and

Solution Stoichiometry

Aqueous Solutions

Water is the dissolving medium, or solvent.

Some Properties of Water

- Water is "bent" or V-shaped.
- The O-H bonds are covalent.
- Water is a polar molecule.
- Hydration occurs when salts dissolve in water.

Figure 4.1: (Left) The water molecule is polar. (Right) A space-filling model of the water molecule.





Figure 4.3: (a) The ethanol molecule contains a polar O—H bond similar to those in the water molecule. (b) The polar water molecule interacts strongly with the polar O—H bond in ethanol. This is a case of "like dissolving like."



A Solute

dissolves in water (or other "solvent")

- changes phase (if different from the solvent)
- is present in lesser amount (if the same phase as the solvent)

A Solvent

- retains its phase (if different from the solute)
- is present in greater amount (if the same phase as the solute)

Ions in Aqueous Solution

Ionic Theory of Solutions • Some molecular compounds dissolve but do not dissociate into ions.

$C_6H_{12}O_6(s)$ (glucose) $\xrightarrow{H_2O}$ $C_6H_{12}O_6(aq)$

 These compounds are referred to as nonelectrolytes. They dissolve in water to give a nonconducting solution.

Ions in Aqueous Solution

Ionic Theory of Solutions

 Many ionic compounds *dissociate* into independent ions when dissolved in water

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

- These compounds that "freely" dissociate into independent ions in aqueous solution are called <u>electrolytes.</u>
- Their aqueous solutions are capable of conducting an electric current.

Figure 4.2: Polar water molecules interact with the positive and negative ions of a salt assisting in the dissolving process.





Electrolytes Strong - conduct current efficiently NaCl, HNO₃

Weak - conduct only a small current vinegar, tap water

Non - no current flows pure water, sugar solution



Ions in Aqueous Solution

Ionic Theory of Solutions

- Electrolytes are substances that dissolve in water to give an electrically conducting solution.
 - Thus, in general, *ionic solids* that dissolve in water are *electrolytes*.
 - Some molecular compounds, such as acids, also dissociate in aqueous solution and are considered electrolytes.

 $HCl (aq) \rightarrow H^+ (aq) + Cl^- (aq)$





Weak acids - dissociate to a slight extent to give H^+ in solution

acetic and formic acid



Bases

Strong bases - react completely with water to give OH^- ions.

sodium hydroxide

Weak bases - react only slightly with water to give OH^- ions.

ammonia







Ions in Aqueous Solution

Ionic Theory of Solutions: Summary
In summary, substances that dissolve in water are either *electrolytes* or *nonelectrolytes*.

- *Nonelectrolytes* form nonconducting solutions because they *dissolve as molecules*.
- *Electrolytes* form conducting solutions because they *dissolve as ions*.

Ions in Aqueous Solution

Ionic Theory of Solutions: Summary • Electrolytes can be *strong* or *weak*.

- Almost all ionic substances that dissolve are *strong electrolytes*.
- Molecular substances that dissolve are either *nonelectrolytes* or *weak electrolytes*.

Working with Solutions

Molar Concentration

• *Molar concentration*, or *molarity (M)*, is defined as the moles of solute dissolved in one liter (cubic decimeter) of solution.

Molarity (M) = $\frac{\text{moles of solute}}{\text{liters of solution}}$

Molarity

Molarity (*M*) = moles of solute per volume of solution in liters:

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

 $3 M HCl = \frac{6 \text{ moles of HCl}}{2 \text{ liters of solution}}$

Common Terms of Solution Concentration

Stock - routinely used solutions prepared in concentrated form.

Concentrated - *relatively* large ratio of solute to solvent. (5.0 *M* NaCl)

Dilute - *relatively* small ratio of solute to solvent. (0.01 *M* NaCl)

Working with Solutions

- The majority of chemical reactions discussed here occur in aqueous solution.
 - When you run reactions in liquid solutions, it is convenient to dispense the amounts of reactants by measuring out *volumes* of reactant solutions.



Figure 4.10: Steps involved in the preparation of a standard aqueous solution.



Working with Solutions Molar Concentration

- When we dissolve a substance in a liquid, we call the substance the **solute** and the liquid the **solvent**.
 - The general term *concentration* refers to the quantity of solute in a standard quantity of solution.



Working with Solutions

How many moles of Cl⁻ are in 1.00 L of 1.36 M FeCl₃ solution?

 $\frac{1.36 \text{ motes EeCh}_3}{1.00 \text{ L-soln}} \times \frac{3 \text{ moles Cl}}{4 \text{ mole FeCl}_3} \times 1.00 \text{ L-soln} = 4.08 \text{ moles Cl}$

 $\operatorname{FeCl}_{3}(s) \rightarrow \operatorname{Fe}^{3+}(\operatorname{aq}) + 3 \operatorname{Cl}^{-}(\operatorname{aq})$ H₂O



From the equation $\frac{\text{moles of solute}}{\text{Liters of solution}} = \text{Molarity}$

 $\underline{Moles} = M$ or, rearranging: L x M = moles L

How many moles of NaCl are in 25.0 ml of 0.100 M NaCl solution?

First, convert ml to L: 25.0 ml x 10⁻³ L/ml = 0.0250 ml (0.0250 L soln) (0.100 moles NaCl/L soln) = 0.00250 moles NaCl

Working with Solutions

Diluting Solutions

 The *molarity* of a solution and its *volume* are inversely proportional. Therefore, adding water makes the solution less concentrated.
 This inverse relationship takes the form of: Moles = Moles

$$M_i \times V_i = M_f \times V_f$$

– So, as water is added, increasing the final volume, $V_{\rm f}$, the final molarity, $M_{\rm f}$, decreases.

How many liters of 0.100 M HCl will be needed to prepare 0.500 L of 0.0750 M HCl? Use $M_iV_i = M_fV_f$ We want to solve for V_i $M_i = 0.100$ M HCl $M_f = 0.0750$ M HCl $V_f = 0.500$ L HCl Solving the equation for V_i $V_i = (M_fV_f)/M_i = (0.0750$ M x 0.500 L)/(0.100 M) = 0.375 L



Types of Solution Reactions

Precipitation reactions $AgNO_3(aq) + NaCl(aq) \rightarrow AgCl(s) +$

 $NaNO_3(aq)$

Acid-base reactions

 $NaOH(aq) + HCl(aq) \rightarrow NaCl(aq) + H_2O(l)$

Oxidation-reduction reactions

 $\operatorname{Fe}_2\operatorname{O}_3(s) + 2\operatorname{Al}(s) \rightarrow 2\operatorname{Fe}(s) + \operatorname{Al}_2\operatorname{O}_3(s)$

Ions in Aqueous Solution

Molecular and Ionic Equations
A molecular equation is one in which the reactants and products are written as if they were molecules, even though they may actually exist in solution as ions.

 $Ca(OH)_2(aq) + Na_2CO_3(aq) \rightarrow CaCO_3(s) + 2NaOH(aq)$

- Note that Ca(OH)₂, Na₂CO₃, and NaOH are all soluble compounds but CaCO₃ is not.

Simple Rules for Solubility

- 1. Most nitrate (NO3-) salts are soluble.
- 2. Most alkali (group 1A) salts and $\mathrm{NH_4^+}$ are soluble.
- 3. Most Cl⁻, Br⁻, and I⁻ salts are soluble (NOT Ag⁺, Pb²⁺, Hg₂²⁺)
- 4. Most sulfate salts are soluble (NOT BaSO₄, PbSO₄, HgSO₄, CaSO₄)
- 5. Most OH⁻ salts are only slightly soluble (NaOH, KOH are soluble, Ba(OH)₂, Ca(OH)₂ are marginally soluble)
- 6. Most S^{2–}, CO₃^{2–}, CrO₄^{2–}, PO₄^{3–} salts are only slightly soluble.

Ions in Aqueous Solution

Molecular and Ionic Equations
An ionic equation, however, represents strong electrolytes as separate independent ions. This is a more accurate representation of the way electrolytes behave in solution.

Ca²⁺(aq) + 2OH⁻(aq) + 2Na⁺(aq) + CO₃²⁻(aq) →CaCO₃(s) ↓ +2Na⁺(aq) + 2OH⁻(aq)

Ions in Aqueous Solution

Molecular and Ionic Equations

Complete and net ionic equations

 A complete ionic equation is a chemical equation in which strong electrolytes (such as soluble ionic compounds) are written as separate ions in solution.

$$Ca (NO_{3})_{2}(aq) + K_{2}CO_{3}(aq) \rightarrow CaCO_{3}(s) + 2KNO_{3}(aq)$$
(strong)
(strong)
(strong)
(arcond)
(strong)
(stro

 Ions in Aqueous Solution Molecular and Ionic Equations

 Complete and net ionic equations.
 A net ionic equation is a chemical equation from which the spectator ions have been removed.
 A spectator ion is an ion in an ionic equation that does not take part in the reaction.

 Ca²⁺(aq) + 2³/₃ (aq) + 2⁴/₄ (aq) + CO₃²⁻(aq) → CaCO₃(s) + 2⁴/₄ (aq) + 2³/₉ (aq)

Ions in Aqueous Solution

Molecular and Ionic Equations

- Complete and net ionic equations
 - Separating the strong electrolytes into separate ions, we obtain the complete ionic equation.

$2H^+(aq) + 2NO_3^-(aq) + Mg(OH)_2(s) \rightarrow$

$2H_2O(l) + Mg^{2+}(aq) + 2NO_3(aq)$

- Note that the nitrate ions did not participate in the reaction. These are *spectator ions*.

Ions in Aqueous Solution

Molecular and Ionic Equations

• Complete and net ionic equations

- Eliminating the spectator ions results in the net ionic equation.

 $2H^+(aq) + 2NO_3^-(aq) + Mg(OH)_2(s) \rightarrow$

$2H_2O(l) + Mg^{2+}(aq) + 2NO_3(aq)$

$2H^+(aq) + Mg(OH)_2(s) \rightarrow 2H_2O(l) + Mg^{2+}(aq)$

This equation represents the "essential" reaction.

TABLE 4.1 Simple Rules for the Solubility of Salts in Water

- 1. Most nitrate (NO3-) salts are soluble.
- 2. Most salts containing the alkali metal ions (Li*, Na*, K*, Cs*, Rb*) and the ammonium ion (NH_4^+) are soluble.
- 3. Most chloride, bromide, and iodide salts are soluble. Notable exceptions are salts containing the ions Ag $^{\rm i}$, Pb^2 $^{\rm i}$, and Hg $_2^{2\,\rm i}$.
- 4. Most sulfate salts are soluble. Notable exceptions are $BaSO_4,\,PbSO_4,\,Hg_2SO_4,\,and\,CaSO_4,$
- Most hydroxide salts are only slightly soluble. The important soluble hydroxides are NaOH and KOH. The compounds Ba(OH)₂, Sr(OH)₂, and Ca(OH)₂ are marginally soluble.
- 6. Most sulfide (S^2-), carbonate (CO₃²⁻), chromate (CrO₄²⁻), and phosphate (PO₄³⁻) salts are only slightly soluble.





Figure 4.17: The reaction of KCl(aq) with $AgNO_3$ to form AgCl(s).



Describing Reactions in Solution

1. Molecular equation (reactants and products as compounds)

 $AgNO_3(aq) + KCl(aq) \rightarrow AgCl(s) + KNO_3(aq)$

2. Complete ionic equation (all strong electrolytes shown as ions)

 $\begin{array}{l} \operatorname{Ag^{+}}(aq) + \operatorname{NO_{3}^{-}}(aq) + \operatorname{K^{+}}(aq) + \operatorname{Cl^{-}}(aq) \rightarrow \\ \operatorname{AgCl}(s) + \operatorname{K^{+}}(aq) + \operatorname{NO_{3}^{-}}(aq) \end{array}$

Describing Reactions in Solution (continued)

3. Net ionic equation (show only components that actually react)

 $Ag^+(aq) + Cl^-(aq) \rightarrow AgCl(s)$

 K^+ and NO_3^- are spectator ions.

Types of Chemical Reactions

Acid-Base Reactions

- The Arrhenius Concept
 - The Arrhenius concept defines *acids* as substances that produce hydrogen ions, H⁺, when dissolved in water.
 - An example is nitric acid, HNO₃, a molecular substance that dissolves in water to give H⁺ and NO₃⁻.

 $HNO_3(aq) \xrightarrow{H_2O} H^+(aq) + NO_3(aq)$

Types of Chemical Reactions

Acid-Base Reactions

- · The Arrhenius Concept
 - The Arrhenius concept defines *bases* as *substances that produce hydroxide ions, OH*, *when dissolved in water.*
 - An example is sodium hydroxide, NaOH, an ionic substance that dissolves in water to give sodium ions and hydroxide ions.

 $NaOH(s) \xrightarrow{H_2O} Na^+(aq) + OH^-(aq)$

Types of Chemical Reactions

Acid-Base Reactions

- The Arrhenius Concept
 - The molecular substance ammonia, NH₃, is a base in the Arrhenius view,

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

because it yields hydroxide ions when it reacts with water.

Types of Chemical Reactions

Acid-Base Reactions

- The Brønsted-Lowry Concept
 - The Brønsted-Lowry concept of acids and bases involves the transfer of a proton (H⁺) from the acid to the base.
 - In this view, acid-base reactions are *proton-transfer reactions*.

Types of Chemical Reactions Acid-Base Reactions

- · The Brønsted-Lowry Concept
 - The Brønsted-Lowry concept defines an acid as the species (molecule or ion) that donates a proton (H⁺) to another species in a protontransfer reaction.
 - A base is defined as the species (molecule or ion) that accepts the proton (H⁺) in a protontransfer reaction.

• The Brønsted-Lowry Concept

In the reaction of ammonia with water,

 $NH_3(aq) + H_2O(l) \stackrel{\leftarrow}{\rightarrow} NH_4^+(aq) + OH^-(aq)$

the H_2O molecule is the acid because it donates a proton. The NH_3 molecule is a base, because it accepts a proton.

Types of Chemical Reactions

Acid-Base Reactions

The Brønsted-Lowry Concept

The $H^+_{(aq)}$ ion associates itself with water to form $H_3O^+_{(aq)}$.

 $H^+(aq) + H_2O(l) \rightarrow H_3O^+(aq)$

This "mode of transportation" for the H⁺ ion is called the hydronium ion.

Types of Chemical Reactions

Acid-Base Reactions

The Brønsted-Lowry Concept

The dissolution of nitric acid, HNO₃, in water is therefore a proton-transfer reaction

 $HNO_3(aq) + H_2O(l) \rightarrow NO_3^{-}(aq) + H_3O^{+}(aq)$

H+

where HNO_3 is an acid (proton donor) and H_2O is a base (proton acceptor).

Types of Chemical Reactions

Acid-Base Reactions

• In summary, the Arrhenius concept and the Brønsted-Lowry concept are essentially the same in aqueous solution.

 The Arrhenius concept acid: proton (H⁺) donor

base: hydroxide ion (OH-) donor

Types of Chemical Reactions

Acid-Base Reactions

- In summary, the Arrhenius concept and the Brønsted-Lowry concept are essentially the same in aqueous solution.
 - The Brønsted-Lowry concept acid: proton (H⁺) donor

base: proton (H⁺) acceptor

Types of Chemical Reactions

• Strong and Weak Acids and Bases

 A strong acid is an acid that ionizes completely in water; it is a strong electrolyte.

 $HNO_3(aq) + H_2O(l) \rightarrow NO_3(aq) + H_3O(aq)$

$HCl(aq) + H_2O(l) \rightarrow Cl^-(aq) + H_3O^+(aq)$

- <u>Table 4.3</u> lists the common strong acids.

Types of Chemical Reactions Acid-Base Reactions

- Strong and Weak Acids and Bases
 - A strong base is a base that is present *entirely* as ions, one of which is OH⁻; it is a *strong* electrolyte.

$NaOH(s) \xrightarrow{H_2O} Na^+(aq) + OH^-(aq)$

 The hydroxides of Group IA and IIA elements, except for beryllium hydroxide, are strong bases. (see <u>Table 4.3</u>)

Strong Acids	Strong Bases
HClO ₄	LiOH
H_2SO_4	NaOH
HI	KOH
HBr	Ca(OH) ₂
HCl	Sr(OH) ₂
HNO ₃	Ba(OH) ₂

Types of Chemical Reactions

Acid-Base Reactions

- Strong and Weak Acids and Bases
 - You will find it important to be able to identify an acid or base as strong or weak.
 - When you write an ionic equation, strong acids and bases are represented as separate ions.
 - Weak acids and bases are represented as undissociated "molecules" in ionic equations.

Types of Chemical Reactions

Acid-Base Reactions

- · Strong and Weak Acids and Bases
 - A weak acid is an acid that only *partially* ionizes in water; it is a *weak* electrolyte.
 - The hydrogen cyanide molecule, HCN, reacts with water to produce a <u>small percentage of</u> <u>ions</u> in solution.

 $HCN(aq) + H_2O(1) \overleftrightarrow{} CN^{-}(aq) + H_3O^{+}(aq)$

Types of Chemical Reactions Acid-Base Reactions

• Strong and Weak Acids and Bases

- A weak base is a base that is only *partially* ionized in water; it is a *weak* electrolyte.
- Ammonia, NH₃, is an example.

 $NH_3(aq) + H_2O(l) \stackrel{\leftarrow}{\rightarrow} NH_4^+(aq) + OH^-(aq)$

Types of Chemical Reactions

Acid-Base Reactions

- Neutralization Reactions – One of the chemical properties of acids and bases is that *they neutralize one another*.
 - A *neutralization reaction* is a reaction of an acid and a base that results in an ionic compound and water.
 - The ionic compound that is the product of a neutralization reaction is called a *salt*.

 $\frac{\text{HCN}(aq) + \text{KOH}(aq) \rightarrow \text{KCN}(aq) + \text{H}_2\text{O}(l)}{\text{salt}}$





Titrant - solution of known concentration used in titration

Analyte - substance being analyzed

Equivalence point - enough titrant added to react exactly with the analyte

Endpoint - the indicator changes color so you can tell the equivalence point has been reached.





Quantitative Analysis

*Volumetric Analysis*Consider the reaction of sulfuric acid, H₂SO₄, with sodium hydroxide, NaOH:

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

- Suppose a beaker contains 35.0 mL of 0.175 M H₂SO₄. How many milliliters of 0.250 M NaOH must be added to completely react with the sulfuric acid?

Quantitative Analysis

Volumetric Analysis

- After balancing the equation, we must convert the 0.0350 L (35.0 mL) to moles of H_2SO_4 (using the molarity of the H_2SO_4).
- Then, convert to moles of NaOH (from the balanced chemical equation).
- Finally, convert to volume of NaOH solution (using the molarity of NaOH).

 $(0.0350L) \times \frac{0.175 \text{ mole } H_2SO_4}{1 \text{ L} \text{ H}_2SO_4 \text{ solution}} \times \frac{2 \text{ mol } \text{ NaOH}}{1 \text{ mol } \text{ H}_2SO_4} \times \frac{1 \text{ L} \text{ NaOH soln.}}{0.250 \text{ mol } \text{ NaOH}} = 0.0490 \text{ L} \text{ NaOH solution} (\text{ or } 49.0 \text{ mL of } \text{ NaOH solution})$

Performing Calculations for Acid-Base Reactions

- 1. List initial species and predict reaction.
- 2. Write balanced net ionic reaction.
- 3. Calculate moles of reactants.
- 4. Determine limiting reactant.
- 5. Calculate moles of required reactant/product.
- 6. Convert to grams or volume, as required.

Types of Chemical Reactions

- Neutralization Reactions
 - The net ionic equation for each acid-base neutralization reaction involves a transfer of a proton.
 - Consider the reaction of the strong acid, HCl(*aq*) and a strong base, LiOH(*aq*).

$HCl(aq) + KOH(aq) \rightarrow KCl(aq) + H_2O(l)$

Types of Chemical Reactions

Acid-Base Reactions

- Neutralization Reactions
 - Writing the strong electrolytes in the form of ions gives the complete ionic equation.

$H^+(aq) + Cl^-(aq) + K^+(aq) + OH^-(aq) \rightarrow$

 $K^+(aq) + Cl^-(aq) + H_2O(l)$

Types of Chemical Reactions Acid-Base Reactions • Neutralization Reactions • Canceling the spectator ions results in the net ionic equation. Note the proton transfer. $H^+(aq) + A^-(aq) + A^+(aq) + OH^-(aq) \rightarrow A^+(aq) + A^-(aq) + H_2O(1)$ $H^+(aq) + OH^-(aq) \rightarrow H_2O(1)$ $H^+(aq) + OH^-(aq) \rightarrow H_2O(1)$

Types of Chemical Reactions

Acid-Base Reactions

- Neutralization Reactions
 - In a reaction involving HCN(*aq*), a weak acid, and KOH(*aq*), a strong base, the product is KCN, a strong electrolyte.
 - The net ionic equation for this reaction is



Types of Chemical Reactions Acid-Base Reactions

- Acid-Base Reactions with Gas Formation
 - Sulfides react with acids to form H₂S, hydrogen sulfide gas.

$Na_2S + 2HCl \rightarrow 2NaCl + H_2S \uparrow$

- These reactions are summarized in Table 4.4.

Acid-Base Reactions

- Acid-Base Reactions with Gas Formation
 - Carbonates react with acids to form CO₂, carbon dioxide gas.

$Na_2CO_3 + 2HCl \rightarrow 2NaCl + H_2O + CO_2$

Sulfites react with acids to form SO₂, sulfur dioxide gas.

 $Na_2SO_3 + 2HCl \rightarrow 2NaCl + H_2O + SO_2$

Acid/Base Reactions

Gas	Example
CO_2	$Na_2CO_3 + 2HCl \longrightarrow 2NaCl + H_2O + CO_3$
SO_2	$Na_2SO_3 + 2HCl \longrightarrow 2NaCl + H_2O + SO_2$
H ₂ S	$Na_2S + 2HCl \longrightarrow 2NaCl + H_2S$
-	
	Gas CO ₂ SO ₂ H ₂ S

Types of Chemical Reactions

- · Oxidation-Reduction Reactions
 - Oxidation-reduction reactions involve the transfer of electrons from one species to another.
 - Oxidation is defined as the loss of electrons.
 - *Reduction* is defined as the gain of electrons.
 - Oxidation and reduction always occur simultaneously.

Figure 4.19: The reaction of solid sodium and gaseous chlorine to form solid sodium chloride.









- Oxidation-Reduction Reactions
 - The reaction of an iron nail with a solution of copper(II) sulfate, CuSO₄, is an oxidationreduction reaction (see Figure 4.10).
 - The molecular equation for this reaction is:

$Fe(s) + CuSO_4(aq) \rightarrow FeSO_4(aq) + Cu(s)$



Types of Chemical Reactions

Oxidation-Reduction Reactions • Oxidation Numbers

- The concept of oxidation numbers is a simple way of keeping track of electrons in a reaction.
- The *oxidation number* (or oxidation state) of an atom in a substance is the actual *charge* of the atom if it exists as a monatomic ion.
- Alternatively, it is *hypothetical charge* assigned to the atom in the substance by simple rules.

Types of Chemical Reactions

Oxidation-Reduction Reactions

 Oxidation Number Rules 				
Rule	Applies to	Statement		
1	Elements	The oxidation number of a		

1	Elements	The oxidation number of an atom in an element is zero. This includes molecular elements.
2	Monatomic ions	The oxidation number of an atom in a monatomic ion equals the charge of the ion.
3	Oxygen	The oxidation number of oxygen is -2 in most of its compounds. (An exception is O in H ₂ O ₂ and other peroxides, where the oxidation number is -1 .)

Types of Chemical Reactions

Oxidation-Reduction Reactions

Oxidation Number Rules		
Rule	Applies to	Statement
4	Hydrogen	The oxidation number of hydrogen is +1 in its covalent compounds.
5	Halogens	Fluorine is -1 in all its compounds. The other halogens are -1 unless the other element is another halogen or oxygen.
6	Compounds and ions	The sum of the oxidation numbers of the atoms in a compound is zero. The sum in a polyatomic ion equals the charge on the ion.

Oxidation-Reduction Reactions

- Describing Oxidation-Reduction Reactions
 - Look again at the reaction of iron with copper(II) sulfate.

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

We can write this reaction in terms of two *half-reactions*.

Types of Chemical Reactions

Oxidation-Reduction Reactions

- Describing Oxidation-Reduction Reactions
 - A *half-reaction* is one of the two parts of an oxidation-reduction reaction. One involves the loss of electrons (oxidation) and the other involves the gain of electrons (reduction).

 $Fe(s) \rightarrow Fe^{2+}(aq) + 2e^{-}$ oxidation half-reaction $Cu^{2+}(aq) + 2e^{-} \rightarrow Cu(s)$ reduction half-reaction

Types of Chemical Reactions

Oxidation-Reduction Reactions

- Describing Oxidation-Reduction Reactions
 - An oxidizing agent is a species that oxidizes another species; it is itself reduced.
 - A *reducing agent* is a species that reduces another species; *it is itself oxidized*.

Loss of 2 e- oxidation

reducing agent

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

Gain of 2 e⁻ reduction

Types of Chemical Reactions

Oxidation-Reduction Reactions

- Balancing Simple Oxidation-Reduction Reactions
 - At first glance, the equation representing the reaction of zinc metal with silver(I) ions might appear to be balanced.

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

- However, a balanced equation must have a *charge balance* as well as a *mass balance*.

Types of Chemical Reactions

Oxidation-Reduction Reactions

- Balancing Simple Oxidation-Reduction Reactions
 - Since the number of electrons lost in the oxidation half-reaction must equal the number gained in the reduction half-reaction,

 $Zn(s) \rightarrow Zn^{2+}(aq) + 2e^{-}$ oxidation half-reaction

 $2 \operatorname{Ag}^+(aq) + 2 \operatorname{e}^- \rightarrow 2 \operatorname{Ag}(s)$ reduction half-reaction we must double the reaction involving the reduction of the silver.





Rules for Assigning Oxidation States

- 1. Oxidation state of an atom in an element = 0
- 2. Oxidation state of monatomic element = charge
- 3. Oxygen = -2 in covalent compounds (except in peroxides where it = -1)
- 4. H = +1 in covalent compounds
- 5. Fluorine = -1 in compounds
- 6. Sum of oxidation states = 0 in compounds Sum of oxidation states = charge of the ion

Balancing by Half-Reaction Method

- 1. Write separate reduction, oxidation reactions.
- 2. For each half-reaction:
 - Balance elements (except H, O)
 - Balance O using H₂O
 - Balance H using H⁺
 - Balance charge using electrons

Half-Reaction Method -Balancing in Base

- 1. Balance as in acid.
- Add OH⁻ that equals H⁺ ions (both sides!)
- 3. Form water by combining H⁺, OH⁻.
- 4. Check elements and charges for balance.

Balancing by Half-Reaction Method (continued)

- 3. If necessary, multiply by integer to equalize electron count.
- 4. Add half-reactions.
- 5. Check that elements and charges are balanced.

Stoichiometry Steps for reactions in solution.

