[\text{H}_3\text{O}^+]$  is equal to the  $[\text{OH}^-]$

End point of an indicator - pH where the indicator changes color.

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