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Unit 1: Atomic Structure & Properties

- The Atom: Consists of protons (+), neutrons (0), electrons (-).
- Mole Concept: Relates the mass of an element to the number of particles. One mole equals Avogadro's number (6.022 x 10²³) particles.
- Molar Mass: The mass of one mole of a substance in grams.
- Isotopes: Atoms of the same element with different numbers of neutrons, leading to different mass numbers.
- Mass Spectroscopy: Identifies the composition of a sample by measuring the mass-tocharge ratio of ions.
- Empirical Formula: The simplest whole number ratio of elements in a compound.
- Electron Configuration: Describes the arrangement of electrons in an atom's orbitals. Electrons fill orbitals from lowest to highest energy.
- Photoelectron Spectroscopy (PES): Measures the ionization energies of electrons to deduce electronic structure.
- Periodic Trends: Atomic radius, ionization energy, and electronegativity trends across periods and down groups.
- Quantum Mechanical Model: Describes electron distribution in atoms. Orbitals (s, p, d, f) with specific shapes and energy levels.
- Heisenberg Uncertainty Principle: Impossible to know both the position and momentum of an electron simultaneously.
- Example Calculation: Calculate the average atomic mass of an element given the isotopic masses and their abundances.



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Unit 2: Molecular & Ionic Bonding

- Intramolecular Forces: Forces within a molecule (ionic, covalent, and metallic bonds).
 - Ionic Bond: Transfer of electrons from a metal to a nonmetal.
 - Polar Covalent Bond: Unequal sharing of electrons between atoms.
 - Nonpolar Covalent Bond: Equal sharing of electrons.
- Ionic Solids: Lattices of cations and anions held together by electrostatic forces.
- Metallic Bonds: Delocalized electrons shared among a lattice of metal atoms.
- Lewis Diagrams: Represent valence electrons and bonds.
- **Resonance Structures:** Different valid Lewis structures for the same molecule.
- VSEPR Theory: Predicts molecular geometry based on electron pair repulsion.
- Bond Energy: Energy required to break a bond in a molecule.
- **Example:** Calculate the formal charge of atoms in a molecule to predict the most stable Lewis structure.
- Hybridization: Mixing of atomic orbitals to form new hybrid orbitals (sp, sp², sp³).



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Unit 3: Intermolecular Forces & Properties

- Intermolecular Forces: Forces between molecules.
 - London Dispersion Forces (LDFs): Weakest, present in all molecules.
 - **Dipole-Dipole Interactions:** Between polar molecules.
 - Hydrogen Bonding: Strongest, occurs when H is bonded to F, O, or N.
 - Ion-Dipole Interactions: Between ions and polar molecules.
- Solids:
 - Amorphous Solids: No long-range order (e.g., glass).
 - Crystalline Solids: Ordered structures (ionic, metallic, covalent network, molecular).
- Liquids:
 - **Properties:** Surface tension, viscosity, capillary action.
 - Laws: Ideal gas law (PV=nRT), law of partial pressures.
- Solutions: Factors affecting solubility (like dissolves like, temperature, pressure).
- **Phase Diagrams:** Graphs showing conditions (temperature and pressure) at which distinct phases occur and coexist.
- **Example:** Explain how intermolecular forces affect boiling and melting points of substances.



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Unit 4: Chemical Reactions

- **Physical vs. Chemical Changes:** Physical changes do not alter chemical composition, chemical changes do.
- Precipitation Reactions: Formation of an insoluble product from soluble reactants.
- Net Ionic Equations: Show only the species that undergo a change.
- Balancing Equations: Conservation of mass and charge.
- Stoichiometry: Calculations based on balanced chemical equations.
- Acid-Base Reactions: Transfer of protons between reactants.
 - Titration: Determines the concentration of an unknown acid/base.
- **Redox Reactions:** Transfer of electrons between reactants.
- **Types of Reactions:** Synthesis, decomposition, single replacement, double replacement, and combustion reactions.
- **Example Problem:** Balance a combustion reaction and calculate the amount of product formed.



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<u>Unit 5: Kinetics</u>

- Rate of Reaction: Change in concentration of reactants/products over time.
- Rate Laws: Express reaction rate as a function of reactant concentrations.

• Rate Law Formula: Rate = $k[A]^{m}[B]^{n}$

- Reaction Mechanisms: Step-by-step sequence of elementary reactions.
- **Collision Model:** Reactions occur when particles collide with sufficient energy and correct orientation.
- Catalysts: Lower the activation energy and speed up reactions without being consumed.
- Factors Affecting Reaction Rate: Concentration, temperature, surface area, and catalysts.
- Arrhenius Equation: $k = Ae^{-Ea}/RT$, relates temperature and rate constant.
- **Example Calculation:** Determine the activation energy given the rate constants at different temperatures.



<u>Unit 6: Thermodynamics</u>

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- **Kinetic vs. Potential Energy:** Kinetic energy relates to motion, potential energy to position or composition.
- Enthalpy (ΔH): Heat absorbed or released in a reaction.
- Endothermic: Absorbs heat (ΔH > 0).
- Exothermic: Releases heat (ΔH < 0).
- Calorimetry: Measures heat changes in reactions.
- Hess's Law: Total enthalpy change is the sum of the enthalpy changes for individual
- steps.
- Standard Enthalpy of Formation (ΔH°f): Enthalpy change when one mole of a compound forms from its elements.
- Second Law of Thermodynamics: Entropy of an isolated system always increases.
- **Example:** Use Hess's Law to calculate the enthalpy change of a reaction from known enthalpy changes of related reactions.



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<u>Unit 7: Equilibrium</u>

- Dynamic Equilibrium: Rate of the forward reaction equals the rate of the reverse reaction.
- Equilibrium Constant (K): Ratio of product concentrations to reactant concentrations at equilibrium.
- Le Chatelier's Principle: A system at equilibrium will adjust to relieve any applied stress.
- ICE Tables: Used to calculate changes in concentrations of reactants/products.
- Le Chatelier's Principle Examples: Predict the shift in equilibrium when concentration, temperature, or pressure changes.
- Equilibrium Expressions: Write Kc and Kp expressions for given reactions.



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<u>Unit 8: Acids & Bases</u>

- Acid-Base Definitions: Bronsted-Lowry (proton transfer), Lewis (electron pair).
- **pH and pOH:** Measures of acidity/basicity.
 - \circ pH = -log[H₃O⁺]
 - \circ pOH = -log[OH-]
- **Titrations:** Determine the concentration of an acid/base by neutralization.
- **Buffers:** Solutions that resist changes in pH when small amounts of acid or base are added.
- Strength of Acids and Bases: Strong acids/bases dissociate completely, weak acids/bases do not.
- **pKa and pKb:** Measures of acid and base strength.
- **Example Calculation:** Calculate the pH of a weak acid solution using the acid dissociation constant (Ka).



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Unit 9: Applications of Thermodynamics

- Entropy (S): Measure of disorder.
- **Gibbs Free Energy (ΔG):** Determines spontaneity of a reaction.
 - $\circ \Delta G = \Delta H T \Delta S$
 - \circ Spontaneous when ΔG < 0.
- Electrochemistry: Study of redox reactions and their applications.
 - Galvanic Cells: Convert chemical energy to electrical energy.
 - Electrolytic Cells: Use electrical energy to drive non-spontaneous reactions.
- **Standard Electrode Potentials:** Determine cell potential from standard reduction potentials.
- Nernst Equation: $E = E^{\circ} \frac{RT}{nF} \ln Q$, relates cell potential to reaction quotient.
- **Example:** Calculate the Gibbs free energy change for a redox reaction from the cell potential.

