

Name \_\_\_\_\_

Date \_\_\_\_\_

## Acids & Bases

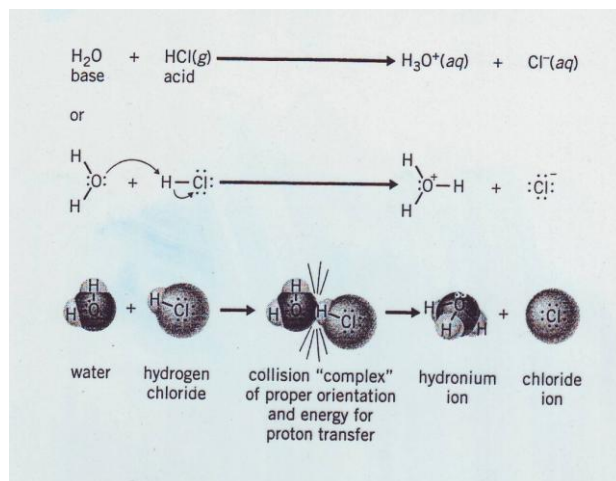
### Properties

	Acid	Base
Taste	sour/tart	bitter
Feel	burn	slippery
Metal rxn	yes	no
Litmus	red	blue
Conductivity	yes	yes

### Definitions

Arrhenius acid (1880's) –

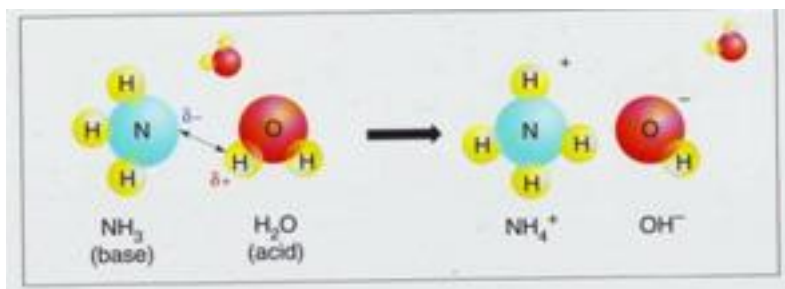
In actuality the reaction occurs by the  $H^+$  attaching itself to  $H_2O$  forming  $H_3O^+$ , the hydronium ion. You may write either symbol, but you need to know the true form.



Some common every day acids are vinegar, citric, gastric (HCl 0.1 M)

Arrhenius base -

Where does the  $OH^-$  really come from?



Some common every day bases are soap, bleach, ammonia ( $NH_3$ ), drano, detergent, milk of magnesia, antacids

**Problem:** The Arrhenius definition couldn't explain neutralization rxns like  $NH_3 + HCl \rightarrow NH_4Cl$ ,

In this rxn, all that happens is that the  $H^+$  from HCl moves to  $NH_3$  & the opposite ions attract. In effect a PROTON is DONATED

Bronsted – Lowry acid

Bronsted – Lowry base

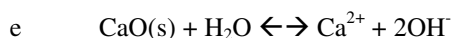
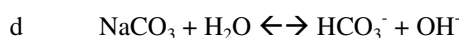
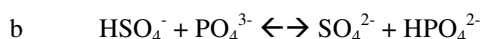
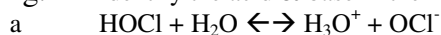
Thus according to Bronsted-Lowry – acid base reactions are proton transfer reactions.

So rxns w/ water are also acid base rxns in which  $H_2O$  acts as acid or base. *Acid Pictures powerpoint slide 3*

Eg. 1 Identify the acid and the base in the reaction  $\text{HCHO}_2 + \text{H}_2\text{O} \rightleftharpoons \text{H}_3\text{O}^+ + \text{CHO}_2^-$



Eg. 2 Identify the acid & base in the following rxns. All of these are weak acids & bases.



*The best way to look at acid base rxns is to view them as an equilibrium. Identify the acid & base in each rxn, forward & reverse. You can identify substances that are related to each other by donating or accepting a proton as a Conjugate acid-base pair. Any Bronsted-Lowry equilibrium has 2 acid-base pairs. An easy way to identify the acid member of the pair is that it has one more  $\text{H}^+$  than the base member.*

Eg 3 Write the conjugate base for nitric acid & hydrogen sulfate ion.

Eg. 4 What are the conjugate acids of  $\text{OH}^-$  &  $\text{PO}_4^{3-}$ ?

Amphoteric, amphiprotic -

## Strength

Autoionization of water –

Write  $K_c$  for autoionization of  $\text{H}_2\text{O}$

$K_w =$

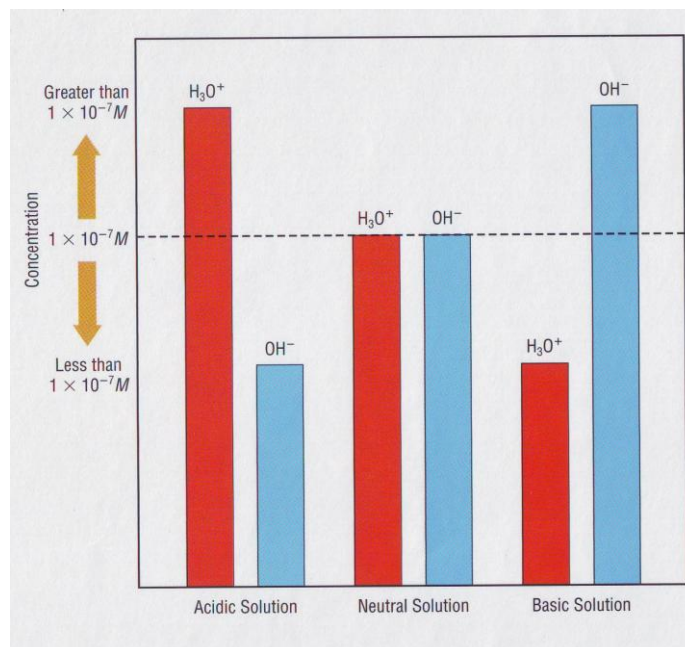
This value holds true for  $25^\circ\text{C}$ . Just like other equilibria  $K_w$  is affected by temperature

$K_w$  is also referred to as the ion-product constant of water.

A neutral solution is

An acid solution has

A basic solution has



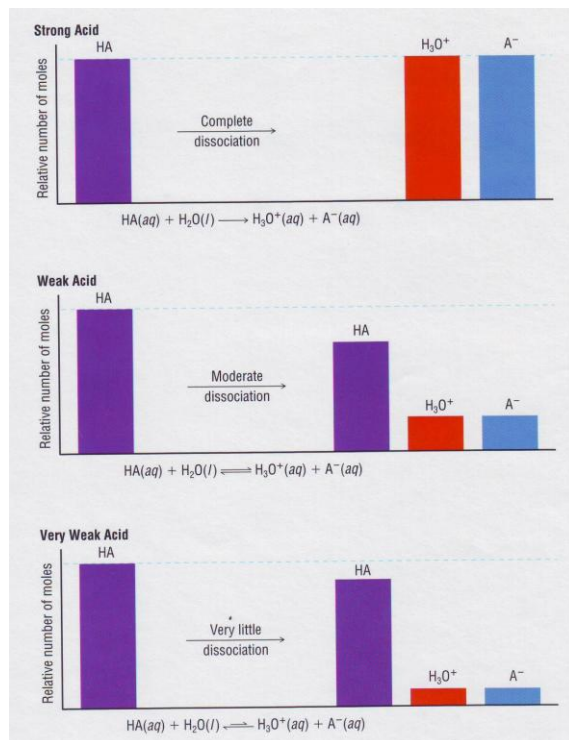
## Acid Strength

What determines the strength of an acid or base?

Strong acids & bases -

Weak acids & bases -

TABLE 14.1 in the book provides properties of weak vs strong acids



The value of K gives an indication of the relative strength of an acid. The smaller the K<sub>a</sub> value the weaker the acid.

Eg. 7 Place the following in order of acid strength from weakest to strongest:

a) HOCl, HI, NH<sub>4</sub><sup>+</sup>, HC<sub>7</sub>H<sub>5</sub>O<sub>2</sub>

b) HF, H<sub>2</sub>SO<sub>4</sub>, H<sub>3</sub>BO<sub>3</sub>, HSO<sub>4</sub>

**Using  $K_w$  to calculate  $[H^+]$  &  $[OH^-]$**

Eg. 5 In a sample of blood at 25°C the  $[H^+] = 4.6 \times 10^{-8}$  M. Find  $[OH^-]$  and decide if the sample is acidic, basic, or neutral.

Eg. 6 An aqueous solution of sodium bicarbonate has  $[OH^-] = 7.8 \times 10^{-6}$ . What is  $[H^+]$ ? Acidic, basic, or neutral?

**pH scale**

Because you take the negative log, pH decreases as  $[H^+]$  increases.

Because it's in a log scale for every 1 unit change in pH the  $[H^+]$  increases by 10 times. If the pH changes by 2 then the  $[H^+]$  changes by \_\_\_\_\_.

Write the expression for  $K_w$ .

$$pH + pOH = 14.00$$

Remember how sig figs work w/ logs.

Eg. 1 Calculate the pOH &  $[H^+]$  of a solution w/ a pH of 11.68.

Eg. 1b Calculate the pH &  $[OH^-]$  of a solution with a pOH of 3.67

## Calculating pH of strong acids

Here we will start to work on a problem solving strategy we will use for the ENTIRE unit.

Eg. 2 Calculate the pH & [OH<sup>-</sup>] of a 0.341 M HCl solution.

With all acid base problems the following steps will help the solution process

1) Identify the major species in the beaker & find their K values

2) Decide which species is contributing significantly to the [H<sup>+</sup>] of the solution & write the equation for it.

3) Determine [H<sup>+</sup>] using the significant species.

4) Calculate pH.

Eg. 3 Calculate the pH & [OH<sup>-</sup>] of a  $2.6 \times 10^{-3}$  M HNO<sub>3</sub> solution.

1)

2)

3)

4)

Eg. 4 Calculate the [H<sup>+</sup>] of a solution of HClO<sub>4</sub> solution that has a pH of 5.78.

Eg. 5 Calculate the pH & [H<sup>+</sup>] of a  $5.3 \times 10^{-10}$  M HBr solution.

Eg. 6 Calculate the pH of a solution made of  $7.2 \times 10^{-5}$  M HClO<sub>4</sub> and  $3.4 \times 10^{-4}$  HI.

## Strong Bases

Strong Bases:

Rxn w/ water

### Calculations w/ strong bases

Similar to calculations with a strong acid but w/ an added step.

Eg. 2 Calculate pH of a solution of 0.20M NaOH.

Eg. 3 Determine the pH of a solution of 0.30 M Ba(OH)<sub>2</sub>

Eg. 4 The pH of a solution of Mg(OH)<sub>2</sub> is 13.65. Determine the concentration of Mg(OH)<sub>2</sub>.

## Weak acids

Weak acids do not



Write the K expression for a weak acid.

$K_a$  = acid ionization constant

Review meaning for  $K_a$  values

### $K_a$ & pH Calculations for weak acids

A similar process is used to calculate pH of weak acids, but the addition of solving an equilibrium problem is added. *Keep in mind that many weak acids have very small  $K_a$  values, so assumptions can be made. The 5% rule for the assumptions made for small K values works because the  $K_a$  values themselves are accurate to within 5%.*

Eg. 1 A 1.0 M solution of acetic acid has a  $K_a = 1.8 \times 10^{-5}$ . Calculate the pH.

- 1) major species (what's in the beaker)
- 2) significant contributor – largest K is the most significant contributor
- 3) write the equilibrium expression for the significant contributor
- 4) using the given info set up the equilibrium problem
- 5) determine concentrations at equilibrium
- 6) Use  $K_a$  to solve for  $[\text{H}^+]$ . Remember assumptions if they apply
- 7) check 5% rule
- 8) calculate pH

Eg. 2 Lactic acid ( $\text{HC}_3\text{H}_5\text{O}_3$ ) which is present in sour milk, also gives sauerkraut its tartness. Calculate the pH of a 0.100 M solution of lactic acid at  $25^\circ\text{C}$ .

- 1) major species (what's in the beaker?)
- 2) significant contributor: largest  $K$  is the most significant contributor
- 3) write the equilibrium expression for the significant contributor
- 4) using the given info set up the equilibrium problem
- 5) determine concentrations at equilibrium
- 6) Use  $K_a$  to solve for  $[\text{H}^+]$ . Remember assumptions if they apply
- 7) check 5% rule
- 8) calculate pH

Eg. 3 Calculate the pH of a solution made of 0.60 M acetic acid & 4.3 M boric acid.

Calculate the concentration of the borate ion.



## Percent dissociation

Formula

The percent of dissociated acid in a solution increases as the solution [ ] decreases.

Why? LeChatelier:

Kinetics:

Mathematics:

*Note: As a solution becomes more dilute, it is less likely that the 5% rule will apply due to the fact that % dissociation increases as [HA] decreases.*

Eg. 4 Calculate the % dissociation of acetic acid in eg. 1 & eg. 3. Compare.

Eg. 5 An 0.400 M acid solution is 4.50% dissociated. Calculate the  $K_a$  for this acid and the pH.