

# Lecture Presentation Chapter 4 Reactions in Aqueous Solution

4.1, 4.2, 4.3, 4.4, 4.6, 4.7,
4.8, 4.12, 4.15, 4.18, 4.20,
4.22, 4.23, 4.24, 4.26, 4.28,
4.31, 4.42, 4.48, 4.52, 4.54,
4.64, 4.68, 4.70, 4.72, 4.90,
4.98, 4.104, 4.110, 4.138

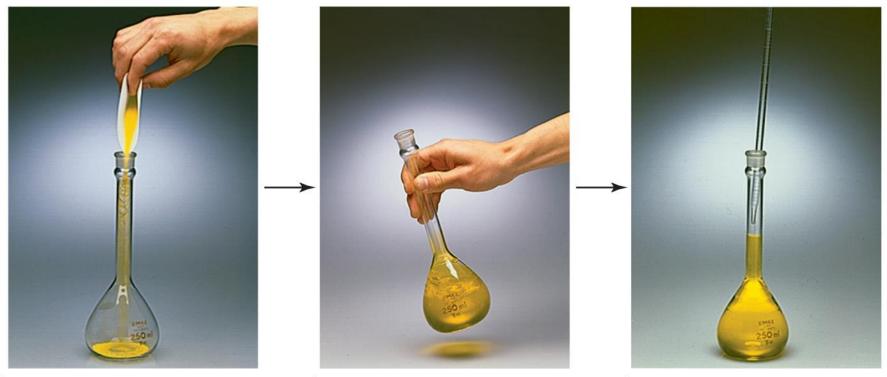
John E. McMurry Robert C. Fay

**Molarity**: The number of moles of a substance dissolved in each liter of solution

Solution: A homogeneous mixture

**Solute**: The dissolved substance in a solution (substance you have less quantity of in solution)

**Solvent**: The major component in a solution (substance you have more of in solution)



A measured number of moles of **solute** is placed in a volumetric flask.

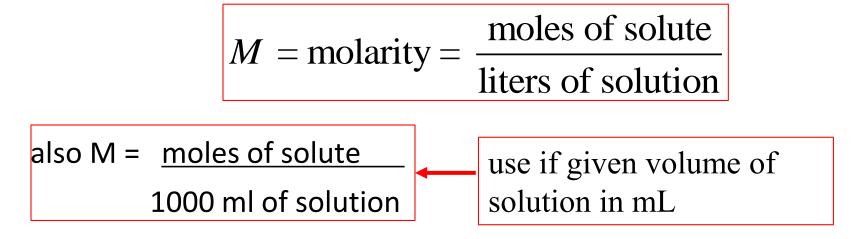
Enough solvent is added to dissolve the solute by swirling.

Further solvent is added to reach the calibration mark on the neck of the flask, and the solution is mixed until uniform.

# **Definition of Molarity**

Molarity (M)

- Commonly used expression for concentration
- Defined as moles of solute per volume of solution in liters



#### **Definition of Molarity**

$$M = \text{molarity} = \frac{\text{moles of solute}}{\text{liters of solution}}$$

Example: What is the molarity if you put 1.00 mol of sodium chloride in enough water to make 1.00 L of solution ? End class D section 9/30

$$\frac{1.00 \text{ mol}}{1.00 \text{ L}} = 1.00 \frac{\text{mol}}{\text{L}} \text{ or } 1.00 \text{ M}$$

End class D section 9/30

 Calculate the molarity of a solution prepared by dissolving 1.56 g of gaseous HCl (36.46 g/mol) in enough water to make 26.8 mL of solution

Doc camera

- Where are we going?
  - To find the molarity of HCl solution
  - What do we know?
    - 1.56 g HCl
    - 26.8 mL solution
  - What information do we need to find molarity?
    - Moles solute

Molarity =  $\frac{\text{mol solute}}{\text{L solution}}$ 

- How do we get there?
  - What are the moles of HCl (36.46 g/mol)?

$$1.56 \text{ g HCI} \times \frac{1 \text{ mol HCl}}{36.46 \text{ g HCl}} = 4.28 \times 10^{-2} \text{ mol HCl}$$

What is the volume of solution (in liters)?

$$26.8 \text{ mL} \times \frac{1 \text{ L}}{1000 \text{ mL}} = 2.68 \times 10^{-2} \text{ L}$$

What is the molarity of the solution?

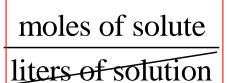
Molarity =  $\frac{4.28 \times 10^{-2} \text{ mol HCl}}{2.68 \times 10^{-2} \text{ L solution}} = 1.60 M \text{ HCl}$ 

- Reality check
  - The units are correct for molarity

Determining Moles of Solute in a Sample

 Use the definition of molarity as a conversion factor (to get moles of solute)

Liters of solution  $\times$  molarity = <u>liters of solution</u>  $\times$ 



Moles of solute = Liters of solution  $\times$  molarity

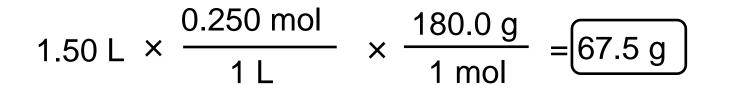
- Calculate the number of moles of Cl<sup>-</sup> ions in 1.75 L of 0.152 *M Na*Cl (M = moles / liter solution)
- On doc camera

How many grams of solute would you use to prepare 1.50 L of 0.250 M glucose,  $C_6H_{12}O_6$ ?

Molar mass  $C_6H_{12}O_6 = 180.0$  g/mol

How many grams of solute would you use to prepare 1.50 L of 0.250 M glucose,  $C_6H_{12}O_6$ ?

Molar mass  $C_6H_{12}O_6 = 180.0$  g/mol



#### Dilution

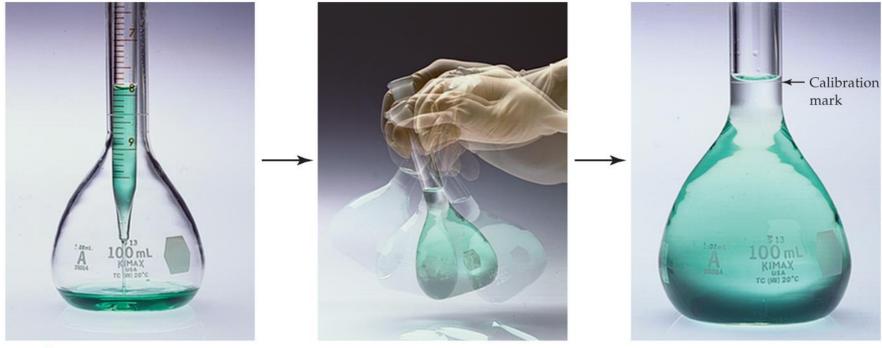
 Process of adding water to a concentrated (stock) solution

moles of solute before dilution = moles of solute after dilution

$$M_i V_i$$
 = moles initial  $M_f V_f$  = moles final

- dilution equation:  $M_i V_i = M_f V_f$
- as long as keep volume unit the same between initial and final, can have both mL or both L

### **Diluting Concentrated Solutions**



The **volume** to be diluted is placed in an empty volumetric flask.

Solvent is added to a level just below the calibration mark, and the flask is shaken.

More solvent is added to reach the calibration mark, and the flask is again shaken.

#### **Diluting Concentrated Solutions**

How would you prepare 250.0 mL of 0.500 M from 18.0 M aqueous  $H_2SO_4$ ?

 $M_i = 18.0 M$   $M_f = 0.500 M$ 

$$V_{\rm i} = ? \,\rm{mL}$$
  $V_{\rm f} = 250.0 \,\rm{mL}$ 

$$V_{\rm i} = \frac{M_{\rm f} V_{\rm f}}{M_{\rm i}} = \frac{0.500 \,\rm M}{18.0 \,\rm M} \times 250.0 \,\rm mL = 6.94 \,\rm mL$$

Add 6.94 mL 18.0 M sulfuric acid to enough water to make 250.0 mL of 0.500 M solution.

# HW: Diluting Concentrated Solutions [do (a) only for HW]

- Describe how you would prepare 2.00 L of each of the following solutions: (on doc camera)
  - a. 0.250 M NaOH from 1.00 M NaOH stock solution

b. 0.100  $M K_2 CrO_4$  from 1.75  $M K_2 CrO_4$  stock solution

HW: Diluting Concentrated Solutions [do (a) only for HW]

End 9/30 F section

- Describe how you would prepare 2.00 L of each of the following solutions: (on doc camera)
  - a. 0.250 M NaOH from 1.00 M NaOH stock solution

Add 500. mL (= 0.500 L) of the 1.00 *M* NaOH stock solution to a 2-L volumetric flask; fill to the mark with

b. 0.100  $M \text{ K}_2 \text{CrO}_4$  from 1.75  $M \text{ K}_2 \text{CrO}_4$  stock solution

Add 114 mL (= 0.114 L) of the 1.75 *M* K<sub>2</sub>CrO<sub>4</sub> stock solution to a 2- L volumetric flask; fill to the mark with water

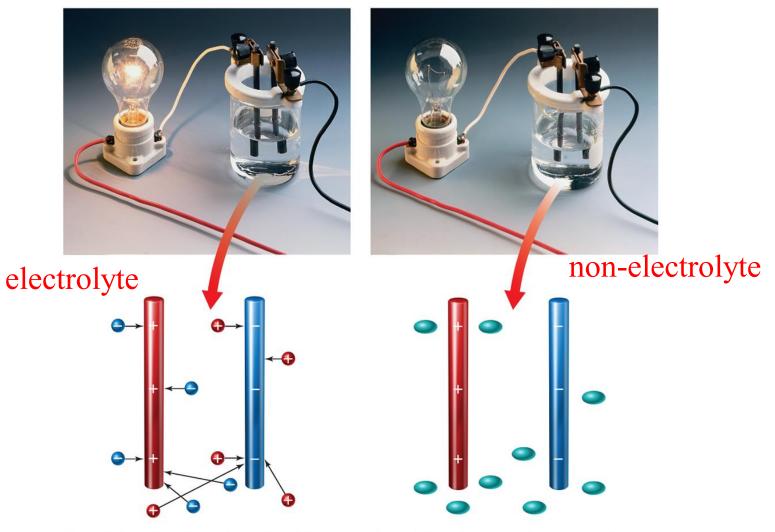
**Electrolytes**: dissociate in water to produce conducting solutions of ions

$$H_2O$$

$$NaCl(s) \longrightarrow Na^+(aq) + Cl^-(aq)$$

**Nonelectrolytes**: do not dissociate to produce ions in water (covalent compounds)

$$H_2O \longrightarrow C_{12}H_{22}O_{11}(s) \longrightarrow C_{12}H_{22}O_{11}(aq)$$



A solution of NaCl conducts electricity because of the movement of charged particles (ions), thereby completing the circuit and allowing the bulb to light. A solution of sucrose does not conduct electricity or complete the circuit because it contains no mobile charged particles. The bulb therefore remains dark.

**Strong Electrolytes**: completely dissociates in water (ionic compounds, strong acids, strong bases)

 $KCI(aq) \longrightarrow K^+(aq) + CI^-(aq)$ 

Weak Electrolytes: incompletely dissociates in water (weak acids, weak bases)

 $CH_3CO_2H(aq) \longrightarrow H^+(aq) + CH_3CO_2(aq)$ 

TABLE 4.1         Electrolyte Classification of Some Common Substances		
Strong Electrolytes	Weak Electrolytes	Nonelectrolytes
HCl, HBr, HI	CH <sub>3</sub> CO <sub>2</sub> H	H <sub>2</sub> O
HClO <sub>4</sub>	HF	CH <sub>3</sub> OH (methyl alcohol)
HNO <sub>3</sub>	HCN	C <sub>2</sub> H <sub>5</sub> OH (ethyl alcohol)
$H_2SO_4$	$\mathbf{N}$	$C_{12}H_{22}O_{11}$ (sucrose)
KBr	$\mathbf{A}$	Most compounds of carbon (organic
NaCl	$\backslash$	compounds)
NaOH, KOH		
Other soluble ionic compounds		

**Strong** Acids: Hydrochloric acid, hydrobromic acid, hydroiodic acid, perchloric acid, nitric acid, sulfuric acid

TABLE 4.1         Electrolyte Classification of Some Common Substances		
Strong Electrolytes	Weak Electrolytes	Nonelectrolytes
HCl, HBr, HI	CH <sub>3</sub> CO <sub>2</sub> H	H <sub>2</sub> O
$HClO_4$	HF	CH <sub>3</sub> OH (methyl alcohol)
HNO <sub>3</sub>	HCN	C <sub>2</sub> H <sub>5</sub> OH (ethyl alcohol)
$H_2SO_4$		$C_{12}H_{22}O_{11}$ (sucrose)
KBr		Most compounds of carbon (organic
NaCl		compounds)
NaOH, KOH		
Other soluble ionic compounds		
Ionic compounds		

TABLE 4.1         Electrolyte Classification of Some Common Substances		
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HCl, HBr, HI	CH <sub>3</sub> CO <sub>2</sub> H	H <sub>2</sub> O
HClO <sub>4</sub>	HF	CH <sub>3</sub> OH (methyl alcohol)
HNO <sub>3</sub>	HCN	C <sub>2</sub> H <sub>5</sub> OH (ethyl alcohol)
$H_2SO_4$		$C_{12}H_{22}O_{11}$ (sucrose)
KBr		Most compounds of carbon (organic
NaCl		compounds)
NaOH, KOH		
Other soluble ionic compounds		
Weak acids		

TABLE 4.1         Electrolyte Classification of Some Common Substances		
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HNO <sub>3</sub>	HCN	C <sub>2</sub> H <sub>5</sub> OH (ethyl alcohol)
$H_2SO_4$		$C_{12}H_{22}O_{11}$ (sucrose)
KBr		Most compounds of carbon (organic
NaCl		compounds)
NaOH, KOH		
Other soluble ionic c	compounds	
Molecular (covalent) compounds		
End class 9/30 Monday G section		

#### **Types of Chemical Reactions in Aqueous Solution**

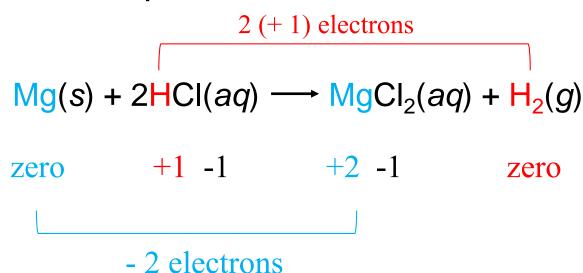
**Precipitation Reactions**: soluble reactants give insoluble solid product (precipitates out)

 $Pb(NO_3)_2(aq) + 2KI(aq) \longrightarrow 2KNO_3(aq) + PbI_2(s)$ 



#### **Types of Chemical Reactions in Aqueous Solution**

**Oxidation–Reduction (Redox) Reactions**: one or more electrons (negative charge) are transferred between reaction partners

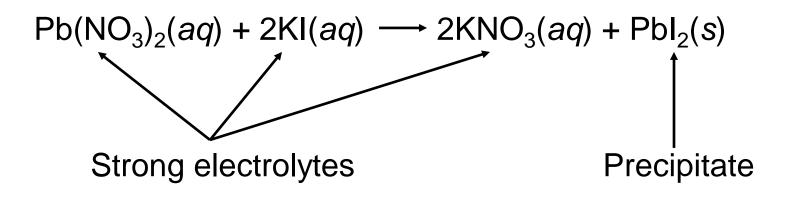


### **Types of Chemical Reactions in Aqueous Solution**

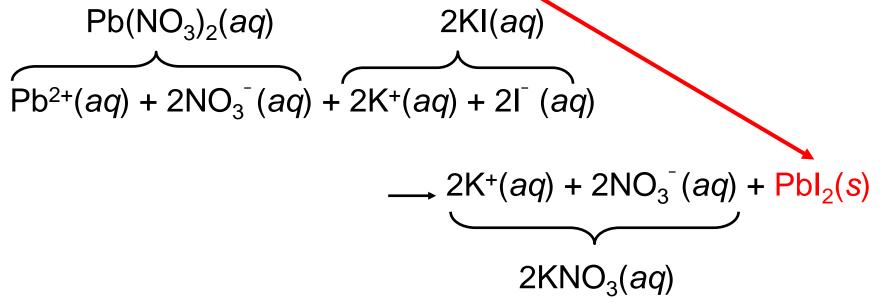
Acid-Base Neutralization Reactions: acid reacts with base to give water & salt (salt is ionic compound, not always NaCl)

HCI(aq	) + NaOH( <i>aq</i> ) -	$\rightarrow$ H <sub>2</sub> O( <i>I</i> ) +	NaCl(aq)
acid	base	water	salt

Molecular Equation: write complete formulas as if *molecules*.



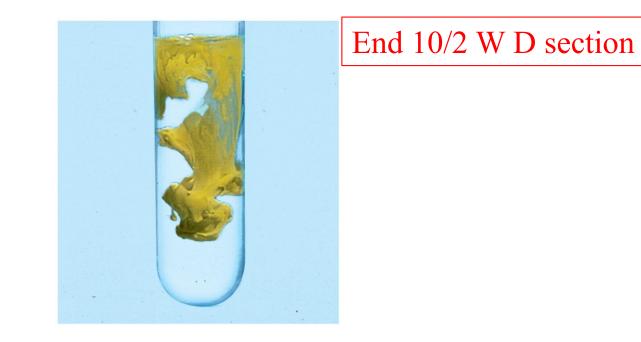
**lonic Equation**: write strong electrolytes as dissociated ions. (solid, liquid, gas, weak electrolyte compounds written as molecular formula-do NOT dissociate)



**Spectator lons**: ions that do not change during the reaction. (& just watch the reaction without doing anything)

**Net Ionic Equation**: Only the ions undergoing change are shown. (leave out spectators)

 $Pb^{2+}(aq) + 2l^{\overline{}}(aq) \longrightarrow Pbl_{2}(s)$ 



# Precipitation Reactions and Solubility Guidelines

Typo L<sup>+</sup> is really Li<sup>+</sup>

#### **TABLE 4.2** Solubility Guidelines for Ionic Compounds in Water

Soluble Compounds	Common Exceptions
$L^+$ , Na <sup>+</sup> , K <sup>+</sup> , Rb <sup>+</sup> , Cs <sup>+</sup> (group 1A cations)	None
$\mathrm{NH_4^+}$ (ammonium ion)	None
Cl <sup>-</sup> , Br <sup>-</sup> , I <sup>-</sup> (halide)	Halides of $Ag^+$ , $Hg_2^{2+}$ , $Pb^{2+}$
$NO_3^{-}$ (nitrate)	None
$ClO_4^-$ (perchlorate)	None
$CH_3CO_2^-$ (acetate)	None
$SO_4^{2-}$ (sulfate)	Sulfates of $Sr^{2+}$ , $Ba^{2+}$ , $Hg_2^{2+}$ , $Pb^{2+}$
Insoluble Compounds	Common Exceptions
$\text{CO}_3^{2-}$ (carbonate)	Carbonates of group 1A cations, $NH_4^+$
S <sup>2–</sup> (sulfide)	Sulfides of group 1A cations, $NH_4^+$ , $Ca^{2+}$ , $Sr^{2+}$ , and $Ba^{2+}$
$PO_4^{3-}$ (phosphate)	Phosphates of group 1A cations, $NH_4^+$
OH <sup>-</sup> (hydroxide)	Hydroxides of group 1A cations, $NH_4^+$ , $Ca^{2+}$ , $Sr^{2+}$ , and $Ba^{2+}$

# HW: Are the following compounds soluble or insoluble in water ?

 $Li_2 CO_3$ Ba  $SO_4$ Mg  $(NO_3)_2$ 

### **Precipitation Reactions and Solubility Guidelines**

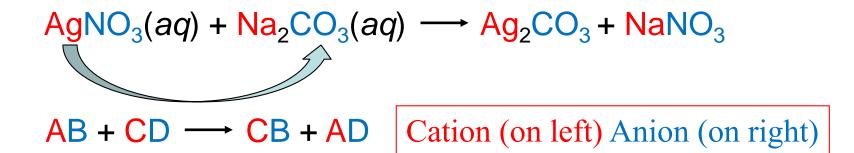
Write the molecular, ionic, and net ionic equations for the reaction that occurs when aqueous solutions of  $AgNO_3$  and  $Na_2CO_3$  are mixed.



#### **Precipitation Reactions and Solubility Guidelines**

Write the molecular, ionic, and net ionic equations for the reaction that occurs when aqueous solutions of  $AgNO_3$  and  $Na_2CO_3$  are mixed.

Write the chemical formulas of the products (use proper ionic rules). (exchange cation / anion partners)

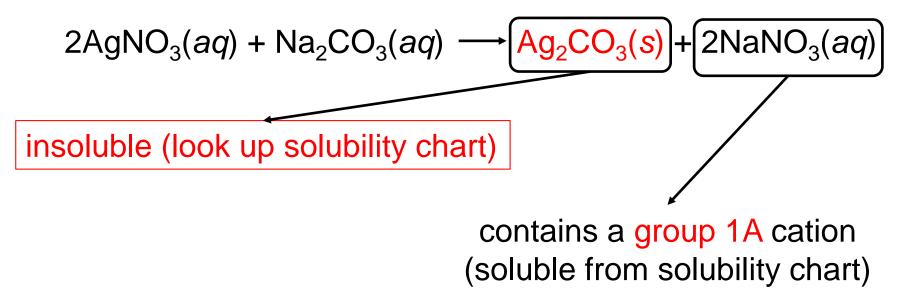


double replacement reaction

## **Precipitation Reactions and Solubility Guidelines**

Write the molecular, ionic, and net ionic equations for the reaction that occurs when aqueous solutions of  $AgNO_3$  and  $Na_2CO_3$  are mixed.

2. **Molecular Equation**: Balance the equation and predict the solubility of each possible product.



## **Precipitation Reactions and Solubility Guidelines**

Write the molecular, ionic, and net ionic equations for the reaction that occurs when aqueous solutions of  $AgNO_3$  and  $Na_2CO_3$  are mixed.

Ionic Equation: dissociate the soluble ionic compounds.
 Do <u>NOT</u> dissociate precipitates.

$$2AgNO_{3}(aq) \qquad Na_{2}CO_{3}(aq)$$

$$2Ag^{+}(aq) + 2NO_{3}^{-}(aq) + 2Na^{+}(aq) + CO_{3}^{2^{-}}(aq)$$

$$\longrightarrow Ag_{2}CO_{3}(s) + 2Na^{+}(aq) + 2NO_{3}^{-}(aq)$$
End 10/2 G section
$$2NaNO_{3}(aq)$$

## **Precipitation Reactions and Solubility Guidelines**

Write the molecular, ionic, and net ionic equations for the reaction that occurs when aqueous solutions of  $AgNO_3$  and  $Na_2CO_3$  are mixed. End F section 10/2

4. Net lonic Equation: Eliminate the spectator ions from the ionic equation.

 $2Ag^{+}(aq) + 2NO_{3}(aq) + 2Na^{+}(aq) + CO_{3}^{2^{-}}(aq)$ 

 $\rightarrow Ag_2CO_3(s) + 2Na^+(aq) + 2NO_3^-(aq)$ 

$$2Ag^{+}(aq) + CO_{3}^{2^{-}}(aq) \longrightarrow Ag_{2}CO_{3}(s)$$

<u>HW: Precipitation Reaction For the following</u>, Write out
(a) molecular reaction (b) total ionic equation and
(c) choose one of the following as the net ionic equation

Lead(II) nitrate + sodium chloride 10/3 end D section

What is the net ionic equation for this reaction?

a. 
$$\operatorname{Pb}^{2+}(aq) + 2\operatorname{NO}_{3}^{-}(aq) \longrightarrow \operatorname{Pb}(\operatorname{NO}_{3})_{2}(s)$$
  
b.  $\operatorname{Na}^{2+}(aq) + \operatorname{Cl}^{-}(aq) \longrightarrow \operatorname{NaCl}(s)$ 

- c.  $Pb^{2+}(aq) + 2Cl^{-}(aq) \longrightarrow PbCl_{2}(s)$
- d.  $\operatorname{Na}^+(aq) + \operatorname{NO}_3^-(aq) \longrightarrow \operatorname{NaNO}_3(s)$

<u>HW: Precipitation Reaction For the following,</u> Write out(a) molecular reaction (b) total ionic equation and(c) choose one of the following as the net ionic equation

Lead(II) nitrate + sodium chloride End 10/4 F, G sect End 10/7 D section

What is the net ionic equation for this reaction?

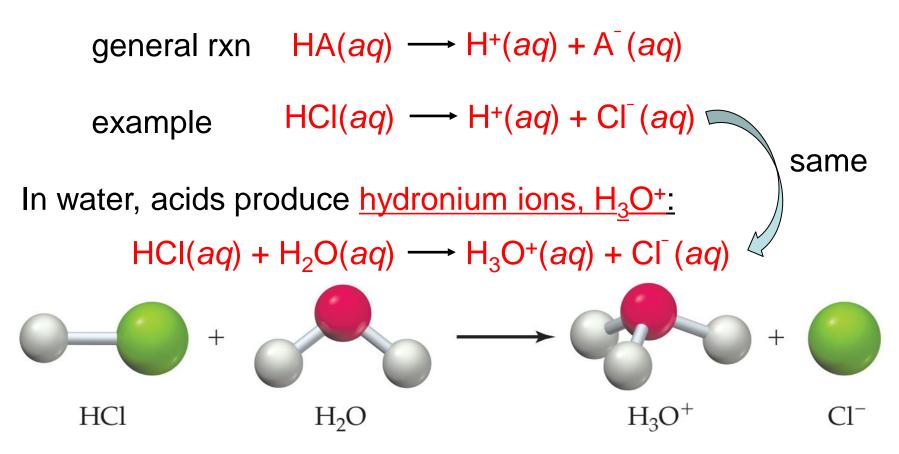
a. 
$$\operatorname{Pb}^{2+}(aq) + 2\operatorname{NO}_{3}^{-}(aq) \longrightarrow \operatorname{Pb}(\operatorname{NO}_{3})_{2}(s)$$
  
b.  $\operatorname{Na}^{2+}(aq) + \operatorname{Cl}^{-}(aq) \longrightarrow \operatorname{NaCl}(s)$   
c.  $\operatorname{Pb}^{2+}(aq) + 2\operatorname{Cl}^{-}(aq) \longrightarrow \operatorname{PbCl}_{2}(s) *$ 

d.  $\operatorname{Na}^+(aq) + \operatorname{NO}_3^-(aq) \longrightarrow \operatorname{NaNO}_3(s)$ 

# Test II ends here.

10/7 Monday was just a review day – <u>no new</u> <u>material covered</u> – only questions & reviewing quiz 2 answer keys (note D section had already seen the question on precipitation molecular, ionic and net ionic reaction on Thursday 10/3 their quiz date but we completed the HW in class on that HW problem for the D section on 10/7 – the answer for the precipitation problem was already posted on 10/5 before the D section did the HW problem in class for points)

Acid (Arrhenius): dissociates in water to produce hydrogen ions, H<sup>+</sup>



**Base (Arrhenius)**: dissociates in water to produce hydroxide ions, OH<sup>-</sup>:

general rxn  $MOH(aq) \rightarrow M^+(aq) + OH^-(aq)$ 

example  $NaOH(aq) \rightarrow Na^+(aq) + OH^-(aq)$ 

Ammonia is a weak base produces ammonium and hydroxide ions: (only weak base commonly seen)

 $NH_3(aq) + H_2O(aq) \longrightarrow NH_4^+(aq) + OH^-(aq)$ 

TABLE 4.3         Some Common Acids and Bases					
Strong acid	HClO <sub>4</sub> H <sub>2</sub> SO <sub>4</sub> $*$ HBr $*$ HCl $*$ HNO <sub>3</sub> $*$	Perchloric acid Sulfuric acid Hydrobromic acid Hydrochloric acid Nitric acid	KOH NaOH Ba(OH) <sub>2</sub> Ca(OH) <sub>2</sub>	Potassium hydroxide Sodium hydroxide Barium hydroxide Calcium hydroxide	Strong base
Weak acid	H <sub>3</sub> PO <sub>4</sub> * HF * HNO <sub>2</sub> CH <sub>3</sub> CO <sub>2</sub> H	Phosphoric acid Hydrofluoric acid Nitrous acid * Acetic acid	NH3	Ammonia *	Weak base

Strong acids and strong bases are strong electrolytes. Weak acids and weak bases are weak electrolytes. Know name, formula, whether strong/weak acid or base for \* (most group 1A & 2A hydroxides are strong bases if soluble)

#### Acids – characterized by # H<sup>+</sup> dissociation

 $H_{3}PO_{4} \rightarrow 3H^{+} + PO_{4}^{-3}$ triprotic  $H_{2}SO_{4} \rightarrow 2H^{+} + SO_{4}^{-2}$ diprotic  $H Br \rightarrow H^{+} + Br^{-}$ monoprotic  $CH_{3}CO_{2}H \rightarrow H^{+} + CH_{3}CO_{2}^{-}$ monoprotic (acetic acid - not 4 H+)

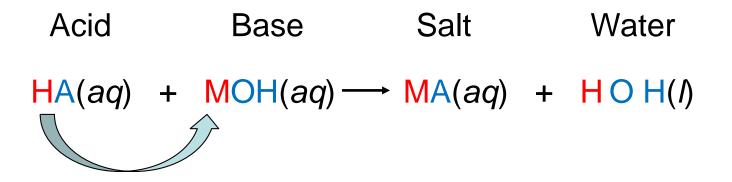
polyprotic acid = more than one hydrogen is acidic

#### Naming Binary Acids (all halogens)

- HCI hydrochloric acid
- HBr hydrobromic acid
- HF hydrofluoric acid (only weak halogen acid)
- HI hydroiodic acid

These <u>acid–base neutralization reactions</u> are doublereplacement reactions just like the precipitation reactions:

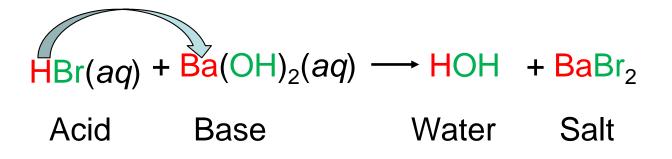
(exchange cation / anion partners)



double replacement reaction

Write the molecular, ionic, and net ionic equations for the reaction of aqueous HBr and aqueous  $Ba(OH)_2$ .

1. Write the chemical formulas of the products (use proper ionic rules for the salt to write neutral formula of the salt).



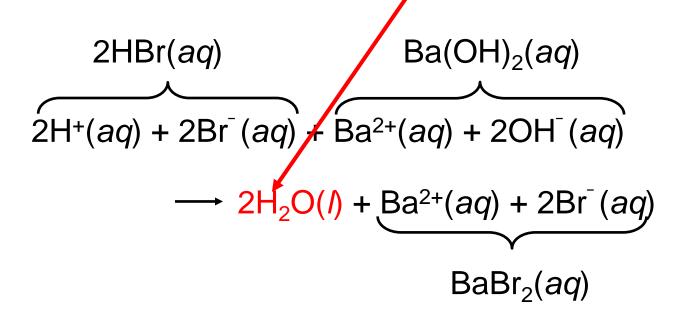
Write the molecular, ionic, and net ionic equations for the reaction of aqueous HBr and aqueous  $Ba(OH)_2$ .

2. Molecular Equation: Balance the equation and predict the solubility of the salt in the products. (from solubility rules table)

$$2\text{HBr}(aq) + \text{Ba}(OH)_2(aq) \longrightarrow 2H_2O(l) + \text{BaBr}_2(aq)$$
  
Use the solubility rules.

Write the molecular, ionic, and net ionic equations for the reaction of aqueous HBr and aqueous  $Ba(OH)_2$ .

3. **Ionic Equation**: Dissociate the strong acid and the soluble ionic compounds. (solid, liquid, gas compounds, weak acid written as molecular formula-do NOT dissociate)



Write the molecular, ionic, and net ionic equations for the reaction of aqueous HBr and aqueous  $Ba(OH)_2$ .

- 4. **Net Ionic Equation**: Eliminate the spectator ions from the ionic equation.
- 2H+(aq) + 2Br<sup>-</sup>(aq) + Ba<sup>2+</sup>(aq) + 2OH<sup>-</sup>(aq)

$$\rightarrow$$
 2H<sub>2</sub>O(*I*) + Ba<sup>2+</sup>(aq) + 2Br<sup>1-</sup>(aq)

 $2H^+(aq) + 2OH^-(aq) \longrightarrow 2H_2O(I)$ 

$$H^+(aq) + OH^-(aq) \longrightarrow H_2O(I)$$

Net ionic equation for acid/base neutralization
is almost always this (unless weak undissociated acid/base)

Write the molecular, ionic, and net ionic equations for the reaction of aqueous NaOH and aqueous HF.

 Write the chemical formulas of the products (use proper ionic rules for the salt to write neutral ionic formulas). (exchange cation anion partners)

> $HF(aq) + NaOH(aq) \longrightarrow HOH + NaF$ Acid Base Water Salt

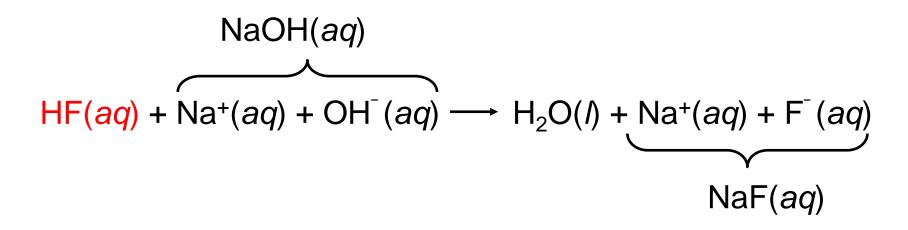
Write the molecular, ionic, and net ionic equations for the reaction of aqueous NaOH and aqueous HF.

2. Molecular Equation: Balance the equation and predict the solubility of the salt in the products.

$$HF(aq) + NaOH(aq) \longrightarrow H_2O(l) + NaF(aq)$$

Write the molecular, ionic, and net ionic equations for the reaction of aqueous NaOH and aqueous HF.

 Ionic Equation: Dissociate the soluble ionic compounds. (HF is a weak acid so does not dissociate)



Write the molecular, ionic, and net ionic equations for the reaction of aqueous NaOH and aqueous HF.

4. **Net lonic Equation**: Eliminate the spectator ions from the ionic equation.

$$HF(aq) + Na^{+}(aq) + OH^{-}(aq) \longrightarrow H_2O(l) + Na^{+}(aq) + F^{-}(aq)$$

$$\left(\mathsf{HF}(aq) + \mathsf{OH}^{-}(aq) \longrightarrow \mathsf{H}_2\mathsf{O}(l) + \mathsf{F}^{-}(aq)\right)$$

Write the molecular, ionic, and net ionic equations for the reaction of aqueous  $C_a$  (OH), and aqueous H I.

molecular equation:

 $Ca (OH)_2 + HI \rightarrow - +$ 

ionic equation

net ionic equation

# **Solution Stoichiometry**(already know how to do combining 2 things you know – stoichiometry & Molarity)

What volume of 0.250 M  $H_2SO_4$  is needed to react with 50.0 mL of 0.100 M NaOH? 10.0 mL  $H_2SO_4$ 

 $H_2SO_4(aq) + 2NaOH(aq) \rightarrow Na_2SO_4(aq) + 2H_2O(l)$ 

End 10/11 F F section

What volume of 0.250 M H Cl is needed to react with 50.0 mL of 0.100 M NaOH? 20.0 mL HCl

 $H CI(aq) + NaOH(aq) \rightarrow Na CI(aq) + H_2O(l)$ 

End 10/10/19 D, G section

#### Solution Stoichiometry (short cut version)

What volume of 0.250 M H Cl is needed to react with 50.0 mL of 0.100 M NaOH?

$$H CI(aq) + NaOH(aq) \rightarrow Na CI(aq) + H_2O(l)$$

Short cut: use  $M_{acid}V_{acid} = M_{base}V_{base}$ 

only works for 1:1 acid/base reaction

 $(0.250 \text{ M HCl}) * (V_{HCl}) = (0.100 \text{ M NaOH}) * (50.0 \text{ mL NaOH})$ 

 $V_{HCl} = (0.100 \text{ M NaOH}) * (50.0 \text{ mL NaOH}) = 20.0 \text{ mL HCl}$ 0.250 M H Cl

## **HW: Solution Stoichiometry**

What volume of 2.25 M Ca  $(OH)_2$  is needed to react with 25.0 mL of 0.100 M H Cl ?

 $2 \operatorname{HCI}(aq) + \operatorname{Ca}(OH)_{2}(aq) \longrightarrow \operatorname{CaCI}_{2}(aq) + 2\operatorname{HOH}(I)$ 

## **HW: Solution Stoichiometry**

What volume of 2.25 M Ca  $(OH)_2$  is needed to react with 25.0 mL of 0.100 M H Cl ?

 $2 \operatorname{HCI}(aq) + \operatorname{Ca(OH)}_{2(}aq) \longrightarrow \operatorname{CaCI}_{2}(aq) + 2\operatorname{HOH}(I)$ 

25.0 mL x 0.100 moles H Cl x  $1 \text{ mol Ca(OH)}_2$  x  $1000 \text{ mL Ca(OH)}_2$ HCl soln 1000 mL HCl soln 2 mol H Cl 2.25 moles Ca(OH) $_2$ = 0.556 mL Ca(OH) $_2$  solution **Measuring the Concentration of a Solution: Titration** 

**Titration**: A procedure for determining the concentration using a second solution with known concentration

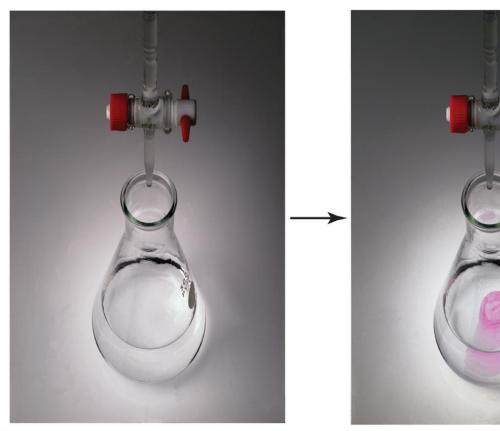
 $HCI(aq) + NaOH(aq) \longrightarrow NaCI(aq) + H_2O(l)$ 

Short cut: use  $M_{acid}V_{acid} = M_{base}V_{base}$ 

How can you tell when the reaction is complete? Equivalence point – point where you have the same amount H<sup>+</sup> and OH<sup>-</sup>

Indicator: Used to mark the equivalence point changes color at the equivalence point

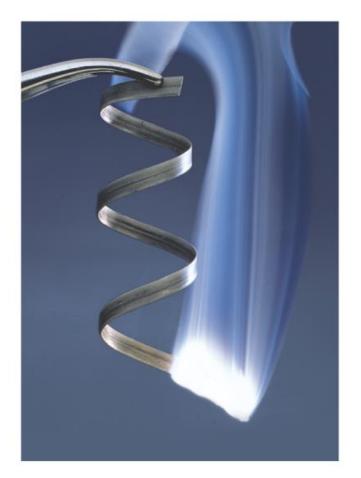
### Measuring the Concentration of a Solution: Titration



A measured volume of acid solution is placed in a flask, and phenolphthalein indicator is added.

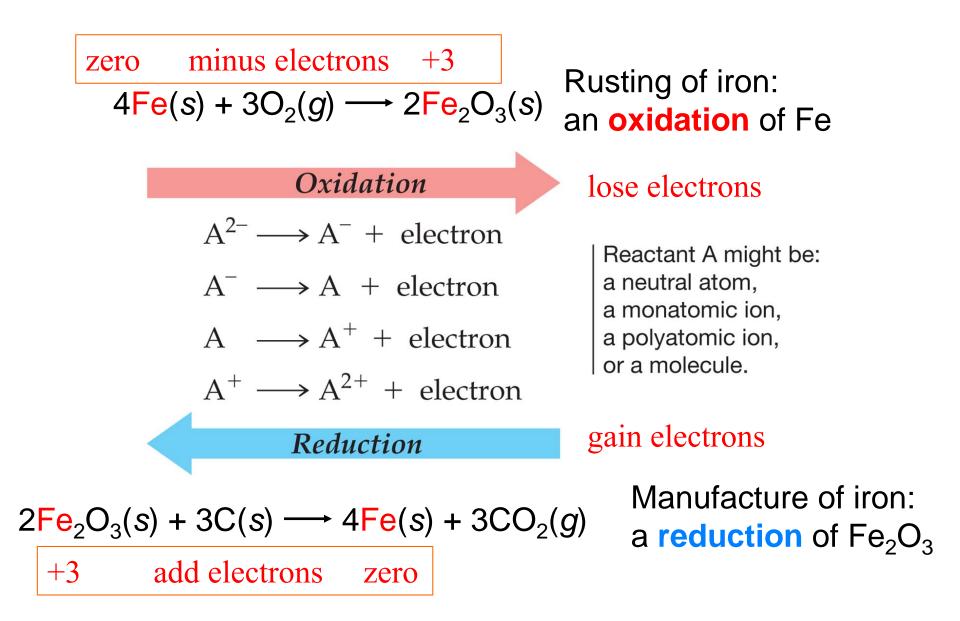
Base solution of known concentration is added from a buret until the indicator changes **color**. Reading the volume of base from the buret allows calculation of the acid concentration.

$$2Mg(s) + O_2(g) \longrightarrow 2MgO(s)$$



$$2P(s) + 3Br_2(l) \longrightarrow 2PBr_3(l)$$





**Oxidation**: The loss of one or more electrons by a substance, whether element, compound, or ion

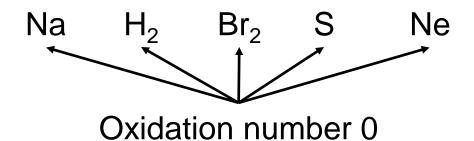
**Reduction**: The gain of one or more electrons by a substance (memorize as add negative – reduced ox number)

**Oxidation–Reduction (Redox) Reaction**: Any process in which electrons are transferred from one substance to another

**Oxidation Number (State)**: A value that indicates whether an atom is neutral, electron-rich, or electron-poor (These rules are on the memorize list on departmental syllabus.)

#### **Rules for Assigning Oxidation Numbers**

1. An atom in its elemental state has an oxidation number of 0 (zero).



2. An atom in a monatomic ion has an oxidation number identical to its charge.

(group 1A to 3A: group #) or

(<u>group 7A to 5A</u>: group # - 8)

- 1A2A3A7A6A5ANa+Ca<sup>2+</sup>Al<sup>3+</sup>Cl<sup>-</sup> $O^{2^-}$  $N^{3^-}$
- +1 +2 +3 -1 -2 -3

3. An atom in a polyatomic ion or in a molecular compound usually has the same oxidation number it would have if it were a monatomic ion.

a) Hydrogen can be either +1 or -1.

b) Oxygen usually has an oxidation number of -2.

$$\begin{array}{cccc} H - O - H & H - O - O - H \\ \swarrow & \uparrow & \uparrow & \uparrow & \uparrow & \uparrow \\ +1 & -2 & +1 & +1 & -1 & -1 & +1 \end{array}$$

3. c) Halogens usually have an oxidation number of -1.

$$\begin{array}{c} H - CI \\ \uparrow \\ +1 \\ -1 \end{array} \qquad \begin{array}{c} CI - O - CI \\ / \\ +1 \\ -2 \\ +1 \end{array}$$

4. The sum of the oxidation numbers is 0 for a neutral compound and is equal to the net charge for a polyatomic ion.

 $H_2SO_4$ 

End 10/16 W D section

$$2(+1) + (?) + 4(-2) = 0$$
 (net charge)  
S = ? = +6

$$Cr_2O_7^{2-}$$
 2(?) + 7(-2) = -2 (net charge)  
/ 1  
2 -2

# What is the oxidation state of the following.

a. Na element zero b. H<sub>2</sub> element zero c. Na Cl Na+1(gp 1A) 7-8 = -1 (gp 7A) d. HNO<sub>3</sub> H+1(gp 1A), O -2 (gp 6A)

- ox state N = variable N, calculate
  - zero = +1 + N + 3\*(-2)

algebra N = +6 - 1 = 5

# HW: What is the oxidation state of the following.

b. Fe

a.  $Cl_2$ 

End 10/16 F section, G section

c. Al  $Cl_3$ 

d.  $PO_4^{-3}$ 

# **Identifying Redox Reactions**

# **Oxidizing Agent (is reduced)**

Causes oxidation Gains one or more electrons Undergoes reduction Oxidation number of atom decreases. (becomes more negative)

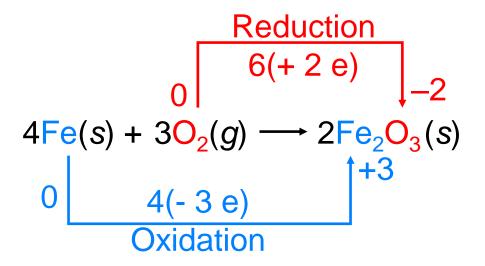
# **Reducing Agent (is oxidized)**

Causes reduction Loses one or more electrons Undergoes oxidation Oxidation number of atom increases. (becomes more positive)

# **Identifying Redox Reactions**

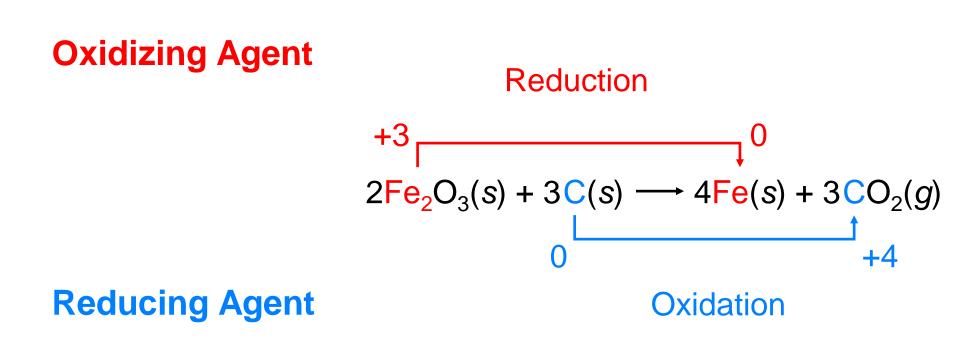
this chapter - only responsible for recognizing oxidation/reduction & oxidizing agent & reducing agent – NOT for balancing redox equations (chapter 18)

# **Oxidizing Agent**



**Reducing Agent** 

# **Identifying Redox Reactions**



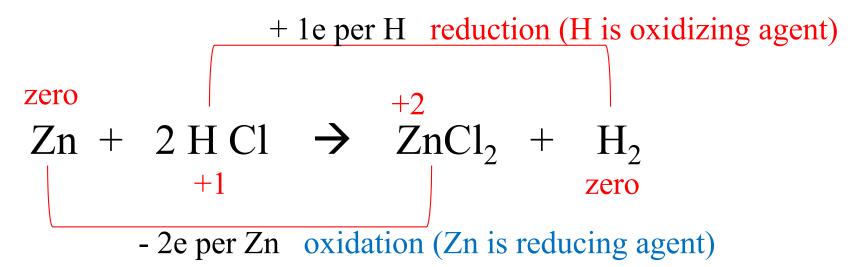
HW: Oxidation or Reduction (reducing agent / oxidizing agent)

- (a) Assign ox states of Zn & H
- (b) assign oxidation or reduction to brackets

# $Zn + 2 H Cl \rightarrow ZnCl_2 + H_2$

HW: Oxidation or Reduction (reducing agent / oxidizing agent)

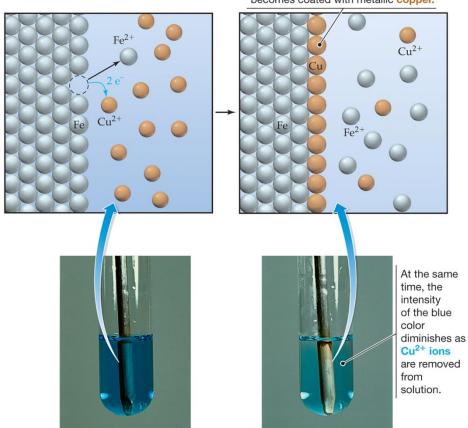
- (a) Assign ox states of Zn & H
- (b) assign oxidation or reduction to brackets



10/17 R - D section

### Iron loses electron to copper

 $Fe(s) + Cu^{2+}(aq) \longrightarrow Fe^{2+}(aq) + Cu(s)$ 



The iron nail reduces Cu<sup>2+</sup> ions and becomes coated with metallic copper.

 $Fe(s) + Cu^{2+}(aq) \longrightarrow Fe^{2+}(aq) + Cu(s)$ 

#### TABLE 4.5 A Partial Activity Series of the Elements

#### **Oxidation Reaction**

Strongly reducing

These elements react rapidly with aqueous H<sup>+</sup> ions (acid) or with liquid H<sub>2</sub>O to release H<sub>2</sub> gas.

These elements react with aqueous  $H^+$  ions or with steam to release  $H_2$  gas.

These elements react with aqueous  $H^+$  ions to release  $H_2$  gas.

These elements do not react with aqueous  $H^+$  ions to release  $H_2$ .

$$\begin{cases} \operatorname{Li} \rightarrow \operatorname{Li}^{+} + e^{-} \\ \operatorname{K} \rightarrow \operatorname{K}^{+} + e^{-} \\ \operatorname{Ba} \rightarrow \operatorname{Ba}^{2+} + 2 e^{-} \\ \operatorname{Ca} \rightarrow \operatorname{Ca}^{2+} + 2 e^{-} \\ \operatorname{Ca} \rightarrow \operatorname{Na}^{+} + e^{-} \end{cases}$$
$$\begin{cases} \operatorname{Mg} \rightarrow \operatorname{Mg}^{2+} + 2 e^{-} \\ \operatorname{Al} \rightarrow \operatorname{Al}^{3+} + 3 e^{-} \\ \operatorname{Mn} \rightarrow \operatorname{Mn}^{2+} + 2 e^{-} \\ \operatorname{Cr} \rightarrow \operatorname{Cr}^{3+} + 3 e^{-} \\ \operatorname{Cr} \rightarrow \operatorname{Cr}^{3+} + 3 e^{-} \\ \operatorname{Fe} \rightarrow \operatorname{Fe}^{2+} + 2 e^{-} \\ \operatorname{Cr} \rightarrow \operatorname{Cr}^{3+} + 3 e^{-} \end{cases}$$
$$\begin{cases} \operatorname{Co} \rightarrow \operatorname{Co}^{2+} + 2 e^{-} \\ \operatorname{Ni} \rightarrow \operatorname{Ni}^{2+} + 2 e^{-} \\ \operatorname{Sn} \rightarrow \operatorname{Sn}^{2+} + 2 e^{-} \\ \operatorname{Sn} \rightarrow \operatorname{Sn}^{2+} + 2 e^{-} \\ \operatorname{Sn} \rightarrow \operatorname{Sn}^{2+} + 2 e^{-} \\ \operatorname{Hg} \rightarrow \operatorname{Hg}^{2+} + 2 e^{-} \\ \operatorname{Au} \rightarrow \operatorname{Au}^{3+} + 3 e^{-} \end{cases}$$

Elements that are higher up in the table (reaction goes  $\rightarrow$ ) are more likely to be oxidized (lose e).

Thus, any element higher in the activity series will reduce (reducing agent) (lose e to) the ion of any element lower in the activity series.

Weakly reducing

# The Activity Series of the Elements $Cu(s) + 2Ag^{+}(aq) \longrightarrow Cu^{2+}(aq) + 2Ag(s)$ (a)

 $2Ag(s) + Cu^{2+}(aq) \rightarrow 2Ag^{+}(aq) + Cu(s)$  (b) Which one of these reactions will occur?

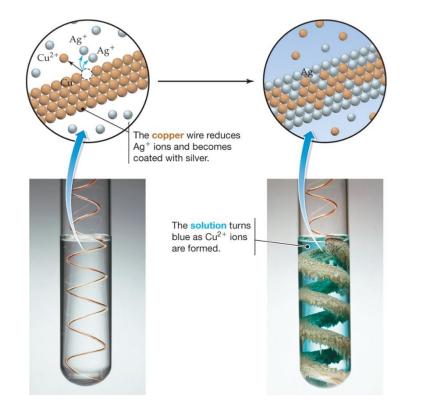
TABLE 4.5	A Partial Activity Series of the Eleme		
Oxidation Rea	action		Cu is above Ag
Strongly reducing Weakly reducing	These elements react rapidly with aqueous $H^+$ ions (acid) or with liquid $H_2O$ to release $H_2$ gas.	$\begin{cases} Li \rightarrow Li^{+} + e^{-} \\ K \rightarrow K^{+} + e^{-} \\ Ba \rightarrow Ba^{2+} + 2e^{-} \\ Ca \rightarrow Ca^{2+} + 2e^{-} \\ Na \rightarrow Na^{+} + e^{-} \end{cases}$	in activity series $Cu \rightarrow Ag \leftarrow$
	These elements react with aqueous $H^+$ ions or with steam to release $H_2$ gas.	$\left\{ \begin{array}{c} Mg \longrightarrow Mg^{2+} + 2 \ e^- \\ Al \longrightarrow Al^{3+} + 3 \ e^- \\ Mn \longrightarrow Mn^{2+} + 2 \ e^- \\ Zn \longrightarrow Zn^{2+} + 2 \ e^- \\ Cr \longrightarrow Cr^{3+} + 3 \ e^- \\ Fe \longrightarrow Fe^{2+} + 2 \ e^- \end{array} \right.$	
	These elements react with aqueous $H^+$ ions to release $H_2$ gas.	$\begin{cases} Co \rightarrow Co^{2+} + 2e^{-} \\ Ni \rightarrow Ni^{2+} + 2e^{-} \\ Sn \rightarrow Sn^{2+} + 2e^{-} \\ H_{2} \rightarrow 2H^{+} + 2e^{-} \end{cases}$	
	These elements do not react with aqueous $H^+$ ions to release $H_2$ .	$\begin{cases} Cu \rightarrow Cu^{2+} + 2e^{-} \\ Ag \rightarrow Ag^{+} + e^{-} \\ Hg \rightarrow Hg^{2+} + 2e^{-} \\ Pt \rightarrow Pt^{2+} + 2e^{-} \\ Au \rightarrow Au^{3+} + 3e^{-} \end{cases}$	

 $Cu(s) + 2Ag^{+}(aq) \longrightarrow Cu^{2+}(aq) + 2Ag(s)$  (a) \*

 $2Ag(s) + Cu^{2+}(aq) \longrightarrow 2Ag^{+}(aq) + Cu(s)$  (b) Which one of these reactions will occur?

TABLE 4.5	A Partial Activity Series of the Elements		
Oxidation Rea	action		Cu is above Ag
Strongly reducing Weakly reducing	These elements react rapidly with aqueous $H^+$ ions (acid) or with liquid $H_2O$ to release $H_2$ gas.	$\begin{cases} \text{Li} \rightarrow \text{Li}^{+} + e^{-} \\ \text{K} \rightarrow \text{K}^{+} + e^{-} \\ \text{Ba} \rightarrow \text{Ba}^{2+} + 2e^{-} \\ \text{Ca} \rightarrow \text{Ca}^{2+} + 2e^{-} \\ \text{Na} \rightarrow \text{Na}^{+} + e^{-} \end{cases}$	in activity series $Cu \rightarrow Ag \leftarrow$
	These elements react with aqueous $H^+$ ions or with steam to release $H_2$ gas.	$\begin{cases} Mg \longrightarrow Mg^{2+} + 2e^{-} \\ Al \longrightarrow Al^{3+} + 3e^{-} \\ Mn \longrightarrow Mn^{2+} + 2e^{-} \\ Zn \longrightarrow Zn^{2+} + 2e^{-} \\ Cr \longrightarrow Cr^{3+} + 3e^{-} \\ Fe \longrightarrow Fe^{2+} + 2e^{-} \end{cases}$	
	These elements react with aqueous $H^+$ ions to release $H_2$ gas.	$\begin{cases} Co \rightarrow Co^{2+} + 2e^{-} \\ Ni \rightarrow Ni^{2+} + 2e^{-} \\ Sn \rightarrow Sn^{2+} + 2e^{-} \\ H_{2} \rightarrow 2H^{+} + 2e^{-} \end{cases}$	
	These elements do not react with aqueous $H^+$ ions to release $H_2$ .	$\begin{cases} Cu \rightarrow Cu^{2+} + 2e^{-} \\ Ag \rightarrow Ag^{+} + e^{-} \\ Hg \rightarrow Hg^{2+} + 2e^{-} \\ Pt \rightarrow Pt^{2+} + 2e^{-} \\ Au \rightarrow Au^{3+} + 3e^{-} \end{cases}$	

 $Cu(s) + 2Ag^{+}(aq) \longrightarrow Cu^{2+}(aq) + 2Ag(s)$ 

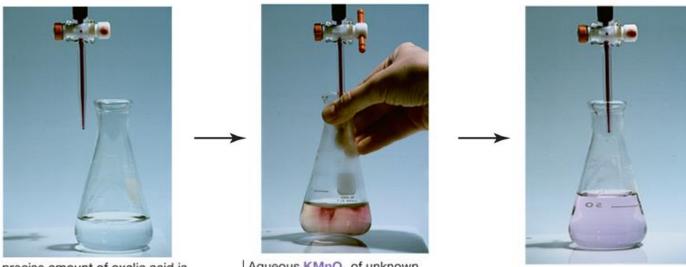


<u>copper</u> is above <u>silver</u> in activity series chart

(copper loses electron and silver gets electron)

# **Redox Titrations**

# $5H_2C_2O_4(aq) + 2MnO_4^{-}(aq) + 6H^{+}(aq)$ $\longrightarrow 10CO_2(g) + 2Mn^{2+}(aq) + 8H_2O(l)$



A precise amount of oxalic acid is weighed and dissolved in aqueous  $H_2SO_4$ .

Aqueous KMnO<sub>4</sub> of unknown concentration is added from a buret until ...

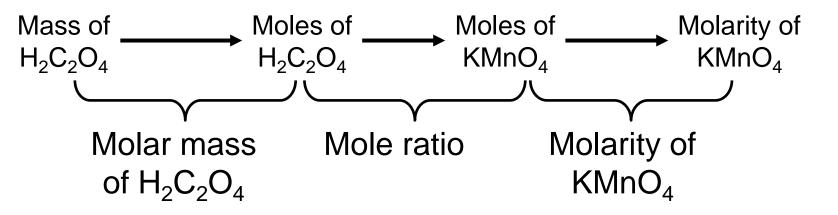
... the purple color persists, indicating that all of the oxalic acid has reacted.

# **Redox Titrations** (probably will not have on exam/quiz)

A solution is prepared with 0.2585 g of oxalic acid,  $H_2C_2O_4$ . 22.35 mL of an unknown solution of potassium permanganate is needed to titrate the solution. What is the concentration of the potassium permanganate solution? (already know how to do this: if given balanced reaction, just stoichiometry & molarity problem)

$$5H_2C_2O_4(aq) + 2MnO_4^{-}(aq) + 6H^{+}(aq)$$

 $\rightarrow 10CO_2(g) + 2Mn^{2+}(aq) + 8H_2O(l)$ 



# **Redox Titrations**

$$5H_2C_2O_4(aq) + 2MnO_4^{-}(aq) + 6H^{+}(aq)$$
$$\longrightarrow 10CO_2(g) + 2Mn^{2+}(aq) + 8H_2O(I)$$

Moles of  $H_2C_2O_4$  available:

$$0.2585 \text{ g H}_2\text{C}_2\text{O}_4 \times \frac{1 \text{ mol}}{90.04 \text{ g}} = 0.002871 \text{ mol H}_2\text{C}_2\text{O}_4$$

Moles of KMnO<sub>4</sub> reacted:

 $0.002871 \text{ mol } \text{H}_2\text{C}_2\text{O}_4 \times \frac{2 \text{ mol } \text{KMnO}_4}{5 \text{ mol } \text{H}_2\text{C}_2\text{O}_4} = 0.001148 \text{ mol } \text{KMnO}_4$ 

# **Redox Titrations**

$$5H_2C_2O_4(aq) + 2MnO_4^{-}(aq) + 6H^+(aq)$$
$$\longrightarrow 10CO_2(g) + 2Mn^{2+}(aq) + 8H_2O(I)$$

### **Concentration of KMnO<sub>4</sub> solution**:

$$\frac{0.001148 \text{ mol KMnO}_4}{22.35 \text{ mL}} \times \frac{1000 \text{ mL}}{1 \text{ L}} = 0.05136 \text{ M KMnO}_4$$