

5.1 Rates, equilibrium and pH

5.1.1 How Fast?

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5.1.1 How Fast?

- Explain and use the terms: rate of reaction, order, overall order, rate constant, half-life, rate-determining step
- Deduction of orders from experimental data
- Deduction of a rate equation from orders of the form: $\text{rate} = k[\text{A}]^m[\text{B}]^n$
- Calculate rate constant k , and related quantities, from a rate equation including determining units
- Deduce the order with respect to a reactant from the shape of a concentration-time graph
- Calculate the reaction rate from the gradient of a concentration-time graph
- Find the constant half-life from a concentration-time graph of a first order reaction
- Determine the rate constant, k , for a first order reaction from the constant half life
- Deduce the order with respect to a reactant from a rate-concentration graph
- Determine the rate constant for a first order reaction from the gradient of a rate-concentration graph
- Describe techniques and procedures used to investigate reaction rates by the initial rates method, including calorimetry
- Predict a rate equation consistent with the rate-determining step of a multi-step reaction
- Predict possible steps in a reaction mechanism for the rate equation and the balanced equation for a multistep
- Qualitative explanation of the affect of temperature change on the rate of a reaction and the rate constant
- Use the Arrhenius equation to find the exponential relationship between the rate constant and temperature
- Use the Arrhenius equation to determine E_a and A graphically

5.1.2 How Far?

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5.1.2 How Far?

- Use the terms mole fraction and partial pressure
- Calculate quantities at equilibrium
- Describe techniques and procedures used to determine quantities present at equilibrium
- Use expressions for K_c and K_p for homogeneous and heterogeneous equilibria
- Calculate K_c and K_p including the determination of units
- Describe the qualitative effect on equilibrium constants of changing temperature for exothermic and endothermic reactions
- Describe the constancy of equilibrium constants with changes in concentration, pressure or the presence of a catalyst
- Explanation of how an equilibrium constant controls the position of equilibrium on changing concentration, pressure and temperature
- Application of above principles for K_c , K_p to other equilibrium constants

5.1.3 Acids, bases and buffers

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5.1.3 Acids, bases and buffers

- Define an Brønsted- lowry acid and base in terms of protons
- Use the term conjugate acid-base pairs
- Use the terms monobasic, dibasic and tribasic
- Describe the role of H^+ in the reaction of acids with metals and bases using ionic equations
- Use the acid dissociation constant K_a for the extent of acid dissociation
- Describe the relationship between K_a and $\text{p}K_a$
- Use the expressions for pH as: $\text{pH} = -\log[\text{H}^+]$; $[\text{H}^+] = 10^{-\text{pH}}$
- Use of the expression for the ionic product of water K_w
- Calculations of pH for: strong monobasic acids and strong bases (using K_w)
- Calculations of pH , K_a or related quantities for a weak monobasic acid using approximations
- Describe the limitations of using approximations to K_a related calculations for 'stronger' weak acids
- Define the term buffer
- Describe the formation of a buffer from a weak acid and salt of the weak acid
- Describe the formation of a buffer from excess of weak acid and a strong alkali
- Explanation of the role of a conjugate acid-base pair in an acid buffer solution
- Calculation of the pH of a buffer solution from the K_a value of a weak acid and the equilibrium concentrations
- Explanation of the control of blood pH by the carbonic acid- hydrogencarbonate buffer system
- Use of pH titration curves for combinations of strong and weak acid
- Describe the techniques and procedures used when measuring pH with a pH meter

5.2 Energy

| 5.2.1 Lattice energy | | R | A | G |
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| 5.2.1 Lattice energy | Explanation of the term lattice energy and its use as a measure of the strength of ionic bonding in a giant ionic lattice | | | |
| | Use of lattice enthalpy of a simple ionic solid and relevant energy terms for the construction of Born-Haber cycles | | | |
| | Use of lattice enthalpy of a simple ionic solid and relevant energy terms for related calculations | | | |
| | Explanation and use of the term enthalpy change of solution | | | |
| | Explanation and use of the term enthalpy change of hydration | | | |
| | Use of the enthalpy change of solution and energy terms for the construction of enthalpy cycles | | | |
| | Use of the enthalpy change of solution and energy terms for related calculations | | | |
| | Qualitative explanation of the effect of ionic charge and ionic radius on lattice enthalpy and enthalpy change of hydration | | | |
| 5.2.2 Enthalpy and entropy | | R | A | G |
| 5.2.2 Enthalpy and entropy | Explanation of term entropy | | | |
| | Explanation of the difference in magnitude of the entropy system of solids, liquids and gases | | | |
| | Explanation of the difference in magnitude of the entropy system for a reaction with a change in number of gaseous molecules | | | |
| | Calculation of the entropy change of a system ΔS and related quantities | | | |
| | Explanation on the feasibility of a process based on $T\Delta S$ and ΔH | | | |
| | Explanation, and related calculations for free energy ΔG | | | |
| | Explanation of process feasibility in terms of ΔG | | | |
| | Limitations of predictions made by ΔG about feasibility in terms of kinetics | | | |
| 5.2.3 Redox and electrode potentials | | R | A | G |
| 5.2.3 Redox and electrode potentials | Explanation and use of the terms oxidising agent and reducing agent | | | |
| | Construction of redox equations using half-equations and oxidation numbers | | | |
| | Interpretation and prediction of reactions involving electron transfer | | | |
| | Describe techniques and procedures used when carrying out redox titrations | | | |
| | Use of structured and non-structured titration calculations, based on experimental results of redox titrations | | | |
| | Use of the term standard electrode (redox) potential including its measurement | | | |
| | Describe techniques and procedures used for the measurement of cell potentials | | | |
| | Calculation of standard cell potential by combining two standard electrode potentials | | | |
| | Prediction of feasibility of a reaction and limitations using standard electrode potentials | | | |
| | Application of principles of electrode potentials to modern storage cells | | | |
| | Explanation of a fuel cell in terms of oxygen | | | |

5.3 Transition elements

| 5.3.1 Transition elements | | R | A | G |
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| 5.3.1 Transition elements | Electron configuration of atoms and ions of the d-block elements of period 4, given atomic number and charge | | | |
| | Know the elements Ti- Cu as transition elements | | | |
| | Illustration using at least two transition elements of: the existence of more than 1 oxidation state | | | |
| | Illustration using at least two transition elements of: the formation of coloured ions | | | |
| | Illustration using at least two transition elements of: the catalytic behaviour of elements and compounds | | | |
| | Explanation of the term ligand in terms of coordinate bonding, including bidentate ligands | | | |
| | Use of terms complex ion and coordination number | | | |
| | Examples of complexes with six-fold coordination with an octohedral shape | | | |
| | Examples of complexes with four-fold coordination with either a planar or tetrahedral shape | | | |
| | Types of stereoisomerism shown by complexes including those associated with bidentate and multidentate ligands | | | |
| | Use of cis-platin as an anticancer drug and its action | | | |
| | Ligand substitution reactions and the accompanying colour changes | | | |
| | Explanation of the biochemical importance of iron in haemoglobin, including ligand substitution | | | |
| | Precipitation reactions, including ionic equations and colour changes for transition element ions with sodium hydroxide and ammonia | | | |
| | Complex formation with sodium hydroxide(aq) and ammonia(aq) | | | |
| | Redox reactions and colour changes for interconversions between Fe^{2+} and Fe^{3+} | | | |
| | Redox reactions and colour changes for interconversions between Cr^{3+} and $\text{Cr}_2\text{O}_7^{2-}$ | | | |
| | Redox reactions and colour changes for reduction of Cu^{2+} to Cu^+ and disproportionation | | | |
| | Interpretation and prediction of unfamiliar reactions including ligand substitution, precipitation and redox | | | |
| 5.3.2 Qualitative analysis | | R | A | G |
| 5.3.2 Qualitative analysis | Qualitative analysis of ions on a test tube scale | | | |

6.1 Aromatic compounds, carbonyls and acids

| 6.1.1 Aromatic compounds | | R | A | G |
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| 6.1.1 Aromatic compounds | Comparison of the Kekulé model of benzene and the subsequent delocalised models | | | |
| | Experimental evidence for delocalised, rather than Kekulé model | | | |
| | Use of IUPAC rules of nomenclature for systematically naming substituted aromatic compounds | | | |
| | Understanding of electrophilic substitution of aromatic compounds with concentrated nitric acid | | | |
| | Understanding of electrophilic substitution of aromatic compounds with a halogen | | | |
| | Understanding of electrophilic substitution of aromatic compounds with a haloalkane or acyl chloride | | | |
| | Understanding of mechanism of electrophilic substitution in arenes for nitration and halogenation | | | |
| | Explanation of the relative resistance to bromination of benzene compared with alkenes | | | |
| | Interpretation of unfamiliar electrophilic substitution reactions of aromatic compounds | | | |
| | Understanding of the weak acidity of phenols | | | |
| | Understanding of the electrophilic substitution reactions of phenol with bromine and dilute nitric acid | | | |
| | Explanation of the relative ease of electrophilic substitution of phenol compared to benzene | | | |
| | Explanation of the 2- and 4-directing effect of electron donating groups in electrophilic substitution of aromatic compounds | | | |
| | Understanding of 3-directing effect of electron withdrawing groups in electrophilic substitution of aromatic compounds | | | |
| | Prediction of substitution products of aromatic compounds | | | |
| 6.1.2 Carbonyl compounds | | R | A | G |
| 6.1.2 Carbonyl compounds | Understand oxidation of aldehydes using $\text{Cr}_2\text{O}_7^{2-}/\text{H}^+$ to form carboxylic acids | | | |
| | Understand nucleophilic addition reactions of carbonyl compounds with NaBH_4 and HCN | | | |
| | Describe the mechanism of nucleophilic addition reactions of aldehydes and ketones with NaBH_4 and HCN | | | |
| | Describe the use of 2,4-dinitrophenylhydrazine to detect the presence of a carbonyl group | | | |
| | Describe the use of 2,4-dinitrophenylhydrazine to identify a carbonyl compound from the melting point of the derivative | | | |
| | Describe the use of Tollens' reagent to detect the presence of an aldehyde group | | | |
| | Describe the use of Tollens' reagent to distinguish between aldehydes and ketones, with explanation | | | |
| 6.1.3 Carboxylic acids and esters | | R | A | G |
| 6.1.3 Carboxylic acids and esters | Explanation of the water solubility of carboxylic acids in terms of hydrogen bonding | | | |
| | Description of the reactions of carboxylic acids with metals and bases | | | |
| | Understanding of esterification of carboxylic acids with alcohols in the presence of an acid catalyst | | | |
| | Understanding of esterification of acid anhydrides with alcohols | | | |
| | Understanding of hydrolysis of esters in hot aqueous acid to form carboxylic acids and alcohols | | | |
| | Understanding of hydrolysis of esters in hot aqueous alkali to form carboxylate salts and alcohols | | | |
| | Understanding of the formation of acyl chlorides from carboxylic acids using SOCl_2 | | | |
| | Use of acyl chlorides in synthesis in formation of esters, carboxylic acids and primary and secondary amides | | | |

6.2 Nitrogen compounds, polymers and synthesis

| 6.2.1 Amines | | R | A | G |
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| 6.2.1 Amines | Explain basicity of amines in terms of protons and reactions with dilute acids | | | |
| | Understand the preparation of aliphatic amines by substitution of haloalkanes with excess ethanolic ammonia and amines | | | |
| | Understand the preparation of aromatic amines by reduction of nitroarenes using tin and concentrated hydrochloric acid | | | |
| 6.2.2 Amino acids, amides and chirality | | | | |
| 6.2.2 Amino acids, amides and chirality | State the general formula for an α -amino acids as $\text{RCH}(\text{NH}_2)\text{COOH}$ | | | |
| | Understand the reactions of the carboxylic acid group with alkalis and in the formation of esters | | | |
| | Understand the reaction of the amine group with acids | | | |
| | Describe the structures of primary and secondary amides | | | |
| | Understanding of optical isomerism | | | |
| | Identification of chiral centres in a molecule of any organic compound | | | |
| 6.2.3 Polyesters and polyamides | | R | A | G |
| 6.2.3 Polyesters and polyamides | Understanding of condensation polymerisation to form polyesters and polyamides | | | |
| | Understanding of the acid and base hydrolysis of ester groups in polyesters and amide groups in polyamides | | | |
| | Prediction from addition and condensation reactions of: the repeat unit, monomer required, type of polymerisation | | | |
| 6.2.4 Carbon-carbon bond formation | | R | A | G |
| 6.2.4 Carbon-carbon bond formation | Describe the use of C-C bond formation in synthesis to increase the length of a carbon chain | | | |
| | Understand the formation of $\text{C}-\text{C}\equiv\text{N}$ by reaction of haloalkanes with CN^- and ethanol, including mechanism | | | |
| | Understand the formation of $\text{C}-\text{C}\equiv\text{N}$ by reaction of carbonyl compounds with HCN, including mechanism | | | |
| | Understand the reaction of nitriles by reduction and acid hydrolysis | | | |
| | Understand the formation of substituted aromatic C-C by alkylation and acylation (Friedel-Crafts reaction) | | | |
| 6.2.5 Organic synthesis | | R | A | G |
| 6.2.5 Organic synthesis | Describe techniques and procedures used for the preparation and purification of organic solids | | | |
| | Describe the synthetic routes for an organic molecule containing several functional groups | | | |
| | Describe the multistage synthetic routes for the preparation of organic compounds | | | |

6.3 Analysis

| 6.3.1 Chromatography and qualitative analysis | | R | A | G |
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| 6.3.1 Chromatography and qualitative analysis | Interpretation of one-way TLC chromatograms in terms of R_f values | | | |
| | Interpretation of gas chromatograms in terms of retention rates and the amounts and proportions of the components | | | |
| | Qualitative analysis of organic functional groups on a test tube scale: alkenes by bromine | | | |
| | Qualitative analysis of organic functional groups on a test tube scale: Halogenoalkanes by aqueous silver nitrate in ethanol | | | |
| | Qualitative analysis of organic functional groups on a test tube scale: Phenols by weak acidity but no reaction with CO_3^{2-} | | | |
| | Qualitative analysis of organic functional groups on a test tube scale: Carbonyl compounds by reaction with 2,4-DNP | | | |
| | Qualitative analysis of organic functional groups on a test tube scale: Aldehydes by reaction with Tollens' reagent | | | |
| | Qualitative analysis of organic functional groups on a test tube scale: Primary and secondary alcohols and aldehydes by reaction with acidified dichromate | | | |
| | Qualitative analysis of organic functional groups on a test tube scale: Carboxylic acids by reaction with CO_3^{2-} | | | |
| 6.3.2 Spectroscopy | | R | A | G |
| 6.3.2 Spectroscopy | Analyse carbon-13 NMR spectrum of an organic molecule to make predictions | | | |
| | Analyse high resolution proton NMR spectrum of an organic molecule to make predictions | | | |
| | Predict the carbon-13 or proton NMR spectrum for a given molecule | | | |
| | Explain the use of tetramethylsilane, TMS, as the standard for chemical shift measurements | | | |
| | Explain the need for deuterated solvents when running an NMR spectrum | | | |
| | Identify O-H and N-H protons by proton exchange using D_2O | | | |
| | Deduce the structures of organic compounds from different analytical data | | | |