Please check the examination details bel	ow before ente	ring your candidate information						
Candidate surname		Other names						
Centre Number Candidate Number								
Pearson Edexcel Level	3 GCE							
Monday 12 June 20	Monday 12 June 2023							
Morning (Time: 1 hour 45 minutes)	Paper reference	9CH0/01						
Chemistry		₾ •						
Advanced								
PAPER 1: Advanced Inorg	PAPER 1: Advanced Inorganic and Physical Chemistry							
You must have: Scientific calculator, Data Booklet, rule	er	Total Marks						

Instructions

- Use **black** ink or ball-point pen.
- If pencil is used for diagrams/sketches/graphs it must be dark (HB or B).
- **Fill in the boxes** at the top of this page with your name, centre number and candidate number.
- Answer **all** questions.
- Answer the questions in the spaces provided
 - there may be more space than you need.

Information

- The total mark for this paper is 90.
- The marks for each question are shown in brackets
 - use this as a guide as to how much time to spend on each question.
- For the question marked with an **asterisk** (*), marks will be awarded for your ability to structure your answer logically, showing the points that you make are related or follow on from each other where appropriate.
- A Periodic Table is printed on the back cover of this paper.

Advice

- Read each question carefully before you start to answer it.
- Show all your working in calculations and include units where appropriate.
- Check your answers if you have time at the end.

Turn over ▶



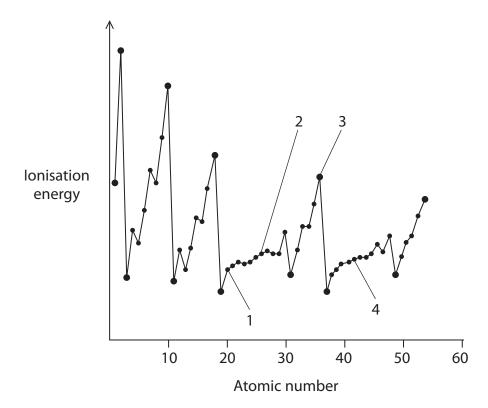




Answer ALL questions.

Some questions must be answered with a cross in a box \boxtimes . If you change your mind about an answer, put a line through the box \boxtimes and then mark your new answer with a cross \boxtimes .

- 1 This question is about d-block elements.
 - (a) Which of the labels 1-4 identifies a d-block element in Period 4?



(1)

- X A 1
- **B** 2
- □ 4

(b) State what is meant by the term d-block element.

(1)

(Total for Question 1 = 2 marks)



2			en use the term 'orbital' when con at is meant by the term orbital.	sidering atomic structure.	(2)
	(b) Draw	dia	grams to show the shape of an s a	and a p orbital.	(2)
			s orbital	p orbital	
	(c) What	is t	he electronic configuration of a co	opper atom?	(1)
	\times	Α	[Ar]4s ¹ 3d ¹⁰		
	X	В	$[Ar]4s^23d^9$		
	X	C	$[Ar]4s^24p^13d^8$		
	X	D	$[Ar]4s^24p^23d^7$		
				(Total for Question 2 = !	5 marks)



3	This questi	on is a	about compounds containing elements from Group 7.	
		_	ge occurs when concentrated sulfuric acid is added to romide?	(1)
	\boxtimes	A bi	romide ions oxidise sulfuric acid forming sulfur	(1)
	\times	B bi	romide ions oxidise sulfuric acid forming sulfur dioxide	
	\boxtimes	C bi	romide ions reduce sulfuric acid forming sulfur	
	\boxtimes	D b	romide ions reduce sulfuric acid forming sulfur dioxide	
	amoun	t of a	n test for the presence of bromide ions in solution by adding a small cidified silver nitrate solution. y of the precipitate in aqueous ammonia is then tested.	
	(i) Whi	ich sta	atement is correct for bromide ions?	(1)
	\boxtimes	Α	a white precipitate forms that dissolves in concentrated ammonia only	(1)
	\boxtimes	В	a white precipitate forms that dissolves in both dilute and concentrated ammonia	
	\boxtimes	C	a cream precipitate forms that dissolves in concentrated ammonia only	
	\boxtimes	D	a cream precipitate forms that dissolves in both dilute and concentrated ammonia	
	(ii) Give	e a rea	ason why the silver nitrate must be acidified.	(1)
			which acid needs to be used to acidify the silver nitrate solution and er acids are unsuitable.	(2)



(c) Iodine trichloride forms a dimer, I_2Cl_6 , in the solid state. When molten, it is suggested that it breaks down as shown.

$$I_2Cl_6 \rightleftharpoons ICl_2^+ + ICl_4^-$$

(i) Draw a labelled diagram of a simple experiment to confirm this dissociation has occurred, stating the positive result.

(2)

Result

(ii) What is the shape of the ICl₄ ion?

(1)

- A octahedral
- **B** square planar
- C tetrahedral
- **D** trigonal bipyramidal
- (iii) The equilibrium position for the dissociation of molten I_2Cl_6 lies to the left.

$$I_2Cl_6 \rightleftharpoons ICl_2^+ + ICl_4^-$$

What is the most likely numerical value of K_c for this equilibrium?

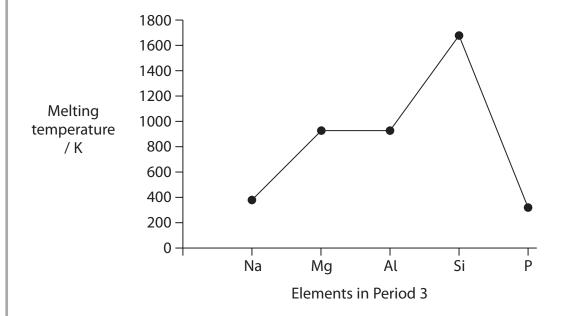
(1)

- **A** 1.0×10^6
- **B** 5.0×10^3
- **D** 5.0×10^{-3}

(Total for Question 3 = 9 marks)

(6)

4 The graph shows the melting temperatures of some elements in Period 3.



Explain the variations in melting temperature across the period in terms of the structure and bonding in these elements.



5 This question is about the decomposition of dinitrogen tetroxide. The reaction eventually reaches equilibrium.

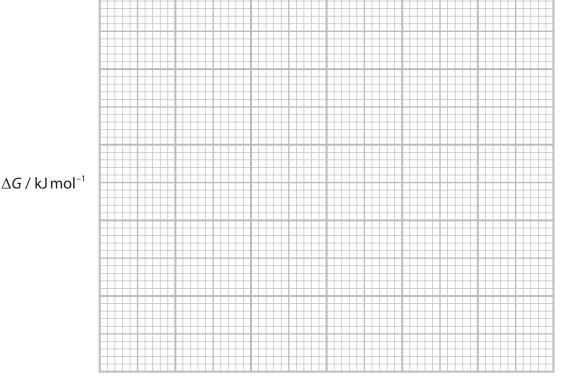
$$N_2O_4(g) \rightleftharpoons 2NO_2(g)$$
 colourless brown

(a) The table shows values of ΔG at different temperatures for this reaction.

Temperature / K	ΔG / kJ mol ⁻¹
350	-4.0
400	-13
450	-22
500	-31
550	-40

(i) Plot a graph of ΔG against temperature.

(2)



Temperature / K

(ii) Calculate the entropy change of the system, ΔS_{system} , in J K⁻¹ mol⁻¹, using your straight line from the graph in (a)(i) and the equation shown.

$$\Delta G = -T\Delta S_{\text{system}} + \Delta H \tag{3}$$

(iii) What feature of the graph in (a)(i) gives the enthalpy change of the reaction?

(1)

- A intercept of the x-axis
- lacksquare **B** (intercept of the x-axis) $\times -1$
- \square **D** (intercept of the y-axis) $\times -1$

(b) What happens to the position of the equilibrium and the colour of the mixture when the pressure is increased?

The volume of the system remains constant.

X

X

X

X

$$N_2O_4(g) \rightleftharpoons 2NO_2(g)$$
 colourless brown

(1)

	Position of equilibrium	Change in colour
Α	moves to the right	mixture gets lighter
В	moves to the right	mixture gets darker
C	moves to the left	mixture gets lighter
D	moves to the left	mixture gets darker

(Total for Question 5 = 7 marks)

6 Benzoic acid is a weak acid found in cranberries.

C₆H₅COOH – benzoic acid

(a) Which of these answers identifies the types of species present when benzoic acid is mixed with nitric acid?

[K_a of benzoic acid = 6.3×10^{-5} mol dm⁻³; K_a of nitric acid = 40 mol dm⁻³]

(1)

$$C_6H_5COOH + HNO_3 \rightleftharpoons C_6H_5COOH_2^+ + NO_3^-$$

- A acid base acid base
- **B** acid base base acid
- C base acid base acid
- □ D base acid acid base
- (b) The ionic salts sodium benzoate and potassium benzoate are both used as food preservatives.

Explain why the melting temperature of sodium benzoate is higher than the melting temperature of potassium benzoate.

(2)

 	 	•••••	 •	 	 	• • • • • • • • • • • • • • • • • • • •	 	• •								
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- (c) The value of K_a for benzoic acid = $6.28 \times 10^{-5} \, \text{mol dm}^{-3}$.
 - (i) Write the expression for the acid dissociation constant, K_a , of benzoic acid.

(1)

(ii) Calculate the mass of benzoic acid needed to prepare $250\,\mathrm{cm}^3$ of a solution with a pH = 3.51

(4)





(d) Weak acids such as benzoic acid can be neutralised by sodium hydroxide solution.

$$C_6H_5COOH(aq) + NaOH(aq) \rightarrow C_6H_5CO_2Na(aq) + H_2O(l)$$

(i) Which of these could be used to show the end-point of a titration of benzoic acid with sodium hydroxide solution?

(1)

- A bromothymol blue
- **B** litmus
- C methyl orange
- **D** phenolphthalein
- (ii) Another weak acid found in cranberries is quinic acid, $C_6H_7(OH)_4COOH$. It is neutralised by sodium hydroxide solution in a similar way to benzoic acid.

A 25.0 cm³ sample of 0.500 mol dm⁻³ quinic acid solution was neutralised under standard conditions in a polystyrene cup using 25.0 cm³ of 0.800 mol dm⁻³ of sodium hydroxide solution.

This resulted in a temperature rise of 2.9 °C.

Calculate the standard enthalpy change of neutralisation, $\Delta_{\text{neut}}H^{\ominus}$, of quinic acid in kJ mol⁻¹.

[Assume the density of both solutions is 1.0 g cm⁻³.

specific heat capacity of solution formed = $4.18 \, \text{Jg}^{-1} \, ^{\circ}\text{C}^{-1}$]

(3)

		(11111111111111111111111111111111111111	,
		(Total for Question 6 = 14 ma	rks)
			(2)
	Explain the difference between these values.		(2)
	is –57.9 kJ mol ⁻¹ .	it of the strong actuation	
(iii) The standard enthalpy change of neutralisation by sodium hydroxide is –11.7 kJ mol ⁻¹ while tha		

7	This question	is about	chromium and	d chromium	compounds.

(a)	Naturally occurring chromium has four isotopes, ⁵⁰ Cr, ⁵² Cr, ⁵³ Cr and ⁵⁴ Cr,
	State what is meant by the term isotopes.

(2)

- (b) Both chromium and calcium can form ions with a +2 charge.
 - (i) Complete the electronic configuration of a Cr^{2+} ion.

(1)

1s².....

(ii) Explain which of chromium or calcium most easily forms a +2 ion using all of the data in the table.

Element	Atomic number	1st ionisation energy / kJ mol ⁻¹	2nd ionisation energy / kJ mol ⁻¹	Metallic radius / nm
Chromium	24	653	1592	0.129
Calcium	20	590	1145	0.197

(3)



(c) Chromium(III) sulfate, $Cr_2(SO_4)_3$, dissolves in water to form the complex ion $[Cr(H_2O)_6]^{3+}$ (aq).				
	(i)	State the colour of this complex ion.	(1)	
	(ii)	Explain why the aqueous solution of this complex ion has an acidic pH by considering the interaction between the metal ion and the ligands.	(2)	

(d) A student researching the role of dichromate(VI) ions, $Cr_2O_7^{2-}$, as an oxidising agent made the statement shown.

'Standard electrode potential data shows that it is never feasible for a $1.00\,\mathrm{mol\,dm^{-3}}$ solution of potassium dichromate(VI) to oxidise the chloride ions in hydrochloric acid.'

Comment on this statement using the data and equilibria shown.

Equilibrium 1

$$Cr_2O_7^{2-}(aq) + 14H^+(aq) + 6e^- \Rightarrow 2Cr^{3+}(aq) + 7H_2O(l)$$
 $E^{\oplus} = +1.33 \text{ V}$

Equilibrium 2

$$Cl_2(aq) + 2e^- \rightleftharpoons 2Cl^-(aq)$$
 $E^{\ominus} = +1.36 V$ (4)

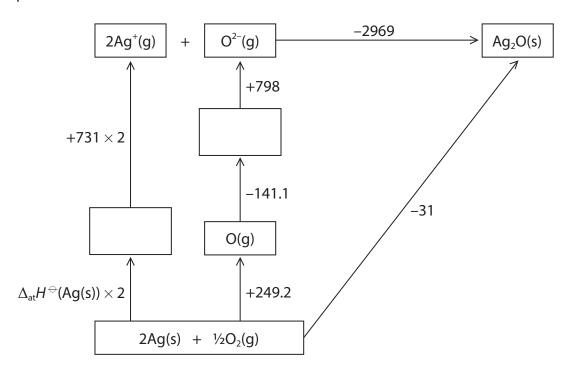


(Total for Question 7 = 13 marks)

∗8	Transition metals and their compounds can act as catalysts in many reactions such a the ones shown:	S
	• platinum, Pt, in the catalytic converters of vehicles	
	• manganese(II) ions, $Mn^{2+}(aq)$, in the oxidation of ethanedioate ions, $C_2O_4^{2-}(aq)$, by manganate(VII) ions, $MnO_4^-(aq)$.	
	Compare and contrast the role of the catalysts in these reactions.	(6)



- **9** This question is about silver compounds.
 - (a) The diagram shows a Born–Haber cycle for the formation of silver(I) oxide, Ag_2O . All quantities are measured in kJ mol⁻¹.



(i) Complete the diagram by adding appropriate species and state symbols to the empty boxes.

(2)

(ii) Explain why the value for the first electron affinity of oxygen is negative and the value for the second electron affinity is positive.

(3)

(iii) Calculate a value for the standard enthalpy change of atomisation of silver, $\Delta_{\rm at}H^{\,\ominus}$, using the Born–Haber cycle.

(3)

(b) Another silver compound is silver chloride, AgCl. Values for its lattice energy can be found by experiment or by theoretical calculation.

Compound	Experimental lattice energy / kJ mol ⁻¹	Theoretical lattice energy / kJ mol ⁻¹	
Silver chloride	-905	-833	

(i)	Give two assumptions used in the model to calculate the
	theoretical lattice energy.

(2)

(ii)	Explain the difference in the two values for the lattice energy of silver chloride
	by considering the possible bonding models.

(3)

(Total for Question 9 = 13 marks)



10 Manganese compounds can be used to determine the amounts of dissolved molecular oxygen in water samples.	
(a) Draw the dot-and-cross diagram for an oxygen molecule, ${\sf O}_2$. Show outer shell electrons only.	(1)
	(1)
(b) The colubility of eyygen in water under standard conditions	
(b) The solubility of oxygen in water under standard conditions is 1.22×10^{-3} mol dm ⁻³ .	
Comment on this value by considering the type and strength of the intermolecular forces in pure water	
pure oxygena mixture of water and oxygen.	
Detailed descriptions of the forces involved are not required.	(4)
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Detailed descriptions of the forces involved are not required.	(4)



- (c) The amount of dissolved oxygen in a sample of river water was found using the process outlined.
 - excess alkaline manganese(II) sulfate, MnSO₄, was added to a 150 cm³ sample of river water
 - the Mn²⁺ ions reacted with the dissolved oxygen forming a precipitate of manganese(IV) oxide hydroxide

$$2Mn^{2+}(aq) + O_2(aq) + 4OH^{-}(aq) \rightarrow 2MnO(OH)_2(s)$$

• the precipitate was then dissolved using excess sulfuric acid, forming $\mathrm{Mn}^{4+}(\mathrm{aq})$ ions

$$MnO(OH)_2(s) + 4H^+(aq) \rightarrow Mn^{4+}(aq) + 3H_2O(l)$$

• excess potassium iodide solution was then added, forming iodine

$$Mn^{4+}(aq) + 2I^{-}(aq) \rightarrow Mn^{2+}(aq) + I_{2}(aq)$$

 the liberated iodine was then titrated with sodium thiosulfate solution, Na₂S₂O₃(aq), of concentration 0.00518 mol dm⁻³

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

- the mean volume of the titre of Na₂S₂O₃(aq) was 34.20 cm³.
- (i) Calculate the concentration of dissolved oxygen in the sample of river water, in g dm⁻³.

(5)



You may use this space to continue your answer to 10(c)(i).

(ii) The concentration of oxygen in water is often expressed in parts per million (ppm), where 1 ppm equals 1 g of solute dissolved in 1×10^6 g of solvent.

Calculate the concentration of the oxygen in the sample of river water in ppm. Assume the density of the river water is $1.00\,\mathrm{g\ cm^{-3}}$.

(1)

(d) Some data is shown for electrode systems involving the Mn³⁺(aq) ion.

Half-cell	Electrode system	E⊕/V
Α	$MnO_2(s) + 4H^+(aq) + e^- \implies Mn^{3+}(aq) + 2H_2O(l)$	+0.95
В	$Mn^{3+}(aq) + e^{-} \rightleftharpoons Mn^{2+}(aq)$	+1.51

Explain why Mn³⁺ ions are unstable in aqueous solution. Include an equation and the type of reaction that occurs.

(4)

TOTAL FOR PAPER = 90 MARKS

(Total for Question 10 = 15 marks)

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Krypton 36 **He**lium 131.3 [222] **Rn** radon 86 20.2 **Ne** argon 18 83.8 Xenon xenon (8) 0 39.9 (18) Αŗ chlorine bromine 35 fluorine 126.9 iodine astatine 35.5 6.62 [210] **Se** selenium **Te** tellurium polonium oxygen 127.6 sulfur 79.0 [209] 32.1 **S** 34 9 N | nitrogen | 7 Bi bismuth 83 hosphorus 121.8 **Sb** antimony **As** arsenic 33 209.0 31.0 **P** 74.9 2 **Ge** germanium Carbon 28.1 **Si** siticon 72.6 118.7 207.2 **Pb** lead 82 **S** # 50 **Al** aluminium **Ga** gallium thallium indium 114.8 10.8 **B** 204.4 69.7 8 Hg mercury 80 Cd Cadmium The Periodic Table of Elements 200.6 Znc zinc 30 48 107.9 197.0 Cu copper 29 **Ag** silver 47 [272] Au gold 79 106.4 **Pd** palladium **Pt** platinum 195.1 **58**.7 **Bi** nickel 28 [271] rhodium iridium 58.9 **Co** cobalt 27 102.9 **Rh** 192.2 [268] **Ru** ruthenium 1.0 **H** hydrogen 1 Os osmium 190.2 101.1 [277] 55.8 **Fe** iron 26 4 8 Re rhenium 75 [98] **Tc**technetium [264] **Bh** bohrium 186.2 43 Mo molybdenum t chromium 24 tungsten 183.8 95.9 [566] 42 9 atomic (proton) number relative atomic mass atomic symbol vanadium 23 [262] **Db**dubnium tantalum niobium 180.9 **>**0.9 92.9 **Nb** titanium 22 zirconium hafnium 178.5 91.2 **Zr** [261] **Rf** 4 9 Sc scandium 21 anthanum yttrium 138.9 88.9 **≻** Ľa* [227] **Ac*** ල Mg magnesium **Be** beryllium **Ca** calcium 20 strontium **Ba** barium 137.3 87.6 **Sr** 40.1 [226] **Ra** 24.3 38 potassium 19 85.5 **Rb** rubidium **CS** caesium **Li** lithium **Na** Sodium 132.9 23.0 39.1 [223] **Fr** 6.9 37

^{*} Actinide series

Elements with atomic numbers 112-116 have been reported

but not fully authenticated

neitnerium damstadtium

hassium

Sg seaborgium

ntherfordium

actinium

radium

francium

6

109

107

^{*} Lanthanide series