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Chapter 4 Reactions in Aqueous Solutions

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4.1 General Properties of Aqueous Solutions

- Solution a homogeneous mixture
 - Solute: the component that is dissolved
 - Solvent: the component that does the dissolving
 - Generally, the component present in the greatest quantity is considered to be the solvent. *Aqueous* solutions are those in which *water* is the solvent.

- Electrolytes and Nonelectrolytes
 - *Electrolyte*: substance that dissolved in water produces a solution that conducts electricity
 - Contains ions

$$\operatorname{NaCl}(s) \xrightarrow{H_2O} \operatorname{Na}^+(aq) + \operatorname{Cl}^-(aq)$$

- Nonelectrolyte: substance that dissolved in water produces a solution that does not conduct electricity
 - Does not contain ions

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

 Dissociation - ionic compounds separate into constituent ions when dissolved in solution

$$\operatorname{NaCl}(s) \xrightarrow{H_2O} \operatorname{Na}^+(aq) + \operatorname{Cl}^-(aq)$$

 Ionization - formation of ions by molecular compounds when dissolved

$$HCl(g) \xrightarrow{H_2O} H^+(aq) + Cl^-(aq)$$

 $NH_3(g) + H_2O(l) \rightleftharpoons NH_4^+(aq) + OH^-(aq)$

- Strong and weak electrolytes
 - Strong Electrolyte: 100% dissociation
 - All water soluble ionic compounds, strong acids and strong bases
 - Weak electrolytes
 - Partially ionized in solution
 - Exist mostly as the molecular form in solution
 - Weak acids and weak bases

TABLE 4.1The Strong Acids

Acid	Ionization Equation
Hydrochloric acid	$\operatorname{HCl}(aq) \longrightarrow \operatorname{H}^+(aq) + \operatorname{Cl}^-(aq)$
Hydrobromic acid	$\operatorname{HBr}(aq) \longrightarrow \operatorname{H}^{+}(aq) + \operatorname{Br}^{-}(aq)$
Hydroiodic acid	$HI(aq) \longrightarrow H^+(aq) + I^-(aq)$
Nitric acid	$HNO_3(aq) \longrightarrow H^+(aq) + NO_3^-(aq)$
Chloric acid	$\text{HClO}_3(aq) \longrightarrow \text{H}^+(aq) + \text{ClO}_3^-(aq)$
Perchloric acid	$\operatorname{HClO}_4(aq) \longrightarrow \operatorname{H}^+(aq) + \operatorname{ClO}_4^-(aq)$
Sulfuric acid*	$H_2SO_4(aq) \longrightarrow H^+(aq) + HSO_4^-(aq)$
	$HSO_4^-(aq) \rightleftharpoons H^+(aq) + SO_4^{2-}(aq)$

*Note that although each sulfuric acid molecule has two ionizable hydrogen atoms, it only undergoes the first ionization completely, effectively producing one H⁺ ion and one HSO₄⁻ ion per H₂SO₄ molecule. The second ionization happens only to a very small extent. Examples of weak electrolytes

- Weak acids \longrightarrow HC₂H₃O_{2(aq)} C₂H₃O_{2⁻ (aq)} + H⁺ (aq)

- Weak bases \implies NH_{3 (aq)} + H₂O_(l) NH₄⁺ (aq) + OH⁻ (aq)

(Note: double arrows indicate a reaction that occurs in both directions - a state of *dynamic equilibrium* exists)

Method to Distinguish Types of Electrolytes







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strong electrolyte

Classify the following as nonelectrolyte, weak electrolyte or strong electrolyte – NaOH

strong electrolyte



nonelectrolyte



weak electrolyte

4.2 Precipitation Reactions

- **Precipitation** (formation of a solid from two aqueous solutions) occurs when product is insoluble
- Produce insoluble ionic compounds
- **Solubility** is the maximum amount of a solid that can dissolve in a given amount of solvent at a specified temperature
- Prediction based on solubility rules

TABLE 4.2	Solubility Guidelines: Soluble Compounds		
Water-Soluble Compounds		Insoluble Exceptions	
Compounds containing an alkali metal cation (Li ⁺ , Na ⁺ , K ⁺ , Rb ⁺ , Cs ⁺) or the ammonium ion (NH ₄ ⁺)			
Compounds contacetate ion (C_2H	taining the nitrate ion (NO_3^-) , ${}_3O_2^-)$, or chlorate ion (ClO_3^-)		
Compounds containing the chloride ion (Cl^-) , bromide ion (Br^-) , or iodide ion (I^-)		Compounds containing Ag^+ , Hg_2^{2+} , or Pb^{2+}	
Compounds cont	taining the sulfate ion (SO_4^{2-})	Compounds containing Ag^+ , Hg_2^{2+} , Pb^{2+} , Ca^{2+} , Sr^{2+} , or Ba^{2+}	

TABLE 4.3Solubility Guidelines: Insoluble Compounds

Water-Insoluble Compounds

Compounds containing the carbonate ion (CO_3^{2-}) , phosphate ion (PO_4^{3-}) , chromate ion (CrO_4^{2-}) , or sulfide ion (S^{2-})

Compounds containing the hydroxide ion (OH⁻)

Soluble Exceptions

Compounds containing Li⁺, Na⁺, K⁺, Rb⁺, Cs⁺, or NH₄⁺

Compounds containing Li^+ , Na^+ , K^+ , Rb^+ , Cs^+ , or Ba^{2+}

• *Hydration*: process by which water molecules remove and surround individual ions from the solid.



Identify the Precipitate

 $Pb(NO_3)_2(aq) + 2Nal(aq) \rightarrow 2NaNO_3(aq) + Pbl_2(s)$

Mixing Solutions of Pb(NO₃)₂ and NaCl



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 $-Ba(NO_3)_2$

soluble

– Agl

insoluble

 $-Mg(OH)_2$

insoluble

• *Molecular equation*: shows all compounds represented by their chemical formulas

 $Na_2SO_4(aq) + Ba(OH)_2(aq) \longrightarrow 2NaOH(aq) + BaSO_4(s)$

 Ionic equation: shows all strong electrolytes as ions and all other substances (nonelectrolytes, weak electrolytes, gases) by their chemical formulas

$$Na_{2}SO_{4}(aq) \longrightarrow 2Na^{+}(aq) + SO_{4}^{2-}(aq)$$
$$Ba(OH)_{2}(aq) \longrightarrow Ba^{2+}(aq) + 2OH^{-}(aq)$$
$$NaOH(aq) \longrightarrow Na^{+}(aq) + OH^{-}(aq)$$

Molecular equation:

 $Na_2SO_4(aq) + Ba(OH)_2(aq) \longrightarrow 2NaOH(aq) + BaSO_4(s)$

Ionic equation:

 $2\mathrm{Na}^{+}(aq) + \mathrm{SO}_{4}^{2-}(aq) + \mathrm{Ba}^{2+}(aq) + 2\mathrm{OH}^{-}(aq) \longrightarrow 2\mathrm{Na}^{+}(aq) + 2\mathrm{OH}^{-}(aq) + \mathrm{Ba}\mathrm{SO}_{4}(s)$

Net lonic equation: shows only the reacting species in the chemical equation

Eliminates spectator ions

 $2\mathrm{Na}^{+}(aq) + \mathrm{SO}_{4}^{2-}(aq) + \mathrm{Ba}^{2+}(aq) + 2\mathrm{OH}^{-}(aq) \longrightarrow 2\mathrm{Na}^{+}(aq) + 2\mathrm{OH}^{-}(aq) + \mathrm{Ba}\mathrm{SO}_{4}(s)$

Net ionic equation:

$$\operatorname{Ba}^{2^+}(aq) + \operatorname{SO}_4^{2^-}(aq) \longrightarrow \operatorname{BaSO}_4(s)$$

- Steps in writing a net ionic equation
 - Write the balanced molecular equation.
 - Predict products by exchanging cations and anions in reactants.
 - Separate strong electrolytes into ions.
 - Cancel spectator ions.
 - Use the remaining species to write the net ionic equation.

Aqueous solutions of silver nitrate and sodium sulfate are mixed. Write the net ionic reaction. Step 1:

 $2AgNO_{3}(aq)+Na_{2}SO_{4}(aq) \rightarrow 2NaNO_{3}(?)+Ag_{2}SO_{4}(?)$

Step 2: Use solubility table; all nitrates are soluble but silver sulfate is insoluble

 $2Ag^{+}(aq) + 2NO_{3}^{-}(aq) + 2Na^{+}(aq) + SO_{4}^{2-}(aq)$ $\rightarrow 2Na^{+}(aq) + 2NO_{3}^{-}(aq) + Ag_{2}SO_{4}(s)$

Step 3: Cancel spectators

 $2Ag^{+}(aq) + 2NO_{3}^{-}(aq) + 2Na^{+}(aq) + SO_{4}^{2-}(aq)$ $\rightarrow 2Na^{+}(aq) + 2NO_{3}^{-}(aq) + Ag_{2}SO_{4}(s)$

Step 4: Write the net ionic reaction

 $2Ag^{+}(aq) + SO_4^{2-}(aq) \rightarrow Ag_2SO_4(s)$

4.3 Acid-Base Reactions

- Termed neutralization reactions.
- Involve an acid and a base.
- A molecular compound (water) is a common product along with a salt (ionic compound).

Common Acids and Bases



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TABLE 4.4	Strong Acids and Strong	Bases	
Strong Acids	Strong Bases	Strong Acids	Strong Bases
HCl	LiOH	HClO ₃	CsOH
HBr	NaOH	$HClO_4$	Ca(OH) ₂
HI	КОН	H_2SO_4	Sr(OH) ₂
HNO ₃	RbOH		Ba(OH) ₂

All the other acids and bases are weak electrolytes (important for net ionic equations).

- Definitions of acids and bases
 - Arrhenius acid produces H⁺ in solution
 - –Arrhenius base produces OH⁻ in solution
 - -More inclusive definitions:
 - Brønsted acid proton donor
 - Brønsted base proton acceptor

- Examples of a weak base and weak acid
 - Ammonia with water:



• Hydrofluoric acid with water:



• Types of acids

- *Monoprotic*: one ionizable hydrogen

 $\mathrm{HCI} + \mathrm{H_2O} \rightarrow \mathrm{H_3O^+} + \mathrm{CI^-}$

- *Diprotic*: two ionizable hydrogens

$$\begin{array}{l} H_2SO_4 + H_2O \rightarrow H_3O^+ \ + HSO_4^- \\ HSO_4^- + H_2O \rightarrow H_3O^+ \ + SO_4^{2-} \end{array}$$

- **Triprotic**: three ionizable hydrogens

$\begin{array}{l} H_{3}PO_{4} + H_{2}O \rightarrow H_{3}O^{+} + H_{2}PO_{4}^{-} \\ H_{2}PO_{4}^{-} + H_{2}O \rightarrow H_{3}O^{+} + HPO_{4}^{2-} \\ HPO_{4}^{2-} + H_{2}O \rightarrow H_{3}O^{+} + PO_{4}^{3-} \end{array}$

 Polyprotic: generic term meaning more than one ionizable hydrogen Types of bases
– Monobasic: One OH⁻ group

$\rm KOH \rightarrow ~K^{+} ~+~OH^{-}$

– **Dibasic**: Two OH[–] groups

$Ba(OH)_2 \rightarrow Ba^{2+} + 2OH^-$

Acid-Base Neutralization

 Neutralization: Reaction between an acid and a base

Acid + Base \rightarrow Salt + Water

Molecular equation: HCI(aq) + NaOH(aq) \rightarrow NaCI(aq) + H₂O(l)

Ionic equation: H⁺(aq)+ Cl⁻(aq) + Ma⁺(aq) + OH⁻(aq) → Na⁺(aq) + Cl⁻(aq) + H₂O(/)

Net ionic equation: $H^+(aq) + OH^-(aq) \rightarrow H_2O(I)$

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Solutions of acetic acid and lithium hydroxide are mixed. Write the net ionic reaction.

 $HC_2H_3O_2(aq) + OH(aq) \longrightarrow C_2H_3O_2(aq) + H_2O(I)$

4.4 Oxidation-Reduction Reactions

- Often called "redox" reactions
- Electrons are transferred between the reactants
 - One substance is oxidized, loses electrons
 - Reducing agent
 - Another substance is reduced, gains electrons
 - Oxidizing agent
- Oxidation numbers change during the reaction

– Example

 $Zn(s) + CuSO_4(aq) \rightarrow ZnSO_4(aq) + Cu(s)$

 $Zn(s) + Cu^{2+}(aq) \rightarrow Zn^{2+}(aq) + Cu(s)$

- Zinc is losing 2 electrons and oxidized.
 - Reducing agent
 - $Zn(s) \rightarrow Zn^{2+}(aq) + 2e^{-}$
- Copper ions are gaining the 2 electrons.
 - Oxidizing agent
 - $Cu^{2+}(aq)$ + $2e^{-} \rightarrow Cu(s)$

Reaction of Cu and Zn^{2+} ions



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- Rules for assigning oxidation numbers
 - 1. Elements (uncombined) are 0. Al, N₂, He, Zn, Ag, Br₂, O₂, O₃
 - 2. Oxidation numbers must sum to the overall charge of the species.

$$SO_4^{2-} = -2$$
 (O is usually -2 so....)
? + 4(-2) = -2

Solve:
$$? - 8 = -2$$
 $? = + 6$ (S)

Guidelines for Assigning Oxidation Numbers

TABLE 4.5	Elements with Reliable Oxidation Numbers in Compounds or Polyatomic Ions		
Element	Oxidation Number	Exceptions	
Fluorine	-1		
Group 1A or 2A metal	+1 or $+2$, respectively		
Hydrogen	+1	Any combination with a Group 1A or 2A metal to form a metal hydride. Examples: LiH and CaH ₂ —the oxidation number of H is -1 in both examples.	
Oxygen	-2	Any combination with something higher on the list that necessitates its having a different oxidation number (see rule 2 for assigning oxidation numbers). Examples: H_2O_2 and KO_2 —the oxidation number of O for H_2O_2 is –1 and for KO_2 is –1/2.	
Group 7A (other than fluorin	-1	Any combination with something higher on the list that necessitates its having a different oxidation number (see rule 2 for assigning oxidation numbers). Examples: CIF, BrO_4^- , and IO_3^- —the oxidation numbers of Cl, Br, and I are +1, +7, and +5, respectively.	
Assign oxidation numbers for all elements in each species MgBr₂ Mg +2, Br -1 CIO_2^- CI +3, O -2

- Displacement reactions
 - A common reaction: active metal replaces (displaces) a metal ion from a solution

 $Mg(s) + CuCl_2(aq) \rightarrow Cu(s) + MgCl_2(aq)$

 The activity series of metals is useful in order to predict the outcome of the reaction.

TABLE 4.6Activity Series

Element	Oxidation Half-Reaction
Lithium	$Li \longrightarrow Li^+ + e^-$
Potassium	$K \longrightarrow K^+ + e^-$
Barium	$Ba \longrightarrow Ba^{2+} + 2e^{-}$
Calcium	$Ca \longrightarrow Ca^{2+} + 2e^{-}$
Sodium	$Na \longrightarrow Na^+ + e^-$
Magnesium	$Mg \longrightarrow Mg^{2+} + 2e^{-}$
Aluminum	$Al \longrightarrow Al^{3+} + 3e^{-}$
Manganese	$Mn \longrightarrow Mn^{2+} + 2e^{-}$
Zinc	$Zn \longrightarrow Zn^{2+} + 2e^{-}$
Chromium	$Cr \longrightarrow Cr^{3+} + 3e^{-}$
Iron	$Fe \longrightarrow Fe^{2+} + 2e^{-}$
Cadmium	$Cd \longrightarrow Cd^{2+} + 2e^{-}$
Cobalt	$Co \longrightarrow Co^{2+} + 2e^{-}$
Nickel	$Ni \longrightarrow Ni^{2+} + 2e^{-}$
Tin	$\operatorname{Sn} \longrightarrow \operatorname{Sn}^{2+} + 2e^{-}$
Lead	$Pb \longrightarrow Pb^{2+} + 2e^{-}$
Hydrogen	$H_2 \longrightarrow 2H^+ + 2e^-$
Copper	$Cu \longrightarrow Cu^{2+} + 2e^{-}$
Silver	$Ag \longrightarrow Ag^+ + e^-$
Mercury	$Hg \longrightarrow Hg^{2+} + 2e^{-}$
Platinum	$Pt \longrightarrow Pt^{2+} + 2e^{-}$
Gold	$Au \longrightarrow Au^{3+} + 3e^{-}$
0	

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- Balancing redox reactions
 - Electrons (charge) must be balanced as well as number and types of atoms
 - -Consider this net ionic reaction:

 $AI(s) + Ni^{2+}(aq) \rightarrow AI^{3+}(aq) + Ni(s)$

 The reaction appears balanced as far as number and type of atoms are concerned, but look closely at the charge on each side. $AI(s) + Ni^{2+}(aq) \rightarrow AI^{3+}(aq) + Ni(s)$

- Divide reaction into two half-reactions $Al(s) \rightarrow Al^{3+}(aq) + 3e^{-}$ $Ni^{2+}(aq) + 2e^{-} \rightarrow Ni(s)$
- Multiply by a common factor to equalize electrons (the number of electrons lost must equal number of electrons gained)

 $2 [Al(s) \rightarrow Al^{3+}(aq) + 3e^{-}]$ 3 [Ni²⁺(aq) + 2e⁻ \rightarrow Ni(s)] Cancel electrons and write balanced net ionic reaction

$$2AI(s) \rightarrow 2AI^{3+}(aq) + 6e^{-}$$

 $3Ni^{2+}(aq) + 6e^{-} \rightarrow 3Ni(s)$

$2AI(s) + 3Ni^{2+}(aq) \rightarrow 2AI^{3+}(aq) + 3Ni(s)$

Predict whether each of the following will occur. For the reactions that do occur, write a balanced net ionic reaction for each.

- Copper metal is placed into a solution of silver nitrate

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

A gold ring is accidentally dropped into a solution of hydrochloric acid
 No reaction occurs, gold is below

hydrogen on the activity series.

- Combination Reactions
 - Many combination reactions may also be classified as redox reactions
 - Consider:

Hydrogen gas reacts with oxygen gas

$2\mathrm{H}_2(g) + \mathrm{O}_2(g) \rightarrow 2\mathrm{H}_2\mathrm{O}(I)$

Identify the substance oxidized and the substance reduced.

- Decomposition reactions
 - Many decomposition reactions may also be classified as redox reactions
 - -Consider:
 - Potassium chlorate is strongly heated 2KClO₃(s) \rightarrow 2KCl(s) + 3O₂(g)

Identify substances oxidized and reduced.

- Disproportionation reactions
 - One element undergoes both oxidation and reduction
 - -Consider:



- Combustion reactions
 - Common example, hydrocarbon fuel reacts with oxygen to produce carbon dioxide and water
 - -Consider:



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1 1A	1																18 8A
1 H +1 -1												10				17	2 He
3 Li +1	2 2A 4 Be +2	(most common in red)									13 3A 5 B +3	14 4A 6 C +4 +2 -4	$ \begin{array}{r} 15 \\ 5A \\ 7 \\ N \\ +5 \\ +4 \\ +3 \\ +2 \\ +1 \\ -3 \\ \end{array} $	$ \begin{array}{c} 16 \\ 6A \\ 8 \\ 0 \\ +2 \\ -\frac{1}{2} \\ -1 \\ -2 \\ \end{array} $	17 7A 9 F -1	10 Ne	
11 Na +1	12 Mg +2	3 3B	4 4B	5 5B	6 6B	7 7B	8	9 —8B—	10	11 1B	12 2B	13 Al +3	14 Si +4 -4	15 P +5 +3 -3	$ \begin{array}{r} 16 \\ S \\ +6 \\ +4 \\ +2 \\ -2 \end{array} $	17 Cl +7 +6 +5 +4 +3 +1 -1	18 Ar
19 K +1	20 Ca +2	21 Sc +3	22 Ti +4 +3 +2	23 V +5 +4 +3 +2	24 Cr +6 +5 +4 +3 +2	25 Mn +7 +6 +4 +3 +2	26 Fe +3 +2	27 Co +3 +2	28 Ni +2	29 Cu +2 +1	30 Zn +2	31 Ga +3	32 Ge +4 -4	33 As +5 +3 -3	34 Se +6 +4 -2	35 Br +5 +3 +1 -1	36 Kr +4 +2
37 Rb +1	38 Sr +2	39 Y +	40 Zr +4	41 Nb +5 +4	42 Mo +6 +4 +3	43 Tc +7 +6 +4	44 Ru +8 +6 +4 +3	45 Rh +4 +3 +2	46 Pd +4 +2	47 Ag +1	48 Cd +2	49 In +3	50 Sn +4 +2	51 Sb +5 +3 -3	52 Te +6 +4 -2	53 I +7 +5 +1 -1	54 Xe +6 +4 +2
55 Cs +1	56 Ba +2	57 La +3	72 Hf +4	73 Ta +5	74 W +6 +4	75 Re +7 +6 +4	76 Os +8 +4	77 Ir +4 +3	78 Pt +4 +2	79 Au +3 +1	80 Hg +2 +1	81 TI +3 +1	82 Pb +4 +2	83 Bi +5 +3	84 Po +2	85 At -1	86 Rn

Oxidation Numbers on the Periodic Table

4.5 Concentration of Solutions

- **Concentration** is the amount of solute dissolved in a given amount of solvent.
- Qualitative expressions of concentration
 - –Concentrated higher ratio of solute to solvent
 - Dilute smaller ratio of solute to solvent

Comparison of a Concentrated and Dilute Solution





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- Quantitative concentration term
 - -*Molarity* is the ratio of moles solute per liter of solution

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

Symbols: *M* or []Different forms of molarity equation

$$M = \frac{mol}{L}$$
 $L = \frac{mol}{M}$ $mol = M \times L$

Calculate the molarity of a solution prepared by dissolving 45.00 grams of KI into a total volume of 500.0 mL. Calculate the molarity of a solution prepared by dissolving 45.00 grams of KI into a total volume of 500.0 mL.

 $\frac{45.00 \text{ g KI}}{500.0 \text{ mL}} \times \frac{1 \text{ mol KI}}{166.0 \text{ g KI}} \times \frac{1000 \text{ mL}}{1 \text{ L}} = 0.5422 M$

How many milliliters of 3.50 *M* NaOH can be prepared from 75.00 grams of the solid?

How many milliliters of 3.50 *M* NaOH can be prepared from 75.00 grams of the solid?

$75.00 \text{ g NaOH} \times \frac{1 \text{ mol NaOH}}{40.00 \text{ g NaOH}} \times \frac{1 \text{ L}}{3.50 \text{ mol NaOH}} \times \frac{1000 \text{ mL}}{1 \text{ L}} = 536 \text{ mL}$

Dilution

Process of preparing a less concentrated solution from a more concentrated one.

moles of solute before dilution = moles of solute after dilution

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

$$M_{\rm c} \times {\rm L}_{\rm c} = M_{\rm d} \times {\rm L}_{\rm d}$$

For the next experiment the class will need 250. mL of $0.10 M \text{CuCl}_2$. There is a bottle of 2.0 $M \text{CuCl}_2$. Describe how to prepare this solution. How much of the 2.0 M solution do we need? Concentrated: 2.0 M use ? mL (L_d) Dilute: 250. mL of 0.10 M

$$M_{\rm c}L_{\rm c} = M_{\rm d}L_{\rm d}$$

$$(2.0 M) (L_c) = (0.10 M) (250.mL)$$

 $L_c = 12.5 mL$

12.5 mL of the concentrated solution are needed; add enough distilled water to prepare 250. mL of the solution.

- Solution Stoichiometry
 - Soluble ionic compounds dissociate completely in solution.
 - -Using mole ratios we can calculate the concentration of all species in solution.
 - NaCl dissociates into Na⁺ and Cl⁻
 - Na_2SO_4 dissociates into $2Na^+$ and SO_4^{2-}
 - AICI₃ dissociates into AI³⁺ and 3CI⁻

Find the concentration of all species in a 0.25 *M* solution of MgCl₂

$$MgCl_2 \rightarrow Mg^{2+} + 2Cl^-$$

Given: $MgCl_2 = 0.25 M$

$$[Mg^{2+}] = 0.25 M (1:1 ratio)$$

 $[Cl^{-}] = 0.50 M (1:2 ratio)$

Using the square bracket notation, express the molar concentration for all species in the following solutions

0.42 *M* Ba(OH)₂ [Ba²⁺] = 0.42 *M* (1:1 ratio) [OH⁻] = 0.84 *M* (2:1 ratio)

4.6 Aqueous Reactions and Chemical Analysis

- Types of quantitative analysis
 - -Gravimetric analysis (mass analysis)
 - Example: precipitation reaction
 - -Volumetric analysis (volume analysis)
 - Example: titration

- Gravimetric Analysis
 - One form: isolation of a precipitate
 - Typical steps:
 - Determine mass of unknown solid
 - Dissolve unknown in water
 - Combine with excess amount of known substance to form a precipitate (excess drives reaction to completion)
 - Filter, dry and weigh the precipitate
 - Use formula and mass of ppt to find % of ion in unknown solid

A 0.825 g sample of an ionic compound containing chloride ions and an unknown metal is dissolved in water and treated with excess silver nitrate. If 1.725 g of AgCI precipitate forms, what is the percent by mass of CI in the original sample?

Steps in solution:

- Find the % of CI in AgCI
- Multiply the % of CI by the mass of the precipitate to obtain the CI in the sample
- Divide the mass of CI in sample by total mass of sample (multiply by 100 for %)







- Volumetric analysis
 - Commonly accomplished by *titration*
 - Addition of a solution of known concentration (standard solution) to another solution of unknown concentration.
 - Standardization is the determination of the exact concentration of a solution.
 - *Equivalence point* represents completion of the reaction.
 - *Endpoint* is where the titration is stopped.
 - An *indicator* is used to signal the endpoint.

Apparatus for a Titration



A student measured exactly 15.0 mL of an unknown monoprotic acidic solution and placed in an Erlenmeyer flask. An indicator was added to the flask. At the end of the titration the student had used 35.0 mL of 0.12 M NaOH to neutralize the acid. Calculate the molarity of the acid.



Calculate the molarity of 25.0 mL of a monoprotic acid if it took 45.50 mL of 0.25 *M* KOH to neutralize the acid.

$$\frac{0.25 \text{ mol KOH}}{L} \times 0.04550 \text{ L} \times \frac{1 \text{ mol acid}}{1 \text{ mol KOH}} = 0.01138 \text{ mol acid}$$

$$\frac{0.01138 \text{ mol acid}}{0.0250 \text{ L}} = 0.455 M$$

Key Points

- Electrolytes (strong, weak, and non)
- Precipitation reactions
 - Solubility rules
- Molecular, ionic, and net ionic reactions
- Acid-base neutralization reactions
- Oxidation-reduction reactions

Key Points

- Balancing redox reactions by the half reaction method
- Various types: decomposition, combination
- Molarity
- Solution stoichiometry
 - Gravimetric analysis
 - Volumetric analysis