

CARIBBEAN
EXAMINATIONS COUNCIL

Caribbean Advanced Proficiency Examination ${ }^{\circledR}$

## SYLLABUS

## CHEMISTRY

CXC A11/U2/17

Effective for examinations from May-June 2019

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Please check the website, www.cxc.org for updates on CXC's syllabuses.

## Introduction

he Caribbean Advanced Proficiency Examination ${ }^{\circledR}$ (CAPE ${ }^{\circledR}$ ) is designed to provide certification of the academic, vocational and technical achievement of students in the Caribbean who, having completed a minimum of five years of secondary education, wish to further their studies. The examinations address the skills and knowledge acquired by students under a flexible and articulated system where subjects are organised in 1-Unit or 2-Unit courses with each Unit containing three Modules. Subjects examined under CAPE ${ }^{\circledR}$ may be studied concurrently or singly.

The Caribbean Examinations Council offers three types of certification at the CAPE ${ }^{\circledR}$ level. The first is the award of a certificate showing each CAPE ${ }^{\circledR}$ Unit completed. The second is the CAPE ${ }^{\circledR}$ Diploma, awarded to candidates who have satisfactorily completed at least six Units, including Caribbean Studies. The third is the $\mathbf{C X C}^{\circledR}$ Associate Degree, awarded for the satisfactory completion of a prescribed cluster of eight CAPE ${ }^{\circledR}$ Units including Caribbean Studies, Communication Studies and Integrated Mathematics. Integrated Mathematics is not a requirement for the CXC ${ }^{\circledR}$ Associate Degree in Mathematics. The complete list of Associate Degrees may be found in the CXC ${ }^{\circledR}$ Associate Degree Handbook.

For the CAPE $^{\circledR}$ Diploma and the CXC $^{\circledR}$ Associate Degree, candidates must complete the cluster of required Units within a maximum period of five years. To be eligible for a CXC ${ }^{\circledR}$ Associate Degree, the educational institution presenting the candidates for the award, must select the Associate Degree of choice at the time of registration at the sitting (year) the candidates are expected to qualify for the award. Candidates will not be awarded an Associate Degree for which they were not registered.

## Chemistry Syllabus

## - RATIONALE

Science plays a major role in the evolution of knowledge. It empowers us to use creative and independent approaches to problem-solving. It arouses our natural curiosity and enables us to meet diverse, and ever expanding, challenges. It enhances our ability to inquire, seek answers, research, and interpret data. These skills use the scientific method which lead to the construction of theories and laws that help us to explain natural phenomena and exercise control over our environment. Science is, thus, an integral component of a balanced education.

Chemistry is a fundamental science that should be included as a part of our science education. Chemical principles are currently applied to societal concerns, such as, the use of pharmaceuticals, communicable diseases, environmental pollution, forensics and depletion of natural resources. As such, chemistry is a major area of scientific study which impinges on and influences every facet of our daily lives - the food we eat, the clothes we wear, our health, environment, and recreational activities.

The CAPE ${ }^{\circledR}$ Chemistry Syllabus is redesigned to allow students to work individually and with others in practical, field, and interactive activities that are related to theoretical concepts in the course. It is expected that students will apply investigative and problem-solving skills, effectively communicate scientific information, and appreciate the contribution that a study of chemistry makes to their understanding of the world. This syllabus places more emphasis on the understanding and application of chemical concepts and principles. Consequently, students will develop skills that will be of long term value in an increasingly technological and entrepreneurial world, rather than focusing on large quantities of factual information. Furthermore, it encourages the use of various student-centred teaching-learning strategies and assessment, while at the same time, catering to the multiple intelligences, and different learning styles and needs.

The most important natural resource in the Caribbean is its people. If the Caribbean is to play an important role in the new global village and survive economically, a sustained development of the scientific and technological resources of its people is essential. This syllabus contributes to the development of the Ideal Caribbean Person as articulated by the CARICOM Heads of Government in the following areas: respect for human life, awareness of the importance of living in harmony with the environment. Students will be given the opportunity to demonstrate multiple literacies, independent and critical thinking, and the innovative application of science and technology to problem-solving. In keeping with the UNESCO Pillars of Learning, on completion of this course of study, students will learn to do, learn to be and learn to transform themselves and society.

## - AIMS

The syllabus aims to:

1. acquire the knowledge and understanding of chemical principles so as to be suitably prepared for employment and for further studies at the tertiary level;
2. develop an ability to communicate scientific information in a logical and structured manner;
3. appreciate, understand and use the scientific method in the solving of problems;
4. assist in the development of critical thinking, analytical, and practical skills;
5. apply chemical knowledge to everyday life situations;
6. recognise that advances in chemistry are constantly influenced by technological, economic, social, cultural, and ethical factors;
7. appreciate that some of the advances in the field of chemistry are the results of the contributions from scientists in other disciplines;
8. further develop the spirit of inquiry in order to continue the search for new ways in which the resources of our environment can be used in a sustainable way;
9. make use of chemical data, concepts, principles, and terminology in communicating chemical information;
10. recognise the power, impact, and influence which chemistry has in a modern scientific world;
11. develop the ability to work independently and collaboratively with others when necessary;
12. appreciate the significance and limitations of science in relation to social and economic development;
13. integrate Information and Communication Technology (ICT) tools and skills into the teaching and learning of chemical concepts; and,
14. contribute to making the Caribbean scientifically literate.

## - SKILLS AND ABILITIES TO BE ASSESSED

The skills, students are expected to have developed on completion of this syllabus, have been grouped under three main headings, namely:

1. Knowledge and Comprehension;
2. Use of Knowledge; and,
3. Experimental Skills.

## 1. Knowledge and Comprehension (KC)

(a) Knowledge - the ability to identify, remember and grasp the meaning of basic facts, concepts and principles.
(b) Comprehension - the ability to select appropriate ideas, match, compare and cite examples and principles in familiar situations.

## 2. Use of Knowledge (UK)

(a) Application

The ability to:
(i) use facts, concepts, principles and procedures in familiar and in novel situations;
(ii) transform data accurately and appropriately; and,
(iii) use formulae accurately for computational purposes.
(b) Analysis and Interpretation

The ability to:
(i) identify and recognise the component parts of a whole and interpret the relationship among those parts;
(ii) identify causal factors and show how they interact with each other;
(iii) infer, predict and draw conclusions; and,
(iv) make necessary and accurate calculations and recognise the limitations and assumptions involved.
(c) Synthesis

The ability to:
(i) combine component parts to form a new and meaningful whole; and,
(ii) make predictions and solve problems.
(d) Evaluation

The ability to:

- make reasoned judgements and recommendations based on the value of ideas and information and their implications.


## 3. Experimental Skills (XS)

(a) Observation, Recording and Reporting The ability to:
(i) use the senses to perceive objects and events accurately;
(ii) record the results of a measurement accurately;
(iii) select and use appropriate formats and presentations, such as tables, graphs and diagrams;
(iv) organise and present a complete report in a clear and logical form using spelling, punctuation and grammar with an acceptable degree of accuracy; and,
(v) report accurately and concisely.
(b) Manipulation and Measurement

The ability to:
(i) handle chemicals carefully and use them economically;
(ii) appropriately prepare materials for observation or investigation; and,
(iii) assemble and use simple apparatus and measuring instruments.
(c) Planning and Designing

The ability to:
(i) recognise the problem and formulate valid hypotheses;
(ii) choose appropriate experimental methods and sampling techniques;
(iii) choose appropriate apparatus;
(iv) plan and execute experimental procedures in a logical and sequential form within the time allotted;
(v) use controls where appropriate; and,
(vi) modify experimental methods after initial work or unexpected outcomes.

## - PREREQUISITES OF THE SYLLABUS

Any person with a good grasp of the Caribbean Secondary Education Certificate (CSEC ${ }^{\circledR}$ ) Chemistry and Mathematics syllabuses, or the equivalent, should be able to pursue the course of study defined by this syllabus. However, successful participation in the course of study will also depend on the possession of good verbal, written communication skills.

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## - STRUCTURE OF THE SYLLABUS

The subject is organised in two (2) Units. A Unit comprises three (3) Modules each requiring 50 hours. The total time for each Unit, is therefore, expected to be 150 hours. Each Unit can independently offer students a comprehensive programme of study with appropriate balance between depth and coverage to provide a basis for further study in this field.

Unit 1: Chemical Principles and Applications I

| Module 1 | - | Fundamentals in Chemistry |
| :--- | :--- | :--- |
| Module 2 | - | Kinetics and Equilibria |
| Module 3 | - | Chemistry of the Elements |

## Unit 2: Chemical Principles and Applications II

Module 1 - The Chemistry of Carbon Compounds
Module 2 - Analytical Methods and Separation Techniques
Module 3 - Industry and the Environment

It is recommended that of the approximately 50 contact hours suggested for each Module, a minimum of about 20 contact hours be spent on laboratory related-activities, such as conducting experiments, making field trips and viewing audio-visual materials.

## - SUGGESTIONS FOR TEACHING THE SYLLABUS

It is recommended that Unit 1, Module 1 be taught first. However, in teaching each section, teachers need not follow the sequence given. SI units and IUPAC conversion of nomenclature should be used throughout. For each Module, there are general and specific objectives. The general and specific objectives indicate the scope of the content, including practical work, on which the examination will be based. However, unfamiliar situations may be presented as stimulus material in a question. Explanatory notes are provided to the right of some specific objectives. These notes provide further guidance to teachers as to the level of detail required. The Suggested Practical Activities indicate those areas of the syllabus that are suitable for practical work. However, practical work should not necessarily be limited to these activities.

Teachers are strongly encouraged to integrate ICT and use inquiry-based strategies to teach chemical concepts. Teachers should ensure that their lessons stimulate the students' curiosity and facilitate critical thinking and problem-solving. This will help students view Chemistry as a dynamic and exciting investigative process. The provision of cooperative and collaborative activities is encouraged to facilitate the development of teamwork and the entrepreneurial spirit.

This syllabus caters to varying teaching and learning styles, with specific attention made to ensure the interrelatedness of concepts. Whenever possible, a variety of teaching and learning strategies suitable to varying learning needs of students should be employed with special attention given to the identification of variables and the use of controls in chemical investigations. The need for repeated investigations and observations to arrive at meaningful conclusions should be emphasised.

In addition to developing a solid foundation of factual information, teachers are encouraged to emphasise the application of scientific concepts and principles and minimise memorisation and rote
learning. In order to make the course as relevant as possible, every opportunity should be taken to help students make the connections between chemistry and their environment.

The role of the teacher is to facilitate students learning accurate and unbiased information that will indirectly contribute to a more scientifically literate citizenry that is capable of making educated decisions regarding the world in which we live.

## - THE PRACTICAL APPROACH

The syllabus is designed to foster the use of inquiry-based learning through the application of the practical approach. Students will be guided to answer scientific (testable) questions by a process of making observations, asking questions, doing experiments and analysing and interpreting data. Students should be made aware of the environmental impact of the improper disposal of waste and the associated safety hazards. Teachers should emphasise the use of appropriate safety gear and the need for acceptable laboratory practices. The CAPE ${ }^{\circledR}$ Chemistry Syllabus focuses on the skills listed below.

## 1. Planning and Designing (PD)

Student's ability to:
(a) Ask questions: how, what, which, why or where. (Students must be guided by their teachers to ask scientific questions based on a stated problem).

Sample Problem: It has been observed that exposed wine usually acquires a sour taste after a few weeks.

Example: Why do wines which are exposed to air acquire a sour taste after a few weeks?
(b) Construct a hypothesis; the hypothesis must be clear, concise and testable.

Example: When wines are exposed to air, they acquire a sour taste after a few weeks.
(c) Design an experiment to test the hypothesis. Experimental reports must include the following:
(i) problem statement;
(ii) aim;
(iii) list of materials and apparatus to be used;
(iv) clear and concise step by step procedure;
(v) manipulated and responding variables;
(vi) controlled variables;
(vii) observations to be made or measurements to be taken;
(viii) suggested display of results (for example, graphs tables);
(ix) proposed use of results;
(x) possible limitations, assumptions; and,
(xi) precautions to be taken.

## 2. Measurement and Manipulation (MM)

Student's ability to:
(a) Handle scientific equipment competently.

The list of equipment includes:
(i) Bunsen burner;
(ii) measuring cylinder;
(iii) beakers;
(iv) thermometer;
(v) ruler;
(vi) stop watch/clock;
(vii) balance;
(viii) boiling tube;
(ix) burette;
(x) pipette;
(xi) conical flask;
(xii) syringe;
(xiii) sintered glass crucible;
(xiv) suction pump;
(xv) voltmeter; and,
(xvi) ammeter.

This list is not exhaustive.
(b) Use appropriate apparatus.
(c) Take accurate measurements.

## 3. Observation, Reporting and Recording (ORR)

(a) Recording

Student's ability to record observations and to collect and organise data. Observations and data may be recorded in the following format.
(i) Prose - Written description of observations in the correct tense.
(ii) Table (Neatly enclosed):

- Numerical: physical quantities in heading, correct units stated in heading, symbols, decimal points.

Non-numerical: headings correct, details present.
(iii) Graph - Axes labelled with units, correct scales, correct plotting, and smooth curves/best fit lines.
(iv) Drawing of apparatus as set up for use.
(b) Reporting

Student's ability to prepare a comprehensive written report on their assignments using the following format:
(i) Date (date of experiment).
(ii) Aim (what is the reason for doing the experiment).
(iii) Apparatus and Materials (all equipment, chemicals and materials used in the experiment must be listed).
(iv) Method/Experimental Procedure (step-by-step procedure written in the past tense, passive voice).
(v) Results and Observations (see (a) above: Recording).
(vi) Discussion (see 4: Analysis and Interpretation).
(vii) Conclusion (should be related to the Aim).

## 4. Analysis and Interpretation

Student's ability to:
(a) identify patterns and trends;
(b) make accurate calculations;
(c) identify limitations and sources of error;
(d) make a conclusion to either support or refute the hypothesis;
(e) compare actual results with expected results if they are different;
(f) suggest alternative methods or modifications to existing methods; and,
(g) analyse and interpret results and observations, and make conclusions.

## - UNIT 1: CHEMICAL PRINCIPLES AND APPLICATIONS I MODULE 1: FUNDAMENTALS IN CHEMISTRY

## GENERAL OBJECTIVES

On completion of this Module, students should:

1. understand that theories in chemistry are subject to change;
2. understand the theory of atoms as a useful construct that explains the structure and behaviour of matter, and the impact of nuclear chemistry on society;
3. understand the development of the periodic table for the classification of elements;
4. appreciate that the forces of attraction between particles influence the properties and behaviour of matter;
5. understand the mole concept;
6. understand redox reactions;
7. understand the kinetic theory;
8. understand concepts associated with energy changes; and,
9. develop the ability to perform calculations involving energy changes.

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES
SUGGESTED PRACTICAL ACTIVITIES

## 1. Atomic Structure and the Periodic Table

Students should be able to:

| 1.1. | discuss the process of theoretical change with respect to Dalton's atomic theory; | The postulates of Dalton's Atomic theory and modifications of the theory. Mention the criteria that are considered when theories are accepted, for example, fit between evidence and theoretical constructs, reliability and accuracy of data, replicability of experiments, consensus within the scientific community, societal |
| :---: | :---: | :---: |

## UNIT 1 <br> MODULE 1: FUNDAMENTALS IN CHEMISTRY (cont'd)

## SPECIFIC OBJECTIVES

## Atomic Structure and the Periodic Table (cont'd)

Students should be able to:
1.2. describe the structure of the atom;

Simple treatment: properties of protons, neutrons and electrons only; their relative masses and charges, location and their behaviour in electric and magnetic fields.
1.3. define the following terms:
(a) mass number;
(b) isotopes; and,
(c) relative atomic and Must include reference to isotopic masses the mass of carbon-12
based on the ${ }_{6}^{12} \mathrm{C}$ scale.
1.4. explain the phenomenon

Write equations of radioactivity;


## UNIT 1 <br> MODULE 1: FUNDAMENTALS IN CHEMISTRY (cont'd)

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES
SUGGESTED PRACTICAL ACTIVITIES

## Atomic Structure and the Periodic Table (cont'd)

## Students should be able to:

1.6. calculate the relative
atomic mass of an
element, given isotopic
masses and abundances;
1.7. explain how data from emission spectra provide evidence for discrete energy levels within the atom;
1.8. $\begin{aligned} & \text { describe the atomic } \\ & \text { orbitals; }\end{aligned}$

Bohr model, simple treatment of the emission spectrum of hydrogen; Lyman series, Balmer series; $\Delta \mathrm{E}$ or $\mathrm{dE}=\mathrm{hv}$.

Principal quantum numbers, $s, p$ and $d$ orbitals; relative energies of 4 s and 3 d orbitals.

Refer to Module, 3 Specific Objective 5.1.
1.9. describe the shapes of the $s$ and $p$ orbitals;
1.10. determine the electronic configurations of atoms and ions in terms of $s, p$ and d orbitals;
1.11. state the factors which influence the first ionisation energy of elements;
1.12. explain how ionisation energy data provide evidence for sub-shells; and,
1.13. derive the electronic configuration of an element from data on successive ionisation energies.

Consider elements from atomic numbers 1 to 30 .

Include atomic radii, nuclear charge, shielding.

Use Period 3 as an example.

## UNIT 1 <br> MODULE 1: FUNDAMENTALS IN CHEMISTRY (cont'd)

## SPECIFIC OBJECTIVES

## 2. Forces of Attraction

Students should be able to:
2.1. state the various forces of attraction between particles;

EXPLANATORY NOTES

Ionic bonds, covalent bonds, hydrogen bonds, metallic bonds, Van der Waals forces. (Permanentpermanent dipole; induced-induced dipole or temporary/instantaneousinduced dipole).
2.2. state the relationship between forces of attraction and states of matter;
2.3. relate physical properties of matter to differences in strength of forces of attraction;

Variation in melting points, boiling points and solubilities.

Conduct melting point and boiling point determinations;
solubilities in polar and non-polar solvents, electrical conductivity. Illustrate practically the properties of ionic and covalent compounds.
$\begin{array}{lll}\text { 2.5. } \begin{array}{l}\text { describe } \\ \text { (dative covalent) bonding; }\end{array} & \begin{array}{l}\text { Use 'dot-cross' diagrams; } \\ \text { refer to simple systems (for } \\ \left.\text { example, } \mathrm{BF}_{3} / \mathrm{NH}_{3}\right) .\end{array}\end{array}$

## SUGGESTED PRACTICAL ACTIVITIES

## UNIT 1 <br> MODULE 1: FUNDAMENTALS IN CHEMISTRY (cont'd)

## SPECIFIC OBJECTIVES

## Forces of Attraction (cont'd)

## Students should be able to:

## 2.6. describe the origin of intermolecular forces;

2.7. predict the shapes of, and bond angles in simple molecules and ions;

EXPLANATORY NOTES

Refer to hydrogen bonding; Van der Waals forces, permanent dipole. Refer to Module 3

Application of the VSEPR theory to include the following systems: trigonal (for example, $\mathrm{BF}_{3}$ ), linear (for example, $\mathrm{BeCl}_{2}$ ), tetrahedral (for example, $\mathrm{NH}_{4}{ }^{+}, \mathrm{CH}_{4}$ ), pyramidal (for example, $\mathrm{H}_{3} \mathrm{O}^{+}, \mathrm{CH}_{3}$, and NH 3 ), non-linear (for example, $\mathrm{H}_{2} \mathrm{O}$ ), octahedral (for example, $\mathrm{SF}_{6}$ ).
2.8. explain the shapes and bond angles of simple organic compounds;
2.9. predict the shapes and bond angles of molecules similar to ethane; and,
2.10. describe qualitatively the lattice structure of crystalline solids and their relation to physical properties.

Ethane, ethene and benzene; apply the concept of hybridisation and resonance. Include $\mathrm{sp}^{2}$ and $\mathrm{sp}^{3}$ hybridisation.

Simple substituted derivatives, for example, dichloroethane.

Simple molecular (for example, $\mathrm{I}_{2}$ ), hydrogen bonded (for example, ice), giant molecular (for example, $\mathrm{SiO}_{2}$ ), ionic (for example, NaCl ), metallic (for example, Cu ), giant atomic (for example, graphite and diamond) structures.

## SUGGESTED PRACTICAL ACTIVITIES

Construct molecular models and measure bond angles.

## UNIT 1 <br> MODULE 1: FUNDAMENTALS IN CHEMISTRY (cont'd)

## SPECIFIC OBJECTIVES

## 3. The Mole Concept

Students should be able to:
3.1. apply Avogadro's law;
3.2. define the mole;
3.3. define the term 'molar mass';
3.4. write balanced molecular and ionic equations;
3.5. $\begin{aligned} & \text { perform } \\ & \text { based on the mole }\end{aligned}$ concept;
3.6. apply the mole concept to molecular and ionic equations;
3.7. calculate empirical and molecular formulae;
3.8. perform titrimetric analyses; and,

## EXPLANATORY NOTES

Perform calculations involving molar volumes.

Relate to masses of substances, volumes of gases, volumes and concentrations of solutions.

Combustion data; absolute masses or relative abundances of elements.

Conduct acid/base titrations and redox titrations. (dichromate (VI)), hydrogen peroxide, iodide thiosulfate, manganate (VII); mean (consecutive accurate values within $0.10 \mathrm{~cm}^{3}$ of each other), significant figures.
3.9. use results from titrimetric analyses to calculate:
(a) mole ratios;
(b) molar concentration; and,
(c) mass concentration.

## SUGGESTED PRACTICAL ACTIVITIES

## UNIT 1 <br> MODULE 1: FUNDAMENTALS IN CHEMISTRY (cont'd)

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES

## 4. Redox Reactions

Students should be able to:
4.1. explain redox reactions in terms of electron transfer and changes in oxidation state (number);
4.2. construct relevant half equations for redox reactions;
4.3. deduce balanced equations for redox reactions from relevant half equations; and,
4.4. order elements in terms of oxidising or reducing ability.

Refer to Module 1, Specific Objective 3.8.

Redox equations should be constructed under both acidic and basic conditions.

## 5. Kinetic Theory

Students should be able to:
5.1. state the basic assumptions of the kinetic theory with reference to an ideal gas;
5.2. explain the differences between real and ideal gases;

## SUGGESTED PRACTICAL ACTIVITIES

Perform simple displacement reactions to order elements in terms of oxidising or reducing ability; addition of zinc to copper (II) sulfate solution; addition of chlorine water to bromide or iodide solutions.

Qualitative treatment only - the conditions which are necessary for a gas to approach ideal behaviour, the limitations of ideality at very high pressures and very low temperatures. Include graphical representations.

## UNIT 1

MODULE 1: FUNDAMENTALS IN CHEMISTRY (cont'd)

## SPECIFIC OBJECTIVES

## Kinetic Theory cont'd

Students should be able to:
5.3. state Boyle's law and Charles' law;
5.4. perform calculations using:
(a) Boyle's law;
(b) Charles' law; and,
(c) the ideal gas equation ( $\mathrm{pV}=\mathrm{nRT}$ ); and,
5.5. explain the following:
(a) the liquid state;
(b) melting; and,
(c) vaporisation.

## 6. Energetics

Students should be able to:
6.1. state that chemical reactions take place through energy changes (usually in the form of heat) associated with the breaking and making of bonds;
6.2. state that energy changes occur in chemical reactions associated with the making and breaking of bonds;
6.3. explain the differences between exothermic and endothermic reactions using energy profile diagrams;

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## EXPLANATORY NOTES

Include graphical representations.

Calculations involving the use of Van der Waals equation of state are not required.

Include calculations of relative molar mass.

## SUGGESTED PRACTICAL

 ACTIVITIES
## UNIT 1 <br> MODULE 1: FUNDAMENTALS IN CHEMISTRY (cont'd)

## SPECIFIC OBJECTIVES

## Energetics cont'd

## Students should be able to:

6.4. explain the term 'bond energy';
6.5. explain how bond energy data may be used to show the relationship between strength of covalent bonds and reactivity of covalent molecules;
6.6. apply concepts associated with enthalpy changes;
6.7. explain the effect of ionic charge and radius on the magnitude of lattice energy;
6.8. state Hess's law of constant heat summation; and,
6.9. calculate enthalpy changes from appropriate experimental data.

## EXPLANATORY NOTES

SUGGESTED PRACTICAL ACTIVITIES

Calculations involving bond energy data.

Lack of reactivity of nitrogen. Consider factors which affect bond energy.

Include enthalpy change of formation, combustion, neutralisation, reaction, hydration, solution, atomisation, ionisation energy, electron affinity and lattice energy.

No calculation needed.

Use standard conditions.
This will require construction of energy cycles including Born Haber cycles. Data may be obtained experimentally or provided.

Experiments may include heats of reaction, solution and neutralisation.

UNIT 1
MODULE 1: FUNDAMENTALS IN CHEMISTRY (cont'd)

## Suggested Teaching and Learning Activities

To facilitate students' attainment of the objectives of this Module, teachers are advised to engage students in the teaching and learning activities listed below.

## Atomic Structure and the Periodic Table

1. Ask students to read A Short History of Nearly Everything by Bill Bryson and discuss the history of the development of the atomic models. (Audiobook available on YouTube).
2. Allow students to carry out practical weighing activities which compare the mass of different objects (for example, coins) in order to develop the concept of relative mass and changing standards of comparison.
3. Ask students to present the story of the discovery of the phenomenon of radioactivity (use video material if available).
4. Have class discussion on the impact of radioactivity in everyday life as cited (from newspaper articles and the electronic media including the Internet).
5. Provide students with appropriate reading material prior to class session. During the class session, teacher and students engage in a discussion on the strengths and weaknesses of the Bohr and Rutherford models of the atom.
6. Have class discussions on the evidence that led to modification of Dalton's atomic theory and on the historical development of the Periodic Table.

## Forces of Attraction

1. Arrange students in small groups, and provide them with appropriate quantitative data and guided questions which will lead them to infer that forces of attraction vary in strength.
2. Ask students to use ball and stick to make models for different molecular shapes.

## The Mole Concept

1. Use appropriate analogies to explain that the mole is a specific amount of particles (atoms, molecules, ions, electrons).
2. Allow students to conduct laboratory work including dilution factor, titration, displacement and yield calculations.

## Redox Reactions, Kinetic Theory and Energetics

1. Use practical activities, diagrams, graphs and guided questions to enhance students' understanding of different concepts.

## UNIT 1 <br> MODULE 1: FUNDAMENTALS IN CHEMISTRY (cont'd)

## RESOURCES

Teachers and students may find reference to the following resource materials useful. The latest editions are recommended.

Amateis, P., and Silberberg, M. Chemistry: The Molecular Nature of Matter and Change. McGraw-Hill Education, 2014.

Cann, P. and Hughes, P. Chemistry, International AS and A Level. London: Hodder Education, 2015.

Clarke, J.
Calculations in AS/A Level Chemistry. Essex: Pearson Education Limited, 2000.

Conoley, C. and Hills, P.
Chemistry, $3^{\text {rd }}$ Edition. London: HarperCollins, 2008.

Hill, G., and Holman, J.
Chemistry in Context. London: Nelson Thorne Limited, 2001.

Lister, T., Renshaw, J.
Understanding Chemistry for Advanced Level. Cheltenham: Trans-Atlantic Publications, 2000.

Maylin-Moseley, V.
Advanced Level Chemistry for Life - Unit 1. Barbados: VHM Publishing, 2017.

Norris, R., Barrett, L., Maynard-Alleyne, A. and Murray, J.

Ramsden, E.
CAPE ${ }^{\circledR}$ Chemistry Study Guide: Cheltenham: Nelson Thorne Limited, 2012.

A-Level Chemistry. Cheltenham: Nelson Thorne Limited, 2000.

## WEBSITES

www.Chemsoc.org
www.Chemguide.co.uk
www.creative-chemistry.org.uk
www.a-levelchemistry.co.uk

## UNIT 1 <br> MODULE 2: KINETICS AND EQUILIBRIA

## GENERAL OBJECTIVES

On completion of this Module, students should:

1. understand the concepts associated with reaction rates;
2. understand the concepts associated with chemical equilibrium;
3. appreciate that equilibrium concepts can be applied to chemical systems; and,
4. appreciate that principles of kinetics and equilibria can be applied to industrial and biological processes.

## SPECIFIC OBJECTIVES

SUGGESTED PRACTICAL

## 1. Rates of Reaction

Students should be able to:
1.1. explain the concepts Include a study of rate associated with reaction constant, order of reaction,
1.2. design suitab experiments for studying the factors which affect rates of reactions;

EXPLANATORY NOTES ACTIVITIES
rates;
1.1. associated with reaction
half-life, rate-determining step, activation energy, collision theory, (simple treatment only), and catalysis. Include enzymes in industrial and biological processes.

Include effects of concentration, temperature and catalysts.
1.3. construct rate equations of the form: Rate $=k[A]^{n}[B]^{m}$ limited to simple cases involving zero, first and second order reactions;

Rate equations may be derived or deduced from experimental data supplied.

Conduct suitable experiments for studying the factors which affect rates of reactions; express results in the form of tables and graphs.
1.4. deduce the order of Include deductions of reaction from appropriate possible reaction data;
mechanisms.

## UNIT 1

MODULE 2: KINETICS AND EQUILIBRIA (cont'd)

## SPECIFIC OBJECTIVES

## EXPLANATORY NOTES

## Rates of Reaction cont'd

Students should be able to:

| 1.5. | interpret concentration |
| :--- | :--- |
| against time and |  |
| concentration against rate |  |
| for zero and first order |  |
| reactions; |  |

1.6. perform calculations from rate data;
1.7. perform simple calculations using half-life data; and,
1.8. explain the effect of temperature and catalysts on the rate of the reaction using Boltzmann distribution of energies (and of collision frequency).

| Qualitative | and |
| :--- | ---: |
| quantitative | treatments |
| required. |  | quantitative treatments required.

Calculate initial rates and rate constants.

Limited to first order reactions.

Include the use of Boltzmann distribution curves.

## 2. Principles of Chemical Equilibrium

## Students should be able to:

2.1. explain the concept of dynamic equilibrium;
2.2. state the characteristics of a system in dynamic equilibrium;
2.3. define the terms $K_{c}$ and $K_{p}$;

Consider examples of static and dynamic equilibrium. Refer to physical and chemical processes.

Write equilibrium constant expressions in terms of $K_{c}$ and $K_{p}$.

Conduct a simple experiment to determine the value of $K_{c}$ for $a$ reaction.
2.4. perform calculations involving equilibrium constants in terms of concentration, $\left(\mathrm{K}_{\mathrm{c}}\right)$ and partial pressure, $\left(K_{p}\right)$;

Conversion of $K_{c}$ to $K_{p}$ is not required. Quadratic equations are not required.

## SUGGESTED PRACTICAL ACTIVITIES

## UNIT 1

MODULE 2: KINETICS AND EQUILIBRIA (cont'd)

## SPECIFIC OBJECTIVES

## Principles of Chemical Equilibrium cont'd

## Students should be able to:

2.5.
state
principle;
2.6. apply Le Chatelier's
principle to explain the effects of changes in temperature, concentration and pressure on a system in equilibrium; and,
2.7. interpret how changes in concentration, pressure, temperature or the presence of a catalyst may affect the value of the equilibrium constant. equilibrium constant.

## 3. $\quad$ Acid/Base Equilibria

Students should be able to:
3.1. explain the differences in
behaviour of strong and
weak acids and bases, using
Bronsted-Lowry theory;
3.1. explain the differences in
behaviour of strong and
weak acids and bases, using
Bronsted-Lowry theory;
3.1. explain the differences in
behaviour of strong and
weak acids and bases, using
Bronsted-Lowry theory;
3.1. explain the differences in
behaviour of strong and
weak acids and bases, using
Bronsted-Lowry theory;
3.2. define the terms $\mathrm{K}_{\mathrm{a}}, \mathrm{pH}, \mathrm{pK}_{\mathrm{a}}$, and $\mathrm{pK}_{\mathrm{b}}, \mathrm{K}_{\mathrm{w}}$ and $\mathrm{pK}_{\mathrm{w}}$;
3.3. perform calculations
involving $\mathrm{pH}, \mathrm{pOH}, \mathrm{K}_{\mathrm{a}}, \mathrm{pK}_{\mathrm{a}} \mathrm{K}_{\mathrm{w}}$
$\begin{array}{ll}\text { 3.3. perform calculations } \\ & \text { involving } \mathrm{pH}, \mathrm{pOH}, \mathrm{K}_{\mathrm{a}}, \mathrm{pK}_{\mathrm{a}} \mathrm{K}_{\mathrm{w}}\end{array}$ and $\mathrm{pK}_{\mathrm{w}}, \mathrm{K}_{\mathrm{b}}$ and $\mathrm{p} \mathrm{K}_{\mathrm{b}}$;
3.4. describe the changes in
3.4. describe the changes in
pH during acid/base titrations;
3.5. explain what is meant by
the pH range of indicator; and,
2.6. principle to explain the
effects of changes in
temperature, concentration
and pressure on a system in
equilibrium; and,

## EXPLANATORY NOTES

## SUGGESTED PRACTICAL ACTIVITIES

Include references to the Haber process and the Contact process.
Include reference to the characteristics of a system in dynamic equilibrium.

Quadratic equations are not required.

Include a study of titration curves.

Perform calculations based on the profitability of these processes on
manufacturing of commercial commodities.

## UNIT 1 <br> MODULE 2: KINETICS AND EQUILIBRIA (cont'd)

## SPECIFIC OBJECTIVES

## Acid/Base Equilibria cont'd

## Students should be able to:

3.6. state the basis for the selection of acid/base indicator for use in titrations.

## 4. Buffers and pH

Students should be able to:
4.1. define the term 'buffer solution';
4.2. explain how buffer solutions control pH ;
4.3. calculate the pH of buffer solutions from appropriate data; and,
4.4. discuss the importance of buffers in biological systems and in industrial processes.

Include phenolphthalein and methyl orange. Titration curves.

EXPLANATORY NOTES

SUGGESTED PRACTICAL ACTIVITIES

Perform experiments to show that the effectiveness of different indicators is related to the pH changes which occur during titration.

Perform simple experiments to determine the pH of buffer solutions.

## 5. Solubility Product

Students should be able to:
5.1. define the term solubility product, $\mathrm{K}_{\mathrm{sp}}$;

Include reference to blood buffer systems such as hydrogencarbonate, phosphate and amino- acid systems, enzyme catalysed reactions and the food processing industry.

| Write | equilibrium |
| :--- | ---: |
| constant | expression for |
| K $_{\text {sp. }}$ |  |

5.2. explain the principles underlying solubility product and the common ion effect;

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## UNIT 1

MODULE 2: KINETICS AND EQUILIBRIA (cont'd)

## SPECIFIC OBJECTIVES

## Solubility Product cont'd

Students should be able to:
5.3. perform calculations involving solubility product; and,
5.4. relate the solubility product principle to the selective precipitation of substances.

## 6. Redox Equilibria

Students should be able to:
6.1. define the terms standard electrode potential and standard cell potential;
6.2. describe the standard hydrogen electrode;
6.3. describe methods used to measure the standard electrode potentials of:
(a) metals or nonmetals in contact with their ions in aqueous solutions; and,
(b) ions of the same element in different oxidation states;
6.4. calculate standard cell potentials from standard electrode potentials of two half cells;

## EXPLANATORY NOTES

Quadratic equations are not required.

Include reference to qualitative analysis and kidney stone formation.

SUGGESTED PRACTICAL ACTIVITIES

## UNIT 1 <br> MODULE 2: KINETICS AND EQUILIBRIA (cont'd)

## SPECIFIC OBJECTIVES

## Redox Equilibria cont'd

Students should be able to:
6.5. use standard electrode potentials of cells:
(a) to determine the direction of electron flow; and,
(b) to determine the feasibility of a reaction;
6.6. predict how the value of an electrode potential varies with concentration; and,
6.7. apply the principles of redox processes to energy storage devices.

EXPLANATORY NOTES
SUGGESTED PRACTICAL ACTIVITIES

> Include cell diagram or notation of the type $\mathrm{Zn}(\mathrm{s}) \mid$ $\mathrm{Zn}^{2+}{ }_{(\text {aq })}| | \mathrm{Cu}^{2+}{ }_{\text {(aq) }} \mid \mathrm{Cu}(\mathrm{s})$.

No treatment of the Nernst equation is required. Apply Le Chatelier's principle.

Include references to two of the following batteries: Leclanche' dry cell, lead acid accumulators (secondary cells); and fuel cells.

## Suggested Teaching and Learning Activities

To facilitate students' attainment of the objectives of this Module, teachers are advised to engage students in the teaching and learning activities listed below.

1. Use appropriate analogies, for example, a moving object on an escalator in motion to distinguish between static and dynamic equilibria so that students get a better understanding of the changes at the microscopic level as opposed to the apparent lack of change at the macroscopic level.
2. Identify suitable practical activities to enhance the theory. It is important that students are conversant with the manipulation of experimental data. In this respect, students should be given the opportunity to develop the various concepts in a stepwise manner. For example, in the determination of rate constant the following sequence of steps can be used:

## UNIT 1 <br> MODULE 2: KINETICS AND EQUILIBRIA (cont'd)

Plot concentration time graph $\rightarrow$ draw tangents to obtain the rates at different concentrations $\rightarrow$ draw rate concentration graphs $\rightarrow$ use slope of graphs to obtain a value for the rate constant.
3. Provide students with appropriate data to work out a variety of problems including:
(a) orders of reactions (practise writing rate equations); and,
(b) rate and equilibrium constant including $\mathrm{K}_{\mathrm{a}}$ and $\mathrm{K}_{\mathrm{b}}, \mathrm{pH} \leftrightarrow\left[\mathrm{H}^{+}\right], \mathrm{pOH} \leftrightarrow\left[\mathrm{OH}^{-}\right]$, and $\mathrm{K}_{\mathrm{w}}$.

It is essential that students be given sufficient practice at these calculations.
4. Emphasise the practical applications of redox reactions to show that the equilibria in electrochemical cells are redox in nature. From here, students may practise writing cell diagrams to determine, for example:
(a) the direction of electron flow;
(b) the nature of the electrodes;
(c) the reaction that may occur; and,
(d) cell potentials.
5. Engage students in a brief discussion on the importance of Kinetics and Equilibria to industrial and biological processes.
6. Ask students to conduct research on kidney stone formation and its prevention.

## UNIT 1 <br> MODULE 2: KINETICS AND EQUILIBRIA (cont'd)

## RESOURCES

Teachers and students may find reference to the following resource materials useful. The latest editions are recommended.

| Amateis, P., and Silberberg, M. | Chemistry: The Molecular Nature of Matter and <br> Change. McGraw-Hill Education, 2014. |
| :--- | :--- |
| Cann, P. and Hughes, P. | Chemistry, International AS and A Level. London: <br> Hodder Education, 2015. |
| Clarke, J. | Calculations in AS/A Level Chemistry. Essex: Pearson <br> Education Limited, 2000. |
| Conoley, C. and Hills, P. | Chemistry, 3 |
| Clugston, Edition. London: HarperCollins, 2008. |  |

## WEBSITES

www.Chemsoc.org
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## www.creative-chemistry.org.uk

www.a-levelchemistry.co.uk

## UNIT 1 <br> MODULE 3: CHEMISTRY OF THE ELEMENTS

## GENERAL OBJECTIVES

On completion of this Module, students should:

1. use fundamental concepts to rationalise the physical and chemical properties of elements and their compounds;
2. appreciate that the properties of elements are related to their compounds and their uses; and,
3. understand the principles underlying the identification of anions and cations.

## SPECIFIC OBJECTIVES

## 1. Period 3: Sodium to Argon

Students should be able to:
1.1. explain the variations in physical properties of the elements in terms of structure and bonding;
1.2. describe the reactions of the elements with oxygen, chlorine and water;
1.3. explain the variation in oxidation number of the oxides and chlorides;
1.4. describe the reactions of the oxides and chlorides with water;
1.5. explain the trend in the acid/base behaviour of the oxides and hydroxides;

EXPLANATORY NOTES
SUGGESTED PRACTICAL ACTIVITIES

Include reference to melting point and electrical conductivity. Atomic and ionic radii, electronegativity and density. Refer to Module 1, Specific Objective 1.11.

No treatment of peroxides or superoxides required.

Include equations.

Include equations.

Conduct experiments to investigate the reactions of the oxides and chlorides with water; include relevant equations.

Conduct experiments to investigate the acid/base behavior of the oxides and hydroxides; include relevant equations.

## UNIT 1

MODULE 3: CHEMISTRY OF THE ELEMENTS (cont'd)

## SPECIFIC OBJECTIVES

## Period 3: Sodium to Argon cont'd

Students should be able to:
1.6. predict the types of chemical bonding present in the chlorides and oxides; and,
1.7. discuss the uses of some of the compounds of aluminium and phosphorous.

## 2. Group II Elements

Students should be able to:
2.1. explain the variations in properties of the elements in terms of structure and bonding;
2.2. describe the reactions of the elements with oxygen, water, and dilute acids;
2.3. explain the variation in the solubility of the sulfates;

EXPLANATORY NOTES

SUGGESTED PRACTICAL ACTIVITIES

Refer to differences in electronegativities and ionic radii of the elements.

Limited to the use of aluminium hydroxide in antacid medication, white phosphorous used in flares and military applications, red phosphorous used at the side of match boxes and argon used in fluorescent and incandescent lighting.

Include reference to atomic and ionic radii and ionisation energies.

Include equations.

Qualitative treatment only is required. Simple explanations in terms of lattice and hydration energies.

Include equations.

Limited to the use of magnesium oxide, calcium oxide, calcium hydroxide and calcium carbonate.

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## UNIT 1

MODULE 3: CHEMISTRY OF THE ELEMENTS (cont'd)

## SPECIFIC OBJECTIVES

## 3. Group IV Elements

Students should be able to:
3.1. explain the variations in physical properties of the elements in terms of structure and bonding;
3.2. describe the bonding of the tetrachlorides;
3.3. explain the reactions of the tetrachlorides with water;
3.4. discuss the trends in:
(a) bonding;
(b) acid/base character; and,
(c) thermal stability of the oxides of oxidation states II and IV;
3.5. discuss the relativ stabilities of the oxides and aqueous cations of the elements in their higher and lower oxidation states; and,
3.6. discuss the uses of ceramics based on silicon (IV) oxide.

## 4. Group VII Elements

Students should be able to:
4.1. explain the variations in physical properties of the elements in terms of structure and bonding;

EXPLANATORY NOTES
Include reference to
variations in metallic
character and electrical
conductivity.

SUGGESTED PRACTICAL ACTIVITIES

Include equations.

Make reference to $E^{\theta}$ values of the elements.

Include equations.

Make reference to $E^{\ominus}$ values of the elements.

Include its use as abrasives, furnace lining, glass and porcelain. Relate use to properties.

Volatility, density, colour, and state. (An explanation of colour is not required).

## UNIT 1 <br> MODULE 3: CHEMISTRY OF THE ELEMENTS (cont'd)

## SPECIFIC OBJECTIVES

## Group VII Elements cont'd

Students should be able to:
4.2. explain the relativ reactivities of the elements as oxidising agents;
4.3. describe the reactions of the elements with hydrogen;
4.4. explain the relative thermal stabilities of the hydrides;
4.5. describe the reactions of the halide ions with:
(a) aqueous solution of $\mathrm{AgNO}_{3}$ followed by aqueous ammonia; and,
(b) concentrated sulfuric acid; and,
4.6. describe the reactions of chlorine with cold and hot aqueous solution of sodium hydroxide.

## EXPLANATORY NOTES

Include reactions with sodium thiosulfate and refer to $E^{\theta}$ values.

Include equations.

Include bond energies in explanations.

Include changes in oxidation number and the process of disproportionation.
Refer to Module 1, Specific Objective 4.1.

## 5. First Row Transition Elements

Students should be able to:
5.1. define the term transition element;
5.2. describe the characteristics transition elements;

D-block elements forming one or more stable ions with incomplete d-orbitals.
the Include variation in of oxidation number, complex formation, coloured compounds, catalytic activity, magnetic properties.

SUGGESTED PRACTICAL ACTIVITIES

Use solutions of the elements with bleach, bromine water, and iodine solution.

Perform experiments of halide ions with aqueous $\mathrm{AgNO}_{3}$ followed by aqueous ammonia.

## UNIT 1 <br> MODULE 3: CHEMISTRY OF THE ELEMENTS (cont'd)

## SPECIFIC OBJECTIVES

## First Row Transition Elements cont'd

Students should be able to:
5.3. discuss qualitatively the properties of transition elements when compared to those of calcium as a typical s-block element;
5.4. determine the electronic configuration of the first row transition elements and of their ions;
5.5. explain the relatively small changes in atomic radii, ionic radii, and ionisation energies of the elements across the period;
5.6. explain the formation of coloured ions by transition elements;
5.7. describe the variation in oxidation states of vanadium;

EXPLANATORY NOTES

Melting point, density, atomic radius, ionic radius, first ionisation energy, and conductivity.

Mention changes in oxidation number.
d-orbital $\quad$ separation
of energy in octahedral
complexes.

## Refer to $E^{\theta}$ values.

Perform experiments to include the use of an acidified solution of ammonium vanadate (V) and granulated zinc.

Octahedral, tetrahedral and square planar.

Refer to Module 1, Specific Objective 4.4.

Stability constants and the $\mathrm{CO} / \mathrm{O}_{2}$ haemoglobin and $\mathrm{NH}_{3(\mathrm{aq})} / \mathrm{Cu}^{2+}{ }_{(\mathrm{aq})}$ systems.

Perform experiments to demonstrate ligand exchange. Include reactions involving $\mathrm{Co}^{2+}{ }_{(\mathrm{aq})}$, $\mathrm{Cu}^{2+}{ }_{(\mathrm{aq})}$.

## UNIT 1 <br> MODULE 3: CHEMISTRY OF THE ELEMENTS (cont'd)

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES

## SUGGESTED PRACTICAL ACTIVITIES

## 6. Identification of Cations and Anions

Students should be able to:
6.1. identify cations: $\mathrm{K}^{+}, \mathrm{Na}^{+}$, $\mathrm{Ca}^{2+}, \mathrm{Ba}^{2+}, \mathrm{Cu}^{2+}$ by their flame tests;
6.2. identify cations $\mathrm{Mg}^{2+}{ }_{(\mathrm{aq})}$, $\mathrm{Al}^{3+}{ }_{(\mathrm{aq})}, \quad \mathrm{Ca}^{2+}{ }_{(\mathrm{aq})}, \quad \mathrm{Cr}^{3+}{ }_{(\mathrm{aq})}$, $\mathrm{Mn}^{2+}{ }_{(\mathrm{aq})}, \quad \mathrm{Fe}^{2+}{ }_{(\mathrm{aq})}, \quad \mathrm{Fe}^{3+}{ }_{(\mathrm{aq})}$, $\mathrm{Cu}^{2+}{ }_{(\mathrm{aq})}, \quad \mathrm{Zn}^{2+}{ }_{(\mathrm{aq})}, \quad \mathrm{Ba}^{2+}{ }_{(\mathrm{aq})}$, $\mathrm{Pb}^{2+}{ }_{(\mathrm{aq})}, \mathrm{NH}_{4}{ }^{+}{ }_{(\mathrm{aq})}$;
6.3. explain the principles upon
which the reactions in Specific Objective 6.2 are based;
6.4. write ionic equations for the reactions in Specific Objective 6.2;
6.5. identify anions: $\mathrm{CO}_{3}{ }^{2-}, \mathrm{NO}_{3}{ }^{-}, \mathrm{SO}_{4}{ }^{2-}, \mathrm{SO}_{3}{ }^{2-}$ (aq), $\mathrm{Cl}^{-}, \mathrm{Br}^{-}, \mathrm{I}^{-}, \mathrm{CrO}_{4}^{-}$; and,

Refer to atomic emission spectra, see Module 1, Specific Objective 1.7.

Include the reactions with $\mathrm{OH}^{-}(\mathrm{aq}), \mathrm{CO}^{2-}{ }_{3(\text { aq })}$ and $\mathrm{NH}_{3(a q)}$ and confirmatory tests.

Refer to equilibrium concepts. Module 2, Specific Objective 5.2.

Basic, amphoteric oxide and complexation.

Include state symbols.

Include the reactions with $\mathrm{HCl}_{(\mathrm{aq})}$, conc $\mathrm{H}_{2} \mathrm{SO}_{4}, \mathrm{~Pb}^{2+}{ }_{(\mathrm{aq})}$, $\mathrm{Ag}^{+}{ }_{(\mathrm{aq})}$, followed by $\mathrm{NH}_{3(\mathrm{aq})}$, $\mathrm{Ca}(\mathrm{OH})_{2(a q)}, \quad \mathrm{Ba}^{2+}{ }_{(\mathrm{aq})}$, followed by dilute acid. For $\mathrm{NO}_{3}{ }^{-}$, use copper turnings and conc. $\mathrm{H}_{2} \mathrm{SO}_{4}$ or add aluminium (powder) or zinc (powder) in the alkaline solution and confirmatory tests for gases where applicable.

Include state symbols.

Perform flame tests on identified cations.

Perform experiments of the identified cations with hydroxide and aqueous ammonia.

Where possible perform confirmatory tests of the identified cations.

Perform experiments to identify the anions $\mathrm{CO}_{3}{ }^{2-}$, $\mathrm{NO}_{3}{ }^{-}, \mathrm{SO}_{4}{ }^{2-}, \mathrm{SO}_{3}{ }^{2-}(a q), \mathrm{Cl}^{-}, \mathrm{Br}^{-}$, l, $\mathrm{CrO}_{4}^{-}$. Where applicable perform confirmatory tests for gases.
6.6. write ionic equations for the reactions in Specific Objective 6.5.

## UNIT 1 <br> MODULE 3: CHEMISTRY OF THE ELEMENTS (cont'd)

## Suggested Teaching and Learning Activities

To facilitate students' attainment of the objectives of this Module, teachers are advised to engage students in the teaching and learning activities listed below.

1. Review fundamental factors which influence the properties of elements and their compounds, for example, ionisation energy, electronegativity, type of bonding.
2. Allow students to use charts and tables when establishing trends and differences in properties of elements and compounds.
3. Allow students to use computer software in simulations to demonstrate the chemistry of the elements and their compounds.
4. Link theory with appropriate laboratory work and real-life applications such as manufacturing, and agriculture.

## RESOURCES

Teachers and students may find reference to the following resource materials useful. The latest editions are recommended.

Cann, P. and Hughes, P. Chemistry, International AS and A Level. London: Hodder Education, 2015.

Conoley, C. and Hills, P.
Maylin-Moseley, V.

Norris, R., Barrett, L., Maynard-Alleyne, A. and Murray, J.

Ramsden, E.

## WEBSITES

www.Chemsoc.org
www.Chemguide.co.uk
www.creative-chemistry.org.uk
www.a-levelchemistry.co.uk

## - UNIT 2: CHEMICAL PRINCIPLES AND APPLICATIONS II MODULE 1: THE CHEMISTRY OF CARBON COMPOUNDS

## GENERAL OBJECTIVES

On completion of this Module, students should:

1. appreciate the scope and nature of carbon-based compounds;
2. understand the processes involved in the formation of carbon compounds;
3. understand the reactions of various functional groups of carbon compounds; and,
4. critically assess the impact of carbon-based compounds on our daily lives.

## SPECIFIC OBJECTIVES

## 1. Structure and Formulae

Students should be able to:
1.1. explain the occurrence of carbon compounds with straight chains, branched chains and rings;
1.2. explain the meaning of the term 'homologous series';
1.3. distinguish among empirical, molecular, and structural formulae;
1.4. determine formulae from experimental data;
1.5. write structural formulae;

EXPLANATORY NOTES
SUGGESTED PRACTICAL ACTIVITIES

Catenation, tetravalency, hybridisation, and resonance of carbon atoms to be used as basis.

Chemical and physical characteristics.

Structural formulae may be written in the following formats:


UNIT 2
MODULE 1: THE CHEMISTRY OF CARBON COMPOUNDS (cont'd)

## SPECIFIC OBJECTIVES

## Structure and Formulae cont'd

Students should be able to:
1.6. apply the IUPAC rules to named organic compounds;
1.7. define, structural isomerism;
1.8. explain stereoisomerism; and,

## EXPLANATORY NOTES

Condensed
$\mathrm{CH}_{3}\left(\mathrm{CH}_{2}\right)_{4} \mathrm{CH}_{3} ; \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{OH}$

Benzene can be represented by

and cyclohexane by


Examples should be given. Include chain, functional group, and positional isomers.

Geometrical (cis/trans) isomers resulting from restricted rotation about

double bond; optical isomerism due to asymmetry in molecules (confined to compounds with one identified chiral centre and drawing the two optical isomers). Optical isomers have an effect on plane polarised light. (No further treatment required).
1.9. determine the possible isomers from given molecular formulae.

SUGGESTED PRACTICAL ACTIVITIES

UNIT 2
MODULE 1: THE CHEMISTRY OF CARBON COMPOUNDS (cont'd)

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES
SUGGESTED PRACTICAL ACTIVITIES
2. Functional Group Analysis, Reactions and Mechanisms

Students should be able to:
2.1. identify homologous serie

Include general formulae.
of organic/carbon
compounds;
2.2. describe selected chemical reactions of alkanes;
2.3. explain the steps involved in the mechanism of free radical substitution;
2.4. describe selected chemical reactions of alkenes;

Halogenation, cracking, and combustion. Equations required.

Steps should include initiation, propagation, and termination. For example, methane and chlorine, homolytic fission.
Include movement of electrons to be indicated by curved arrows and fish hook notations.
2.4. reactions of alkenes;

$\mathrm{Br}_{2(\text { aq) })}$ (addition reaction)
Cold - $\mathrm{KMnO}_{4(a q)} / \mathrm{H}^{+}(a q)$ (oxidation and diol formation),
Hot - $\mathrm{KMnO}_{4(a q)} / \mathrm{H}^{+}(a q)$ (cleavage).
Hydrogen halides.
(Markovnikov rule for addition to asymmetric alkenes).
Include catalytic
hydrogenation of fats to
produce trans-fats which are harmful, for example, margarine.

Equations for reactions of alkenes with $\mathrm{KMnO}_{4(\mathrm{aq})}$ $/ \mathrm{H}^{+}(\mathrm{aq)}$ are not required.

## UNIT 2

MODULE 1: THE CHEMISTRY OF CARBON COMPOUNDS (cont'd)

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES

## Functional Group Analysis, Reactions and Mechanisms cont'd

Students should be able to:
2.5. explain the steps involved in the mechanism of selected chemical reactions of alkene functional group;
2.6. describe selected chemical reactions of alcohols;
2.7. describe selected reactions of halogenoalkanes;

Electrophilic addition of bromine and hydrogen bromide to alkenes.

Include movement of electrons. To be indicated by curved arrows and fish hook notation.

Reactions of alcohols to include primary, secondary and tertiary $\left(1^{0}, 2^{0}\right.$, and $3^{0}$ respectively) with $\mathrm{KMnO}_{4(\text { (aq })} / \mathrm{H}^{+}{ }_{(a q)}$ purple to colourless with primary and secondary; no reaction with tertiary;
$\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7(\text { aq })} / \mathrm{H}^{+}{ }_{(\text {aq })}$ orange to green with primary and secondary; no reaction with tertiary; Carboxylic acid (R-COOH); conc. sulfuric acid. A solution of $\mathrm{I}_{2}$ and NaOH (iodoform test).
Equations for reactions of alcohols with carboxylic acid and conc. sulfuric acid only are required.

Limited to hydrolysis of primary and tertiary halogenoalkanes using $\mathrm{NaOH}_{(\text {(aq) })}$ ( No elimination reaction required).

SUGGESTED PRACTICAL ACTIVITIES

Perform experiments examining the fragrances produced when different esters are formed.

UNIT 2
MODULE 1: THE CHEMISTRY OF CARBON COMPOUNDS (cont'd)

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES

## Functional Group Analysis, Reactions and Mechanisms cont'd

Students should be able to:
2.8. explain the steps involved in the mechanism of selected reactions of halogenoalkanes;

Nucleophilic substitution of halogenoalkanes with hydroxyl ions.

Include movement of electrons to be indicated by curved arrows and fish hook notation. Illustrate bond breaking or bond making as occurring in either a stepwise or concerted manner.
2.9. describe selected chemical reactions of carbonyl compounds


$$
\begin{array}{lr}
\mathrm{NaCN} / \mathrm{HCl}_{(\text {aq) }) ;} 2,4-\text { DNP } \\
\left(\mathrm{Brady}^{\prime} \mathrm{Reagent),}\right. & \text { Tollens' } \\
\text { reagent/Fehling's } & \text { solution; } \\
\mathrm{KMnO}_{4(\mathrm{aq})} / \mathrm{H}^{+}(\mathrm{aq}) ; & \text { LiAlH }_{4} ; \\
\mathrm{H}_{2} / \mathrm{Pt} . &
\end{array}
$$

No equations required
2.10. explain the steps involved in mechanisms of selected chemical reactions of carbonyl compounds;

Nucleophilic addition. Include reaction of carbonyl compounds with hydrogen cyanide as an example. Include movement of electrons to be indicated by curved arrows and fish hook notation.
2.11. describe selected chemical reaction of carboxylic acids ( $\mathrm{R}-\mathrm{COOH}$ );
$\mathrm{NaOH}, \mathrm{NaHCO}_{3}$, metals, alcohols $\mathrm{PCl}_{5} / \mathrm{PCl}_{3}$ or $\mathrm{SOCl}_{2}$. Equations are not required for reaction with $\mathrm{PCl}_{5}, \mathrm{PCl}_{3}$ or $\mathrm{SOCl}_{2}$.
2.12. describe selected chemical reactions of esters;

Acid and base hydrolysis. Simple soap production Include saponification, using vegetable oil and transesterification biodiesel NaOH . production.

SUGGESTED PRACTICAL ACTIVITIES

## UNIT 2

MODULE 1: THE CHEMISTRY OF CARBON COMPOUNDS (cont'd)

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES
SUGGESTED PRACTICAL ACTIVITIES

## Functional Group Analysis, Reactions and Mechanisms cont'd

Students should be able to:

### 2.13. perform suitable laboratory tests for functional groups in carbon compounds referred to above;

2.14. describe the chemical
$\begin{array}{llr}\text { 2.15. } & \text { describe selected chemical } \\ \text { reactions of } & \text { benzene } \\ \text { methylbenzene } & \text { and } \\ & \text { nitrobenzene; }\end{array}$
2.16. explain the steps involved in the mechanism of selected chemical reactions of benzene;
2.17. describe selected chemical reactions of phenol;
2.18. describe the formation of an azo compound; and,
2.19. state uses of azo compounds.
reaction of primary amines ( $\mathrm{RNH}_{2}$ ) with dilute acid;
$\mathrm{PCl}_{3}$ should be used in a fume cupboard. Theoretical considerations are satisfactory for $\mathrm{SOCl}_{2}$.

UNIT 2
MODULE 1: THE CHEMISTRY OF CARBON COMPOUNDS (cont'd)

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES
SUGGESTED PRACTICAL ACTIVITIES

## 3. Acidic and Basic Character of Organic Compounds

Students should be able to:
3.1. explain the difference in acidity of alcohols, phenols and carboxylic acids;
3.2. explain differences in basic character of aliphatic amines, amides, and aromatic amines; and,
3.3. explain the acid-base properties of amino acids.

Include chlorosubstituted acids. Reference should be made to the relationship between acid strength, pH and pK .

Inductive and conjugative effects.
pH and pK .
Inductive and conjugative effects.

Formation of zwitterions.

## 4. Macromolecules

Students should be able to:
4.1. describe the characteristics of addition polymerisation;

Examples to include polyethene; polyvinyl chloride; and, polytetrafluoroethene.

Include monomers for each polymer respectively.

Terylene; nylon 6.6, (specific monomers are required); proteins, starch, (representative structure required).

UNIT 2
MODULE 1: THE CHEMISTRY OF CARBON COMPOUNDS (cont'd)

## SPECIFIC OBJECTIVES

## Macromolecules cont'd

Students should be able to:
4.3. predict types of polymer formed from given monomers;
4.4. deduce the repeat unit of a polymer;
4.5. identify proteins as naturally occurring macromolecules;
4.6. identify carbohydrates as naturally occurring macromolecules; and,

EXPLANATORY NOTES
SUGGESTED PRACTICAL ACTIVITIES

Treat amino acids as monomeric molecules.

Generalised monomer and linkage required.

Include the following: cellulose, starch, and pectin. Treat simple sugars as monomeric materials.

Generalised monomer and linkage required.
4.7. illustrate the connection between carbohydrates and their monomers.

## Suggested Teaching and Learning Activities

To facilitate students' attainment of the objectives of this Module, teachers are advised to engage students in the teaching and learning activities listed below.

1. Arrange for students to view videos and interactive materials on industrial carbon chemistry. This should be followed by class or group discussion.
2. Arrange site visits, wherever possible, to industrial institutions, for example, (refineries, breweries, forensic laboratories, petrochemical plants).
3. Ask students to create a product which is marketable using simple organic reactions.
4. Ask students to construct and use models in sections dealing with structures.
5. Discuss articles in relevant reference journals and periodicals, for example, Chemistry in Education.

## UNIT 2 MODULE 1: THE CHEMISTRY OF CARBON COMPOUNDS (cont'd)

## RESOURCES

Teachers and students may find reference to the following resource materials useful. The latest editions are recommended.

Brown, T. and Le May, H. Chemistry, The Central Science. New Jersey: PrenticeHall, Incorporated, 2014.

Cann, P. and Hughes, P. Chemistry, International AS and A Level. London: Hodder Education, 2015.

Conoley, C. and Hills, P.
Chemistry, $3^{\text {rd }}$ Edition. London: HarperCollins, 2008.

Clugston, M. and Flemming, R.
Advanced Chemistry. London: Oxford University Press, 2000.

Hill, G., and Holman, J. Chemistry in Context. London: Nelson Thorne Limited, 2001.

Lister, T., Renshaw, J. Understanding Chemistry for Advanced Level. Cheltenham: Trans-Atlantic Publications, 2000.

Norris, R., Barrett, L., Maynard-Alleyne, A. and Murray, J.

Ramsden, E.
CAPE ${ }^{\circledR}$ Chemistry Study Guide. Cheltenham: Nelson Thorne Limited, 2012.

A-Level Chemistry. Cheltenham: Nelson Thorne Limited, 2000.

## WEBSITES

## www.Chemsoc.org

www.Chemguide.co.uk
www.creative-chemistry.org.uk
www.a-levelchemistry.co.uk

UNIT 2
MODULE 2: ANALYTICAL METHODS AND SEPARATION TECHNIQUES

## GENERAL OBJECTIVES

On completion of this Module, students should:

1. appreciate that all measurements have some degree of uncertainty that is related to both the measuring device and the skills of the operator;
2. understand the basic theoretical principles of selected methods of analysis and separation techniques;
3. demonstrate a knowledge of the basic equipment and operational procedures, as well as carry out experiments associated with selected methods of analysis and separation procedures;
4. use experimental data to quantify substances or elucidate partial structure; and,
5. demonstrate an awareness of the wide applications of various methods of analysis and separation techniques in industry and in medicine by citing examples.

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES

## 1. Uncertainty in Measurements

Students should be able to:
1.1. apply appropriate concepts to the analysis of scientific data;

Definitions required for: mean, standard deviation, precision, accuracy, calibration curves, and standards.

Calculation of the mean and standard deviation from data provided will be required.
$S=\frac{\sqrt{\sum(\mathrm{x}-\overline{\mathrm{x}})^{2}}}{\mathrm{n}-1}$
1.2. carry out experiments to assess the degree of uncertainty in measurements associated with the use of certain common pieces of laboratory equipment; and,

Perform experiments which should include the use of pipettes, burettes, volumetric flasks, thermometers, top-loading balances and analytical balances.

## SUGGESTED PRACTICAL ACTIVITIES

UNIT 2
MODULE 2: ANALYTICAL METHODS AND SEPARATION TECHNIQUES (cont'd)

## SPECIFIC OBJECTIVES

## Uncertainty in Measurements cont'd

Students should be able to:
1.3. select appropriate pieces of equipment to make measurements, depending upon the degree of accuracy required.

EXPLANATORY NOTES

Selection of equipment limited to those in Specific Objective 1.2 above.

## 2. Titrimetric (Volumetric) Methods of Analysis

## Students should be able to:

2.1. explain the basic principles upon which titrimetric analyses are based;
2.2. discuss the criteria used in selecting primary standards;
$\mathrm{NaHCO}_{3}, \quad \mathrm{Na}_{2} \mathrm{CO}_{3}, \quad \mathrm{KIO}_{3}$, $(\mathrm{COOH})_{2}$ and its salts as primary standards.

Definition of primary standards required.
2.3. use data obtained from potentiometric, thermometric, and conductrometric methods for titration which do not require the use of indicators;
2.4. perform experiments based on titrimetric analyses;
2.5. perform calculations based on data obtained from titrimetric analyses; and,
2.6. cite examples of the use of titrimetric analysis in the quantification of various substances.

Refer to Unit 1, Module 2, Specific Objective 3.6.

Refer to vinegar, household cleaners, vitamin C tablets, aspirin, antacids.

SUGGESTED PRACTICAL ACTIVITIES

UNIT 2
MODULE 2: ANALYTICAL METHODS AND SEPARATION TECHNIQUES (cont'd)

## SPECIFIC OBJECTIVES

## EXPLANATORY NOTES

## 3. Gravimetric Methods of Analysis

Students should be able to:
3.1. explain the principles upon which gravimetric analyses are based;
3.2. describe the functions of the various pieces of basic equipment used in gravimetric analyses;
3.3. perform experiments based on gravimetric determinations;

Limited to a discussion on precipitation and volatilisation methods.

Limited to suction flasks, suction funnels, silica crucibles, sinter glass crucibles, ovens, and furnaces.
3.4. perform calculations based on data obtained from gravimetric analyses; and,
3.5. cite examples of the use of gravimetric analysis in quality control.

Use data from actual experiments carried out or from the literature.

Examples include determination of $\mathrm{SO}_{2}$ in the air, in wine or fruit drink; determination of the amount of elements such as phosphorous in fertilizers; the chloride ion presence in water supply.

SUGGESTED PRACTICAL ACTIVITIES

Limited to
experiments involving volatilisation methods, such as, moisture content of soils; water of crystallisation.

## 4. Spectroscopic Methods of Analysis

Students should be able to:
4.1. explain the nature of Calculations using the electromagnetic radiation;
equation:
$\mathrm{E}=\mathrm{hv}=\mathrm{hc} / \lambda$ are required.

UNIT 2
MODULE 2: ANALYTICAL METHODS AND SEPARATION TECHNIQUES (cont'd)

## SPECIFIC OBJECTIVES

EXPLANATORY NOTES

SUGGESTED PRACTICAL ACTIVITIES

## Spectroscopic Methods of Analysis cont'd

Students should be able to:

## 4.2. state the approximate wavelength ranges of the X-ray, UV/VIS, IR and radiofrequency regions of the electromagnetic spectrum; and,

4.3. recall that the energy levels in atoms and molecules are quantised.

Relative energies and dangers associated with exposure to high energy wavelengths.

Insert actual ranges for $x$ ray, UV, VIS, IR and RF.
Refer to Unit 1, Module 1.

## 5. Ultra Violet-Visible (UV/VIS) Spectroscopy

Students should be able to:
5.1. explain the origin of absorption in UV/VIS spectroscopy;
5.2. explain why some species will absorb light in the UV/VIS region whereas others will not;
5.3. describe the basic steps involved in analysing samples by UV/VIS spectroscopy;
5.4. use Beer-Lambert's Law to calculate the concentration of a given species in solution; and,
5.5. list examples of the use of UV/VIS spectroscopy in the quantitation of substances.

Simple treatment based on Unit 1, Module 1.
Consideration of sigma ( $\sigma$ ), pi $(\pi)$, anti-bonding ( $\sigma^{*}$, $\pi^{*}$ ) and non-bonding ( $n$ ) orbitals.

Brief mention should be made of the use of complexing reagents to form coloured compounds. Sensitivity and detection limits.

Use of standards and calibration curves.

Iron tablets; glucose and urea in blood; cyanide in water.

UNIT 2
MODULE 2: ANALYTICAL METHODS AND SEPARATION TECHNIQUES (cont'd)

## SPECIFIC OBJECTIVES

## 6. Infrared Spectroscopy

Students should be able to:
6.1. explain the origin of absorption in IR spectroscopy;
6.2. describe the basic steps involved in analysing samples by IR spectroscopy;
6.3. comment on the limitations associated with the use of IR spectroscopy;
6.4. deduce the functional groups present in organic compounds from IR spectra; and,
6.5. cite examples of the use of IR spectroscopy in the monitoring of air pollutants.

## 7. Mass Spectroscopy

Students should be able to:
7.1. explain the basic principles of mass spectrometry;

EXPLANATORY NOTES

Include reference to preparation of solids, use of $\mathrm{KBr}, \mathrm{NaCl}$ discs/pellets, and nujol mulls.

The usefulness of IR data when used in conjunction with other data (for example, mass spectrometry).

Groups to be identified include: $-\mathrm{OH},-\mathrm{NH}_{2}$,

$-\mathrm{CO}_{2} \mathrm{H} \quad-\mathrm{CONH}_{2}$
Use should be made of IR spectral data. (See Table 6 in the data booklet in Appendix 2).
$\mathrm{CO}_{2}, \mathrm{SO}_{2}$.

Include block diagram.
(Simple schematic diagram of the process).

SUGGESTED PRACTICAL ACTIVITIES

Use IR spectra along with absorption tables to deduce the presence or absence of particular bonds or functional groups.

UNIT 2
MODULE 2: ANALYTICAL METHODS AND SEPARATION TECHNIQUES (cont'd)

## SPECIFIC OBJECTIVES

## Mass Spectroscopy cont'd

Students should be able to:
7.3. use mass spectral data to:
(a) determine relative isotopic masses; and relative isotopic abundance; and,
(b) distinguish between molecules of similar relative molecular mass.

## EXPLANATORY NOTES

Use should be made of mass spectral sheets. Refer to the ( $M+1$ and $M+2$ peaks).

## 8. Chromatographic Methods of Separation

Students should be able to:
8.1. explain the theoretical principles upon which chromatographic methods are based;

These should be explained in terms of adsorption and partition between the mobile and stationary phases; refer to paper, column, thin layer, and gas-liquid chromatography.

SUGGESTED PRACTICAL ACTIVITIES

UNIT 2
MODULE 2: ANALYTICAL METHODS AND SEPARATION TECHNIQUES (cont'd)

## SPECIFIC OBJECTIVES

## Chromatographic Methods of Separation cont'd

Students should be able to:
8.4. name examples of commonly used stationary phases;
8.5. separate the components of mixtures; and,

EXPLANATORY NOTES

Include reference to cellulose, silica gel, alumina.

Suitable mixtures which could be used include amino acids, plant pigments, food colouring.

Perform simple experiments to separate the components of mixtures using paper and column chromatographic techniques.
8.6. cite the wide applications of chromatographic methods of separation.

## 9. Phase Separations

Students should be able to:
9.1. discuss the chemical principles upon which simple distillation and fractional distillation are based;

Refer to pesticide analysis, forensic testing, purification of natural products.

State Raoult's Law. Perform simple experiments Definition of ideal and non- to illustrate the concept of ideal mixture required.

The interpretation of boiling point composition curves of both ideal and non-ideal mixtures is required. A qualitative treatment of boiling point composition curves of azeotropic mixtures is required.

## SUGGESTED PRACTICAL ACTIVITIES

UNIT 2
MODULE 2: ANALYTICAL METHODS AND SEPARATION TECHNIQUES (cont'd)

## SPECIFIC OBJECTIVES

## Phase Separations cont'd

Students should be able to:
9.3. discuss the chemical
principles and use of steam
distillation;
9.4. discuss the principles upon which solvent extraction is based;
9.5. select appropriate methods of separation, given the physical and chemical properties of the components of a mixture;
9.6. perform distillation experiments; and,

EXPLANATORY NOTES

Laboratory work on the extraction of essential oils from plant materials. Purification of nitrobenzene and phenylamine. Simple calculations are required.

A discussion on partition coefficient and simple calculations are required.

These should include (but not be limited to) a comparison of the efficiency of separation of ethanol in beer or rum by simple and fractional distillation.

Include petroleum, rum, and the fragrance industries.

SUGGESTED PRACTICAL ACTIVITIES

Conduct simple separation experiments based on solute partitioning between two immiscible solvents. These could include the separation of an acid/base mixture.
9.7. cite examples of the applications of the distillation methods used in various industries.

## UNIT 2 <br> MODULE 2: ANALYTICAL METHODS AND SEPARATION TECHNIQUES (cont'd)

## Suggested Teaching and Learning Activities

To facilitate students' attainment of the objectives of this Module, teachers are advised to engage students in the teaching and learning activities listed below.

1. Establish contact with industries and institutions in your locality which make use of the spectroscopic methods or separation techniques in their operations in order to:
(a) facilitate site visits with your students;
(b) make use of personnel from such industries and institutions to come in and give lectures and demonstrations; and
(c) access appropriate written, audio or visual material available at such industries and institutions.
2. Prepare handouts compiled from data in literature (readily available) for use by students. This is especially important where spectral data are concerned. Students should be able to work with spectral data sheets.

## RESOURCES

Teachers and students may find reference to the following resource materials useful. The latest editions are recommended.

| Cann, P. and Hughes, P. | Chemistry, International AS and A Level. London: <br> Hodder Education, 2015. |
| :--- | :--- |
| Conoley, C. and Hills, P. | Chemistry, 3 3rd Edition. London: HarperCollins, 2008. |
| Clugston, M. and Flemming, R. | Advanced Chemistry. London: Oxford University Press, <br> 2000. |
| Ramsden, E. | A-Level Chemistry. Cheltenham: Nelson Thorne <br> Limited, 2000. |

UNIT 2
MODULE 2: ANALYTICAL METHODS AND SEPARATION TECHNIQUES (cont'd)

## WEBSITES

www.Chemsoc.org
www.Chemguide.co.uk
www.creative-chemistry.org.uk
www.a-levelchemistry.co.uk

## UNIT 2

MODULE 3: INDUSTRY AND THE ENVIRONMENT

## GENERAL OBJECTIVES

On completion of this Module, students should:

1. appreciate that chemical principles can be applied to industry;
2. appreciate that using Green Chemistry principles in industry helps to produce a sustainable world that supports a healthy economy;
3. understand the sources of pollution and strategies which assist in reduction of pollutants;
4. understand that there are physical and chemical changes occurring in the environment and assess their impact on it;
5. recognise the influence of industrial processes on social and economic life; and,
6. appreciate the impact of man's activities on the environment.

SPECIFIC OBJECTIVES
EXPLANATORY NOTES

## 1. Locating and Operating Industrial plants: Benefits and Risk

Students should be able to:
1.1. discuss factors which Factors should be related to influence the location of an the industrial plant. industrial plant; and,

Environmental impact assessment should be carried out.
1.2. discuss general safety requirements for industry; and,
1.3. assess the processes of Brief explanation of the select industries using Green principles of Green Chemistry principles. Chemistry.
Selected industries ammonia and crude oil.

## SUGGESTED PRACTICAL ACTIVITIES

UNIT 2
MODULE 3: INDUSTRY AND THE ENVIRONMENT (cont'd)

## SPECIFIC OBJECTIVES

## 2. Aluminium

Students should be able to:
2.1. describe the processes involved in the production of aluminium from its ores;
2.2. explain the uses of aluminium in relation to its physical and chemical properties; and,
2.3. assess the impact of the aluminium industry on the environment.
3. Crude Oil

Students should be able to:
3.1. explain the method used in the separation of the components of crude oil;
3.2. discuss the uses of the components of crude oil as fuels and as raw materials for the petro-chemical industry; and,
3.3. assess the impact of the petroleum industry on the environment.

EXPLANATORY NOTES

## SUGGESTED

 ACTIVITIESInclude purification of the ore. Technical details are not required. Include equations.

High energy consumption in the production of aluminium.

Refer to Specific Objective 1.3.

Refer to Module 2 Specific Objective 9.6. Include fractional distillation, catalytic cracking, and reforming techniques.

Refer to Specific Objective 1.3.

UNIT 2
MODULE 3: INDUSTRY AND THE ENVIRONMENT (cont'd)

## SPECIFIC OBJECTIVES

## 4. Ammonia

Students should be able to:
4.1. outline the steps in the manufacture of ammonia from its elements, by the Haber process;
4.2. discuss the uses of ammonia; and,
4.3. assess the impact of the ammonia industry on the environment.
5. Ethanol

Students should be able to:
5.1. explain the process of fermentation and distillation in the manufacture of alcoholic beverages;
5.2. discuss the uses of ethanol;
5.3. discuss the social and economic impact of ethanol production and consumption; and,
5.4. assess the impact of the ethanol industry on the environment.

EXPLANATORY NOTES

SUGGESTED ACTIVITIES

PRACTICAL
Activies

Include the production of the starting materials and manufacturing conditions. Include equations.

Apply the principles of chemical equilibrium and kinetics.

Including in agriculture and chemical industry.

Refer to Specific Objective 1.3.

Include equations and sources of carbohydrates.

Include fuel, pharmaceutical industry.

Include physiological changes.

Refer to Specific Objective 1.3.

UNIT 2

## MODULE 3: INDUSTRY AND THE ENVIRONMENT (cont'd)

## SPECIFIC OBJECTIVES

## 6. Chlorine

Students should be able to:
6.1. describe the chemical processes involved in the electrolysis of brine using the diaphragm cell;
6.2. discuss the economic advantages of chlorine production by the diaphragm cell method;
6.3. discuss the industrial importance of the halogens and their compounds; and,

## 6.4. assess the impact of the chlor-alkali industry on the environment

## 7. Sulfuric Acid

Students should be able to:
7.1. describe the Contac process for the manufacture of sulfuric acid;
7.2. discuss the industria importance of compounds of sulfur; and,

Bleaches, PVC, halogenated hydrocarbons, solvents, aerosols, refrigerants, anaesthetics.

Refer to Specific Objective 1.3

Include the principles of chemical equilibrium and kinetics. Include equations.
$\mathrm{SO}_{2}$ in food preservation and $\mathrm{H}_{2} \mathrm{SO}_{4}$ manufacture.
7.3. assess the impact of the Refer to Specific Objective 1.3. sulfuric acid industry.

## UNIT 2 <br> MODULE 3: INDUSTRY AND THE ENVIRONMENT (cont'd)

## SPECIFIC OBJECTIVES

## 8. Water

Students should be able to:
8.1. describe the importance of the water cycle;
8.2. discuss methods of water purification;
8.3. discuss the importance of dissolved oxygen to aquatic life;
8.4. discuss the sources of water pollution;

Definition of the terms persistent and bioaccumulation in relation to all forms of pollution. Refer to Specific Objectives 9.2 and 10.3.

Sources of water pollution may include nitrates, phosphates, heavy metals (lead and mercury), cyanides, trace metals, pesticides, herbicides, petroleum residue, suspended particles.

EXPLANATORY NOTES Include desalination.






## SUGGESTED PRACTICAL ACTIVITIES

 suspended particles.8.5. perform experiments to test for some pollutants specified in Specific Objective 8.4; and,
8.6. assess the impact of the pollutants in Specific Objective 8.4 and Specific Objective 9.2 on the aquatic environment.

UNIT 2
MODULE 3: INDUSTRY AND THE ENVIRONMENT (cont'd)

## SPECIFIC OBJECTIVES

## 9. The Atmosphere

Students should be able to:
9.1. explain how the concentration of ozone in the atmosphere is maintained;
9.2. discuss the
environmental significance of CFCs in the ozone layer;
9.3. discuss the effects of ozone on human life;
9.4. explain the importance of maintaining the balance of carbon dioxide concentration in the atmosphere;
9.5. explain the following terms: green-house effect, global warming, photochemical smog;
9.6. discuss the effects of the products of combustion of hydrocarbon-based fuels;

EXPLANATORY NOTES

Photodissociation.

Include reference to free radical reactions in the upper atmosphere.

Refer to both stratosphere and troposphere, and Specific Objective 9.2.

Equilibrium concepts, carbon cycle and reforestation.

Brief analysis on the impact of climate change.

Include re-radiation of energy from the infrared region.

Consider $\mathrm{CO}, \mathrm{SO}_{2}$, oxides of nitrogen, lead compounds and volatile organic compounds.

Primary and secondary pollutants, for example, NO and $\mathrm{NO}_{2}$, respectively.

Nitrogen cycle and acid rain.

SUGGESTED PRACTICAL ACTIVITIES

## UNIT 2

## MODULE 3: INDUSTRY AND THE ENVIRONMENT (cont'd)

## SPECIFIC OBJECTIVES

9.8. discuss methods of control and prevention of atmospheric pollution.

EXPLANATORY NOTES

For prevention include alternative and cleaner fuels improved technology and mass transit. For control include sequestering, filters, washers and scrubbers.
10. Solid Waste

Students should be able to:
10.1. distinguish among the terms reduce, reuse, recycle.
10.2. describe the processes involved in waste reduction; and,
10.3. assess the impact of solid wastes on the terrestrial environment.

Consider reusing and recycling of glass, paper, plastic, steel and aluminium. (A simple process approach is adequate).

Include reference to iron, glass, plastic, paper, lead, biodegradable and nonbiodegradable materials, proper and improper disposal techniquesdumps and sanitary landfills.

## SUGGESTED PRACTICAL ACTIVITIES

Visit a landfill, bauxite mines, alumina plant, mineral quarry.

## UNIT 2 <br> MODULE 3: INDUSTRY AND THE ENVIRONMENT (cont'd)

## Suggested Teaching and Learning Activities

To facilitate students' attainment of the objectives of this Module, teachers are advised to engage students in the teaching and learning activities listed below.

1. Arrange visits to industrial plants and view video materials. This should be followed by class/group discussions.
2. Invite personnel from environmental groups (Non-Governmental Organisations, Community Based Organisations) and the Environmental Management Authority (EMA) in their territories to be guest lecturers on environmental issues.
3. Have students collect newspaper articles and journal articles, and conduct web-based searches on issues relating to industry and environment (include in the study the benefits of Green Chemistry). Engage in discussion on the main issues of the article collected.
4. Conduct class discussions and debates on the social and economic issues that arise from degradation of the environment.
5. Have students do a survey on community awareness about environmental issues.

## RESOURCES

Teachers and students may find reference to the following resource materials useful. The latest editions are recommended.

| Cann, P. and Hughes, P. | Chemistry, International AS and A Level. London: Hodder Education, 2015. |
| :---: | :---: |
| Conoley, C. and Hills, P. | Chemistry, 3 ${ }^{\text {rd }}$ Edition. London: HarperCollins, 2008. |
| Clugston, M. and Flemming, R. | Advanced Chemistry. London: Oxford University Press, 2000. |
| Hill, G., and Holman, J. | Chemistry in Context. London: Nelson Thorne Limited, 2001. |
| Norris, R., Barrett, L., Maynard-Alleyne, A. and Murray, J. | CAPE ${ }^{\circledR}$ Chemistry Study Guide: Cheltenham: Nelson Thorne Limited, 2012. |
| Ramsden, E. | A-Level Chemistry. Cheltenham: Nelson Thorne Limited, 2000. |

UNIT 2
MODULE 3: INDUSTRY AND THE ENVIRONMENT (cont'd)

## WEBSITES

www.Chemsoc.org
www.Chemguide.co.uk
www.creative-chemistry.org.uk
www.a-levelchemistry.co.uk
www.asc.org
www.epa.gov
www.rsc.org

## - OUTLINE OF ASSESSMENT

## EXTERNAL ASSESSMENT

(80\%)

## Paper 01 <br> Forty-five multiple-choice items, 15 from each Module. <br> 40\%

(1 hour 30 minutes)

## Paper 02

(2 hours 30 minutes)

Paper 032
Three compulsory structured essay questions, one from each
40\% Module. Each question is worth 30 marks.

For private candidates only (2 hours)

Three questions, one from each Module, as follows:
(a) a practical-based question to be executed by the candidate;
(b) a question based on data analysis; and
(c) a data analysis/a planning and design exercise.

## SCHOOL-BASED ASSESSMENT

The School-Based Assessment will consist of selected practical laboratory exercises and one research project aligned to any Unit of the CAPE ${ }^{\circledR}$ Sciences (Biology, Chemistry or Physics).

## MODERATION OF SCHOOL-BASED ASSESSMENT

The reliability (consistency) of the marks awarded by teachers on the School-Based Assessment is an important characteristic of high quality assessment. To assist in this process, the Council undertakes on-site moderation of the School-Based Assessment during Term 2/3. This is conducted by visiting External Moderators who will visit the centre.

Teachers are required to present to the Moderator ALL Assessment Sheets (Record of Marks), ALL lab books, Mark Schemes and the project or evidence of the project. This is also required when marks are being transferred from one Unit/subject to another. Candidates marks are to be recorded on the School-Based Assessment Record Sheets which are available online via the CXC ${ }^{\text {®'s }}$ s website www.cxc.org. All candidates' marks are to be submitted electronically using the SBA data capture module of the Online Registration System (ORS). Teachers are NOT required to submit to CXC ${ }^{\circledR}$ samples of candidates' work, unless specifically requested to do so by the Council.

The Moderator will re-mark the skills and projects for a sample of five candidates using the guidelines below. This is only applicable if the candidates selected in the sample are not using transferred marks for the projects.

1. Candidates' total marks on the SBA are arranged in descending order (highest to lowest).
2. The sample comprises the work of the candidates scoring the:
(a) highest Total Mark;
(b) middle Total Mark;
(c) lowest Total Mark;
(d) mark midway between the highest and middle Total Mark; and,
(e) mark midway between the middle and lowest Total Mark.
3. The Moderator will also re-mark the laboratory practical activities for the other skills (ORR, AI and PD) that are recorded in the lab books for the five candidates in the sample.
4. The Moderator will re-mark the skills for ALL the candidates where the total number of candidates is five or less than five.
5. The Moderator will provide teachers with feedback. Please note that Candidates' marks may be adjusted as a result of the moderation exercise.

The Moderators are required to submit the moderated marks (Moderation of SBA Sample Form), the Moderation Feedback Report and the External Moderator Report to the Local Registrar by 30 June of the year of the examination.

A copy of the Assessment Sheets and all candidates' work must be retained by the school for three months after the examination results are published by CXC ${ }^{\circledR}$.

## ASSESSMENT DETAILS

Each Unit of the syllabus is assessed as outlined below.

## External Assessment by Written Papers (80\% of Total Assessment)

1. Paper 01 consists of 45 multiple-choice items. There will be a combined question paper and answer booklet for Paper 02.
2. S.I. Units will be used on all examination papers.
3. The use of silent, non-programmable calculators will be allowed in the examination. Candidates are responsible for providing their own calculators.
4. Data not specifically required to be recalled, defined or stated will be made available for this examination.

## Paper 01 (1 hour 30 minutes - 40\% of Total Assessment)

## 1. Composition of the Paper

This paper will consist of 45 multiple-choice items, 15 from each Module.
All questions are compulsory and knowledge of the entire Unit is expected. The paper will assess the candidate's knowledge across the breadth of the Unit.

## 2. Mark Allocation

The paper will be worth 45 marks, which will be weighted to 90 marks.

## 3. Question Type

Questions may be presented using diagrams, data, graphs, prose or other stimulus material.

## Paper 02 ( 2 hours 30 minutes - 40\% of Total Assessment)

## 1. Composition of Paper

This paper will consist of three questions, one from each module. All questions are compulsory.

Questions on this paper test all three skills KC, UK and XS.

Knowledge of the entire Unit is expected.

## 2. Mark Allocation

The paper will be worth 90 marks, 30 marks per question and distributed across the question sub-parts.

## 3. Question Type

Questions will be presented in structured essay format. The questions will test the skills of $K C$, $U K$ and $X S$. Answers are to be written in the question booklet.

## School-Based Assessment (20\%)

School-Based Assessment is an integral part of student assessment in the course covered by this syllabus. It is intended to assist students in acquiring certain knowledge, skills and attitudes that are associated with the subject. Students are encouraged to work in groups.

During the course of study for the subject, students obtain marks for the competence they develop and demonstrate in undertaking their School-Based Assessment assignments. These marks contribute to the final marks and grades that are awarded to students for their performance in the examination.

School-Based Assessment provides an opportunity to individualise a part of the curriculum to meet the needs of students. It facilitates feedback to the student at various stages of the experience. This helps to build the self- confidence of students as they proceed with their studies. School-Based Assessment also facilitates the development of the critical skills and abilities emphasised by this CAPE ${ }^{\circledR}$ subject and enhances the validity of the examination on which candidate performance is reported.

School-Based Assessment, therefore, makes a significant and unique contribution to both the development of relevant skills and the testing and rewarding of students for the development of those skills.

The Caribbean Examinations Council seeks to ensure that the School-Based Assessment scores that contribute to the overall scores of candidates are valid and reliable estimates of accomplishment. The guidelines provided in this syllabus are intended to assist in doing so.

## Award of Marks

The following skills will be assessed through the laboratory practical activities:

1. Analysis and Interpretation;
2. Manipulation and Measurement;
3. Observation, Recording and Reporting; and,
4. Planning and Designing.

The candidates are also required to do an investigative project in any one Unit of the CAPE ${ }^{\oplus}$ Sciences. The table below shows how the marks are allocated for each Unit.

Table 1
School-Based Assessment Skills

| Skill | Unit 1 | Unit 2 |
| :--- | :---: | :---: |
| Observation, Recording and Reporting | 12 | 12 |
| Manipulation and Measurement | 12 | 12 |
| Analysis and Interpretation* | 12 | 12 |
| Planning and Designing* | 12 | 12 |
| TOTAL | 48 marks | 48 marks |

*Includes an investigative project

## Teachers are required to provide criteria which clearly indicate how they award marks.

Please note that candidates will be required to do one investigative project in any Unit of any of the CAPE ${ }^{\circledR}$ Sciences (Biology, Chemistry or Physics) in the first sitting, and can use that mark for the other Units of the Sciences. So for example, a candidate may do the investigative project in Unit 2 Physics in the first sitting, and then (transfer) use the AI and PD marks for Unit 1 Physics, Units 1 and 2 Chemistry and Units 1 and 2 Biology.

Each Module will carry a maximum of 16 marks.
Each candidate's total School-Based Assessment mark for any Unit should be divided in three and allocated to each Module equally.

Fractional marks should not be awarded. Wherever the Unit mark is not divisible by three, then
(a) when the remainder mark is 1, it should be allocated to Module 1; and,
(b) when the remainder is 2 , one of the marks should be allocated to Module 2 and the other mark to Module 3.

Appropriate practical exercises for assessing any skill may be selected from any Module in the relevant Unit.

## - INVESTIGATIVE PROJECT

## Objectives of the Investigative Project

The Investigative Project must focus on a challenge to be addressed within the environment or society. On completion of the Investigative Project students should:

1. Appreciate the use of the scientific method for discovery of new knowledge and to the solution of problems;
2. Communicate accurately and effectively the purpose and results of research;
3. Apply experimental skills and theory to the solution of problems; and,
4. Synthesise information based on data collected.

Students are encouraged to work collaboratively. Where collaborative work is done, group sizes must not exceed six (6) persons per group. The teacher is expected to use the group mark for the project and add it to the marks for the other skills for each individual candidate within the group.

## CRITERIA FOR ASSESSING INVESTIGATIVE SKILLS

A. PLANNING AND DESIGN

| - HYPOTHESIS |  | 1 |  |
| :---: | :---: | :---: | :---: |
| - AIM |  | 1 |  |
| - MATERIALS AND APPARATUS |  | 1 |  |
| VARIABLES STATED <br> - Controlled <br> - Manipulated <br> - Responding | 1 1 1 | 3 |  |

CAPE ${ }^{\circ}$


| B. | ANALYSIS AND INTERPRETATION |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
|  | - RESULTS | 2 | 2 |  |
|  | - Complete set of results from quantities mentioned in method. |  |  |  |
|  | - DISCUSSION |  | 4 |  |
|  | - Complete set of calculations or statement of observations or trends. | 2 |  |  |
|  | - Interpretations of calculated values, observations or trends linked to data in results. | 2 |  |  |
|  | - LIMITATIONS AND SOURCES OF ERROR <br> Limitation stated <br> Source of error stated |  | 2 |  |
|  |  | 11 |  |  |
|  |  |  |  |  |
|  | - REFLECTIONS <br> - Relevance of experiment to real life. <br> - Impact of knowledge gained from experiment. <br> - How can experiment be changed and improved. |  | 3 |  |
|  |  | 111 |  |  |
|  |  |  |  |  |
|  |  |  |  |  |
|  | - CONCLUSION <br> - Clearly stated and related to Aim in PD. | 1 | 1 |  |
|  |  |  |  |  |
|  | TOTAL |  |  | (12) |

## SCHOOL-BASED ASSESSMENT - GENERAL GUIDELINES FOR TEACHERS

1. Each candidate is required to keep a laboratory workbook which is to be marked by the teacher. Teachers are also expected to assess candidates as they perform practical exercises in which Manipulation and Measurement skills are required.
2. A maximum of two skills may be assessed by any one experiment.
3. The mark awarded for each skill assessed by practical exercises should be the average of at LEAST TWO separate assessments. The average mark for AI and PD must include the mark from the investigative project. In each Unit, total marks awarded at the end of each Module will be 0 to 16 .
4. The maximum mark for any skill will be 12. The mark awarded for each skill assessed by practical exercises should be the average of at LEAST TWO separate assessments. In each Unit, total marks awarded at the end of each Module will be 0 to 16.
5. Candidates who do not fulfil the requirements for the School-Based Assessment will be considered absent from the whole examination.

Candidates' laboratory books should contain all practical work undertaken during the course of study. Those exercises which are selected for use for the School-Based Assessment should be clearly identified. The skill(s) tested in these selected practical exercises, the marks assigned and the scale used must be placed next to the relevant exercises.

## - REGULATIONS FOR PRIVATE CANDIDATES

1. Candidates who are registered privately will be required to sit Papers 01,02 and 032 . Detailed information on Papers 01, 02 and 032 is given on page 64 of this syllabus.
2. Paper 032 will constitute 20 per cent of the overall assessment of the candidates' performance on the Unit.

## - REGULATIONS FOR RESIT CANDIDATES

1. Candidates may reuse any moderated SBA score within a two-year period. In order to assist candidates in making decisions about whether or not to reuse a moderated SBA score, the Council will continue to indicate on the preliminary results if a candidate's moderated SBA score is less than 50 per cent in a particular Unit.
2. Candidates reusing SBA scores should register as "Resit candidates" and must provide the previous candidate number when registering.
3. Resit candidates must complete Papers 01 and 02 of the examination for the year in which they register.

## - ASSESSMENT GRID

The Assessment Grid for each Unit contains marks assigned to papers and to Modules and percentage contribution of each paper to total scores.

| Paper | Module 1 | Module 2 | Module 3 | Paper Total <br> (Weighted Total) | \% <br> Weighting <br> of Papers |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Paper 01 | 15 <br> $(30)$ | 15 <br> $(30)$ | 15 <br> $(30)$ | 45 <br> $(90)$ | 40 |
| Paper 02 | 30 | 30 | 30 | 90 | 40 |
| Paper 031 | 16 <br> $(15)$ | 16 <br> $(15)$ | 16 <br> Paper 032 | 15 | 15 |

CAPE ${ }^{\circledR}$

## - GLOSSARY OF EXAMINATIONS TERMS

## KEY TO ABBREVIATIONS

KC - Knowledge and Comprehension
UK - Use of Knowledge
XS - Experimental Skills

## WORD

Annotate

Apply

Assess

## Calculate

Cite

Classify

Comment

## Compare

## Construct

## DEFINITION

requires a brief note to be added to a label.
requires the use of knowledge or principles to solve problems.
requires the inclusion of reasons for the importance of particular structures, relationships or processes.
requires a numerical answer for which working must be shown.
requires a quotation or a KC reference to the subject.
requires a division into groups according to observable and stated characteristics.
requires a statement of an opinion or a view, with reason supporting.
requires a statement about similarities and differences. UK

UK

## NOTES

Simple phrase or a few words only; KC

Make references or conclusions; UK

Compare the advantages and disadvantages or the merits and demerits of a particular structure, relationship or process; UK

Steps should be shown; units must be included; UK

An example of a significance of each similarity and the difference stated may be required for comparisons which are other than structural; UK

Such representations should normally bear a title, appropriate headings and legend; UK

Deduce

Define

Demonstrate

Derive

Describe

Design

## Determine

DEFINITION
practical investigations, or building of models or the drawing of scale diagrams.
the making of logical connections between pieces of information.
requires a formal statement or an equivalent paraphrase, such as defining equation with symbols identified.
show; direct attention to.
implies a deduction, determination or extraction of some relationship, formula or result from data by a logical set of steps.
requires a statement in words (using diagrams where appropriate) of the main points of the topic. This can also imply the inclusion of reference to (visual) observations associated with particular phenomena or experiments. The amount of description intended should be interpreted from the context.
includes planning and presentation with appropriate practical detail.
implies that the quantity concerned should not be measured directly but should be obtained by calculator or derivation.
implies an expansion or elaboration of an idea or argument with supporting evidence.

## NOTES

UK

This should include the defining equation or formula where relevant; UK

KC

UK

Description may be words, drawings or diagrams or an appropriate combination. Drawings or diagrams should be annotated to show appropriate detail where necessary; KC

UK/XS

Where hypotheses are stated or when tests are to be conducted, possible outcomes should be clearly shown or the way in which data will be analysed and presented; XS

KC/UK

## Develop

WORD

Differentiate or Distinguish (between or among)

DEFINITION
requires a statement and brief explanation of the differences between or among items.
requires a critical account of the points involved in the topic.
requires a line representation of the item, showing accurate relationship between the parts.
implies a reasoned order of magnitude statement or calculation of the quantity concerned, using such simplifying assumptions as may be necessary about points of principle and about the values of quantities not otherwise included.
requires the weighing of evidence and judgements based on stated criteria.
implies that a definition or a description should be given, together with some relevant comment on the significance or context of the term or situation concerned. The amount of supplementary comment intended should be interpreted from the context.

Find

Formulate

Identify
requires the location of $a$ UK feature or the determination as from a graph.
implies the articulation of a UK hypothesis.
requires the naming of specific KC components or features. Implies a clear demonstration, using appropriate examples or diagrams.

## WORD

| Label | implies the inclusion of names to identify structures or parts as indicated by pointers. | KC/XS |
| :---: | :---: | :---: |
| List | requires a number of points with no elaboration. Where a given number of points is specified, this should not be exceeded. | KC |
| Measure | implies that the quantity concerned can be directly obtained from a suitable measuring instrument. | XS |
| Name | requires only the identification of the item. | No additional information is required; KC |
| Note | implies the writing down of observations. | XS |
| Observe | implies the direction of attention to details which characterize reaction or change taking place and examination of scientific notations. | Observation may involve all the senses and extensions of them but would normally exclude the sense of taste; XS |
| Outline | requires basic steps only. |  |
| Plan | implies preparation to conduct an exercise or operation. | XS |
| Predict | implies the use of information to arrive at a likely conclusion or the suggestion of possible outcomes. | UK |
| Record | implies an accurate account or description of the full range of observations made during a given procedure. | This includes the values for any variable being investigated; where appropriate, record data may be depicted in graphs, histograms or tables; XS |
| Relate | implies the demonstration of connections between sets of facts or data. | UK |


| WORD | DEFINITION | NOTES |
| :---: | :---: | :---: |
| Show | see Demonstrate. |  |
| Sketch | in relation to graphs, implies that the shape or position of the curve need only be qualitatively correct and, depending on the context, some quantitative aspects may need to be included. In relation to diagrams, implies that a simple, freehand drawing is acceptable, provided proportions and important details are made clear. | KC/UK/XS |
| State | implies a concise statement with little or no supporting argument. | KC |
| Suggest | could imply either that there is no unique response or the need to apply general knowledge to a novel situation. | No correct or incorrect solution is presumed but suggestions must be acceptable within the limits of scientific knowledge; UK |
| Test | implies the determination of a result by following set procedures. | XS |
| Use | implies the need to recall and apply in order to come to a conclusion. | UK |

## Western Zone Office

9 August 2018

# CARIBBEAN ADVANCED PROFICIENCY EXAMINATION (CAPE®) 

## CHEMISTRY DATA BOOKLET

# CARIBBEAN EXAMINATIONS COUNCIL CARIBBEAN ADVANCED PROFICIENCY EXAMINATION ${ }^{\circledR}$ 

## CHEMISTRY

DATA BOOKLET

DO NOT TAKE AWAY FROM THE EXAMINATION ROOM

TABLE 1: SOME IMPORTANT CONSTANTS

| The Avogadro constant | L | $=6.02 \times 10^{23} \mathrm{~mol}^{-1}$ |
| :---: | :---: | :---: |
| Electronic charge | $e$ | $=-1.60 \times 10^{-19} \mathrm{C}$ |
| The Faraday constant | F | $=9.65 \times 10^{4} \mathrm{C} \mathrm{mol}^{-1}$ |
| Ionic product of water | $K_{w}$ | $=1.00 \times 10^{-14} \mathrm{~mol}^{2} \mathrm{dm}^{-6}($ at 298 K$)$ |
| Molar gas constant | $R$ | $=8.31 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}$ |
| Molar volume of gas | $V_{m}$ | $=22.4 \mathrm{dm}^{3} \mathrm{~mol}^{-1}$ at s.t.p <br> $=24 \mathrm{dm}^{3} \mathrm{~mol}^{-1}$ under room conditions |
| The Planck constant | $h$ | $=6.63 \times 10^{-34} \mathrm{~J} \mathrm{~s}$ |
| Rest mass of electron, ${ }_{-1}^{0} \mathrm{e}$ | $m_{e}$ | $=9.11 \times 10^{-31} \mathrm{~kg}$ |
| Rest mass of neutron, ${ }_{0}^{1} \mathrm{n}$ | $m_{n}$ | $=1.67 \times 10^{-27} \mathrm{~kg}$ |
| Rest mass of proton, ${ }_{1}^{1} \mathrm{H}$ | $m_{p}$ | $=1.67 \times 10^{-27} \mathrm{~kg}$ |
| Specific heat capacity of water | $C_{\text {water }}$ | $=4.18 \mathrm{~kJ} \mathrm{~kg}^{-1} \mathrm{~K}^{-1}$ |
| Speed of light in a vacuum | c | $=3.00 \times 10^{8} \mathrm{~m} \mathrm{~s}^{-1}$ |

TABLE 2: IONISATION ENERGIES OF SELECTED ELEMENTS

| Element | Proton Number | Ionisation Energies ( $\mathrm{kJ} \mathrm{mol}^{-1}$ ) |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | First | Second | Third | Fourth |
| H | 1 | 1310 | - | - | - |
| He | 2 | 2370 | 5250 | - | - |
| Li | 3 | 519 | 7300 | 11800 | - |
| Be | 4 | 900 | 1760 | 14800 | 21000 |
| B | 5 | 799 | 2420 | 3660 | 25000 |
| C | 6 | 1090 | 2350 | 4610 | 6220 |
| N | 7 | 1400 | 2860 | 4590 | 7480 |
| O | 8 | 1310 | 3390 | 5320 | 7450 |
| F | 9 | 1680 | 3370 | 6040 | 8410 |
| Ne | 10 | 2080 | 3950 | 6150 | 9290 |
| Na | 11 | 494 | 4560 | 6940 | 9540 |
| Mg | 12 | 736 | 1450 | 7740 | 10500 |
| Al | 13 | 577 | 1820 | 2740 | 11600 |
| Si | 14 | 786 | 1580 | 3230 | 4360 |
| P | 15 | 1060 | 1900 | 2920 | 4960 |
| S | 16 | 1000 | 2260 | 3390 | 4540 |
| Cl | 17 | 1260 | 2300 | 3850 | 5150 |
| Ar | 18 | 1520 | 2660 | 3950 | 5770 |
| K | 19 | 418 | 3070 | 4600 | 5860 |
| Ca | 20 | 590 | 1150 | 4940 | 6480 |
| Sc | 21 | 632 | 1240 | 2390 | 7110 |
| Ti | 22 | 661 | 1310 | 2720 | 4170 |
| V | 23 | 648 | 1370 | 2870 | 4600 |
| Cr | 24 | 653 | 1590 | 2990 | 4770 |
| Mn | 25 | 716 | 1510 | 3250 | 5190 |
| Fe | 26 | 762 | 1560 | 2960 | 5400 |
| Co | 27 | 757 | 1640 | 3230 | 5100 |
| Ni | 28 | 736 | 1750 | 3390 | 5400 |
| Cu | 29 | 745 | 1960 | 3350 | 5690 |
| Zn | 30 | 908 | 1730 | 3828 | 5980 |
| Ge | 32 | 762 | 1540 | 3300 | 4390 |
| Br | 35 | 1140 | 2080 | 3460 | 4850 |
| Sr | 38 | 548 | 1060 | 4120 | 5440 |
| Sn | 50 | 707 | 1410 | 2940 | 3930 |
| I | 53 | 1010 | 1840 | 2040 | 4030 |
| Ba | 56 | 502 | 966 | 3390 | - |
| Pb | 82 | 716 | 1450 | 3080 | 4080 |

TABLE 3: ATOMIC AND IONIC RADII OF SELECTED ELEMENTS


TABLE 4: SELECTED BOND ENERGIES

|  | Bond | Energy/kJ mol ${ }^{-1}$ |
| :---: | :---: | :---: |
| (a) | Diatomic molecules |  |
|  | H-H | 436 |
|  | D-D | 442 |
|  | $\mathrm{N} \equiv \mathrm{N}$ | 994 |
|  | $\mathrm{O}=\mathrm{O}$ | 496 |
|  | F-F | 158 |
|  | $\mathrm{Cl}-\mathrm{Cl}$ | 244 |
|  | $\mathrm{Br}-\mathrm{Br}$ | 193 |
|  | I-I | 151 |
|  | H-F | 562 |
|  | $\mathrm{H}-\mathrm{Cl}$ | 431 |
|  | $\mathrm{H}-\mathrm{Br}$ | 366 |
|  | H-I | 299 |
| (b) | Polyatomic molecules |  |
|  | $\mathrm{C}-\mathrm{C}$ | 350 |
|  | $\mathrm{C}=\mathrm{C}$ | 610 |
|  | $\mathrm{C} \equiv \mathrm{C}$ | 840 |
|  | $\mathrm{C}-\mathrm{C}$ ( (benzene) | 520 |
|  | $\mathrm{C}-\mathrm{H}$ | 410 |
|  | $\mathrm{C}-\mathrm{Cl}$ | 340 |
|  | $\mathrm{C}-\mathrm{Br}$ | 280 |
|  | $\mathrm{C}-\mathrm{I}$ | 240 |
|  | $\mathrm{C}-\mathrm{N}$ | 305 |
|  | $\mathrm{C}=\mathrm{N}$ | 610 |
|  | $\mathrm{C} \equiv \mathrm{N}$ | 890 |
|  | $\mathrm{C}-\mathrm{O}$ | 360 |
|  | $\mathrm{C}=\mathrm{O}$ | 740 |
|  | $\mathrm{N}-\mathrm{H}$ | 390 |
|  | $\mathrm{N}-\mathrm{N}$ | 160 |
|  | $\mathrm{N}=\mathrm{N}$ | 410 |
|  | $\mathrm{O}-\mathrm{H}$ | 460 |
|  | $\mathrm{O}-\mathrm{O}$ | 150 |
|  | $\mathrm{Si}-\mathrm{Cl}$ | 359 |
|  | Si-H | 320 |
|  | Si-O | 444 |
|  | $\mathrm{Si}-\mathrm{Si}$ | 222 |
|  | $\mathrm{S}-\mathrm{Cl}$ | 250 |
|  | $\mathrm{S}-\mathrm{H}$ | 347 |
|  | S-S | 264 |

TABLE 5: STANDARD ELECTRODE AND REDOX POTENTIALS

| Electrode Reaction |  | $\mathbf{E}^{\ominus} / \text { volts }$ |
| :---: | :---: | :---: |
| $\mathrm{Ag}^{+}+\mathrm{e}^{-} \rightleftharpoons$ | Ag | +0.80 |
| $\mathrm{Al}^{3+}+3 \mathrm{e}^{-} \rightleftharpoons$ | Al | -1.66 |
| $\mathrm{Ba}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Ba | -2.90 |
| $\mathrm{Br}_{2}+2 \mathrm{e}^{-} \rightleftharpoons$ | $2 \mathrm{Br}^{-}$ | +1.07 |
| $\mathrm{Ca}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Ca | -2.87 |
| $\mathrm{Cl}_{2}+2 \mathrm{e}^{-} \rightleftharpoons$ | $2 \mathrm{Cl}^{-}$ | +1.36 |
| $2 \mathrm{HOCl}+2 \mathrm{H}^{+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Cl}_{2}+2 \mathrm{H}_{2} \mathrm{O}$ | +1.64 |
| $\mathrm{Co}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Co | -0.28 |
| $\mathrm{Co}^{3+}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Co}^{2+}$ | +1.82 |
| $\left[\mathrm{Co}\left(\mathrm{NH}_{3}\right)_{6}{ }^{2+}+2 \mathrm{e}^{-} \rightleftharpoons\right.$ | $\mathrm{Co}+6 \mathrm{NH}_{3}$ | -0.43 |
| $\mathrm{Cr}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Cr | -0.91 |
| $\mathrm{Cr}^{3+}+3 \mathrm{e}^{-} \rightleftharpoons$ | Cr | -0.74 |
| $\mathrm{Cr}^{3+}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Cr}^{2+}$ | -0.41 |
| $\mathrm{Cr}_{2} \mathrm{O}^{2-}{ }_{7}+14 \mathrm{H}^{+}+6 \mathrm{e}^{-} \rightleftharpoons$ | $2 \mathrm{Cr}^{3+}+7 \mathrm{H}_{2} \mathrm{O}$ | +1.33 |
| $\mathrm{Cu}^{+}+\mathrm{e}^{-} \rightleftharpoons$ | Cu | +0.52 |
| $\mathrm{Cu}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Cu | +0.34 |
| $\mathrm{Cu}^{2+}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Cu}^{+}$ | +0.15 |
| $\left[\mathrm{Cu}\left(\mathrm{NH}_{3}\right)_{4}\right]^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Cu}+4 \mathrm{NH}_{3}$ | -0.05 |
| $\mathrm{F}_{2}+2 \mathrm{e}^{-} \rightleftharpoons$ | $2 \mathrm{~F}^{-}$ | +2.87 |
| $\mathrm{Fe}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Fe | -0.44 |
| $\mathrm{Fe}^{3+}+3 \mathrm{e}^{-} \rightleftharpoons$ | Fe | -0.04 |
| $\mathrm{Fe}^{3+}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Fe}^{2+}$ | +0.77 |
| $[\mathrm{Fe}(\mathrm{CN})]^{3-}+\mathrm{e}^{-} \rightleftharpoons$ | $[\mathrm{Fe}(\mathrm{CN})]_{6}{ }^{4}$ | +0.36 |
| $\mathrm{Fe}(\mathrm{OH})_{3}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Fe}(\mathrm{OH})_{2}+\mathrm{OH}^{-}$ | -0.56 |
| $2 \mathrm{H}^{+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{H}_{2}$ | 0.00 |
| $\mathrm{I}_{2}+2 \mathrm{e}^{-} \rightleftharpoons$ | $2 \mathrm{I}^{-}$ | +0.54 |
| $\mathrm{K}^{+}+\mathrm{e}^{-} \rightleftharpoons$ | K | -2.92 |
| $\mathrm{Li}^{+}+\mathrm{e}^{-} \rightleftharpoons$ | Li | -3.04 |
| $\mathrm{Mg}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Mg | -2.38 |
| $\mathrm{Mn}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Mn | -1.18 |

TABLE 5: STANDARD ELECTRODE AND REDOX POTENTIALS (CONT'D)

| Electrode Reaction |  | $\mathrm{E}^{\boldsymbol{\theta}} / \mathrm{volts}$ |
| :---: | :---: | :---: |
| $\mathrm{Mn}^{3+}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Mn}^{2+}$ | +1.49 |
| $\mathrm{MnO}_{2}+4 \mathrm{H}^{+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Mn}^{2+}+2 \mathrm{H}_{2} \mathrm{O}$ | +1.23 |
| $\mathrm{MnO}_{4}^{-}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{MnO}^{2-}$ | +0.56 |
| $\mathrm{MnO}_{4}^{-}+4 \mathrm{H}^{+}+3 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{MnO}_{2}+2 \mathrm{H}_{2} \mathrm{O}$ | +1.67 |
| $\mathrm{MnO}_{4}^{-}+8 \mathrm{H}^{+}+5 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Mn}^{2+}+4 \mathrm{H}_{2} \mathrm{O}$ | +1.52 |
| $\mathrm{NO}_{3}^{-}+2 \mathrm{H}^{+}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{NO}_{2}+\mathrm{H}_{2} \mathrm{O}$ | +0.81 |
| $\mathrm{NO}_{3}^{-}+3 \mathrm{H}^{+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{HNO}_{2}+\mathrm{H}_{2} \mathrm{O}$ | +0.94 |
| $\mathrm{NO}_{3}^{-}+10 \mathrm{H}^{+}+8 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{NH}_{4}^{-}+3 \mathrm{H}_{2} \mathrm{O}$ | +0.87 |
| $\mathrm{Na}^{+}+\mathrm{e}^{-} \rightleftharpoons$ | Na | -2.71 |
| $\mathrm{Ni}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Ni | -0.25 |
| $\left[\mathrm{Ni}\left(\mathrm{NH}_{3}\right)_{6}\right]^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Ni}+6 \mathrm{NH}_{3}$ | -0.51 |
| $\mathrm{H}_{2} \mathrm{O}_{2}+2 \mathrm{H}^{+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $2 \mathrm{H}_{2} \mathrm{O}$ | +1.77 |
| $\mathrm{O}_{2}+4 \mathrm{H}^{+}+4 \mathrm{e}^{-} \rightleftharpoons$ | $2 \mathrm{H}_{2} \mathrm{O}$ | +1.23 |
| $\mathrm{O}_{2}+2 \mathrm{H}_{2} \mathrm{O}+4 \mathrm{e}^{-} \rightleftharpoons$ | $4 \mathrm{OH}^{-}$ | +0.40 |
| $\mathrm{O}_{2}+2 \mathrm{H}^{+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{H}_{2} \mathrm{O}_{2}$ | +0.68 |
| $2 \mathrm{H}_{2} \mathrm{O}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{H}_{2}+2 \mathrm{OH}^{-}$ | -0.83 |
| $\mathrm{Pb}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Pb | -0.13 |
| $\mathrm{Pb}^{4+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Pb}^{2+}$ | +1.69 |
| $\mathrm{PbO}_{2}+4 \mathrm{H}^{+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Pb}^{2+}+2 \mathrm{H}_{2} \mathrm{O}$ | +1.47 |
| $\mathrm{SO}_{4}^{2-}+4 \mathrm{H}^{+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{SO}_{2}+2 \mathrm{H}_{2} \mathrm{O}$ | +0.17 |
| $\mathrm{S}_{2} \mathrm{O}^{2-}+2 \mathrm{e}^{-} \rightleftharpoons$ | $2 \mathrm{SO}_{4}^{2-}$ | +2.01 |
| $\mathrm{S}_{4} \mathrm{O}^{2-}{ }_{6}+2 \mathrm{e}^{-} \rightleftharpoons$ | $2 \mathrm{~S}_{2} \mathrm{O}^{2-}$ | +0.09 |
| $\mathrm{Sn}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Sn | -0.14 |
| $\mathrm{Sn}^{4+}+2 \mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{Sn}^{2+}$ | +0.15 |
| $\mathrm{V}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | V | -1.20 |
| $\mathrm{V}^{3+}+\mathrm{e}^{-} \rightleftharpoons$ |  | -0.26 |
| $\mathrm{VO}^{2+}+2 \mathrm{H}^{+}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{V}^{3+}+\mathrm{H}_{2} \mathrm{O}$ | +0.34 |
| $\mathrm{VO}_{2}^{+}+2 \mathrm{H}^{+}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{VO}^{2+}+\mathrm{H}_{2} \mathrm{O}$ | +1.00 |
| $\mathrm{VO}_{3}^{-}+4 \mathrm{H}^{+}+\mathrm{e}^{-} \rightleftharpoons$ | $\mathrm{VO}^{2+}+2 \mathrm{H}_{2} \mathrm{O}$ | +1.00 |
| $\mathrm{Zn}^{2+}+2 \mathrm{e}^{-} \rightleftharpoons$ | Zn | -0.76 |

TABLE 6: SELECTED INFRARED ABSORPTION SPECTROSCOPIC DATA

| Bond |  | Characteristic Absorption Ranges* <br> (Wave Number, $\mathbf{c m}^{-1}$ ) |
| :--- | :--- | :--- |
| $\mathrm{O}-\mathrm{H}$ | 'free' | 3580 to 3670 |
| $\mathrm{~N}-\mathrm{H}$ | primary amines |  |
| $\mathrm{O}-\mathrm{H}$ | 'hydrogen-bonded' in alcohols, phenols | 3350 to 3500 |
| $\mathrm{C}-\mathrm{H}$ | alkanes, alkenes, arenes | 3230 to 3550 |
| $\mathrm{O}-\mathrm{H}$ | 'hydrogen-bonded' in acids | 2840 to 3095 |
| $\mathrm{C} \equiv \mathrm{N}$ |  | 2500 to 3300 |
| $\mathrm{C} \equiv \mathrm{C}$ |  | 2200 to 2280 |
| $\mathrm{C}=\mathrm{O}$ | aldehydes, ketones, acids, esters | 2070 to 2250 |
| $\mathrm{C}=\mathrm{C}$ |  | 1680 to 1750 |
| $\mathrm{C}-\mathrm{O}$ | alcohols, ethers, esters | 1610 to 1680 |
| $\mathrm{C}-\mathrm{Cl}$ |  | 1000 to 1300 |

[^0]The Periodic Table

| Group |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| I | II |  |  |  |  |  |  |  |  |  |  | III | IV | v | VI | VII | 0 |
| $\begin{gathered} 1.0 \\ \mathrm{H} \\ 1 \end{gathered}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | 4.0 He 2 |
| $\begin{gathered} 6.9 \\ \mathrm{Li} \\ 3 \end{gathered}$ | 9.0 Be 4 |  |  |  |  |  |  |  |  |  |  | $\begin{gathered} 10.8 \\ \text { B } \\ 5 \end{gathered}$ | $\begin{gathered} 12.0 \\ \text { C } \\ 6 \end{gathered}$ | $\begin{gathered} 14.0 \\ \mathrm{~N} \\ 7 \end{gathered}$ | $\begin{gathered} 16.0 \\ \mathrm{O} \\ 8 \end{gathered}$ | $\begin{gathered} 19.0 \\ \text { F } \\ 9 \end{gathered}$ | $\begin{aligned} & 0.2 \\ & \mathrm{Ne} \\ & 10 \end{aligned}$ |
| $\begin{gathered} 23.0 \\ \mathrm{Na} \\ 11 \end{gathered}$ | $\begin{aligned} & 24.3 \\ & \mathrm{Mg} \\ & 12 \end{aligned}$ |  |  |  |  |  |  |  |  |  |  | $\begin{gathered} 27.0 \\ \mathrm{Al} \\ 13 \end{gathered}$ | $\begin{gathered} 28.1 \\ \mathrm{Si} \\ 14 \end{gathered}$ | $\begin{gathered} 31.0 \\ \mathrm{p} \\ 15 \end{gathered}$ | $\begin{gathered} 32.1 \\ \text { S } \\ 16 \end{gathered}$ | $\begin{gathered} 35.5 \\ \mathrm{Cl} \\ 17 \end{gathered}$ | $\begin{gathered} 39.9 \\ \text { Ar } \\ 18 \end{gathered}$ |
| $\begin{gathered} 39.1 \\ K \\ 19 \end{gathered}$ | $\begin{gathered} 40.1 \\ \mathrm{Ca} \\ 20 \end{gathered}$ | $\begin{gathered} 45.0 \\ \mathrm{Sc} \\ 21 \end{gathered}$ | $\begin{gathered} 47.9 \\ \mathrm{Ti}_{2} \\ 2 \end{gathered}$ | $\begin{gathered} 50.9 \\ \mathrm{~V} \\ 23 \end{gathered}$ | $\begin{gathered} 520 \\ \mathrm{Cr} \\ 24 \end{gathered}$ | $\begin{aligned} & 54.9 \\ & \mathrm{Mn} \\ & 25 \end{aligned}$ | $\begin{gathered} 55.8 \\ \mathrm{Fe} \\ 26 \end{gathered}$ | $\begin{gathered} 58.9 \\ \text { Co } \\ 27 \end{gathered}$ | $\begin{gathered} 58.7 \\ \mathrm{Ni} \\ 28 \end{gathered}$ | $\begin{gathered} 63.5 \\ \mathrm{Cu} \\ 29 \end{gathered}$ | $\begin{aligned} & 65.4 \\ & \mathrm{Zn} \\ & 30 \end{aligned}$ | $\begin{gathered} 69.7 \\ \mathrm{Ga} \\ 31 \end{gathered}$ | $\begin{aligned} & 72.6 \\ & \mathrm{Ge} \\ & 32 \end{aligned}$ | $\begin{gathered} 74.9 \\ \text { As } \\ 33 \end{gathered}$ | $\begin{aligned} & 79.0 \\ & \text { Se } \\ & 34 \end{aligned}$ | $\begin{gathered} 79.9 \\ \mathrm{Br} \\ 35 \end{gathered}$ | $\begin{gathered} 83.8 \\ \mathrm{Kr} \\ 36 \end{gathered}$ |
| $\begin{gathered} 85.5 \\ \mathrm{Rb} \\ 37 \end{gathered}$ | $\begin{gathered} 87.6 \\ \mathrm{Sr} \\ 38 \end{gathered}$ | $\begin{gathered} 88.9 \\ \text { Y } \\ 39 \end{gathered}$ | $\begin{gathered} 91.2 \\ \mathrm{Zr} \\ 40 \end{gathered}$ | $\begin{gathered} 92.9 \\ \mathrm{Nb} \\ 41 \end{gathered}$ | $\begin{gathered} 95.9 \\ \text { Mo } \\ 42 \end{gathered}$ | $\begin{aligned} & \mathrm{Tc} \\ & 43 \end{aligned}$ | $\begin{aligned} & 101 \\ & \text { Ru } \\ & 44 \end{aligned}$ | $\begin{aligned} & 103 \\ & \text { Rh } \\ & 45 \end{aligned}$ | $\begin{aligned} & 106 \\ & \text { Pd } \\ & 46 \end{aligned}$ | $\begin{aligned} & 108 \\ & \mathrm{Ag} \\ & 47 \end{aligned}$ | $\begin{aligned} & 112 \\ & \mathrm{Cd} \\ & 48 \end{aligned}$ | $\begin{aligned} & 115 \\ & \text { In } \\ & 49 \end{aligned}$ | $\begin{aligned} & 119 \\ & \mathrm{Sn} \\ & 50 \end{aligned}$ | $\begin{aligned} & 122 \\ & \mathrm{Sb} \\ & 51 \end{aligned}$ | $\begin{aligned} & 128 \\ & \mathrm{Te} \\ & 52 \\ & 52 \end{aligned}$ | $\begin{gathered} 127 \\ \text { I } \\ 53 \end{gathered}$ | $\begin{aligned} & 131 \\ & \text { Xe } \\ & 54 \end{aligned}$ |
| $\begin{aligned} & 133 \\ & \mathrm{Cs} \\ & 55 \end{aligned}$ | $\begin{aligned} & 137 \\ & \mathrm{Ba} \\ & 56 \end{aligned}$ | $\begin{aligned} & \mathrm{La}^{*} \\ & \text { to } \\ & \mathrm{Lu} \end{aligned}$ | $\begin{aligned} & 178 \\ & \mathrm{Hf} \\ & 72 \end{aligned}$ | $\begin{aligned} & 181 \\ & \text { Ta } \\ & 73 \end{aligned}$ | $\begin{aligned} & 184 \\ & \mathrm{~W} \\ & 74 \end{aligned}$ | $\begin{aligned} & 186 \\ & \text { Re } \\ & 75 \end{aligned}$ | $\begin{aligned} & 190 \\ & \text { Os } \\ & 76 \end{aligned}$ | $\begin{aligned} & 192 \\ & \text { Ir } \\ & 77 \end{aligned}$ | $\begin{aligned} & 195 \\ & \mathrm{Pt} \\ & 78 \end{aligned}$ | $\begin{aligned} & 197 \\ & \mathrm{Au} \\ & 79 \end{aligned}$ | $\begin{aligned} & 201 \\ & \mathrm{Hg} \\ & 80 \end{aligned}$ | $\begin{gathered} 204 \\ \mathrm{Tl} \\ \mathrm{T1} \\ 81 \end{gathered}$ | $\begin{aligned} & 207 \\ & \mathrm{~Pb} \\ & 82 \end{aligned}$ | $\begin{gathered} 209 \\ \text { Bi } \\ 83 \end{gathered}$ | $\begin{aligned} & - \\ & \text { Po } \\ & 84 \end{aligned}$ | $\begin{aligned} & -\overline{A t} \\ & \begin{array}{l} \text { A } \end{array} \end{aligned}$ | $\overline{\mathrm{Rn}}$ |
| $\begin{aligned} & - \\ & \mathrm{Fr} \\ & 87 \end{aligned}$ | $\begin{aligned} & \mathrm{Ra} \\ & 88 \end{aligned}$ | $\begin{gathered} \mathrm{Ac}^{* *} \\ \text { to } \\ \mathrm{Lr} \end{gathered}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |


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# CARIBBEAN EXAMINATIONS COUNCIL 

## Caribbean Advanced Proficiency Examination® CAPE ${ }^{\circledR}$

## CHEMISTRY

## Specimen Papers and Mark Schemes/Keys

## Specimen Papers:

Unit 1 Paper 01
Unit 1 Paper 02
Unit 1 Paper 32
Unit 2 Paper 01
Unit 2 Paper 02
Unit 2 Paper 32

Mark Schemes and Key:
Unit 1 Paper 01
Unit 1 Paper 02
Unit 1 Paper 32
Unit 2 Paper 01
Unit 2 Paper 02
Unit 2 Paper 32

# CARIBBEAN EXAMINATIONS COUNCIL <br> CARIBBEAN ADVANCED PROFICIENCY EXAMINATION <br> <br> CHEMISTRY 

 <br> <br> CHEMISTRY}

## SPECIMEN 2017

## TABLE OF SPECIFICATIONS

## Unit 1 - Paper 02

| Module | Question | Specific Objective | Content | Cognitive Level Marks |  |  | Total |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | KC | UK | XS |  |
| 1 | 1 | $\begin{gathered} 3.1,3.2,3.3,3.5,3.6, \\ 3.8,3.9,6.1,6.2,6.4, \\ 6.6,6.7,6.16 \end{gathered}$ | The Mole Concept Energetics | 10 | 15 | 5 | 30 |
| 2 | 2 | $\begin{gathered} 3.1,3.2,3.3,4.1,4.2, \\ 6.3,6.4,6.5 \end{gathered}$ | Redox Equilibria Principles of Chemical Equilibria, Buffer Solutions | 10 | 15 | 5 | 30 |
| 3 | 3 | 2.1, 3.4, 5.2, 5.6, 5.10 | Transition Elements Group II Group IV | 10 | 15 | 5 | 30 |
| Total |  |  |  | 30 | 45 | 15 | 90 |

Unit 1 - Paper 032 (Alternative to SBA)

| Module | Question |  | Content | Cognitive Level Marks |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | KC | UK | XS |  |
| 1 | 1 | Practical | Module 1-3.8, 3.9 | 0 | 5 | 10 | 15 |
| 2 | 2 | Data Analysis | Module 2-1.4-1.6 | 0 | 5 | 10 | 15 |
| 3 | 3 | Planning \& Design | Module 2-6.26 | 0 | 5 | 10 | 15 |
| Total |  |  |  | 0 | 15 | 30 | 45 |

# CARIBBEAN EXAMINATIONS COUNCIL <br> CARIBBEAN ADVANCED PROFICIENCY EXAMINATION <br> <br> CHEMISTRY 

 <br> <br> CHEMISTRY}

## SPECIMEN 2017

## TABLE OF SPECIFICATIONS

Unit 2-Paper 02

| Module | Question | Specific <br> Objective | Content | Cognitive Level Marks |  |  | Total |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | KC | UK | XS |  |
| 1 | 1 | $\begin{gathered} 1.8,2.1,2.2,2,3 \\ 2.6,2.7,2.13 \\ 2.11,2.14 \end{gathered}$ | Structures and Formulae Functional Group Analysis Alcohols, Acids, Ketones Haloalkanes | 10 | 15 | 5 | 30 |
| 2 | 2 | $\begin{aligned} & 7.1,7.2,8.1, \\ & 8.2,8.3,8.5 \end{aligned}$ | Chromatography <br> Mass Spectroscopy | 10 | 15 | 5 | 15 |
| 3 | 3 | $\begin{aligned} & 1.3,4.1,8.4, \\ & 8.5,9.6,9.7 \end{aligned}$ | Water and the Atmosphere Manufacture of Ammonia | 10 | 15 | 5 | 30 |
| Total |  |  |  | 30 | 45 | 15 | 90 |

## Unit 2 - Paper 032

Alternative to SBA

| Module | Question | Specific Objective | Content |  | itive Mark |  | Total |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | KC | UK | XS |  |
| 1 | 1 | Practical | Module 2 | 0 | 5 | 10 | 15 |
| 2 | 2 | Data Analysis | Module 2 | 0 | 5 | 10 | 15 |
| 3 | 3 | Planning \& Design | Module 1 | 0 | 5 | 10 | 15 |
| Total |  |  |  | 0 | 15 | 30 | 45 |

# CARIBBEAN EXAMINATIONS COUNCIL 

CARIBBEAN ADVANCED PROFICIENCY EXAMINATION ${ }^{\circledR}$ CHEMISTRY

Unit 1 - Paper 01

## 1 hour 30 minutes

## SPECIMEN PAPER

## READ THE FOLLOWING INSTRUCTIONS CAREFULLY.

1. This test consists of 45 items. You will have 1 hour and 30 minutes to answer them.
2. In addition to this test booklet, you should have an answer sheet.
3. Do not be concerned that the answer sheet provides spaces for more answers than there are items in this test.
4. Each item in this test has four suggested answers lettered (A), (B), (C), (D). Read each item you are about to answer and decide which choice is best.
5. On your answer sheet, find the number which corresponds to your item and shade the space having the same letter as the answer you have chosen. Look at the sample item below.

## Sample Item

Which of the following is the highest energy transition in an organic compound?
(A) $\quad \mathrm{n}$ to $\sigma^{*}$

Sample Answer
(B) $\quad \mathrm{n}$ to $\pi^{*}$
(C) $\sigma$ to $\sigma^{*}$
(D) $\quad \pi$ to $\pi^{*}$

The correct answer to this item is " $\sigma$ to $\sigma^{*}$ ", so (C) has been shaded.
6. If you want to change your answer, erase it completely before you fill in your new choice.
7. When you are told to begin, turn the page and work as quickly and as carefully as you can. If you cannot answer an item, go on to the next one. You may return to that item later.
8. You may do any rough work in this booklet.
9. Figures are not necessarily drawn to scale.
10. You may use a silent, non-programmable calculator to answer items.

1. When a Group I metal atom, Q , forms ion $\mathrm{Q}^{+}$, the
(A) positive charge on the nucleus increases
(B) number of protons increases
(C) number of occupied electron shells decreases by one
(D) radius of the particle increases
2. Radioactivity is hazardous because it
(A) can cause cancer and sterilization in human beings
(B) is accumulative and interacts with matter
(C) needs to be transported and disposed of properly
(D) has particulate as well as high energy emissions
3. In which of the following compounds will hydrogen bonding NOT be present?
(A) $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{I}$
(B) $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$
(C) $\mathrm{CH}_{3} \mathrm{NH}_{2}$
(D) $\quad \mathrm{H}_{2} \mathrm{SO}_{3}$
4. Which of the following species has a structure with a bond angle less than $109^{\circ}$ $28^{\prime}$ ?
(A) $\mathrm{NF}_{3}$
(B) $\mathrm{NH}_{4}^{+}$
(C) $\mathrm{CH}_{4}$
(D) $\quad \mathrm{BF}_{3}$
5. The molar volume of a gas at room temperature and pressure (r.t.p) is $24 \mathrm{dm}^{3}$. If 0.032 g of a gas occupies $48 \mathrm{~cm}^{3}$ at r.t.p., what is the molar mass?
(A) 0.016 g
(B) 2 g
(C) 16 g
(D) 64 g
6. Propane is a fuel used in lamps. The equation for the combustion of propane is
$\mathrm{C}_{3} \mathrm{H}_{8}+5 \mathrm{O}_{2} \rightarrow 3 \mathrm{CO}_{2}+4 \mathrm{H}_{2} \mathrm{O}$
What volume of $\mathrm{CO}_{2}$ is produced when 8.8 g of propane is burnt in oxygen at r.t.p.?
(The molar volume of gas at r.t.p $=24 \mathrm{dm}^{3}$ )
(A) $1.44 \mathrm{dm}^{3}$
(B) $4.8 \mathrm{dm}^{3}$
(C) $\quad 14.4 \mathrm{dm}^{3}$
(D) $72 \mathrm{dm}^{3}$
7. Which of the following are redox reactions?
I. $2 \mathrm{H}_{2} \mathrm{O}_{2}(\mathrm{aq}) \rightarrow 2 \mathrm{H}_{2} \mathrm{O}(l)+\mathrm{O}_{2}(\mathrm{~g})$
II. $\mathrm{Fe}_{2} \mathrm{O}_{3}(\mathrm{~s})+2 \mathrm{Al}(\mathrm{s}) \rightarrow \mathrm{Al}_{2} \mathrm{O}_{3}(\mathrm{~s})+2 \mathrm{Fe}(\mathrm{s})$
III. $\mathrm{BaCl}_{2}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{SO}_{4}(\mathrm{aq}) \rightarrow \mathrm{BaSO}_{4}(\mathrm{~s})+2 \mathrm{HCl}(\mathrm{aq})$
(A) I and II only
(B) I and III only
(C) II and III only
(D) I, II and III
8. In the reaction between potassium manganate (VII) and potassium iodide in acid solution, the following relevant half equations can be written:
$2 \mathrm{I}^{-} \rightarrow \mathrm{I}_{2}+2 \mathrm{e}-$
$\mathrm{MnO}_{4}{ }^{-}+8 \mathrm{H}^{+}+5 \mathrm{e}^{-} \rightarrow \mathrm{Mn}^{2+}+4 \mathrm{H}_{2} \mathrm{O}$
Which of the following is the overall balanced equation?
(A) $\quad \mathrm{MnO}_{4}^{-}+8 \mathrm{H}^{+}+2 \mathrm{I}^{-} \rightarrow \mathrm{Mn}^{2+}+4 \mathrm{H}_{2} \mathrm{O}+\mathrm{I}_{2}$
(B) $\quad \mathrm{MnO}_{4}^{-}+8 \mathrm{H}^{+}+2 \mathrm{I}^{-} \rightarrow \mathrm{Mn}^{2+}+4 \mathrm{H}^{2} \mathrm{O}+\mathrm{I}_{2}+2 \mathrm{e}^{-}$
(C) $2 \mathrm{MnO}_{4}^{-}+16 \mathrm{H}^{+}+10 \mathrm{I}^{-} \rightarrow 2 \mathrm{Mn}^{2+}+8 \mathrm{H}_{2} \mathrm{O}+5 \mathrm{I}_{2}$
(D) $\quad 2 \mathrm{MnO}_{4}^{-}+16 \mathrm{H}^{+}+4 \mathrm{I}^{-} \rightarrow 2 \mathrm{Mn}^{2+}+8 \mathrm{H}_{2} \mathrm{O}+2 \mathrm{I}_{2}$
9. Which of the following is NOT true about a dynamic equilibrium?
(A) It is reversible.
(B) Macroscopic properties are constant.
(C) Microscopic processes are in balance.
(D) It cannot be achieved in a closed system.

Item 10 refers to the following equation

$$
\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{NH}_{2}(\mathrm{~g})+\mathrm{HNO}_{2}(\mathrm{aq}) \rightarrow \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}(l)+\mathrm{H}_{2} \mathrm{O}(l)+\mathrm{N}_{2}(\mathrm{~g})
$$

10. The volume of nitrogen gas collected at r.t.p. when 0.01 mol of ethylamine completely reacts with an excess of nitrous acid is
(A) $0.224 \mathrm{dm}^{3}$
(B) $0.24 \mathrm{dm}^{3}$
(C) $2.24 \mathrm{dm}^{3}$
(D) $2.40 \mathrm{dm}^{3}$
11. A mass of 25 g of an unknown gas, X , is pumped through an industrial pipeline of volume $60 \mathrm{~cm}^{3}$ at a pressure of 120 kPa and temperature of $150^{\circ} \mathrm{C}$. (Molar gas constant $=8.31 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}$.)

What is the relative molecular mass of the gas?
(A) $1.2 \times 10^{-2}$
(B) $4.3 \times 10^{-2}$
(C) 4.3
(D) $\quad 12.2$
12. The melting points and boiling points of two pure substances, X and Y , are given below:

## Melting point $/ \mathrm{K} \quad$ Boiling point $/ \mathrm{K}$

| X | 273 | 373 |
| :--- | :--- | :--- |
| Y | 317 | 400 |

Which of the following statements about X and $Y$ are correct?
I. At 298 K , the particles of X can occupy the volume of its container.
II. At 298 K , the particles of Y are in fixed positions.
III. At 380 K , the particles of X have more energy than the particles of Y.
(A) I and II only
(B) I and III only
(C) II and III only
(D) I, II and III

Items 13 and 14 refers to the following options.
(A) Enthalpy of formation
(B) Lattice energy
(C) Enthalpy of solution
(D) Enthalpy of hydration

Match EACH item below with one of the options above, each of which may be used once, more than once or not at all.
13. $\mathrm{Y}(\mathrm{s})+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{YO}_{2}(\mathrm{~g})$
14. $X Y(s)+$ water $\rightarrow X Y(a q)$

Item 15 refers to the diagram below which shows a Born-Haber Cycle for NaCl .

15. What is the lattice energy of sodium chloride?
(A) $\quad-727 \mathrm{~kJ} \mathrm{~mol}^{-1}$
(B) $\quad-95 \mathrm{~kJ} \mathrm{~mol}^{-1}$
(C) $\quad+663 \mathrm{~kJ} \mathrm{~mol}^{-1}$
(D) $\quad+727 \mathrm{~kJ} \mathrm{~mol}^{-1}$
16. The general form of a rate equation is given by: Rate $=k[A]^{\mathrm{a}}[\mathrm{B}]^{\mathrm{b}}$

The rate constant, $k$, is defined as the rate of reaction when the concentration of each reactant (in $\mathrm{mol} \mathrm{dm}^{-3}$ ) is
(A) 0
(B) $1 / 2$
(C) 1
(D) 2

Item 17 refers to the data in the table, obtained for the reaction between molecules X and Y at 500 K . The stoichiometric equation for the reaction is
$2 \mathrm{X}(\mathrm{g})+\mathrm{Y}_{2}(\mathrm{~g}) \rightarrow \mathrm{X}_{2} \mathrm{Y}_{2}(\mathrm{~g})$

| Experiment <br> Number | Initial concentration/ <br> mol dm |  | Inital rate/ <br> mol $\mathbf{d m}^{-3} \mathbf{s}^{-1}$ |
| :---: | :---: | :---: | :---: |
|  | $\mathbf{X}$ | $\mathbf{Y}$ |  |
| 1 | 0.01 | 0.02 | $2.4 \times 10^{-6}$ |
| 2 | 0.02 | 0.02 | $4.8 \times 10^{-6}$ |
| 3 | 0.01 | 0.01 | $1.2 \times 10^{-6}$ |

17. The correct orders of reaction with respect to X and Y are

|  | $X$ | $Y$ |
| :---: | :---: | :---: |
| $(A)$ | 0 | 2 |
| (B) | 2 | 0 |
| $(C)$ | 1 | 0 |
| $(D)$ | 1 | 1 |

Item 18 refers to the following data for the reaction between propanone, iodine and hydrogen ions.
$\mathrm{CH}_{3} \mathrm{COCH}_{3}+\mathrm{H}^{+}+\mathrm{I}_{2} \rightarrow \mathrm{ICH}_{2} \mathrm{COCH}_{3}+2 \mathrm{H}^{+}+\mathrm{I}_{2}$

| Expt. <br> no | Concentration/mol dm |  |  | Rate $/ \mathbf{m o l}$ <br> $\mathbf{d m}^{-3} \mathbf{s}^{\mathbf{- 1}}$ |
| :---: | :---: | :---: | :---: | :---: |
|  | $\mathbf{C H}_{\mathbf{3}} \mathbf{C O C H}_{\mathbf{3}}$ | $\mathbf{H}^{+}$ | $\mathbf{I}_{\mathbf{2}}$ |  |
| 1 | 0.1 | 0.1 | 0.1 | 0.04 |
| 2 | 0.2 | 0.1 | 0.1 | 0.08 |
| 3 | 0.1 | 0.2 | 0.1 | 0.08 |
| 4 | 0.1 | 0.2 | 0.2 | 0.08 |

The rate of reaction is given by
Rate $=\mathrm{k}\left[\mathrm{CH}_{3} \mathrm{COCH}_{3}\right]^{\mathrm{x}}\left[\mathrm{H}^{+}\right]^{\mathrm{y}}\left[\mathrm{I}_{2}\right]^{\mathrm{z}}$
18. The values of $x, y$ and $z$ are
(A) $\mathrm{x}=1, \mathrm{y}=1, \mathrm{z}=1$
(B) $\quad \mathrm{x}=0, \mathrm{y}=1, \mathrm{z}=1$
(C) $\mathrm{x}=1, \mathrm{y}=1, \mathrm{z}=0$
(D) $\mathrm{x}=0, \mathrm{y}=0, \mathrm{z}=1$

Item 19 refers to the graph below which shows the Boltzmann distribution of kinectic energies in a gas.

No. of molecules with a given energy

19. Which of the following statements are correct for the Boltzmann distribution?

Increasing temperature
I. causes the maximum of the curve to move to the right
II. increases the number of molecules
III. increases the number of molecules with energy greater than the activation energy
(A) I and II only
(B) I and III only
(C) II and III only
(D) I, II and III
20. Based on Le Chatelier's principle, which combination of obervations is NOT true?

|  | Type of Reaction | Temperature <br> Change | Effect on $\mathbf{K}_{\mathbf{c}}$ | Effect on Reactants |
| :--- | :--- | :---: | :---: | :---: |
| (A) | Endothermic | decrease | decrease | increase |
| (B) | Endothermic | increase | increase | decrease |
| (C) | Exothermic | increase | increase | decrease |
| (D) | Exothermic | decrease | increase | decrease |

Item 21 refers to the data in the table below

| Gas | Equilibrium Partial Pressure |
| :---: | :---: |
| $\mathrm{N}_{2}$ | a |
| $\mathrm{H}_{2}$ | b |
| $\mathrm{NH}_{3}$ | c |

21. What is the expression for $\mathrm{K}_{\mathrm{p}}$ for the equilibruim $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \underset{\rightleftarrows}{\rightleftarrows} \mathrm{NH}_{3}(\mathrm{~g})$ ?
(A) $\frac{c^{2}}{a^{3} b}$
(B) $\frac{a^{3} b}{c^{2}}$
(C) $\frac{a b^{3}}{c^{2}}$
(D) $\frac{c^{2}}{a b^{3}}$
22. Which of the following are BronstedLowry acids?
I. $\quad \mathrm{NH}_{4}^{+}$
II. $\quad \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{OH}$
III. $\quad \mathrm{CH}_{3} \mathrm{COO}^{-}$
(A) I and II only
(B) I and III only
(C) II and III only
(D) I, II and III
23. What is the $\mathrm{K}_{\mathrm{w}}$ value of $2.0 \mathrm{~mol} \mathrm{dm}^{-3}$ $\mathrm{NaOH}(\mathrm{aq})$ at 298 K ?
(A) $2.0 \times 1.0 \times 10^{-14}$
(B) $1.0 \times 10^{-14} / 2.0$
(C) $1.0 \times 10^{-14}$
(D) $4.0 \times 10^{-14}$
24. Which of the following indicators would be suitable for use in titrating a strong acid against a weak base?
Indicator $\quad \mathrm{pH}$ range
I. Bromophenol blue
2.8-4.6
II. Methyl red
4.2-6.3
III. Alizaren yellow 10.1-13.0
(A) I only
(B) I and II only
(C) II and III only
(D) I, II and III
25. A buffer solution consists of ethanoic acid and sodium ethanoate solution.

Which equations show how the buffer maintains constant pH in solution?
I. $\mathrm{CH}_{3} \mathrm{COO}^{-}+\mathrm{NH}_{4}^{+} \rightarrow \mathrm{CH}_{3} \mathrm{COONO}_{4}$
II. $\mathrm{CH}_{3} \mathrm{COO}^{-}+\mathrm{H}^{+} \rightarrow \mathrm{CH}_{3} \mathrm{COOH}$
III. $\mathrm{CH}_{3} \mathrm{COOH}+\mathrm{OH}^{-} \rightarrow \stackrel{3}{\mathrm{CH}_{3} \mathrm{COO}^{-}+\mathrm{H}_{2} \mathrm{O}}$
(A) I and II only
(B) I and III only
(C) II and III only
(D) I, II and III

Item 26 refers to the following reaction:
$\mathrm{CH}_{3} \mathrm{COOH}(\mathrm{aq}) \rightleftarrows \mathrm{H}^{+}(\mathrm{aq})+\mathrm{CH}_{3} \mathrm{COO}^{-}(\mathrm{aq})$
[sodium ethanoate] $=0.5 \mathrm{~mol} \mathrm{dm}^{-3}$ and [ethanoic acid] $=0.3 \mathrm{~mol} \mathrm{dm}^{-3}$
26. Given that $\mathrm{K}_{\mathrm{a}}=1.8 \times 10^{-5}$, what is the pH of the buffer solution?
(A) 3.25
(B) 4.97
(C) 6.50
(D) 9.94
27. When solid silver chloride is added to a solution of potassium iodide, a yellow precipitate of silver iodide forms because
(A) chlorine is more electronegative than iodine
(B) potassium chloride and potassium iodide are both soluble
(C) silver chloride has a lower $\mathrm{K}_{\text {sp }}$ value than silver iodide
(D) silver iodide has a lower $\mathrm{K}_{\text {sp }}$ value than silver chloride
28. Which of the following is NOT a standard condition for measurement of electrode potentials?
(A) Solutions have a concentration of $1 \mathrm{~mol} \mathrm{dm}^{-3}$.
(B) Gases have a pressure of 1 atmosphere.
(C) Temperature is $25^{\circ} \mathrm{C}$.
(D) Metal alloys are used as electrodes
29. Which pair of half cell potentials would produce an $\mathrm{E}^{\Theta} / \mathrm{V}$ of 1.1 when combined to form a cell?
(A) $\mathrm{Cu}^{2+}(\mathrm{aq}) / \mathrm{Cu}(\mathrm{s})$ and $\mathrm{Fe}^{3+}(\mathrm{aq}) / \mathrm{Fe}^{2+}(\mathrm{aq})$
(B) $\quad \mathrm{Cu}^{2+}(\mathrm{aq}) / \mathrm{Cu}(\mathrm{s})$ and $\mathrm{Zn}^{2+}(\mathrm{aq}) / \mathrm{Zn}(\mathrm{s})$
(C) $\quad \mathrm{Al}^{3+}(\mathrm{aq}) / \mathrm{Al}(\mathrm{s})$ and $\mathrm{Zn}^{2+}(\mathrm{aq}) / \mathrm{Zn}(\mathrm{s})$
(D) $\quad \mathrm{Ag}^{+} / \mathrm{Ag}(\mathrm{s})$ and $\mathrm{Cu}^{2+}(\mathrm{aq}) / \mathrm{Cu}$ (s)
30. The standard electrode potentials for dichromate and iodide are:

$$
\begin{array}{ll}
1 / 2 \mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}(\mathrm{aq})+7 \mathrm{H}^{+}(\mathrm{aq})+3 \mathrm{e} \rightleftarrows \mathrm{Cr}^{3+}(\mathrm{aq})+7 / 2 \mathrm{H}_{2} \mathrm{O} & +1.33 \mathrm{~V} \\
\mathrm{I}_{2}(\mathrm{aq})+2 \mathrm{e}^{-}+2 \mathrm{e}^{-} \rightleftarrows 2 \mathrm{I}^{-}(\mathrm{aq}) & \\
& +0.54 \mathrm{~V}
\end{array}
$$

From this it can be deduced that
(A) acidified dichromate ions would oxidise iodide ions to iodine
(B) iodine would oxidise chromium (III) ions to dichromate
(C) combining both half cells would produce a cell potential of 1.87
(D) electrons would flow from the dichromate half cell to the iodine half cell if the two are connected

Items 31 and 32 refer to the following graphs.
(A)
(B)

(C)


In answering Items 31 and 32, match each item with one of the graphs.
Each graph may be used once, more than once or not at all.
31. Atomic radius
32. First ionization energy

Items 33 and 34 refer to the following oxidation states.
(A) +1
(B) +3
(C) +6
(D) +7

Match each element below with one of the options above, each of which may be used more than once, once or not at all
33. $\mathrm{P}_{4} \mathrm{O}_{6}$
34. $\mathrm{Na}_{2} \mathrm{O}$
35. Which of the following has a simple molecular structure?
(A) Sodium oxide
(B) Magnesium chloride
(C) Aluminium chloride
(D) Silicon (IV) oxide
36. In descending Group II of the periodic table, the first ionization energy value of the elements decreases. This is due to which two of the following?
I. Atomic radius increases down the group.
II. Electrons are more delocalised as ionic radius decreases.
III. Each succeeding atom has one more quantum shell than the other.
(A) I and II only
(B) I and III only
(C) II and III only
(D) I, II and III
37. Group IV elements show an increasing tendency towards metallic character as atomic number increases. Which of the following may account for this?
I. The +2 state becomes more stable than the +4 state.
II. The +4 state becomes more stable than the +2 state.
III. The valence electrons in the +4 state do not take part in bonding.
(A) I only
(B) II only
(C) III only
(D) II and III only
38. Which of the following oxides of elements in Group IV is the LEAST acidic in character?
(A) $\mathrm{SnO}_{2}$
(B) PbO
(C) $\quad \mathrm{CO}_{2}$
(D) $\quad \mathrm{SiO}_{2}$
39. The decrease in volatility going down Group VII CANNOT be described by which of the following?
(A) Strength of van der Waals forces increases
(B) Electropositivity increases
(C) Size of atoms increases
(D) Molecular mass increases
40. A student tests an unknown sample with a few $\mathrm{cm}^{3}$ of $\mathrm{AgNO}_{3}(\mathrm{aq})$ followed by a few $\mathrm{cm}^{3}$ of dilute $\mathrm{NH}_{3}(\mathrm{aq})$. Which of the following observations would confirm that the sample contains the $\mathrm{Br}^{-}$ion?

|  | Precipitate <br> with AgNO 3 | Dissolves in dilute <br> $\mathbf{N H}_{\mathbf{3}}$ |
| :---: | :---: | :---: |
| (A) | white | sparingly |
| (B) | white | readily |
| (C) | off-white | sparingly |
| (D) | off-white | readily |

41. Transition metal complexes contain a central metal atom or ion surrounded by ligands. Which of the following does NOT explain why these complexes are formed?
(A) The presence of lone pairs of electrons on ligands
(B) The presence of vacant d-orbitals on the metal atom or ion
(C) The formation of covalent bonds between the metal and ligands
(D) The complex ion formed is stable with respect to its constituents
42. Which of the following represents the electronic configuration of $\mathrm{Mn}^{3+}$ ?

Note: $[\mathrm{Ar}]=1 \mathrm{~s}^{2}, 2 \mathrm{~s}^{2} 2 \mathrm{p}^{6}, 3 \mathrm{~s}^{2} 3 \mathrm{p}^{6}$
(A)
[Ar]
3d

|  | $\uparrow$ | $\uparrow$ | $\uparrow$ |
| :--- | :--- | :--- | :--- | $\begin{array}{r}\text { 4S } \\ \uparrow \downarrow \\ \hline\end{array}$

(B) $[\mathrm{Ar}]$
$\begin{array}{r}4 S \\ \square \\ \hline\end{array}$
(C)
[Ar]

| $4 S$ |
| ---: |
| $\square$ |

(D)
[Ar]

| 4 S |
| :---: |
| $\uparrow \downarrow$ |

43. Transition metals can exhibit more than one oxidation state because they
(A) have more than one free electron
(B) exist in different colours
(C) have similar energy levels for 3d and 4 s
(D) have incomplete d-orbitals

Item 44 refers to the following information.

An unknown salt sample, $F A l$, is subjected to analysis and the results are shown in the table.

| Reaction | Results |
| :--- | :--- |
| $\mathrm{KI}(\mathrm{aq})+\mathrm{FA} l(\mathrm{aq})$ | No visible reaction |
| $\mathrm{NH}_{3}(\mathrm{aq})+\mathrm{FA} l(\mathrm{aq})$ | White precipitate, <br> insoluble in excess <br> $\mathrm{NH}_{3}(\mathrm{aq})$ |

44. It can be deduced that the cation is MOST likely
(A) $\mathrm{Al}^{3+}$
(B) $\mathrm{Zn}^{2+}$
(C) $\mathrm{Na}^{+}$
(D) $\mathrm{Pb}^{2+}$
45. When silver nitrate solution followed by dilute ammonia is added to a solution containing chloride ions, which of the following is the formula for the complex formed?
(A) $\quad \mathrm{Ag}\left(\mathrm{NH}_{4}\right)^{+}$
(B) $\quad \mathrm{AgNH}_{2}$
(C) $\quad\left[\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}\right]^{+}$
(D) $\quad\left[\mathrm{Ag}(\mathrm{NH})_{4}\right]^{2+}$

## END OF TEST

IF YOU FINISH BEFORE TIME IS CALLED, CHECK YOUR WORK ON THIS TEST.

| Question | Syllabus Reference | Profile | Key |
| :---: | :---: | :---: | :---: |
| 1. | 1.1.13 | KC | C |
| 2. | 1.1.5 | KC | D |
| 3. | 1.2.6 | UK | A |
| 4. | 1.2.7 | UK | A |
| 5. | 1.3.1 | UK | C |
| 6. | 1.3.5 | UK | C |
| 7. | 1.4.1 | UK | A |
| 8. | 1.4.3 | UK | C |
| 9 | 1.2.1 | KC | D |
| 10. | 1.3.5 | UK | B |
| 11. | 1.3.6 | UK | D |
| 12. | 1.2.3 | KC | C |
| 13. | 1.6.6 | KC | A |
| 14. | 1.6 .6 | KC | C |
| 15. | 1.6.9 | UK | A |
| 16. | 2.1.3 | UK | C |
| 17. | 2.1.6 | UK | D |
| 18. | 2.1.4 | UK | C |
| 19. | 2.1.8 | KC | A |
| 20. | 2.2.6 | KC | C |
| 21. | 2.2.4 | UK | D |
| 22. | 2.3.1 | KC | A |
| 23. | 2.3.3 | UK | C |
| 24. | 2.3.6 | KC | B |
| 25. | 2.4.2 | KC | C |
| 26. | 2.4.3 | UK | B |
| 27. | 2.5.2 | KC | D |
| 28. | 2.6.3 | KC | D |
| 29. | 2.6.4 | UK | B |
| 30. | 2.6.5 | UK | A |
| 31. | 3.1.3 | UK | B |
| 32. | 3.1.3 | UK | A |
| 33. | 3.1.6 | KC | C |
| 34. | 3.1.1 | KC | C |
| 35. | 3.1.1 | KC | A |
| 36. | 3.2.1 | KC | B |
| 37. | 3.3.1 | KC | A |
| 38. | 3.3.1 | KC | B |
| 39. | 3.4.1 | KC | B |
| 40. | 3.6.5 | UK | C |
| 41. | 3.5.8 | KC | D |
| 42. | 3.5.4 | KC | B |
| 43. | 3.5.2 | KC | C |
| 44. | 3.6.1 | UK | A |
| 45. | 3.6.5 | KC | C |



## CANDIDATE'S RECEIPT

## INSTRUCTIONS TO CANDIDATE:

1. Fill in all the information requested clearly in capital letters.

TEST CODE:

| 0 | 2 | 1 | 1 | 2 | 0 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

SUBJECT:
CHEMISTRY - UNIT 1 - Paper 01

PROFICIENCY:
ADVANCED

REGISTRATION NUMBER:

|  |  |  |  |  |  |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

FULL NAME: $\qquad$
(BLOCK LETTERS)

Signature: $\qquad$

Date: $\qquad$
2. Ensure that this slip is detached by the Supervisor or Invigilator and given to you when you hand in this booklet.
3. Keep it in a safe place until you have received your results.

## INSTRUCTION TO SUPERVISOR/INVIGILATOR:

Sign the declaration below, detach this slip and hand it to the candidate as his/her receipt for this booklet collected by you.

I hereby acknowledge receipt of the candidate's booklet for the examination stated above.

Signature: $\qquad$
Supervisor/Invigilator

Date: $\qquad$

Unit 1 Paper 01 Keys

| Question | Syllabus Reference | Profile | Key |
| :---: | :---: | :---: | :---: |
| 1. | 1.1.13 | KC | C |
| 2. | 1.1.5 | KC | D |
| 3. | 1.2 .6 | UK | A |
| 4. | 1.2.7 | UK | A |
| 5. | 1.3.1 | UK | C |
| 6. | 1.3.5 | UK | C |
| 7. | 1.4.1 | UK | A |
| 8. | 1.4.3 | UK | C |
| 9 | 1.2.1 | KC | D |
| 10. | 1.3.5 | UK | B |
| 11. | 1.3.6 | UK | A |
| 12. | 1.2.3 | KC | C |
| 13. | 1.6.6 | KC | A |
| 14. | 1.6.6 | KC | C |
| 15. | 1.6.9 | UK | A |
| 16. | 2.1.3 | UK | C |
| 17. | 2.1.6 | UK | D |
| 18. | 2.1.4 | UK | C |
| 19. | 2.1.8 | KC | B |
| 20. | 2.2.6 | KC | C |
| 21. | 2.2.4 | UK | D |
| 22. | 2.3.1 | KC | A |
| 23. | 2.3.3 | UK | C |
| 24. | 2.3.6 | KC | B |
| 25. | 2.4.2 | KC | C |
| 26. | 2.4.3 | UK | B |
| 27. | 2.5.2 | KC | D |
| 28. | 2.6.3 | KC | D |
| 29. | 2.6.4 | UK | B |
| 30. | 2.6.5 | UK | A |
| 31. | 3.1.3 | UK | C |
| 32. | 3.1.3 | UK | A |
| 33. | 3.1.6 | KC | B |
| 34. | 3.1.1 | KC | A |
| 35. | 3.1.1 | KC | C |
| 36. | 3.2.1 | KC | B |
| 37. | 3.3.1 | KC | A |
| 38. | 3.3.1 | KC | B |
| 39. | 3.4.1 | KC | B |
| 40. | 3.6 .5 | UK | C |
| 41. | 3.5.8 | KC | D |
| 42. | 3.5.4 | KC | B |
| 43. | 3.5.2 | KC | C |
| 44. | 3.6.1 | UK | A |
| 45. | 3.6.5 | KC | C |

CARIBBEAN EXAMINATIONS COUNCIL
CARIBBEAN ADVANCED PROFICIENCY EXAMINATION ${ }^{\circledR}$

FILL IN ALL THE INFORMATION REQUESTED CLEARLY IN CAPITAL LETTERS.

TEST CODE

| 0 | 2 | 1 | 1 | 2 | 0 | 2 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

SUBJECT $\qquad$ CHEMISTRY - UNIT 1 - Paper 02

PROFICIENCY $\qquad$ ADVANCED REGISTRATION NUMBER


| NAME OF SCHOOL/CENTRE |
| :---: |
|  |


| CANDIDATE'S FULL NAME (FIRST, MIDDLE, LAST) |
| :---: |
|  |

DATE OF BIRTH


SIGNATURE $\qquad$

| "*"Barcode Area"*" |
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| Sequential Bar Code |



TEST CODE $\mathbf{0 2 1 1 2 0 2 0}$

## CARIBBEAN EXAMINATIONS COUNCIL <br> CARIBBEAN ADVANCED PROFICIENCY EXAMINATION ${ }^{\circledR}$ <br> CHEMISTRY

SPECIMEN PAPER
UNIT 1 - Paper 02
2 hours 30 minutes

## READ THE FOLLOWING INSTRUCTIONS CAREFULLY.

1. This paper consists of THREE questions. Answer ALL questions.
2. Write your answers in the spaces provided in this booklet.
3. Do NOT write in the margins.
4. Where appropriate, ALL WORKING MUST BE SHOWN in this booklet.
5. A data booklet is provided.
6. You may use a silent, non-programmable calculator to answer questions.
7. If you need to rewrite any answer and there is not enough space to do so on the original page, you must use the extra lined page(s) provided at the back of this booklet. Remember to draw a line through your original answer.
8. If you use the extra page(s) you MUST write the question number clearly in the box provided at the top of the extra page(s) and, where relevant, include the question part beside the answer.
do not turn this page until you are told to do so.
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## Answer ALL questions. <br> MODULE 1 <br> FUNDAMENTALS IN CHEMISTRY

1. (a) Define the terms
(i) Mole
$\qquad$
$\qquad$
$\qquad$
(ii) Molar mass
$\qquad$
$\qquad$
$\qquad$
(iii) State the units of molar mass
$\qquad$
$\qquad$
$\qquad$
(iv) State Avogadro's Law.
$\qquad$
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(b) $20 \mathrm{~cm}^{3}$ of a solution of phosphorous (V) acid containing $1.96 \mathrm{~g} \mathrm{dm}^{-3}$ reacted with $25 \mathrm{~cm}^{3}$ of a solution containing $1.28 \mathrm{~g} \mathrm{dm}^{-3}$ sodium hydroxide.

Relative atomic masses: $\mathrm{P}=31.0, \mathrm{H}=1.0, \mathrm{Na}=23.0, \mathrm{O}=16.0$
Calculate the number of moles of
(i) phosphorous (V) acid that reacted
(ii) sodium hydroxide that reacted
(iii) sodium hydroxide that reacted with 1 mole of the acid.
[1 mark]
(iv) Derive the equation for the reaction that occurred in Part (b) above.
$\qquad$
(v) Outline the steps involved in carrying out the reaction described in (b) above.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
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(c) The following diagram represents the Born Haber cycle for the formation of compound MX(s).


The associated enthalpy changes in $\mathrm{kJ} \mathrm{mol}^{-1}$ for the cycle above are as follows:

$$
\Delta \mathrm{HM}(\mathrm{~s})=+86 ; \Delta \mathrm{H} \mathrm{M}(\mathrm{~g})=+408 ; \Delta \mathrm{H}(\mathrm{MX}(\mathrm{~s}))=-431 ; \Delta \mathrm{H} \mathrm{X}_{2}=+122 ; \Delta \mathrm{HX}(\mathrm{~g})=-372
$$

(i) State Hess' Law of heat summation.
$\qquad$
$\qquad$
(ii) Define standard enthalpy change of formation.
$\qquad$
$\qquad$
(iii) Which enthalpy values correspond to EACH of the stages I, III and IV in the cycle above?
$\qquad$
$\qquad$
(iv) Define lattice energy.
$\qquad$
$\qquad$
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(v) Calculate the lattice energy of MX(s).
(d) Account for the difference in the lattice energy for $\mathrm{MgCl}_{2}(\mathrm{~s})$ and $\mathrm{NaCl}(\mathrm{s})$.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(e) Explain why the molar enthalpy changes for the following reactions have identical values.
$\mathrm{HCl}(\mathrm{aq})+\mathrm{NaOH}(\mathrm{aq}) \rightarrow \mathrm{NaCl}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}$ (l)
$\mathrm{H}_{2} \mathrm{SO}_{4}(\mathrm{aq})+2 \mathrm{KOH}(\mathrm{aq}) \rightarrow \mathrm{K}_{2} \mathrm{SO}_{4}(\mathrm{aq})^{2}+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$
$\qquad$
$\qquad$
$\qquad$

## MODULE 2

## KINETICS AND EQUILIBRIA

2. A student is attempting to find the cell potential of a Daniell $(\mathrm{Zn} / \mathrm{Cu})$ cell.
(a) (i) Outline the steps he has to follow to obtain a reading of approximately 1.10 V on his voltmeter.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) Write the ionic equation for the reaction occurring at EACH of the electrodes.
$\qquad$
$\qquad$
(iii) Identify the anode and the cathode.

Anode: $\qquad$

Cathode: $\qquad$
(iv) Explain the direction of electron flow.
(v) Write the cell diagram.
(vi) Write the equation to represent the cell reaction.
$\qquad$
$\qquad$
(b) Use the $\mathrm{E}^{\ominus}$ value for each electrode (in the data booklet) to determine the $\mathrm{E}^{\ominus}$ cell. Does your calculated value concur with the measured value of $\mathrm{E}^{\theta}$ cell $=1.10 \mathrm{~V}$ given on page 8 ?
(c) Suggest TWO changes which could be made to the cell in (a) to cause the cell potential to be greater than 1.10 V .
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
[2 marks]
(d) Account for the observations in the statement below.

Pure water is a poor conductor of electricity whereas pure water in the presence of a trace of $\mathrm{H}^{+}(\mathrm{aq})$ is a good conductor.
$\qquad$
$\qquad$
$\qquad$
(e) (i) Derive the relationship for the ionic product of water, Kw.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) State how the value of Kw varies with temperature.
$\qquad$
$\qquad$
(f) (i) Calculate the pH of EACH of the solutions $\mathrm{A}, \mathrm{B}$ and C , given that their concentrations are as shown in the table below.

| Solution | Concentration of $\mathbf{H}^{+}(\mathbf{a q}) \mathbf{m o l}$ dm $^{-3}$ |
| :---: | :---: |
| A | $2 \times 10^{-5}$ |
| B | $1 \times 10^{-2}$ |
| C | $1 \times 10^{-14}$ |

(i) Place the solutions $\mathrm{A}, \mathrm{B}$ and C in 5 (c) (i) on a pH scale relative to the pH of pure water.

(g) A student is asked to prepare a buffer solution using an aqueous solution of sodium hydroxide and one of the aqueous solutions, $\mathrm{X}(\mathrm{aq})$ or $\mathrm{Y}(\mathrm{aq})$. The dissociation constants for X is $\mathrm{K}_{\mathrm{a}}=1.8 \times 10^{-5}$, and for Y is $\mathrm{K}_{\mathrm{b}}=1.8 \times 10^{-5}$.
(i) Define the term 'buffer solution'.
$\qquad$
$\qquad$
(ii) Which of the solutions, X or Y , would you use with the sodium hydroxide to prepare a buffer solution? Justify your answer.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(iii) Explain how small additions of $\mathrm{H}^{+}$and $\mathrm{OH}^{-}$ions are accommodated in the buffer solution prepared above.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$

Total 30 marks
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## MODULE 3

## CHEMISTRY OF THE ELEMENTS

3. (a) Transition elements form coloured compounds.

State THREE other characteristic properties of transition elements
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(b) What are the colours of aqueous
(i) $\mathrm{CO}^{2+}$ ?
$\qquad$
(ii) $\mathrm{Mn}^{2+}$ ?
(c) If aqueous $\mathrm{CO}^{2+}$ is heated to dryness, what colour is observed?
$\qquad$
(d) Explain what is meant by the term 'ligand'.
$\qquad$
$\qquad$
(e) Write the formula of the species formed, and describe what occurs when
(i) ammonia solution is added to aqueous copper (II) sulphate (IV)
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) an excess of ammonia solution is added to (e) (i).
$\qquad$
$\qquad$
$\qquad$
(f) The presence of carbon monoxide in the blood can prevent oxygen from reaching the tissues.

Use the ligand exchange theory to account for this occurrence.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
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(g) Calculate the oxidation number of the first mentioned element in EACH of the following compounds, and explain the variation in these oxidation numbers:
$\mathrm{MgCl}_{2} \rightarrow\left[\mathrm{Al}(\mathrm{OH})_{4}\right]^{-} \rightarrow \mathrm{SiF}_{6}{ }^{2-} \rightarrow \mathrm{PO}_{3}^{3-}$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(h) (i) Explain the terms 'atomic radius' and 'ionic radius'.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) Account for the variation in atomic and ionic radii of the elements in Group II.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(i) Explain the variation in the acid/base character of the oxides of oxidation state +2 of the elements of Group IV.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$

Total 30 marks

## END OF TEST

## IF YOU FINISH BEFORE TIME IS CALLED, CHECK YOUR WORK ON THIS TEST.

## EXTRA SPACE

## If you use this extra page, you MUST write the question number clearly in the box provided.

 Question No. $\square$$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
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SUBJECT:
CHEMISTRY - UNIT 1 - Paper 02

PROFICIENCY:
ADVANCED

REGISTRATION NUMBER:

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CHEMISTRY

UNIT 1 - PAPER 02
MARK SCHEME
SPECIMEN

## Question 1

S.O.: Module: 1 - $3.1,3.2,3.3,3.5,3.6,3.8,3.9,6.1,6.2,6.4,6.6,6.7$, 6.16
(a) (i) mole: an amount of a substance that contains as many particles as C atoms in 12 g of $\mathrm{C}-12$ isotope [2 marks] [C amount of a substance containing $6 \times 10^{23}$ particles 1 mark only]
(ii) Mass of 1 mole of a substance in grams
[1 mark]
(iii) Unit of molar mass is $g \mathrm{~mol}^{-1}$
[1 mark]
(iv) Avogadro's Law: Equal volumes of all gases contain the same number of molecules under the same conditions of temperature and pressure.
[1 mark]
(b) (i) Molar mass $\mathrm{H}_{3} \mathrm{PO}_{4}=3+31+64=98 \mathrm{~g}$

98 g of $\mathrm{H}_{3} \mathrm{PO}_{4}$ is the mass of 1 mole
1.96 g of $\mathrm{H}_{3} \mathrm{PO}_{4}$ is the $1 / 98 \mathrm{x} 1.96 \mathrm{~mole}=0.02 \mathrm{~mole}$
[1 mark]
$1000 \mathrm{~cm}^{3} \mathrm{H}_{3} \mathrm{PO}_{4}$ soln contains 0.02 mole
$\therefore 20 \mathrm{~cm}^{3} \mathrm{H}_{3} \mathrm{PO}_{4}$ soln contains $\frac{0.02}{1000} \times 20$
$=0.0004 \mathrm{~mole}$
[1 mark]
(ii) No of mole $\mathrm{NaOH}=\frac{1}{40} \times 1.28=0.032$ mole
$25 \mathrm{~cm}^{3} \mathrm{Na}$ OH solution contains $\frac{0.032}{1000} \times 25$
$=0.0008 \mathrm{~mole}$
[1 mark]
(iii) 0.0004 mole acid reacts with 0.0008 mole NaOH
$\therefore 1$ mole acid reacts with $\frac{0.0008}{0.0004}$
$=2$ moles
[1 mark]
(iv) $\mathrm{H}_{3} \mathrm{PO}_{4}(\mathrm{aq})+2 \mathrm{NaOH}(\mathrm{aq}) \longrightarrow \mathrm{Na}_{2} \mathrm{HPO}_{4}(\mathrm{aq})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$
[1 mark]
(v)

- Measure $25 \mathrm{~cm}^{3} \mathrm{NaOH}$ solution with a clean pipette
- Deliver this solution in a clean $250 \mathrm{~cm}^{3}$ conical flask
- Add a few drops of phenolphthalein indicator
- Fill a clean burette to the zero mark with phosphoric acid
- Add acid dropwise to the alkali until the solution goes colourless
[5 marks]

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## CHEMISTRY

UNIT 01 - PAPER 02
MARK SCHEME
(c) (i) Hess's law of constant heat summation states that for any extensive property (enthalpy), the $\Delta H$ in going from reactants to products is a constant regardless of the path taken, as long as the reaction goes from the same initial state to the same final state.
[1 mark]
(ii) Standard enthalpy of formation is the enthalpy change when one mole of a compound is formed from the elements in their standard states.
[1 mark]
(iii) Stage I: $\quad \Delta H^{\theta}$
[1 mark]
Stage III: $\triangle H^{\theta_{D}}$
[1 mark]
Stage IV: $\triangle H^{\theta_{E}}$
[1 mark]
(iv) Lattice energy is the enthalpy of formation for one mole of the ionic compound from gaseous ions under standard conditions.
[1 mark]
(v) Lattice energy of $M X(s)=\Delta H^{\theta}{ }_{F}-\left(\Delta H^{\theta}+\Delta H^{\theta}+\Delta H^{\theta}+\Delta H^{\theta}{ }_{\mathrm{E}}\right)$
[1 mark]
$=-431-(+86+408+122-372)$
[1 mark]
$=-675 \mathrm{~kJ} \mathrm{~mol}^{-1}$
[1 mark]
(d) The difference in lattice energies

- For Mg, one has to take into consideration the $1^{\text {st }}$ and $2^{\text {nd }}$ ionization energies of Mg. (+736, 1450).
[2 marks]
- For Cl, the two electrons that are released are given one each to 2 chlorine atoms. Adding one electron as a time - you need to multiply E.A by 2.
[2 marks]
(e) The same overall reaction for each since a strong acid and a strong base are reacting.

$$
\begin{aligned}
& \mathrm{H}^{+}+\mathrm{OH}^{-} \longrightarrow \mathrm{H}_{2} \mathrm{O} \\
& 2 \mathrm{H}^{+}+\mathrm{OH}^{-} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}
\end{aligned}
$$

Since one is dealing with molar enthalpy change the values of $\Delta H$ will be the same.
[2 marks]

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S.O.: Module: $1-3.1,3.2,3.3,4.1,4.2,6.3,6.4,6.5$
(a) (i) Place a strip of zinc in 1 mol $\mathrm{dm}^{-3} \mathrm{Zn}^{2+}(\mathrm{aq})$ in a beaker - Place a strip of copper in $1 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{Cu}^{2+}(a q)$ in a beaker

- Connect the two metal strips to a high resistance voltmeter
- Connect the two solutions by means of a salt bridge a strip filter soaked in $\mathrm{KNO}_{3}(a q)$ or $\mathrm{KCI}(a q)$ (or solutions in the tube fitted with porous plugs)
- Try to maintain temperature if $25^{\circ} \mathrm{C}$
(ii) $\quad \mathrm{Zn}(\mathrm{s}) \leftrightarrows \mathrm{Zn}^{2+}+2 \mathrm{e}$
[1 mark]
$\mathrm{Cu}^{2+}(\mathrm{aq})+2 \mathrm{e} \leftrightarrows \mathrm{Cu}(\mathrm{s})$
[1 mark]
(iii) The anode is Zn ; the cathode is Cu
[1 mark]
(iv) The electrons flow from zinc to the copper half cell since the zinc gives up electrons more readily than copper.
[1 mark]
(v) $\quad \mathrm{Zn}(\mathrm{s}) \backslash \mathrm{Zn}^{2+}(\mathrm{aq}) \backslash \backslash \mathrm{Cu}^{2+}(\mathrm{aq})+\mathrm{Cu}(\mathrm{s})$
[1 mark]
(vi) $\mathrm{Cu}^{2+}(\mathrm{aq})+\mathrm{Zn}(\mathrm{s}) \longleftrightarrow \mathrm{Zn}^{2+}(\mathrm{aq})+\mathrm{Cu}(\mathrm{s})$
[1 mark]

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(d) Pure water is a poor conductor of electricity due to the following equilibrium which lies to the far left $\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \leftrightarrows \mathrm{H}^{+}(\mathrm{aq})+\mathrm{OH}^{-}(\mathrm{aq})$.

Since the degree of ionization is very small, it is a poor conductor.

In the presence of $\mathrm{H}+$, pure water is a good conductor as the concentration of free ions present increases.
[1 mark]
(e) (i) The $K_{c}$ expression for water becomes

$$
\mathrm{K}_{\mathrm{c}}=\frac{\left[\mathrm{H}^{+}(\mathrm{aq})\right]\left[\mathrm{OH}^{-}(\mathrm{aq})\right]}{\left[\mathrm{H}_{2} \mathrm{O}(\mathrm{l})\right]}
$$

Since the $\mathrm{H}_{2} \mathrm{O}$ is effectively constant, this is incorporated into the $K_{c}$ constant giving a new constant

$$
\mathrm{K}_{\mathrm{w}}=\left[\mathrm{H}^{+}(\mathrm{aq})\right]\left[\mathrm{OH}^{-}(\mathrm{aq})\right]
$$

[2 marks]
(ii) $K_{w}$ increases with temperature.

As $T$ increases, more energy is available to ionize $\mathrm{H}_{2} \mathrm{O}$. [ $\mathrm{H}^{+}$] increases, hence $\mathrm{K}_{\mathrm{w}}$ increases.
[1 mark]
(f) (i) pH of $\operatorname{soln} \mathrm{A}=-\log \left(2 \mathrm{x} 10^{-5}\right)=4.7$
[1 mark]
pH of $\operatorname{soln} \mathrm{B}=-\log \left(1 \times 10^{-2}\right)=2$
[1 mark]
pH of $\operatorname{soln} C=-\log \left(1 \times 10^{-14}\right)=14$
[1 mark]
(ii) pH scale relative to water: solution
pH
C
Pure Water
14
A 4.7
B 2.0
[1 mark]
(g) (i) A buffer solution is one that resists changes in pH when small quantities of acid or base are added to it.
[1 mark]

```
CHEMISTRY
UNIT 01 - PAPER 02
MARK SCHEME
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(ii) From the dissociation constants, solution $X$ is a solution of weak acid while solution $Y$ is a solution of a weak base.

Sodium hydroxide will therefore react with $X$ to form the sodium salt. The mixture will consist of the weak acid and the salt of that acid, which contains the conjugate base of the acid. This will act as a buffer solution.

```
Weak acid dissociating: HX = H+ + X-
```

Salt dissociating: NaX = Na+ + X-
[2 marks]
(iii) On addition of small amounts of $\mathrm{H}^{+}$, the excess $\mathrm{X}^{-}$ions from the salt will react with the added $\mathrm{H}^{+}$and maintain the pH almost constant, hence buffering action.
[1 mark]

On addition of small amounts of base, e.g. $\mathrm{OH}^{-}$ions $\mathrm{H}^{+}$ from the acid will react and the acid equilibrium will then shift to the right in this way almost nullifying the effect of the added $\mathrm{H}^{+}$and the pH remains effectively constant, hence buffering action.
[1 mark]

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## CHEMISTRY <br> UNIT 01 - PAPER 02 <br> MARK SCHEME

Question 3
S.O: Module 3: - $2.1,3.4,5.2,5.6,5.10$
(a) (i) variable oxidation number
(ii) form complex ions

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| [1 mark] |  |  |  |
| [1 mark] |  |  |  |
| [1 mark] | 3 |  |  |
| [1 mark] | 2 |  |  |
| [1 mark] |  |  |  |
| [1 mark] |  |  |  |

(c) Blue
[1 mark]
(d) Ligands are groups that can donate electron pairs ions thus forming a co-ordinate bond.
to metal
(e) (i) blue precipitate
$\mathrm{Cu}(\mathrm{OH})_{2}$
(ii) blue ppt dissolves forming deep blue solution
[1 mark]
[1 mark]
(f) Oxygen is transported to the tissues as a complex with the haemoglobin that is found in the red blood cells.
[1 mark]

CO can also form a complex with haemoglobin.
[1 mark]

The stability constant for the haemoglobin - CO complex is much higher than that of the haemoglobin - $O_{2}$ complex, so CO competes successfully with oxygen.
[2 marks]
Therefore, the presence of $C O$ can prevent oxygen from reaching the tissues.
[1 mark]

```
\(\mathrm{O}_{2}+\) haemoglobin \(\leftrightarrows\) oxhaemoglobin
    CO + haemoglobin \(\rightarrow\) carboxyhaemoglobin
```

5
1
$\mathrm{Mg}=+2$

| $\left[\mathrm{A} 1 \mathrm{OH}_{4}\right]^{-}$ | $\mathrm{SiF}_{6}{ }^{2-}$ | $\mathrm{PO}^{3-}{ }_{3}$ |
| :--- | :--- | :--- |
| $\mathrm{X}-4=-1$ | $\mathrm{X}-6=-2$ | $\mathrm{X}-6=-3$ |
| $\mathrm{X}=+3$ | $\mathrm{X}=+4$ | $\mathrm{X}=+3$ |
| $\mathrm{~A} 1=+3$ | $\mathrm{Si}=+4$ | $\mathrm{P}=+3$ |

[4 correct $=4$ marks]
[3 correct $=3$ marks]
[2 correct $=2$ marks]
[1 correct = 1 mark]

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SUBJECT $\qquad$ CHEMISTRY - UNIT 1 - Paper 032

PROFICIENCY $\qquad$ ADVANCED
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CARIBBEAN ADVANCED PROFICIENCY EXAMINATION ${ }^{\circledR}$
CHEMISTRY

UNIT 1 - Paper 032

## ALTERNATIVE TO SCHOOL-BASED ASSESSMENT

## 2 hours

## READ THE FOLLOWING INSTRUCTIONS CAREFULLY.

1. This paper consists of THREE questions. Answer ALL questions.
2. Write your answers in the spaces provided in this booklet.
3. Do NOT write in the margins.
4. Where appropriate, ALL WORKING MUST BE SHOWN in this booklet.
5. A data booklet is provided.
6. You may use a silent, non-programmable calculator to answer questions.
7. You are advised to take some time to read through the paper and plan your answers.
8. If you need to rewrite any answer and there is not enough space to do so on the original page, you must use the extra lined page(s) provided at the back of this booklet. Remember to draw a line through your original answer.
9. If you use the extra page(s) you MUST write the question number clearly in the box provided at the top of the extra page(s) and, where relevant, include the question part beside the answer.
dO NOT TURN THIS PAGE UNTIL YOU ARE TOLD TO DO SO.

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## Answer ALL questions.

1. You are provided with TWO solutions:
(i) a solution of ammonium iron (II) sulphate hexahydrate, $\left(\mathrm{NH}_{4}\right)_{2} \mathrm{SO}_{4} \mathrm{FeSO}_{4} 6 \mathrm{H}_{2} \mathrm{O}$ containing 39.00 g of the salt in $1 \mathrm{dm}^{3}$ of solution labelled R.
(ii) a solution of potassium manganate(VII), $\mathrm{KMnO}_{4}$, labelled S .

The ionic equation for the reaction is
$5 \mathrm{Fe}^{2+}(\mathrm{aq})+\mathrm{MnO}_{4-}(\mathrm{aq})+8 \mathrm{H}^{+}(\mathrm{aq}) \rightarrow 5 \mathrm{Fe}^{3+}(\mathrm{aq})+\mathrm{Mn}^{2+}(\mathrm{aq})+4 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$.
Carry out the following procedure to determine the concentration for potassium manganate (VII) solution S .

The results are to be entered into Table 1 .

## (a) Procedure

A. Pipette $25 \mathrm{~cm}^{3}$ of the solution R into a clean conical flask.
B. Add an approximately equal volume of dilute sulphuric acid to the conical flask.
C. Fill the burette with the solution S.
D. Titrate the mixture of R with the solution S .
E. Record both your initial burette reading and reading at the end point to two decimal places.
F. Repeat steps A-E until consistent results are obtained.

TABLE 1: DATA FOR EXPERIMENTAL PROCEDURE

|  | $\mathbf{1}$ | $\mathbf{2}$ | $\mathbf{3}$ |
| :--- | :--- | :--- | :--- |
| Final burette reading $\left(\mathrm{cm}^{3}\right)$ |  |  |  |
| Initial burette reading $\left(\mathrm{cm}^{3}\right)$ |  |  |  |
| Volume of $\mathrm{KMnO}_{4}$ used $\left(\mathrm{cm}^{3}\right)$ |  |  |  |

[6 marks]
(b) Describe the colour of the mixture at the end point of the titration.
$\qquad$
$\qquad$
(c) Calculate the volume of $\mathrm{KMnO}_{4}$ used for the titration.

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(d) Calculate the number of moles of iron (II) ions in $1 \mathrm{dm}^{3}$ of solution, R.
(e) Calculate the number of moles of iron (II) ions in the volume of $25 \mathrm{~cm}^{3}$ of R .
(f) Calculate the number of moles of potassium manganate (VII) in $1 \mathrm{dm}^{3}$ of solution.
(g) (i) State TWO reagents that can be used in testing for the presence of $\mathrm{Fe}^{2+}$ ions.
$\qquad$
$\qquad$
(ii) Describe what should be observed when any ONE of the reagents in (g) (i) is added to a solution containing only $\mathrm{Fe}^{2+}$ cations.
2. A student is required to investigate the rate of reaction in which a fixed mass of magnesium metal $(0.12 \mathrm{~g})$ is added to different volumes of $1.5 \mathrm{~mol} \mathrm{dm}^{-3}$ hydrochloric acid. The acid is added from a burette and water added to make the final volume of $50 \mathrm{~cm}^{3}$. The time taken for the magnesium ribbon to disappear is recorded. Figure 1 below shows the burette reading for the volume of acid added and the time taken for the magnesium to disappear for each reaction. The initial burette reading is always $0.0 \mathrm{~cm}^{3}$.


Figure 1
(a) From the results shown in Figure 1, construct a table to show experiment number, volume of acid added from the burette, volume of water added to the acid, and the time taken for the magnesium to disappear.
(b) On the graph paper on page 7, plot a graph of time taken for the magnesium ribbon to disappear against volume of acid added from the burette.

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(iii) Explain the shape of the graph.
$\qquad$
$\qquad$
(iv) From the graph, determine the rate of reaction at 10, 20 and 30 seconds.

Rate at 10 s :

Rate at 20 s : $\qquad$

Rate at 30 s :
[3 marks]

Total 15 marks
3. Four bottles labelled A, B, C and D are found in the laboratory. One bottle contains a strong monobasic acid, $1 \mathrm{~mol} \mathrm{dm}{ }^{-3}$; another contains a weak monobasic acid, also of concentration $1 \mathrm{~mol} \mathrm{dm}^{-3}$; the third bottle contains a reducing agent of $\mathrm{E}^{\theta}=-1.39 \mathrm{~V}$, and the fourth contains another reducing agent of $\mathrm{E}^{e}=-0.14 \mathrm{~V}$. All are colourless liquids.

You have access to all laboratory equipment, and reagents including $\mathrm{H}_{2} \mathrm{O}_{2}$.
(i) Hypothesis:
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(ii) Apparatus and materials:
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(iii) Method:
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(iv) Variables:
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(v) Expected results:
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(vi) Chemical principles:
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(vii) TWO sources of error:
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CHEMISTRY - UNIT 1 - Paper 032

PROFICIENCY:
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CHEMISTRY
UNIT 1 - PAPER 032
MARK SCHEME

SPECIMEN

## Question 1

S.O.: Module: 1 -3.8, 3.9
(a)

| Burette readings $\mathrm{KMnO}_{4} / \mathrm{H}^{+}$ |  |  |  |
| :--- | :---: | :---: | :---: |
| Reading/cm ${ }^{3}$ | 1 | 2 | 3 |
| Final |  |  |  |
| Initial |  |  |  |
| Difference |  |  |  |

Correct burette readings
[6 marks]

| (If one incorrect | $=5$ marks) |
| :--- | :--- |
| (If two incorrect | $=4$ marks) |
| (If THREE incorrect | $=3$ marks) |
| (If FOUR incorrect | $=2$ marks) |
| (If FIVE incorrect | $=1$ mark) |

(More than FIVE incorrect $=0$ marks)
Volumes recorded to 2 decimal places - 1 mark
(b)

Pink tinge
[1 mark]
(c)

Candidate's average of closest values
(d)

No. of moles of $\mathrm{Fe}^{2+}(\mathrm{aq})$ in $1 \mathrm{dm}^{3}$

$$
\begin{aligned}
& =\frac{39}{392} \\
& =\quad 0.1 \mathrm{~mol} \mathrm{dm}^{-3}
\end{aligned}
$$

(e)

No. of moles $\mathrm{Fe}^{2+}$ in $25 \mathrm{~cm}^{3}=(0.1 \times 0.025)=0.0025$ moles [1 mark]


## CHEMISTRY

UNIT 1 - PAPER 032
MARK SCHEME
(f) \# of Moles $\mathrm{KMnO}_{4}$ reacted $=0.0025 \div 5=0.0005$ moles [1 mark] \# of Moles $\mathrm{KMnO}_{4}$ in $1 \mathrm{dm}^{3}$
$=0.0005$ moles $\div$ candidate's average volume x 1000 [1 mark]
(g)
(i) $\quad \mathrm{NH}_{3}(\mathrm{aq}), \mathrm{NaOH}$ (aq)
[2 marks]
(ii) Green ppt.
[1 mark]

```
CHEMISTRY
UNIT 1 - PAPER 032
MARK SCHEME
```


## Question 2

S.O.: Module: 2 -1.4 - 1.6
(i)

| Expt. \# | Volume of Acid <br> $\left(\mathbf{c m}^{3}\right)$ | Volume of <br> water (cm $)^{2}$ | Time <br> (seconds) |
| :---: | :---: | :---: | :---: |
| 1 | 9.8 | 40.2 | 85 |
| 2 | 11.4 | 38.6 | 55 |
| 3 | 16.2 | 33.8 | 29 |
| 4 | 22.5 | 27.5 | 16 |
| 5 | 30.3 | 19.7 | 8 |
| 6 | 42.1 | 7.9 | 4 |

$$
\begin{array}{rlll}
\text { Table headings } & = & 1 \text { mark } \\
4 \text { or more rows correct } & = & 4 \text { marks } \\
3 \text { rows correct } & & = & 3 \text { marks } \\
2 \text { rows correct } & & = & 2 \text { marks } \\
1 \text { row correct } & & = & 1 \text { mark } \\
& \\
\text { Axes correct } & = & 2 \text { marks } \\
\text { Shape of graph } & = & 1 \text { mark } \\
4-6 \text { points correct } & = & 2 \text { marks } \\
<4 \text { points correct } & = & 1 \text { mark }
\end{array}
$$

[Total 5 marks]
(ii)
[Total 5 marks]
(iii) As the volume of acid increases the reaction time increases. Any other reasonable expression.
[2 marks]
(iv) Use the candidates' graphs to determine rate of reaction at 10,20 , and 30 seconds.
[1 mark each = 3 marks]

Total 15 marks


Question 3
S.O.: Module: $2-6.26$
(i) Hypothesis

Any correct answer, for example:
A is a strong monobasic acid
B is ...
C is ...
D is ...
[1 mark]
(ii) Apparatus and materials
[2 marks]
(iii) Method

Any reasonable workable set of procedures, for example:

- use of pH meter for weak and strong acids;
- react acids with metal or carbonate and record time taken for reaction to stop;
- for each reducing agent, set up a cell with $\mathrm{H}_{2} \mathrm{O}_{2}$ as one half-cell and the reducing agent in the other halfcell. Measure the voltage on a high resistant voltmeter.
[3 marks]
(iv) Variables

Correct and relevant
[2 marks]
(v) Expected results

Any reasonable answer
[2 marks]
(vi) Chemical principles

- Weak acid has low $\left[\mathrm{H}^{+}\right]$and strong acid has high [ $\mathrm{H}^{+}$]. $\mathrm{pH}=-\log _{10}\left[\mathrm{H}^{+}\right]$. Therefore, weak acid has high pH , and strong acid has low pH.
- Low $\left[\mathrm{H}^{+}\right]$in solution produces slow rate of reaction and vice versa.
- $\mathrm{H}_{2} \mathrm{O}_{2}$ half-cell reaction:

$$
\mathrm{H}_{2} \mathrm{O}_{2}+2 \mathrm{H}^{+}+2 \mathrm{e} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}(\mathrm{I}) \quad \mathrm{E}^{\theta}=+1.77 \mathrm{~V}
$$

- Reducing agent half-cell reaction:

$$
\begin{array}{ll}
-e \rightarrow \text { Product } & \mathrm{E}^{\theta}=+1.39 \mathrm{~V} \\
& \mathrm{E}_{\text {cell }}=+3.16 \mathrm{~V}
\end{array}
$$

- $E_{\text {cell }}$ is produced for the weak reducing agent.
(vii) Sources of error/assumptions/limitations

Any correct answer, for example:

- It is assumed that the reducing agents are nonacidic.
[2 marks]

| KC | UK | XS |
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## CARIBBEAN EXAMINATIONS COUNCIL

CARIBBEAN ADVANCED PROFICIENCY EXAMINATION ${ }^{\circledR}$ CHEMISTRY

UNIT 2 - Paper 01
1 hour 30 minutes

## SPECIMEN PAPER

## READ THE FOLLOWING INSTRUCTIONS CAREFULLY.

1. This test consists of 45 items. You will have 1 hour and 30 minutes to answer them.
2. In addition to this test booklet, you should have an answer sheet and a data booklet.
3. Each item in this test has four suggested answers lettered (A), (B), (C), (D). Read each item you are about to answer and decide which choice is best.
4. On your answer sheet, find the number which corresponds to your item and shade the space having the same letter as the answer you have chosen. Look at the sample item below.

## Sample Item

Which of the following energy transitions is the HIGHEST in an organic compound?

Sample Answer
(A) $\quad \mathrm{n}$ to $\sigma^{*}$

(B) $\quad \sigma$ to $\sigma^{*}$
(C) n to $\pi^{*}$
(D) $\pi$ to $\pi^{*}$

The correct answer to this item is " $\sigma$ to $\sigma *$ ", so (B) has been shaded.
5. If you want to change your answer, erase it completely before you fill in your new choice.
6. When you are told to begin, turn the page and work as quickly and as carefully as you can. If you cannot answer an item, go on to the next one. You may return to that item later.
7. You may do any rough work in this booklet.
8. Figures are not necessarily drawn to scale.
9. You may use a silent, non-programmable calculator to answer items.

DO NOT TURN THIS PAGE UNTIL YOU ARE TOLD TO DO SO.

1. Which of the following may be responsible for the wide diversity of organic compounds?
2. Carbon forms strong bonds with itself.
3. Carbon forms four bonds.
4. Carbon exists in three allotropic forms.
5. Carbon acquires a full octet of electrons.
(A) I and II only
(B) I, II and III only
(C) I, II and IV only
(D) I, II, III and IV
6. Which of the following is the structure of 2-phenyl butanoic acid?

(A)

(B)

(C)

(D)

7. Which of the following compounds can be resolved into optical isomers?
(A) $\left(\mathrm{CH}_{3}\right)_{2} \mathrm{CHCH}_{2} \mathrm{OH}$
(B) $\mathrm{NH}_{2} \mathrm{CH}_{2} \mathrm{COOH}$
(C) $\mathrm{NH}_{2} \mathrm{CH}(\mathrm{CH} 3) \mathrm{COOH}$
(D) $\left(\mathrm{CH}_{3}\right)_{2} \mathrm{CHOCH}_{3}$
8. Which of the following features may be characteristic of condensation polymers?
9. Presence of OH and COOH groups in monomers
10. Elimination of a small molecule
11. Region of unsaturation in the monomers
(A) I only
(B) II only
(C) I and II only
(D) I, II and III
12. Qiana is a polymer that feels like silk and has the following generalized partial structure:


The pair of monomers in Qiana is
(A)


(B)


(C)

and

(D)

and

6. Which of the following represent naturally occuring macromolecules?
(I)

(II)

(III)

(IV) $-\mathrm{CO}-\langle \rangle_{-} \mathrm{CO}-\mathrm{O}-\mathrm{CH}_{2}-\mathrm{CH}_{2}-\mathrm{O}-$
(A) I and II only
(B) II and III only
(C) III and IV only
(D) I and IV only
7. To which carbon atom in the compound below will an electrophile be attached?

(A) A
(B) B
(C) C
(D) D
8. Chlorine reacts with methane to form chloromethane. The organic intermediate in the reaction is
(A) an ion
(B) an electrophile
(C) a nucleophile
(D) a free radical
9. Compounds $\mathrm{X}, \mathrm{Y}$ and Z below are isomers.

Y

Z


The order of decreasing ease of removal of the bromide ion by hydrolysis is
(A) $\mathrm{X}>\mathrm{Y}>\mathrm{Z}$
(B) $\quad$ Z $>$ Y $>$ X
(C) $\quad$ Y $>\mathrm{X}>\mathrm{Z}$
(D) $\quad \mathrm{Y}>\mathrm{Z}>\mathrm{X}$

Items 10-11 refer to the following compounds:
(A) $\quad \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{COCH}_{3}$
(B) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CONH}_{2}$
(C) $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CHO}$
(D) $\quad \mathrm{CH}_{3} \mathrm{COOCH}_{2} \mathrm{CH}_{3}$

Match EACH item below with one of the options above, each of which may be used once, more than once or not at all.
10. Aldehyde
11. Ester
12. Which of the following is NOT oxidised by $\mathrm{KMnO}_{4} / \mathrm{H}^{+}(\mathrm{aq})$ ?
(A)

(B)

(C)

(D)

13. Which pair of compounds is formed when benzene and methyl benzene undergo nitration?
(A)

(B)

(C)

(D)

and

14. Ethanoic acid turns blue litmus red but ethanol does not, because ethanoic acid
(A) has a polar - OH bond but ethanol does not
(B) has a higher pka value than ethanol
(C) is a stronger acid than ethanol
(D) is more miscible with water than ethanol
15. Which of the following statements about aminoethanoic acid, $\mathrm{NH}_{2} \mathrm{CH}_{2} \mathrm{COOH}$, is NOT true?
(A) Its aqueous solution has a pH of 7
(B) It has a high melting point
(C) It is soluble in base only
(D) It is soluble in both acid and base
16. In an iron determination, a student obtains a mean value of $35.50 \%$ for the iron content and a standard deviation of 0.01 . The actual value for the percentage of iron in the sample is 39.45 . In terms of precision and accuracy, which option best describes the student's results?
(A) High accuracy and high precision
(B) High accuracy and low precision
(C) Low accuracy and high precision
(D) Low accuracy and low precision
17. Which of the following is NOT a characteristic of primary standards used in titrimetric analysis?
(A) High purity
(B) Stable in air
(C) Deliquescent
(D) Soluble
18. For which pair of titrations is methyl orange (indicator range 2.9-4.6) BEST suited?
(A) Strong acid - strong base AND strong acid - weak base
(B) Strong base - weak acid AND weak acid - weak base
(C) Strong acid - strong base AND weak acid - weak base
(D) Strong acid - weak base AND strong base - weak acid
19. $25.0 \mathrm{~cm}^{3}$ of an aqueous iron (II) salt is acidified with an equal volume of dilute sulphuric acid and titrated against $0.02 \mathrm{~mol} \mathrm{dm}-3$ potassium manganate (VII) solution. The endpoint is 30.0 cm 3 .
$5 \mathrm{Fe}^{2+}(\mathrm{aq})+\mathrm{MnO}_{4}^{-}(\mathrm{aq})+8 \mathrm{H}^{+}(\mathrm{aq}) \rightarrow 5 \mathrm{Fe}^{3+}(\mathrm{aq})+\mathrm{Mn}^{2+}(\mathrm{aq})+4 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$

The correct concentration of the iron (II) ions, in $\mathrm{mol} \mathrm{dm}^{-3}$, is
(A) $\frac{5 \times 25.0}{0.02 \times 30.0}$
(B) $5 \times 0.02 \times 30.0$ 25.0
(C) $\frac{5 \times 0.02 \times 25.0}{30.0}$
(D) $\quad 25.0$
$\overline{5 \times 0.02 \times 30.0}$
20. Treatment of an 8.00 g sample containing a lead ore with excess sulphuric acid yields 4.75 g of lead (II) sulphate.

The correct expression for the percentage of lead in the sample is
(A) $\frac{207 \times 8.00}{303 \times 4.75} \times 100$
(B) $\frac{303 \times 4.75}{303 \times 8.00} \times 100$
(C) $\frac{207 \times 4.75}{303 \times 8.00} \times 100$
(D) $\frac{303 \times 8.00}{207 \times 4.75} \quad \times 100$
21. Sintered glass crucibles are used in gravimetric analysis because they
(A) have negligible weight
(B) can tolerate high temperatures
(C) have a small coefficient of expansion
(D) are porous and allow mixtures to be filtered directly
22. In the determination of the purity of a sample of magnesium carbonate by a back titration method, which of the following statements is INCORRECT?
(A) Phenolphthalein or methyl orange are suitable indicators.
(B) The amount of excess acid re maining is determined by titration with standard alkali.
(C) The sample is reacted with excess alkali.
(D) Unreacted magnesium carbonate is filtered off and weighed.
23. Which of the following substances would be produced by fractionating a $20 \%$ solution of nitric acid in water?
(A) Azeotrope then nitric acid
(B) Water then azeotrope
(C) Nitric acid only
(D) Water only

Item 24 refers to the following structure of caesalpinin.


The compound caesalpinin exhibits absorp tion at wave numbers $3400 \mathrm{~cm}^{-1}$ and 1750 $\mathrm{cm}^{-1}$ in the IR spectrum.
24. Which functional groups are responsible for these absorptions?
(A) $\quad-\mathrm{OH}$ and $\stackrel{\rightharpoonup}{\mathrm{C}}=\mathrm{O}$
(B) $\mathrm{C}-\mathrm{H}$ and -OH
(C)

(D)

25. The use of IR Spectroscopy in identifying structure of organic molecules is limited because
(A) it helps to determine the presence or absence of some functional groups
(B) the spectrum of molecules can be complex where some absorption frequencies are superimposed
(C) the position of the peaks makes it difficult for the identification of the functional groups in the molecule
(D) vibrations due to a certain bond occur in the same region of the spectrum, which makes identification difficult

Item 26 refers to the following gas/liquid chromatogram.

26. Based on the chromatogram which of the liquids is present in the LARGEST quantity?
27. Which of the following atomic nuclei pairs both have nuclear spin?
(A) ${ }^{2} \mathrm{H}$ and ${ }^{13} \mathrm{C}$
(B) ${ }^{1} \mathrm{H}$ and ${ }^{24} \mathrm{Mg}$
(C) ${ }^{19} \mathrm{~F}$ and ${ }^{23} \mathrm{Na}$
(D) $\quad{ }^{32} \mathrm{P}$ and ${ }^{31} \mathrm{p}$
28. Which of the following equations governs the determination of an unknown compound, X , in a solution by UV/VIS spectroscopy?
(A) $\quad \log _{10}\left(\mathrm{I}_{\mathrm{o}} / \mathrm{I}\right)=\in \mathrm{cL}$
(B) $\quad \log _{10}\left(\mathrm{I} /{ }_{0}\right)=\mathrm{cL}$
(C) $\quad \log _{10}\left(I_{0} / I\right)=\frac{\in \mathrm{c}}{\mathrm{L}}$
(D) $\quad \log _{10}\left(\mathrm{I} / \mathrm{I}_{\mathrm{o}}\right)=\frac{\in \mathrm{C}}{\mathrm{L}}$
29. Whcih of the following are all examples of commonly used stationary phases in chromatography?
(A) Cellolose, alumina, starch
(B) Silica gel, alumina, cellulose
(C) Silica gel, cellulose, sodium chloride
(D) Alumina, silica gel, aluminium chloride
30. A solution of 5 g of an organic compound X in $200 \mathrm{~cm}^{3}$ of water is shaken with 100 cm 3 of choloroform. 1.3 g of the acid remain in the aqueous solution at equilibrium. The partition coefficient of X between water and chloroform is
(A) 3.7
6.5
(B) $\frac{3.7 \times 10^{-2}}{6.5 \times 10^{-3}}$
(C) $\quad 6.5$
3.7
(D) $\frac{6.5 \times 10^{-2}}{3.7 \times 10^{-2}}$
31. In the conversion of bauxite to pure aluminium oxide, what is the compound obtained after filtering and adding carbon dioxide to the impure bauxite?
(A) $\quad \mathrm{A} 1(\mathrm{OH})_{3}$
(B) $\mathrm{Al}(\mathrm{OH})_{4}^{-}$
(C) $\mathrm{SiO}_{3}{ }^{2-}$
(D) $\quad \mathrm{Al}_{2}\left(\mathrm{CO}_{3}\right)_{3}$
32. Crude oil is separated into petroleum products by use of fractional distillation.

Which fractions would be obtained between $20-200{ }^{\circ} \mathrm{C}$ and $275-375^{\circ} \mathrm{C}$ respectively?

$$
20-200^{\circ} \mathrm{C} \quad 275-375^{\circ} \mathrm{C}
$$

(A) Bitumen refinery gas
(B) Diesel kerosene
(C) Naphtha refinery gas
(D) Diesel gasoline
33. Purified $\mathrm{N}_{2}$ gas and $\mathrm{H}_{2}$ gas are reacted together to form ammonia as shown below.

\[

\]

Based on Le Chatelier's principle, which combination of pressure and temperature would give the highest yield of ammonia?
(A) Low temperature, low pressure
(B) Low temperature, high pressure
(C) High temperature, high pressure
(D) High temperature, low pressure
34. Ethanol formed by fermentation will destroy the enzyme-producing yeast. What is the lowest percentage of alcohol at which this occurs?
(A) 3
(B) 13
(C) 20
(D) 30
35. The oxygen gas used in the contact process must be dust free to avoid poisoning the vanadium (V) oxide catalyst. Which reaction would be affected if dust is present?
(A) $\quad \mathrm{S}(\mathrm{g})+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{SO}_{2}(\mathrm{~g})$
(B) $\quad 2 \mathrm{~S}(\mathrm{~g})+3 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{SO}(\mathrm{g})$
(C) $\quad 2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{SO}_{3}(\mathrm{~g})$
(D) $\quad 2 \mathrm{~S}(\mathrm{~g})+3 \mathrm{O}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \rightarrow$ $2 \mathrm{H}_{2} \mathrm{SO}_{4}(\mathrm{aq})$
36. What is the purpose of green chemistry?
(A) Removing wastes from industrial reactions
(B) Designing chemical products that maximize profits
(C) Inventing technologies that will promote recycling of products
(D) Designing safer chemical products and processes that reduce or eliminate the use of and generation of hazardous substances
37. A polluted river near a paint factory is tested with acidified potassium iodide and a bright yellow precipitate is observed. Which of the following pollutants is likely to be present?
(A) $\quad \mathrm{PO}_{4}{ }^{3-}$
(B) $\mathrm{CN}^{-}$
(C) $\mathrm{Cr}^{3+}$
(D) $\mathrm{Pb}^{2+}$
38. The substance formed in combustion engines of vehicles, which may lead to the formation of acid rain is
(A) C
(B) NO
(C) CO
(D) $\quad \mathrm{C}_{4} \mathrm{H}_{10}$
39. It is estimated that one chlorine radical can break down 100000 ozone molecules.

Which feature of chlorine radicals may account for this?
(A) Regeneration by reacting with ozone
(B) Ionization of oxygen gas
(C) Chlorine monoxide radicals formed with oxygen
(D) Reaction with ozone to form more oxygen gas
40. Which of the following CANNOT be used to control atmospheric pollution?
(A) Using cleaner fuels in industrial plants
(B) Using catalytic converters in vehicles
(C) Decreasing vegetation in urban areas
(D) Using chemical scrubbers
41. Water can be purified by all of the following EXCEPT
(A) desalination
(B) reverse osmosis
(C) distillation
(D) sublimation
42. The ozone layer is a thick blanket over the earth's surface. It prevents ultraviolet radiation from entering the earth's atomosphere. Where is the ozone layer located?
(A) Troposphere
(B) Stratosphere
(C) Ionosphere
(D) Mesosphere
43. Which properties of aluminium make it suitable for constructing airplanes?
I. Low density
II. Imperneable oxide
III. Conducts electricity
(A) I and II only
(B) I and III only
(C) II and III only
(D) I, II and III
44. The term 'residence time' for gases in the atmosphere refers to the length of time the gas
(A) takes to move from the earth's surface to the upper regions of the atmosphere
(B) remains in the atmosphere until it is removed in a sink
(C) takes to move from the troposphere to the stratosphere
(D) reaches a concentration to become a pollutant in the atmosphere
45. Which of the following chemicals forms a non-toxic substance when it reacts with nitrogen monoxide?
(A) $\mathrm{CH}_{4}$
(B) $\mathrm{H}_{2} \mathrm{O}$
(C) $\mathrm{SO}_{2}$
(D) CO

## END OF TEST



## CANDIDATE'S RECEIPT

## INSTRUCTIONS TO CANDIDATE:

1. Fill in all the information requested clearly in capital letters.

TEST CODE:

| 0 | 2 | 2 | 1 | 2 | 0 | 1 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

SUBJECT: CHEMISTRY - UNIT 2 - Paper 01

PROFICIENCY:
ADVANCED

REGISTRATION NUMBER:

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| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

FULL NAME: $\qquad$
(BLOCK LETTERS)

Signature: $\qquad$

Date: $\qquad$
2. Ensure that this slip is detached by the Supervisor or Invigilator and given to you when you hand in this booklet.
3. Keep it in a safe place until you have received your results.

## INSTRUCTION TO SUPERVISOR/INVIGILATOR:

Sign the declaration below, detach this slip and hand it to the candidate as his/her receipt for this booklet collected by you.

I hereby acknowledge receipt of the candidate's booklet for the examination stated above.

Signature: $\qquad$
Supervisor/Invigilator

Date: $\qquad$

Unit 1 Paper 02 Keys

| Question | Module/Syllabus Reference | Profile | Key |
| :---: | :---: | :---: | :---: |
| 1. | 1.1.1 | KC | A |
| 2. | 1.1.5 | KC | A |
| 3. | 1.1.8 | UK | C |
| 4. | 1.4.2 | KC | C |
| 5. | 1.4.3 | UK | A |
| 6. | 1.4.5 | KC | B |
| 7. | 1.2 .5 | UK | B |
| 8. | 1.2.2 | KC | D |
| 9 | 1.2.7 | UK | C |
| 10. | 1.2.1 | KC | C |
| 11. | 1.2.1 | KC | D |
| 12. | 1.2.6 | UK | B |
| 13. | 1.2.15 | UK | D |
| 14. | 1.3.1 | KC | C |
| 15. | 1.3.3 | UK | C |
| 16. | 2.1.1 | UK | C |
| 17. | 2.2.2 | KC | C |
| 18. | 2.2.3 | UK | A |
| 19. | 2.2.5 | UK | B |
| 20. | 2.3.4 | UK | C |
| 21. | 2.3.2 | KC | B |
| 22. | 2.2.4 | KC | B |
| 23. | 2.9.1 | UK | C |
| 24. | 2.6.4 | UK | A |
| 25. | 2.6.1 | KC | B |
| 26. | 2.8.5 | UK | C |
| 27. | 2.8.3 | UK | C |
| 28. | 2.5.3 | KC | A |
| 29. | 2.8.4 | KC | B |
| 30. | 2.9.4 | UK | B |
| 31. | 3.2.1 | KC | A |
| 32. | 3.3.2 | KC | D |
| 33. | 3.4.1 | UK | B |
| 34. | 3.5.1 | KC | B |
| 35. | 3.7.1 | UK | C |
| 36. | 3.1.3 | KC | D |
| 37. | 3.8.5 | UK | D |
| 38. | 3.9.6 | KC | B |
| 39. | 3.9.1 | KC | A |
| 40. | 3.9.8 | KC | C |
| 41. | 3.8.2 | KC | D |
| 42. | 3.9.2 | KC | B |
| 43. | 3.2.2 | KC | A |
| 44. | 3.9.1 | KC | B |
| 45. | 3.9.6 | KC | D |

## CARIBBEAN EXAMINATIONS COUNCIL

## CARIBBEAN ADVANCED PROFICIENCY EXAMINATION ${ }^{\circledR}$

FILL IN ALL THE INFORMATION REQUESTED CLEARLY IN CAPITAL LETTERS.

TEST CODE

| 0 | 2 | 2 | 1 | 2 | 0 | 2 | 0 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |

SUBJECT $\qquad$ CHEMISTRY - UNIT 2 - Paper 02

PROFICIENCY $\qquad$ ADVANCED REGISTRATION NUMBER


| NAME OF SCHOOL/CENTRE |
| :---: |
|  |


| CANDIDATE'S FULL NAME (FIRST, MIDDLE, LAST) |
| :---: |
|  |

DATE OF BIRTH


SIGNATURE $\qquad$

| "*"Barcode Area"*" |
| :--- |
| Sequential Bar Code |



## CARIBBEAN <br> EXAMINATIONS <br> COUNCIL

## CARIBBEAN ADVANCED PROFICIENCY EXAMINATION ${ }^{\circledR}$ <br> CHEMISTRY

## SPECIMEN PAPER

UNIT 2 - Paper 02
2 hours 30 minutes

## READ THE FOLLOWING INSTRUCTIONS CAREFULLY.

1. This paper consists of THREE questions. Answer ALL questions.
2. Write your answers in the spaces provided in this booklet.
3. Do NOT write in the margins.
4. Where appropriate, ALL WORKING MUST BE SHOWN in this booklet.
5. A data booklet is provided.
6. You may use a silent, non-programmable calculator to answer questions.
7. If you need to rewrite any answer and there is not enough space to do so on the original page, you must use the extra lined page(s) provided at the back of this booklet. Remember to draw a line through your original answer.
8. If you use the extra page(s) you MUST write the question number clearly in the box provided at the top of the extra page(s) and, where relevant, include the question part beside the answer.
dO NOT TURN THIS PAGE UNTIL YOU ARE TOLD TO DO SO.

## Answer ALL questions.

## MODULE 1

## THE CHEMISTRY OF CARBON COMPOUNDS

1. Dopamine is a neurotransmitter found in the human brain.


Figure 1. Dopamine
(a) (i) Name AND write THREE functional groups in the dopamine molecule.
$\qquad$
$\qquad$
$\qquad$
(ii) Explain whether or not the dopamine molecule would exhibit optical activity.
(b) Draw the structural formula of the organic product formed when dopamine is dissolved in excess aqueous sodium hydroxide.
(c) A sample of dopamine is first dissolved in dilute hydrochloric acid and then subjected to an electric current.
(i) Draw the structure of the organic species formed on the addition of the acid.
(i) Draw
(ii) To which electrode would the species formed be expected to migrate?
(d) Draw the fully displayed structural formula of the organic product formed when dopamine is treated with ethanol in the presence of concentrated sulphuric acid and refluxed.
(e) Dopamine is subjected to chemical analysis and some of the results are presented in Table 1 below. Complete Table 1 by writing the missing test, observation and inference.

TABLE 1: RESULTS OF CHEMICAL ANALYSIS OF DOPAMINE

| Test | Observation | Inference |
| :--- | :--- | :--- |
| $\mathrm{PCl}_{3}$ or $\mathrm{PCl}_{5}$ is added to dopamine. | • |  |
| • |  |  |
|  | White crystals are produced on <br> heating. | $\bullet$ |

[5 marks]

| "*"Barcode Area"*" |
| :---: |
| Sequential Bar Code |

(f) State THREE characteristic properties of members of a homologous series.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(g) A reaction of 2-methylpropane with chlorine in sunlight results in the production of different substituted species.
(i) By examination of the reaction mechanism, account for the production of the different monosubstituted species.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) Name the primary monosubstituted species. Justify your answer.
$\qquad$
$\qquad$
$\qquad$
$\qquad$

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(h) 2-chloro-2-methylpropane (tertiary butyl chloride) can be converted to the corresponding alcohol by reacting with aqueous sodium hydroxide. Outline the mechanism for the reaction.

## MODULE 2

## ANALYTICAL METHODS AND SEPARATION TECHNIQUES

2. The chromatogram in Figure 2 is obtained when a mixture of organic compounds, $\mathrm{A}-\mathrm{D}$ is analysed by gas-liquid chromatography.


Figure 2: Chromatogram
(a) Define EACH of the following terms:
(i) Retention time
$\qquad$
$\qquad$
(ii) Mobile phase
$\qquad$
$\qquad$

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(iii) Name TWO commonly used mobile phase in gas-liquid chromatography.
$\qquad$
$\qquad$
(b) Explain the chemical principle upon which the separation of $\mathrm{A}-\mathrm{D}$ can be achieved.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(c) (i) Which of the compounds, $\mathrm{A}-\mathrm{D}$, is present in the highest concentration?
(ii) For how long was the compound identified in (c) (i) retained on the column?
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(d) Assuming that the mixture contained methanol, methanal, propane and propanoic acid, suggest the identities of $\mathrm{A}, \mathrm{B}, \mathrm{C}, \mathrm{D}$. Give reasons for your response.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
Reasons:
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(e) Outline how a mixture of plant pigments could be separated by column chromatography.
$\qquad$
$\qquad$
$\qquad$
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Sequential Bar Code
(f) Explain the principles upon which mass spectrometry is based.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(g) An organic compound, A, contained only carbon, hydrogen and oxygen. The following mass spectrogram was obtained upon analysis.

(i) State the relative molecular mass of A.
(ii) State the name of the $\mathrm{M}^{+1}$ peak in the spectrum.

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(iii) Suggest a possible identity for EACH of the species corresponding to peaks.

B: $\qquad$

C: $\qquad$

F: $\qquad$
(iv) Calculate the relative abundance of the species corresponding to peak E.
(v) What is the significance of $100 \%$ intensity for peak F?
$\qquad$
$\qquad$
$\qquad$

## MODULE 3

## INDUSTRY AND THE ENVIRONMENT

3. (a) In industrial laboratories, the cadmium reduction method is widely used, instead of the chemical tests that are normally used in school laboratories, to determine if nitrate ions are present as a pollutant in water samples.
(i) Give TWO sources of nitrate ions in water.
$\qquad$
(ii) Describe a chemical test that is often used in school laboratories to detect the presence of aqueous nitrate ions.
$\qquad$
$\qquad$
$\qquad$
(iii) State ONE ion that cannot be distinguished from the $\mathrm{NO}_{3}^{-}$by the chemical test that you described in 3 (a) (ii) above.
$\qquad$
(iv) Suggest ONE possible reason why the laboratory test that you described in 3 (a) (ii), for the determination of nitrate ions as a pollutant in water samples, is not as appropriate as the cadmium reduction method.
$\qquad$
$\qquad$
$\qquad$
$\qquad$

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(b) Figure 3 shows the average concentration of pollutants NO and $\mathrm{NO}_{2}$ in an urban area during a 24 -hour period.


Figure 3: Concentration of NO and $\mathrm{NO}_{2}$
(i) Using the following equilibrium reaction
$\mathrm{N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \longleftrightarrow 2 \mathrm{NO}(\mathrm{g}) ; \Delta \mathrm{H}=+180 \mathrm{~kJ} \mathrm{~mol}^{-1}$ account for the formation of $\mathrm{NO}(\mathrm{g})$ as a pollutant in urban areas.
$\qquad$
$\qquad$
$\qquad$
(ii) Suggest a reason for the time lapse between maximum concentrations of $\mathrm{NO}(\mathrm{g})$ and $\mathrm{NO}_{2}(\mathrm{~g})$ shown in Figure 3.
$\qquad$
$\qquad$
$\qquad$
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(iii) Indicate, by drawing a curve on Figure 3, how you would expect the concentration of ozone $\left(\mathrm{O}_{3}\right)$ in an urban area to vary during the course of the day. Explain your answer.
$\qquad$
$\qquad$
$\qquad$
(c) The chemical equation for the formation of ammonia by the Haber Pprocess is as follows.
$\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g}) \Delta \mathrm{H}=-92 \mathrm{~kJ} / \mathrm{mol}$
(i) State the source of nitrogen and name the process by which it is obtained.
$\qquad$
$\qquad$
(ii) Use the equation above to predict the conditions of temperature and pressure which would maximize the yield of ammonia.
$\qquad$
$\qquad$
(iii) State whether the conditions used in industry are similar to those you predicted in (c) (ii). Account for any variations in these conditions.
(iv) Write a balanced chemical equation to show how hydrogen is obtained from methane, which is a source of hydrogen.
$\qquad$
$\qquad$
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(d) (i) Identify TWO Green Chemistry principles that are implemented in the manufacture of ammonia by the Haber Process.
$\qquad$
$\qquad$
(ii) Choose one of the Green Chemistry principles identified in (d) (i), and explain how that principle is adhered to in the process.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(iii) Using a named Green Chemistry principle not already utilized in the Haber Process, explain how the manufacture of ammonia may be improved.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(e) Air serves as an important chemical resource for the biosphere. State ONE process by which nitrogen in the atmosphere can enter the nitrogen cycle.
$\qquad$

## END OF TEST

## IF YOU FINISH BEFORE TIME IS CALLED, CHECK YOUR WORK ON THIS TEST.

## EXTRA SPACE

If you use this extra page, you MUST write the question number clearly in the box provided. Question No. $\square$
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## CANDIDATE'S RECEIPT

## INSTRUCTIONS TO CANDIDATE:

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SUBJECT:
CHEMISTRY - UNIT 2 - Paper 02

PROFICIENCY:
ADVANCED

REGISTRATION NUMBER: $\square$

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Supervisor/Invigilator

Date: $\qquad$

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CHEMISTRY
UNIT 2 - PAPER 02
MARK SCHEME

SPECIMEN

## Question 1

S.O.: Module: 1 -1.8, 2.1, 2.2, 2,3 2.6, 2.7, 2.13, 2.11, 2.14
(a) (i)

(ii) Yes (1) - A chiral carbon is present (1)
(b)

( 1 mark)
(c) (i)

(ii) Negative electrode or cathode (1)
(d)


| KC | UK | XS |
| :---: | :---: | :---: |
| 3 |  |  |
| 1 | 2 |  |
|  |  |  |
|  |  |  |

(e)
(g)
(h)

| (i) | Colourless gas evolves (1) which forms dense whi with $\mathrm{NH}_{3}(1)$ | te fumes <br> 2 marks] |
| :---: | :---: | :---: |
| (ii) | COOH/Carboxylic acid group present | [1 mark] |
| (iii) | Dil. HCl is added and the resulting solution gently | heated [1 mark] |
| (iv) | The $\mathrm{NH}_{2}$ and amine group present | [1 mark] |

- They show a steady gradation in physical properties
- Chemical properties are similar
- Relative molecular mass of consecutive members increase by H or $\mathrm{CH}_{2}$
[1 mark each]

$\mathrm{Cl} \bullet+\mathrm{CH}_{3} \mathrm{CHCH}_{3} \mathrm{CH}_{3} \rightarrow \mathrm{CH}_{3} \stackrel{\bullet}{\mathrm{C}} \mathrm{CH}_{3} \mathrm{CH}_{3}+\mathrm{HCl}$
$\mathrm{Cl} \bullet+\mathrm{CH}_{3} \mathrm{CHCH}_{3} \mathrm{CH}_{3} \rightarrow \dot{\mathrm{C}} \mathrm{H}_{2} \mathrm{CHCH}_{3} \mathrm{CH}_{3}+\mathrm{HCl}$
$\mathrm{CH}_{3} \dot{\mathrm{C}} \mathrm{CH}_{3} \mathrm{CH}_{3}+\dot{\mathrm{C}} 1 \rightarrow \mathrm{CH}_{3} \mathrm{CClCH}_{3} \mathrm{CH}_{3}$
[1 mark]
$\bullet \mathrm{CH}_{2} \mathrm{CHCH}_{3} \mathrm{CH}_{3}+\mathrm{Cl} \rightarrow \mathrm{CH}_{2} \mathrm{ClCHCH}_{3} \mathrm{CH}_{3}$ [1 mark]
[1 mark]
[1 mark] (propagation steps also acceptable)
(ii) $\mathrm{CH}_{3} \mathrm{CClCH}_{3} \mathrm{CH}_{3}$

Stability of the tertiary free radical in the intermediate stage [2 marks]

For 2 - methylpropene, the mechanism is electrophilic addition (1). The double bond causes polarization of the Chlorine molecule which produces the electrophile by heterolytic fission (1). The mechanism for the 2 -methyl propane is free radical substitution (1) involving homolytic fission. (1)
[4 marks]

| 10 | 15 | 5 |
| :--- | :--- | :--- |

## Question 2

S.O.: Module: $2-7.1,7.2,8.1,8.2,8.3,8.5$
(a) (i) Length of time a solute remains in the column, before entering the detector
[1 mark]
(ii) Mobile phase: The solvent or solvent mixture which moves over the stationary phase carrying the solutes with it
[1 mark]
(iii) Commonly used mobile phase in GLC: nitrogen, helium, argon. Any two
[2 marks]
(b) Partitioning [1 mark]

Substances A-D have different solubilities in the solvent being used. Based on the differences in their solubilities the substances have different speeds as they pass through the column.
[2 marks]

| KC | UK | XS |
| :---: | :---: | :---: |
|  |  |  |
|  |  |  |

(c) (i) B
[1 mark]
(ii) Approximately 6.8 minutes
0.1 minute
[1 mark]
(d) A - methanal

B - propane
C - methanol
D - propanoic acid
[1 mark each]

Methanol has lowest density
Propane is more volatile than methanol and propanoic acid Propanoic acid is the heaviest or densest
[Any 2]
(e)

- Pack a glass column with a paste of $\mathrm{SiO}_{2}, \mathrm{CaCO}_{3}$ or $\mathrm{Al}_{2} \mathrm{O}^{3}$ Ensure there are no air spaces.
- Using a Pasteur pipette, place about $5 \mathrm{~cm}^{3}$ of liquid plant pigment mixture in the column.
- Add the solvent (water) slowly to the mixture ensuring that the paste is not disturbed.
- Open the tap at the base of the column and collect fractions of equal volumes into separate conical flasks.
- Evaporate contents of flask and analyse via TLC.
(f) - Vapourised sample may be broken into positively charged fragments by bombardment with high-speed electrons.
- Fragments are deflected in a magnetic field, proportionate to the mass/charge ratio.
- These fragments can be used to deduce the structure of the original molecule.
[1 mark each]
(g)
(i) 46
[1 mark]
(ii) J
(iii) B: $\mathrm{CH}_{3}$
$\mathrm{C}: \mathrm{OH}$
$\mathrm{F}: \mathrm{CH}_{2} \mathrm{OH}$
(iv) $\frac{3}{7} \times 100 \%=43 \%$
[3 marks]
(v) Most stable fragment
[1 mark]

Total 30 marks


## Question 3

S.O: Module 3: - $1.3,4.1,8.4,8.5,9.6,9.7$
(a) (i) Fertilisers, decaying plants or animals
(ii) Add aluminum metal or zinc metal followed by sodium hydroxide solution and warm. If nitrate ions are present, ammonia gas is produced OR Add copper turnings followed by conc. Sulphuric acid poured carefully down the side of the testy tube. If nitrate ions are present a brown ring is produced. [2 marks]
(iii) $\mathrm{NO}_{2}{ }^{-}$
[1 mark]
(iv) Pollutants are usually present in trace (small) quantities. School laboratory tests are not As sensitive as the cadmium reduction method to small quantities of $\mathrm{NO}_{3}{ }^{-}$.
[2 marks]
(b) (i) In urban centres, there is a heavy flow of traffic. The burning of fuels in the internal combustion engine is an exothermic reaction, and the high temperatures produced provide the conditions that favour the forward endothermic reaction for the production of NO from nitrogen and oxygen that are present in the air.
[2 marks]
(ii) NO is produced first as the primary pollutant after which $\mathrm{NO}_{2}$ is formed by the oxidation of NO . [1 mark]
(iii)


The ozone concentration begins to increase after $\mathrm{NO}_{2}$ has been formed. If $\mathrm{NO}_{2}$ absorbs a quantum of light the molecule can decompose into NO and atomic oxygen, a radical. The very reactive radical can then react with molecular oxygen to form ozone. The concentration of the ozone falls as it reacts with NO to re-form $\mathrm{NO}_{2}$ and molecular oxygen.
[2 marks] [1 mark for graph]


## CHEMISTRY

UNIT 2 - PAPER 02
MARK SCHEME
(c) (i) Air; obtained by liquefication of air
(ii) Conditions for maximum yield:

High pressures and low temperatures
[2 marks]
(iii) Conditions used in industry:

No, a set of compromise conditions are used
Award marks based on a discussion of the applications of the principles of equilibrium coupled with economic factors.
[2 marks]
(iv) Hydrogen from Methane
$\mathrm{CH}_{4}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{CO}+3 \mathrm{H}_{2}$
Accept also
$\mathrm{CH}_{4}+2 \mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{CO}_{2}+4 \mathrm{H}_{2}$
$\mathrm{CO}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{CO}_{2}+\mathrm{H}_{2}$
[2 marks]
(d) (i) Prevent waste

Use catalysts not the stoichiometric reagents
Maximise atom efficiency
[2 marks]
(ii) Prevent waste: the raw materials, nitrogen gas and hydrogen gas are recycled through the reactor. In the reactor only $15 \%$ is converted to ammonia, by continued recycling of the reactants there is $98 \%$ conversion of the reactants very little is wasted by recycling the reactants there is very little waste generated in the Haber process.

Use of catalysts: the iron catalyst (aided by the potassium hydroxide as a promoter overcomes the inefficiency in the stoichiometric equation. The reaction is favored by low temperature but the rate of the reaction slows down significantly under these conditions. The catalyst speeds up the reaction thereby overcoming this.

Maximize atom efficiency: the synthesis of the ammonia requires just the materials (elements) that are in the raw material. There are no side reactions or loss of elements during the reaction. 98\% of the $N_{2}$ and $H_{2}$ end up in the final $\mathrm{NH}_{3}$ molecule.
(iii) Use renewable feedstocks: the $H_{2}$ currently is obtained from natural gas which is a non-renewable resource. The hydrogen can be obtained from a renewable/alternative source. For example, Electrolysis of water using electricity from hydroelectric or some other renewable source of energy. Or any other reliable process.
(e)

Nitrogen fixation
Electrical discharges (thunderstorms)
Accept also combustion of fuels from aircrafts
[1 mark]
TOTAL 30 marks


CHEMISTRY

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FILL IN ALL THE INFORMATION REQUESTED CLEARLY IN CAPITAL LETTERS.

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SUBJECT $\qquad$ CHEMISTRY - UNIT 2 - Paper 032

PROFICIENCY $\qquad$ ADVANCED REGISTRATION NUMBER
"*"Barcode Area"

## SPECIMEN PAPER

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| NAME OF SCHOOL/CENTRE |
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## CARIBBEAN ADVANCED PROFICIENCY EXAMINATION ${ }^{\circledR}$ <br> CHEMISTRY

SPECIMEN PAPER
UNIT 2 - Paper 032
2 hours

## READ THE FOLLOWING INSTRUCTIONS CAREFULLY.

1. This paper consists of THREE questions. Answer ALL questions.
2. Write your answers in the spaces provided in this booklet.
3. Do NOT write in the margins.
4. Where appropriate, ALL WORKING MUST BE SHOWN in this booklet.
5. A data booklet is provided.
6. You may use a silent, non-programmable calculator to answer questions.
7. If you need to rewrite any answer and there is not enough space to do so on the original page, you must use the extra lined page(s) provided at the back of this booklet. Remember to draw a line through your original answer.
8. If you use the extra page(s) you MUST write the question number clearly in the box provided at the top of the extra page(s) and, where relevant, include the question part beside the answer.
dO NOT TURN THIS PAGE UNTIL YOU ARE TOLD TO DO SO.

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## Answer ALL questions.

1. You are provided with Sample A, which is a mixture of fine sand and $\mathrm{CuSO}_{4} \cdot \mathrm{nH}_{2} \mathrm{O}$ crystals in a crucible. You are required to determine the value of $n$ by following the procedure outlined below and recording your data in Table 1.

## (a) Procedure

A. Weigh the crucible and Sample A (M1).
B. Heat the sample over a bunsen burner for approximately 15 minutes.

Record any changes in appearance of the sample.
C. Remove the sample from the heat, cool in a dessicator for five minutes and reweigh the crucible and sample (M2).
D. Remove the sample as completely as possible in $250 \mathrm{~cm}^{3}$ of water. Record your observation.
E. Filter the mixture from Step D above through Whatman \#4 filter paper. Wash and dry the crucible.
F. Wash the residue in the filter paper with two separate $100 \mathrm{~cm}^{3}$ aliquots of water. Record the colour of the filtrate from each of these washings.
G. Transfer the washed residue to the previously dried crucible.
H. Dry the transferred residue over a bunsen flame. Record your observations.
I. Cool the dried residue in a dessicator for five minutes.
J. Weigh the dried cool residue in the crucible (M3)
(b) Outline the procedure that would have been followed to obtain the results in (a).
$\qquad$
$\qquad$
$\qquad$
$\qquad$

TABLE 1: DATA FOR EXPERIMENTAL PROCEDURE

| STEP | DATA |
| :---: | :---: |
| A | Initial mass of Sample A in crucible, M1... |
| B | Change in appearance of Sample A. ........................... |
| C | Mass of Sample A in crucible after the first heating process, M2 |
| D | Colour of mixture. ..... |
| E | Colour of filtrate obtained from washing residue: <br> first washing $\qquad$ <br> second washing $\qquad$ |
| F | Change in appearance of sand |
| G | Mass of dried residue and crucible, M3........................ |

(b) Determine EACH of the following:
(i) The mass of the water of crystallization in $\mathrm{CuSO}_{4} \cdot \mathrm{n}_{2} \mathrm{O}$
$\qquad$
$\qquad$
$\qquad$
(ii) The mass of anhydrous copper sulphate
$\qquad$
$\qquad$
$\qquad$
(iii) The value of $n$
$\qquad$
$\qquad$
$\qquad$

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2. After being opened and used, a bottle of $A R$ iron (II) sulphate was left on the laboratory shelf for some time. 10 g of the sample were then dissolved in some dilute suphuric acid and $1 \mathrm{dm}^{3}$ of solution was prepared. $20 \mathrm{~cm}^{3}$ of this solution was placed in a conical flask and titrated with $0.02 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{KMnO}_{4}(\mathrm{aq}) / \mathrm{H}^{+}(\mathrm{aq})$. The ionic equation for the reaction is $5 \mathrm{Fe}^{2+-}(\mathrm{aq})+\mathrm{MnO}_{4-}(\mathrm{aq})+8 \mathrm{H}^{+}(\mathrm{aq}) \rightarrow 5 \mathrm{Fe}^{3+}(\mathrm{aq})+\mathrm{Mn}^{2+}(\mathrm{aq})+4 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$.

Figure 1 below shows the readings on the burette before and after each titration.


Figure 1. Readings on the burette
(a) (i) What colour change would you expect at the end point of the reaction?
$\qquad$
(ii) Complete Table 2 to record the tritration results. You should include the initial and final burette readings and the volumes of $\mathrm{KMnO}_{4}(\mathrm{aq}) / \mathrm{H}^{+}(\mathrm{aq})$ used.

TABLE 2: TITRATION RESULTS

|  | $\mathbf{1}$ | $\mathbf{2}$ | $\mathbf{3}$ |
| :--- | :--- | :--- | :--- |
| Final burette reading $\left(\mathrm{cm}^{3}\right)$ |  |  |  |
| Initial burette reading $\left(\mathrm{cm}^{3}\right)$ |  |  |  |
| Volume of $\mathrm{KMnO}_{4}$ used $\left(\mathrm{cm}^{3}\right)$ |  |  |  |

[4 marks]

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(iii) What is the volume of $\mathrm{KMnO}_{4}(\mathrm{aq}) / \mathrm{H}^{+}(\mathrm{aq})$ to be used for calculation?
$\qquad$
$\qquad$
(v) Calculate the concentration of the $\mathrm{Fe}^{2+}$ ions in the given solution in $\mathrm{g} \mathrm{dm}^{-3}$.
$\qquad$
$\qquad$
(vi) Calculate the percentage purity of the sample.
$\qquad$
$\qquad$
$\qquad$
(b) Outline the steps that you would take to prepare $1 \mathrm{dm}^{3}$ of the iron (II) sulphate solution.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(c) State why it is not usually good practice to store iron (II) sulphate for extended periods of time after its bottle has been opened.
$\qquad$
$\qquad$
$\qquad$
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3. Plan and design an experiment to determine whether the protein in beef and chicken contains the same amino acids.

Your answer should include the following:
(i) Hypothesis
$\qquad$
$\qquad$
(ii) Aim
$\qquad$
$\qquad$
(iii) Procedure
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(iv) Variables to be controlled
$\qquad$
$\qquad$
(v) Variables to be manipulated and responding variables
$\qquad$
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(vi) Discussion of results
$\qquad$
$\qquad$
$\qquad$
(vii) TWO possible sources of error
$\qquad$
$\qquad$
$\qquad$
(viii) TWO precautions
$\qquad$
$\qquad$
$\qquad$

Total 15 marks

## END OF TEST <br> IF YOU FINISH BEFORE TIME IS CALLED, CHECK YOUR WORK ON THIS TEST.

## EXTRA SPACE

If you use this extra page, you MUST write the question number clearly in the box provided. Question No. $\square$
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## CANDIDATE'S RECEIPT

## INSTRUCTIONS TO CANDIDATE:

1. Fill in all the information requested clearly in capital letters.

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SUBJECT:
CHEMISTRY - UNIT 2 - Paper 032

PROFICIENCY:
ADVANCED

REGISTRATION NUMBER:

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CHEMISTRY
UNIT 2 - PAPER 032
MARK SCHEME
SPECIMEN

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UNIT 2 - PAPER 032
MARK SCHEME
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## Question 1

S.O.: Module: $2-3.3$
(a)

| Data |  |
| :--- | :--- |
| Candidate's reading for $M_{1}$. | [1 mark] |
| Colour changes from blue/green to white. | [2 marks] |
| Candidate's reading for $M_{2}$. | [1 mark] |
| Blue/green mixture. | [1 mark] |
| First washing: light blue/green | [1 mark] |
| Second washing: colourless |  |
| Candidate's reading for $M_{3}$. | [1 mark] |
| 2 marks for 3 correct units |  |
| [1 mark for 1 - 2 correct units] | [2 marks] |


| KC | UK | XS |
| :---: | :---: | :---: |
|  |  |  |

[1 mark]
[1 mark]
$\frac{M_{2}-M_{3}}{63.5}=Q$
[1 mark]
$\frac{\mathrm{P}}{\mathrm{Q}} / \frac{\mathrm{Q}}{\mathrm{P}}$
[2 marks]

$$
\begin{aligned}
& \text { (b) } \begin{array}{l}
\text { (i) } M_{1}-M_{2}= \\
\text { (ii) } M_{2}-M_{3}= \\
\text { (iii) } \frac{M_{1}-M_{2}}{18}=P
\end{array}=P \text { (in } \quad l
\end{aligned}
$$

[1 mark]
[1 mark]
[1 mark]

```
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```

(a)

## Question 2

S.O.: Module: 2 -2.4, 2.5
(i) Pale yellow $\rightarrow$ pink tinge
(ii)

| Burette readings $\mathrm{KMnO}_{4} / \mathrm{H}^{+}$ |  |  |  |
| :--- | :---: | :---: | :---: |
| Reading/cm ${ }^{3}$ | $\mathbf{1}$ | $\mathbf{2}$ | 3 |
| Final | 11.50 | 17.20 | 31.50 |
| Initial | 0.50 | 6.70 | 20.90 |
| Difference | 11.00 | 10.50 | 10.60 |

Correct burette readings $=$ [4 marks]

$$
\begin{array}{lll}
\text { (If one incorrect } & = & 3 \text { marks) } \\
\text { (If two incorrect } & = & 2 \text { marks) } \\
\text { (If THREE incorrect } & =1 \text { mark) } \\
\text { (More than THREE incorrect } & =0 \text { marks) } \\
\text { Volumes recorded to } 2 \text { decimal places }
\end{array}
$$

(iii) $\frac{10.5+10.6}{2}=10.55 \mathrm{~cm}^{3}$
[1 mark]
(iv) Conc. of $\mathrm{MnO}_{4}^{-}(\mathrm{aq})=0.02 \mathrm{~mol} \mathrm{dm}{ }^{-3}$

No. of moles of $\mathrm{MnO}_{4}^{-}=\left(10.55 \times 0.02 \times 10^{-3}\right)$
No. of moles of $\mathrm{Fe}^{2+}(\mathrm{aq})=\left(5 \mathrm{x} 10.55 \mathrm{x} 0.02 \mathrm{x} 10^{-3}\right)$
[1 mark]

No. of moles of $\mathrm{Fe}^{2+}(\mathrm{aq})$ in $1 \mathrm{dm}^{3}$

$$
\begin{aligned}
& =\frac{5 \times 10.55 \times 0.02 \times 10^{-3}}{20} \\
& =\quad 5.275 \times 10^{-2} \mathrm{~mol} \mathrm{dm}
\end{aligned} \quad[1 \text { mark }] \quad . \quad\left[\begin{array}{l}
-3
\end{array}\right.
$$

Conc. of $\mathrm{Fe}^{2+}=\left(5.275 \times 10^{-2} \times 56\right)=2.95 \mathrm{~g} \mathrm{dm}^{-3} \quad[1$ mark]

(v) \% purity $=\left(\frac{2.95}{10} \mathrm{x} 100\right)=29.5 \%$

## Steps

Dissolve the $10 g$ of $\mathrm{FeSO}_{4}$ in minimum volume of $\mathrm{H}_{2} \mathrm{SO}_{4}$ in a beaker.
Transfer quantitatively to a $1 \mathrm{dm}^{3}$ volumetric flask.
(b) Make up to mark with distilled water.

Stopper and shake/invert to ensure thorough mixing.
[4 marks]

It is oxidised in air from $\mathrm{Fe}^{2+}$ to $\mathrm{Fe}^{3+}$.
[2 marks]
(C)

Question 3
S.O.: Module: 1 -3.3
(i) Hypothesis: For example, chromatograms will be identical/different from beef and chicken.
(ii) Aim
[1 mark]
(iii) Apparatus and materials (Subtract one mark if fume hood missing)
[2 marks]
(iv) Procedure

- Hydrolyse with dilute acid
- At least 30 mins
- Use of visualising agent
- Use of appropriate solvent
- Separate using chromatography
- Compare 2 chromatograms
[2 marks]
(v) Variables to control: amount of each protein, time of hydrolysis, Vol. acid.
[2 marks]
(vi) Variables to be manipulated: proteins
[1 mark]
(vii) Responding variable
[1 mark]
(viii) Discussion of results as it relates to Hypothesis: compare $R_{f}$ values of different components of the 2 proteins.
(x) Possible sources error:

For example: proteins could be incompletely hydrolysed - introduction of extraneous protein from improper handling.
[2 marks]
(xi) Precautions

For example: use of fume hood; be careful not to touch chromatography paper with fingers; introduce protein to sample, etc.
[2 marks]



[^0]:    *due to stretching vibrations

