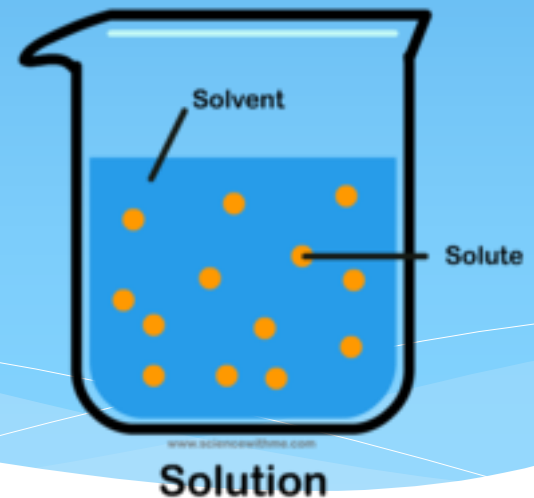


# CHEMISTRY 20

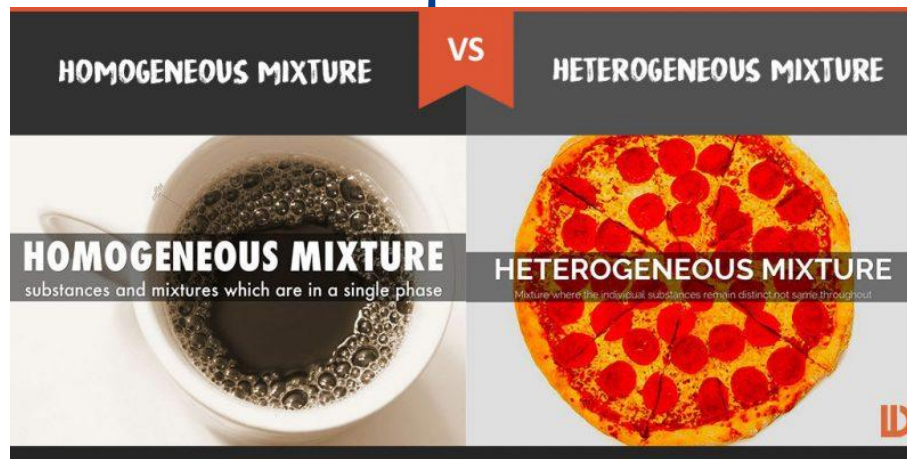
## UNIT 2: Solution Chemistry



Lesson 1  
DEFINITIONS &  
PROPERTIES OF  
SOLUTIONS

# A. Definitions

- \* Mixture: a combination of 2 or more substances
- \* Solution/Homogenous: uniform mixture where the parts are not visible
  - \* Solvent (large part - water) + Solute (small part) → Solution
  - \* Alloy: a solution of two metals like bronze
- \* Mechanical/Heterogeneous: non-uniform mixture where the parts are visible.

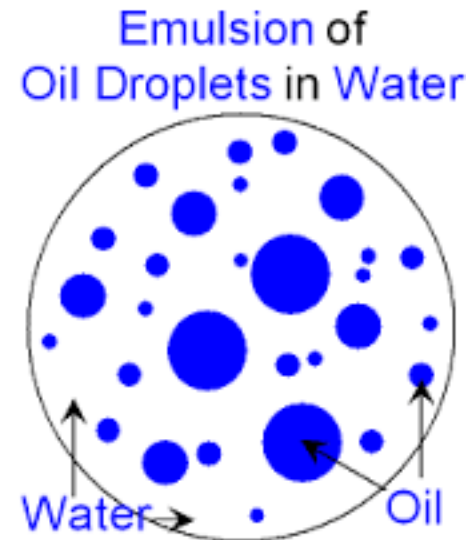
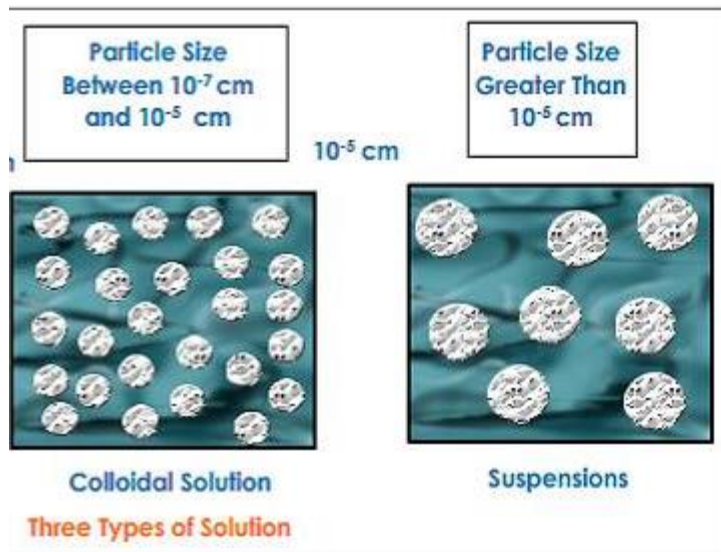


le) Coffee

ie) Pizza

# A. Definitions

- \* Suspension: a heterogeneous mixture containing large particles that settle
- \* Colloid: a heterogeneous mixture containing small particles that do not settle.
- \* Emulsion: a heterogeneous mixture made of two liquids



# A. Definitions Continued

- \* Miscible: liquids that dissolve in each other. (ethanol & water)
- \* Immiscible: liquids that are insoluble in each other - do not dissolve. (oil & water)
- \* Electrolyte: a aqueous or molten compound that conducts electricity. Only aqueous ionic compounds are electrolytes



Immiscible  
Oil & water

Miscible  
alcohol & water

## B. Solubility

Definition: Maximum amount of solute that dissolves in a given quantity of solvent at a given temperature ie) NaCl

31.6g/100mL at 0C

36.2g/100mL at 25C

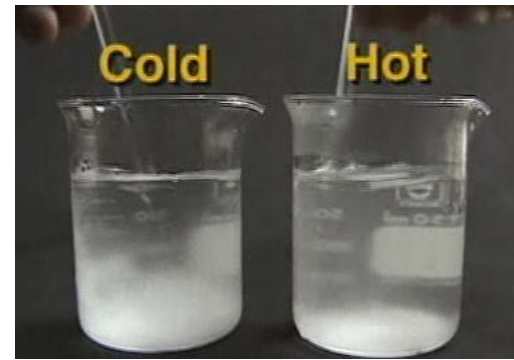
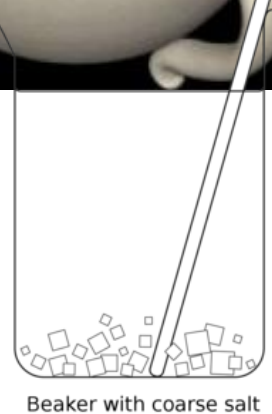
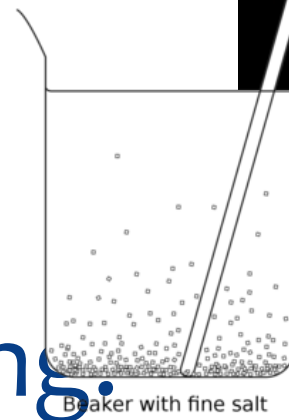
39.2g/100ml at 100C

## B. Solubility Rate depends on:

1. Agitation: shaking brings solvent into contact with solute & results in faster dissolving. (Does not affect the amount)

2. Particle size: larger surface area (powder) results in faster dissolving.

3. Temperature: higher temperature results in faster dissolving.



## B. Solubility Amount depends on

1. Temperature: the higher the temp. the greater the amount Exception: Gases

2. Pressure: the greater the pressure the greater the amount.

le) chromatography: separates compounds depending on their:

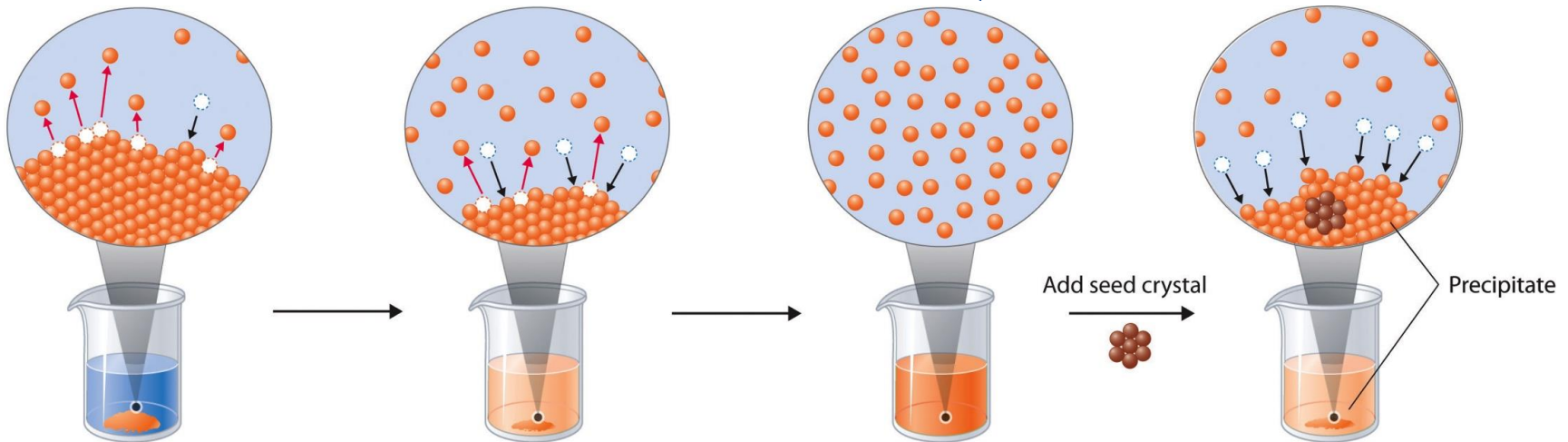
a) size (smaller particles on top)

b) solubility (most soluble parts on top)



# C. Saturation

- \* Saturated: maximum amount of solute at given conditions.
- \* Unsaturated: maximum amount of solute not reached
- \* Supersaturated: higher amount of solute than expected (heat & then



(a) Unsaturated solution

(b) Saturated solution

(c) Supersaturated solution

(d) Precipitate forms

# D. Dissolving: Solute + Solvent

Polar dissolves polar (not a chemical reaction)

## 1. Molecular compound



+ve hydrogen from water surrounds the oxygens

## 2. Ionic compound



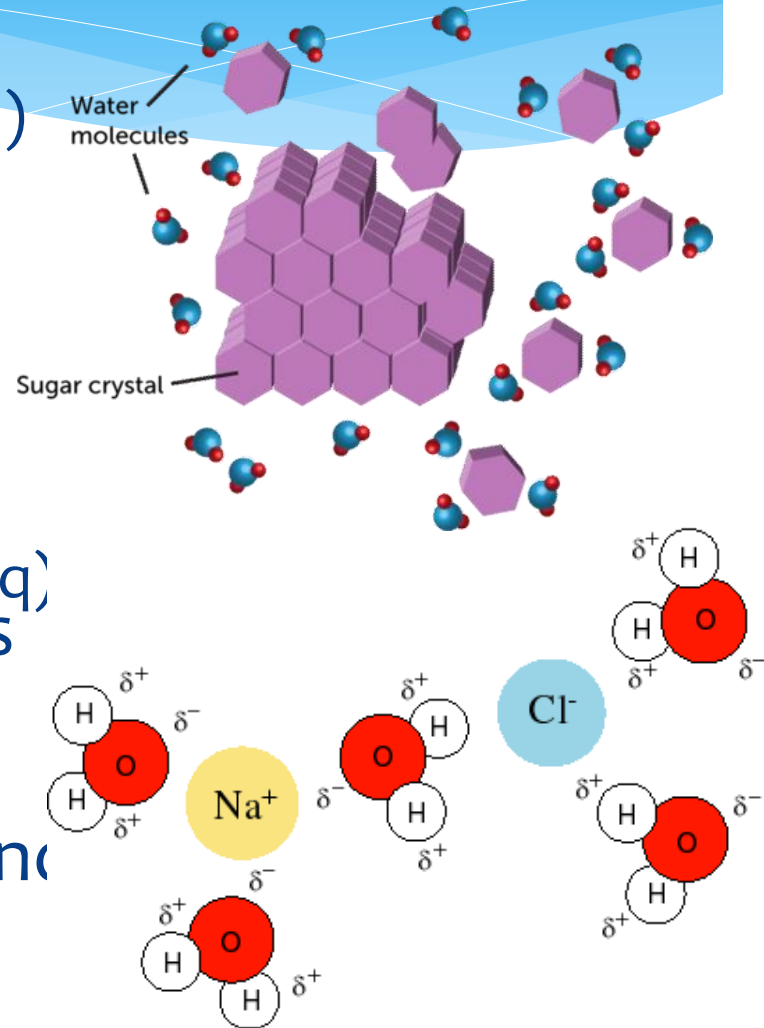
-ve oxygen from water surrounds

the positive sodium ion.

+ve hydrogen from water surround

The negative chloride ion.

How Sugar Dissolves in Water



Lesson 2  
MOLAR  
CONCENTRATION

## A. Definitions

The moles of solute in a given volume of solution (liters)

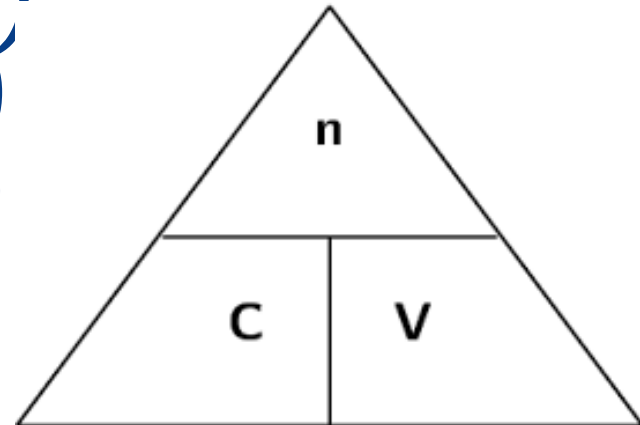
- \* Symbol is  $C$  or  $[ ]$
- \* Unit is mol/L or  $M$

\* Formula:  $C = n/V$  ( $n = CV$ )

$C$  = concentration (mol/L)

$n$  = moles of solute (mol)

$V$  = volume of solvent (L)



## B. Examples

1. Calculating concentration from moles and volume.

\* What is the concentration of 0.100 mol of 1.00 L solution

$$C = n/V$$

$$C = 0.100 \text{ mol}/1.00\text{L}$$

$$\underline{C = 0.100 \text{ mol/L}}$$

## B. Examples

2. Calculating concentration from mass and volume.

\* What is the concentration of 40.0g of NaOH in 1000 mL of water

$$\text{Step 1) } n = m/M \qquad n = 40.0\text{g}/40.00\text{g/mol}$$

$$n = 1.00 \text{ mol}$$

$$\text{Step 2) } C = n/V \qquad C = 1.00\text{mol}/1.00\text{L}$$

$$\underline{C = 1.00 \text{ mol/L}}$$

## B. Other forms of concentration

a)  $\text{ppm} = \frac{\text{mass solute}}{\text{mass solvent}} \times 10^6$

ie)  $2.00\text{E-}3\text{g}$  of Pb in  $1.0\text{kg}$  of water =  $2\text{ppm}$

NOTE:  $1.0\text{ mL}$  of water =  $1.0\text{ g}$

b)  $\% \text{ mass} = \frac{\text{mass solute}}{\text{mass solute} + \text{solvent}} \times 100$

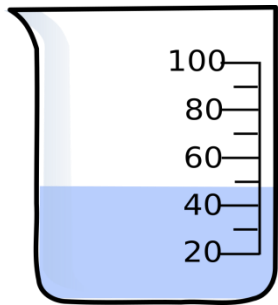
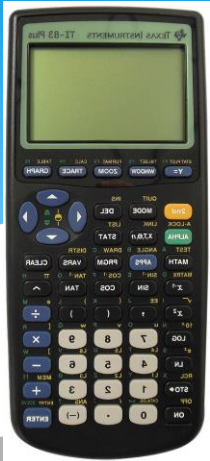
ie)  $0.84\text{g}$  of tin in a  $7.0\text{g}$  loonie =  $12\%$  tin

c)  $\% \text{ volume} = \frac{\text{volume solute}}{\text{volume solution}} \times 100$

ie)  $1000\text{mL}$  of  $2\%$  milk has  $20\text{ mL}$  of fat

## B. Steps to prepare a solution

- 1) Calculate the moles of solute needed. Use  $n=CV$
- 2) Calculate & measure (with a scale) the mass of solute needed. Use  $m=nM$
- 3) In a beaker, mix solute in about half of the solvent.





## B. Steps to prepare a solution

4) Pour into a volumetric flask with the desired volume. Add more solvent (water). Use an eye dropper to add solvent to the calibration line.



5) Cap the flask, invert and mix

## B. Example

Prepare 100 mL of 0.400 M  
magnesium sulfate

1) Calculate moles.  $n = CV$

$$n = 0.100 \text{ L} \times 0.400 \text{ mol/L}$$

$$n = 0.0400 \text{ mol}$$

2) Calculate mass. Use scale  $m = nM$

$$m = 0.0400 \text{ mol} \times 120.338 \text{ g/mol}$$

$$m = 4.82 \text{ g (3 sig figs)}$$

Weight in with a scale

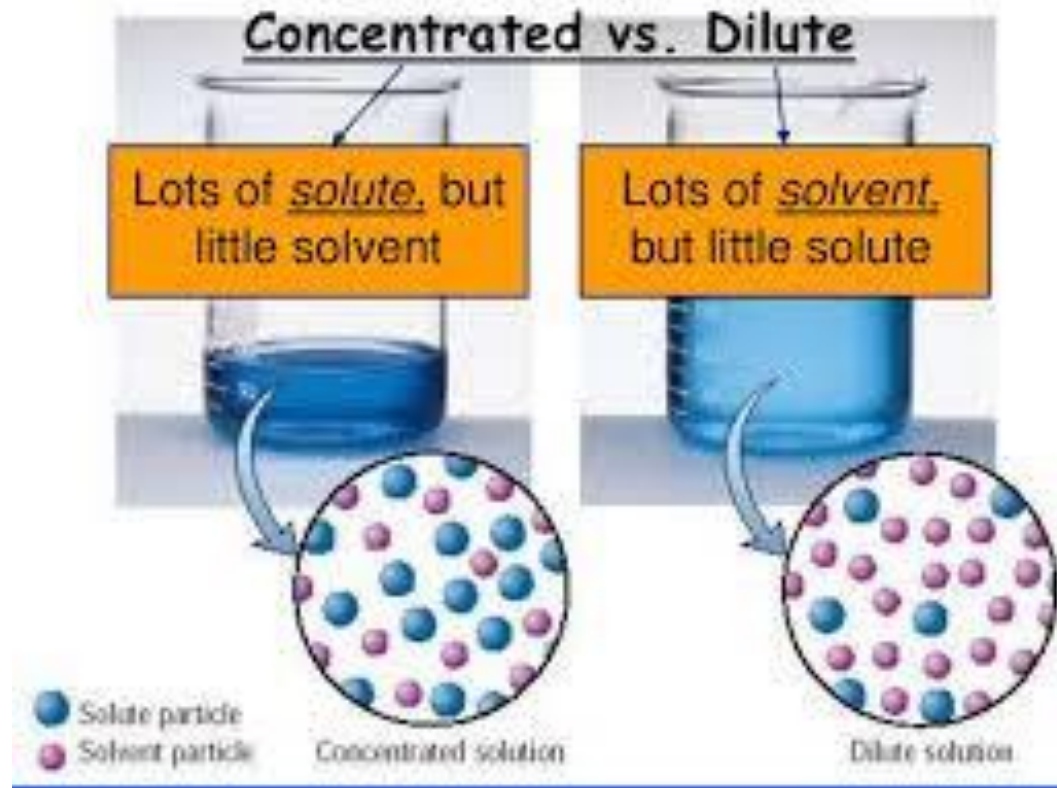
## B. Example

- 3) In 100mL beaker, mix 4.82 g in about 50 mL of water.
- 4) Pour into a 100 mL volumetric flask and fill to the calibration line. Use eye dropper near the end.
- 5) Cap the flask, invert and mix

# Lesson 3

## DILUTIONS

- \* **Dilution:** process of decreasing the concentration of a solution by adding more solvent.
- \* **Concentrated:** a solution containing a large amount of solute when compared to another solution
- \* **Dilute:** a solution containing a small amount of solute when compared to another solution.



## B. Mathematical expression

$$n_1 = n_2; \quad (n = CV)$$

$$C_1 V_1 = C_2 V_2$$

HINTS:  $V_2 = V_1 + V_{\text{water}}$

1 refers to concentrated, original or pure solutions

2 refers to diluted or prepared solutions

$C_1$  is larger than  $C_2$

$V_1$  is smaller than  $V_2$

## C. Examples

Steps to a dilution

- 1) Find volume  $V_1 = C_2 V_2 / C_1$
- 2) Remove volume with a volumetric or graduated pipet
- 3) Place in a \_\_\_ mL volumetric flask and fill with water
- 4) Cap, Invert & Mix

## C. Example 1)

How do you prepare 100mL 0.40 M magnesium sulphate from 100mL of 2.0M solution

$$1) V_1 = C_2 V_2 / C_1$$

$$V = 100\text{mL} \times 0.40\text{M} / 2.0\text{M}$$

$$V = 0.020\text{L} \text{ or } 20 \text{ mL}$$

2) Remove 20 mL with pipet

3) Fill volumetric flask with 80 mL of water

4) Cap, Invert & Mix



## C. Example 2)

If a student begins with 1000 ml of 1.00 M solution and dilutes it to 0.100 M, what volume of water was added?

$$1) V_2 = C_1 V_1 / C_2$$

$$V = 1.0 M \times 1.000 L / 0.10 = 10.0 L$$

$$\mathbf{V_{water} = V_2 - V_1}$$

$$= 10.0 - 1.000 = 9.0 L$$

## C. Example 3)

If 100 mL of 1.0 M solution is added to 900 mL of 0.10 M solution what is the new C

$$C = n_{\text{total}}/V_{\text{total}}$$

$$= \frac{(0.1\text{L} \times 1.0\text{M}) + (0.9\text{L} \times 0.1\text{M})}{(0.100 + 0.900)}$$

$$= 0.19 \text{ M}$$

NOTE: Cannot add concentrations

## C. Example 4)

If 100 mL ( $V_w$ ) of water was added to 1.0 M solution and the resulting solution was 0.250 M.

What is the initial volume? 1.00 L solution was made

$$V_1 = C_2 V_2 / C_1$$

$$V_1 = 0.250 \times 1.00 \text{ L} / 1.0 \text{ M}$$

# Lesson 4

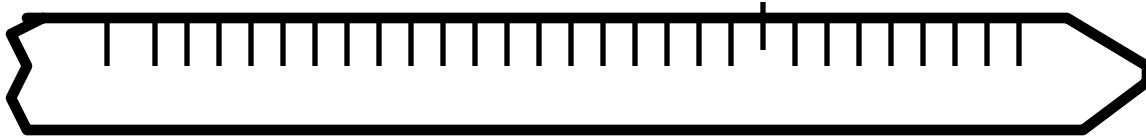
# Pipetting

## A. Definitions

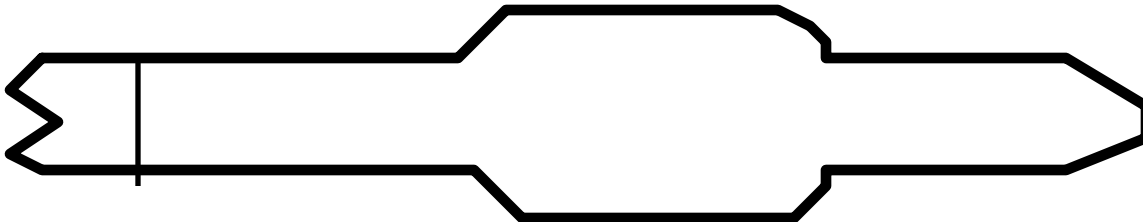
Pipetting: Technique used to measure out a small accurate volume of liquid (25 mL or less) to high precision (0.1 mL to 0.01 mL)

## B. Types of pipets

- \* graduated: Measures a range of volumes from 0.1 to 10 mL



- \* volumetric: delivers one specific volume accurate to within 0.04 mL



## C. Technique

Step 1: Rinse with distilled water to remove any residue or liquid

Step 2: Rinse with sample

**WHY?** Distilled water lowers the concentration

## C. Technique

Step 3: Hold the pipet near the top with one (left) hand leave your index finger free

**WHY?** more accurate, easier to read





## C. Technique

Step 4: Place the pipet into the sample, resting the tip on the bottom of the container

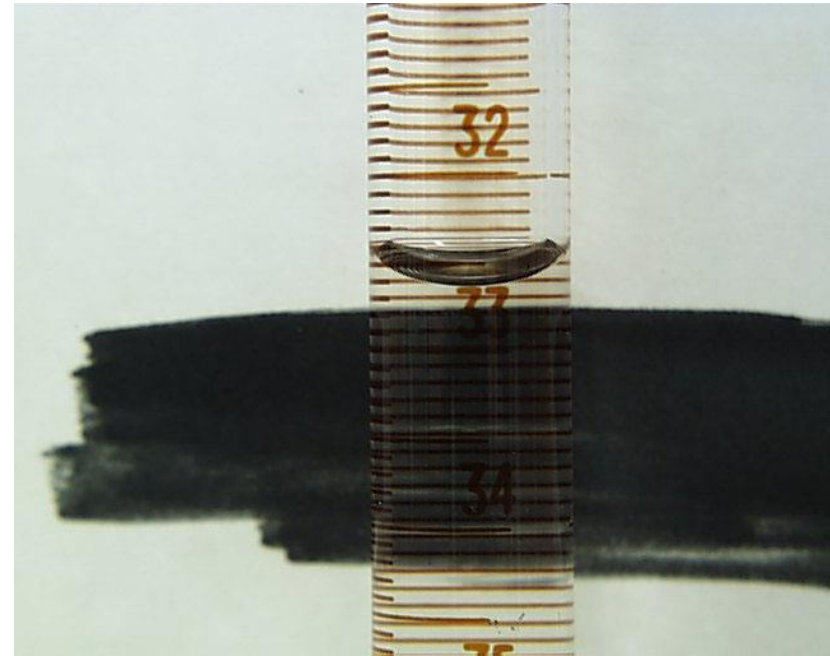
Step 5: Squeeze the bulb with your right hand and place the bulb/dispenser firmly and squarely on the end of the pipet



## C. Technique

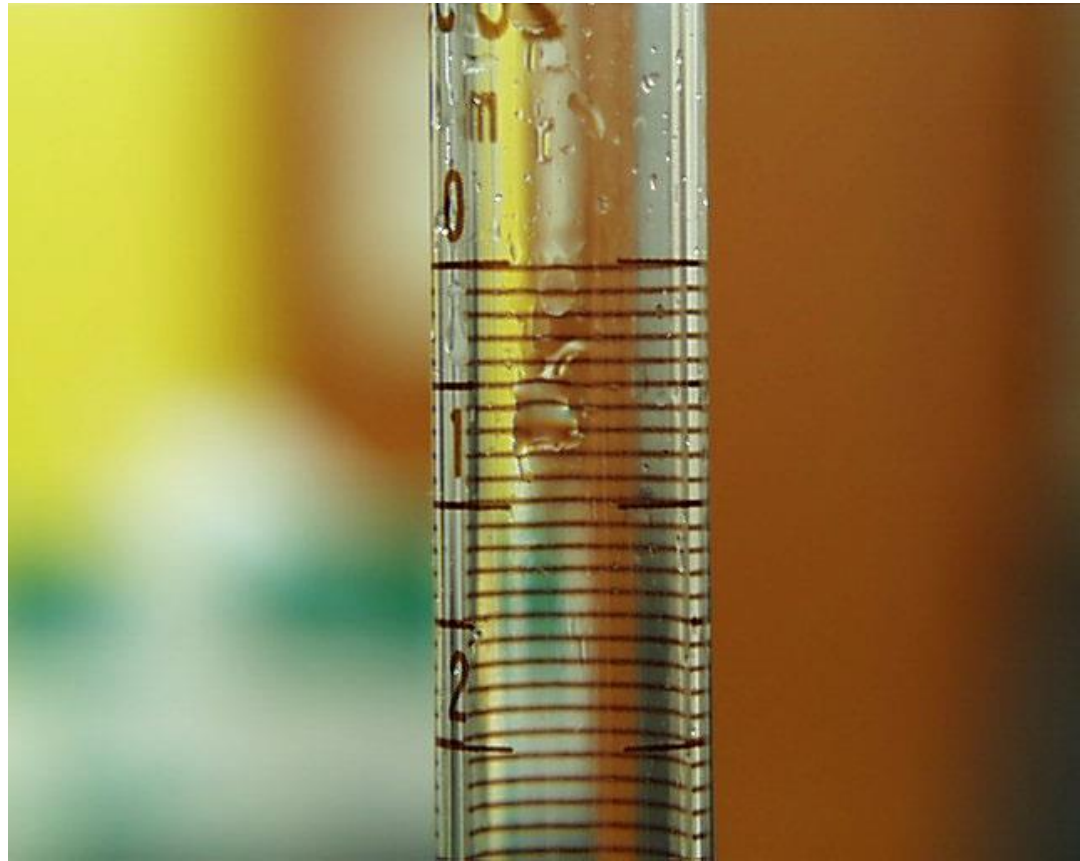
Step 6: Release the bulb until the liquid has risen above the desired calibration line

Step 7: Remove the bulb and quickly put your index finger of your left hand over the top



## C. Technique

Step 8: remove from the solution and wipe excess solution. **WHY?** Changes Volume

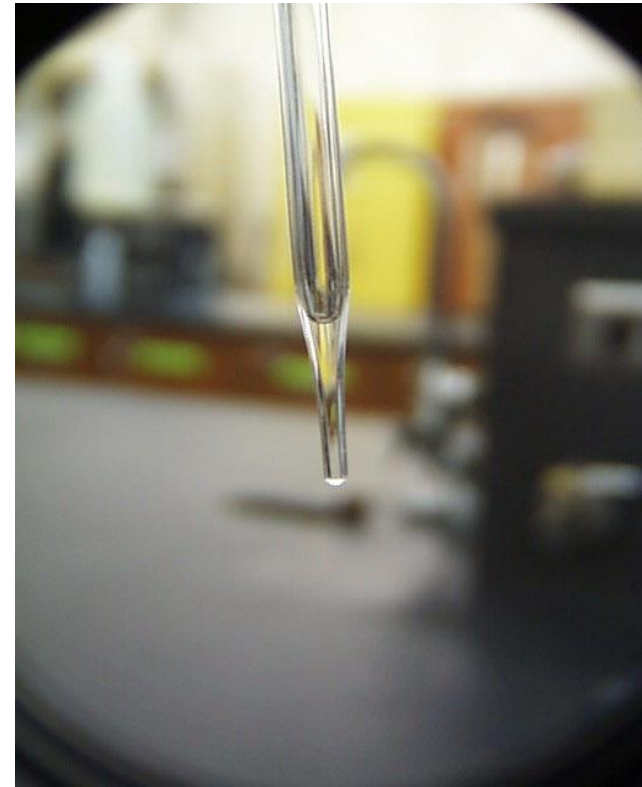


## C. Technique

Step 9: while touching the tip of the pipet to the inside of the waste beaker, gently roll your index finger off to allow the liquid layer to drop. Stop when meniscus reaches the calibration line. (at eye level)

## C. Technique

Step 10) Touch the tip of the pipet to the inside wall of the receiving container and remove finger to allow liquid to flow freely. A small volume is expected to stay in the tip.



# Lesson 5

## Dissociation

# A. Solubility Table

\* Look for the ions in row 1 of your solubility table.

A) if the other ion is in row 2 it is soluble and is aqueous (aq) in the presence of water

B) If the other ion is in row 3 or is not on the table it will form a precipitate (non-soluble) (s)

## A. Other terminology

- \* Hydronium ion:  $\text{H}_3\text{O}^+_{(\text{aq})}$  formed when water gains a  $\text{H}^+_{(\text{aq})}$  (acidic)



- \* Solvation: A process that occurs when an ionic solute dissolves; in solution, the ions are surrounded by solvent molecules
- \* Hydration: addition of water



## B. Dissociation & Ionization

Definition: Separation of an compound into individual ***ions*** when placed in a solvent

\* ie)  $\text{NaOH}_{(s)}$  ***dissociates*** into  $\text{Na}^+_{(aq)} + \text{OH}^-_{(aq)}$



\* NOTE:  $\text{NaOH}$  ***decomposes*** into Na,  $\text{O}_2$  and  $\text{H}_2$  elements



## B. Dissociation & Ionization

- \* **Electrolyte:** a compound that conducts electricity (separates into ions) ionic compounds and acids
- \* **Non- electrolyte:** compounds that do not conduct electricity

## B. Dissociation & Ionization

- \* **Dissociation equations:** breaking of ionic compounds into ions
- \* **Ionization equations:** breaking of molecular gases and acids into ions when water is added

## B. Dissociation & Ionization

### CAUTIONS:

- \* Write a balanced equation.
- \* Remember all ions are (aq).
- \* Show correct ionic charges.

# B. Dissociation & Ionization

## EXAMPLES

### \* a) Ionic compounds



Atoms must be conserved. The net charge must be zero.

## B. Dissociation & Ionization

### b) Molecular Gases



### c) Bases - the only one



## B. Dissociation & Ionization

### \*d) Acids



Only the six following acids completely dissociate: HCl, HBr, HI, HNO<sub>3</sub>, H<sub>2</sub>SO<sub>4</sub>, HClO<sub>4</sub>

**Strong Acid - completely dissociates**



**Weak Acid - do not completely dissociate**



## C. Equilibrium

- \* Definition: a state of a closed system in which all measurable properties are constant.
- \* There is a dynamic equilibrium that exists in saturated solution since the rate of crystallization is equal to the rate of dissolving. You observe no change
- \* l.e)  $\text{NaCl}_{(s)} \rightleftharpoons \text{NaCl}_{(aq)}$
- \* rate of salt crystals forming = rate of salt dissolving



# Lesson 6

## Net Ionic Equations

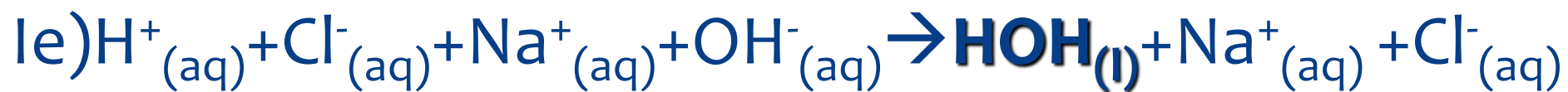
# A. Definitions

Non Ionic equation - an equation without any ions



## A. Definitions

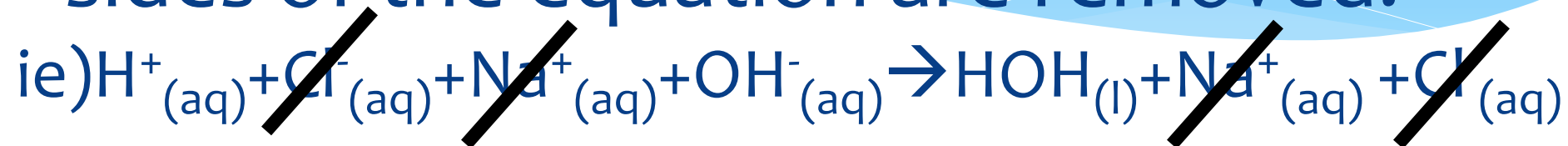
Total Ionic equation - an equation where only the aqueous solutions are broken into their ions.



NOTE: liquids, solids and gases do not break into ions

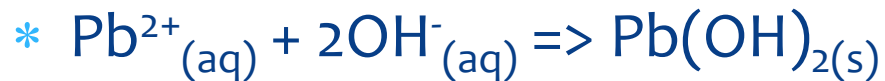
# A. Definitions

Net Ionic equation - a summary where spectator ions that appear on both sides of the equation are removed.



# A. Example

Lead(II)nitrate & sodium hydroxide react.



### B. Purpose of ionic equations

- \* A net ionic equation is a summary of what occurs. Most reactions do not occur unless they are in water.
- \* A net ionic equation is used when you want to know the amount (concentration) of an ion.

## B. Steps to write ionic equations

1. Write the non ionic equation
2. Write the total ionic equation. Do not change solids, gases or liquids.
3. Write the net ionic equation - cross out ions that repeat & reduce

# Lesson 7

# Solution Stoichiometry



# STEPS

1. Write a balanced equation-non ionic, dissociation or net ionic
2. Find the moles of known using one of the 4 formulas (if 2 reactants;  $\times 2$ )
3. Multiply by mole ratio ( R/G)
4. Change to desired unit using one of the 4 formulas
5. Calculate %yield &/or %error

## Example #1

If 200 mL of NaOH reacts with 100mL of 0.150 M HCl what is the concentration of the NaOH?



?  $n = CV = 0.100\text{L} \times 0.150\text{M}$

$n = 0.0150\text{ mol}$

$0.0150 \times 1\text{mol}/1\text{mol} = 0.0150\text{mol}$

$C = n/V = 0.0150\text{mol}/0.200\text{L} = \mathbf{0.0750\text{ mol/L}}$

## Example #2

In the previous question, what is the concentration of the solution produced?



\*  $n = CV \quad ?$

$$n = 0.0150 \text{ mol}$$

\*  $0.0150 \times 1 \text{ mol} / 1 \text{ mol} = 0.0150 \text{ mol}$

\*  $C = n/V = 0.0150 \text{ mol} / 0.300 \text{ L} = \mathbf{0.0500 \text{ mol/L}}$   
(total volume)

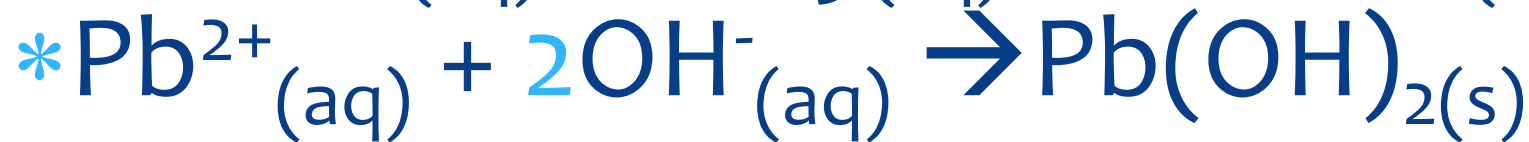
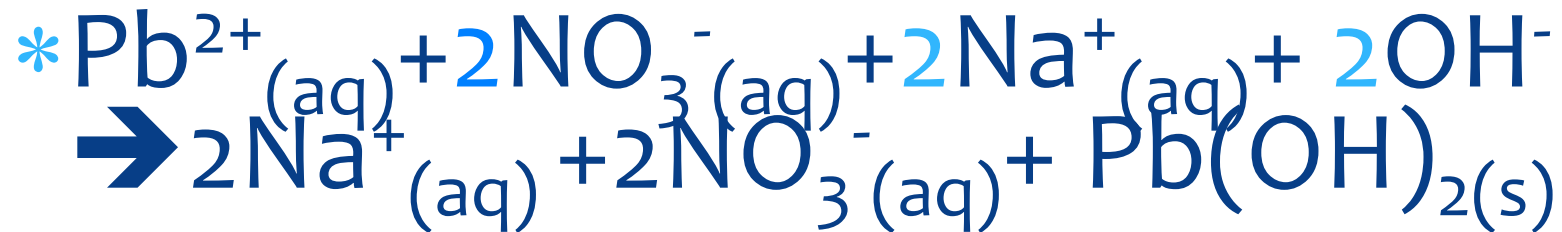
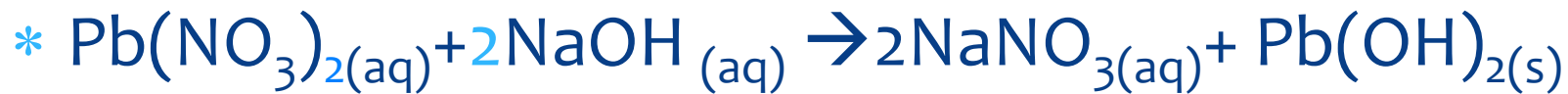
## Example #3

What is the concentration of ammonium ions if there is 100 g of ammonium sulphate solid placed in 100 mL of water?

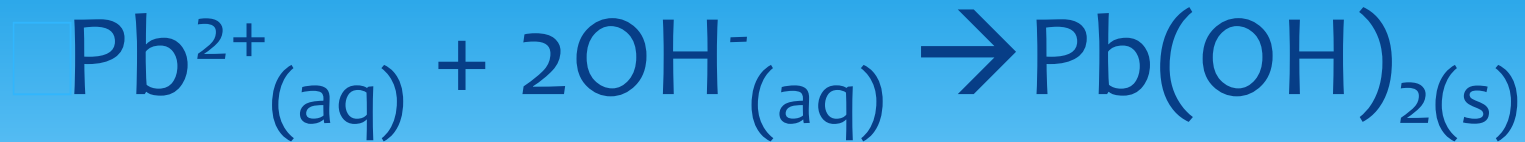
1.  $(\text{NH}_4)_2\text{SO}_4(s) \Rightarrow 2\text{NH}_4^+(aq) + \text{SO}_4^{2-}(aq)$
2.  $n = m/M$  ?  
 $n = 100/132.16 = 0.756658\dots\text{mol}$
3.  $0.756\text{mol} \times 2\text{mol}/1\text{mol} = 1.513\text{mol}$
4.  $C = n/V = 1.513\text{mol}/0.100\text{L} = 15.1\text{ M}$

## Example #4

Lead (II) nitrate solution reacts with sodium hydroxide. Using net ionic equation, how many grams of solid form if  $6.02 \times 10^{23}$  particles of OH ion are present?



## Example #4 con' t



$n = p/P$                       ? g

$n = 1.00 \text{ mol}$

$1.00 \text{ mol} \times 1 \text{ mol} / 2 \text{ mol} = 0.500 \text{ mol}$

$m = nM = 0.500 \text{ mol} \times 241.21 \text{ g/mol}$

**$m = 121 \text{ g}$**

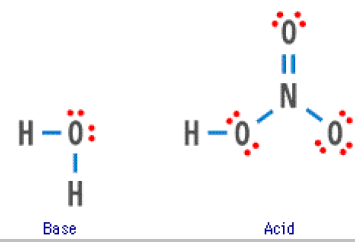
# Chemistry 20: Acid and Bases



# I) Definitions



# Definitions - Acid



#1) Arrhenius def: substance that ionizes to form hydrogen ions

MODIFIED-forms hydronium ions

(H<sub>3</sub>O<sup>+</sup>)

Bronstead Lowry:

Donates hydrogen

(proton)

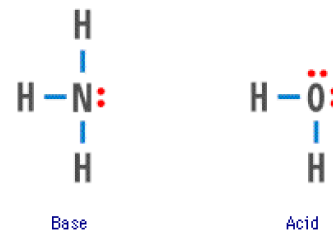


## Definitions - Acid Con't

#2) Operational def: turns blue litmus paper red & has a pH below 7

Bromothymol blue turns yellow

# Definitions - Base



#1) Arrhenius def: substance that dissociates to form hydroxide ions

Bronstead Lowry: accepts hydrogen ions (protons)

#2) Operational def: turns red litmus paper blue & has a pH above 7

Bromothymol blue turns green

# Definitions - neutral

#1) Arrhenius def: substance that forms equal amounts of hydroxide and hydronium ions; Bronstead Lowry def: does not donate or accept protons

#2) Operational def: red litmus & blue litmus do not change; pH = 7 (distilled water);

bromothymol blue turns green

## Definitions - solutions

**Acid Solutions:** A solution where the hydronium concentration is greater than the hydroxide concentration.

**Basis Solutions:** A solution where the hydroxide concentration is greater than the hydronium concentration

## Definitions - indicators

- \* **Indicator:** a chemical (weak acids) substance that changes color when added to an acid or base. (pg. 10 of databook)
- \* **PURPOSE:** determine the approximate pH range of a solution
- \* **Common indicators:**
  - \* Bromothymol blue:  $\text{HBb}(\text{aq})/\text{Bb}^-(\text{aq})$
  - \* Phenolphthalein:  $\text{HPh}/\text{Ph}^-(\text{aq})$

# Use of the Indicator table

- Bromothymol blue is an ideal indicator in its pH range, 6.0-7.6 where it is yellow + blue or green.
- Below 6.0 HBb is yellow
- Above 7.6 HBb is blue.
- Indicators can be combined to narrow the pH.
- Don't worry about  $K_a$  until Chemistry 30

Acid-Base Indicators at 298.15 K

Indicator	Suggested Abbreviations	pH Range	Colour Change as pH Increases	$K_a$
methyl violet	HMv(aq) / Mv <sup>-</sup> (aq)	0.0 – 1.6	yellow to blue	$-2 \times 10^{-1}$
cresol red	H <sub>2</sub> Cr(aq) / HCr <sup>-</sup> (aq)	0.0 – 1.0	red to yellow	$-3 \times 10^{-1}$
	HCr <sup>-</sup> (aq) / Cr <sup>2-</sup> (aq)	7.0 – 8.8	yellow to red	$3.5 \times 10^{-9}$
thymol blue	H <sub>2</sub> Tb(aq) / HTb <sup>-</sup> (aq)	1.2 – 2.8	red to yellow	$2.2 \times 10^{-2}$
	HTb <sup>-</sup> (aq) / Tb <sup>2-</sup> (aq)	8.0 – 9.6	yellow to blue	$6.3 \times 10^{-10}$
orange IV	HOr(aq) / Or <sup>-</sup> (aq)	1.4 – 2.8	red to yellow	$-1 \times 10^{-2}$
methyl orange	HMo(aq) / Mo <sup>-</sup> (aq)	3.2 – 4.4	red to yellow	$3.5 \times 10^{-4}$
bromocresol green	HBg(aq) / Bg <sup>-</sup> (aq)	3.8 – 5.4	yellow to blue	$1.3 \times 10^{-5}$
methyl red	HMr(aq) / Mr <sup>-</sup> (aq)	4.8 – 6.0	red to yellow	$1.0 \times 10^{-5}$
chlorophenol red	HCh(aq) / Ch <sup>-</sup> (aq)	5.2 – 6.8	yellow to red	$5.6 \times 10^{-7}$
bromothymol blue	HBb(aq) / Bb <sup>-</sup> (aq)	6.0 – 7.6	yellow to blue	$5.0 \times 10^{-8}$
phenol red	HPr(aq) / Pr <sup>-</sup> (aq)	6.6 – 8.0	yellow to red	$1.0 \times 10^{-8}$
phenolphthalein	HPh(aq) / Ph <sup>-</sup> (aq)	8.2 – 10.0	colourless to pink	$3.2 \times 10^{-10}$
thymolphthalein	HTh(aq) / Th <sup>-</sup> (aq)	9.4 – 10.6	colourless to blue	$1.0 \times 10^{-10}$
alizarin yellow R	HAY(aq) / Ay <sup>-</sup> (aq)	10.1 – 12.0	yellow to red	$6.9 \times 10^{-12}$
indigo carmine	Hlc(aq) / Ic <sup>-</sup> (aq)	11.4 – 13.0	blue to yellow	$-6 \times 10^{-12}$
1,3,5-trinitrobenzene	HNb(aq) / Nb <sup>-</sup> (aq)	12.0 – 14.0	colourless to orange	$-1 \times 10^{-13}$

# Definitions - pH & pOH

**pH:** Power/parts of Hydrogen  
(hydronium ion/proton) measured as a  
negative log of the hydronium  
concentration.  $[\text{H}_3\text{O}^+]$ ; no units

**pOH:** Parts of Hydroxide ion

Measured as a negative log of the  
hydroxide concentration.  $[\text{OH}^-]$ ; no  
units



# Definitions - pH meter

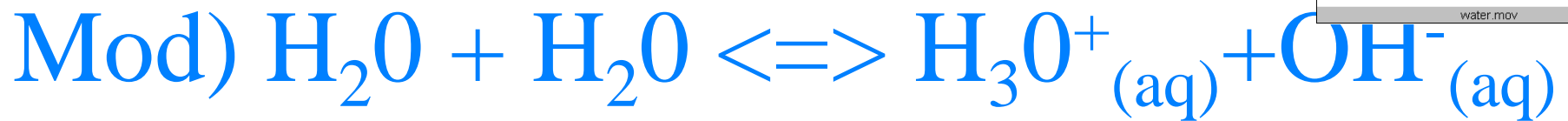
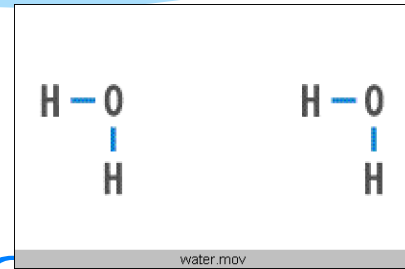
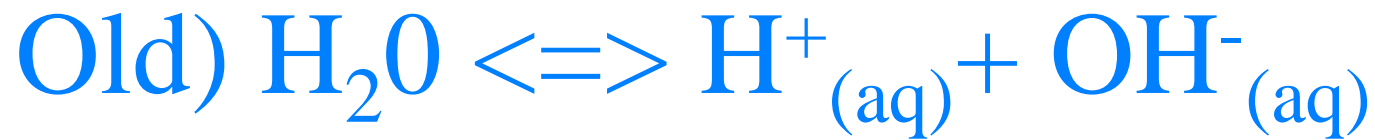
pH meter: instrument used to measure the pH

dynamic equilibrium: a balance between the forward and reverse processes to achieve a steady state

Ie) self-ionization of water (next slide)

# Definitions: self-ionization of water

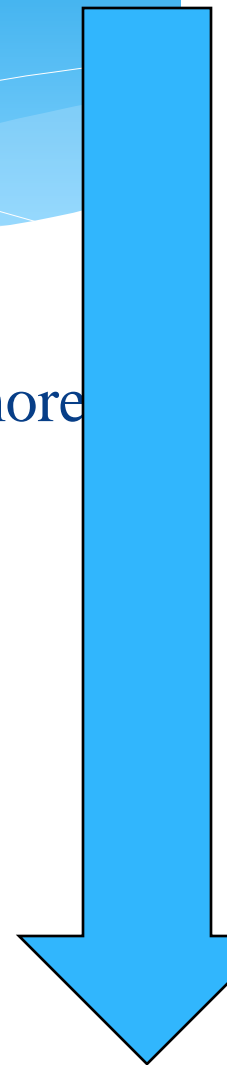
self-ionization: when two water molecules react & break up into ions



NOTE: fewer than 2 in one billion ionize

# 7 acids completely ionize(p. 8)

- \* perchloric acid( $\text{HClO}_4$ ) – Strongest Acid
  - \* monoprotic: gives away one H
- \* hydroiodic acid (HI)
- \* hydrobromic acid (HBr)
- \* hydrochloric acid(HCl)
- \* \*sulphuric acid ( $\text{SO}_3(\text{g}) + \text{H}_2\text{O} \rightarrow \text{H}_2\text{SO}_4$ ) polyprotic: gives more than one H
  - \*  $\text{H}_2\text{SO}_{4(\text{aq})} + 2\text{H}_2\text{O}_{(\text{l})} \rightarrow 2\text{H}_3\text{O}^+_{(\text{aq})} + \text{SO}_4^{2-}_{(\text{aq})}$
- \* \*nitric acid ( $\text{NO}_2(\text{g}) + \text{H}_2\text{O} \rightarrow \text{HNO}_3$ )
- \* hydronium ion ( $\text{H}_3\text{O}^+$ ) SA



# Relative Strengths of Acids and Bases at 298.15 K

SA

Common Name IUPAC / Systematic Name	Acid Formula	Conjugate Base Formula
perchloric acid aqueous hydrogen perchlorate	HClO <sub>4</sub> (aq)	ClO <sub>4</sub> <sup>-</sup> (aq)
hydroiodic acid aqueous hydrogen iodide	HI(aq)	I <sup>-</sup> (aq)
hydrobromic acid aqueous hydrogen bromide	HBr(aq)	Br <sup>-</sup> (aq)
hydrochloric acid aqueous hydrogen chloride	HCl(aq)	Cl <sup>-</sup> (aq)
sulfuric acid aqueous hydrogen sulfate	H <sub>2</sub> SO <sub>4</sub> (aq)	HSO <sub>4</sub> <sup>-</sup> (aq)
nitric acid aqueous hydrogen nitrate	HNO <sub>3</sub> (aq)	NO <sub>3</sub> <sup>-</sup> (aq)
hydronium ion	H <sub>3</sub> O <sup>+</sup> (aq)	H <sub>2</sub> O(l)
oxalic acid	HOOC <sup>-</sup> COOH(aq)	HOOC <sup>-</sup> COO <sup>-</sup> (aq)
sulfurous acid aqueous hydrogen sulfite	H <sub>2</sub> SO <sub>3</sub> (aq)	HSO <sub>3</sub> <sup>-</sup> (aq)
hydrogen sulfate ion	HSO <sub>4</sub> <sup>-</sup> (aq)	SO <sub>4</sub> <sup>2-</sup> (aq)
phosphoric acid aqueous hydrogen phosphate	H <sub>3</sub> PO <sub>4</sub> (aq)	H <sub>2</sub> PO <sub>4</sub> <sup>-</sup> (aq)
citric acid 2-hydroxy-1,2,3-propanetricarboxylic acid	C <sub>3</sub> H <sub>5</sub> O(COOH) <sub>3</sub> (aq)	C <sub>3</sub> H <sub>5</sub> O(COOH) <sub>2</sub> COO <sup>-</sup>
hydrofluoric acid aqueous hydrogen fluoride	HF(aq)	F <sup>-</sup> (aq)
nitrous acid aqueous hydrogen nitrite	HNO <sub>2</sub> (aq)	NO <sub>2</sub> <sup>-</sup> (aq)
formic acid methanoic acid	HCOOH(aq)	HCOO <sup>-</sup> (aq)
hydrogen oxalate ion	HOOC <sup>-</sup> COO <sup>-</sup> (aq)	OOC <sup>-</sup> COO <sup>2-</sup> (aq)
lactic acid 2-hydroxypropanoic acid	C <sub>2</sub> H <sub>5</sub> OCOOH(aq)	C <sub>2</sub> H <sub>5</sub> OCOO <sup>-</sup> (aq)
ascorbic acid 2(1,2-dihydroxyethyl)-4,5-dihydroxy-furan-3-one	C <sub>6</sub> H <sub>8</sub> O <sub>6</sub> (aq)	HC <sub>6</sub> H <sub>7</sub> O <sub>6</sub> <sup>-</sup> (aq)

Acid  
Rain



salts

WB

benzoic acid benzenecarboxylic acid	C <sub>6</sub> H <sub>5</sub> COOH(aq)	C <sub>6</sub> H <sub>5</sub> COO <sup>-</sup> (aq)
acetic acid ethanoic acid	CH <sub>3</sub> COOH(aq)	CH <sub>3</sub> COO <sup>-</sup> (aq)
dihydrogen citrate ion	C <sub>3</sub> H <sub>5</sub> O(COOH) <sub>2</sub> COO <sup>-</sup> (aq)	C <sub>3</sub> H <sub>5</sub> O(COOH)(COO) <sub>2</sub> <sup>2-</sup> (aq)
butanoic acid	C <sub>3</sub> H <sub>7</sub> COOH(aq)	C <sub>3</sub> H <sub>7</sub> COO <sup>-</sup> (aq)
propanoic acid	C <sub>2</sub> H <sub>5</sub> COOH(aq)	C <sub>2</sub> H <sub>5</sub> COO <sup>-</sup> (aq)
carbonic acid (CO <sub>2</sub> + H <sub>2</sub> O) aqueous hydrogen carbonate	H <sub>2</sub> CO <sub>3</sub> (aq)	HCO <sub>3</sub> <sup>-</sup> (aq)
hydrogen citrate ion	C <sub>3</sub> H <sub>5</sub> OCOOH(COO) <sub>2</sub> <sup>2-</sup> (aq)	C <sub>3</sub> H <sub>5</sub> O(COO) <sub>3</sub> <sup>3-</sup> (aq)
hydrosulfuric acid aqueous hydrogen sulfide	H <sub>2</sub> S(aq)	HS <sup>-</sup> (aq)
hydrogen sulfite ion	HSO <sub>3</sub> <sup>-</sup> (aq)	SO <sub>3</sub> <sup>2-</sup> (aq)
dihydrogen phosphate ion	H <sub>2</sub> PO <sub>4</sub> <sup>-</sup> (aq)	HPO <sub>4</sub> <sup>2-</sup> (aq)
hypochlorous acid aqueous hydrogen hypochlorite	HOCl(aq)	OCl <sup>-</sup> (aq)
hydrocyanic acid aqueous hydrogen cyanide	HCN(aq)	CN <sup>-</sup> (aq)
ammonium ion	NH <sub>4</sub> <sup>+</sup> (aq)	NH <sub>3</sub> (aq)
hydrogen carbonate ion	HCO <sub>3</sub> <sup>-</sup> (aq)	CO <sub>3</sub> <sup>2-</sup> (aq)
hydrogen ascorbate ion	HC <sub>6</sub> H <sub>6</sub> O <sub>6</sub> <sup>-</sup> (aq)	C <sub>6</sub> H <sub>6</sub> O <sub>6</sub> <sup>2-</sup> (aq)
hydrogen phosphate ion	HPO <sub>4</sub> <sup>2-</sup> (aq)	PO <sub>4</sub> <sup>3-</sup> (aq)
water	H <sub>2</sub> O(l)	OH <sup>-</sup> (aq)

vinegar

WA

SB

# One type of base completely dissociates

- \* Any Base with OH- completely dissociates and is considered a strong base (see the right side of pg 8 & 9 of your data book)
  - \* NaOH (mono hydroxide);  $\text{NaOH(s)} \rightarrow \text{Na}^+_{\text{(aq)}} + \text{OH}^-_{\text{(aq)}}$
  - \* Ba(OH)<sub>2</sub> (di hydroxide);  $\text{Ba(OH)}_2 \rightarrow \text{Ba}^{2+}_{\text{(aq)}} + 2 \text{OH}^-_{\text{(aq)}}$
  - \* Al(OH)<sub>3</sub> (poly hydroxide);  $\text{Al(OH)}_3 \rightarrow \text{Al}^{3+}_{\text{(aq)}} + 3 \text{OH}^-_{\text{(aq)}}$
- \* The acids above the OH get weaker as you move up the table until water, which is the weakest base. The substances above water are called salts.
- \* NOTE: often the bases have a spectator ion like Na

# Neutralization

\* S Acid + S Base => Salt + Water



- \* If you spill an acid neutralize it with baking soda (harmless)
- \* If you spill a base neutralize it with acetic acid (harmless vinegar)



Safety - so no one gets hurt.

WHMIS (Workplace Hazardous  
Materials Information System)

symbols often used for acids and  
bases. a) corrosive materials

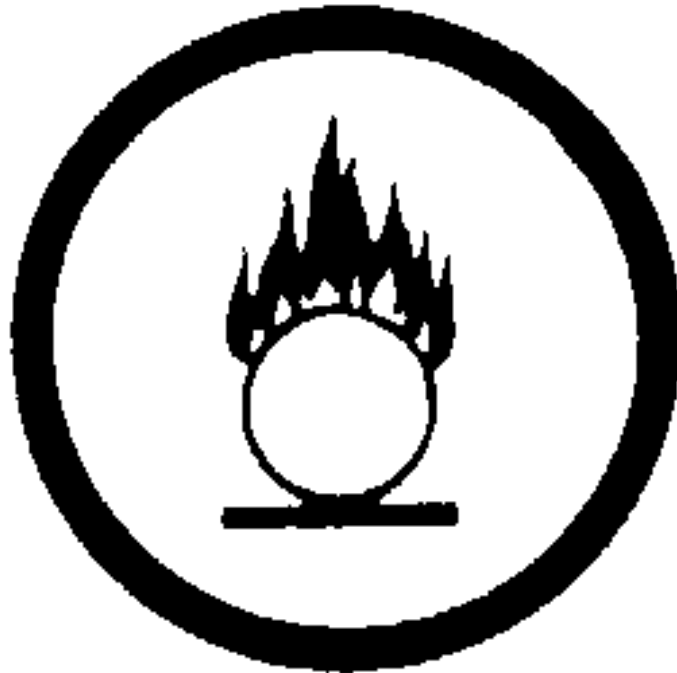


b) Materials causing immediate and serious toxic effects





## c) Oxidizing materials



# Safety practices

- \* Report all accidents to a supervisor
- \* Wash off all acids and bases with cold water
- \* SAFELY Neutralize an acid with baking soda (sodium hydrogen carbonate)
- \* SARFELY Neutralize a base with vinegar (ethanoic acid)
- \* Always add acid to water (A to W)

## II) Properties

# Properties of Acids & Bases

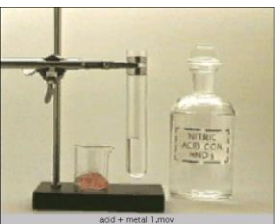
## Acid

## Base

<input type="checkbox"/> Feel:	like water	slippery
<input type="checkbox"/> Taste	sour	bitter
<input type="checkbox"/> Electrolyte:	Y	Y
<input type="checkbox"/> State	s/aq	s/aq
<input type="checkbox"/> Neutralize	bases	acids
<input type="checkbox"/> $[\text{H}_3\text{O}^+_{(\text{aq})}]$	high	low
<input type="checkbox"/> $[\text{OH}^-_{(\text{aq})}]$	low	high

# Properties of Acids & Bases

	<u>Acid</u>	<u>Base</u>
*Common	HCl	NaOH
*Reacts with _ to produce _	metals H <sub>2</sub> gas	fat soap
*Bromothymol	Yellow	Blue
*Phenolphthalein	Colorless	Pink
*Litmus	Red	Blue



# III) Formulas

# 1) Ion-product constant

**Definition:** The product of the concentrations of hydroxide(OH<sup>-</sup>) ions and the hydronium(H<sub>3</sub>O<sup>+</sup>) ions is the ion constant for water

**Symbol:**  $K_w = 1.000 \times 10^{-14} (\text{mol/L})^2$

**Formula:**  $K_w = [\text{H}_3\text{O}^+] \times [\text{OH}^-]$

le)  $1.0 \times 10^{-14} = 1.0 \times 10^{-7} \times 1.0 \times 10^{-7}$

# K<sub>w</sub> example #1

What is the concentration of the hydroxide ions in a 0.15 mol/L solution of hydrochloric acid?



Step 2: C=0.15 mol/L (given)

Step 3: Multiply by mole ratio

$$[\text{H}_3\text{O}^+_{(\text{aq})}] = 0.15 \text{ mol/L} \times 1 \text{ mol}/1 \text{ mol} = 0.15 \text{ mol/L}$$

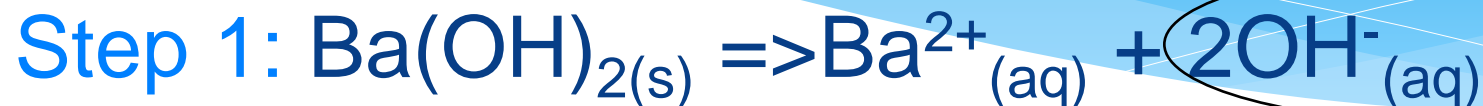
Step 4: Use the ion product to Solve:

$$\begin{aligned} [\text{OH}] &= K_w / [\text{H}_3\text{O}^+_{(\text{aq})}] \\ &= 1.000 \times 10^{-14} \text{ M}^2 / 0.15 \text{ M} \\ &= 6.7 \times 10^{-14} \text{ mol/L of OH}^-(\text{aq}) \end{aligned}$$



# K<sub>w</sub> example #2

Calculate the hydronium concentration in a 0.25 mol/L solution of barium hydroxide



Step 2: C=0.25 mol/L of barium hydroxide

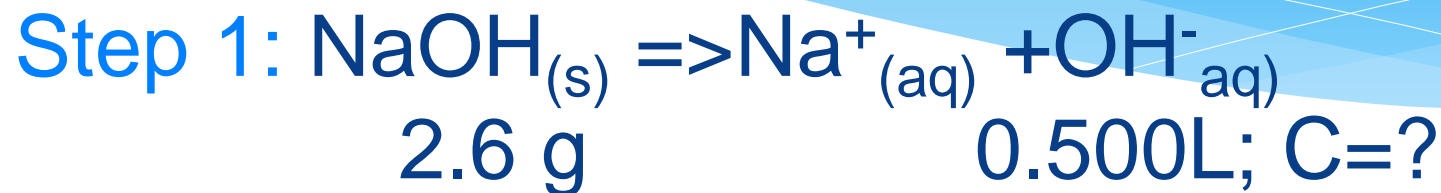
Step 3: Multiply by mole ratio

$$[\text{OH}^{-}_{(\text{aq})}] = 0.25 \text{ M} \times 2/1 = 0.50 \text{ M}$$

Step 4:  $[\text{H}_3\text{O}^{+}_{(\text{aq})}] = K_w / [\text{OH}^{-}_{(\text{aq})}]$   
 $= 1.000 \times 10^{-14} \text{ M}^2 / 0.50 \text{ M}$   
 $= 2.0 \times 10^{-14} \text{ mol/L of H}_3\text{O}^{+}$

# K<sub>w</sub> example #3

Calculate the  $[\text{H}_3\text{O}^+_{(\text{aq})}]$  and  $[\text{OH}^-]$  in 500 mL of water that has 2.6g of sodium hydroxide.



Step 2:  $n = m/M$      $n = 2.6\text{g}/40.00\text{g/mol} = 0.065\text{mol}$

Step 3:  $[\text{OH}^-_{(\text{aq})}] = 0.065 \text{ mol} \times 1/1 = 0.065\text{mol}$

Step 4:  $C = n/V$      $C = 0.065\text{mol}/0.500\text{L} = 0.13\text{mol/L}$

Use the ion product formula

$$\begin{aligned} [\text{H}_3\text{O}^+_{(\text{aq})}] &= K_w / [\text{OH}^-_{(\text{aq})}] \\ &= 1.000 \times 10^{-14} \text{ M}^2 / 0.13 \text{ M} \\ &= 7.7 \times 10^{-14} \text{ mol/L} \end{aligned}$$

## 2) Calculating pH/pOH

pH calculations are the negative log of concentrations.

Formulas:  $\text{pH} = -\log [\text{H}_3\text{O}^+_{(\text{aq})}]$   
 $\text{pOH} = -\log [\text{OH}^-_{(\text{aq})}]$

## 2) Calculating pH/pOH

Significant digits: FOR pH & pOH values only - the digit before the decimal is NOT significant. This number represents the exponent.

# pH example #1

pH of  $4.7 \times 10^{-11}$  mol/L of  $[\text{H}_3\text{O}^+_{(\text{aq})}]$

$$\text{pH} = - \log [4.7 \times 10^{-11}]$$

NOTE: For programmable calculators punch in the exact order. For other calculators punch the number in first, then log, then +/-

$$\text{pH} = 10.\underline{33} \text{ (2 significant digits)}$$

## pH example #2

What is the pOH of 0.0030 mol/L of hydroxide ions?

$$\text{pOH} = -\log [0.0030]$$

$$\text{pOH} = \underline{2.52} \text{ (2 significant digits)}$$

### 3) Calculating $[\text{H}_3\text{O}^+_{(\text{aq})}]$

Formulas:  $[\text{H}_3\text{O}^+_{(\text{aq})}] = 10^{\times(-\text{pH})}$   
 $= 10^{-\text{pH}}$

$$[\text{OH}^-_{(\text{aq})}] = 10^{\times(-\text{pOH})}$$
$$= 10^{-\text{pOH}}$$

$10^{\times}$  may be called antilog, inverse log or 2nd log.

# [H<sub>3</sub>O<sup>+</sup>] example #1

What is the hydronium

concentration of a pH of 10.33.

NOTE: 2 sig digs

$$\begin{aligned}[\text{H}_3\text{O}^+] &= 2^{\text{nd}} \log (-\text{pH}) \\ &= 2^{\text{nd}} \log (-10.33) \\ &= \underline{4.7} \times 10^{-11} \text{ mol/L}\end{aligned}$$



## [H<sub>3</sub>O<sup>+</sup>] example #2

Calculate the hydroxide ion concentration if the pOH is 2.31

NOTE: 2 sig digs

$$\begin{aligned}[\text{OH}^-] &= 2^{\text{nd}} \log (-\text{pOH}) \\ &= 2^{\text{nd}} \log (-2.31) \\ &= 4.91 \times 10^{-3} \text{ mol/L} \\ &= 4.9 \times 10^{-3} \text{ mol/L}\end{aligned}$$

## 4) pH & pOH relationship

Formula:  $14.00 = \text{pH} + \text{pOH}$

Example:

What is the pOH in example 1?

$$\text{pOH} = 14.00 - 10.33 = 3.67$$

What is the pH in example 2?

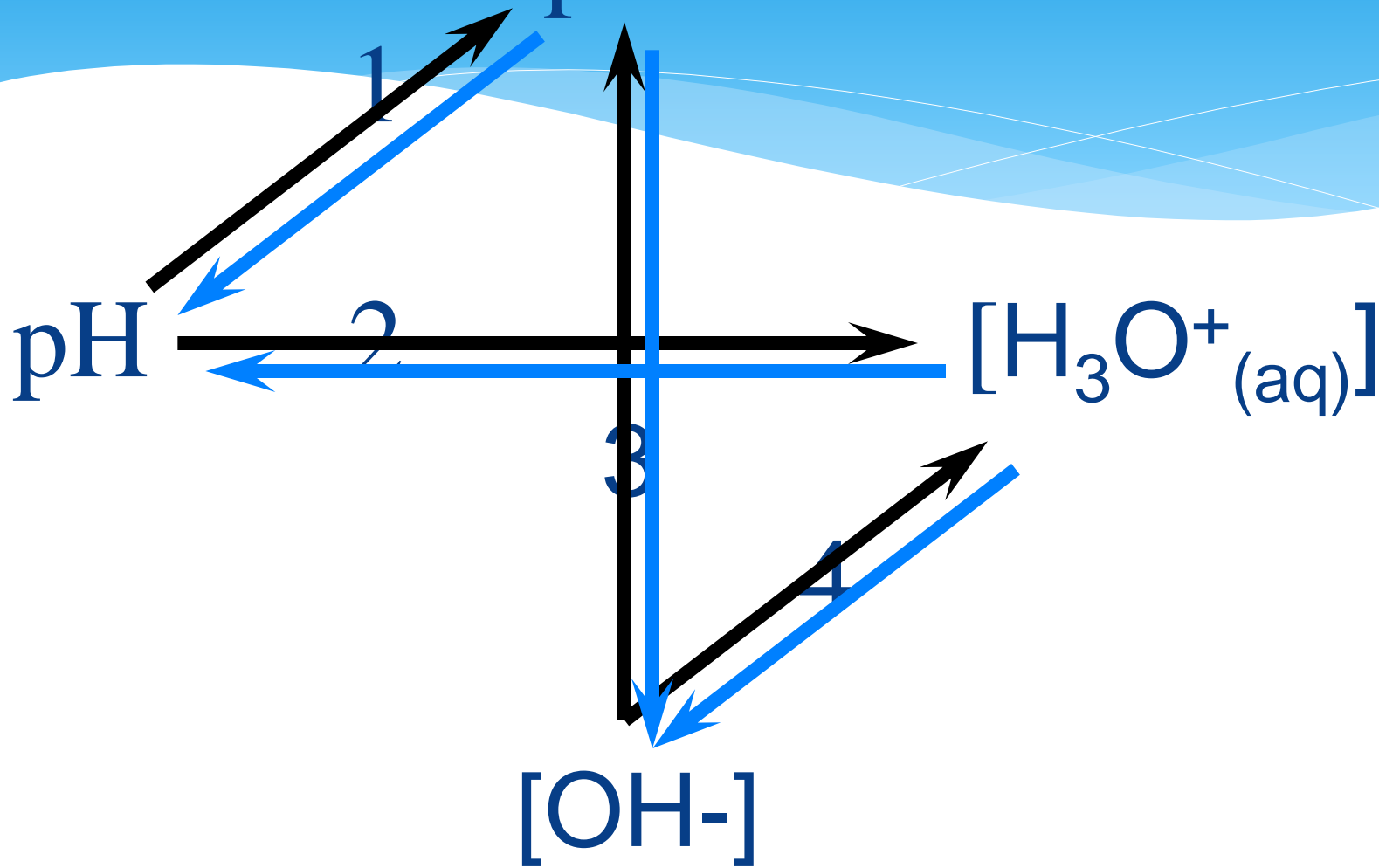
$$\text{pH} = 14.00 - 2.52 = 11.48$$

# Summary

- \* Defined acids, bases, indicators, pH, pOH
- \* Reviewed dissociation, self-ionization, neutralization
- \* Learned WHMIS & safety
- \* Properties of acids & bases

# Summary

pOH



# Summary

1)  $\text{pH} = 14.00 - \text{pOH}$

$$\text{pOH} = 14.00 - \text{pH}$$

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

$$[\text{H}_3\text{O}^+] = 2^{\text{nd}} \log (-\text{pH})$$

3)  $[\text{OH}] = 2^{\text{nd}} \log (-\text{pOH})$

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

4)  $[\text{OH}^-_{(\text{aq})}] = K_w / [\text{H}_3\text{O}^+_{(\text{aq})}]$

$$[\text{H}_3\text{O}^+_{(\text{aq})}] = K_w / [\text{OH}^-_{(\text{aq})}]$$

# Acid – Base Stoichiometry

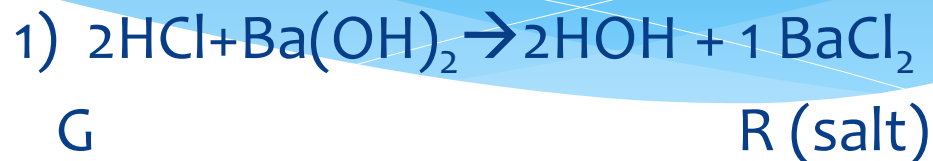
1. Balanced reaction or equation containing a Strong Acid or Strong Base
2. Change the given (limiting) to moles: Often you will need to find the concentration using one of the four acid-base formulas and then find moles using  $n=CV$ .
3. Multiply by the mole ratio (R/G)
4. Convert into desired units.

# Example

200 mL solution of HCl with a pH of 1.45 neutralizes 300 mL of  $\text{Ba}(\text{OH})_2$ . Determine the concentration of salt formed.

A 200 mL solution of  $\text{HCl}_{(\text{aq})}$  with a pH of 1.45 neutralizes 300 mL of  $\text{Ba}(\text{OH})_{2(\text{aq})}$ . Determine the concentration of salt formed.

\* Step 1: Reaction



\* Step 2: Find moles;  
First find concentration of HCl using pH formula

$$2) [\text{A}] = 10^{-1.45} = 3.548... \text{E-2 mol/L}$$

$$n = CV; n = 7.096... \text{E}^{-3} \text{ mol}$$

$$3) 7.096... \text{E}^{-3} \times \frac{1}{2} = 3.548... \text{E}^{-3} \text{ mol}$$

$$4) C = n/V_{\text{total}}; C = 3.548\text{E-3}/0.500\text{L}$$

$$\mathbf{C = 7.096\text{E-3 mol/L}}$$

\* Step 3: mole ratio

\* Step convert to units



# More Information

- \*Chapter 18 in your text book.
- \*Notes & Worksheets
- \*Tutorials after class