

Curriculum Plans: Year 12 (Chemistry)

	Unit	Knowledge: By the end of the unit students will know:	Skills: What skills will students have developed by the end of this unit?	Key terms: What new key terms and vocabulary will be learnt in this unit?	Summative Assessment: How will pupils be assessed in this unit?
Michaelmas 1	1.1 Atomic Structure 1.5 Kinetics 3.1 Introduction to Organic Chemistry 1.3 Bonding and Structure	<ul style="list-style-type: none"> - The structure of an atom, including protons, neutrons, and electrons, along with their relative charges and masses. - The definitions of mass number (A) and atomic number (Z), and how these relate to the number of fundamental particles. - The existence and concept of isotopes. - The principles of mass spectrometry, including ionization, acceleration, ion drift, detection, and data analysis. - How to interpret mass spectra to determine isotopic abundances and calculate relative atomic mass. - The principles of collision theory and how it affects reaction rates. - The concept of activation energy and its significance in determining whether reactions occur. - How the Maxwell-Boltzmann distribution explains the distribution of molecular energies in gases. 	<ul style="list-style-type: none"> - Ability to calculate the number of protons, neutrons, and electrons in atoms and ions from atomic and mass numbers. - Understanding and explaining the concept of isotopes. - Interpretation of mass spectra to deduce the relative isotopic mass and abundance. - Use of significant figures and weighted means in calculations. - Writing electron configurations for atoms and ions up to $Z=36$. - Defining and calculating ionization energies. - Ability to explain the relationship between molecular collisions and reaction rates. - Interpreting Maxwell-Boltzmann distribution curves and explaining temperature effects on reaction rates. - Investigating and explaining the effect of temperature, concentration, 	<ul style="list-style-type: none"> - Isotope - Mass number (A) - Atomic number (Z) - Ionization energy - Electron configuration - Time of flight (TOF) mass spectrometer - Relative atomic mass - Isotopic abundance - Collision theory - Activation energy - Maxwell-Boltzmann distribution - Rate of reaction - Catalyst - Reaction rate - Concentration - Pressure 	Each unit throughout the year is assessed using both peer assessment of Homework and Teacher assessment of end of unit tests

Curriculum Plans: Year 12 (Chemistry)

		<ul style="list-style-type: none"> - The effect of temperature on the rate of reaction and how a small temperature change can lead to a significant increase in reaction rate. - The impact of concentration and pressure on collision frequency and reaction rates. - How catalysts work by providing an alternative reaction route with a lower activation energy. - The basics of organic chemistry, including the diversity of organic compounds. - The importance of carbon in organic chemistry and how organic compounds range from fuels to biological molecules. - How to use the IUPAC system for naming organic compounds. - The different ways of representing the structure of organic compounds (empirical, molecular, general, structural, displayed, and skeletal formulas). - The concept of homologous series and functional groups. - How different types of chemical bonds (ionic, covalent, and metallic) are formed and the principles behind each type of bonding. - The concept of electrostatic 	<p>and pressure changes on reaction rates.</p> <ul style="list-style-type: none"> - Practical skills in monitoring reaction rates using different methods (e.g., continuous monitoring, initial rate method). - Drawing and interpreting Maxwell-Boltzmann distribution curves. - Drawing and interpreting various formula representations (empirical, molecular, structural, displayed, skeletal). - Applying IUPAC rules to name organic compounds with up to six carbon atoms. - Drawing the structure of organic compounds based on IUPAC names. - Explaining reaction mechanisms, including free-radical mechanisms and the use of curly arrows in non-radical mechanisms. - Ability to draw Lewis structures and predict the shapes of molecules using electron pair repulsion theory. - Identifying bond types in different compounds based on their formulas. 	<ul style="list-style-type: none"> - IUPAC nomenclature - Homologous series - Empirical formula - Molecular formula - Structural formula - Displayed formula - Skeletal formula - Free-radical mechanism - Curly arrow - Structural isomerism - Stereoisomerism - E-Z isomerism - Electronegativity - Bond polarity - Intermolecular forces - Dipole-dipole forces - Van der Waals forces 	
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Curriculum Plans: Year 12 (Chemistry)

		<p>attraction between ions in ionic bonding and between nuclei and electrons in covalent bonding.</p> <ul style="list-style-type: none"> - The nature of metallic bonding and the concept of delocalized electrons. - The relationship between bonding and the physical properties of materials (e.g., melting point, conductivity). - The different types of crystal structures (ionic, metallic, macromolecular, molecular) and examples of each. - The concept of electronegativity and how it affects bond polarity. 	<ul style="list-style-type: none"> - Predicting and explaining the physical properties of substances from their bonding and structure. - Drawing and interpreting diagrams to represent various crystal structures. - Calculating bond polarity and explaining the effects of electronegativity on molecular polarity. 	<p>(London dispersion forces)</p> <ul style="list-style-type: none"> - Hydrogen bonding - Macromolecular structure - Crystal structure 	
Michaelmas 2	<p>1.3 Bonding and Structure (Continued)</p> <p>1.2 Amount of Substance</p> <p>3.2 to 3.3 Alkanes and Halogenoalkanes</p> <p>3.4 to 3.5 Alkenes and Alcohols</p>	<ul style="list-style-type: none"> - The concept of the mole as a unit for amount of substance and how it applies to atoms, molecules, and ions. - The use of the Avogadro constant in relation to the number of particles in a mole. - How to calculate relative atomic mass (Ar) and relative molecular mass (Mr). - The ideal gas equation, $pV=nRT$, and its application in calculating properties of gases under varying conditions. - The relationship between empirical and molecular formulas and how to calculate each from 	<ul style="list-style-type: none"> - Ability to perform calculations using the mole concept, including converting between mass, number of moles, and number of particles using the Avogadro constant. - Using the ideal gas equation to calculate unknown quantities (e.g., pressure, volume, or temperature of gases). - Performing percentage yield and atom economy calculations. - Writing and balancing chemical equations and using them to carry out stoichiometric calculations. - Determining empirical and molecular formulas from experimental data. 	<ul style="list-style-type: none"> - Mole - Avogadro constant - Relative atomic mass (Ar) - Relative molecular mass (Mr) - Ideal gas equation $pV=nRT$ - Empirical formula - Molecular formula - Stoichiometry - Percentage yield - Atom economy 	
Lent 1	<p>1.2 Amount of Substance (Continued)</p> <p>3.4 to 3.5 Alkenes and</p>				

Curriculum Plans: Year 12 (Chemistry)

	Alcohols (Continued)	<p>experimental data.</p> <ul style="list-style-type: none"> - The importance of balanced equations in determining the amounts of reactants and products. - The principles behind the combustion and chlorination of alkanes. - The reactivity of halogenoalkanes and their substitution reactions with nucleophiles like OH⁻, CN⁻, and NH₃. - The mechanism of ozone depletion and the role of chlorofluorocarbons (CFCs) in this process. - The structure and reactivity of alkenes, including electrophilic addition reactions and polymerization. - The production and oxidation of alcohols, and their uses in industry, particularly ethanol as a biofuel. 	<ul style="list-style-type: none"> - Writing and interpreting mechanisms for free-radical substitution, nucleophilic substitution, and electrophilic addition reactions. - Predicting the products of combustion and reactions of alkanes, halogenoalkanes, and alkenes. - Testing organic compounds for unsaturation using bromine water. - Practical skills in synthesizing alcohols and separating products using techniques like distillation and recrystallization. 	<ul style="list-style-type: none"> - Alkanes - Halogenoalkanes - Electrophilic addition - Polymerization - Nucleophilic substitution - Ozone depletion - Free radical - Biofuel 	
Lent 2	1.6 Equilibria and Redox 3.6 Organic Analysis	<ul style="list-style-type: none"> - The concept of dynamic equilibrium and how, in a reversible reaction, the rates of the forward and reverse reactions are equal at equilibrium. - Le Chatelier's principle and how 	<ul style="list-style-type: none"> - Applying Le Chatelier's principle to predict the effects of changing temperature, pressure, and concentration on the position of equilibrium. - Constructing and interpreting the 	<ul style="list-style-type: none"> - Dynamic equilibrium - Le Chatelier's principle - Equilibrium constant K_c - Homogeneous 	

Curriculum Plans: Year 12 (Chemistry)

		<p>changes in temperature, pressure, and concentration affect the position of equilibrium.</p> <ul style="list-style-type: none"> - How a catalyst affects the rate of reaction but not the position of equilibrium. - The definition and mathematical expression of the equilibrium constant K_c for homogeneous systems. <p>- The definitions of oxidation and reduction in terms of electron transfer.</p> <ul style="list-style-type: none"> - The role of oxidising agents as electron acceptors and reducing agents as electron donors. - How to assign oxidation states to elements in compounds and ions. - How to write and balance redox half-equations for oxidation and reduction processes. - How to combine half-equations to form an overall redox equation. <p>- The use of test-tube reactions to identify organic functional groups such as alcohols, aldehydes,</p>	<p>expression for the equilibrium constant K_c from balanced equations.</p> <ul style="list-style-type: none"> - Performing calculations involving K_c values from equilibrium concentrations. - Understanding how changes in temperature influence the value of K_c - Using spectroscopic techniques to identify organic compounds and analyse their structures. <p>- Ability to determine oxidation states for elements in compounds and ions.</p> <ul style="list-style-type: none"> - Writing redox half-equations for oxidation and reduction reactions. - Combining half-equations to form balanced overall redox equations. - Applying redox concepts to various chemical reactions. <p>- Conducting test-tube reactions to identify organic compounds.</p> <ul style="list-style-type: none"> - Using mass spectrometry to deduce the molecular formula of organic 	<p>equilibrium</p> <ul style="list-style-type: none"> - Reversible reaction <ul style="list-style-type: none"> - Oxidation - Reduction - Oxidising agent - Reducing agent - Redox reaction - Oxidation state - Half-equation <ul style="list-style-type: none"> - Mass spectrometry - Molecular ion peak 	
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Curriculum Plans: Year 12 (Chemistry)

		<p>alkenes, and carboxylic acids.</p> <ul style="list-style-type: none"> - The principles behind mass spectrometry and how it can be used to determine the molecular formula of a compound. - How infrared (IR) spectroscopy is used to identify functional groups based on absorption of infrared radiation by chemical bonds. 	<p>compounds.</p> <ul style="list-style-type: none"> - Interpreting infrared spectra to identify functional groups and determine molecular structures. - Applying organic analysis techniques to determine the presence of impurities in compounds. 	<ul style="list-style-type: none"> - Infrared (IR) spectroscopy - Functional group - Fingerprint region - Wavenumber - Electromagnetic spectrum 	
Trinity 1	<p>2.1 periodicity 2.2 and 2.3 Group 2 and Group 7 1.4 Energetics</p>	<ul style="list-style-type: none"> - The structure and classification of the Periodic Table, including the classification of elements as s, p, d, or f block based on their proton number. - Trends in atomic radius, first ionization energy, and melting points across Period 3 (from sodium to argon) and the underlying reasons for these trends in terms of atomic structure and bonding. - How these properties reflect the electronic configurations of the elements in Period 3. - The properties and trends of Group 2 elements, including atomic radius, first ionization energy, and melting points of the elements from magnesium (Mg) to barium (Ba). - The reactions of Group 2 elements with water, including the general 	<ul style="list-style-type: none"> - Ability to explain and predict trends in atomic radius, ionization energy, and melting points across a period. - Explaining the melting points of the elements based on their bonding type (metallic, covalent, or ionic). - Identifying the relationships between an element's position in the Periodic Table and its physical and chemical properties. - Explaining and predicting trends in reactivity, atomic size, ionization energy, and melting points for Group 2 elements. - Performing laboratory experiments to test the reactions of Group 2 metals with water and their 	<ul style="list-style-type: none"> - Atomic radius - Ionization energy - Melting point - S, p, d, f block elements - Trends in periodicity - Alkaline earth metals - Ionization energy - Reactivity - Hydroxides - Sulfates - Solubility 	

Curriculum Plans: Year 12 (Chemistry)

		<ul style="list-style-type: none"> - The significance of standard enthalpy changes, including the standard enthalpy of combustion ($\Delta_c H^\ominus$) and standard enthalpy of formation ($\Delta_f H^\ominus$). - The calorimetry equation $q = mc\Delta T$ for calculating heat changes in reactions, where m is the mass, c is the specific heat capacity, and ΔT is the temperature change. - The applications of Hess's law to calculate enthalpy changes for reactions from known enthalpies of combustion or formation. - The concept of mean bond enthalpy and how it can be used to estimate reaction enthalpies. 	<p>unknown enthalpy changes.</p> <ul style="list-style-type: none"> - Estimating enthalpy changes from bond enthalpies and understanding the limitations of this method. - Reporting results with appropriate significant figures and units. 	<ul style="list-style-type: none"> - Standard enthalpy of combustion ($\Delta_c H^\ominus$) - Standard enthalpy of formation ($\Delta_f H^\ominus$) - Calorimetry - Hess's law - Mean bond enthalpy 	
Trinity 2	1.9 Rate Equations 1.10 Equilibria in Gases	<ul style="list-style-type: none"> - The mathematical relationship between the rate of a chemical reaction and the concentration of reactants, expressed through the rate equation - The significance of the orders of reaction in relation to the reaction mechanism and how they are determined experimentally. - The Arrhenius equation - How changes in temperature affect the rate constant and, consequently, the rate of reaction. 	<ul style="list-style-type: none"> - Defining and calculating the order of reaction from concentration-time data. - Using concentration-time graphs to deduce rates and determine orders of reaction (0, 1, or 2) with respect to reactants. - Performing calculations using the rate equation and the Arrhenius equation to find unknown values 	<ul style="list-style-type: none"> - Rate equation - Rate constant - Order of reaction - Activation energy - Arrhenius equation - Initial rate - Concentration-time graph 	

Curriculum Plans: Year 12 (Chemistry)

		<ul style="list-style-type: none">- The definition of the equilibrium constant K and its significance in reversible gas phase reactions.- How to derive partial pressures from mole fractions and total pressure.- The mathematical expression for K_p for a homogeneous system in equilibrium, relating to the partial pressures of reactants and products.- How changes in temperature and pressure affect the position of equilibrium and the value of K_p- The concept that a catalyst affects the rate of reaching equilibrium but does not change the value of the equilibrium constant	<ul style="list-style-type: none">- Constructing and interpreting the expression for K_p based on balanced equations for gas phase reactions.- Performing calculations involving K_p and deriving partial pressures.- Predicting qualitative effects of changes in temperature and pressure on the position of equilibrium and K_p.- Understanding the application of K_p in industrial processes to optimize conditions for maximum yield.	<ul style="list-style-type: none">- Equilibrium constant K_p- Partial pressure- Homogeneous equilibrium- Reversible reaction- Catalyst	
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