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PHYSICAL SCIENCES

PAPER 2 (CHEMISTRY)

GRADE 12

TERMS & DEFINITIONS

QUESTIONS & ANSWERS PER TOPIC

2017



This document consists of 114 pages.

Terms, definitions, questions and answers



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HOW TO USE THIS DOCUMENT

Dear Grade 12 learner

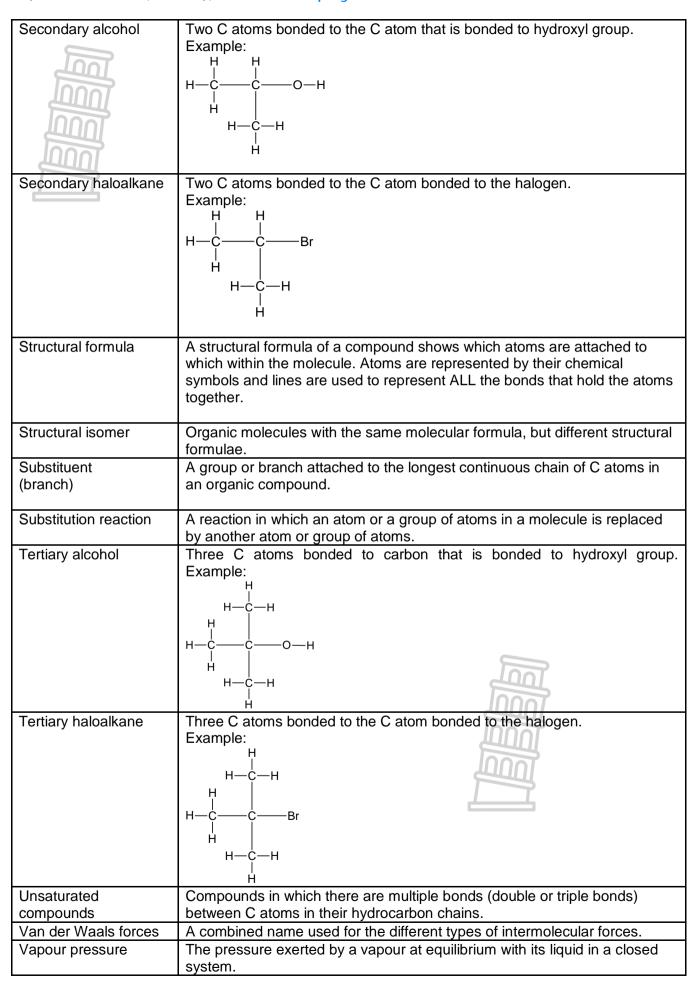
- 1. This document was compiled as an extra resource to help you to perform well in Physical Sciences.
- 2. Firstly you must make sure that you study the terms and definitions provided for each topic. Theory always forms part of any test or examination and you should ensure that you obtain full marks for ALL theory questions. Always be prepared to write a test on terms and definitions as soon as a topic is completed in class. Revise terms and definitions of topics already completed frequently so that you know them by the time you are sitting for a test or an examination.
- 3. Answer all the questions on a certain topic in your homework book as soon as the topic is completed. DO NOT look at the answers before attempting the questions. First try it yourself. Compare your answers with the answers at the back of the document. Mark your work with a pencil and do corrections for your incorrect answers. If you do not know how to answer a question, the answers are there to guide you. Acquaint yourself with the way in which a particular type of question should be answered. Answers supplied are from memoranda used to mark the questions in previous years.
- 4. Your teacher can, for example, give you two of the questions in this document as homework. The following day he/she will just check whether you answered them and whether you marked your answers. The teacher will only discuss those questions in which you do not understand the answers supplied in the document. Therefore a lot of time will be saved.
- 5. You are probably thinking about the point behind the answers at the back of the document. It is intended to help you to prepare for your tests and examinations. If you choose to copy answers into your homework book without trying them out yourself, you will be the losing party in the end! Not your teacher or anybody else!
- 6. Your teacher can also decide to give you a test on one of the questions given for homework. If you just copied the answers without any understanding, surely he/she will catch you! None of us want to be branded as dishonest, do we?
- 7. Work through all the questions and answers of a particular topic before you sit for an examination, even if you answered the questions before.
- 8. Any additional resource is only of help when used correctly. Ensure that you make use of all help provided in the correct way to enable you to be successful. All the best for 2017 and may you perform very well in Physical Sciences.

TERMS AND DEFINITIONS

MA MA	TTER AND MATERIALS: ORGANIC MOLECULES
Addition reaction	A reaction in which a double bond in the starting material is broken and elements are added to it.
Addition polymer	A polymer formed when monomers (usually containing a double bond) combine through an addition reaction.
Addition polymerisation	A reaction in which small molecules join to form very large molecules by adding on double bonds.
Alcohol	An organic compound in which H atoms in an alkane have been substituted with hydroxyl groups (-OH groups). General formula: $C_nH_{2n+1}OH$
Aldehydes	Organic compounds having the general structure RCHO where R = H or alkyl. General formula: RCHO (R = alkyl group)
Alkane	An organic compound containing only C-H and C-C single bonds. General formula: C_nH_{2n+2}
Alkene	A compound of carbon and hydrogen that contains a carbon-carbon double bond. General formula: C_nH_{2n}
Alkyl group	A group formed by removing one H atom from an alkane.
Alkyne	A compound of carbon and hydrogen that contains a carbon-carbon triple bond.
Boiling point	The temperature at which the vapour pressure of a liquid equals atmospheric pressure.
Carbonyl group	Functional group of ketones (>C=O)
Carboxyl group	Functional group of carboxylic acids (-COOH)
Carboxylic acid	An organic compound containing a carboxyl group (-COOH group). General formula: $C_nH_{2n+1}COOH$ (or RCOOH)
Chain isomers	Compounds with the same molecular formula, but different types of chains.
Condensation polymer	A polymer formed by two monomers with different functional groups that are linked together in a condensation reaction in which a small molecule, usually water, is lost.
Condensation polymerisation	Molecules of two monomers with different functional groups undergo condensation reactions with the loss of small molecules, usually water
Condensed structural formula	A formula that shows the way in which atoms are bonded together in the molecule, but DOES NOT SHOW ALL bond lines.
Cracking	The chemical process in which longer chain hydrocarbon molecules are broken down to shorter more useful molecules.
Dehydration	Elimination of water from a compound usually such as an alcohol.
Dehydrohalogenation	The elimination of hydrogen and a halogen from a haloalkane.
Dipole-dipole force	Intermolecular forces found between polar molecules i.e. molecules in which there is an uneven distribution of charge so that the molecule has a positive and a negative side.
Elimination reaction	A reaction in which elements of the starting material are "lost" and a double bond is formed.
Esterification	The preparation of an ester from the reaction of a carboxylic acid with an alcohol.
Functional group	A bond or an atom or a group of atoms that determine(s) the physical and chemical properties of a group of organic compounds.
Functional isomers	Compounds with the same molecular formula, but different functional groups.
Haloalkane (Alkyl halide)	An organic compound in which one or more H atoms in an alkane have been replaced with halogen atoms. General formula: $C_nH_{2n+1}X$ (X = F, C ℓ , Br or I)

Halogenation	The reaction of a halogen (Br_2 , Cl2) with a compound.
Homologous series	A series of organic compounds that can be described by the same general formula.
	OR A series of organic compounds in which one member differs from the next with a CH_2 group.
Hydration	The addition of water to a compound.
Hydrocarbon	Organic compounds that consist of hydrogen and carbon only.
Hydrogenation	The addition of hydrogen to an alkene
Hydrogen bond	A strong intermolecular force found between molecules in which an H atom is covalently bonded to wither an N, O or F atom.
Hydrohalogenation	The addition of a hydrogen halide to an alkene.
Hydrolysis	The reaction of a compound with water.
Intermolecular force	Forces between molecules that determine physical properties of compounds.
IUPAC name	A chemical nomenclature (set of rules) created and developed by the International Union of Pure and Applied Chemistry (IUPAC) to generate systematic names for chemical compounds.
London force	A weak intermolecular force between non-polar molecules.
Macromolecule	A molecule that consists of a large number of atoms.
Melting point	The temperature at which the solid and liquid phases of a substance are at equilibrium.
Molecular formula	A chemical formula that indicates the type of atoms and the correct number of each in a molecule, e.g. CH_4 .
Monomer	Small organic molecules that can be covalently bonded to each other in a repeating pattern.
Organic chemistry	Chemistry of carbon compounds.
Polymer	A large molecule composed of smaller monomer units covalently bonded to each other in a repeating pattern
Polymerisation	A chemical reaction in which monomer molecules join to form a polymer
Positional isomer	Compounds with the same molecular formula, but different positions of the side chain, substituents or functional groups on the parent chain.
Primary alcohol	One C atom is bonded to the C atom bonded to hydroxyl group. Example: H H H-C-C-O-H H H H H
Primary haloalkane	One C atom is bonded to the C atom bonded to the halogen. Example: H H H—C——C——Br H H H H
Saturated compounds	Compounds in which there are no multiple bonds between C atoms in their hydrocarbon chains. OR Compounds with only single bonds between C atoms in their hydrocarbon chains.

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CHEMICAL	CHANGE: QUANTITATIVE ASPECTS OF CHEMICAL CHANGE
Mole	One mole of a substance is the amount of substance having the same number of particles as there are atoms in 12 g carbon-12.
Molar gas volume at STP	The volume of one mole of a gas. (1 mole of any gas occupies 22,4 dm ³ at 0 °C (273 K) and 1 atmosphere (101,3 kPa).
Molar mass	The mass of one mole of a substance. Symbol: M Unit: $g \cdot mol^{-1}$
Avogadro's Law	Under the same conditions of temperature and pressure, the same number of moles of all gases occupy the same volume.
Concentration	The amount of solute per litre/cubic decimeter of solution.
	In symbols: $c = \frac{n}{V}$ Unit: mol·dm ⁻³
Empirical formula	The simplest positive integer ratio of atoms present in a compound.
Percentage yield	Yield is the amount of product obtained from a reaction. percentage yield = $\frac{\text{actual mass obtained}}{\text{calculated mass}} \times 100$
Percentage purity	percentage purity = $\frac{\text{mass of pure chemical}}{\text{total mass of sample}} \times 100$
Percentage composition	The percentage of each of the components in a substance. Percentage of component = $\frac{\text{mass contribute d by component}}{\text{mass of all components}} \times 100$
Limiting reagents	The substance that is totally consumed when the chemical reaction is complete.

	CHEMICAL CHANGE: ENERGY AND CHANGE
Heat of reaction (ΔH)	The energy absorbed or released in a chemical reaction.
Exothermic reactions	Reactions that release energy. ($\Delta H < 0$)
Endothermic reactions	Reactions that absorb energy. ($\Delta H > 0$)
Activation energy	The minimum energy needed for a reaction to take place.
Activated complex	The unstable transition state from reactants to products.

	CHEMICAL CHANGE: RATE OF REACTION
Reaction rate	The change in concentration of reactants or products per unit time.
	Rate = $\frac{\Delta c}{\Delta t}$ Unit: mol·dm ⁻³ ·s ⁻¹
Collision theory	a model that explains reaction rate as the result of particles colliding with a certain minimum energy.
Catalyst	A substance that increases the rate of a chemical reaction without itself undergoing a permanent change. (A catalyst increases the rate of a reaction by providing an alternative path of lower activation energy. It therefore decreases the net/total activation energy.)
Factors that affect reaction rate	Nature of reacting substances, surface area, concentration (pressure for gases), temperature and the presence of a catalyst.

	CHEMICAL CHANGE: CHEMCIAL EQUILIBRIUM
Open system	A system which continuously interacts with the environment – it exchanges matter and energy with its environment.
Closed system	A system that only exchanges energy with its surroundings, but it does not exchange matter with its surroundings.
Reversible reaction	A reaction is reversible when products can be converted back to reactants.
Chemical equilibrium	Dynamic equilibrium when the rate of the forward reaction equals the rate of the reverse reaction.
Factors that influence the equilibrium position	Pressure (gases only), concentration and temperature.
Le Chatelier's principle	When the equilibrium in a closed system is disturbed, the system will re- instate a new equilibrium by favouring the reaction that will oppose the disturbance.

	CHEMICAL CHANGE: ACIDS AND BASES
Acid-base indicator	A dye used to distinguish between acidic and basic solutions by means of
	the colour changes it undergoes in these solutions.
Amphiprotic	A substance that can act as either an acid or a base.
substance/ampholyte	
Arrhenius theory	An acid is a substance that produces hydrogen ions (H ⁺)/ hydronium ions
	(H_3O^+) when it dissolves in water.
	A base is a substance that produces hydroxide ions (OH) when it dissolves
	in water.
Auto-ionisation of water	A reaction in which water reacts with itself to form ions (hydronium ions and hydroxide ions).
Concentrated acids/bases	Contain a large amount (number of moles) of acid/base in proportion to the volume of water.
Conjugate acid-base	A pair of compounds or ions that differ by the presence of one H^+ ion.
pair	Example: CO_3^2 and HCO_3^2 OR HCl and Cl^2
Conjugate acid and	A conjugate acid has one H^+ ion more than its conjugate base.
base	Example: HCO_3^{-1} is the conjugate acid of base $CO_3^{2^{-1}}$.
	5
	CO_3^{2-} is the conjugate base of acid HCO_3^{-} .
Dilute acids/bases	Contain a small amount (number of moles) of acid/base in proportion to the volume of water.
Diprotic acid	An acid that can donate two protons. Example: H_2SO_4
Dissociation	The process in which ionic compounds split into ions.
Endpoint	The point in a titration where the indicator changes colour.
Equivalence point	The point in a reaction where equivalent amounts of acid and base have reacted completely.
Hydrolysis	The reaction of a salt with water.
	The reaction of an ion with water to produce the conjugate acid and a
	hydroxide ion or the conjugate base and a hydronium ion.
Ionisation	The process in which ions are formed during a chemical reaction.
lon product of water	The product of the ions formed during auto-ionisation of water i.e. $[H_3O^+][OH^-]$ at 25 °C.
lonisation constant of water (K_w)	The equilibrium value of the ion product $[H_3O^+][OH^-]$ at 25 °C.
K _a value	Ionisation constant for an acid.
K _b value	Dissociation or ionisation constant for a base.

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Lowry-Brønsted theory	An acid is a proton (H^+ ion) donor.
	A base is a proton (H ⁺ ion) acceptor.
Monoprotic acid	An acid that can donate one proton.
	Example: HCl
Neutralisation	The reaction of an acid with a base to form a salt (ionic compound) and water.
pH	The negative of the logarithm of the hydronium ion concentration in mol dm ⁻³ .
	In symbols: $pH = -log[H_3O^+]$ Unit: None
pH scale	A scale from $0 - 14$ used as a measure of the acidity and basicity of solutions where pH = 7 is neutral, pH > 7 is basic and pH < 7 is acidic.
Salt	The ionic compound that is the product of a neutralisation reaction.
Standard solution	A solution of precisely known concentration.
Strong bases	Dissociate completely in water to form a high concentration of OH ⁻ ions. Examples: sodium hydroxide (NaOH) and potassium hydroxide (KOH)
Strong acids	Ionise completely in water to form a high concentration of H_3O^+ ions. Examples: hydrochloric acid (HCl), sulphuric acid (H_2SO_4) and nitric acid (HNO ₃)
Titration	The procedure for determining the amount of acid (or base) in a solution by determining the volume of base (or acid) of known concentration that will completely react with it.
Weak acids	Ionise incompletely in water to form a low concentration of H_3O^+ ions. Examples: ethanoic acid (CH ₃ COOH) and oxalic acid (COOH) ₂
Weak bases	Dissociate/ionise incompletely in water to form a low concentration of OH ⁻ ions. Examples: ammonia (NH ₃), sodium hydrogen carbonate (NaHCO ₃), sodium carbonate (Na ₂ CO ₃), potassium carbonate (K ₂ CO ₃), calcium carbonate (CaCO ₃)

CHEMICAL CHANGE: ELECTROCHEMICAL REACTIONS	
Galvanic cell	A cell in which chemical energy is converted into electrical energy. A galvanic (voltaic) cell has self-sustaining electrode reactions.
Electrolytic cell	A cell in which electrical energy is converted into chemical energy.
Redox reaction	A reaction in which an electron transfer takes place.
Oxidation	A loss of electrons./An increase in oxidation number.
Reduction	A gain of electrons./A decrease in oxidation number.
Oxidising agent	A substance that is reduced/gains electrons/whose oxidation number decreases.
Reducing agent	A substance that is oxidised/loses electrons/whose oxidation number increases.
Anode	The electrode where oxidation takes place.
Cathode	The electrode where reduction takes place.
Electrolyte	A solution that conducts electricity through the movement of ions.
Electrolysis	The chemical process in which electrical energy is converted to chemical energy OR the use of electrical energy to produce a chemical change.
Salt bridge	The connection between two half-cells needed to ensure electrical neutrality in the cell. OR A component used in a galvanic cell to complete the circuit.
Electrodes	An electrical conductor used in a galvanic cell to make contact with a non- metallic part of the circuit e.g. the electrolyte.

Cell notation	A short way to represent a galvanic cell.
	 When writing cell notation, the following convention should be used: The H₂ H⁺ half-cell is treated just like any other half-cell. Cell terminals (electrodes) are written on the outside of the cell notation. Active electrodes: reducing agent oxidised species oxidising agent reduced species Inert electrodes (usually Pt or C): Pt reducing agent oxidised species oxidising agent reduced species Pt Example: Pt Cl⁻(aq) Cl₂(g) F₂(g) F⁻(aq) Pt
Overall cell reaction	The reaction obtained by combining two half-reactions.
Positive value of the standard emf	The reaction is spontaneous under standard conditions.
Standard conditions for a galvanic cell	Temperature: 25 °C / 298 K Concentration: 1 mol·dm ⁻³ Pressure (gases only): 101,3 kPa / 1 atmosphere
Standard hydrogen electrode	The reference electrode used to compile the Table of Standard Reduction Potentials. The hydrogen half-cell was given a standard reduction potential of 0 V. Half-cell notation: Pt $H_2(g)$ $H^+(aq)$ Half-reaction: $2H^+ + 2e^- \Rightarrow H_2$
Electroplating	The covering of an object with a metal by making it the cathode in an electrolytic cell.
Bauxite	The ore from which aluminium is recovered.
Cryolite	An aluminium compound in which aluminium oxide is dissolved to reduce the cost of the extraction of aluminium. Cryolite has a lower melting point than aluminium oxide.

CHEMICAL SYSTEMS: FERTILISERS	
Non-mineral nutrients	C, H and O
in plants	Obtained from the atmosphere (CO_2) and rain (H_2O)
Primary nutrients for plants	Nitrogen (N), phosphorous (P) and potassium (K)
	Function:
	N: essential for plant growth, green leaves
	P: development of roots, stems and seeds
	K: resistance against diseases and production of flowers/fruits
NPK fertilisers	Fertilisers containing the three primary nutrients for plants. They contain ammonium nitrate (NH_4NO_3), ammonium phosphate [(NH_4) ₃ PO ₄] and potassium chloride (KC ℓ).
N : P : K ratio	The percentage nitrogen, phosphorous and potassium in a fertiliser.
Fractional distillation of liquid air	Industrial preparation of nitrogen from air.
Steam reforming	Preparation of hydrogen from earth gas (methane)
	Reaction: $CH_4 + H_2O \rightarrow 3H_2 + CO$
Haber process	Industrial preparation of ammonia
	Reaction: $N_2 + 3H_2 \Rightarrow 2NH_3$
	Iron catalyst

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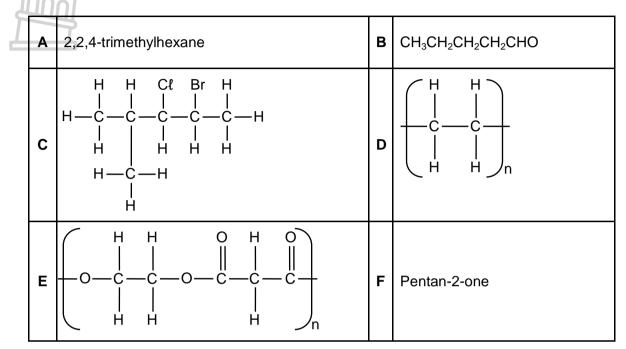
Ostwald process	Industrial preparation of nitric acid
	Reactions: Catalytic oxidation of ammonia; catalyst: Pt $4NH_3 + 5O_2 = 4NO + 6H_2O$
	$2NO + O_2 \rightarrow 2NO_2$ $3NO_2 + H_2O \rightarrow 2HNO_3 + NO OR 4NO_2 + 2H_2O + O_2 \rightarrow 4HNO_3$
Contact process	Industrial preparation of sulphuric acid Reactions: $S + O_2 \rightarrow SO_2$
	$2SO_2 + O_2 \rightarrow 2SO_3$ Contact catalyst: vanadium pentoxide (V ₂ O ₅)
	$SO_3 + H_2SO_4 \rightarrow H_2S_2O_7$ (H ₂ S ₂ O ₇ : fuming sulphuric acid OR pyro sulphuric acid OR oleum)
	$H_2S_2O_7 + H_2O \rightarrow 2H_2SO_4$
Preparation of ammonium sulphate	ammonia + sulphuric acid \rightarrow ammonium sulphate 2NH ₃ + H ₂ SO ₄ \rightarrow (NH ₄) ₂ SO ₄
Preparation of ammonium nitrate	ammonia + nitric acid \rightarrow ammonium nitrate NH ₃ + HNO ₃ \rightarrow NH ₄ NO ₃
Eutrophication	The process by which an ecosystem, e.g. a river or dam, becomes enriched with inorganic plant nutrients, especially phosphorus and nitrogen, resulting in excessive plant growth. As plant growth becomes excessive, the amount of dead and decaying plant material increases rapidly.





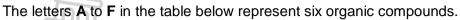
ORGANIC MOLECULES: NOMENCLATURE

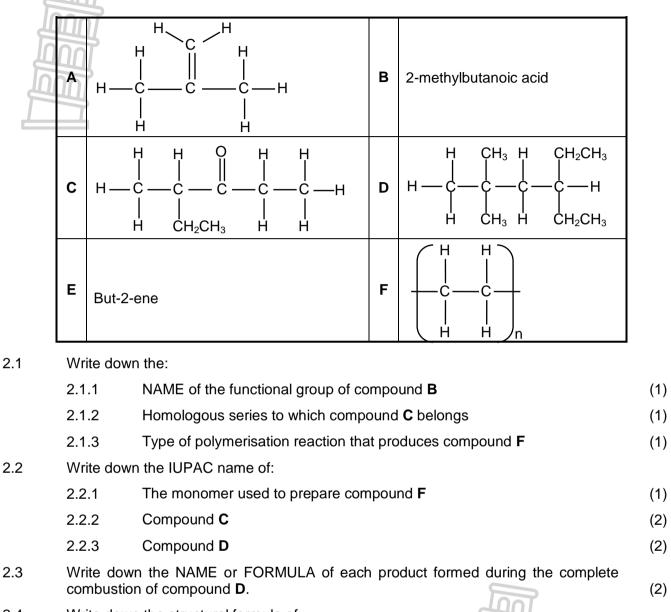
Consider the organic compounds represented by the letters A to F in the table below.



1.1 Write down the LETTER that represents the following:

	1.1.1	An aldehyde	(1)
	1.1.2	A condensation polymer	(1)
	1.1.3	A compound which has a carbonyl group bonded to two carbon atoms as its functional group	(1)
1.2	Write down	the IUPAC name of:	
	1.2.1	Compound C	(3)
	1.2.2	The monomer of compound D	(1)
1.3	Write down	the structural formula of:	
	1.3.1	Compound A	(2)
	1.3.2	Compound F	(2)
1.4	The table c	ontains compounds which are functional isomers.	
	1.4.1	Define the term functional isomer.	(2)
	1.4.2	Write down the LETTERS that represent two compounds that are functional isomers.	(1) [14]





- 2.4 Write down the structural formula of:
 - 2.4.1 Compound **B**
 - 2.4.2 A CHAIN ISOMER of compound A
- 2.5 A laboratory assistant uses bromine water to distinguish between compounds **D** and **E**. She adds bromine water to a sample of each in two different test tubes. She observes that the one compound decolourises the bromine water immediately, whilst the other one only reacts after placing the test tube in direct sunlight. Write down the:
 - 2.5.1 Letter (**D** or **E**) of the compound that will immediately decolourise the bromine water
 - 2.5.2 Name of the type of reaction that takes place in the test tube containing compound **D**
 - 2.5.3 Structural formula of the organic product formed in the test tube containing compound **E**

(2) **[18]**

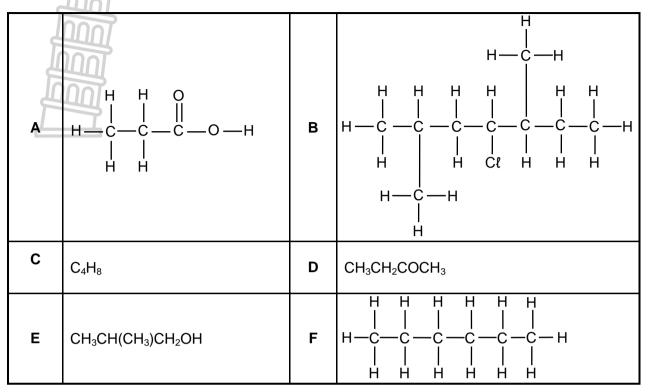
(1)

(1)

(2)

(2)

The letters A to F in the table below represent six organic compounds.

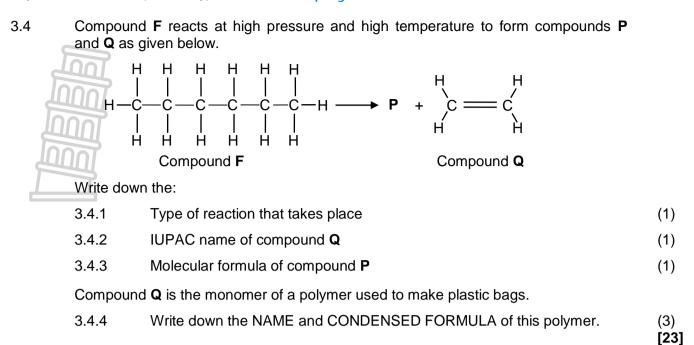


Use the information in the table (where applicable) to answer the questions that follow.

3.1	Write down the LETTER that represents a compound that:
	(A compound may be used more than once.)

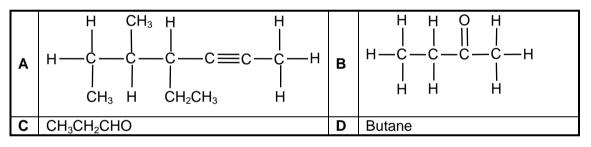
	3.1.1	Is a haloalkane	(1)
	3.1.2	Has a hydroxyl group as functional group	(1)
	3.1.3	Belongs to the same homologous series as ethanoic acid	(1)
3.2	Write dow	n the:	
	3.2.1	IUPAC name of compound B	(3)
	3.2.2	IUPAC name of compound E	(2)
	3.2.3	Structural formula of the functional group of compound D	(1)
3.3	Compoun	d C has CHAIN and POSITIONAL isomers.	
	3.3.1	Define the term positional isomer.	(2)
	3.3.2	Write down the IUPAC name of each of the TWO positional isomers of compound ${f C}.$	(4)
	3.3.3	Write down the structural formula of a chain isomer of compound ${f C}.$	(2)

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QUESTION 4

The letters **A** to **D** in the table below represent four organic compounds.



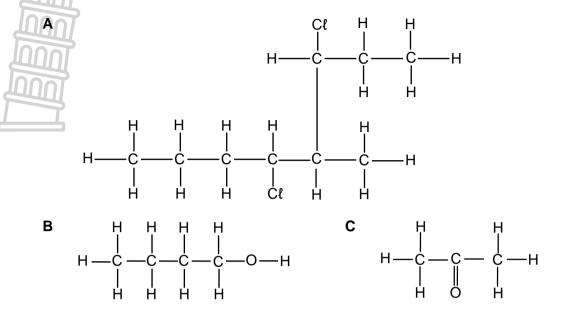
4.1 Write down the:

4.2

4.3

4.1.1	Letter that represents a ketone	(1)
4.1.2	Structural formula of the functional group of compound ${f C}$	(1)
4.1.3	General formula of the homologous series to which compound A belongs	(1)
4.1.4	IUPAC name of compound A	(3)
4.1.5	IUPAC name of compound B	(2)
Compound	D is a gas used in cigarette lighters.	
4.2.1	To which homologous series does compound D belong?	(1)
4.2.2	Write down the STRUCTURAL FORMULA and IUPAC NAME of a structural isomer of compound D .	(4)
4.2.3	Is the isomer in QUESTION 4.2.2 a CHAIN, POSITIONAL or FUNCTIONAL isomer?	(1)
•		
4.3.1	Homologous series to which 2-bromobutane belongs	(1)
4.3.2	Type of reaction that takes place	(1) [16]
	 4.1.2 4.1.3 4.1.4 4.1.5 Compound 4.2.1 4.2.2 4.2.3 Compound Write down 4.3.1 	 4.1.2 Structural formula of the functional group of compound C 4.1.3 General formula of the homologous series to which compound A belongs 4.1.4 IUPAC name of compound A 4.1.5 IUPAC name of compound B Compound D is a gas used in cigarette lighters. 4.2.1 To which homologous series does compound D belong? 4.2.2 Write down the STRUCTURAL FORMULA and IUPAC NAME of a structural isomer of compound D. 4.2.3 Is the isomer in QUESTION 4.2.2 a CHAIN, POSITIONAL or FUNCTIONAL isomer? Compound D reacts with bromine (Br₂) to form 2-bromobutane. Write down the name of the: 4.3.1 Homologous series to which 2-bromobutane belongs

5.1 Consider the organic compounds represented by the letters **A** to **C** below.



Write down the:

5.2

5.1.1	Name of the homologous series to which compound ${f C}$ belongs	(1)
5.1.2	IUPAC name of compound A	(3)
5.1.3	Structural formula of a tertiary alcohol that is a structural isomer of compound ${\bf B}$	(2)
An alcohol acid to forr	and methanoic acid are heated in the presence of concentrated sulphuric n an ester.	
5.2.1	What is the role of the concentrated sulphuric acid in this reaction?	(1)
5.2.2	Write down the NAME or FORMULA of the inorganic product formed.	(1)
	contains 6,67% hydrogen (H), 40% carbon (C) and 53,33% oxygen (O). mass of the ester is 60 g⋅mol ⁻¹ . Use a calculation to determine its:	
5.2.3	Empirical formula	(5)
5.2.4	Molecular formula	(3)
Write dowr	n the:	
5.2.5	Structural formula of methanoic acid	(1)
5.2.6	IUPAC name of the ester	(2) [19]

(2) **[14]**

QUESTION 6

6.2

6.3

Consider the organic compounds A to F below.

	$CH_3 \rightarrow C = C < H_{CH_3}$	В	H H H C C C C C H H CH ₃
C	CH ₃ CH ₂ CH ₂ CH ₂ OH	D	2,2-dimethylpropane
E	H O H H H C O O C C C H H H H	F	CH ₃ CHC(CH ₃) ₂

6.1 Write down the LETTER that represents a compound that	at:
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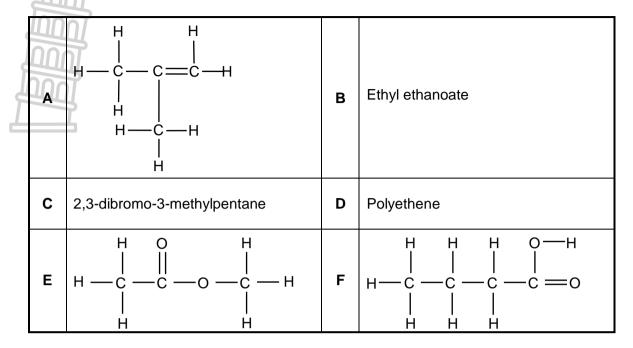
	6.1.1	Has a carbonyl group	(1)
	6.1.2	Is an alcohol	(1)
	6.1.3	Is a CHAIN ISOMER of CH ₃ (CH ₂) ₃ CH ₃	(1)
2	Write dow	n the:	
	6.2.1	IUPAC name of compound B	(2)
	6.2.2	Structural formula of compound F	(2)
	6.2.3	IUPAC name of a POSITIONAL isomer of compound A	(3)
3	Compound organic co	,	
	Write dow	n the:	
	6.3.1	Homologous series to which compound E belongs	(1)
	6.3.2	NAME or FORMULA of the catalyst used for the preparation of compound E	(1)

6.3.3 IUPAC name of compound **E**

7.2

7.3

The letters A to F in the table below represent six organic compounds.



7.1 Write down the LETTER that represents the following:

7.1.1	A hydrocarbon	(1)
7.1.2	A functional isomer of compound F	(1)
7.1.3	A compound which belongs to the same homologous series as compound B	(1)
7.1.4	A plastic	(1)
Write dowr	n the STRUCTURAL FORMULA of EACH of the following:	
7.2.1	Compound C	(3)
7.2.2	The acid used to prepare compound B	(2)
7.2.3	The monomer used to make compound D	(2)
Compound	A reacts with an unknown reactant, X , to form 2-methylpropane.	
Write dowr	n the:	
7.3.1	NAME of reactant X	(1)
7.3.2	Type of reaction that takes place	(1) [13]

ORGANIC MOLECULES: PHYSICAL PROPERTIES

QUEST	ON 1		
1.1	Give a rea	ason why alkanes are saturated hydrocarbons.	(1)
1.2	Write dow	n the structural formula of:	
	1.2.1	The functional group of alcohols	(1)
	1.2.2	A tertiary alcohol that is a structural isomer of butan-1-ol	(2)
1.3		investigate factors that influence the boiling points of alkanes and alcohols. f the investigations they determine the boiling points of the first three	
	1.3.1	Write down an investigative question for this investigation.	(2)
	1.3.2	Fully explain why the boiling point increases from methane to propane.	(3)
1.4	Explain th	ers find that the boiling point of propan-1-ol is higher than that of propane. his observation by referring to the TYPE of INTERMOLECULAR FORCES each of these compounds.	(3) [12]

QUESTION 2

Learners use compounds **A** to **C**, shown in the table below, to investigate a factor which influences the boiling point of organic compounds.

Α	CH ₃ CH ₂ CH ₂ CH ₃
В	CH ₃ CH ₂ CH ₂ CH ₂ CH ₃
С	CH ₃ CH ₂ CH ₂ CH ₂ CH ₂ CH ₃

2.1	Which ONE of the compounds (A , B or C) has the highest boiling point?	(1)
~ ~		

- 2.2 For this investigation, write down the:
 - 2.2.1 Independent variable
 - 2.2.2 Dependent variable
- 2.3 Write down the name of the type of Van der Waals force that occurs between the molecules of compound **B**. (1)
- 2.4 How will the vapour pressure of 2-methylpentane compare to that of compound **C**? Write down only HIGHER THAN, LOWER THAN or EQUAL TO. (1)

The learners now compare the boiling points of compounds **D** and **E**, shown in the table below.

D	CH ₃ CH ₂ CH ₂ CH ₂ CH ₂ OH
Е	CH ₃ CH ₂ CH ₂ COOH

2.5 How does the boiling point of compound **D** compare to that of compound **E**? Write down HIGHER THAN, LOWER THAN or EQUAL TO. Fully explain the answer.

(4) **[9]**

(1)

(1)

(1)

(1)

(4)

(1) **[12]**

(2)

(1)

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QUESTION 3

The table below shows five organic compounds represented by the letters A to E.

Α	CH ₄
В	CH ₃ CH ₃
С	CH ₃ CH ₂ CH ₃
D	CH ₃ CH ₂ CH ₂ CH ₃
E	CH ₃ CH ₂ OH

3.1 Is compound **B** SATURATED or UNSATURATED? Give a reason for the answer. (2)

Consider the boiling points of compounds **A** to **E** given in random order below and use them, where applicable, to answer the questions that follow.

0 °C	- 162 °C	- 42 °C	- 89 °C	78 °C
------	----------	---------	---------	-------

3.2 Write down the boiling point of:

- 3.2.2 Compound E
- 3.3 Explain the difference in boiling points of compounds **C** and **E** by referring to the TYPE of intermolecular forces present in EACH of these compounds. (3)
- 3.4 Does vapour pressure INCREASE or DECREASE from compounds **A** to **D**? Fully explain the answer.
- 3.5 How will the vapour pressure of 2-methylpropane compare to the vapour pressure of compound **D**? Write down only HIGHER THAN, LOWER THAN or EQUAL TO.

QUESTION 4

Four compounds of comparable molecular mass are used to investigate the effect of functional groups on vapour pressure. The results obtained are shown in the table below.

	COMPOUND	VAPOUR PRESSURE (kPa at 20 °C)
Α	Butane	204
В	Propan-2-one	24,6
С	Propan-1-ol	2
D	Ethanoic acid	1,6

- 4.1 Define the term *functional group* of an organic compound.
- 4.2 Which ONE of the compounds (**A**, **B**, **C** or **D**) in the table has the:
 - 4.2.1 Highest boiling point (Refer to the vapour pressures in the table to give a reason for the answer.) (2)
 - 4.2.2 Weakest intermolecular forces

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- 4.3 Refer to the type of intermolecular forces to explain the difference between the vapour pressure of compound **A** and compound **B**.
- 4.4 The vapour pressures of compounds **C** and **D** are much lower than those of compounds **A** and **B**. Name the type of intermolecular force in **A** and **B** that is responsible for this difference.
- 4.5 Briefly explain the difference in vapour pressure between compound **C** and compound **D**.
- 4.6 During a combustion reaction in a closed container of adjustable volume, 8 cm³ of compound **A** (butane) reacts in excess oxygen according to the following balanced equation:

$$2C_4H_{10}(g) + 13O_2(g) \rightarrow 8CO_2(g) + 10H_2O(g)$$

If the initial volume of the oxygen in the container was 60 cm³, calculate the TOTAL volume of the gases that are present in the container at the end of the reaction. All the gases in the container are at the same temperature and pressure.

QUESTION 5

- 5.1 Define the term *boiling point*.
- 5.2 What is the relationship between strength of intermolecular forces and boiling point? (1)

The relationship between strength of intermolecular forces and boiling point is investigated using four organic compounds from different homologous series. The compounds and their boiling points are given in the table below.

	COMPOUND	BOILING POINT (°C)
Α	Propane	-42
В	Propan-2-one	56
С	Propan-1-ol	97
D	Propanoic acid	141

- 5.3 Refer to the TYPE and the STRENGTH of intermolecular forces to explain the difference in boiling points between:
 - 5.3.1Compounds A and B(3)5.3.2Compounds C and D(3)
- 5.4 Is compound **B** a GAS or a LIQUID at room temperature?

(1)

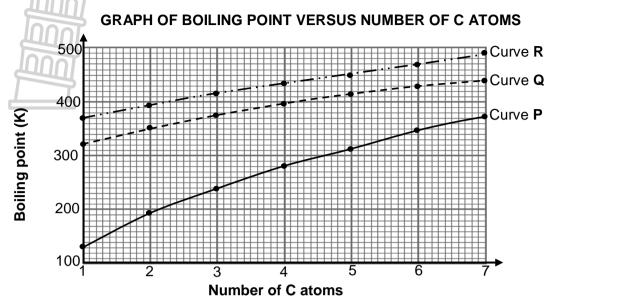
(2)

(5) **[16]**

(2)

(1) [**10**]

The relationship between boiling point and the number of carbon atoms in straight chain molecules of alkanes, carboxylic acids and alcohols is investigated. Curves **P**, **Q** and **R** are obtained.



6.1 Define the term *boiling point*.

6.2 For curve P , write down a conclusion that can be drawn from the above results.	(2)
--	-----

- 6.3 Identify the curve (**P**, **Q** or **R**) that represents each of the following:
 - 6.3.1 Alkanes
 - 6.3.2 Carboxylic acids
 - Explain the answer to QUESTION 6.3.2 by referring to the:
 - Types of intermolecular forces present in alkanes, carboxylic acids and alcohols
 - Relative strengths of these intermolecular forces
 - Energy needed

QUESTION 7

6.4

The boiling points of three isomers are given in the table below.

	ISOMERS	BOILING POINT (°C)
Α	2,2-dimethylpropane	9
В	2-methylbutane	28
С	pentane	36

- 7.1 Define the term *structural isomer*.
- 7.2 What type of isomers (POSITIONAL, CHAIN or FUNCTIONAL) are these three compounds? (1)
- 7.3 Explain the trend in the boiling points from compound **A** to compound **C**.
- 7.4 Which ONE of the three compounds (**A**, **B** or **C**) has the highest vapour pressure? Refer to the data in the table to give a reason for the answer. (2)
- 7.5 Use MOLECULAR FORMULAE and write down a balanced equation for the complete combustion of compound **B**.

(2)

(1)

(1)

(5) **[11]**

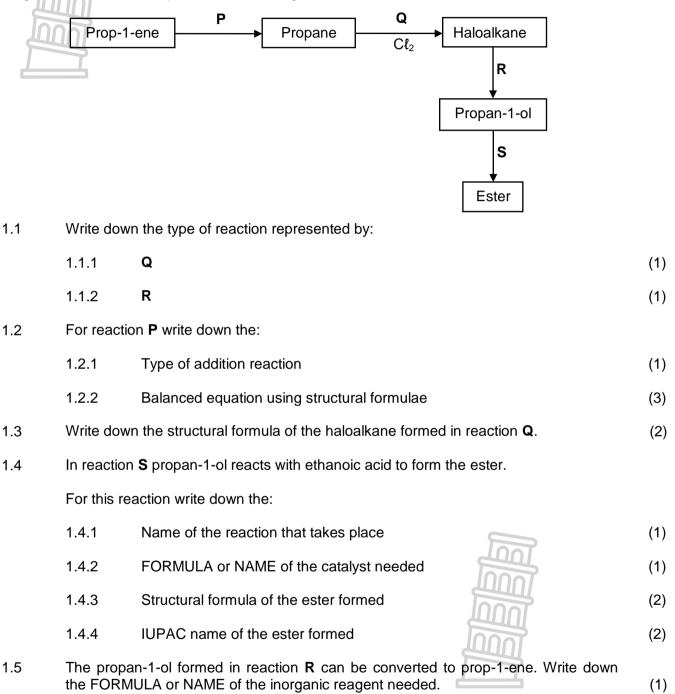
(2)

(3)

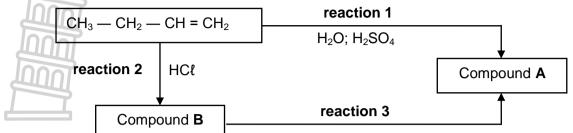
ORGANIC MOLECULES: ORGANIC REACTIONS

QUESTION 1

The flow diagram below shows the preparation of an ester using prop-1-ene as a starting reagent. **P**, **Q**, **R** and **S** represent different organic reactions.



In the flow diagram below, but-1-ene is used as starting material in the preparation of compound A.



- 2.1 Is but-1-ene a SATURATED or UNSATURATED compound? Give a reason for the answer. (2)
- 2.2 Compound **A** is the major product formed in **reaction 1**.

Write down the:

2.2.1	Structural formula of compound A	(2)
2.2.2	Type of reaction that takes place	(1)
For compo	aund B , write down the:	
2.3.1	IUPAC name	(2)
2.3.2	Structural formula of the positional isomer	(2)
For reaction	on 3, write down:	
2.4.1	TWO reaction conditions needed	(2)
2.4.2	The type of reaction that occurs	(1)
2.4.3	A balanced equation, using molecular formulae	(3) [15]

QUESTION 3

2.3

2.4

Consider the incomplete equations of two reactions below.

X represents the organic product formed in **reaction 1**, which is a SUBSTITUTION REACTION. In **reaction 2**, X reacts with reactant Y as shown.

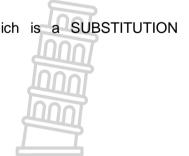
Reaction 1: $C_2H_5Br \xrightarrow{\text{strong base}} NaBr + X$

Reaction 2: $X + Y \xrightarrow{\text{Concentrated } H_2SO_4} \rightarrow C_3H_6O_2 + H_2O$

3.1 Consider **reaction 1**. Write down the:

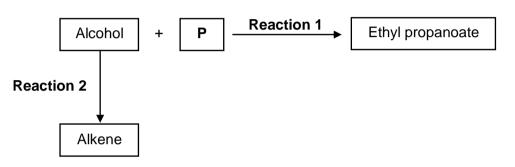
3.1.1	Type of substitution reaction that takes place	(1)
3.1.2	TWO reaction conditions	(2)

3.1.3 IUPAC name of compound **X**



(1)

4.1 The flow diagram below shows two organic reactions. The letter **P** represents an organic compound.



Use the information in the flow diagram to answer the questions that follow.

Write down the:

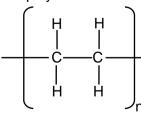
QUESTION 4

4.1.1	Type of reaction of which Reaction 1 is an example	(1)
4.1.2	STRUCTURAL FORMULA of the functional group of ethyl propanoate	(1)

4.1.3 IUPAC name of compound P

Reaction 2 takes place in the presence of an acid catalyst and heat. Write down the:

- 4.1.4 Type of reaction of which **Reaction 2** is an example (1)
- 4.1.5 NAME or FORMULA of the acid catalyst
- 4.1.6 STRUCTURAL FORMULA of the alkene
- 4.2 The condensed formula of a polymer is shown below.





Write down the:

- 4.2.1 STRUCTURAL FORMULA of the monomer that is used to prepare the above polymer
- 4.2.2 Type of polymerisation reaction (ADDITION or CONDENSATION) that is used to prepare this polymer

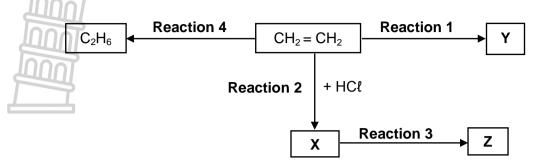
(2)

(1)

(1)

(2)

The flow diagram below shows different organic reactions using $CH_2 = CH_2$ as the starting reactant. **X**, **Y** and **Z** represent different organic compounds.



5.1 During **Reaction 1**, $CH_2 = CH_2$ undergoes polymerisation to form compound **Y**.

For this reaction, write down the:

5.1.1	Type of polymerisation	(1)

- 5.1.2 NAME of compound \mathbf{Y} (1)
- 5.2 For **Reaction 2**, write down the:
 - 5.2.1 IUPAC name of compound \mathbf{X} (2)
 - 5.2.2 Type of addition reaction of which this is an example (1)
- 5.3 During **Reaction 3**, compound **X** reacts with excess hot water.

Write down the:

- 5.3.1 STRUCTURAL FORMULA of compound **Z** (2)
- 5.3.2 NAME or FORMULA of the INORGANIC product (1)

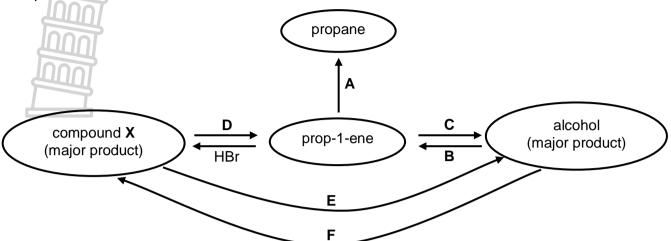
5.4 **Reaction 4** is an addition reaction.

5.4.1 Is C₂H₆ a SATURATED or an UNSATURATED compound? Give a reason for the answer. (2)
5.4.2 Write down the NAME or FORMULA of the INORGANIC reactant needed for this reaction. (1)
5.4.3 Using molecular formulae, write down a balanced equation for the complete combustion of C₂H₆. (3) [14]

[15]

QUESTION 6

The flow diagram below shows how prop-1-ene can be used to prepare other organic compounds.



6.1 Write down the type of reaction represented by:

6.1.1	Α		(1)
6.1.2	D		(1)

- 6.1.3 **F** (1)
- 6.2 Write down the:

6.2.1	NAME or FORMULA of the catalyst needed for reaction A	(1)
6.2.2	NAME or FORMULA of the inorganic reagent needed for reaction ${f B}$	(1)
6.2.3	Type of addition reaction represented by reaction ${f C}$	(1)
6.2.4	IUPAC name of compound X	(2)
Use struct	ural formulae to write down a balanced equation for reaction B .	(5)
	ctions D and E take place in the presence of a strong base.	(2)

QUESTION 7

6.3

6.4

Butane (C_4H_{10}) is produced in industry by the THERMAL cracking of long-chain hydrocarbon molecules, as shown in the equation below. **X** represents an organic compound that is produced.

$$C_{10}H_{22} \ \to \ {\bm X} + C_4 H_{10}$$

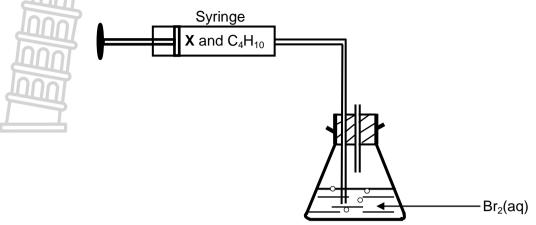
7.1 Write down:

7.1.1	ONE condition required for THERMAL cracking to take place	(1)
7.1.2	The molecular formula of compound X	(1)

7.1.3 The homologous series to which compound **X** belongs (1)

(4)

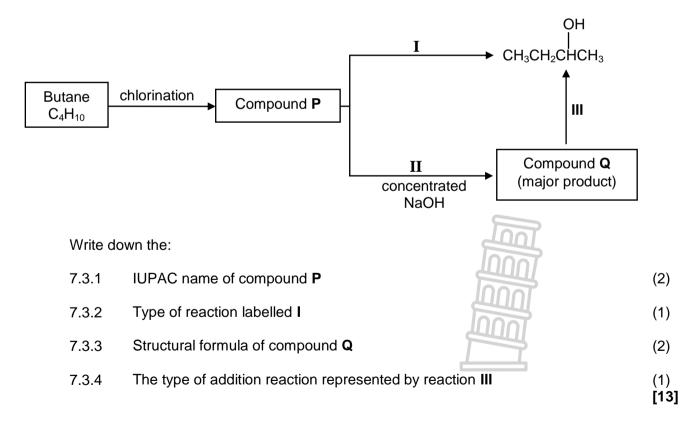
7.2 A mixture of the two gases, compound **X** and butane, is bubbled through bromine water, Br₂(aq), in a conical flask, as illustrated below. THE REACTION IS CARRIED OUT IN A DARKENED ROOM.



The colour of the bromine water changes from reddish brown to colourless when the mixture of the two gases is bubbled through it.

Which ONE of the gases (X or BUTANE) decolorises the bromine water? Explain the answer.

7.3 Study the flow diagram below, which represents various organic reactions, and answer the questions that follow.



REACTION RATE AND ENERGY IN CHEMICAL REACTIONS

QUESTION 1

1.1 Define the term reaction rate in words.

Learners use the reaction between IMPURE POWDERED calcium carbonate and excess hydrochloric acid to investigate reaction rate. The balanced equation for the reaction is:

 $CaCO_3(s) + 2HC\ell(aq) \rightarrow CaC\ell_2(aq) + H_2O(\ell) + CO_2(g)$

They perform four experiments under different conditions of concentration, mass and temperature as shown in the table below. They use identical apparatus in the four experiments and measure the volume of gas released in each experiment.

	EXPERIMENT			
	1	2	3	4
Concentration of acid (mol·dm ⁻³)	1	0,5	1	1
Mass of impure calcium carbonate (g)	15	15	15	25
Initial temperature of acid (°C)	30	30	40	40

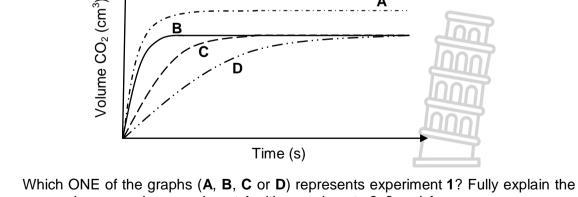
1.2 The results of experiments **1** and **3** are compared in the investigation.

Write down the:

1.4

- 1.2.1 Independent variable
 - 1.2.2 Dependent variable
- 1.3 Use the collision theory to explain why the reaction rate in experiment **4** will be higher than that in experiment **3**. (3)

The learners obtain graphs **A**, **B**, **C** and **D** below from their results.



- answer by comparing experiment 1 with experiments 2, 3 and 4.
- 1.5 When the reaction in experiment **4** reaches completion, the volume of gas formed is 4,5 dm³. Assume that the molar gas volume at 40 °C is equal to 25,7 dm³.

Calculate the mass of the impurities present in the calcium carbonate.

(6)

(5) **[18]**

(1)

(2)

A group of learners uses the reaction of EXCESS hydrochloric acid (HCl) with zinc (Zn) to investigate factors which influence reaction rate. The balanced equation for the reaction is:

$$Zn(s) + 2HC\ell(aq) \rightarrow ZnC\ell_2(aq) + H_2(g)$$

They use the same volume of hydrochloric acid and 1,2 g of zinc in each of five experiments. The reaction conditions and temperature readings before and after completion of the reaction in each experiment are summarised in the table below.

	REACTION CONDITIONS				Time
Experiment	Concentration of	Temperature (°C)		State of division	Time (s)
	HCℓ (mol·dm⁻³)	Before	After	of the 1,2 g of Zn	(3)
1	0,5	20	34	granules	50
2	0,5	20	35	powder	10
3	0,8	20	36	powder	6
4	0,5	35	50	granules	8
5	0,5	20	34	granules	11

2.1 Is the reaction between hydrochloric acid and zinc EXOTHERMIC or ENDOTHERMIC? Give a reason for the answer by referring to the data in the table. (2)

2.2 Give a reason for the difference in reaction rate observed for **Experiments 1** and **2**. (1)

- 2.3 The learners compare the results of **Experiments 1** and **3** to draw a conclusion regarding the effect of concentration on reaction rate. Give a reason why this is not a fair comparison.
- 2.4 How does the rate of the reaction in **Experiment 5** compare to that in **Experiment** 1? Write down FASTER THAN, SLOWER THAN or EQUAL TO.

Write down the factor responsible for the difference in the rate of reaction and fully explain, by referring to the collision theory, how this factor affects reaction rate.

2.5 Calculate the rate at which the hydrochloric acid reacts in **Experiment 4** in mol \cdot s⁻¹.

QUESTION 3

A group of learners uses the reaction of clean magnesium ribbon with dilute hydrochloric acid to investigate factors that influence reaction rate. The balanced equation for the reaction is:

$$Mg(s) + 2HC\ell(aq) \rightarrow MgC\ell_2(aq) + H_2(g) \qquad \Delta H < 0$$

- 3.1 Is the above reaction EXOTHERMIC or ENDOTHERMIC? Give a reason for the answer.
- 3.2 In one of the experiments 5 g magnesium ribbon was added to the hydrochloric acid solution.
 - 3.2.1 If 30 cm³ dilute hydrochloric acid solution of concentration 1,5 mol·dm⁻³ is USED UP in 1 minute, calculate the average reaction rate in mol·s⁻¹.

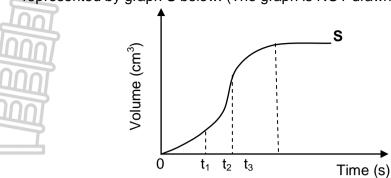
(5)

(2)

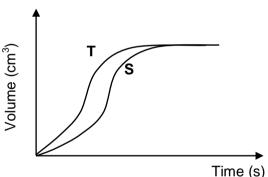
(1)

(5)

(6) **[15]** The volume of hydrogen gas produced as a function of time in this experiment is represented by graph **S** below. (The graph is NOT drawn to scale.)



- 3.2.2 How does the rate of the reaction change between: (Write down INCREASES, DECREASES or NO CHANGE.)
 - (a) t₁ and t₂
 Use the collision theory to explain the answer.
 - (b) t_2 and t_3 Give a reason for the answer without referring to the graph. (2)
- 3.3 In another experiment they add 5 g of magnesium to 30 cm³ of dilute hydrochloric acid of concentration 1,5 mol·dm⁻³. They obtained graph **T** below. (The graph is NOT drawn to scale.)



Give TWO possible reasons why graph T differs from graph S.

QUESTION 4

Dilute acids, indicated in the table below, react with EXCESS zinc in each of the three experiments to produce hydrogen gas. The zinc is completely covered with the acid in each experiment.

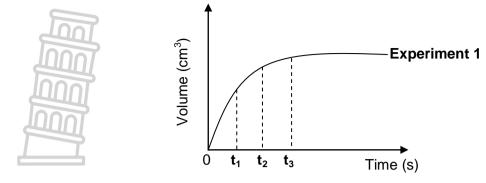
EXPERIMENT	DILUTE ACID
1	100 cm ³ of 0,1 mol·dm ⁻³ H ₂ SO ₄
2	50 cm ³ of 0,2 mol·dm ⁻³ H ₂ SO ₄
3	100 cm ³ of 0,1 mol·dm ⁻³ HCł

The volume of hydrogen gas produced is measured in each experiment.

4.1 Name TWO essential apparatuses needed to determine the rate of hydrogen production.

(4)

(2) [**15**] The graph below was obtained for Experiment 1.



Use this graph and answer the questions that follow.

4.2 At which time $(\mathbf{t}_1, \mathbf{t}_2 \text{ or } \mathbf{t}_3)$ is the:

4.2.1	Reaction rate the highest
-------	---------------------------

- 4.2.2 Mass of zinc present in the flask the smallest
- 4.3 In which time interval, between t_1 and t_2 OR between t_2 and t_3 , does the largest volume of hydrogen gas form per second? (1)
- 4.4 Redraw the graph for Experiment 1 in the ANSWER BOOK.
 On the same set of axes, sketch the graphs that will be obtained for Experiments 2 and 3. Clearly label the three graphs as EXPERIMENT 1, EXPERIMENT 2 and EXPERIMENT 3.
- 4.5 The initial mass of zinc used in each experiment is 0,8 g. The balanced equation for the reaction in **Experiment 3** is:

$$Zn(s) + 2HC\ell(aq) \rightarrow ZnC\ell_2(aq) + H_2(g)$$

- 4.5.1 Calculate the mass of zinc present in the flask after completion of the reaction in **Experiment 3**.
- 4.5.2 How will the mass of zinc present in the flask after completion of the reaction in **Experiment 2** compare to the answer to QUESTION 5.5.1? Write down only LARGER THAN, SMALLER THAN or EQUAL TO.

QUESTION 5

Methanol and hydrochloric acid react according to the following balanced equation:

 $CH_3OH(aq) + HC\ell(aq) \rightarrow CH_3C\ell(aq) + H_2O(\ell)$

- 5.1 State TWO factors that can INCREASE the rate of this reaction.
- 5.2 Define the term *reaction rate*.
- 5.3 The rate of the reaction between methanol and hydrochloric acid is investigated. The concentration of HCl(aq) was measured at different time intervals. The following results were obtained:

TIME (MINUTES)	HCℓ CONCENTRATION (mol·dm ⁻³)
0	1,90
15	1,45
55	1,10
100	0,85
215	0,60

(5)

(1)

(1)

(4)

(1)

[15]

(2)

(2)

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5.3.1	Calculate the average reaction rate, in (mol·dm ⁻³)·min ⁻¹ during the 15 minutes.	first	(3)
5.3.2	Use the data in the table to draw a graph of concentration versus tim a graph paper.	e on	(3)
5.3.3	From the graph, determine the concentration of HCl(aq) at 40 th minute.	the	(1)
5.3.4	Use the collision theory to explain why the reaction rate decreases time. Assume that the temperature remains constant.	with	(3)
5.3.5	Calculate the mass of $CH_3Cl(aq)$ in the flask at the 215 th minute. volume of the reagents remains 60 cm ³ during the reaction.	The	(5) [19]

The reaction between dilute hydrochloric acid and sodium thiosulphate $(Na_2S_2O_3)$ is used to investigate one of the factors that influences reaction rate. The balanced equation for the reaction is:

 $Na_2S_2O_3(aq) + 2HC\ell(aq) \rightarrow 2NaC\ell(aq) + S(s) + H_2O(\ell) + SO_2(g)$

The hydrochloric acid solution is added to the sodium thiosulphate solution in a flask. The flask is placed over a cross drawn on a sheet of white paper, as shown in the diagram below. The time that it takes for the cross to become invisible is measured to determine the reaction rate.

flask Na $_2S_2O_3(aq) + HC\ell(aq)$ white paper

Four experiments, **A** to **D**, are conducted during this investigation. The volumes of reactants used in each of the four experiments and the times of the reactions are summarised in the table below.

Experiment	Volume of Na₂S₂O₃(aq) (cm³)	Volume of H₂O(ℓ) (cm³)	Volume of HCℓ(aq) (cm³) Time (s)	
Α	25	0	5 50,0	
В	20	5	5 62,5	
С	15	10	5 83,3	
D	10	15	5 125,0	

6.1 State TWO factors that can influence the rate of the reaction above.

(2)

(1)

(1)

(2)

(1)

- 6.2 Write down the NAME or FORMULA of the product that causes the cross to become invisible.
- 6.3 Give a reason why water is added to the reaction mixture in experiments **B** to **D**.
- 6.4 Write down an investigative question for this investigation.
- 6.5 In which experiment (**A**, **B**, **C** or **D**) is the reaction rate the highest?

- 6.6 Use the collision theory to explain the difference in reaction rate between experiments **B** and **D**.
- 6.7 The original $Na_2S_2O_3$ solution was prepared by dissolving 62,50 g $Na_2S_2O_3$ crystals in distilled water in a 250 cm³ volumetric flask.

Calculate the mass of sulphur, S, that will form in experiment **D** if $Na_2S_2O_3$ is the limiting reactant.

QUESTION 7

Hydrogen peroxide, H_2O_2 , decomposes to produce water and oxygen according to the following balanced equation:

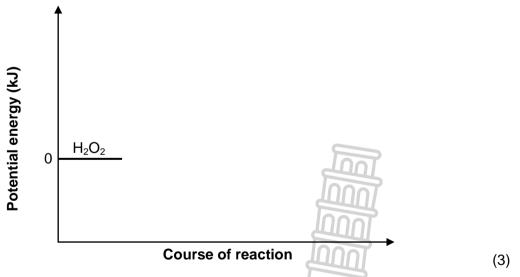
$$2H_2O_2(\ell) \rightarrow 2H_2O(\ell) + O_2(g)$$

- 7.1 The activation energy (E_A) for this reaction is 75 kJ and the heat of reaction (ΔH) is 196 kJ.
 - 7.1.1 Define the term *activation energy*.
 - 7.1.2 Redraw the set of axes below in your ANSWER BOOK and then complete the potential energy diagram for this reaction.

Indicate the value of the potential energy of the following on the y-axis:

- Activated complex
- Products

(The graph does NOT have to be drawn to scale.)



When powdered manganese dioxide is added to the reaction mixture, the rate of the reaction increases.

- 7.1.3 On the graph drawn for QUESTION 7.1.2, use broken lines to show the path of the reaction when the manganese dioxide is added.
- 7.1.4 Use the collision theory to explain how manganese dioxide influences the rate of decomposition of hydrogen peroxide.

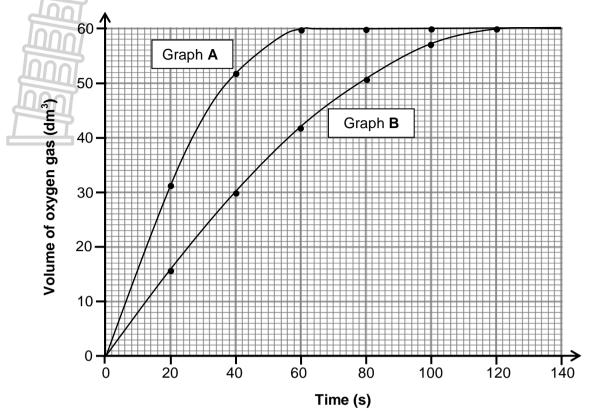
(3)

(2)

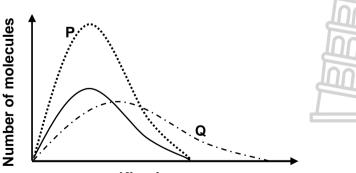
(2)

(3)

(7) **[17]** 7.2 Graphs **A** and **B** below were obtained for the volume of oxygen produced over time under different conditions.



- 7.2.1 Calculate the average rate of the reaction (in $dm^3 \cdot s^{-1}$) between t = 10 s and t = 40 s for graph **A**.
- 7.2.2 Use the information in graph **A** to calculate the mass of hydrogen peroxide used in the reaction. Assume that all the hydrogen peroxide decomposed. Use 24 dm³⋅mol⁻¹ as the molar volume of oxygen.
- 7.2.3 How does the mass of hydrogen peroxide used to obtain graph **B** compare to that used to obtain graph **A**? Choose from GREATER THAN, SMALLER THAN or EQUAL TO.
- 7.3 Three energy distribution curves for the oxygen gas produced under different conditions are shown in the graph below. The curve with the solid line represents 1 mol of oxygen gas at 90 °C.



Kinetic energy

Choose the curve (\mathbf{P} or \mathbf{Q}) that best represents EACH of the following situations:

- 7.3.1 1 mol of oxygen gas produced at 120 °C
- 7.3.2 2 moles of oxygen gas produced at 90 °C

(3)

(4)

(1)

(1)

(1) **[20]**

CHEMICAL EQUILIBRIUM

QUESTION 1

A certain amount of nitrogen dioxide gas (NO₂) is sealed in a gas syringe at 25 °C. When equilibrium is reached, the volume occupied by the reaction mixture in the gas syringe is 80 cm³. The balanced chemical equation for the reaction taking place is:

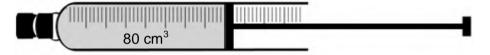


 $\begin{array}{rcl} 2NO_2(g) &\rightleftharpoons & N_2O_4(g) & \Delta H < 0 \\ \text{dark brown} & \text{colourless} \end{array}$

- 1.1 Define the term *chemical equilibrium*.
- 1.2 At equilibrium the concentration of the $NO_2(g)$ is 0,2 mol·dm⁻³. The equilibrium constant for the reaction is 171 at 25 °C.

Calculate the initial number of moles of $NO_2(g)$ placed in the gas syringe.

1.3 The diagram below shows the reaction mixture in the gas syringe after equilibrium is established.



The pressure is now increased by decreasing the volume of the gas syringe at constant temperature as illustrated in the diagram below.



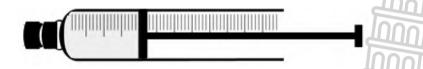
1.3.1 IMMEDIATELY after increasing the pressure, the colour of the reaction mixture in the gas syringe appears darker than before. Give a reason for this observation.

(1)

(2)

(8)

After a while a new equilibrium is established as illustrated below. The colour of the reaction mixture in the gas syringe now appears lighter than the <u>___initial</u> colour.



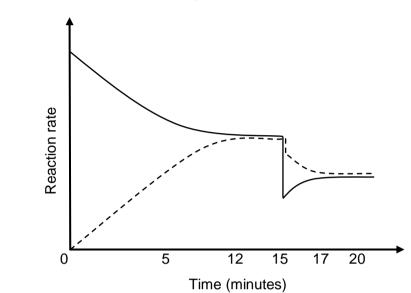
- 1.3.2 Use Le Chatelier's principle to explain the colour change observed in the gas syringe. (3)
- 1.4 The temperature of the reaction mixture in the gas syringe is now increased and a new equilibrium is established. How will each of the following be affected?
 - 1.4.1Colour of the reaction mixture
Write down only DARKER, LIGHTER or REMAINS THE SAME.(1)
 - 1.4.2 Value of the equilibrium constant (K_c) Write down only INCREASES, DECREASES or REMAINS THE SAME. (1)

[16]

Pure hydrogen iodide, sealed in a 2 dm³ container at 721 K, decomposes according to the following balanced equation:

- $2HI(g) \rightleftharpoons H_2(g) + I_2(g)$ $\Delta H = + 26 \text{ kJ} \cdot \text{mol}^{-1}$

The graph below shows how reaction rate changes with time for this reversible reaction.



2.1 Write down the meaning of the term *reversible reaction*.

(1)

- 2.2 How does the concentration of the reactant change between the 12th and the 15th minute? Write down only INCREASES, DECREASES or NO CHANGE.
- 2.3 The rates of both the forward and the reverse reactions suddenly change at t = 15 minutes.
 - 2.3.1 Give a reason for the sudden change in reaction rate. (1)
 - 2.3.2 Fully explain how you arrived at the answer to QUESTION 2.3.1. (3)

The equilibrium constant (K_c) for the forward reaction is 0,02 at 721 K.

- 2.4 At equilibrium it is found that 0,04 mol HI(g) is present in the container. Calculate the concentration of $H_2(g)$ at equilibrium. (6)
- 2.5 Calculate the equilibrium constant for the reverse reaction.
- 2.6 The temperature is now increased to 800 K. How will the value of the equilibrium constant (K_c) for the forward reaction change? Write down only INCREASES, DECREASES or REMAINS THE SAME. (1)

[14]

(1)

Initially excess NH₄HS(s) is placed in a 5 dm³ container at 218 °C. The container is sealed and the reaction is allowed to reach equilibrium according to the following balanced equation:

> $NH_4HS(s) \Rightarrow NH_3(g) + H_2S(g)$ $\Delta H > 0$

- State Le Chatelier's principle. 3.1
- 3.2 What effect will each of the following changes have on the amount of $NH_3(g)$ at equilibrium? Write down only INCREASES, DECREASES or REMAINS THE SAME.

3.2.1 More NH₄HS(s) is added (1)

- 3.2.2 The temperature is increased
- 3.3 The equilibrium constant for this reaction at 218 °C is 1.2×10^{-4} . Calculate the minimum mass of NH₄HS(s) that must be sealed in the container to obtain equilibrium. (6)

The pressure in the container is now increased by decreasing the volume of the container at constant temperature.

3.4 How will this change affect the number of moles of $H_2S(g)$ produced? Fully explain the answer.

QUESTION 4

An unknown gas, $X_2(g)$, is sealed in a container and allowed to form $X_3(g)$ at 300 °C. The reaction reaches equilibrium according to the following balanced equation:

$$3X_2(g) \Rightarrow 2X_3(g)$$

4.1 How will the rate of formation of $X_3(g)$ compare to the rate of formation of $X_2(g)$ at equilibrium? Write down only HIGHER THAN, LOWER THAN or EQUAL TO. (1)

The reaction mixture is analysed at regular time intervals. The results obtained are shown in the table below.

[X₂]

(mol·dm⁻³)

0,4

0,22

0.08

		6	0,06	0,226	
		8	0,06	0,226	
		10	0,06	0,226	
4.2	Calculate the equ	uilibrium con	stant, K _c , for this	s reaction at 300 °	C.

TIME

(s) 0

2

4

4.3 More $X_3(g)$ is now added to the container.

> 4.3.1 How will this change affect the amount of $X_2(g)$? Write down INCREASES, DECREASES or REMAINS THE SAME. (1)

[X₃]

(mol·dm⁻³)

0

0.120

0.213

4.3.2 Use Le Chatelier's principle to explain the answer to QUESTION 4.3.1. (2)

(2)

(1)

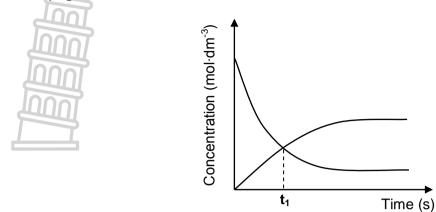
(3) [13]

(4)

(1)

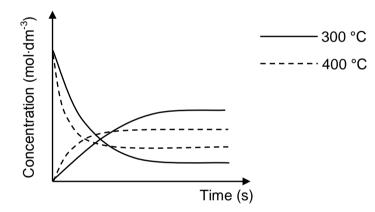
(4)

The curves on the set of axes below (not drawn to scale) was obtained from the results in the table on page 10.



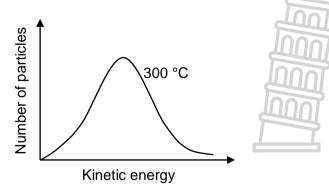
4.4 How does the rate of the forward reaction compare to that of the reverse reaction at t_1 ? Write down only HIGHER THAN, LOWER THAN or EQUAL TO.

The reaction is now repeated at a temperature of 400 °C. The curves indicated by the dotted lines below were obtained at this temperature.



4.5 Is the forward reaction EXOTHERMIC or ENDOTHERMIC? Fully explain how you arrived at the answer.

The Maxwell-Boltzmann distribution curve below represents the number of particles against kinetic energy at 300 °C.



4.6 Redraw this curve in the ANSWER BOOK. On the same set of axes, sketch the curve that will be obtained at 400 °C. Clearly label the curves as **300** °C and **400** °C respectively.

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QUESTION 5

Initially, 2,2 g of pure $CO_2(g)$ is sealed in an empty 5 dm³ container at 900 °C.

5.1	Calculate the initial concentration of $CO_2(g)$.	(4)

5.2 Give a reason why equilibrium will not be established.

 $CaCO_3(s)$ is now added to the 2,2 g $CO_2(g)$ in the container and after a while equilibrium is established at 900 °C according to the following balanced equation:

 $CaCO_3(s) \rightleftharpoons CaO(s) + CO_2(g)$

The equilibrium constant for this reaction at 900 °C is 0,0108.

- 5.3 Give a reason why this reaction will only reach equilibrium in a SEALED container. (1)
- 5.4 Calculate the minimum mass of CaCO₃(s) that must be added to the container to achieve equilibrium. (7)
- 5.5 How will EACH of the following changes affect the amount of CO₂(g)? Write down only INCREASES, DECREASES or REMAINS THE SAME.

(1)

- 5.5.2 The pressure is increased
- 5.6 It is found that the equilibrium constant (K_c) for this reaction is 2,6 x 10⁻⁶ at 727 °C. Is the reaction EXOTHERMIC or ENDOTHERMIC? Fully explain how you arrived at the answer.

QUESTION 6

Carbon dioxide reacts with carbon in a closed system to produce carbon monoxide, CO(g), according to the following balanced equation:

$$CO_2(g) + C(s) \rightleftharpoons 2CO(g)$$
 $\Delta H > 0$

6.1 What does the double arrow indicate in the equation above? (1)
6.2 Is the above reaction an EXOTHERMIC reaction or an ENDOTHERMIC reaction? Give a reason for the answer. (2)
Initially an unknown amount of carbon diaxide is exposed to bet earbon at 200 °C in a scaled

Initially an unknown amount of carbon dioxide is exposed to hot carbon at 800 °C in a sealed 2 dm^3 container. The equilibrium constant, K_c, for the reaction at this temperature is 14. At equilibrium it is found that 168,00 g carbon monoxide is present.

6.3 How will the equilibrium concentration of the product compare to that of the reactants? Choose from LARGER THAN, SMALLER THAN or EQUAL TO.

Give a reason for the answer. (No calculation is required.)

- 6.4 Calculate the initial amount (in moles) of CO₂(g) present.
- 6.5 State how EACH of the following will affect the yield of CO(g) at equilibrium. Choose from INCREASES, DECREASES or REMAINS THE SAME.

6.5.1 More carbon is added at constant temperature.		(1)
6.5.2	The pressure is increased.	(1)

6.5.2The pressure is increased.(1)6.5.3The temperature is increased.(1)[17]

(1)

(1)

(4) **[19]**

(2)

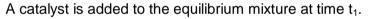
(9)

Hydrogen gas, $H_2(g)$, reacts with sulphur powder, S(s), according to the following balanced equation:

 $H_2(g) + S(s) \rightleftharpoons H_2S(g) \qquad \Delta H < 0$

The system reaches equilibrium at 90 °C.

- 7.1 Define the term *chemical equilibrium*.
- 7.2 How will EACH of the following changes affect the number of moles of $H_2S(g)$ at equilibrium? Choose from INCREASES, DECREASES or REMAINS THE SAME.
 - 7.2.1 The addition of more sulphur
 - 7.2.2An increase in temperature
Use Le Chatelier's principle to explain the answer.(4)
- 7.3 The sketch graph below was obtained for the equilibrium mixture.



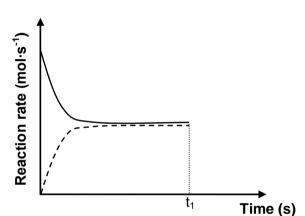
Redraw the graph above in your ANSWER BOOK. On the same set of axes, complete the graph showing the effect of the catalyst on the reaction rates.

Initially 0,16 mol $H_2(g)$ and excess S(s) are sealed in a 2 dm³ container and the system is allowed to reach equilibrium at 90 °C.

An exact amount of $Pb(NO_3)_2$ solution is now added to the container so that ALL the $H_2S(g)$ present in the container at EQUILIBRIUM is converted to PbS(s) according to the following balanced equation: $Pb(NO_3)_2(aq) + H_2S(g) \rightarrow PbS(s) + 2HNO_3(aq)$

The mass of the PbS precipitate is 2,39 g.

7.4 Calculate the equilibrium constant K_c for the reaction $H_2(g) + S(s) \Rightarrow H_2S(g)$ at 90 °C.



(2)

(1)

(9) **[18]**

(2)

ACIDS AND BASES

1.1

Nitric acid (HNO_3), an important acid used in industry, is a strong acid.

Give a reason why nitric acid is classified as a strong acid. (1)



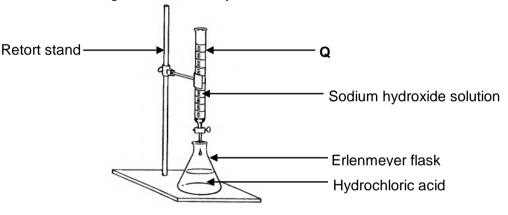
1.2.3

QUESTION 1

Write down the NAME or FORMULA of the conjugate base of nitric acid. (1)

- Calculate the pH of a 0,3 mol·dm⁻³ nitric acid solution.
- 1.2 A laboratory technician wants to determine the percentage purity of magnesium oxide. He dissolves a 4,5 g sample of the magnesium oxide in 100 cm³ hydrochloric acid of concentration 2 mol·dm⁻³.
 - 1.2.1 Calculate the number of moles of hydrochloric acid added to the magnesium oxide.

He then uses the apparatus below to titrate the EXCESS hydrochloric acid in the above solution against a sodium hydroxide solution.



1.2.2 Write down the name of apparatus **Q** in the above diagram.

The following indicators are available for the titration:

INDICATOR	pH RANGE	
Α	3,1 – 4,4	
В	6,0-7,6	
С	8,3 – 10,0	

(1)

(3)

(1)

(3)

(3)

(3)

- Which ONE of the above indicators (A, B or C) is most suitable to indicate the exact endpoint in this titration? Give a reason for the answer.
- 1.2.4 During the titration, the technician uses distilled water to wash any sodium hydroxide spilled against the sides of the Erlenmeyer flask into the solution. Give a reason why the addition of distilled water to the Erlenmeyer flask will not influence the results.
- 1.2.5 At the endpoint of the titration he finds that 21 cm³ of a 0.2 mol dm⁻³ sodium hydroxide solution has neutralised the EXCESS hydrochloric acid. Calculate the number of moles of hydrochloric acid in excess.
- 1.2.6 The balanced equation for the reaction between hydrochloric acid and $MgO(s) + 2HC\ell(aq) \rightarrow MgC\ell_2(aq) + H_2O(\ell)$ magnesium oxide is:

Calculate the percentage purity of the magnesium oxide. Assume that only the magnesium oxide in the 4.5 g sample reacted with the acid.

(5) [21]

(1)

(1)

(6)

(2)

QUESTION 2

2.1.1

2.1.2

2.1 Sulphuric acid is a diprotic acid.

Define an *acid* in terms of the Lowry-Brønsted theory. (2)

Give a reason why sulphuric acid is referred to as *a diprotic acid*. (1)

2.2 The hydrogen carbonate ion <u>can act as both an acid and a base</u>. It reacts with water according to the following balanced equation:

$$HCO_{3}^{-}(aq) + H_{2}O(\ell) \rightleftharpoons H_{2}CO_{3}(aq) + OH^{-}(aq)$$

- 2.2.1 Write down ONE word for the underlined phrase.
- 2.2.2 $HCO_3^-(aq)$ acts as base in the above reaction. Write down the formula of the conjugate acid of $HCO_3^-(aq)$.
- 2.3 A learner accidentally spills some sulphuric acid of concentration 6 mol·dm⁻³ from a flask on the laboratory bench. Her teacher tells her to neutralise the spilled acid by sprinkling sodium hydrogen carbonate powder onto it. The reaction that takes place is: (Assume that the H_2SO_4 ionises completely.)

$$H_2SO_4(aq) + 2NaHCO_3(s) \rightarrow Na_2SO_4(aq) + 2H_2O(\ell) + 2CO_2(g)$$

The fizzing, due to the formation of carbon dioxide, stops after the learner has added 27 g sodium hydrogen carbonate to the spilled acid.

2.3.1 Calculate the volume of sulphuric acid that spilled. Assume that all the sodium hydrogen carbonate reacts with all the acid.

The learner now dilutes some of the 6 mol \cdot dm⁻³ sulphuric acid solution in the flask to 0,1 mol \cdot dm⁻³.

2.3.2 Calculate the volume of the 6 mol·dm⁻³ sulphuric acid solution needed to prepare 1 dm³ of the dilute acid.

During a titration 25 cm³ of the 0,1 mol·dm⁻³ sulphuric acid solution is added to an Erlenmeyer flask and titrated with a 0,1 mol·dm⁻³ sodium hydroxide solution.

- 2.3.3 The learner uses bromothymol blue as indicator. What is the purpose of this indicator?
- 2.3.4 Calculate the pH of the solution in the flask after the addition of 30 cm³ of sodium hydroxide. The endpoint of the titration is not yet reached at this point.

(8) **[22]**

(1)

Anhydrous oxalic acid is an example of <u>an acid that can donate two protons</u> and thus ionises in two steps as represented by the equations below:

I:

$$H(COO)_{2}^{-}(aq) + H_{2}O(\ell) \Rightarrow H_{3}O^{+}(aq) + H(COO)_{2}^{-}(aq)$$

 $H(COO)_{2}^{-}(aq) + H_{2}O(\ell) \Rightarrow H_{3}O^{+}(aq) + (COO)_{2}^{2-}(aq)$

3.1 Write down:

3.2

3.3

3.1.1	ONE word for the underlined phrase in the above sentence	(1)
3.1.2	The FORMULA of each of the TWO bases in reaction II	(2)
3.1.3	The FORMULA of the substance that acts as ampholyte in reactions I and II . Give a reason for the answer.	(2)
Give a rea	son why oxalic acid is a weak acid.	(1)
A standar dissolving	d solution of $(COOH)_2$ of concentration 0,20 mol·dm ⁻³ is prepared by a certain amount of $(COOH)_2$ in water in a 250 cm ³ volumetric flask.	

Calculate the mass of $(COOH)_2$ needed to prepare the standard solution. (4)

3.4 During a titration 25 cm³ of the standard solution of $(COOH)_2$ prepared in QUESTION 7.3 is neutralised by a sodium hydroxide solution from a burette.

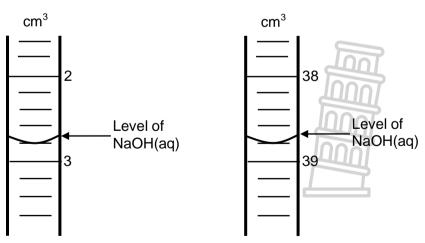
The balanced equation for the reaction is:

 $(COOH)_2(aq) + 2NaOH(aq) \rightarrow (COONa)_2(aq) + 2H_2O(l)$

The diagrams below show the burette readings before the titration commenced and at the endpoint respectively.

Before the titration

At the endpoint



- 3.4.1 Use the burette readings and calculate the concentration of the sodium hydroxide solution.
- 3.4.2 Write down a balanced equation that explains why the solution has a pH greater than 7 at the endpoint.

(5)

(3) **[18]**

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QUESTION 4

4.1 Ammonium chloride crystals, NH₄Cl(s), dissolve in water to form ammonium and chloride ions. The <u>ammonium ions react with water</u> according to the balanced equation below:

$$NH_4^+$$
 (aq) + $H_2O(\ell) \rightleftharpoons NH_3(aq) + H_3O^+(aq)$

Write down the name of the process described by the underlined sentence.

- Is ammonium chloride ACIDIC or BASIC in aqueous solution? Give a reason for the answer.
- 4.2 A certain fertiliser consists of 92% ammonium chloride. A sample of mass x g of this fertiliser is dissolved in 100 cm³ of a 0,10 mol·dm⁻³ sodium hydroxide solution, NaOH(aq). The NaOH is in excess. The balanced equation for the reaction is:

$$NH_4Cl(s) + NaOH(aq) \rightarrow NH_3(g) + H_2O(l) + NaCl(aq)$$

4.2.1 Calculate the number of moles of sodium hydroxide in which the sample is dissolved.

During a titration, 25 cm³ of the excess sodium hydroxide solution is titrated with a 0,11 mol·dm⁻³ hydrochloric acid solution, HC ℓ (aq). At the endpoint it is found that 14,55 cm³ of the hydrochloric acid was used to neutralise the sodium hydroxide solution according to the following balanced equation:

$$HC\ell(aq) + NaOH(aq) \rightarrow NaC\ell(aq) + H_2O(\ell)$$

4.3 Calculate the pH of a 0,5 mol·dm⁻³ sodium hydroxide solution at 25 °C.

QUESTION 5

- 5.1 Define an *acid* in terms of the Lowry-Brønsted theory. (2)
- 5.2 Carbonated water is an aqueous solution of carbonic acid, H_2CO_3 . $H_2CO_3(aq)$ ionises in two steps when it dissolves in water.

5.2.1	Write down the FORMULA of the conjugate base of $H_2CO_3(aq)$.	(1)
-------	---	-----

- 5.2.2 Write down a balanced equation for the first step in the ionisation of carbonic acid. (3)
- 5.2.3 The pH of a carbonic acid solution at 25 °C is 3,4. Calculate the hydroxide ion concentration in the solution. (5)
- 5.3 X is a monoprotic acid.
 - 5.3.1 State the meaning of the term *monoprotic*.
 - 5.3.2 A sample of acid **X** is titrated with a standard sodium hydroxide solution using a suitable indicator. At the endpoint it is found that 25 cm³ of acid **X** is neutralised by 27,5 cm³ of the sodium hydroxide solution of concentration 0,1 mol·dm⁻³. Calculate the concentration of acid **X**.
 - 5.3.3 The concentration of H_3O^+ ions in the sample of acid **X** is 2,4 x 10⁻⁴ mol·dm⁻³. Is acid **X** a WEAK or a STRONG acid? Explain the answer by referring to the answer in QUESTION 5.3.2.

Terms, definitions, questions and answers

4.1.1

(3)

(4) **[18]**

(1)

(2)

)

(1)

(5)

....

(3) **[20]**

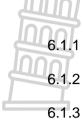
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(2)

QUESTION 6

6.1 Hydrogen carbonate ions react with water according to the following balanced equation:

$$HCO_{3}^{-}(aq) + H_{2}O(\ell) \rightleftharpoons H_{2}CO_{3}(aq) + OH^{-}(aq)$$



Define an acid according to the Lowry-Brønsted theory.

Write down the FORMULAE of the two acids in the equation above. (2)

- Write down the formula of a substance in the reaction above that can act as an ampholyte. (1)
- 6.2 During an experiment 0,50 dm³ of a 0,10 mol·dm⁻³ HCl solution is added to 0,80 dm³ of a NaHCO₃ solution of concentration 0,25 mol·dm⁻³. The balanced equation for the reaction is:

 $NaHCO_3(aq) + HCl(aq) \rightarrow NaCl(aq) + CO_2(g) + H_2O(l)$

Calculate the:

6.	2.1	Concentration of the hydroxide ions in the solution on completion of the reaction	(8)
6.	2.2	pH of the solution on completion of the reaction	(4) [17]

QUESTION 7

- 7.1 A learner dissolves ammonium chloride ($NH_4C\ell$) crystals in water and measures the pH of the solution.
 - 7.1.1Define the term *hydrolysis* of a salt.(2)
 - 7.1.2 Will the pH of the solution be GREATER THAN, SMALLER THAN or EQUAL TO 7? Write a relevant equation to support your answer. (3)
- 7.2 A sulphuric acid solution is prepared by dissolving 7,35 g of $H_2SO_4(l)$ in 500 cm³ of water.

7.2.1 Calculate the number of moles of H_2SO_4 present in this solution.

Sodium hydroxide (NaOH) pellets are added to the 500 cm³ H₂SO₄ solution. The balanced equation for the reaction is:

 $H_2SO_4(aq)$ + 2NaOH(s) \rightarrow Na₂SO₄(aq) + 2H₂O(ℓ)

After completion of the reaction, the pH of the solution was found to be 1,3. Assume complete ionisation of H_2SO_4 .

7.2.2 Calculate the mass of NaOH added to the H_2SO_4 solution. Assume that the volume of the solution does not change.

(9) **[16]**

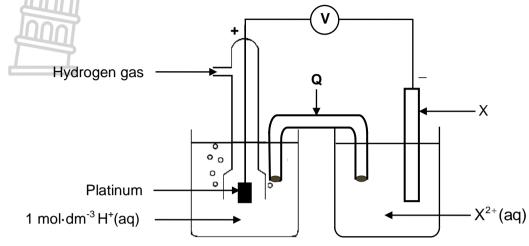
(2)

(2)

GALVANIC CELLS

QUESTION 1

A standard electrochemical cell is set up using a standard hydrogen half-cell and a standard $X|X^{2+}$ half-cell as shown below. A voltmeter connected across the cell, initially registers 0,31 V.



- 1.1 Besides concentration write down TWO conditions needed for the hydrogen half-cell to function under standard conditions.
- 1.2 Give TWO reasons, besides being a solid, why platinum is suitable to be used as electrode in the above cell. (2)
- 1.3 Write down the:

1.5

1.3.1	NAME of component Q	(1)

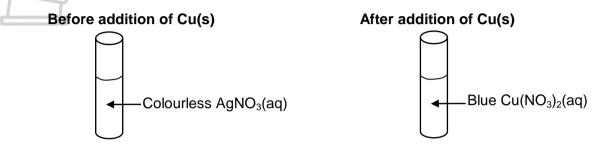
- 1.3.2 Standard reduction potential of the $X|X^{2+}$ half-cell (1)
- 1.3.3 Half-reaction that takes place at the cathode of this cell (2)
- 1.4 The hydrogen half-cell is now replaced by a $\mathbf{M}|\mathbf{M}^{2+}$ half-cell. The cell notation of this cell is:

	M (s) M ²⁺ (aq) X ²⁺ (aq) X(s)	
The initia	I reading on the voltmeter is now 2,05 V.	
1.4.1	Identify metal M . Show how you arrived at the answer.	(5)
1.4.2	Is the cell reaction EXOTHERMIC or ENDOTHERMIC?	(1)
	ding on the voltmeter becomes zero after using this cell for several hours. eason for this reading by referring to the cell reaction.	(1) [15]

A learner conducts two experiments to investigate the reaction between copper (Cu) and a silver nitrate solution, $AgNO_3(aq)$.

EXPERIMENT 1

The learner adds a small amount of copper (Cu) powder to a test tube containing silver nitrate solution, $AgNO_3(aq)$. The solution changes from colourless to blue after a while.



2.1 Define the term *oxidising agent*.

(2)

2.2 Explain why the solution turns blue by referring to the relative strength of oxidising agents. (4)

EXPERIMENT 2

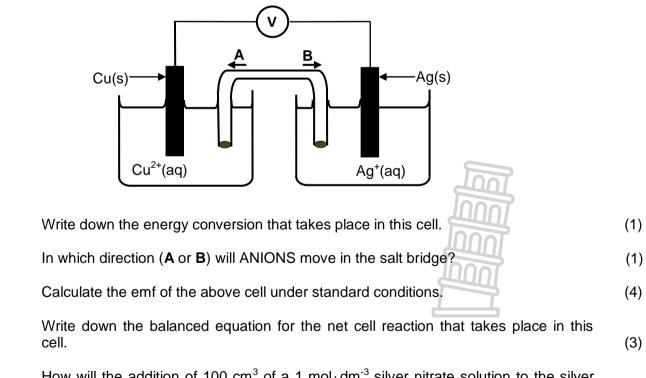
2.3

2.4

2.5

2.6

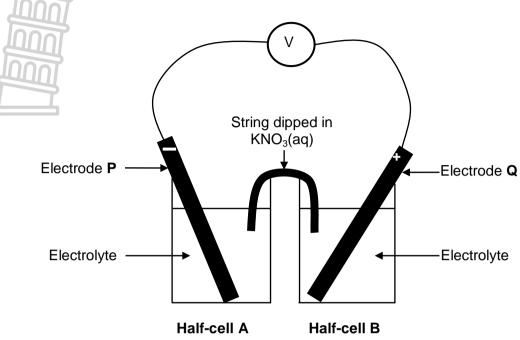
The learner now sets up a galvanic cell as shown below. The cell functions under standard conditions.



2.7 How will the addition of 100 cm³ of a 1 mol·dm⁻³ silver nitrate solution to the silver half-cell influence the initial emf of this cell? Write down only INCREASES, DECREASES or REMAINS THE SAME.

(1) **[16]**

Learners set up an electrochemical cell, shown in the simplified diagram below, using magnesium and lead as electrodes. Nitrate solutions are used as electrolytes in both half-cells.



3.1	What type in this cell	of reaction (NEUTRALISATION, REDOX or PRECIPITATION) takes place ?	(1)
3.2	Which ele	ctrode, P or Q , is magnesium? Give a reason for the answer.	(2)
3.3	Write dow	n the:	
	3.3.1	Standard conditions under which this cell functions	(2)
	3.3.2	Cell notation for this cell	(3)
	3.3.3	NAME or FORMULA of the oxidising agent in the cell	(1)
3.4	Calculate	the initial emf of the cell above under standard conditions.	(4)
3.5		ne voltmeter reading change if the: vn only INCREASES, DECREASES or REMAINS THE SAME.)	
	3.5.1	Size of electrode P is increased	(1)
	3.5.2	Initial concentration of the electrolyte in half-cell B is increased	(1) [15]

(3)

QUESTION 4

Learners are given the following two unknown half-cells:

Half-cell 1: $Q^{2+}(aq) | Q(s)$ Half-cell 2: Pt | $R_2(g) | R^{-}(aq)$

During an investigation to identify the two half-cells, the learners connect each half-cell alternately to a $Cd^{2+}(aq) | Cd(s)$ half-cell under standard conditions. For each combination of two half-cells, they write down the net cell reaction and measure the cell potential. The results obtained for the two half-cell combinations are given in the table below.

COMBINATION	NET CELL REACTION	CELL POTENTIAL
I	$\mathbf{Q}^{2+}(aq) + Cd(s) \rightarrow Cd^{2+}(aq) + \mathbf{Q}(s)$	0,13 V
II	$\mathbf{R}_2(g)$ + Cd(s) \rightarrow Cd ²⁺ (aq) + 2 \mathbf{R}^- (aq)	1,76 V

Write down THREE conditions needed for these cells to function as standard cells.

			()
4.2	For Con	nbination I, identify:	
	4.2.1	The anode of the cell	(1)
	4.2.2	Q by using a calculation	(5)
4.3	For Con	nbination II, write down the:	
	4.3.1	Oxidation half-reaction	(2)
	4.3.2	NAME or FORMULA of the metal used in the cathode compartment	(1)
4.4	Arrange	the following species in order of INCREASING oxidising ability:	
	Q^{2+} ; R_2	; Cd ²⁺	
	Explain	fully how you arrived at the answer. A calculation is NOT required.	(4) [16]

QUESTION 5

4.1

An electrochemical cell consisting of half-cells **A** and **B** is assembled under standard conditions as shown below.

Half-cell A	Pt, Cl ₂ (101,3 kPa) Cl ⁻ (1 mol·dm ⁻³)
Half-cell B	Mg ²⁺ (1 mol·dm ⁻³) Mg(s)

5.1	At which half-cell, A or B,	are electrons released into the external circuit?	(1)

- 5.2 Write down the:
 - 5.2.1 Reduction half-reaction that takes place in this cell (2)
 - 5.2.2 NAME or FORMULA of the substance whose oxidation number DECREASES (1)
- 5.3 Calculate the initial cell potential of this cell when it is in operation.
- 5.4 Write down an observation that will be made in half-cell **B** as the cell operates. Give a reason for the answer. (2)

<u>[10]</u>

(4)

Magnesium (Mg) reacts with a dilute hydrochloric acid solution, HCl(aq), according to the following balanced equation:

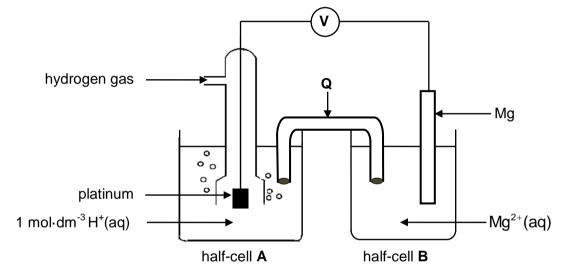
 $Mg(s) + 2HC\ell(aq) \rightarrow MgC\ell_2(aq) + H_2(q)$

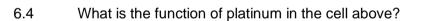
6.1 Give a reason why the reaction above is a redox reaction. (1) 6.2 Write down the FORMULA of the oxidising agent in the reaction above. (1)

It is found that silver does not react with the hydrochloric acid solution.

6.3 Refer to the relative strengths of reducing agents to explain this observation. (3)

The reaction of magnesium with hydrochloric acid is used in an electrochemical cell, as shown in the diagram below. The cell functions under standard conditions.





Write down the: 6.5

6.6

6.7

6.5.1	Energy conversion that takes place in this cell	(1)	
6.5.2	Function of Q	(1)	
6.5.3	Half-reaction that takes place at the cathode	(2)	
6.5.4	Cell notation of this cell	(3)	
Calcu	ulate the initial emf of this cell.	(4)	
How will the addition of concentrated acid to half-cell A influence the answer to			

QUESTION 6.6? Choose from INCREASES, DECREASES or REMAINS THE SAME.

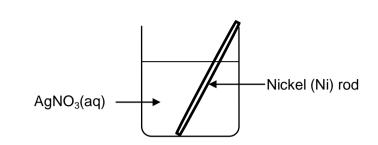
(1)

(2)

QUESTION 7

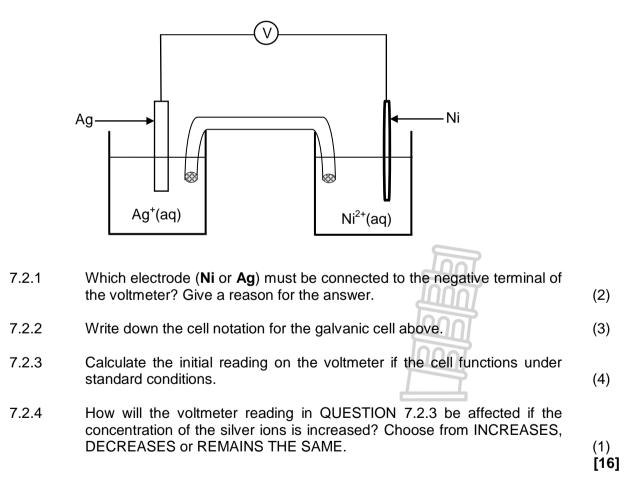


A nickel (Ni) rod is placed in a beaker containing a silver nitrate solution, $AgNO_3(aq)$ and a reaction takes place.



Write down the:

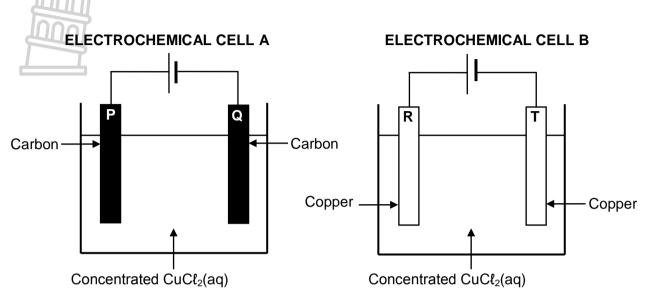
- 7.1.1 NAME or FORMULA of the electrolyte (1)
- 7.1.2 Oxidation half-reaction that takes place
- 7.1.3 Balanced equation for the net (overall) redox reaction that takes place (3)
- 7.2 A galvanic cell is now set up using a nickel half-cell and a silver half-cell.



ELECTROLYTIC CELLS



The simplified diagrams below represent two electrochemical cells, **A** and **B**. A concentrated copper(II) chloride solution is used as electrolyte in both cells.



- 1.1 Are **A** and **B** ELECTROLYTIC or GALVANIC cells?
- 1.2 Which of the electrodes (**P**, **Q**, **R** or **T**) will show a mass increase? Write down a halfreaction to motivate the answer. (4)
- 1.3 Write down the NAME or FORMULA of the product formed at:

1.3.1	Electrode P	(1)	
-------	-------------	-----	--

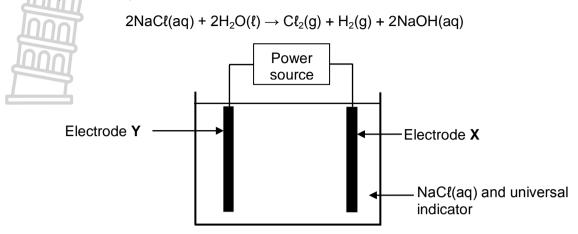
- 1.3.2 Electrode **R** (1)
- 1.4
 Fully explain the answer to QUESTION 1.3.2 by referring to the relative strengths of the reducing agents involved.
 (3)

 [10]
 [10]



(1)

The apparatus below is used to demonstrate the electrolysis of a concentrated sodium chloride solution. Both electrodes are made of carbon. A few drops of universal indicator are added to the electrolyte. The equation for the net cell reaction is:



Initially the solution has a green colour. Universal indicator becomes red in acidic solutions and purple in alkaline solutions.

2.1 Define the term *electrolyte*.

When the power source is switched on, the colour of the electrolyte around electrode **Y** changes from green to purple.

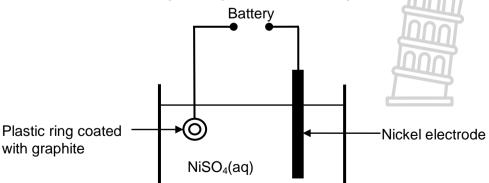
2.2 Write down the:

2.2.1	Half-reaction that takes place at electrode Y	(2)
2.2.2	NAME or FORMULA of the gas released at electrode ${\bf X}$	(1)

2.3 Refer to the Table of Standard Reduction Potentials to explain why hydrogen gas, and not sodium, is formed at the cathode of this cell. (2) [7]

QUESTION 3

The diagram below shows a simplified electrolytic cell that can be used to electroplate a plastic ring with nickel. Prior to electroplating the ring is covered with a graphite layer.



- 3.1 Define the term *electrolyte*.
- 3.2 Give ONE reason why the plastic ring must be coated with graphite prior to electroplating. (1)

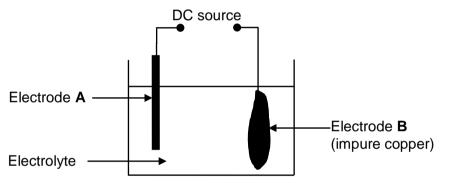
(2)

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3.3 Write down the:				
3.3.1 Half-reaction that occurs at the plastic ring	(2)			
NAME or FORMULA of the reducing agent in the cell. Give a reason for the answer.	(2)			
3.4 Which electrode, the RING or NICKEL , is the cathode? Give a reason for answer.	or the (2)			
The nickel electrode is now replaced with a carbon rod.				

3.5 How will the concentration of the electrolyte change during electroplating? Write down only INCREASES, DECREASES or NO CHANGE. Give a reason for the answer.

QUESTION 4

The simplified diagram below represents an electrochemical cell used for the purification of copper.



- 4.1 Define the term *electrolysis*.
- 4.2 Give a reason why a direct-current (DC) source is used in this experiment. (1)
- 4.3 Write down the half-reaction which takes place at electrode **A**.
- 4.4 Due to small amounts of zinc impurities in the impure copper, the electrolyte becomes contaminated with Zn²⁺ ions.

Refer to the attached Table of Standard Reduction Potentials to explain why the Zn^{2+} ions will not influence the purity of the copper obtained during this process. (3)

4.5 After the purification of the impure copper was completed, it was found that $2,85 \times 10^{-2}$ moles of copper were formed.

The initial mass of electrode **B** was 2,0 g. Calculate the percentage of copper that was initially present in electrode **B**.

(4) **[12]**

(2)

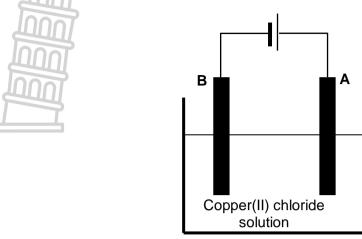
(2)

(2) [11]

(1)

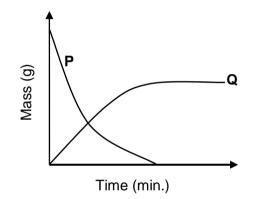
QUESTION 5

The electrochemical cell below is set up to demonstrate the purification of copper.



5.1 Write down the type of electrochemical cell illustrated above.

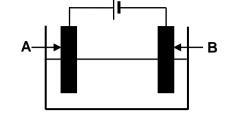
The graphs below show the change in mass of the electrodes whilst the cell is in operation.



5.2	Define a reducing agent in terms of electron transfer.	(2)
5.3	Which graph represents the change in mass of electrode A ?	(1)
5.4	Write down the half-reaction that takes place at electrode A.	(2)
5.5	Electrodes A and B are now replaced by graphite electrodes. It is observed that chlorine gas (Cl_2) is released at one of the electrodes.	
	At which electrode (A or B) is chlorine gas formed? Fully explain how it is formed.	(3) [9]

The diagram below shows an electrochemical cell used to purify copper. <u>A solution that</u> <u>conducts electricity</u> is used in the cell.





6.1 Write down:

6.1.1	ONE word for the underlined phrase above the diagram	(1)
6.1.2	The type of electrochemical cell illustrated above	(1)
In which circuit?	direction (from A to B or from B to A) will electrons flow in the external	(1)
Which e	lectrode (A or B) is the:	
6.3.1	Cathode	(1)
6.3.2	Impure copper	(1)
	I the mass of electrode A change as the reaction proceeds? Choose from SES, DECREASES or REMAINS THE SAME. Give a reason for the answer.	(2)

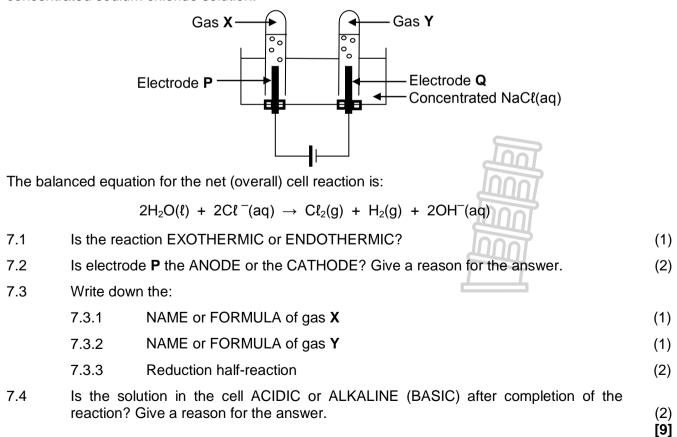
QUESTION 7

6.2

6.3

6.4

In the electrochemical cell below, carbon electrodes are used during the electrolysis of a concentrated sodium chloride solution.

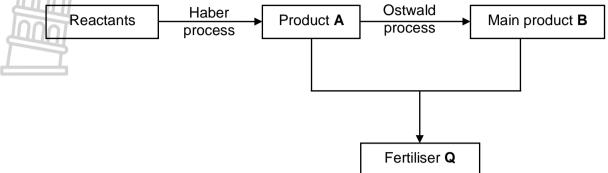


[7]

FERTILISERS

QUESTION 1

1.1 The flow diagram below shows the processes involved in the industrial preparation of fertiliser **Q**.



Write down the:

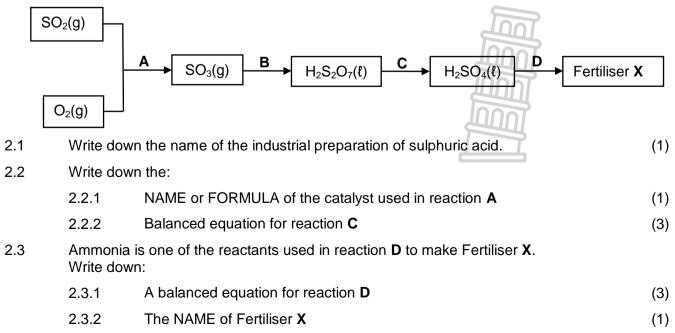
- 1.1.1 NAMES or FORMULAE of the reactants used in the Haber process (2)
- 1.1.2 Balanced equation for the formation of fertiliser **Q**
- 1.2 The diagram below shows a bag of NPK fertiliser.



Calculate the mass of nitrogen in the bag.

QUESTION 2

Reactions A, B, C and D in the flow diagram below represent the manufacturing of Fertiliser X.

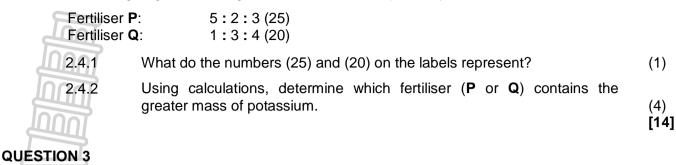


(4) **[9]**

(3)

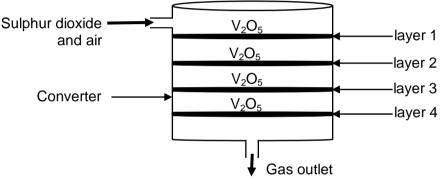
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2.4 Two 50 kg bags, containing fertilisers **P** and **Q** respectively, are labelled as follows:



The industrial process for the preparation of sulphuric acid involves a series of stages.

The second stage in this process involves the conversion of sulphur dioxide into sulphur trioxide in a converter as illustrated below. In the converter the gases are passed over vanadium pentoxide (V_2O_5) placed in layers as shown below.



3.1 Write down the:

3.1.1 Balanced equation for the reaction taking place in the converter

3.1.2 Function of the vanadium pentoxide

The table below shows data obtained during the second stage.

VANADIUM PENTOXIDE LAYER	TEMPERATURE OF GAS BEFORE THE REACTION (°C)	TEMPERATURE OF GAS AFTER THE REACTION (°C)	PERCENTAGE OF REACTANT CONVERTED TO PRODUCT
1	450	600	66
2	450	518	85
3	450	475	93
4	450	460	99,5

- 3.2 Is the reaction in the second stage EXOTHERMIC or ENDOTHERMIC? Refer to the data in the table to give a reason for the answer. (2)
- 3.3 After the conversion at each layer the gases are cooled down to 450 °C. Fully explain why the gases must be cooled to this temperature. (3)
- 3.4 During the third stage sulphur trioxide is dissolved in sulphuric acid rather than in water to produce oleum.
 - 3.4.1 Write down the FORMULA of oleum.
 - 3.4.2 Give a reason why sulphur trioxide is not dissolved in water. (1)
- 3.5 Sulphuric acid reacts with ammonia to form a fertiliser. Write down a balanced equation for this reaction.

(1)

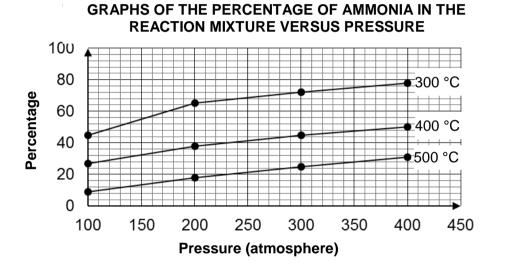
(3)

(1)

Ammonia is an important fertiliser. Large amounts are prepared from hydrogen and nitrogen in industry.

4.1	For the inc	dustrial preparation of ammonia, write down:	
Ë	4.1.1	The name of the process used	(1)
4	4.1.2	A balanced equation for the reaction that occurs	(3)
	4.1.3	The source of nitrogen	(1)
4.0	The state	of expression of a property with the property of a property of a line in the industrial	

4.2 The yield of ammonia changes with temperature and pressure during its industrial preparation. The graphs below show how the percentage of ammonia in the reaction mixture that leaves the reaction vessel varies under different conditions.



- 4.2.1 Use the appropriate graph to estimate the percentage of ammonia present in the reaction mixture at 240 atmosphere and 400 °C. (1)
- 4.2.2 State TWO advantages of using high pressure in the preparation of ammonia. (2)
- 4.2.3 The advantage of using a low temperature is the large percentage of ammonia formed. What is the disadvantage of using a low temperature? (1)
- 4.3 Ammonia is also used in the preparation of other fertilisers such as ammonium nitrate. Calculate the mass of nitrogen in a 50 kg bag of pure ammonium nitrate fertiliser.

QUESTION 5

Ammonium nitrate is an important fertiliser. It is produced by reacting nitric acid with ammonia. Both nitric acid and ammonia are prepared on a large scale in industry.

- 5.1 Write down the name of the industrial preparation of nitric acid.
- 5.2 The catalytic oxidation of ammonia is one of the steps in the process named in QUESTION 5.1. Write down the NAMES or FORMULAE of the TWO products formed in this step.
- 5.3 Write down a balanced equation for the preparation of ammonium nitrate.

Terms, definitions, questions and answers

(1)

(3) **[12]**

(2)

(3)

(3)

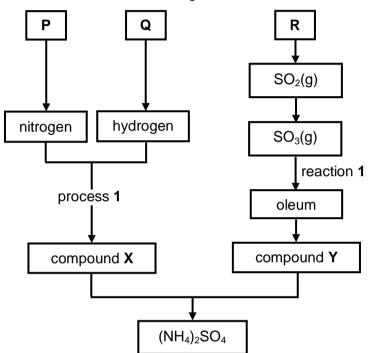
(1) **[10]**

- 5.4 Calculate the mass, in kilogram, of ammonium nitrate that can be made from 6.8×10^4 kg of ammonia and excess nitric acid. (One mole of ammonia produces one mole of ammonium nitrate.)
- 5.5 Ammonium nitrate is often mixed with potassium chloride and ammonium phosphate. Give a reason why it is mixed with these compounds.

QUESTION 6

6.2

A chemical company produces ammonium sulphate, $(NH_4)_2SO_4$, starting from the raw materials **P**, **Q** and **R**, as shown in the flow diagram below.



6.1 Write down the NAME of raw material:

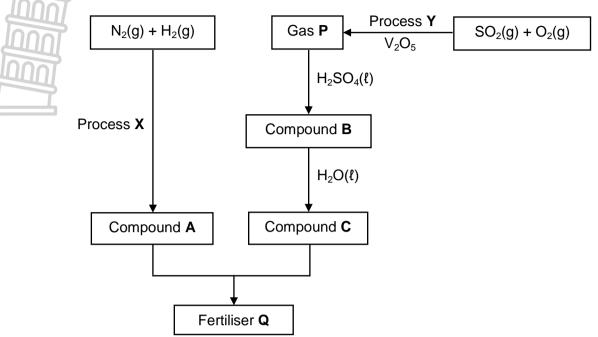
6.1.1	P	(1)
6.1.2	Q	(1)
6.1.3	R	(1)
Write dow	In the:	
6.2.1	NAME of process 1	(1)
6.2.2	NAME of compound X	(1)
6.2.3	FORMULA of compound Y	(1)
6.2.4	Balanced equation for reaction 1	(3)

- 6.3 The company compares the nitrogen content of ammonium sulphate with that of ammonium nitrate, NH_4NO_3 .
 - 6.3.1 Determine, by performing the necessary calculations, which ONE of the two fertilisers has the higher percentage of nitrogen per mass.
 - 6.3.2 Write down the name of the process that should be included in the flow diagram above if the company wants to prepare ammonium nitrate instead of ammonium sulphate.

(1) **[14]**

(4)

7.1 The flow diagram below shows the processes involved in the industrial preparation of fertiliser Q.



Write down the:

7.1.1	Name of process X	(1)
7.1.2	Name of process Y	(1)
7.1.3	NAME or FORMULA of gas P	(1)

- 7.1.4 Balanced equation for the formation of compound B (3)
- 7.1.5 Balanced equation for the formation of fertiliser Q (4)
- 7.2 The diagram below shows a bag of NPK fertiliser of which the NPK ratio is unknown. It is found that the mass of nitrogen in the bag is 4,11 kg and the mass of phosphorus is 0,51 kg.

NAME or FORMULA of gas P

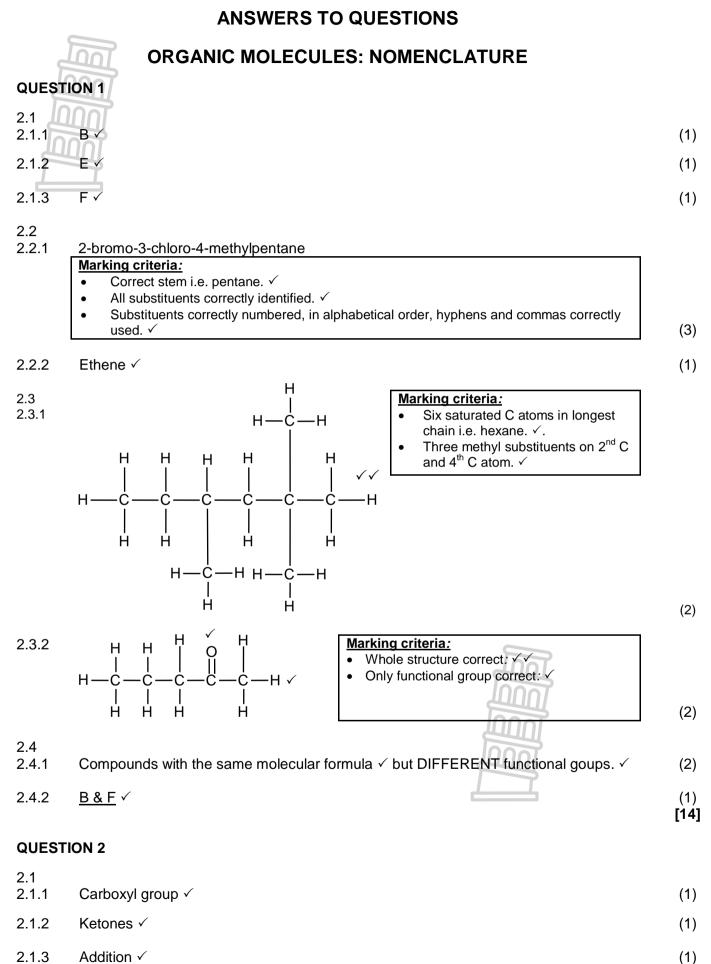


Calculate the NPK ratio of the fertiliser.

(4) [14]

(1)

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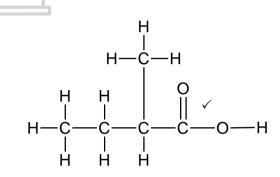


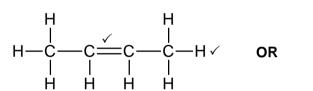
2.1.3 Addition ✓

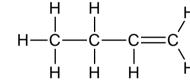
Terms, definitions, questions and answers

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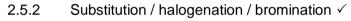


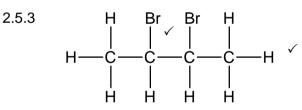






 \checkmark





QUESTION 3

2.4 2.4.1

2.4.2

QULUI		ЩППІ	
3.1 3.1.1	B√		(1)
3.1.2	E✓		(1)
3.1.3 3.2	A ✓		(1)
3.2.1	4-chloro-2,5-dimethylheptane		
	Marking criteria:		

Correct stem i.e. <u>heptane</u>. ✓
 All substituents (chloro and dimethyl) correctly identified. ✓ Substituents correctly numbered, in alphabetical order, hyphens and commas correctly used. ✓

3.2.2 <u>2-methyl</u>√propan-1-ol √

(2)

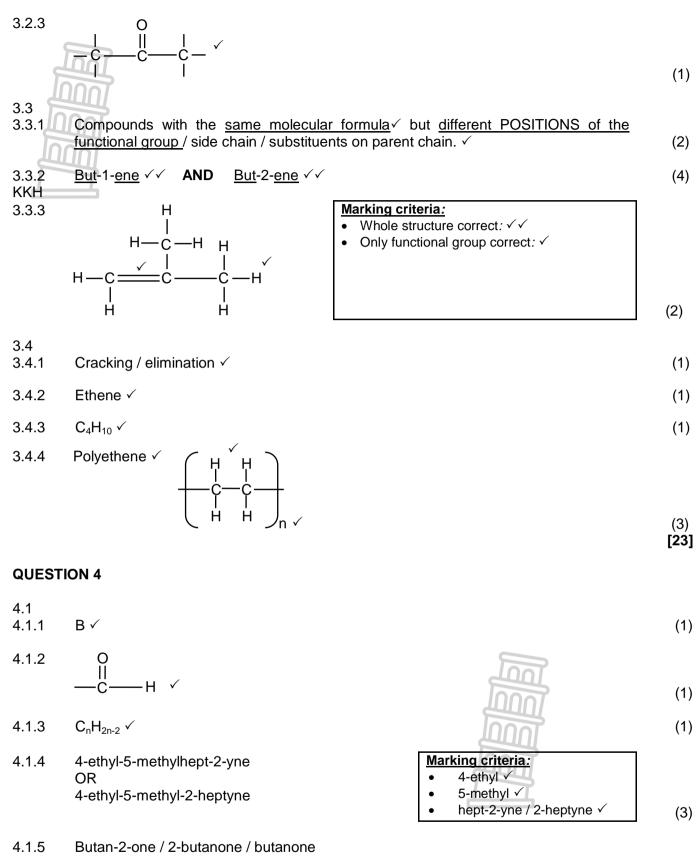
(2)

(1)

(2) **[18]**

(3)

(2)

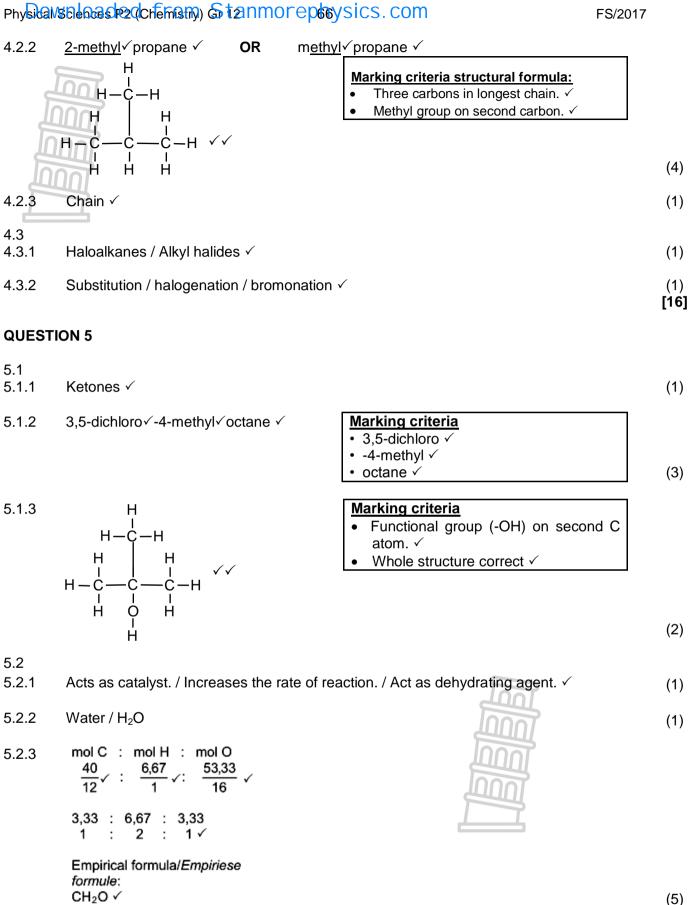


Ma	rking criteria:	
•	Functional group ✓	
•	Whole name correct ✓	(2

4.2

4.2.1 Alkanes √

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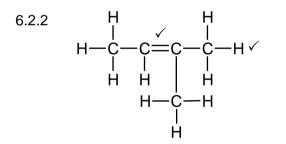


5.2.4 $M(CH_2O) = 30 \text{ g} \cdot \text{mol}^{-1} \checkmark$ Formula-units/Formule-eenhede: $\frac{60}{30} = 2 \checkmark$	
Molecular formula/ <i>Molekulêre</i> formule: C ₂ H ₄ O ₂ ✓	(3)
5.2.5 O H C H	(1)

5.2.6 Methyl v methanoate v

QUESTION 6

- 6.1 6.1.1 E ✓
- 6.1.2 C ✓
- 6.1.3 D ✓
- 6.2
- 6.2.1 Pent-2√-yne√ **OR** 2-pentyne



M	arking criteria <i>:</i>
٠	Whole structure correct: VV
٠	Only functional group correct: \checkmark

(2)

(2) **[19]**

(1)

(1)

(1)

(2)

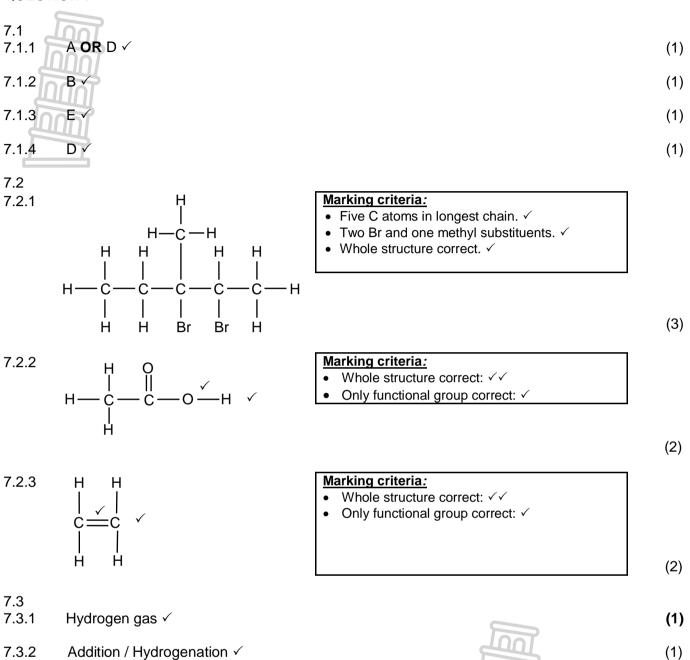
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6.2.3 2-methylbut-1-ene OR 3-methylbut-1-ene

	 Marking criteria: Correct stem i.e. <u>but-1-ene/1-butene</u>. √ Only one type of substituent, methyl, correctly identified. √ Entire name correct. √ 	(3)
6.3 6.3.1	Esters ✓	(1)
6.3.2	Sulphuric acid / $H_2SO_4 \checkmark$	(1)
6.3.3	Methyl√ propanoate √	(2) [14]

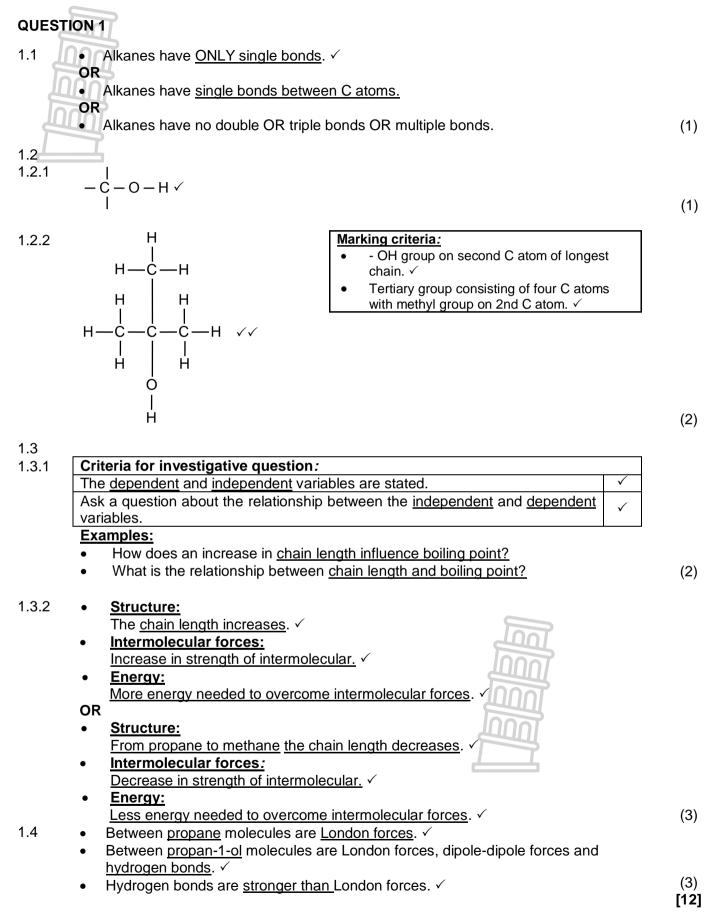
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QUESTION 7





ORGANIC MOLECULES: PHYSICAL PROPERTIES



(1) **[12]**

QUESTION 2

2.1	C-✓	(1)
2.2	Chain length √	(1)
2.2.2	Boiling point ✓	(1) (1)
2.3	London forces ✓	(1)
2.4	Higher than ✓	(1)
		(.)

2.5 Lower than √

- Both compounds **D** and **E** have hydrogen bonding between molecules. \checkmark
- Compound **D** has one site for hydrogen bonding whilst compound **E** has two sites for hydrogen bonding OR can form dimers. ✓
- More energy needed to overcome intermolecular forces in compound E. √ (4)
 [9]

QUESTION 3

3.1	Saturated/√ ANY ONE: • B has <u>ONLY single bonds</u> . √	
	B has <u>single bonds between C atoms.</u> B has no double OD triple hands OD multiple hands	$\langle 0 \rangle$
	B has no double OR triple bonds OR multiple bonds.	(2)
3.2		
3.2.1	- 42 (°C) ✓	(1)
3.2.2	78 (°C) ✓	(1)
3.3	 Between molecules of <u>C/propane</u> are <u>London forces / induced dipole forces</u>. ✓ Between molecules of <u>E/ethanol</u> are (London forces / induced dipole forces and) <u>hydrogen bonds</u>. ✓ <u>Hydrogen bonds</u> / Forces between alcohol molecules are <u>stronger</u>. ✓ 	(3)
3.4	Decrease ✓	
	From A to D:	
	Chain length increases. ✓	
	Strength of intermolecular forces increases. ✓	(4)
	 More energy needed to overcome intermolecular forces. 	(4)

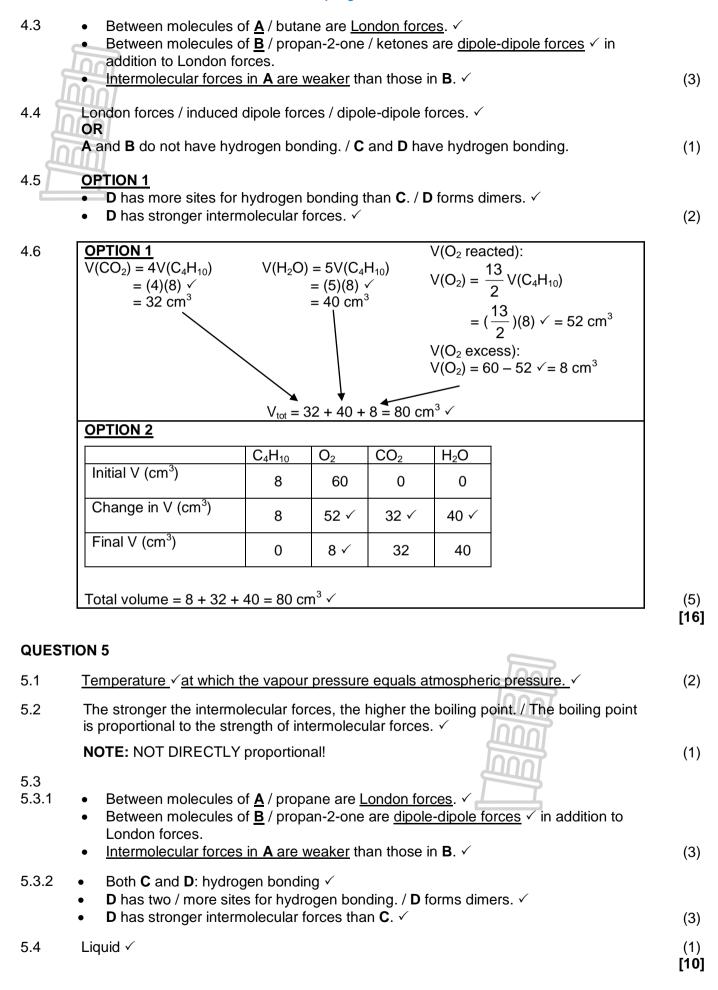
3.5 Higher than √

QUESTION 4

4.1	<u>A bond OR an atom OR a group of atoms</u> \checkmark that determine(s) the (physical and chemical) properties of a group of organic compounds. \checkmark	(2)
4.2 4.2.1	D / ethanoic acid ✓ Lowest vapour pressure. ✓	(2)
4.2.2	A / butane ✓	(1)

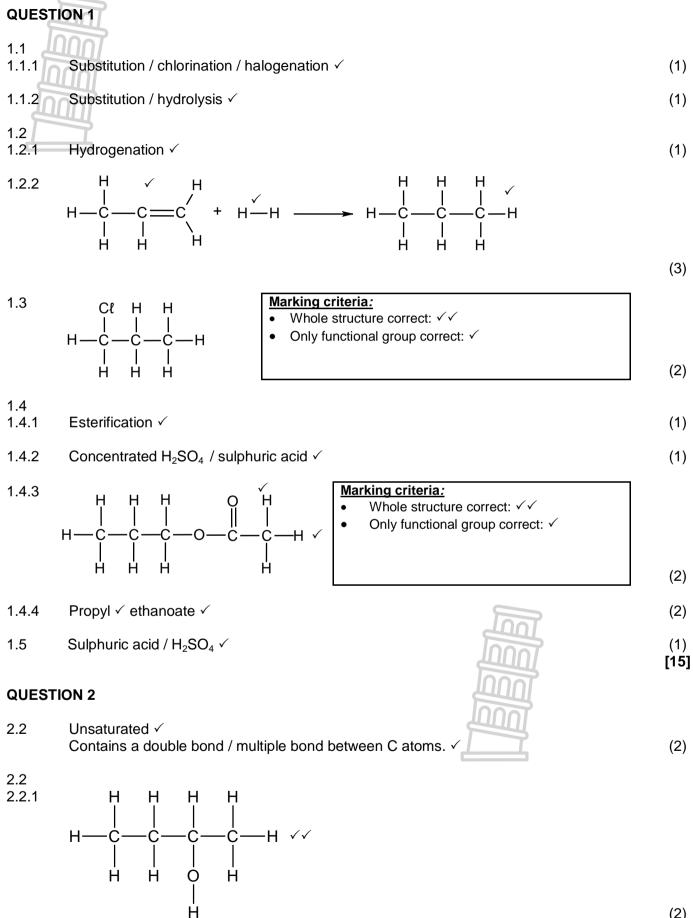
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QULUI		
6.1	Temperature \checkmark at which the vapour pressure equals atmospheric pressure. \checkmark	(2)
6.2	Criteria for conclusion: Image: Conclusion: Dependent and independent variables correctly identified. Image: Conclusion: Relationship between the independent and dependent variables correctly istated. Image: Conclusion: Examples: Image: Conclusion: Boiling point increases with increase in chain length. Image: Conclusion: Boiling point decreases with decrease in chain length. Image: Conclusion: Boiling point is proportional to chain length. Image: Conclusion: NOTE: Boiling point is NOT DIRECTLY proportional to chain length.	(2)
6.3 6.3.1	P✓	(1)
6.3.2	R✓	(1)
6.4	 Between <u>alkane molecules</u> are <u>London forces</u>. ✓ In addition to London forces and dipole-dipole forces each <u>alcohol</u> molecule has one site for <u>hydrogen bonding</u>. ✓ In addition to London forces and dipole-dipole forces each <u>carboxylic acid</u> molecule has two sites for <u>hydrogen bonding</u>. ✓ Intermolecular forces in carboxylic acids are <u>stronger</u> than intermolecular forces in alkanes and alcohols. ✓ More energy is needed to overcome intermolecular forces in carboxylic acids than in the other two compounds. ✓ 	(5) [11]
QUEST	ION 7	
7.1	Compounds with the same molecular formula \checkmark but different structural formulae. \checkmark	(2)
7.2	Chain ✓	(1)
7.3	 From A to C: Structure: <u>Less branched</u> / less compact / larger surface area (over which intermolecular forces act).√ Intermolecular forces: Stronger intermolecular forces. ✓ Energy: More energy needed to overcome intermolecular forces. ✓ 	(3)
7.4	A ✓	(0)
1.4	Lowest boiling point. ✓	(2)
7.5	C_5H_{12} + 8 O_2 ✓ → 5 CO_2 + 6 H_2O ✓ Bal ✓	(3) [11]

ORGANIC MOLECULES: ORGANIC REACTIONS



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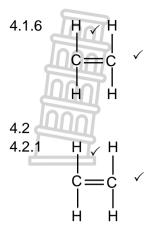
Physical	/ടെപക്ഷിജ്ജർഗ്രഹ്ത്നുണ്ണ) ക്ഷോനനാലാനും sics.com	FS/2017
2.2.2	Addition / hydration \checkmark	(1)
2.3 2.3.1	2-chloro√butane √	(2)
2.3.2	н н н н C − C − C − − C − − н ✓✓ C н н н	(1)
2.4 2.4.1	H₂O OR dilute NaOH / KOH ✓ Mild heat ✓	(1)
2.4.2	Substitution / hydrolysis ✓	(1)
2.4.3	$C_4H_9C\ell$ + NaOH ✓ → $C_4H_{10}O$ + NaCℓ ✓ bal. ✓ OR	
	$C_4H_9C\ell + H_2O \checkmark \rightarrow C_4H_{10}O + HC\ell \checkmark$ bal. \checkmark	(3) [15]
QUEST	ION 3	
3.1 3.1.1	Hydrolysis ✓	(1)
3.1.2	(Mild) heat ✓ <u>Dilute</u> (strong) base/aqueous base ✓	(2)
3.1.3	Ethanol 🗸	(1)
3.2 3.2.1	Esterification ✓	(1)
3.2.2	O ✓ H C O H ✓ ✓ ✓ ✓ H C O H ✓	(2)
3.2.3	Ethyl√ methanoate √	(2) [9]
QUEST	ION 4	
4.1 4.1.1	Esterification / Condensation ✓	(1)
4.1.2	ooi ✓	
4.1.3	Propanoic acid ✓	(1) (1)

(1)

4.1.4 Dehydration / elimination \checkmark

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4.1.5 (Concentrated) H_2SO_4 / sulphuric acid / H_3PO_4 / phosphoric acid



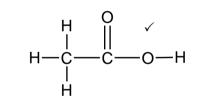
PO ₄	′ phosphoric acid ✓	(
Ma	king criteria:	
•	Only functional group correct: ✓	
•	Whole structure correct: $\checkmark\checkmark$	
Ma	king criteria:	7
<u>Ma</u> ●	<pre>'king criteria: Functional group correct: ✓ Whole structure correct: ✓✓</pre>	7

4.2.2 Addition ✓

QUESTION 5

5.1 5.1.1	Addition 🗸	(1)
5.1.2	Polyethene ✓	(1)
5.2 5.2.1	Chloro√ethane √	(2)
5.2.2	Hydrohalogenation/hydrochlorination 🗸	(1)

5.3 5.3.1



5.3.2	HCℓ / hydrogen chloride ✓		(1)
5.4 5.4.1	Saturated \checkmark There are no double / multiple bonds between C atoms. \checkmark		(2)
5.4.2	H₂ / hydrogen gas√	<u>han</u>	(1)
5.4.3	$2C2H_6 + 7O_2 \checkmark \rightarrow 4CO_2 + 6H_2O \checkmark$ Bal. \checkmark		(3)

QUESTION 6

6.1 6.1.1	Addition / Hydrogenation ✓	(1)
6.1.2	Elimination / Dehydrohalogenation / Dehydrobromination \checkmark	(1)
6.1.3	Substitution / Halogenation / Bromination \checkmark	(1)
Terms, c	definitions, questions and answers	

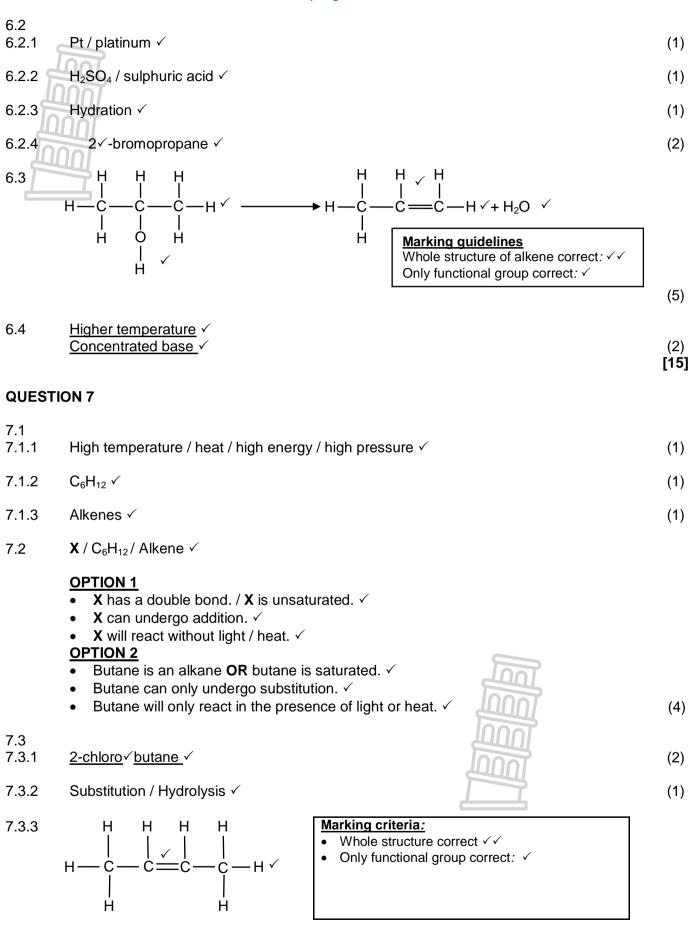
(2)

(1) [**10**]

(2)

[14]

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7.3.4 Hydration √

(1) [**13**]

REACTION RATE AND ENERGY IN CHEMICAL REACTIONS

QUESTION 1

1.1 Change in co	oncentration of products / reactants \checkmark per unit time. \checkmark	(2)
1.2 1.2.1 Temperature	9 √	(1)
1.2.2 Rate of reac	tion / <u>Volume of gas</u> formed <u>per unit time</u> \checkmark	(1)

1.3 • Larger mass / amount / surface area. ✓

• <u>More effective collisions per unit time</u>. / More particles collide with sufficient kinetic energy & correct orientation per unit time. √√

(3)

1.4 Marking criteria

Compare Exp.1 with	The reaction in <u>exp. 1 is faster</u> than in <u>exp. 2</u> due to the <u>higher</u> <u>acid concentration</u> .	\checkmark	
Exp. 2:	Therefore the <u>gradient</u> of the graph representing <u>exp. 1</u> is <u>greater / steeper</u> than that of <u>exp. 2</u> .	~	
Compare Exp. 1 with	The reaction in <u>exp. 3 is faster</u> than that in <u>exp. 1</u> due to the <u>higher temperature</u> .	\checkmark	
Exp 3 & 4:	The reaction in <u>exp. 4 is faster</u> than that in <u>exp. 1</u> due to the <u>higher temperature / larger surface area</u> . OR Graph <u>A</u> represents <u>exp. 4 due to the greater mass</u> of CaCO ₃ - greater yield of CO ₂ at a faster rate.	~	
	Therefore the <u>gradient</u> of the graphs of <u>exp. 3 & 4 are</u> greater/steeper than that of <u>exp. 1</u> .	\checkmark	
Final answer	С	\checkmark	

1.5 Marking criteria:

• Divide volume by: $25, 7 \checkmark$ • Use ratio: $n(CO_2) = n(CaCO_3) = 1:1 \checkmark$ • Substitute 100 in $n = \frac{m}{M} \cdot \checkmark$ • Subtraction. \checkmark • Final answer: 7,00 g to 7,5 g \checkmark n(CO₂) = $\frac{V}{V_m}$ $= \frac{4,5}{25,7} \checkmark$ = 0,18 mol $n(CaCO_3) = n(CO_2) = 0,18 \text{ mol} \checkmark$ $n(CaCO_3) = \frac{m}{M}$ $0,18 = \frac{m}{100} \therefore m(CaCO_3) = 18 \text{ g}$ $m(CaCO_3) \text{ not reacted: } 25 - 18 \checkmark = 7,00 \text{ g} \checkmark$ (6)

(5) **[18]**

(2)

QUESTION 2

2.1	Exothermic \checkmark <u>Temperature increases during reaction.</u> / T _i < T _f \checkmark	(2)
2.2	Larger surface area in experiment 2. \checkmark	(1)
2.3	More than one independent variable. \checkmark OR Different concentrations and state of division.	(1)
2.4	 Faster than ✓ A catalyst is used in experiment 5. ✓ A catalyst provides an alternative pathway of <u>lower activation energy</u>. ✓ More molecules have sufficient kinetic energy. ✓ More effective collisions per unit time. ✓ 	(5)
2.5	$n(Zn) = \frac{m}{M} \checkmark = \frac{1,2}{65} \checkmark = 0,018 \text{ mol}$ $n(HC\ell)_{\text{reacted}} = 2n(Zn) = 2(0,018) = 0,037 \text{ mol} \checkmark$ Rate = $\frac{\Delta n}{\Delta t} = \frac{0,037}{8} \checkmark = 4,63 \times 10^{-3} \text{ mol} \cdot \text{s}^{-1} \checkmark$	(6) [15]

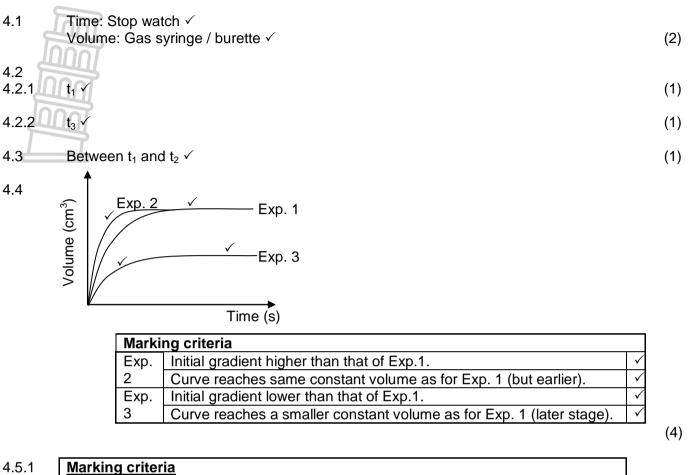
QUESTION 3

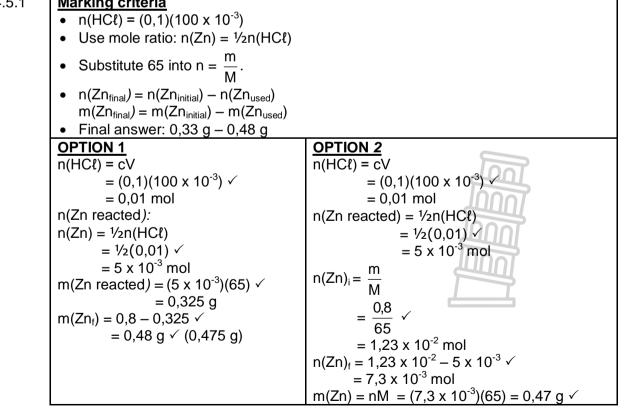
3.1 Exothermic \checkmark $\Delta H < 0 / Energy is released. <math>\checkmark$

3.2 3.2.1	$\frac{\text{OPTION 1}}{n(\text{HC}\ell) = cV} = (1,5) \checkmark (30 \times 10^{-3}) \checkmark = 0,045 \text{ mol}$ Ave rate = $-\frac{\Delta n}{\Delta t}$ $= -\frac{(0-45)}{(60-0)} \checkmark$ = 7,5 x 10 ⁻⁴ mol·s ⁻¹ ✓	$\frac{\text{OPTION 2}}{\text{Ave rate}} = -\frac{\Delta c}{\Delta t}$ $= -\frac{(0-1,5)}{(60-0)} \checkmark$ $= 0,025 \text{ mol·dm}^{-3} \cdot \text{s}^{-1}$ ∴ average rate = (0,025)(30 x 10^{-3}) \checkmark \checkmark $= 7,5 x 10^{-4} \text{ (mol·s}^{-1}) \checkmark$	(5)
3.2.2(a)	 Increases √ The reaction is <u>exothermic</u>, resulting in the reaction is <u>exothermic</u>. 	n an <u>increase in temperature</u> . ✓	

3.Z.Z(a)	increases *	
	 The reaction is <u>exothermic</u>, resulting in an <u>increase in temperature</u>. ✓ 	
	 More molecules have sufficient kinetic energy. ✓ 	
	More effective collisions per unit time. ✓	(4)
	Innat	
3.2.2(b)	Decreases ✓	
	Concentration of acid decreases. ✓	
	OR The surface area of magnesium decreases.	(2)
3.3	ANY TWO	
	 Higher temperature ✓ 	
	• Larger surface area/state of division/contact area of Mg. / Use Mg powder. ✓	
	Addition of a catalyst.	(2)
	·	[15]

QUESTION 4





4.5.2 Smaller than ✓

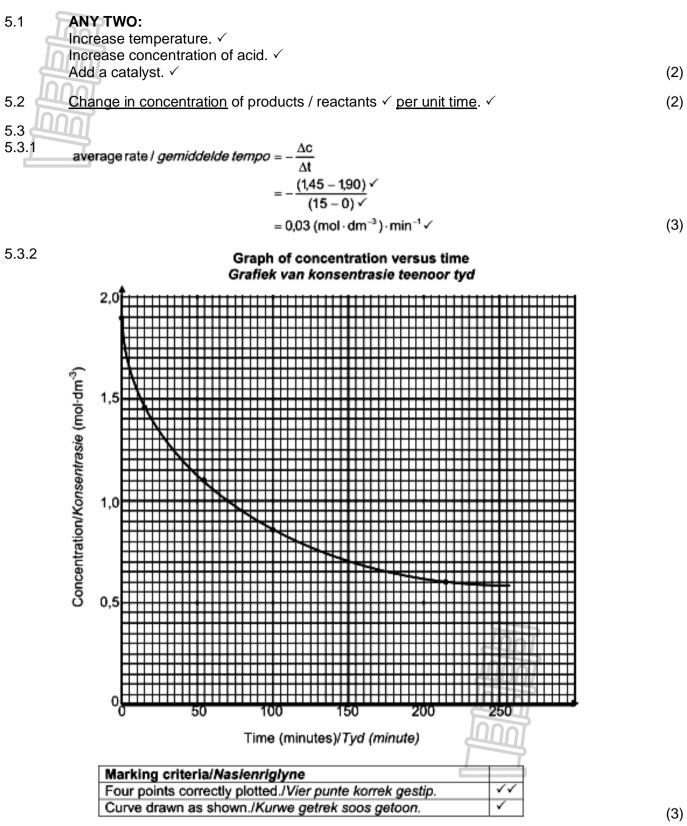
(1) **[15]**

(5)

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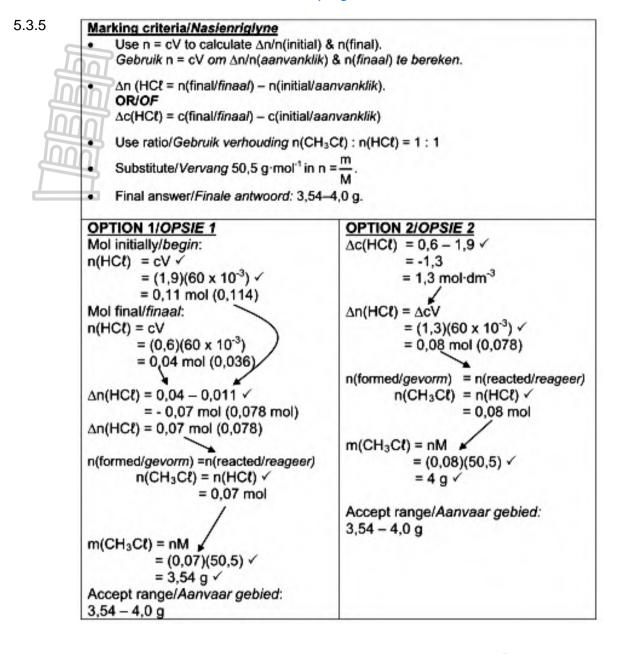
5.3.3 1,15 to 1,25 mol·dm⁻³ \checkmark

- 5.3.4 Concentration of reactants decreases. \checkmark
 - Less particles per unit volume. ✓
 - Less effective collisions per unit time. ✓

(3)

(1)

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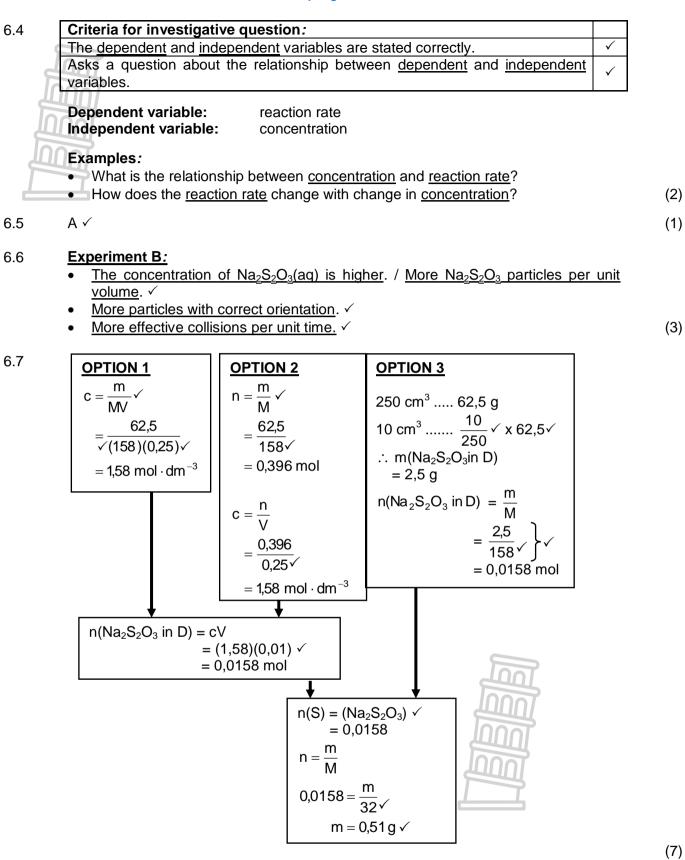
QUESTION 6

- 6.1 ANY TWO: Temperature of reaction mixture ✓ Addition of a catalyst ✓ Concentration of reactants
 6.2 Sulphur / S ✓
- 6.3 Water is used to dilute/change the concentration of the $Na_2S_2O_3(aq)$

(5) **[19]**

(1)

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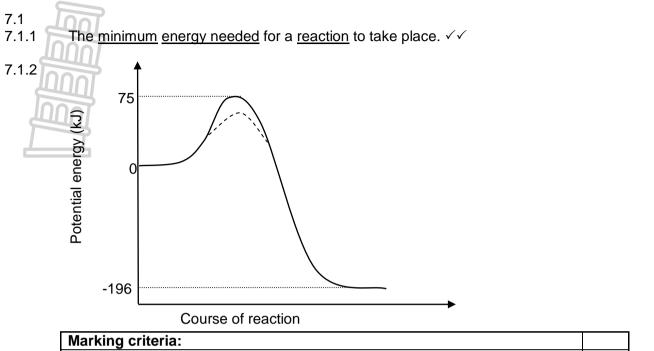


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[17]

(2)

QUESTION 7



Marking criteria:	
Shape of curve for exothermic reaction as shown.	\checkmark
Energy of activated complex shown as 75 kJ in line with the peak.	\checkmark
Energy of products shown as - 196 kJ below the zero.	\checkmark
IF: Wrong shape, e.g. straight line.	$\frac{0}{3}$

(3)

7.1.3 Marking criteria

- Dotted line (---) on graph in QUESTION 7.1.2 showing lower energy for activated complex. ✓
- Dotted curve starts at/above energy of reactants and ends at/above energy of products on the inside of the original curve. ✓
- 7.1.4 A catalyst provides an alternative pathway of <u>lower activation energy</u>. ✓
 - More molecules have sufficient kinetic energy. ✓
 - More effective collisions per unit time. ✓

7.2

7.2.1

Ave rate =
$$\frac{\Delta v}{\Delta t}$$

= $\frac{52 - 16}{40 - 10}$
= 1,2 (dm³·s⁻¹) \checkmark

 ΔV



(3)

(3)

(2)

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7.2.2	Marking criteria:			
5	• $V(O_2) = 60 \text{ dm}^3 \text{ AND divide volume by } 24. \checkmark$			
4	• Use ratio: $n(H_2O_2) = 2n(O_2)$	_,		
	• Use 34 g·mol ⁻¹ in $n = \frac{m}{M}$	or in ratio calculation. ✓		
Ľ.				
In	 Final answer: 170 g ✓ 			
	OPTION 1	OPTION 2	OPTION 3	
	$n(O_2) = \frac{V}{V}$	24 dm ³ : 1 mol	$p(O_{2}) = V$	
	\overline{V}_{M}	60 dm³ : 2,5 mol √	$n(O_2) = \frac{V}{V_M}$	
	60		60	
	$=\frac{60}{24}\checkmark$	$n(H_2O_2) = 2n(O_2)$	$=\frac{60}{24}\checkmark$	
		$= 2(2,5) \checkmark$	- 25 mol	
	= 2,5 mol	= 5 mol	= 2,5 mol	
	$n(H_2O_2) = 2n(O_2)$			
	$= 2(2,5) \checkmark$	34 g √: 1 mol	$n(O_2) = \frac{m}{M}$ $\therefore 2,5 = \frac{m}{32}$	
	= 5 mol	x : 5 mol	M	
	- 5 1161	x = 170 g ✓	$\therefore 2.5 = \frac{m}{m}$	
	m		32	
	$n(H_2O_2) = \frac{m}{M}$		∴ m = 80 g	
			\checkmark	
	$\therefore 5 = \frac{m}{34}$		$2(34) g\sqrt{H_2O_2} \dots 32 g O_2$ x g H ₂ O ₂ \ldots 80 g O ₂	
			x g H ₂ O ₂ 80 g O ₂	
	∴ m = 170 g ✓			
			m(H ₂ O ₂) = 170 g √	

- Equal to \checkmark 7.2.3
- 7.3
- 7.3.1 Q√
- 7.3.2 P√

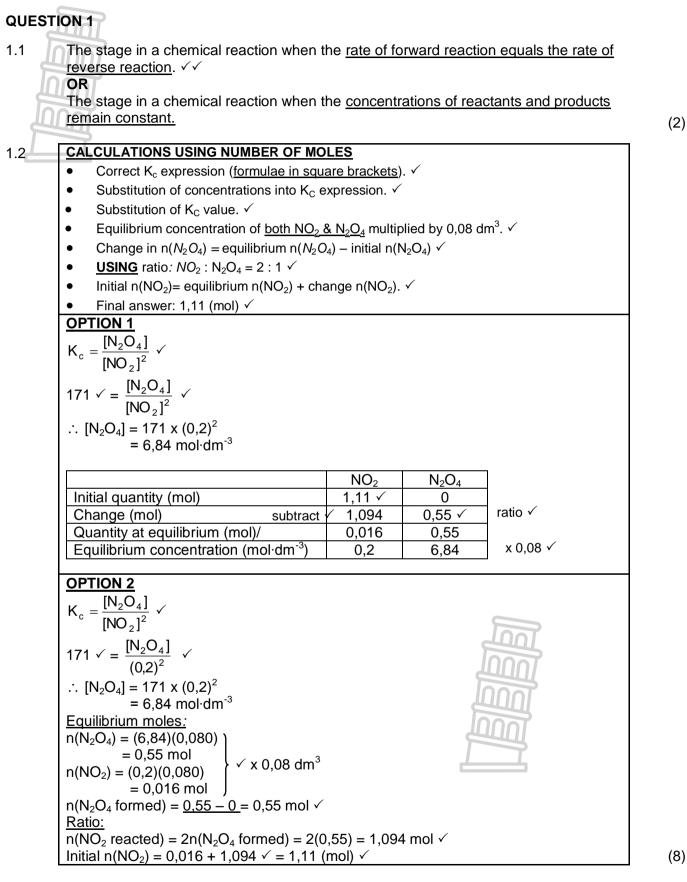


(1)

(4)

- (1)
- (1) **[20]**

CHEMICAL EQUILIBRIUM



1.3

1.3.1 Concentration (of the gases) increases. / Molecules become more condensed or move closer to each other. ✓

Terms, definitions, questions and answers

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1.3.2	 Increase in pressure favours the reaction that leads to smaller number of moles of gas. ✓ Forward reaction is favoured. ✓ Number of moles/amount of N₂O₄ / colourless gas increases. ✓ OR Number of moles/amount of NO₂ / brown gas decreases. ✓ 					
1.4		of NO_2 / I	brown gas	decreases	5. v	(3)
1.4.1	Darker √					(1)
1.4.2	Decreases ✓					(1) [16]
QUEST	ION 2					[]
2.1	A reaction is reversible when produ	ucts can b	e converte	d back to	reactants. √	(1)
2.2	No change ✓					(1)
2.3						<i></i>
2.3.1	Temperature decreases ✓	a		, ,		(1)
2.3.2	 Decrease in temperature decrea reactions. ✓ 	ises the ra	ate of both	forward ar	nd reverse	
	 Decrease in temperature favours The rate of the reverse (exothern favoured. √ 				verse reaction is	(3)
	OPTION 1/OPSIE 1At equilibrium/by ewewig: $[H_2] = [I_2] \checkmark$ $K_c = \frac{[H_2][I_2]}{[HI]^2} \checkmark$ Divide by 2 dm ³ \checkmark $\therefore 0,02 \checkmark = \frac{(x)(x)}{\left(\frac{0,04}{2}\right)^2} \checkmark$ Divide by 2 dm ³ \checkmarkDivide by 2 dm ³ ✓Divide by 2 dm ³ ✓					
		HI	H ₂	I2		
	Initial quantity (mol) Aanvangshoeveelheid (mol)	x	0	0		
	Change (mol) Verandering (mol)	x - 0,04	$\frac{x-0,04}{2}$	$\frac{x-0,04}{2}$	ratio ✓ verhouding	
	Quantity at equilibrium (mol)/ Hoeveelheid by ewewig (mol)	0,04	$\frac{x-0,04}{2}$	$\frac{x-0,04}{2}$	0001	
	Equilibrium concentration (mol·dm ⁻³)	0,02	<u>x - 0,04</u>	<u>x - 0,04</u>	Divide by 2 dm ³ \checkmark Deel deur 2 dm ³	
	Ewewigskonsentrasie (mol·dm ⁻³)	0,02	4	4		
	$K_{c} = \frac{[H_{2}][I_{2}]}{[HI]^{2}} \checkmark$ ∴ 0,02 \sigma = $\frac{(\frac{x - 0,04}{4})(\frac{x - 0,04}{4})}{(0,02)^{2}} \checkmark$		ession, corre , <i>korrekte sub</i>			
	$\therefore 0,02 \checkmark = \underbrace{-4 4}_{(0,02)^2} \checkmark$ $\therefore x = 0,05$	Wrong K _c (Max./Maks	expression/Ve · 2/6	erkeerde K _c -i	uitdrukking:	
	$[H_2] = \frac{x - 0.04}{2}$ $= \frac{0.05 - 0.04}{2}$					
	2 = 2,83 x 10 ⁻³ mol⋅dm ⁻³ ✓				(6)	(6)

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 $K_{c} = \frac{1}{0.02} = 50 \checkmark$

2.5

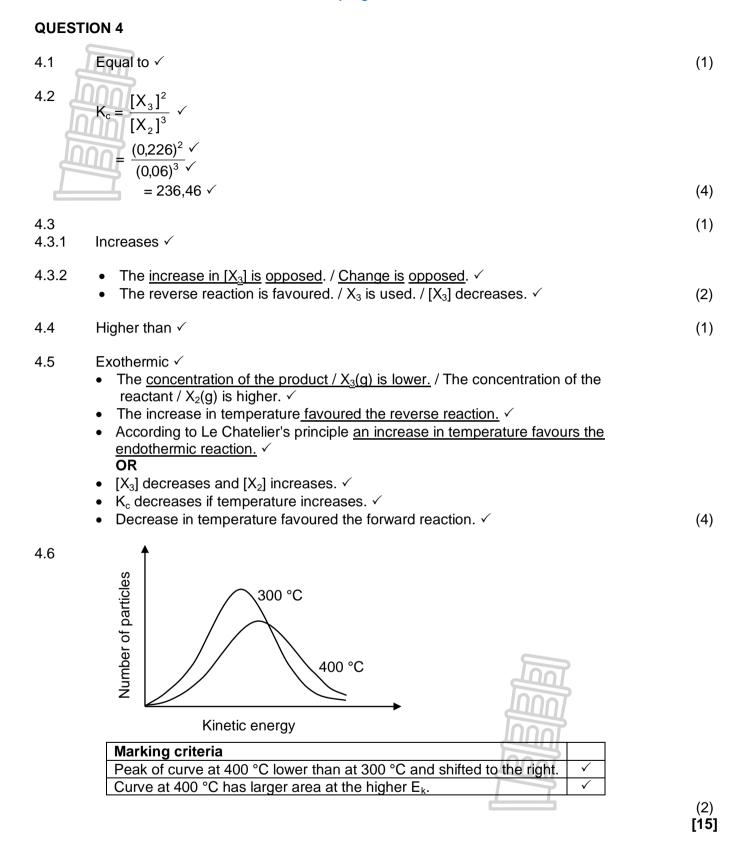
10N 3				
When the <u>equilibrium</u> (in a closed system <u>new equilibrium</u> ✓ by favouring the reaction				
Remains the same \checkmark				
Increases 🗸				
$\begin{array}{l} \hline \textbf{OPTION 1} \\ K_c = [NH_3][H_2S] \checkmark \\ \therefore 1,2 \times 10^{-4} \checkmark = [NH_3][H_2S] \\ \therefore [NH_3] = [H_2S] \\ = 0,011 \text{ mol} \cdot dm^{-3} \\ n(NH_3) = cV \\ = (0,011)(5) \checkmark \\ = 0,06 \text{ mol} (0,06 \text{ mol}) \\ n(NH_4HS) = n(NH_3) = 0,06 \text{ mol} \checkmark \\ m(NH_4HS) = nM \\ = (0,06)(51) \checkmark \\ = 2,81 \text{ g} \checkmark \end{array}$	 K_c expl Substit x volun <u>Use</u> mo Substit 	 x volume (5 dm³) <u>Use</u> mole ratio: 1:1 		
OPTION 2	NH₄HS	NH ₃	ЦС	П
Initial quantity (mol)		0	H_2S	-
Change (mol)	x	x	x	-
Quantity at equilibrium (mol)	-	X	X	-
Equilibrium concentration (mol·dm ⁻³)	-	x 5	x 5	Divide by 5 ✓
K _c = [NH ₃][H ₂ S] ✓ ∴ 1,2 x 10 ⁻⁴ ✓ = $(\frac{x}{5})(\frac{x}{5})$ ∴ x = 0,0547 mol m(NH ₄ HS) = nM		Ĩ		

- Increase in pressure favours the reaction that leads to the smaller number of • moles of gas. VV
- The reverse reaction is favoured. \checkmark .

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(3) [13]

FS/2017



(1)

(1)

QUESTION 5

5.1 OPTION 1/OPSIE 1	OPTION 2/OPSIE 2	7
$G = \frac{m}{MV}$	$n = \frac{m}{M}$	
	$=\frac{2.2}{44}$ Both formulae/ albei formules	
0,01 mol · dm ⁻³ ✓	= 0,05 mol	
	$c = \frac{n}{V}$	
	$=\frac{0.05}{5}\checkmark$	
	= 0,01 mol · dm ⁻³ ✓	(4)

5.2 For equilibrium, a forward and a reverse reaction are needed. ✓
 OR Without CaO(s), the reverse reaction is not possible.
 OR If only CO₂ is present, the reverse reaction cannot take place.

5.3
$$CO_2$$
 is a gas and will escape if the container is not sealed. \checkmark

5.4 **OPTION 1** (POSITIVE MARKING FROM Q6.1) K_c = [CO₂] √ = 0,0108 ∴ [CO2] = 0,0108 (mol·dm⁻³) √ n(CO2 at equilibrium/by ewewig) = cV = (0,0108)(5) < = 0,054 mol n(CO₂ formed/gevorm) n(CO₂ equilibrium/by at ewewig) n(CO2initially/begin) = 0,054 - 0,05 ✓ = 0,004 mol n(CaCO3) = n(CO2 formed) = 0,004 mol </ m(CaCO₃) = nM = (0,004)(100) ✓ = 0,4 g √

5	$\frac{\text{OPTION 2}}{\text{K}_{6} = [CO_{2}]} \checkmark$ $= 0.0108 \checkmark$ (POSITIVE M	ARRING		Q0.1)	
	.:.[CO ₂] = 0,0108 (mol·dm ⁻³)				-
ĥ	Initial quantity (mol) Aanvangshoeveelheid (mol)	CaCO ₃	CaO 0	CO ₂ 0,05	-
	Change (mol) Verandering (mol)	0,004	x	0,004 🗸	√Ratio/ Verhouding
	Quantity at equilibrium (mol) Hoeveelheid by ewewig (mol)			0,054 🗸	
	Equilibrium concentration (mol·dm ⁻³) Ewewigskonsentrasie (mol·dm ⁻³)			0,0108	
5 5.1	= (0,004)(100) ✓ = 0,4 g ✓ Remains the same ✓				
5.2	Decreases √				
	Endothermic 🗸				
	 K_c decreases with decrease in tem Therefore the product of the con reaction is favoured. ✓ A decrease in temperature favours OR K_c increases with increase in temp Increase in temperature favours th 	the exotheration	of pro iermic i	reaction. v	
	 Increase in temperature favours th 				



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(1)

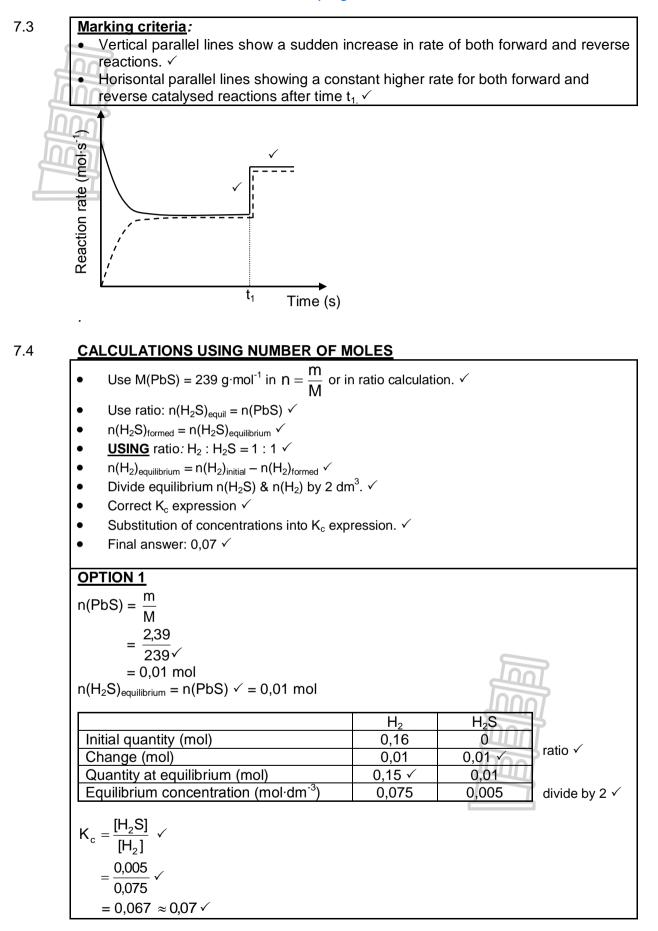
(2)

(2)

QUEST	ION 6			
6.1	Reversible reaction ✓			
6.2	Endothermic \checkmark ΔH is positive. / $\Delta H > 0 \checkmark$			
6.3	Larger than \checkmark K _c > 1 \checkmark			
6.4	OPTION 1			
	$n = \frac{m}{M}$			
	$=\frac{168}{28}$			
	= 6 mol			
	Initial quantity (mal)		CO 0	
	Initial quantity (mol) Change (mol)	<u>х</u> З	0 6 √	ratio ✓
	Quantity at equilibrium (mol)	$x - 3 \checkmark$	6	-
	Equilibrium concentration (mol·dm ⁻³)	$\frac{x-3}{2}$	3	Divide by 2 ✓
	$K_{c} = \frac{[CO]^{2}}{[CO_{2}]} \checkmark$ $14 \checkmark = \frac{(3)^{2}}{\frac{x-3}{2}} \checkmark$ ∴ x = 4,29 mol ✓			
	OPTION 2 $n = \frac{m}{M}$ $c = \frac{n}{V}$ $= \frac{168}{28}$ $= \frac{6}{2}$ Divide by 2 \checkmark			7
	$= 6 \text{ mol}$ $= 3 \text{ mol} \cdot \text{dm}^{-3}$			7
	Initial concentration (mol·dm ⁻³)	CO ₂ x	CO 0	
	Change (mol·dm ⁻³)	1,5	3✓	ratio ✓
	Equilibrium concentration (mol·dm ⁻³)	x – 1,5 √	3	
	$K_{c} = \frac{[CO]^{2}}{[CO_{2}]} \checkmark$			
	14 √ = $\frac{[3]^2}{x - 1.5}$ √ ∴ x = 2.14 mol·dm ⁻³			
	$n(CO_2) = cV = (2,14)(2) = 4,29 \text{ mol } \sqrt{2}$			

	OPTION 3				
۶	$n = \frac{m}{M}$				
	$1 = \frac{168}{28}$				
Г	= 6 mol				
6				- I	
- H	Initial quantity (mol)	CO ₂ 4,28√	<u> </u>	-	
	Change (mol)	3	6	ratio √	
	Quantity at equilibrium (mol)	1,28 ✓	6√		
	Equilibrium concentration (mol·dm ⁻³)	0,64	3	x 2 √	
	$K_{c} = \frac{[CO]^2}{[CO_2]} \checkmark$				
	$14 \checkmark = \frac{[3]^2}{[CO_2]} \checkmark$				
	[CO₂] ∴ [CO₂] = 0,64 mol·dm ⁻³				(0)
	$1.1 [CO_2] = 0.04 \text{ morely}$				(9)
6.5 6.5.1	Remains the same \checkmark				(1)
6.5.2	Decreases √				(1)
6.5.3	Increases v				(1)
					[17]
QUEST	ION 7				
7.1	The stage in a chemical reaction when the	rate of forward	reaction eq	uals the rate of	
	reverse reaction. √√ OR				
	The stage in a chemical reaction when the	ne <u>concentratior</u>	ns / quantiti	es of reactants	
	and products remain constant.				(2)
7.2					
7.2.1	Remains the same \checkmark		To	3	(1)
7.2.2	Decreases ✓			3	
	• When the temperature is increased th		will oppose	this increase /	
	decrease the temperature will be favou OR	ired. ✓			
	The forward reaction is exothermic.			7	
	• An increase in temperature favours the	e endothermic re	eaction. 🗸	5	
	• The reverse reaction is favoured. \checkmark			3	(4)

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OPTION 2	
$n(PbS) = \frac{m}{M}$ $= \frac{2,39}{239}\checkmark$ $= 0,01 \text{ mol}$ $n(H_2S)_{\text{reacted}} = n(PbS) \checkmark = 0,01 \text{ mol}$ $= n(H_2S)_{\text{equilibrium}}$ $n(H_2S)_{\text{formed}} = n(H_2S)_{\text{equilibrium}} - n(H_2)$ $= 0,01 - 0 \checkmark$ $= 0,01 \text{ mol}$	
$n(H_2)_{\text{reacted}} = n(H_2S)_{\text{formed}} \checkmark = 0,01 \text{ r}$ $n(H_2)_{\text{equilibrium}} = n(H_2)_{\text{initial}} - n(H_2)_{\text{reacte}}$ $= 0,16 - 0,01 \checkmark$ $= 0,15 \text{ mol}$	
$c(H_{2}) = \frac{n}{V}$ = $\frac{0.15}{2}$ = 0.075 mol·dm ⁻³ $K_{c} = \frac{[H_{2}S]}{[H_{2}]}$ = $\frac{0.005}{0.075}$ \checkmark = 0.067 ≈ 0.07 \checkmark	$c(H_2S) = \frac{n}{V}$ = $\frac{0.01}{2}$ = 0.005 mol·dm ⁻³



(9) **[18]**

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ACIDS AND BASES

QUEST	ION 1		
1.1	000		
1.1.1	lonises / dissociates completely in wate	r. ✓	(1)
1.1.2	NO $_{3}^{-}$ / Nitrate ion \checkmark		(1)
1.1.3	$pH = -\log[H_3O^+] / -\log[H^+] \checkmark = -\log(0,3)$	✓ = 0,52 ✓	(3)
1.2			
1.2.1	$c = \frac{n}{V} \checkmark$		
	$2 = \frac{n}{0.1} \checkmark$		
	0,1 ∴ n(HCℓ) = 0,2 mol √		(3)
1.2.2	Burette 🗸		(1)
1.2.3	В√		
	Titration of strong acid and strong base	_ < <	(3)
1.2.4	The <u>number of moles</u> of acid in the flash	< <u>remains constant</u> . ✓	(1)
1.2.5	$c = \frac{n}{V} \checkmark \therefore 0, 2 = \frac{n}{0.021} \checkmark \therefore n = 4, 2$	x 10 ⁻³ mol √	
	$n(HC\ell)_{excess} = n(NaOH) = 4,2 \times 10^{-3} \text{ mol}$		(3)
1.2.6	POSITIVE MARKING FROM QUESTIO	N 1.2.1 AND 1.2.5.	
	 Marking criteria: n(HCl reacted) = initial (from Q1.2.1 	$-$ excess (from 01.2.5) \checkmark	
	 Use mol ratio of acid: base = 2 : 1. 		
	• Substitute 40 into n = $\frac{m}{M}$ \checkmark		
	• $\frac{m(MgO)_{reacted}}{4,5} \times 100 \checkmark$		
	 Final answer: 87,11 % ✓ 	<u> </u>	
	<u>OPTION 1</u> n(HCℓ reacted):	n(HCl reacted):	
	$0.2 - 4.2 \times 10^{-3} \checkmark = 0.196 \text{ mol}$	$0.2 - 4.2 \times 10^{-3} \checkmark = 0.196 \text{ mol}$	
	n(MgO reacted):	n(HCl reacted) = $\frac{m}{M}$	
	$\frac{1}{2}n(HC\ell) = \frac{1}{2}(0,196)$ = 9,8 x 10 ⁻² mol \checkmark	$0,196 = \frac{m}{36.5}$	
	n(MgO reacted) = $\frac{m}{M}$	∴ m(HCl reacted) = 7,154 g	
	$\therefore 0,098 = \frac{m}{40}$	40 g MgO ✓	
	m = 3.02 m	∴ x = 3,92 g	
	% purity = $\frac{3,92}{4,5 \times 100}$ \checkmark	% purity = $\frac{3,92}{4,5 \times 100}$ \checkmark	
		= 87,11% ✓	
	= 87,11% ✓		(5) [21]

(5) **[21]**

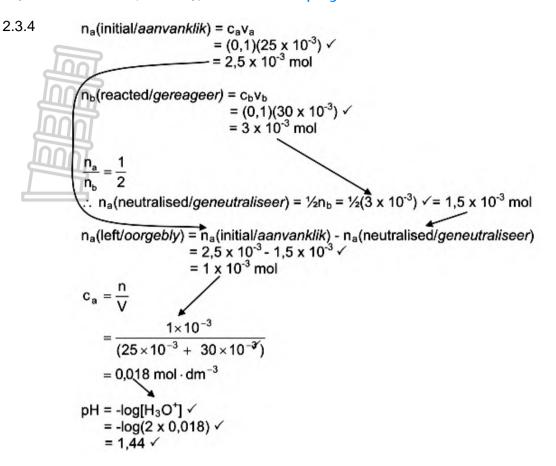
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QUESTION 2

2.1 2.1.1	An acid is a proton (H⁺ ion) donor. ✓	(1)
2.1.2	It ionises to form 2 protons for each H_2SO_4 molecule. / It ionises to form 2 moles of H^+ ions. \checkmark OR It donates 2 H^+ ions per H_2SO_4 molecule.	(1)
2.2 2.2.1	Amphiprotic substance / Ampholyte ✓	(1)
2.2.2	H₂CO₃ ✓	(1)
2.3 2.3.1	n(NaHCO ₃) = $\frac{m}{M} \checkmark$ = $\frac{27}{84} \checkmark$ = 0,32 mol (0,0321485 mol)	
	$n(H_2SO_4) = \frac{1}{2}n(NaHCO_3) = \frac{1}{2}(0,32) \checkmark = 0,16 \text{ mol}$ (0,01607142 mol)	
	$c = \frac{n}{V}$	
	$6 = \frac{0.16}{V}$	
	\therefore V = 0,03 dm ³ \checkmark (30 cm ³ /0,027 dm ³ /27 cm ³)	(6)
2.3.2	n _a (initial <i>/aanvanklik</i>) = n _a (final <i>/finaal)</i> c _a v _a (initial <i>/aanvanklik</i>) = c _a v _a (final <i>/finaal</i>) ∴ (6)v _a = (0,1)(1) ✓	
	$\therefore v_a = 0.02 \text{ dm}^3 \checkmark (20 \text{ cm}^3/0.0167 \text{ dm}^3/16.7 \text{ cm}^3)$	(2)
2.3.3	Shows end point of titration. / Shows when neutralisation occurs. \checkmark	(1)



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QUESTION 3

3.1 3.1.1	Diprotic 🗸	(1)
3.1.2	$H_2O \checkmark$ (COO) $_2^{2-} \checkmark$	(2)

3.1.3
$$HC_2O_4^-/H(COO)_2^- \checkmark$$

It acts as base (in reaction I) and as acid (in reaction II). \checkmark

3.2 Ionises / dissociates incompletely / partially. ✓

3.3	OPTION 1	OPTION 2	Marking guidelines:	
	$c = \frac{m}{MV} \checkmark$	$c = \frac{n}{M} \checkmark$	Any formula of:	
	MV	v	$c = \frac{m}{MV} / c = \frac{n}{V} / n = \frac{m}{M}$	
	$\therefore 0,2 = \frac{m}{0,25\checkmark \times 90}\checkmark$ $\therefore m = 4,5 g\checkmark$	$0,2 = \frac{n}{0.05} \sqrt{10}$	MV M	
	,25√×90√	0,25	 Substitution of V as 0,25 dm³. 	
	∴ m = 4,5 g √	m		
		$n = \frac{m}{M}$	 Substitution of 90 g⋅mol⁻¹. 	
			- Final answer: 4.5 g	
		$0,05 = \frac{m}{90}\checkmark$ $m = 4,5 g \checkmark$	• Final answer: 4,5 g	
		m = 4,5 g √		(4)

(8) [22]

(2)

(2)

(1)

3.4 3.4.1	OPTION 1	Marking guidelines:	
	$\frac{c_a V_a}{c_b V_b} = \frac{n_a}{n_b} \checkmark$ $\frac{0.2 \times 25}{c_b \times 36} \checkmark \frac{1}{2} \checkmark$ $\therefore c_b = 0.28 \text{ mol} \cdot \text{dm}^{-3} \checkmark$	 Formula Substitution of 0,2 x 25. Use V_b = 36 cm³. Use mol ratio 1:2. Final answer: 0,28 mol·dm⁻³ 	
	$\frac{\text{OPTION 2}}{n((COOH)_2) = cV \checkmark} = (0,2)(0,025) \checkmark = 0,005 \text{ mol}$ $n(NaOH) = \underline{2}(0,005) \checkmark = 0,01 \text{ mol}$ $c = \frac{n}{V} = \frac{0,01}{0036} \checkmark = 0,28 \text{ mol} \cdot \text{dm}^{-3} \checkmark$	Marking guidelines:• Any ONE of formulae.• Substitution of $0,2 \ge 0,025$.• Use mol ratio 1:2.• Use $V_b = 0,036 \text{ dm}^3$.• Final answer: $0,28 \text{ mol} \cdot \text{dm}^{-3}$	(5)
3.4.2	$(\text{COO})_2^{2-}(\text{aq}) + 2\text{H}_2\text{O}(\ell) \checkmark \rightleftharpoons (\text{COOH})_2(\text{aq})$) + 2OH⁻(aq) ✓ Bal. ✓	(3)
QUEST	ION 4		[18]
4.1 4.1.1	Hydrolysis 🗸		(1)

 4.1.2 Acidic ✓ Forms H₃O⁺ ions during hydrolysis. ✓ OR Salt of strong acid and weak base.

4.2

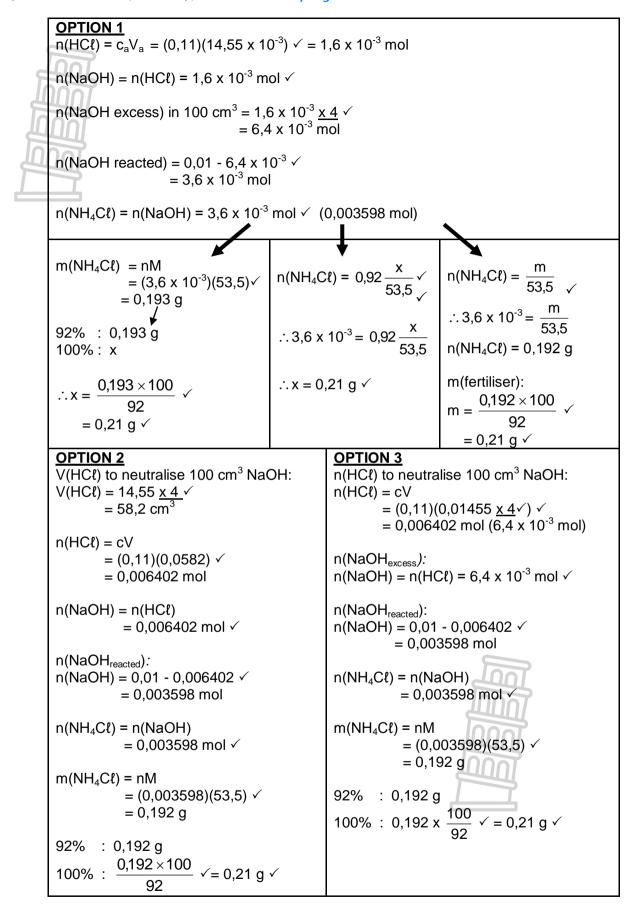
4.2.1 $n = cV \checkmark$ = (0,1)(0,1) \checkmark = 0,01 mol \checkmark

larking criteria	
Substitute volume and concentration to calculate n(HCl) \checkmark	
Use mole ratio: n(NaOH) = n(HCℓ) = 1:1 ✓	Innat
n(NaOH) x 4 OR V(HCℓ) x 4 OR n(HCℓ) x 4 ✓	
Subtraction: n(NaOH _{initial}) - n(NaOH _{excess}) ✓	
Use mole ratio: $n(NaOH) = n(NH_4C\ell) = 1:1 \checkmark$	
Substitute 53,5 g·mol ⁻¹ in n = $\frac{M}{M}$.	
Percentage calculation ✓	
Final answer: 0,11 g - 0,21 g ✓	

(2)

(3)

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4.3 OPTION 1	OPTION 2	
[OH ⁻] = [NaOH] = 0,5 mol·dm ⁻³	$\overline{\text{pOH}} = -\log[\text{OH}^{-}] \checkmark$	
	= -log(0,5) ✓	
$K_{w} = [H_{3}O^{+}][OH^{-}]$	= 0,301	
$K_w = [H_3O^+][OH^-]$ 1 x 10 ⁻¹⁴ = [H_3O^+]0,5		
$[H_3O^+] = 2 \times 10^{-14} \text{ mol} \cdot \text{dm}^{-3}$	pH + pOH = 14	
24101	pH = 14 − 0,301 ✓	
pH = -log[H ⁺] ✓	= 13,7 < (13,699)	
$pH = -log[H^+] \checkmark$ = $-log(2 \times 10^{-14}) \checkmark$		
= 13,7 ✓		(4)
		[18

QUESTION 5

- 5.1 It is a proton donor. $\checkmark\checkmark$ (2)
- 5.2
- 5.2.1 CO₃^{2−} (aq) √

5.2.2
$$H_2CO_3 + H_2O \checkmark \rightleftharpoons HCO_3^-(aq) + H_3O^+(aq) \checkmark$$
 Bal. \checkmark

5.2.3	OPTION/OPSIE 1	OPTION/OPSIE 2	7
0.2.0	pH =-log[H [*]] ✓	pH + pOH = 14 ✓	
	3,4 =-log[H ⁺] ✓	3,4 + pOH = 14 √	
	[H ⁺] = 10 ^{-3,4} / 3,98 x 10 ⁻⁴ mol·dm ⁻³	pOH = 11,6	
	[H ⁺][OH ⁻] = 10 ⁻¹⁴ ✓	pOH =-log[OH'] ✓	
	1×10 ⁻¹⁴	11,6 =-log[OH] ✓	
	$\therefore [OH] = \frac{1 \times 10^{-14}}{3,98 \times 10^{-4}} \checkmark$	[OH ⁻] = 10 ^{-11,6/} 2,51x10 ⁻¹¹ mol·dm ⁻³ ✓	
	= 2.51x10 ⁻¹¹ mol·dm ⁻³		(5)

5.3

5.3.1 An acid that donates ONE proton / H^+ ion / H_3O^+ ion. \checkmark

OR

An acid of which ONE mol ionises to form ONE mol of protons / H^+ ions / H_3O^+ ions. (1)

5.3.2	$\frac{\text{OPTION/OPSIE 1}}{\begin{array}{c} c_a \times V_a \\ \hline c_b \times V_b \end{array}} = \frac{n_a}{n_b} \checkmark$ $\underbrace{\begin{array}{c} c_a \times 25 \\ \hline 0.1 \times 27.5 \end{array}}_{c_a} = \frac{1}{1} \checkmark$ $c_a = 0.11 \text{ mol} \cdot \text{dm}^{-3} \checkmark$	$\frac{OPTION/OPSIE 2}{n(NaOH) = cV \checkmark}$ = 0,1 x 0,0275 \leftarrow = 0,00275 mol n(acid X) = n(NaOH) = 0,00275 mol \leftarrow = 0,00275 mol \leftarrow	(5)
-------	---	---	-----

5.3.3 Weak ✓

The [H⁺] OR [H₃O⁺] is lower than the concentration of acid X. \checkmark Therefore the acid is incompletely ionised. \checkmark

(1)

(3)

FS/2017

(2)

(3)

QUESTION 6

QUESTION	
6.1 6.1.1 An acid is a proton/ H ⁺ donor. $\sqrt{4}$ NOTE: NOT H ₃ O ⁺	(2)
	()
6.1.2 $H_2O \checkmark$ $H_2CO_3 \checkmark$	(2)
6.1.3 $H_2O \checkmark OR HCO_3^-$	(1)
6.2	
6.2.1 $n(HC\ell) = cV \checkmark$ = (0,1)(0,5) \sqcap = 0,05 mol	
n(NaHCO ₃) = cV = (0,25) (0,8) ✓	
= 0,2 mol	
$n(NaHCO_3)_{reacted} = n(HC\ell)$ = 0,05 mol \checkmark	
$n(NaHCO_3)_{excess} = \frac{0.2 - 0.05}{0.2 - 0.05} \checkmark$ = 0.15 mol	
$n(OH) = n(NaHCO_3) \checkmark$	
= 0,15 mol	
$c(OH^-) = \frac{n}{V}$	
$=\frac{0,15}{1,3}\checkmark$	
$=$ 0,12 mol \cdot dm ⁻³ \checkmark	(8)
6.2.2 POSITIVE MARKING FROM QUESTION 6.2.1	
$K_w = [H_3O^+][OH^-] = 1 \times 10^{-14}$ $pOH = -log[OH^-] \checkmark$ $1 \times 10^{-14} = [H_3O^+](0,12) \checkmark$ $= -log(0,12) \checkmark$	

	$1 \times 10^{-14} = [H_3O^+](0,12) \checkmark$ $[H_3O^+] = 8,33 \times 10^{-14} \text{ mol} \cdot \text{dm}^{-3}$	$= -\log(0, 12) \checkmark$ = 0,92	
	$pH = -\log [H_3O^+] \checkmark$	pH + pOH = 14	
	$= -\log(8,33 \times 10^{-14})$	pH + 0,92 = 14 ✓	
	= 13,08 ✓	pH = 13,08 ✓	(4)
			[17]
QUEST	ION 7		
7.1			

7.1.1 Hydrolysis is the reaction of a salt with water. $\checkmark\checkmark$

7.1.2 Smaller than (7) \checkmark

• $NH_4^+ + H_2O \checkmark \rightarrow NH_3 + H_3O^+ \checkmark$

7.2		
	g guidelines:	
	stitution of 98 g·mol⁻¹. ✓	
	al answer: 0,08 mol ✓	
OPTIO	<u>N 1</u>	OPTION 2
m		98 g √: 1 mol
		7,35 :0,08 mol ✓
7,3	35	OPTION 3
9	8	$c = \frac{m}{m}$
= 0,0	8 mol ✓ (0,075 mol)	$C = \frac{1}{MV}$
		7,35
		$=\frac{1}{98 \times 0.5}$
		= 0,15 mol·dm ⁻³
		n = cV
		$= 0,15 \times 0,5$
		= 0,08 mol √

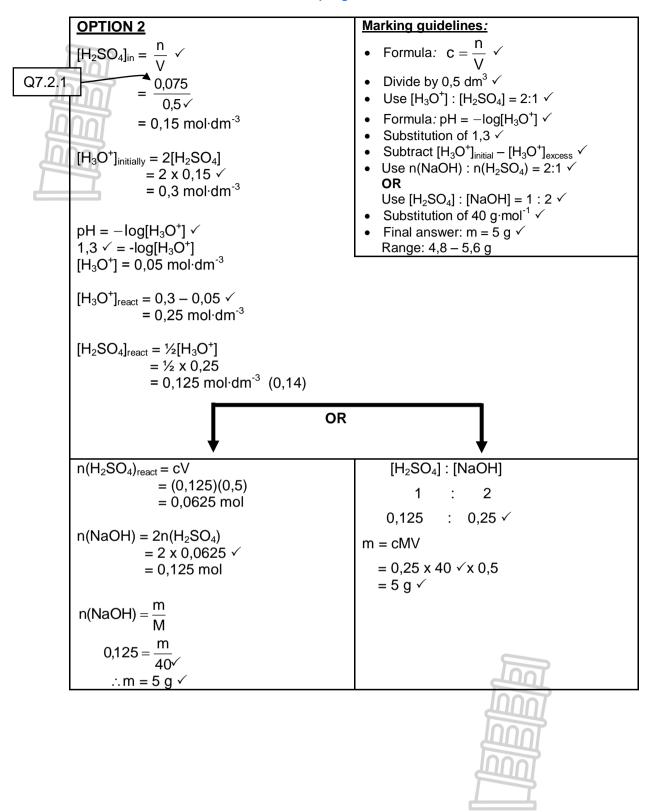
7.2.2 **POSITIVE MARKING FROM QUESTION 7.2.1.** Marking guidelines: **OPTION 1** • Formula: $pH = -log[H_3O^+] \checkmark$ $pH = -log[H_3O^+] \checkmark$ Substitution of 1,3 ✓ • $1,3 \checkmark = -\log[H_3O^+]$ Use $[H_2SO_4]$: $[H_3O^+] = 1 : 2 \checkmark$ • $[H_3O^+] = 0,05 \text{ mol} \cdot \text{dm}^{-3}$ Formula: $c = \frac{n}{V}$ ٠ $[H_2SO_4] = \frac{1}{2}[H_3O^+]$ Multiply by 0,5 dm³ \checkmark $= \frac{1}{2} \times 0.05 \checkmark$ ٠ Subtract $n_{initial} - n_{excess} \checkmark$ • $= 0.025 \text{ mol} \cdot \text{dm}^{-3}$ • Use n(NaOH) : n(H₂SO₄) = 2:1 √ Substitution of 40 g·mol⁻¹ \checkmark • $n(H_2SO_4)_{excess} = cV \checkmark$ Final answer: $m = 5 g \checkmark$ • = 0,025 x 0,5 √ Range: 4,8 – 5,6 g = 0,0125 mol Q7.2.1 $n(H_2SO_4)_{react} = 0,075 - 0,0125 \checkmark$ = 0,0625 mol $n(NaOH) = 2n(H_2SO_4)$ = 2 x 0,0625 √ = 0,125 mol n(NaOH) = 1 mol : 40 g √ M 0,125 mol : 5 g m ∴ 0,125 = OR 40√ m = 5 g √

(2)

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(9)

[16]



QUEST	ION 1	
1.1	 Pressure: <u>1 atmosphere (atm)</u> / 101,3 kPa / 1,013 x 10⁵ Pa ✓ Temperature: <u>25 °C</u> / 298 K ✓ 	(2)
1.2	Platinum is inert / does not react with the H ⁺ ions OR acid. \checkmark Platinum is a conductor (of electricity). \checkmark	(2)
1.3 1.3.1	Salt bridge ✓	(1)
1.3.2	-0,31 V 🗸	(1)
1.3.3	$2H^+ + 2e^- \rightarrow H_2 \checkmark \checkmark$	(2)
1.4 1.4.1	POSITIVE MARKING FROM QUESTION 1.3.2. $E_{cell}^{\theta} = E_{reduction}^{\theta} - E_{oxidation}^{\theta} \checkmark$ 2,05 \checkmark = -0,31 \checkmark - $E_{M/M^{2+}}^{\theta}$ $E_{M/M^{2+}}^{\theta}$ = -2,36 (V) \checkmark	
	M is magnesium/ Mg. ✓	(5)
1.4.2	Exothermic 🗸	(1)
1.5	The cell reaction reaches equilibrium. \checkmark	(1) [15]
QUEST	ION 2	
2.1	 A substance that is being reduced. √√ OR A substance that gains/accepts electrons. OR A substance whose oxidation number decreases. 	(2)
2.2	Ag ⁺ is a stronger oxidising agent \checkmark than Cu ²⁺ \checkmark and will oxidise Cu \checkmark to (blue) Cu ²⁺ ions. \checkmark OR Cu ²⁺ is a weaker oxidising agent \checkmark than Ag ⁺ \checkmark and Cu will be oxidised \checkmark to Cu ²⁺ ions. \checkmark	(4)
2.3	Chemical energy to electrical energy ✓	(1)
2.4	A✓	(1)
2.5	$E_{cell}^{\theta} = E_{reduction}^{\theta} - E_{oxidation}^{\theta} \checkmark$ = +0,80 \sqcap - 0,34 \sqcap = +0,46 V \sqcap	(4)
2.6	Cu + 2Ag ⁺ (aq) $\checkmark \rightarrow$ Cu ²⁺ (aq) + 2Ag(s) \checkmark Balancing \checkmark	(3)
2.7	Remains the same \checkmark	(1) [16]

GALVANIC CELLS

Terms, definitions, questions and answers

FS/2017

QUESTION 3

3.1	Redox reaction ✓	(1)
3.2	P Negative electrode. / Mg is a stronger reducing agent/is oxidized/release electrons.	(2)
3.3 3.3.1	(Temperature): <u>25 °C/298 K</u> ✓ (Concentration): <u>1 mol·dm⁻³</u> ✓	(2)
3.3.2	Mg(s) Mg ²⁺ (aq) √ Pb ²⁺ (aq) Pb(s) √ OR Mg Mg ²⁺ Pb ²⁺ Pb	(3)
3.3.3	$Pb^{2+}/Pb(NO_3)_2/Iead(II)$ ions \checkmark	(1)
3.4	$E_{cell}^{\theta} = E_{reduction}^{\theta} - E_{oxidation}^{\theta} \checkmark$ = -0,13 \sqrt{-(-2,36)} = 2,23 V	(4)
3.5 3.5.1	Remains the same \checkmark	(1)
3.5.2	Increases ✓	(1) [15]
3.5.2 QUEST		
QUEST	ION 4	
QUEST	Temperature: <u>25 °C /</u> 298 K ✓	
QUEST	TON 4 Temperature: 25 °C / 298 K ✓ Pressure: 101,3 kPa / 1,013 x 10 ⁵ Pa / 1 atm / 100 kPa ✓	[15]
QUEST 4.1 4.2	ION 4Temperature: $25 \circ C / 298 \text{ K} \checkmark$ Pressure: $101,3 \text{ kPa} / 1,013 \times 10^5 \text{ Pa} / 1 \text{ atm} / 100 \text{ kPa} \checkmark$ Concentration: $1 \text{ mol·dm}^{-3} \checkmark$ Cd(s) / Cadmium / Cd Cd ²⁺ / Cd ²⁺ Cd ✓ $E_{cell}^{\theta} = E_{cathode}^{\theta} - E_{anode}^{\theta} \checkmark$ $0,13 \checkmark = E_{cathode}^{\theta} - (-0,40) \checkmark$	[15] (3)
QUEST 4.1 4.2 4.2.1 4.2.2	HON 4 Temperature: $25 \circ C / 298 \text{ K} \checkmark$ Pressure: $101,3 \text{ kPa} / 1,013 \times 10^5 \text{ Pa} / 1 \text{ atm} / 100 \text{ kPa} \checkmark$ Concentration: $1 \text{ mol·dm}^{-3} \checkmark$ Cd(s) / Cadmium / Cd Cd ²⁺ / Cd ²⁺ Cd ✓ $E_{cell}^{\theta} = E_{cathode}^{\theta} - E_{anode}^{\theta} \checkmark$	[15] (3)
QUEST 4.1 4.2 4.2.1	ION 4 Temperature: $25 \circ C / 298 \text{ K} \checkmark$ Pressure: $101,3 \text{ kPa} / 1,013 \times 10^5 \text{ Pa} / 1 \text{ atm} / 100 \text{ kPa} \checkmark$ Concentration: $1 \text{ mol·dm}^{-3} \checkmark$ Cd(s) / Cadmium / Cd Cd ²⁺ / Cd ²⁺ Cd \checkmark $E_{cell}^{\theta} = E_{cathode}^{\theta} - E_{anode}^{\theta} \checkmark$ $0,13 \checkmark = E_{cathode}^{\theta} - (-0,40) \checkmark$ $E_{cathode}^{\theta} = -0,27 \forall \checkmark$	[15] (3) (1)

(2)

(1)

(2) [10]

(1)

(1)

(3)

4.4 **OPTION 1**

9	Compare Q ²⁺ & Cd ²⁺	Q^{2+} is reduced / Cd is oxidised and therefore Q^{2+} is a stronger oxidising agent than Cd ²⁺ .	\checkmark
L	Compare R ₂ &	R_2 is reduced / Cd is oxidised and therefore <u>R_2 is a stronger</u>	\checkmark
7	Cd ²⁺	oxidising agent than Cd^{2+} .	
Ц	Compare R ₂ &	The cell potential of combination II is higher than that of	
6	Q ²⁺	combination I, therefore \mathbf{R}_2 is a stronger oxidising agent than	\checkmark
		<u>Q²⁺</u> .	
	Final answer	$Cd^{2+}; Q^{2+}; R_2 OR Cd^{2+}; Ni^{2+}; C\ell_2$	\checkmark

OPTION 2

The reduction potential of <u>Cl</u> | <u>Cl</u>₂ = 1,36 V ✓ because the cell potential of combination II is 1,76 V and the reduction potential of Cd | Cd²⁺ is 0,4 V. OR

<u>**R**</u>₂ is <u>**C**</u><u>1</u>₂ because the cell potential of combination II is 1,76 V and the reduction potential of Cd[Cd²⁺ is 0,4 V.

- CdICd²⁺ has the lowest reduction potential (-0,4 V) and therefore Cd²⁺ is the weakest oxidising agent. \checkmark
- Cℓ ICl₂ has the highest reduction potential and therefore Cl₂ is the strongest oxidising agent. ✓
- Final answer: Cd^{2+} ; Q^{2+} ; $R_2 \checkmark OR Cd^{2+}$; Ni^{2+} ; Cl_2 (4) [16]

QUESTION 5

5.1	B✓	(1)	

5.2

- 5.2.1 $C\ell_2(g) + 2e^- \rightarrow 2C\ell(aq) \checkmark \checkmark$
- 5.2.2 $C\ell_2$ / chlorine \checkmark
- 5.3 $\begin{array}{c} \begin{array}{c} \begin{array}{c} \textbf{OPTION 1/OPS/E 1} \\ E_{cell}^{\theta} = E_{callbode}^{\theta} E_{anode}^{\theta} \\ \end{array} \\ = 1,36 \checkmark -(-2,36) \checkmark \\ \end{array} \\ = 3,72 \lor \checkmark \end{array}$ (4)
- 5.4 The Mg electrode becomes smaller. / The mass of the Mg electrode decreases. / The Mg electrode being corroded. ✓
 - Magnesium is oxidised. / Mg \rightarrow Mg²⁺ + 2e⁻ \checkmark

QUESTION 6

- 6.1 <u>Electrons are transferred</u>. ✓ OR The oxidation number of Mg / H changes. OR Mg is oxidised / H⁺ is reduced.
 6.2 H⁺ ions / HCℓ / H⁺(aq) / HCℓ(aq) ✓
- Ag is a weaker reducing agent ✓ than H₂ and will not be oxidised ✓ to Ag⁺ ✓
 OR
 H₂ is a stronger reducing agent ✓ than Ag and will be oxidised ✓ to H⁺.✓

Physical selected and tanmore physics.com	FS/2017
6.4 Electrode / Conductor of electrons in hydrogen half-cell ✓	(1)
 6.5 6.5.1 Chemical energy to electrical energy ✓ 	(1)
6.5.2 Provides path for movement of ions./Completes the circuit./Ensures el neutrality in cell. \checkmark	lectrical (1)
$6.5.3 \qquad 2H^+ + 2e^- \rightarrow H_2 \checkmark \checkmark$	(2)
6.5.4 Mg(s) $Mg^{2+}(aq) \parallel H^{+}(aq) \mid H_2(g) \mid Pt$	
OR Mg(s) Mg ²⁺ (1 mol·dm ⁻³) H ⁺ (1 mol·dm ⁻³) H ₂ (g) Pt	(3)
6.6 $E_{cell}^{\theta} = E_{reduction}^{\theta} - E_{oxidation}^{\theta} \checkmark$ $= 0,00 \checkmark - (-2,36) \checkmark$ $= 2,36 \lor \checkmark$	(4)
6.7 Increases ✓	(1) [18]
QUESTION 7	
 7.1 7.1.1 AgNO₃ / Silver nitrate ✓ 	(1)
7.1.2 Ni \rightarrow Ni ²⁺ + 2e ⁻ $\checkmark \checkmark$	(2)
7.1.3 Ni + 2Ag ⁺ $\checkmark \rightarrow$ Ni ²⁺ + 2Ag \checkmark Bal \checkmark	
OR Ni + 2 AgNO ₃ \rightarrow Ni(NO ₃) ₂ + 2Ag	(3)
 7.2 7.2.1 Ni √ Ni is a stronger reducing agent. / Ni is the anode. / Ni loses electrons. oxidised. √ 	/ Ni is (2)
7.2.2 Ni (s) Ni ²⁺ (aq) Ág ⁺ (aq) Ág ^(s)	
OR Ni (s) Ni ²⁺ (1 mol·dm ⁻³) Ag ⁺ (1 mol·dm ⁻³) Ag(s)	(3)
7.2.3 $E_{cell}^{\theta} = E_{reduction}^{\theta} - E_{oxidation}^{\theta} \checkmark$ = 0,80 \sqcap - (-0,27) \sqcap = 1,07 V \sqcap \sqcap =	(4)
7.2.4 Increases ✓	(1) [16]

Terms, definitions, questions and answers

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ELECTROLYTIC CELLS

1.2 Q * & T × (13 Ct ² + 2e → Cu × × 1.3.1 Ct ² / chorine gas × (1.3.2 Cu ^{2*} ions / copper(II) ions / CuCt ² / copper(II) chloride × (1.4 Cu is a stronger reducing agent × than Ct ions × and Cu will be oxidised × to Cu ^{2*} . OR Ct ions is a weaker reducing agent × than Cu × and Cu will be oxidised × to Cu ^{2*} . OR Ct ions is a weaker reducing agent × than Cu × and Cu will be oxidised × to Cu ^{2*} . (14 Cu is a stronger reducing agent × than Cu × and Cu will be oxidised × to Cu ^{2*} . (15 Culestion 2 2.1 A solution that conducts electricity through the movement of ions. × × (15 Culestion 2 2.2.1 2H ₂ O + 2e ⁻ → H ₂ + 2OH ⁻ × × 2.2.2 Chlorine gas / Ct ² × (12.3 H ₂ O is a stronger oxidising agent × than Na ⁺ and will be reduced × to H ₂ . (12.3 H ₂ O is a stronger oxidising agent × than Na ⁺ and will be reduced × to H ₂ . (2) 3.1 A solution that conducts electricity. / Graphite is a conductor. × (2) 3.3 Ni ^{2*} (aq) + 2e ⁻ → Ni(s) × × (3.3 3.3.1 Ni ^{2*} (aq) + 2e ⁻ → Ni(s) × × (3.4 Ring × Reduction takes place at the cathode. ×			
1.2 Q $A = T \checkmark$ 1.3 C $L_2 \land C L_2 \land C L_2 \land C L C L_2 \land C C C L_2 \land C C C L_2 \land C C L C L_2 \land C C L C L C L C L C L C L C L C L C L$	QUEST	ION 1	
Image: Second Secon	1.1	Electrolytic ✓	(1)
1.3 $C_{I_2}/chlorine gas \checkmark$ (1.3.1 $C_{I_2}/chlorine gas \checkmark$ (1.3.2 Cu^{2^*} ions / copper(II) ions / $CuC_{I_2}/copper(II)$ chloride \checkmark (1.3.2 Cu^{2^*} ions / copper(II) ions / $CuC_{I_2}/copper(II)$ chloride \checkmark (1.3.2 Cu^{2^*} ions is a stronger reducing agent \checkmark than CI ions \checkmark and Cu will be oxidised \checkmark to Cu^{2^*} . (1.3.2 Cu^{2^*} ions is a weaker reducing agent \checkmark than $Cu \checkmark$ and Cu will be oxidised \checkmark to Cu^{2^*} . (1.3.2 QUESTION 2 2.1 A solution that conducts electricity through the movement of ions. \checkmark (2.2.2 (1.2.1) (1.2.2) 2.2.1 $2H_2O + 2e^- \rightarrow H_2 + 2OH^- \checkmark$ (2.2.2) (1.2.2) (1.2.2) 2.2.2 Chlorine gas / $Ct_2 \checkmark$ (2.3.3) H_2O is a stronger oxidising agent \checkmark than Na^+ and will be reduced \checkmark to H_2 . (2.2.2) QUESTION 3 3.1 A solution that conducts electricity through the movement of ions. \checkmark (2.3.3) (2.3.4) 3.1 A solution that conducts electricity. / Graphite is a conductor. \checkmark (1.3.3) (2.3.3) 3.3.1 Ni ^{2^*} (aq) + 2e ⁻ \rightarrow Ni(S) \checkmark (2.3.4) 3.3.2 Ni / nickel \checkmark (2.3.4) Ni is the anode. (2.3.4) (2.3.4) OR Ni is the anode. (2.3.4) OR Ni is the	1.2	QY&TV	
1.3.1 $Cl_2 / chlorine gas \checkmark$ ((1.3.2 Cu^{2*} ions / copper(II) ions / $CuCl_2 / copper(II)$ chloride \checkmark ((1.3.2 Cu is a stronger reducing agent \checkmark than Ct ions \checkmark and Cu will be oxidised \checkmark to Cu^{2*} . ((1.3.2) OR Ct ions is a weaker reducing agent \checkmark than Cu \checkmark and Cu will be oxidised \checkmark to Cu^{2*} . (1.3.2) OR Ct ions is a weaker reducing agent \checkmark than Cu \checkmark and Cu will be oxidised \checkmark to Cu^{2*} . (1.3.2) OR Ct ions is a weaker reducing agent \checkmark than Cu \checkmark and Cu will be oxidised \checkmark to Cu^{2*} . (1.3.2) OR Ct ions is a weaker reducing agent \checkmark than Cu \checkmark and Cu will be oxidised \checkmark to Cu^{2*} . (1.3.2) $QUESTION 2$ 2.1 A solution that conducts electricity through the movement of ions. $\checkmark \checkmark$ (2.2.2) $2.2.1$ $2H_2O + 2e^{-} \rightarrow H_2 + 2OH^{-} \checkmark \checkmark$ (2.2.2) (2.2.2) (2.2.2) $Chlorine gas / Cl_2 \checkmark$ (2.2.2) (2.2.2) (2.2.2.2) (2.2.2.2) (2.2.2.2.2.2.2.2.2.2.2.2.2.2.2.2.2.2.2.		$Cu^{2+} + 2e \rightarrow Cu \checkmark \checkmark$	(4)
1.4 Cu is a stronger reducing agent than Ct ions and Cu will be oxidised to Cu ²⁺ . Ct ions is a weaker reducing agent than Cu and Cu will be oxidised to Cu ²⁺ . CUESTION 2 1 A solution that conducts electricity through the movement of ions. (1 2.1 A solution that conducts electricity through the movement of ions. (1 2.2 2.1 2H ₂ O + 2e ⁻ → H ₂ + 2OH ⁻ (1 2.2.2 Chlorine gas / Ct ₂ (1 2.3 H ₂ O is a stronger oxidising agent than Na ⁺ and will be reduced to H ₂ . 3.1 A solution that conducts electricity through the movement of ions. (2 3.2 Plastic is a non-conductor of electricity. / Graphite is a conductor. (1) 3.3 Ni ²⁺ (aq) + 2e ⁻ → Ni(s) (2) 3.3.2 Ni / nickel (2) 3.3.2 Ni / nickel (2) Ni is the anode. (2) OR Ni is the anode. (2) 3.4 Ring Reduction takes place at the cathode. OR Ni is the anode. (2) OR Ni is the cathode. (2) <td></td> <td>Cℓ₂ / chlorine gas ✓</td> <td>(1)</td>		Cℓ₂ / chlorine gas ✓	(1)
OR Cf ions is a weaker reducing agent \checkmark than Cu \checkmark and Cu will be oxidised \checkmark to Cu ^{2*} .(1QUESTION 22.1A solution that conducts electricity through the movement of ions. \checkmark (12.22.2.1 $2H_2O + 2e^{\circ} \rightarrow H_2 + 2OH^{\circ} \checkmark$ (22.2.2Chlorine gas / $Cl_2 \checkmark$ (12.3 H_2O is a stronger oxidising agent \checkmark than Na* and will be reduced \checkmark to H_2 .(2QUESTION 33.1A solution that conducts electricity through the movement of ions. \checkmark (2)3.2Plastic is a non-conductor of electricity. / Graphite is a conductor. \checkmark (1)3.33.3.1Ni ²⁺ (aq) + 2e^{\circ} \rightarrow Ni(s) \checkmark (2)3.3.2Ni / nickel \checkmark Ni is oxidised. \checkmark OR 	1.3.2	Cu^{2+} ions / copper(II) ions / $CuCl_2$ / copper(II) chloride \checkmark	(1)
Cr ions is a weaker reducing agent than Cu and Cu will be oxidised to Cu ²⁺ .QUESTION 22.1A solution that conducts electricity through the movement of ions. (2.22.2.1 $2H_2O + 2e^{-} \rightarrow H_2 + 2OH^{-} \checkmark \checkmark$ (2.2.2Chlorine gas / $Ct_2 \checkmark$ (2.3 H_2O is a stronger oxidising agent than Na* and will be reduced to H_2.CUESTION 33.1A solution that conducts electricity through the movement of ions. (3.2Plastic is a non-conductor of electricity. / Graphite is a conductor. (3.3 $Ni^{2*}(aq) + 2e^{-} \rightarrow Ni(s) \checkmark$ (2)3.3.1 $Ni^{2*}(aq) + 2e^{-} \rightarrow Ni(s) \checkmark$ (2)3.3.2Ni / nickel (Ni is oxidised. (ORNi is the anode.(ORNi is the positive electronde.(2)3.4Ring Reduction takes place at the cathode. (ORNegative electrode.(2)	1.4		
2.1A solution that conducts electricity through the movement of ions. \checkmark ((2.22.2.1 $2H_2O + 2e^- \rightarrow H_2 + 2OH^- \checkmark$ ((2.2.2Chlorine gas / $Ct_2 \checkmark$ ((2.3 H_2O is a stronger oxidising agent \checkmark than Na ⁺ and will be reduced \checkmark to H_2 .((QUESTION 33.1A solution that conducts electricity through the movement of ions. \checkmark (2)3.2Plastic is a non-conductor of electricity. / Graphite is a conductor. \checkmark (1)3.33.3.1Ni ²⁺ (aq) + 2e ⁻ \rightarrow Ni(s) \checkmark (2)3.3.2Ni / nickel \checkmark (2)3.3.2Ni / nickel \checkmark (2)3.4Ring \checkmark Reduction takes place at the cathode. \checkmark OR Negative electrode.(2)3.4			(3) [10]
2.2 2.2.1 $2H_2O + 2e^{-} \rightarrow H_2 + 2OH^{-} \checkmark \checkmark$ ((2) 2.2.2 Chlorine gas / $Cl_2 \checkmark$ ((2) 2.3 H_2O is a stronger oxidising agent ✓ than Na ⁺ and will be reduced ✓ to H_2 . ((2) QUESTION 3 3.1 A solution that conducts electricity through the movement of ions. ✓ √ (2) 3.2 Plastic is a non-conductor of electricity. / Graphite is a conductor. ✓ (1) 3.3 3.3.1 Ni ²⁺ (aq) + 2e ⁻ → Ni(s) ✓ ✓ (2) 3.3.2 Ni / nickel ✓ (2) OR Ni is set electrons. (2) OR Ni is the positive electrode. (2) 3.4 Ring ✓ (2) Reduction takes place at the cathode. ✓ OR (2) 3.4 Ring ✓ (2) 3.4 Ring ✓ (2) 3.4 Ring ✓ (2) Negative electrode. (2)	QUEST	ION 2	
2.2.1 $2H_2O + 2e^- \rightarrow H_2 + 2OH^- \checkmark \checkmark$ (2.2.2Chlorine gas / $Cl_2 \checkmark$ (2.3 H_2O is a stronger oxidising agent \checkmark than Na* and will be reduced \checkmark to H_2 .(QUESTION 33.1A solution that conducts electricity through the movement of ions. $\checkmark \checkmark$ (2)3.2Plastic is a non-conductor of electricity. / Graphite is a conductor. \checkmark (1)3.33.3.1Ni ²⁺ (aq) + 2e^- \rightarrow Ni(s) $\checkmark \checkmark$ (2)3.3.2Ni / nickel \checkmark Ni is oxidised. \checkmark OR Ni is the anode. 	2.1	A solution that conducts electricity through the movement of ions. $\checkmark\checkmark$	(2)
2.3 H_2O is a stronger oxidising agent \checkmark than Na ⁺ and will be reduced \checkmark to H_2 . ([] QUESTION 3 3.1 A solution that conducts electricity through the movement of ions. \checkmark (2) 3.2 Plastic is a non-conductor of electricity. / Graphite is a conductor. \checkmark (1) 3.3 $Ni^{2+}(aq) + 2e^{-} \rightarrow Ni(s) \checkmark$ (2) 3.3.1 $Ni^{2+}(aq) + 2e^{-} \rightarrow Ni(s) \checkmark$ (2) 3.3.2 Ni / nickel \checkmark (2) 3.3.2 Ni / nickel \checkmark (2) 3.3.2 Ni is notes electrons. (2) OR Ni is the anode. (2) 3.4 Ring \checkmark Reduction takes place at the cathode. \checkmark OR Ni is the cathode. (2) 3.4 Ring \checkmark Reduction takes place at the cathode. \checkmark (2) 3.4 Ring \checkmark Reduction takes place at the cathode. \checkmark (2) OR Negative electrode. (2) (2)		$2H_2O + 2e^- \rightarrow H_2 + 2OH^- \checkmark\checkmark$	(2)
QUESTION 3 3.1 A solution that conducts electricity through the movement of ions. \checkmark (2) 3.2 Plastic is a non-conductor of electricity. / Graphite is a conductor. \checkmark (1) 3.3 3.3.1 Ni ²⁺ (aq) + 2e ⁻ \rightarrow Ni(s) \checkmark (2) 3.3.2 Ni / nickel \checkmark (2) 3.3.2 Ni / nickel \checkmark (2) 3.3.4 Ni loses electrons. (2) OR Ni is the anode. (2) OR Ni is the positive electrode. (2) 3.4 Ring \checkmark Reduction takes place at the cathode. \checkmark OR Negative electrode. (2)	2.2.2	Chlorine gas / $Cl_2 \checkmark$	(1)
3.1 A solution that conducts electricity through the movement of ions. \checkmark (2) 3.2 Plastic is a non-conductor of electricity. / Graphite is a conductor. \checkmark (1) 3.3 $3.3.1$ Ni ²⁺ (aq) + 2e ⁻ \rightarrow Ni(s) \checkmark (2) 3.3.1 Ni ²⁺ (aq) + 2e ⁻ \rightarrow Ni(s) \checkmark (2) 3.3.2 Ni / nickel \checkmark (2) 3.3.2 Ni / nickel \checkmark (2) OR Ni loses electrons. (2) OR Ni is the anode. (2) 3.4 Ring \checkmark Reduction takes place at the cathode. \checkmark (2) OR Negative electrode. (2)	2.3	H_2O is a stronger oxidising agent \checkmark than Na^{+} and will be reduced \checkmark to $H_2.$	(2) [7]
3.2 Plastic is a non-conductor of electricity. / Graphite is a conductor. (1) 3.3 $3.3.1$ Ni ²⁺ (aq) + 2e ⁻ \rightarrow Ni(s) (2) 3.3.2 Ni / nickel \checkmark (2) 3.3.2 Ni / nickel \checkmark (2) 3.3.2 Ni / nickel \checkmark (2) 3.3.4 Ring \checkmark (2) 3.4 Ring \checkmark Reduction takes place at the cathode. (2) 3.4 Ring \checkmark Reduction takes place at the cathode. (2) (2) (2) (2)	QUEST	ION 3	
3.3 $3.3.1$ $Ni^{2^+}(aq) + 2e^- \rightarrow Ni(s) \checkmark \checkmark$ (2) 3.3.2 Ni / nickel \checkmark (2) 3.3.4 Ring \checkmark Reduction takes place at the cathode. \checkmark (2) 3.4 Ring \checkmark Reduction takes place at the cathode. \checkmark (2) 3.4 Ring \checkmark Reduction takes place at the cathode. \checkmark (2)	3.1	A solution that conducts electricity through the movement of ions. $\checkmark\checkmark$	(2)
3.3.1 $Ni^{2^+}(aq) + 2e^- \rightarrow Ni(s) \checkmark \checkmark$ (2)3.3.2Ni / nickel \checkmark Ni is oxidised. \checkmark OR OR 	3.2	Plastic is a non-conductor of electricity. / Graphite is a conductor. \checkmark	(1)
Ni is oxidised. \checkmark ORNi loses electrons.Image: Construction of the positive electrode.Image: Construction of the positive e		$Ni^{2+}(aq) + 2e^{-} \rightarrow Ni(s) \checkmark \checkmark$	(2)
Reduction takes place at the cathode. ✓ OR Negative electrode. (2)	3.3.2	Ni is oxidised. ✓ORNi loses electrons.ORNi is the anode.	(2)
3.5 Decreases √	3.4	Ring ✓ Reduction takes place at the cathode. ✓ OR	(2)
Ni ²⁺ ions from the electrolyte will be reduced (to Ni). ✓ OR Ni ²⁺ changes to Ni (2)	3.5	Ni ²⁺ ions from the electrolyte will be reduced (to Ni). \checkmark OR	(2) [11]

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QUESTION 4

4.1	ANY ONE: The chemical process in which electrical energy is converted to			
	 <u>chemical energy</u>. The <u>use</u> of <u>electrical energy to produce a chemical change</u>. The <u>process</u> during which an <u>electric current passes through a solution / molten</u> <u>ionic</u> compound. 	(2)		
4.2	 ANY ONE: To keep the polarity of the electrodes constant. √ To prevent the anode and cathode from swopping. DC provides a <u>one way flow of electrons</u> ensuring that the same chemical reaction occurs all the time at the same electrodes. 	(1)		
4.3	$Cu^{2+}(aq) + 2e^{-} \rightarrow Cu(s) \checkmark \checkmark$	(2)		
4.4	 <u>Cu²⁺ is a stronger oxidising agent</u> √ <u>than Zn²⁺</u>. √ <u>Cu²⁺ will be reduced</u> to Cu. √ 			
	 OR <u>Zn is a stronger reducing agent than Cu</u>. <u>Cu²⁺ will be reduced</u> to Cu. 	(3)		
QUEST	ION 5			
5.1	Electrolytic cell ✓	(1)		
5.2	The substance/species which loses electrons. $\checkmark\checkmark$	(2)		
5.3	P✓	(1)		
5.4	$Cu(s) \rightarrow Cu^{2+}(aq) + 2e^{-\sqrt{\sqrt{2}}}$			
5.5	A \checkmark Ct ions move to the positive electrode / anode where they are oxidised to Ct $_2$. $\checkmark\checkmark$	(3) [12]		
QUEST	ION 6	[]		
6.1 6.1.1	Electrolyte ✓	(1)		
6.1.2	Electrolytic cell ✓	(1)		
6.2	A to B ✓	(1)		
6.3 6.3.1	B √	(1)		
6.3.2	A 🗸	(1)		
6.4	Decreases ✓			
	Copper (Cu) is oxidised to Cu ²⁺ / Oxidation takes place at A. \checkmark	(2) [7]		

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QUESTION 7

7.1	Endothermic 🗸	(1)
7.2	Anode 🗸	
La	Connected to the positive terminal of the battery. \checkmark	(2)
7.3	וחם	
7.3.1	Chlorine gas / Cl₂ ✓	(1)
7.3.2	Hydrogen gas /H₂ ✓	(1)
7.3.3	$2H_2O(l) + 2e^- \rightarrow H_2(g) + 2OH^-(aq) \checkmark \checkmark$	(2)
7.4	Basic / alkaline ✓ OH⁻ ions form / NaOH forms ✓	(2) [9]



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FERTILISERS

QUESTION 1			
1.1 1.1.1 Nitrogen / $N_2 \checkmark$ Hydrogen / $H_2 \checkmark$		(2))
1.1.2 NH_3 + HNO ₃ $\checkmark \rightarrow \operatorname{NH}_4\operatorname{N}$	$O_3 \checkmark$ Bal. \checkmark	(3))
1.2 Marking criteria: Use ratio: $\frac{3}{9} \checkmark$ x 20 kg \checkmark x 36 / 36 % \checkmark Final answer: 2,4 kg \checkmark OPTION 1: % N = $\frac{3}{9} \checkmark$ (x 36) \checkmark = 12 % \therefore m(N): $\frac{12}{100}$ (× 20 kg) = 2,4 kg \checkmark	$\frac{\text{OPTION 2:}}{\text{m(nutrients):}}$ $\frac{36}{100} \checkmark (\text{x 20}) = 7,2 \text{ kg}$	$ \frac{\text{OPTION 3:}}{m(N):} \\ \frac{3^{\checkmark}}{9} \times (\times 20) (\times \frac{36^{\checkmark}}{100}) = 2,4 \text{ kg} $ (4)	

QUESTION 2

2.1 Contact process ✓ (1) 2.2 Vanadium pentoxide / Vanadium(V) oxide / V₂O₅ ✓ 2.2.1 (1) 2.2.2 $H_2S_2O_7(\ell) + H_2O(\ell) \checkmark \rightarrow 2H_2SO_4(\ell) \checkmark$ Bal. \checkmark (3) 2.3 2.3.1 $H_2SO_4 + 2NH_3 \checkmark \rightarrow (NH_4)_2SO_4 \checkmark Bal. \checkmark$ (3) 2.3.2 Ammonium sulphate ✓ (1) 2.4 2.4.1 Total percentage of fertiliser √ (1)Mass of fertiliser in P/Massa kunsmis in P: $\frac{25}{100} \times 50 = 12,5 \text{ kg}$ 2.4.2 Mass of fertiliser in Q/Massa kunsmis in Q: $\frac{20}{100} \times 50 = 10$ kg Amount of potassium in P/Massa kalium in P: $\frac{3}{10}$ x 12,5 = 3,75 kg \checkmark Amount of potassium in Q/Massa kalium in Q: $\frac{4}{8}$ x 10 = 5 kg \checkmark Fertiliser Q has more potassium per mass than fertiliser P. </ Kunsmis Q het meer kalium per massa as kunsmis P. (4) [14]

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QUESTION 3

3.1 3.1.1	$2SO_2(g) + O_2(g) \checkmark \rightarrow 2SO_3(g) \checkmark$ Bal. \checkmark	(3)
3.1.2	Catalyst ✓	
	OR Increase the reaction rate	(1)
3.2	Exothermic ✓ The temperature increases ✓	(2)
3.3	 An exothermic reaction is favoured by a decrease in temperature. ✓ The forward reaction is favoured. ✓ Higher yield (of SO₃). ✓ OR An endothermic reaction is favoured by an increase in temperature. ✓ The reverse reaction is favoured. ✓ Lower yield (of SO₃). ✓ 	(3)
3.4 3.4.1	$H_2S_2O_7 \checkmark$	(1)
3.4.2	A mist will form (which is difficult to collect). ✓ OR The reaction is too exothermic.	(1)
3.5	$H_2SO_4 + 2NH_3 \checkmark \rightarrow (NH_4)_2SO_4 \checkmark Bal. \checkmark$	(3) [14]

QUESTION 4

4.1 4.1.1	Haber process ✓	(1)
4.1.2	$N_2 + 3H_2 \checkmark \rightleftharpoons 2NH_3 \checkmark$ bal \checkmark	(3)
4.1.3	Air ✓	(1)
4.2 4.2.1	40% ✓	(1)
4.2.2	 High yield ✓ <u>High rate</u> due to higher concentration. ✓ 	(2)
4.2.3	Low reaction rate ✓	(1)

OPTION 1	OPTION 2	
% N in NH ₄ NO ₃ = $\frac{28}{80}$ \checkmark ×100	$\overline{\mathrm{m(N in NH}_4\mathrm{NO}_3)} = \frac{28}{80} \checkmark \times 50 \checkmark$	
= 35%	= 17,5 kg √	
m(N) in 50 kg:		
$\frac{35}{100} \times 50 \checkmark = 17,5 \text{ kg} \checkmark$		
$\frac{100}{100}$ 300 $^{-17,3}$ kg 10		

(3)

(1) **[10]**

QUESTION 5

5.1	Ostwald process ✓		(1)
5.2	NO / nitrogen monoxide \checkmark Water / H ₂ O \checkmark		(2)
5.3	$NH_3 + HNO_3 \checkmark \rightarrow NH4NO_3 \checkmark$	bal 🗸	(3)
5.4	$\frac{\text{OPTION 1/OPTION 1}}{n(\text{NH}_3) = \frac{\text{m}}{\text{M}}}$ $= \frac{6.8 \times 10^7}{17 \sqrt{2}}$ $= 4 \times 10^6 \text{ mol}$	$\frac{\text{OPTION } 2/\text{OPSIE 2}}{\text{m}(\text{NH}_4\text{NO}_3)} = \underbrace{\begin{array}{c} 6,8 \times 10^4 \\ 17^{\checkmark} \times 80^{\checkmark} \end{array}}_{= 3,2 \times 10^5 \text{ kg}^{\checkmark}}$	
	$ \int \\ n(NH_4NO_3) = n(NH_3) \\ = 4 \times 10^6 \text{ mol} \\ m(NH_4NO_3) = nM \\ = (4 \times 10^6)(80) \checkmark \\ = 3,2 \times 10^8 \text{ g} \\ = 3,2 \times 10^5 \text{ kg} \checkmark $	OPTION 3/OPS/E 3 17 g \checkmark NH ₃ forms/vorm 80 g \checkmark NH ₄ NO ₃ 6,8 x 10 ⁴ kg forms/vorm x g NH ₄ NO ₃ x = 6,8 x 10 ⁴ x $\frac{80}{17}$ = 3,2 x 10 ⁵ kg \checkmark	(3)

5.5 To make a NPK fertiliser . / To make fertilisers which contain all three primary nutrients. \checkmark

QUESTION 6

6.3.2	Ostwald process ✓	(1) [14]
	Ammonium nitrate (has the highest percentage of nitrogen). \checkmark	(4)
6.3 6.3.1		
6.2.4	$SO_3 + H_2SO_4 \checkmark \rightarrow H_2S_2O_7 \checkmark$ Bal. \checkmark	(3)
6.2.3	H₂SO₄ ✓	(1)
6.2.2	Ammonia ✓	(1)
6.2 6.2.1	Haber process ✓	(1)
6.1.3	Sulphur / iron pyrite / iron sulphide ✓	(1)
6.1.2	Natural gas / methane / oil / coal ✓	(1)
6.1 6.1.1	Air ✓	(1)

QUESTION 7

7.1			
7.1.1	Haber process ✓		(1)
7.1.2	Contact process / Catalytic oxidation of SO2	\sim	(1)
7.1.3	Sulphur trioxide / SO $_3$ \checkmark		(1)
7.1.4	$SO_3 + H_2SO_4 \checkmark \rightarrow H_2S_2O_7 \checkmark$	Bal. ✓	(3)
7.1.5	$H_2SO_4\checkmark + 2NH_3\checkmark \rightarrow (NH_4)_2SO_4\checkmark$	Bal. ✓	(4)
7.2	 Marking guidelines: Calculate the mass of fertiliser. Add %N and %P OR mass N and mass P. Subtraction: 100 - (%N + %P) OR m(fertiliser) - [m(N) + m(P)] OR %fertiliser - [%N + %P] Final answer: 8:1:5 		
	$\frac{\text{OPTION 1}}{\text{m(fertiliser)}} = \frac{36}{100} \times 20 \checkmark$ $= 7.2 \text{ kg}$	$\frac{\text{OPTION 2}}{\text{m(fertiliser)}} = \frac{36}{100} \times 20 \checkmark$ $= 7.2 \text{ kg}$	
	$\%$ N = $\frac{4,11}{7,2}$ x 100 = 57,08%	$m(K) = \frac{7.2 - \checkmark}{2.58} (4.11 + 0.51) \checkmark$ = 2.58 kg	

4,11	:	0,51	:	2,58
8	:	1	:	5 √

= 7,08%%K = $\underline{100} - \checkmark (57,08 + 7,08) \checkmark$ = 35,84%57,08 : 7,08 : 35,848 : 1 : $5\checkmark$ $\overline{OPTION 3}$ %N = $\frac{4,11}{20} \times 100 = 20,55\%$ %P = $\frac{0,51}{20} \times 100 = 2,55\%$ %K = $\underline{36} - \checkmark (20,55 + 2,55) \checkmark = 12,9\%$ 20,55 : 2,55 : 12,9 8 : 1 : $5\checkmark$

(4) **[14]**

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 $\% P = \frac{0.51}{7.2} \times 100$