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A-level CHEMISTRY

Paper 1 Inorganic and Physical Chemistry

Tuesday 2 June 2020

Afternoon

Time allowed: 2 hours

Materials

For this paper you must have:

- the Periodic Table/Data Booklet, provided as an insert (enclosed)
- a ruler with millimetre measurements
- a scientific calculator, which you are expected to use where appropriate.

Instructions

- Use black ink or black ball-point pen.
- Fill in the boxes at the top of this page.
- Answer all questions.
- You must answer the questions in the spaces provided. Do **not** write outside the box around each page or on blank pages.
- If you need extra space for your answer(s), use the lined pages at the end of this book. Write the question number against your answer(s).
- All working must be shown.
- Do all rough work in this book. Cross through any work you do not want to be marked.

Information

- The marks for questions are shown in brackets.
- The maximum mark for this paper is 105.

For Examiner's Use		
Question	Mark	
1		
2		
3		
4		
5		
6		
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8		
9		
10		
11		
TOTAL		



Answer all questions in the spaces provided.

- 0 1 This question is about enthalpy changes.
- 0 1. 1 Figure 1 shows a Born–Haber cycle for the formation of strontium chloride, SrCl₂

Figure 1

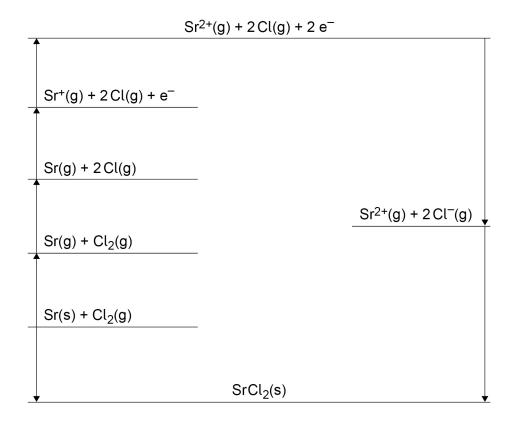


Table 1 shows some thermodynamic data.

Table 1

	Enthalpy change / kJ mol ⁻¹
First ionisation energy of strontium	+548
Second ionisation energy of strontium	+1060
Enthalpy of atomisation of chlorine	+121
Enthalpy of atomisation of strontium	+164
Enthalpy of formation of strontium chloride	-828
Enthalpy of lattice formation of strontium chloride	-2112

	Use the data in Table 1 to ca	alculate a value for the e	lectron affinity of chlorine. [3 mark	(s]
	Ele	ectron affinity	kJ mol ^{_1}	1
0 1.2	Draw a line from each substa	ance to the enthalpy of la	attice formation of that substand	
	Substance		Enthalpy of lattice formation / kJ mol ⁻¹	
	MgCl ₂		-2018	
	MgO		-2493	
	BaCl ₂		-3889	
	Question 1 c	continues on the next p	page	





Table 2 shows the theoretical lattice enthalpy, based on a perfect ionic model, and an experimental value for the enthalpy of lattice formation of silver chloride.

Table 2

	Theoretical	Experimental
Enthalpy of lattice formation / kJ mol ⁻¹	–770	– 905

Table 3 Li ⁺ (g) Na ⁺ (g) K ⁺ (g) Enthalpy of hydration / kJ mol ⁻¹ -519 -406 -322 Explain why the enthalpy of hydration becomes less exothermic from Li ⁺ to K ⁺	Table 3 Li ⁺ (g) Na ⁺ (g) K ⁺ (g) Enthalpy of hydration / kJ mol ⁻¹ -519 -406 -322 Explain why the enthalpy of hydration becomes less exothermic from Li ⁺ to K ⁺	3 State why there is a difference between	en the theoretic	cal and experim	nental value [
Table 3 Li ⁺ (g) Na ⁺ (g) K ⁺ (g) Enthalpy of hydration / kJ mol ⁻¹ -519 -406 -322 Explain why the enthalpy of hydration becomes less exothermic from Li ⁺ to K ⁺	Table 3 Li ⁺ (g) Na ⁺ (g) K ⁺ (g) Enthalpy of hydration / kJ mol ⁻¹ -519 -406 -322 Explain why the enthalpy of hydration becomes less exothermic from Li ⁺ to K ⁺				
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Enthalpy of hydration / kJ mol ⁻¹	Enthalpy of hydration / kJ mol ⁻¹			·	
Enthalpy of hydration / kJ mol ⁻¹	Enthalpy of hydration / kJ mol ⁻¹	[l i ⁺ /a\	No ⁺ /a)	K+(a)
		Enthalpy of hydration / kJ mol ⁻¹			
• • • • • • • • • • • • • • • • • • •		Explain why the enthalpy of hydration	becomes less	exothermic from	m Li⁺ to K⁺ [2



0 1. 5 Calcium bromide dissolves in water.

Table 4 shows some enthalpy data.

Table 4

	Enthalpy change / kJ mol ⁻¹
Enthalpy of solution of calcium bromide	–110
Enthalpy of lattice formation of calcium bromide	-2176
Enthalpy of hydration of calcium ions	-1650

Use the data in **Table 4** to calculate the enthalpy of hydration, in kJ mol⁻¹, of bromide ions.

[3 marks]

Enthalpy of hydration of bromide ions kJ mol⁻¹

10

Turn over for the next question



0 2	This question is about the isotopes of chromium.	
0 2.1	Give the meaning of the term relative atomic mass.	[2 marks]
		[Z IIIdi KS]
0 2.2	A sample of chromium containing the isotopes 50 Cr, 52 Cr and 53 Cr has a relative atomic mass of 52.1	
	The sample contains 86.1% of the ⁵² Cr isotope.	
	Calculate the percentage abundance of each of the other two isotopes.	[4 marks]
		[
	Abundance of ⁵⁰ Cr % Abundance of ⁵³ Cr	%



0 2.3	State, in terms of the numbers of fundamental particles, one similarity and one difference between atoms of ⁵⁰ Cr and ⁵³ Cr
	[2 marks]
	Similarity
	Difference
0 2.4	The sample of chromium is analysed in a time of flight (TOF) mass spectrometer. Give two reasons why it is necessary to ionise the isotopes of chromium before they can be analysed in a TOF mass spectrometer.
	[2 marks]
	1
	2
	Question 2 continues on the next page



0 2.5 A ⁵³Cr⁺ ion travels along a flight tube of length 1.25 m The ion has a constant kinetic energy (KE) of 1.102 × 10⁻¹³ J

$$KE = \frac{mv^2}{2}$$

m = mass of the ion / kg $v = \text{speed of ion / m s}^{-1}$

Calculate the time, in s, for the 53Cr+ ion to travel down the flight tube to reach the detector.

The Avogadro constant, $L = 6.022 \times 10^{23} \text{ mol}^{-1}$

[5 marks]

15

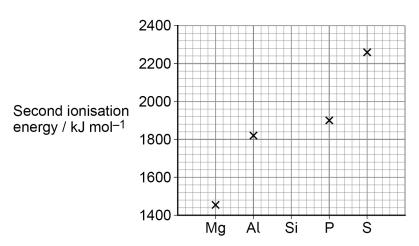
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0 3 This question is about Period 3 elements.

Figure 2 shows the **second** ionisation energies of some elements in Period 3.

Figure 2



0 3 . 1 Draw a cross (x) on **Figure 2** to show the **second** ionisation energy of silicon.

[1 mark]

0 3. 2 Identify the element in Period 3, from sodium to argon, that has the highest **second** ionisation energy.

Give an equation, including state symbols, to show the process that occurs when the **second** ionisation energy of this element is measured.

If you were unable to identify the element you may use the symbol **Q** in your equation. **[2 marks]**

Element _____

Equation

0 3. Explain why the atomic radius decreases across Period 3, from sodium to chlorine.

[2 marks]



0 3.4	Identify the element in Period 3, from sodium to chlorine, that has the highes electronegativity.	t [1 mark]	outside the box
0 3.5	Phosphorus burns in air to form phosphorus(V) oxide. Give an equation for this reaction.	[1 mark]	7

Turn over for the next question

0 4	Propanoic acid (C₂H₅COOH) is a weak acid.
	The acid dissociation constant (K_a) for propanoic acid is 1.35 × 10 ⁻⁵ mol dm ⁻³ at 25 °C
0 4.1	State the meaning of the term weak acid. [1 mark]
0 4.2	Give an expression for the acid dissociation constant for propanoic acid. [1 mark]
	\mathcal{K}_{a}
0 4.3	A student dilutes 25.0 cm 3 of 0.500 mol dm $^{-3}$ propanoic acid by adding water until the total volume is 100.0 cm 3
	Calculate the pH of this diluted solution of propanoic acid.
	Give your answer to 2 decimal places. [4 marks]
	pH



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0	4		4
		-	

A buffer solution with a pH of 4.50 is made by dissolving x g of sodium propanoate (C_2H_5COONa) in a solution of propanoic acid. The final volume of buffer solution is 500 cm³ and the final concentration of the propanoic acid is 0.250 mol dm⁻³

Calculate x in g

For propanoic acid, $K_a = 1.35 \times 10^{-5} \text{ mol dm}^{-3}$

[6 marks]

ç g

12



0 5	Some reactions of the $[Al(H_2O)_6]^{3+}(aq)$ ion are shown.
	Colourless solution containing complex ion A Na ₄ EDTA(aq) [Al(H ₂ O) ₆] ³⁺ (aq) Na ₂ CO ₃ (aq) White precipitate NaOH(aq) White precipitate that reacts to form a colourless solution containing complex ion C
0 5.1	Give the formula of the white precipitate B .
	State one other observation when $Na_2CO_3(aq)$ is added to a solution containing $[Al(H_2O)_6]^{3+}(aq)$ ions.
	Give an equation for this reaction. [3 marks]
	Formula of B
	Observation
	Equation
0 5.2	Give the formula of the complex ion C .
	State one condition needed for the formation of C from $[Al(H_2O)_6]^{3+}(aq)$ and NaOH(aq).
	Give an equation for this reaction. [3 marks]
	Formula of C
	Condition
	Equation



Do not write
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box

0 5 . 3	Deduce the formula of the complex ion A . [1 mark]	Do out
0 5.4	Explain, with the use of an equation, why a solution containing $[Al(H_2O)_6]^{3+}$ has a pH <7	
	Equation	
	Explanation_	
		1

Turn over for the next question



0 6

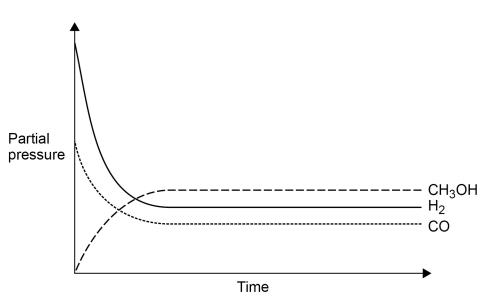
Methanol can be manufactured in a reversible reaction as shown.

$$CO(g) + 2H_2(g) \rightleftharpoons CH_3OH(g)$$

 $\Delta H^{\circ} = -91 \text{ kJ mol}^{-1}$

Figure 3 shows how the partial pressures change with time at a constant temperature.

Figure 3



0 6 . 1

Draw a cross (x) on the appropriate axis of **Figure 3** when the mixture reaches equilibrium.

[1 mark]

0 6 . 2

A 0.230 mol sample of carbon monoxide is mixed with hydrogen in a 1:2 mol ratio and allowed to reach equilibrium in a sealed flask at temperature T.

At equilibrium the mixture contains 0.120 mol of carbon monoxide.

The total pressure of this mixture is 1.04×10^4 kPa

Calculate the partial pressure, in kPa, of hydrogen in the equilibrium mixture.

[4 marks]

Partial pressure of hydrogen

kPa



0 6.3	Give an expression for the equilibrium constant (K_p) for this reaction.	Do not v outside box
	State the units. [2 marks]	
	\mathcal{K}_p	
	Units	
0 6.4	Some more carbon monoxide is added to the mixture in Question 06.2 . The new mixture is allowed to reach equilibrium at temperature T .	
	State the effect, if any, on the partial pressure of methanol and on the value of \mathcal{K}_p [2 marks]	
	Effect on partial pressure of methanol	
	Effect on value of K _p	
0 6.5	State the effect, if any, of the addition of a catalyst on the value of K_p for this equilibrium. Explain your answer. [2 marks]	
	Effect on value of K _p	
	Explanation	
		11
		<u>'''</u>

Turn over for the next question



0 7	The melting point of XeF ₄ is higher than the melting point of PF ₃	
	Explain why the melting points of these two compounds are different.	
	In your answer you should give the shape of each molecule, explain why each molecule has that shape and how the shape influences the forces that affect the melting point.	
	[6 mar	ˈks]



	_
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0 8

A student does an experiment to determine the percentage by mass of sodium chlorate(I), NaClO, in a sample of bleach solution.

Method:

- Dilute a 10.0 cm³ sample of bleach solution to 100 cm³ with distilled water.
- Transfer 25.0 cm³ of the diluted bleach solution to a conical flask and acidify using sulfuric acid.
- Add excess potassium iodide to the conical flask to form a brown solution containing l₂(aq).
- Add 0.100 mol dm⁻³ sodium thiosulfate solution (Na₂S₂O₃) to the conical flask from a burette until the brown solution containing I₂(aq) becomes a colourless solution containing I⁻(aq).

The student uses 33.50 cm³ of sodium thiosulfate solution.

The density of the original bleach solution is 1.20 g cm⁻³

The equations for the reactions in this experiment are

$$ClO^{-}(aq) + 2H^{+}(aq) + 2I^{-}(aq) \rightarrow Cl^{-}(aq) + H_2O(I) + I_2(aq)$$

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

0 8 . 1

Use all the information given to calculate the percentage by mass of NaClO in the original bleach solution.

Give your answer to 3 significant figures.

[7 marks]

Percentage by mass

0 8 . 2	The total uncertainty from two readings and an end point error in using a burette is $\pm \ 0.15 \ \text{cm}^3$	Do not write outside the box
	What is the total percentage uncertainty in using the burette in this experiment?	
	Tick (✓) one box.	
	0.45%	
	0.90%	
	1.34%	8

Turn over for the next question

0 9	This question is about sodium halides.	
0 9 . 1	State what is observed when silver nitrate solution is added to sodium fluoric solution.	
		[1 mark]
0 9.2	State one observation when solid sodium chloride reacts with concentrated sulfuric acid.	
	Give an equation for the reaction.	
	State the role of the chloride ions in the reaction.	[3 marks]
	Observation	
	Equation	
	Role	
0 9 . 3	Give an equation for the redox reaction between solid sodium bromide and concentrated sulfuric acid.	
	Explain, using oxidation states, why this is a redox reaction.	[3 marks]
	Equation	
	Explanation	
0 9 . 4	State what is observed when aguacus oblering is added to addium bromide	colution
0 9 . 4	State what is observed when aqueous chlorine is added to sodium bromide	Solution.
	Give an ionic equation for the reaction.	[2 marks]
	Observation	
	Ionic equation	



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Methanol is formed when carbon dioxide and hydrogen react.

$$CO_2(g) + 3H_2(g) \rightleftharpoons CH_3OH(g) + H_2O(g)$$

Table 5 contains enthalpy of formation and entropy data for these substances.

Table 5

	CO ₂ (g)	H ₂ (g)	CH₃OH(g)	H₂O(g)
Δ _f H / kJ mol ⁻¹	-394	0	-201	-242
S / J K ⁻¹ mol ⁻¹	214	131	238	189

1 0. 1 Use the equation and the data in **Table 5** to calculate the Gibbs free-energy change (Δ*G*), in kJ mol⁻¹, for this reaction at 890 K

[6 marks]

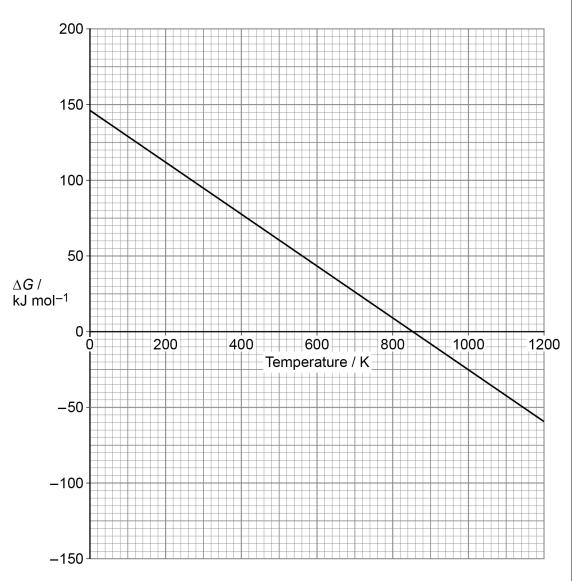
 ΔG kJ mol⁻¹



Figure 4 shows how the Gibbs free-energy change varies with temperature in a different gas phase reaction.

The straight line graph for this gas phase reaction has been extrapolated to zero Kelvin.

Figure 4





re 4 to calculate), in J K ⁻¹ mol ⁻¹ , for	nt from the graph in Figure the entropy change (ΔS),	Use the values of the intercept an the enthalpy change (ΔH), in kJ m this reaction.	0 . 2
[4 marks]			
kJ mol ⁻¹	ΔH		
J K ⁻¹ mol ⁻¹	ΔS		
	ibility of the reaction.	State what Figure 4 shows about	0 . 3

Turn over ▶

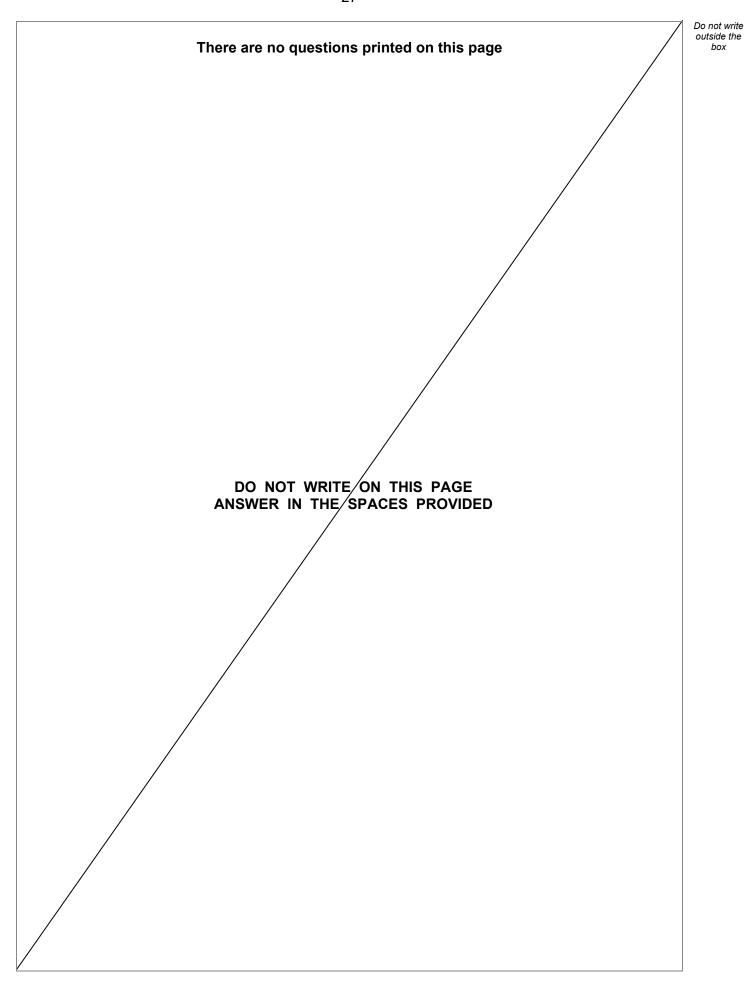
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1 1	This question is about a glucose–oxygen fuel cell.		outside t
	When the cell operates, the glucose ($C_6H_{12}O_6$) molecules react with water a negative electrode to form carbon dioxide and hydrogen ions.	t the	
	Oxygen gas reacts with hydrogen ions to form water at the positive electrode	Э.	
1 1.1	Deduce the half-equation for the reaction at the negative electrode.	[1 mark]	
1 1.2	Deduce the half-equation for the reaction at the positive electrode.	[1 mark]	
1 1.3	Give the equation for the overall reaction that occurs in the Glucose–oxygen fuel cell.	[1 mark]	
1 1.4	The negative electrode is made of carbon and the positive electrode is made platinum. Give the conventional representation for the glucose–oxygen fuel cell.	e of [2 marks]	
1 1.5	State what must be done to maintain the EMF of this fuel cell when in use.	[1 mark]	6
	END OF QUESTIONS		







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