



Chapter 15 Acid Base Diary

At one time acids and bases were just liquids, or solutions, with no chemical understanding. In the late 1880's a man in Sweden named Svante Arrhenius (on left) developed his Arrhenius Theory of Acids and Bases. It covers nearly 99% of all acids and 98% of all bases in chemistry. He won the Nobel Prize in Chemistry in 1903 for this. He stated that an acid was a substance that had excess hydrogen ions in aqueous solution, and that a base had excess hydroxide ions in solution.

Arrhenius Acids

A list of acids you need to be familiar with are located in Table K in your reference table. They are listed with the strongest at the top and weakest at the bottom.

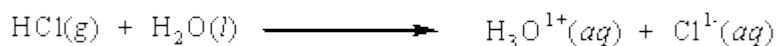
HCl hydrochloric acid	dissociates very well	LOTS of H ⁺	<p>The stronger the acid, the more H⁺ ions (and more other ions as well). The more ions, the better it conducts electricity.</p> <p>Strong acids are good electrolytes. Weak acids are poor electrolytes.</p> <p>Acetic acid is written "normally" and as an ORGANIC acid as well. Both are the same thing, but with different naming styles.</p>
HNO ₃ nitric acid	dissociates very well	LOTS of H ⁺	
H ₂ SO ₄ sulfuric acid	dissociates well	many H ⁺	
H ₃ PO ₄ phosphoric acid	dissociates less well	less H ⁺	
H ₂ CO ₃ or CO ₂ (AQ) carbonic acid	dissociates poorly	few H ⁺	
HC ₂ H ₃ O ₂ or CH ₃ COOH acetic acid or ethanoic acid	dissociates quite poorly	very few H ⁺	

Arrhenius acid is a substance that when added to water increases the concentration of H¹⁺ ions present. The chemical formulas of Arrhenius acids are written with the acidic hydrogens first. The concentration of hydrogen ions can be written in this way: [H⁺] The higher the concentration of hydrogen ions, the more acidic the solution is. A strong acid dissociates well and is a good electrolyte.



The H¹⁺ ion produced by an Arrhenius acid is always associated with a water molecule to form the hydronium ion, H₃O¹⁺(aq). Arrhenius acids are frequently referred to as proton donors, hydrogen ion donors, or hydronium ion donors, depending on whether we are trying to emphasize the species liberated by the acid (proton or hydrogen ion) or the species present in solution (hydronium ion). To represent the

transfer of the H^{1+} ion to water to form the hydronium ion, we must include H_2O in the chemical equation for acid ionization.

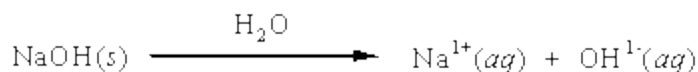


Arrhenius Bases

The "opposite" of acids are called bases. A basic solution is that any substance that produces excess hydroxide ions in solution. The list of bases in table L in your reference charts is expanded just below.

NaOH sodium hydroxide	dissociates very well	LOTS of OH^{-1}	<p>The stronger the base, the more OH^{-1} ions (and more other ions as well). The more ions, the better it conducts electricity.</p> <p>Strong bases are good electrolytes. Weak bases are poor electrolytes.</p> <p>Ammonia is a weak base & a common household chemical. The Brønsted-Lowry Theory can explain it.</p>
KOH potassium hydroxide	dissociates very well	LOTS of OH^{-1}	
$\text{Ca}(\text{OH})_2$ calcium hydroxide	dissociates somewhat	Less OH^{-1}	
NH_3 ammonia	an exception to the Arrhenius Theory, read more below.	"NO" OH^{-1}	

An **Arrhenius base** is a substance that when added to water increases the concentration of OH^{1-} ions present.



Neutralization

Arrhenius Theory also explains how and why when acids and bases combine that they can "neutralize" each other.

Acids and bases combine to ALWAYS form water + a salt.

Salts are chemical substances that are IONIC COMPOUNDS:

IONIC COMPOUNDS are: metal cation + nonmetal anion.

The neutralization reaction is summarized as:

acid + base yields water + a salt

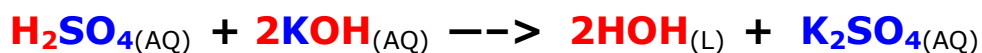
This is a type of **double replacement reaction**, as you will see below, but because it also neutralizes the acid and base, that is the appropriate name.

example reaction:



The hydrogen ions from the acid combine to the hydroxide ions in the base, and they make water. The other ions, the cation from the base (here that is the Na) plus the anion from the acid (here that is the Cl), combine to form a salt (an ionic compound, here that is table salt).

Another example could be:



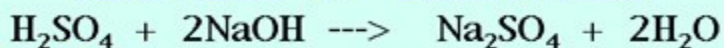
Acids have excess $[\text{H}^{+1}]$ while bases have excess $[\text{OH}^{-1}]$

When a solution has equal numbers of hydrogen and hydroxide ions, they combine PERFECTLY into water, and the combined solution is no longer acid or base, it is said to be **NEUTRAL**. The acid is neutralized by the base. The base is neutralized by the acid. There are still many ions in solution (the anions of the acid and the cations of the base), so the resulting neutral solution is still a good electrolyte.

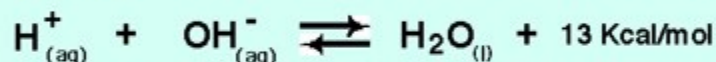
Neutralization

Acid-Base Neutralization

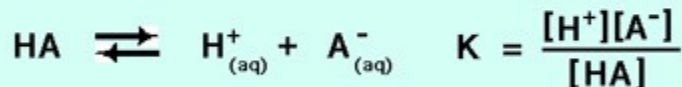
Acid + Base \longrightarrow a Salt + Water



Ionic Equations show the predominant reacting species.



Strength of Acids, the Equilibrium Constant:



A large K means a strong acid. It's highly ionized.

A Salt is a compound derived from the reaction of an acid plus a base. Or from ions from opposite sides of the periodic chart.

A Special base case: Ammonia (no apparent OH⁻¹ ions)

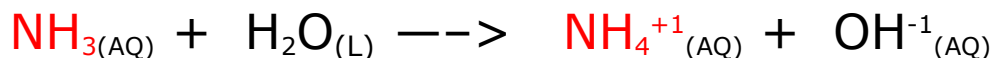
In chemistry there are many exceptions to many "rules", including the Arrhenius Theory of Acids and Bases. Ammonia is a common substance used for cleaning houses (bathrooms mostly) and also used in fertilizer production. We use the reversible reaction of ammonia synthesis when examining LeChatelier's principle, and the economics of ammonia cannot be understated. It's important stuff. It is also an exception to the Arrhenius Theory of bases.

There are NO hydroxide ions in ammonia. In fact, there are NO IONS at all. How then can it act like a base? An alternate theory, called the Brønsted-Lowry Theory describes acids and bases a totally different way than Arrhenius did. This theory is actually much more comprehensive, but it is more complicated.

The **Brønsted-Lowry (B/L)** theory states:

*B/L bases will be hydrogen ion "acceptors"
B/L acids will be hydrogen ion "donors"*

Examine this equation, and see if you can spot how ammonia "accepts" a H⁺ ion and how water "donates" it to the ammonia.

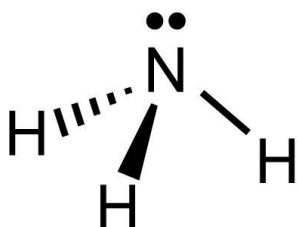


Imagine this way:
NH₃ plus H⁺¹ equals NH₄⁺¹
H₂O minus H⁺¹ equals OH⁻¹

**NH₃ becomes NH₄⁺¹ in solution (RED),
and that hydrogen ion that it gains comes from the water.**

The end result is the key, there are hydroxides in solution to provide the base properties.

Technically, by definition, ammonia is the base and the hydroxide is the conjugate.



pH scale

The measure of the concentration of hydrogen ions is called the pH.

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

which means that the pH is equal to the negative logarithm of the concentration of hydrogen ions in solution.

This scale runs from zero (very high H^+ concentration, very strong acid) up to 14 (very low H^+ concentration, very strong base).

At exactly pH 7.0 the concentration of hydrogen ions equals the concentration of hydroxide ions, and they all turn to water.

$$[\text{H}^+] = [\text{OH}^-] \text{ at NEUTRAL, neither acid or base}$$

At a pH of 5.0, there are more H^+ ions than OH^- ions in solution.

At a pH of 12.8, there are more OH^- ions than H^+ in solution.

Many solutions have equal numbers of both ions, water for example is one. Any aqueous solution with molecules (sugars or alcohols) are also neutral.

	Environmental Effects	pH Value	Examples
ACIDIC		pH = 0	Battery acid
		pH = 1	Sulfuric acid
		pH = 2	Lemon juice, Vinegar
		pH = 3	Orange juice, Soda
	All fish die (4.2)	pH = 4	Acid rain (4.2-4.4) Acidic lake (4.5)
	Frog eggs, tadpoles, crayfish, and mayflies die (5.5)	pH = 5	Bananas (5.0-5.3) Clean rain (5.6)
NEUTRAL		pH = 6	Healthy lake (6.5) Milk (6.5-6.8)
	Rainbow trout begin to die (6.0)	pH = 7	Pure water
BASIC		pH = 8	Sea water, Eggs
		pH = 9	Baking soda
		pH = 10	Milk of Magnesia
		pH = 11	Ammonia
		pH = 12	Soapy water
		pH = 13	Bleach
		pH = 14	Liquid drain cleaner

So,

Acids have $[\text{H}^+] > [\text{OH}^-]$ bases have $[\text{H}^+] < [\text{OH}^-]$

Acid Base Neutralization in Lab

To neutralize an acid with a base, or do the reverse, to neutralize a base with an acid, you have to EXACTLY match up the hydrogen ions with the hydroxide ions to balance them and end up with no excess of either. In lab we will use a buret, which is a long glass tube with MANY lines. Burets are expensive and very accurate, so always be careful handling them and reading them.

We use a buret of acid to neutralize a beaker of base.

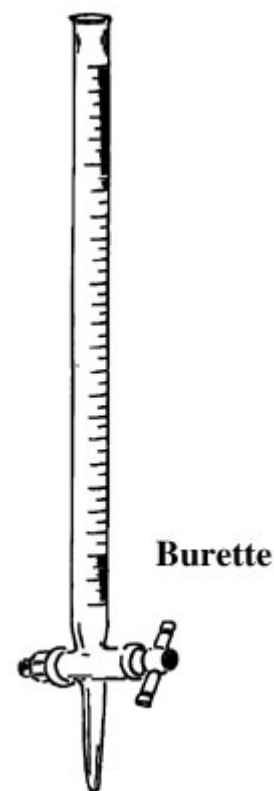
We use a buret of base to neutralize a beaker of acid.

In that photo below, the acid is RED and the base is BLUE

We read a buret by the accurately drawn lines on it. When it is "FULL", it reads 0.0 mL (none missing). When you open the valve at the bottom, solution flows out.

At the end you can read the final reading in the buret, which is always higher than when you started.

The difference between initial and final readings is how many mL of solution you ran out of the buret.



At some point you will have put in enough acid to the base, or base to the acid to perfectly neutralize the solutions. How will you know when they are at neutral?

If you add an acid base indicator into the beaker there will be a color change at neutral to show you when the hydrogen ions equal the hy-



Acid Base Titration Math

Titration is the process to measure how acidic or basic a solution of unknown concentration is. We use a known molar strength acid and drip in the unknown base until we reach the neutral point. This point is a color change due to the presence of an acid base indicator (we'll use phenolphthalein in lab).

The formula, which is on the back of your reference table is:

$$(M_A)(V_A) = (M_B)(V_B)$$

It stands for the molarity of the acid multiplied by the volume of the acid is equal to the molarity of the base multiplied by the volume of the base. Since we'll always know both volumes and the molarity of either the acid or base, we can solve for the missing molarity.

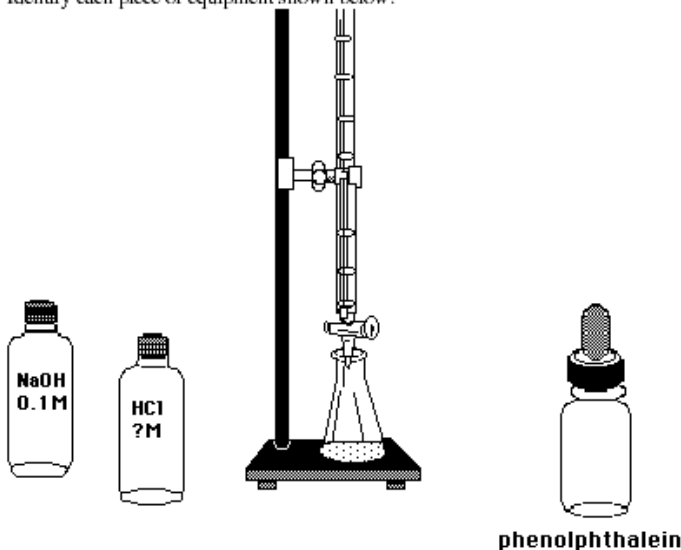
At neutral (or as close as we can get in lab) when the acid and base are neutralized, we can use our chemistry math to calculate unknown molarity of the base. Volume units can be any units, as long as they are the same on both sides of the equal sign.

An example problem:

If it takes 35.6 mL of acid of 3.40 Molarity to completely neutralize 88.5 mL of KOH, what is the molarity of the base?

$$\begin{aligned}(M_A)(V_A) &= (M_B)(V_B) \\ (3.40)(35.6) &= (M_B)(88.5) \\ \text{Solve for molarity of base} \\ 13.676\dots &= (M_B) \\ 13.7 \text{ M} &= (M_B) \text{ with 3 significant figures}\end{aligned}$$

5 Identify each piece of equipment shown below.



Unusual Acid Base Neutralization Situations

When we neutralize HCl and NaOH using the "normal" titration math formula of $M_A V_A = M_B V_B$, often we overlook the exact formula meaning. M_A means the molarity of the acid, or more importantly, the molarity of the H^+ ions.

1.0 moles of HCl have 1.0 moles of H^+ , and 0.50 moles of HCl have 0.50 moles of H^+ .

But what if it's a "double acid" or "triple acid"? H_2SO_4 has 2 H^+ ions per molecule, or two moles of H^+ ions per mole of molecules. Phosphoric acid, H_3PO_4 has 3 H^+ ions per molecule, or three moles of H^+ per mole of molecules.

The M_A in these problems needs to be looked at closely, because with H_2SO_4 and NaOH, there is NOT a 1:1 ratio of H^+ hydrogen ions to OH^{-1} hydroxide ions. This changes the math somewhat, and you need to watch for this.

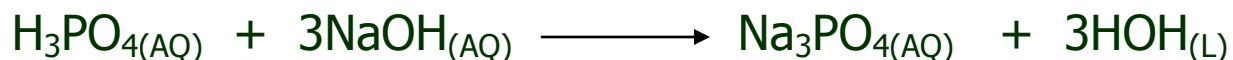
One mole of hydrochloric acid and one mole of sodium hydroxide neutralize to one mole of sodium chloride and one mole of water



Here it's clear it takes 2 moles of sodium hydroxide to neutralize one mole of the sulfuric acid, because the acid is a "double H^+ " acid rather than a single like HCl above.



Here, it takes three moles of sodium hydroxide to neutralize one mole of phosphoric acid, because the acid is a "triple H^+ " acid and the base is a "single OH^{-1} " base.



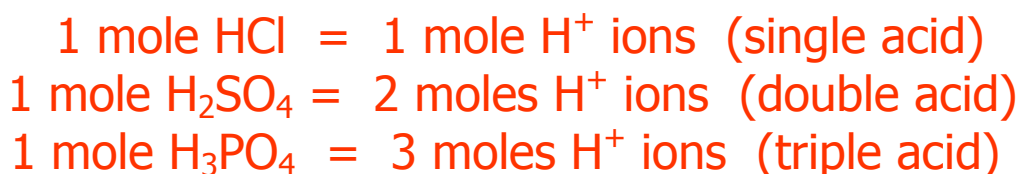
The same works in "reverse" when the acid is single, but the base is doubled.



It takes 2 moles of the "single H^+ " acid to neutralize the "double OH^{-1} " base.

The M_A or the M_B stands for the molarity of the H^+ ions,
or the molarity of the OH^{-1} ions.

They do not stand for the molarity of the molecules of acid,
or formula units of base.



Make sure you don't always "assume" that the
 $\text{H}^+:\text{OH}^{-1}$ ratio is 1:1,
because it's not and you need to adjust for that.

[Animation of Titration](#)



Use of an Indicator for Neutralization

In lab we will use an indicator in our beakers to show us with color when we have neutralized (or come very close to) our acid base combination.

There are several indicators on table M that we could use, litmus and bromthymol blue both have some color shift through the pH value of 7.0 which is neutral. Both would give us a fairly qualitative measure and would not be 100% accurate.

If we use phenolphthalein instead, which we do use, there is the clear to pink color change which is much easier to see. Of course that change happens at 8.2 rather than 7.0 pH.

We won't be "exactly" neutralizing in lab, but we'll be close enough for a high school chemistry lab. In order to be closer to true neutral we'd need to use an expensive pH meter, which we do not have (or need). Color indicators are pretty accurate, considering that they are qualitative and not quantitative.

Acid Base Indicators of Table M

On Table M there are six chemical indicators we can use to give us a handle on the approximate pH values of a solution. They will show, by color, a range of pH values that could be for a solution. They never can give an EXACT pH reading, but they can show a qualitative measure of concentration easily.

Methyl Orange is the first indicator. The table shows us that for any solution with a pH value of 3.2 or less, the solution would be red with this indicator. If the pH of the solution changed to 4.4 or higher, then the color of the solution would be yellow. Only in that 3.2 to 4.4 range of pH would the colors be a blur between red and yellow.

By using this indicator you could be able to accurately say that the solution has a pH of less than 3.2 (if it were red), or over 4.4 (if it were yellow), or between those values (if it were "orange-y" in color). This is not a quantitative measure of concentration, but it is quick and the colors are fun.

Thymol blue is the last indicator in table M. That shows as yellow if the pH of the solution is less than 8.0, or blue if it's over 9.6. Between those values the color is a blurry greenish color.

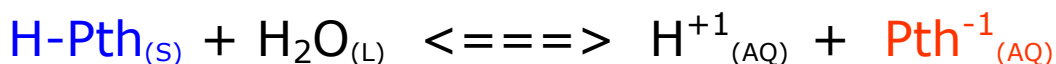
Each indicator has its own range and will be helpful at different times. There are many other acid base indicators (methyl red, indigo carmine, etc.) that we could use, but the reference table lists just six.

How do acid base indicators actually work?

Most indicators are weak acids or bases. The example I will use is phenolphthalein indicator. It has a chemical formula of: $C_{20}H_{14}O_4$ but as you will learn, these types of formulas for ORGANIC MOLECULES like this can hide what the compound really is, which is a weak acid.

The phenolphthalein molecule is colorless. When it ionizes in water it will turn pink, which is what we see. When a base is added to the solution (pH rises above 8.2 on table M), LeChatelier's predicts correctly that there will be a shift forward as these hydroxide ions combine with the loose hydrogen ions, letting more and more phenolphthalein ions in solution.

In this "equation" phenolphthalein is written as H-Pth. The H stands for the hydrogen that gets dissociated from the rest of the "Pth" phenolphthalein molecule. This puts $H^{+1} + Pth^{-1}$ ions into solution. Since it is a weak acid, not many molecules dissociate (like vinegar or acetic acid). This molecule is colorless and there are too few "Pth⁻¹" anions to pink up the solution.



"phenolphthalein" plus water in dynamic equilibrium with hydrogen ions plus "phenolphthalein ions"

add OH^{-1} shifts equilibrium forward as the hydroxides + hydrogen ions = WATER, putting more and more of the PINK "phenolphthalein ions" into the solution.

Adding more acid, or H^{+} ions shifts the equilibrium to the reverse, which lowers the amount of pink ions on the loose, making the solution colorless at LOWER pH.

As acids are added, the excess H^{+1} ions actually force the Ph^{-1} ions into reverse. Less Ph^{-1} ions means even less pink (too few to see becomes less).

When a base is added to the solution, these OH^{-1} ions immediately attach to the loose hydrogen ions, which forces the reversible reaction to shift it's equilibrium forward, putting more and more of these pink Pth^{-1} ions into solution. That gives the solution a pink color we see (at pH above 8.2, which has more and more OH^{-1} ions!).

[Titration with Phenolphthalein video](#)

Table M practice

Using table M from the reference tables, determine what color each solution should be with the specific indicator in it. Answers on the NEXT PAGE (don't peek)

pH	indicator	Color???
1.5	litmus	
2.3	thymol blue	
3.1	methyl orange	
4.7	bromthymol blue	
5.9	bromcresol green	
6.6	methyl orange	
7.0	thymol blue	
7.2	bromcresol green	
8.4	phenolphthalein	
9.6	litmus	
10.2	bromthymol blue	
11.8	phenolphthalein	
12.0	thymol blue	
14.0	phenolphthalein	

Table M practice **ANSWERS**

Using table M from the reference tables, determine what color each solution should be with the specific indicator in it. Answers on the NEXT PAGE (don't peek)

pH	indicator	Color???
1.5	litmus	red
2.3	thymol blue	yellow
3.1	methyl orange	red
4.7	bromthymol blue	yellow
5.9	bromcresol green	blue
6.6	methyl orange	yellow
7.0	thymol blue	yellow
7.2	bromcresol green	blue
8.4	phenolphthalein	hint of pink
9.6	litmus	blue
10.2	bromthymol blue	blue
11.8	phenolphthalein	pink
12.0	thymol blue	blue
14.0	phenolphthalein	pink