CANDIDATE NAME	CT GROUP	22\$
	J	

CHEMISTRY 9729/03

Paper 3 Free Response

28 September 2022

55 min

Candidates answer on the Question Paper.

Additional Materials: Data Booklet

READ THESE INSTRUCTIONS FIRST

Write your name and CT group on all the work you hand in.

Write in dark blue or black pen.

You may use an HB pencil for any diagrams or graphs.

Do not use staples, paper clips, glue or correction fluid.

Answer **all** questions in the spaces provided on the Question Paper. If additional space is required, you should use the pages at the end of this booklet. The question number and part must be clearly shown.

Section A

Question 1 is a compulsory question.

Section B

Answer only one question from this section.

A Data Booklet is provided.

The use of an approved scientific calculator is expected, where appropriate.

You are reminded of the need for good English and clear presentation in your answers.

At the end of the examination, fasten all your work securely together.

The number of marks is given in brackets [] at the end of each question or part question.

FOR EXAMINER'S USE

Question No.	Marks obtained
1	/ 20
^Circle the question atte	empted below.
^2/3	/ 15
Total Marks	35

Section A

Question 1 is a **compulsory** question. All candidates should answer this question.

1	(a)	Volcanology is the scientific study of volcanoes. At the Earth's core, there is a molten mixture of rock, gases and crystals called magma. During a volcanic eruption, magma makes its way to the surface and flows out as lava.		
		The most abundant elements in magma are oxygen and silicon.		
		(i)	State the electronic configuration of an atom of silicon. [1]	
		(ii)	Describe one similarity and one difference between a 3s orbital and a 3p orbital. [2]	
		(iii)	Describe one difference between a 2p orbital and a 3p orbital. [1]	

(b) Silica, SiO_2 , is a major component of lava. At the high temperatures during eruptions, silica in the lava can react with other metals present to form a variety of salts, such as basalt.

Analysis of a sample of basalt produced the following composition:

element	mass percentage (%)
Ca	18.51
Mg	11.22
Si	25.95
0	44.32

(i)	Use the information provided above to determine the empirical formula of basalt. [2]
(ii)	Basalt can be thought of as a mixture of a salt of calcium and a salt of magnesium, both containing the same anion. Based on your answer to (b)(i) , state the formula of this anion. [1]

(c) Hydrocarbon reservoirs can often be found in volcanic zones. Pentenes, one of the byproducts of thermal cracking of hydrocarbons, are alkenes with the formula C₅H₁₀. Pentenes are generally very volatile and boil just above room temperature. (i) D, E, and F are three straight-chain pentenes with different structures. E and F are a pair of stereoisomers. Draw the structures of **D**, **E** and **F**, and name each of them. [3] (ii) Explain how the isomerism between **E** and **F** arises. [2] Hydrogenation is a useful way to convert unsaturated hydrocarbons, such as pentenes, into saturated hydrocarbons. $C_5H_{10}(g) + H_2(g) \rightarrow C_5H_{12}(g)$ This conversion is usually conducted in the presence of a finely divided metal catalyst. Without the catalyst, pentenes do not react readily with hydrogen gas. (iii) Explain, with the aid of a labelled Maxwell-Boltzmann distribution diagram, the effect on a rate constant of adding a catalyst. [3] (iv) At very low pressures, the reaction rate is directly proportional to the partial pressure of each reactant, while at higher pressures, the reaction rate appears to be independent of the partial pressure of each reactant. Suggest an explanation for this observation.

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MARGIN

d)		s an alkene with the formula C_6H_{12} . An isolated sample of G is able to rotate ne-polarised light. Draw a possible structure for hydrocarbon G . [1]
e)		a hydrocarbon with the formula C_xH_y . J is a liquid at room temperature, but a gas at ${}^{\circ}C$.
	diox	cm 3 of hydrocarbon J was exploded in exactly 95 cm 3 of oxygen and a mixture of carbon kide and steam were formed. Both J and the oxygen were completely used up. These times were measured at 150 °C and 1.0 atm.
	The	complete combustion of J is shown below.
		$C_xH_y + (x + \frac{y}{4})O_2 \rightarrow xCO_2 + \frac{y}{2}H_2O$
		en the gaseous product mixture was cooled down at 1.0 atm from 150 °C to 20 °C, 5 cm³ of gas was left after discarding the liquid water.
	(i)	Calculate the volume of CO ₂ at 150 °C. [1]
	(ii)	Hence, deduce the molecular formula of J , showing your working clearly. [1]

[Total: 201

Section B

Answer **one** question from this section.

2 (a) 2.10 g of a Group 1 metallic salt ($M_r < 500$) containing the $I_x C l_y^-$ anion decomposed when heated to form iodine vapour and the metallic chloride as the residue.

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The iodine evolved was bubbled into excess potassium iodide solution. The iodine required 25.56 cm³ of 0.500 mol dm⁻³ aqueous sodium thiosulfate for complete reaction. The ionic equation for the reaction between iodine and thiosulfate is shown below:

$$I_2 + 2S_2O_3^{2-} \rightarrow S_4O_6^{2-} + 2I^{-}$$

0.92 g of solid AgCl was obtained when the metallic chloride residue was added to an excess of aqueous silver nitrate.

- (i) Calculate the amount in moles of I atoms present in the sample of metallic salt. [1]
- (ii) Calculate the amount in moles of Cl atoms present in the sample of metallic salt. [1]

(iii) Hence, determine the formula of the I_xCl_y anion and deduce the identity of the

Group 1 metal present in the metallic salt.	[2]

as	DO NOT WRITE IN THIS MARGIN
[1]	
[1]	
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[2]	
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(b)	I_2 molecules and I^- ions react to form tri-iodide ions, I_3^- . A <i>dynamic equilibrium</i> is set up as shown in equilibrium 1.				
		equilibrium 1	$I_2(aq) + I^-(aq) \rightleftharpoons I_3^-(aq)$	Δ <i>H</i> < 0	
	(i)	Explain what is meant by	dynamic equilibrium.		[1]
	(ii)	The shape of I_3^- is linear.	. Draw a dot-and-cross diagrar	m for I ₃	[1]
	(iii)	KI(aq). With reference to	e is not very soluble in water, o the shift in position of equil n why molecular iodine is solul	ibrium 1 and forces of attract	
	Some solid I_2 is dissolved in 100 cm ³ of KI(aq) and the reaction mixture is allowed to rea equilibrium at 25 °C, according to equilibrium 1. The resulting concentrations of I_2 (aq), I^- (a and I_3^- (aq) are 0.040 mol dm ⁻³ , 0.040 mol dm ⁻³ and 0.850 mol dm ⁻³ respectively.				
	(iv)	Calculate a value of K_c for	or equilibrium 1.		[1]
	(v)	Explain the significance of	of the magnitude of your answe	er in (b)(iv) .	[1]
	When more solid $\rm I_2$ is added to the equilibrium mixture at 25 °C, the amount of $\rm I_3^-$ increases to 0.086 mol at the new equilibrium.			ses	
	(vi)	Use your value of K_c to equilibrium concentration	calculate the amount in moles of I_2 (aq).	s of ${ m I_2}$ added and hence the ${ m r}$	new [3]
	(vii)		sign of ΔS with respect to ecopontaneity of the forward reac		

[Tot	al: 15]

 $\begin{tabular}{ll} {\bf 3} & Nitrogen dioxide, NO_2, is an oxide of nitrogen that is an air pollutant. \end{tabular}$



nitrogen dioxide

(a)	(i)	Two molecules of NO_2 can dimerise to form another oxide of nitrogen, N_2O_4 . Draw the structure of N_2O_4 and state the shape about each N atom.	ne 2]
	(ii)	The bond angle in the NO ₂ molecule is 134°, much larger than 120°. Explain why. [7]	1]

The	dimerisation of NO ₂ is a	reversible reacti	on.		
equ	ilibrium 2	$2NO_2(g) \rightleftharpoons$	$N_2O_4(g)$		
(i)	Write the K_p expression	for equilibrium 2			[1]
(ii)	x kPa of NO ₂ was intro equilibrate at a consta decreased by 100 kPa	nt temperature o	of 25 °C. The		
	Determine the partial pr	essure of N ₂ O ₄ a	t equilibrium.		[1]
(iii)	The K_p of equilibrium 2	is 6.25 × 10 ⁻² kP	a⁻¹ at 25 °C.		
	Use your answers in (b))(i) and (b)(ii) to	determine the v	alue of x.	[1]
(iv)	Deduce the effect on th decreased. Explain you		ilibrium 2 when	the volume of t	the vessel used is [2]

(b)

- (c) (i) Sketch a graph of pV vs p for a fixed mass of an ideal gas at constant temperature, where p is the pressure and V is the volume of the gas. [1]
 - (ii) When real gas molecules collide, the kinetic energy during collision is not necessarily conserved.

State **two** other properties of real gas molecules which could lead to the product *pV* for a fixed mass of real gas at constant temperature being different from that for an ideal gas.

[2]

The plotted points in Fig. 3.1 are experimental values of the product pV for a mass m of a gas **A** at a constant temperature of T.

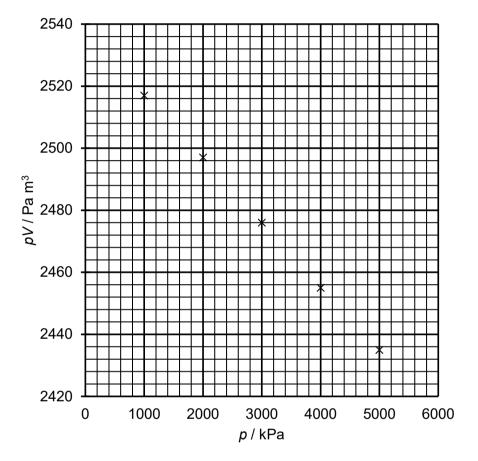


Fig. 3.1

- (iii) In Fig. 3.1, pV of gas **A** decreases as p increases, which differs from your sketch in (c)(i) for an ideal gas. Which one of the properties that you have stated in (c)(ii) results in the decrease in pV as p increases? Explain your answer briefly.
- (iv) A real gas can approach ideal behavior under certain conditions, such as high temperature. State another condition under which a real gas behaves more ideally. [1]
- (v) Use the graph in Fig. 3.1 and your answer in (c)(iv) to estimate the pV value for an ideal gas. [1]
- (vi) Hence, use the ideal gas equation to determine the relative molecular mass of gas A to 1 decimal place, given that the experimental results in Fig. 3.1 were obtained with a sample of 56 g of gas A at 150 K.

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[Total: 15]	

Additional answer space

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