

Please write clearly in block capitals.

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I declare this is my own work.

# A-level CHEMISTRY

## Paper 1 Inorganic and Physical Chemistry

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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.

For Examiner's Use	
Question	Mark
1	
2	
3	
4	
5	
6	
7	
8	
9	
<b>TOTAL</b>	

### Information

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



J U N 2 1 7 4 0 5 1 0 1

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**7405/1**

Answer **all** questions in the spaces provided.

**0 | 1**

This question is about enthalpy changes for calcium chloride and magnesium chloride.

**0 | 1 . 1**

State the meaning of the term enthalpy change.

[1 mark]

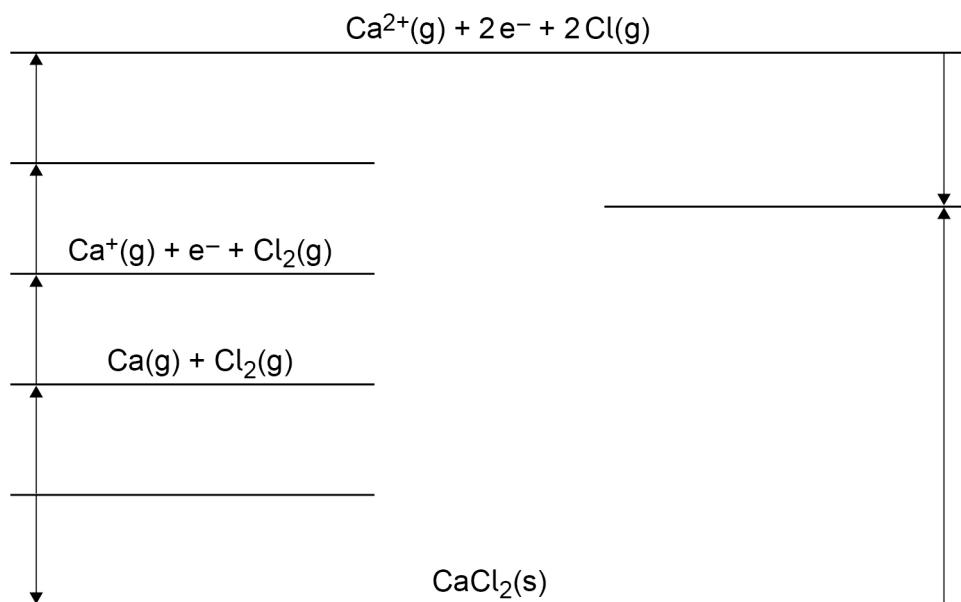
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**Figure 1** shows an incomplete Born–Haber cycle for the formation of calcium chloride.

**Figure 1**



**0 | 1 . 2**

Complete **Figure 1** by writing the formulas, including state symbols, of the appropriate species on each of the three blank lines.

[3 marks]



0 2

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**0 | 1 . 3** **Table 1** shows some enthalpy data.

Do not write outside the box

**Table 1**

	<b>Enthalpy change / kJ mol<sup>-1</sup></b>
Enthalpy of formation of calcium chloride	−795
Enthalpy of atomisation of calcium	+193
First ionisation energy of calcium	+590
Second ionisation energy of calcium	+1150
Enthalpy of atomisation of chlorine	+121
Electron affinity of chlorine	−364

Use **Figure 1** and the data in **Table 1** to calculate a value for the enthalpy of lattice dissociation of calcium chloride.

**[2 marks]**

Enthalpy of lattice dissociation \_\_\_\_\_ kJ mol<sup>-1</sup>

**Question 1 continues on the next page**

**Turn over ►**



0 3

IB/M/Jun21/7405/1

**0 1 . 4** Magnesium chloride dissolves in water.

Give an equation, including state symbols, to represent the process that occurs when the enthalpy of solution of magnesium chloride is measured.

[1 mark]

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**0 1 . 5** **Table 2** shows some enthalpy data.

**Table 2**

	Enthalpy change / kJ mol <sup>-1</sup>
Enthalpy of lattice dissociation of MgCl <sub>2</sub>	+2493
Enthalpy of hydration of Mg <sup>2+</sup> (g)	-1920
Enthalpy of hydration of Cl <sup>-</sup> (g)	-364

Use your answer to Question **01.4** and the data in **Table 2** to calculate a value for the enthalpy of solution of magnesium chloride.

[2 marks]

Enthalpy of solution \_\_\_\_\_ kJ mol<sup>-1</sup>

**0 1 . 6** The enthalpy of hydration of Ca<sup>2+</sup>(g) is -1650 kJ mol<sup>-1</sup>

Suggest why this value is less exothermic than that of Mg<sup>2+</sup>(g)

[2 marks]

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11



**0 2**

This question is about atomic structure.

**0 2 . 1**

Define the mass number of an atom.

**[1 mark]**


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**0 2 . 2**

Complete **Table 3** to show the numbers of neutrons and electrons in the species shown.

**[2 marks]****Table 3**

	<b>Number of protons</b>	<b>Number of neutrons</b>	<b>Number of electrons</b>
$^{46}\text{Ti}$	22		
$^{49}\text{Ti}^{2+}$	22		

**0 2 . 3**

A sample of titanium contains four isotopes,  $^{46}\text{Ti}$ ,  $^{47}\text{Ti}$ ,  $^{48}\text{Ti}$  and  $^{49}\text{Ti}$

This sample has a relative atomic mass of 47.8

In this sample the ratio of abundance of isotopes  $^{46}\text{Ti}$ ,  $^{47}\text{Ti}$  and  $^{49}\text{Ti}$  is 2:2:1

Calculate the percentage abundance of  $^{46}\text{Ti}$  in this sample.

**[3 marks]**

Abundance of  $^{46}\text{Ti}$  \_\_\_\_\_ %

**6****Turn over ►**

0 5

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**0 3**

This question is about elements in Period 3 and their compounds.

**0 3 . 1**

When a piece of sodium is added to 200 cm<sup>3</sup> of water in a large beaker a vigorous reaction occurs. The temperature of the water increases by 25 °C

Give an equation, including state symbols, for the reaction of sodium with water.

Suggest why it is dangerous to react a similar piece of sodium with 10 cm<sup>3</sup> of water in a boiling tube.

**[2 marks]**

Equation

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Why it is dangerous

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**0 3 . 2**

Give an equation for the reaction of phosphorus(V) oxide with water.

Suggest a pH for the solution formed.

**[2 marks]**

Equation

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pH

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**0 3 . 3**

Explain, in terms of crystal structure and bonding, why silicon(IV) oxide has a higher melting point than phosphorus(V) oxide.

**[4 marks]**

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- 0 3 . 4** An element in Period 3 forms an oxide that is insoluble in water.  
This oxide reacts with sulfuric acid and with aqueous potassium hydroxide.

Give the formula for this oxide.

Give an equation for the reaction of this oxide with sulfuric acid.

**[2 marks]**

Formula \_\_\_\_\_

Equation \_\_\_\_\_

- 0 3 . 5** Give the formula of a hydroxide of an element in Period 3 used in medicine.

**[1 mark]**

- 0 3 . 6** Identify the element in Period 3, from sodium to chlorine, that has the largest atomic radius.

**[1 mark]**

12

**Turn over for the next question**

**Turn over ►**



0 7

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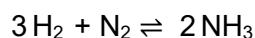
0 | 4

This question is about iron and its ions.

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0 | 4

1 Discuss the role of iron as a heterogeneous catalyst in the Haber process.



Your answer should include:

- the meaning of the term heterogeneous catalyst
  - how iron acts as a heterogeneous catalyst
  - the factors that affect the efficiency and lifetime of the catalyst.

[6 marks]



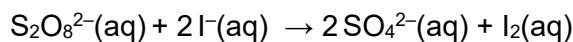
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0 9

**0 4 . 2** Fe<sup>2+</sup> ions catalyse the reaction between peroxodisulfate(VI) ions and iodide ions in aqueous solution.



Explain why this reaction is slow before the catalyst is added.  
Give **two** equations to show how Fe<sup>2+</sup> ions catalyse this reaction.

**[4 marks]**

Why reaction is slow before catalyst added \_\_\_\_\_

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Equation 1

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Equation 2

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**0 4 . 3** Give a reason why Zn<sup>2+</sup> ions do **not** catalyse the reaction in Question **04.2**.

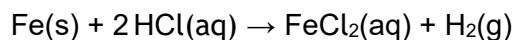
**[1 mark]**

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**0 4 . 4** Iron reacts with dilute hydrochloric acid to form iron(II) chloride and hydrogen.



A 0.998 g sample of pure iron is added to 30.0 cm<sup>3</sup> of 1.00 mol dm<sup>-3</sup> hydrochloric acid.

One of these reagents is in excess and the other reagent limits the amount of hydrogen produced in the reaction.

Calculate the maximum volume, in m<sup>3</sup>, of hydrogen gas produced at 30 °C and 100 kPa.

Give your answer to 3 significant figures.

In your answer you should identify the limiting reagent in the reaction.

The gas constant,  $R = 8.31 \text{ J K}^{-1} \text{ mol}^{-1}$

**[6 marks]**

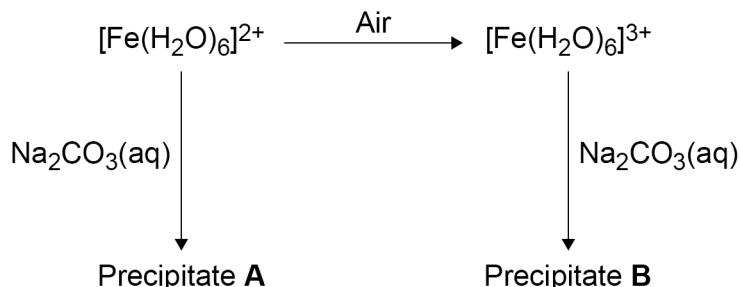
Volume of hydrogen \_\_\_\_\_ m<sup>3</sup>

Turn over ►



**Figure 2** shows some reactions of iron ions in aqueous solution.

**Figure 2**



- 0 4 . 5** Identify **A** and state its colour.

**[2 marks]**

Identity \_\_\_\_\_

Colour \_\_\_\_\_

- 0 4 . 6** Give the formula of **B** and state its colour.

Give an ionic equation for the reaction of  $[\text{Fe}(\text{H}_2\text{O})_6]^{3+}$  with aqueous  $\text{Na}_2\text{CO}_3$  to form **B**.

**[3 marks]**

Formula \_\_\_\_\_

Colour \_\_\_\_\_

Ionic equation \_\_\_\_\_



*Do not write outside the box*

**0 4 . 7** Explain why an aqueous solution containing  $[\text{Fe}(\text{H}_2\text{O})_6]^{3+}$  ions has a lower pH than an aqueous solution containing  $[\text{Fe}(\text{H}_2\text{O})_6]^{2+}$  ions.

[3 marks]

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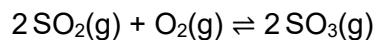
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**0 5**

This question is about the equilibrium

**0 5 . 1**

State and explain the effect, if any, of a decrease in overall pressure on the equilibrium yield of  $\text{SO}_3$

**[3 marks]**

Effect \_\_\_\_\_

Explanation \_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

**0 5 . 2**

A 0.460 mol sample of  $\text{SO}_2$  is mixed with a 0.250 mol sample of  $\text{O}_2$  in a sealed container at a constant temperature.

When equilibrium is reached at a pressure of 215 kPa, the mixture contains 0.180 mol of  $\text{SO}_3$

Calculate the partial pressure, in kPa, of  $\text{SO}_2$  in this equilibrium mixture.

**[4 marks]**

Partial pressure of  $\text{SO}_2$  \_\_\_\_\_ kPa



**Question 5 continues on the next page**

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1 5

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**0 5 . 3** A different mixture of SO<sub>2</sub> and O<sub>2</sub> reaches equilibrium at a different temperature.

**Table 4** shows the partial pressures of the gases at equilibrium.

**Table 4**

Gas	Partial pressure / kPa
SO <sub>2</sub>	$1.67 \times 10^2$
O <sub>2</sub>	$1.02 \times 10^2$
SO <sub>3</sub>	$1.85 \times 10^2$

Give an expression for the equilibrium constant ( $K_p$ ) for this reaction.

Calculate the value of the equilibrium constant for this reaction and give its units.

**[3 marks]**

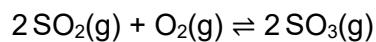
$K_p$

$K_p$  \_\_\_\_\_

Units \_\_\_\_\_



- 0 | 5 . 4** What is the effect on the value of  $K_p$  if the pressure of this equilibrium mixture is increased at a constant temperature?



[1 mark]

Tick ( $\checkmark$ ) **one** box.

The value of  $K_p$

increases.

stays the same.

decreases.

**11**

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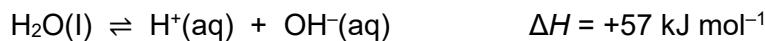
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**0 6**

This question is about pH.

Pure water dissociates slightly.



$$\text{The equilibrium constant, } K_c = \frac{[\text{H}^+][\text{OH}^-]}{[\text{H}_2\text{O}]}$$

$$\text{The ionic product of water, } K_w = [\text{H}^+][\text{OH}^-]$$

**0 6 . 1** Explain why  $[\text{H}_2\text{O}]$  is not shown in the  $K_w$  expression.

[1 mark]

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**Table 5** shows how  $K_w$  varies with temperature.

**Table 5**

Temperature / °C	$K_w / \text{mol}^2 \text{ dm}^{-6}$
10	$2.93 \times 10^{-15}$
20	$6.81 \times 10^{-15}$
25	$1.00 \times 10^{-14}$
30	$1.47 \times 10^{-14}$
50	$5.48 \times 10^{-14}$

**0 6 . 2** Explain why the value of  $K_w$  increases as the temperature increases.

[2 marks]

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1 8

**0 6 . 3** Give the expression for pH.

Calculate the pH of pure water at 50 °C  
Give your answer to 2 decimal places.

Explain why water is neutral at 50 °C

**[4 marks]**

Expression \_\_\_\_\_

Calculation

pH \_\_\_\_\_

Explanation \_\_\_\_\_  
\_\_\_\_\_

**Question 6 continues on the next page**

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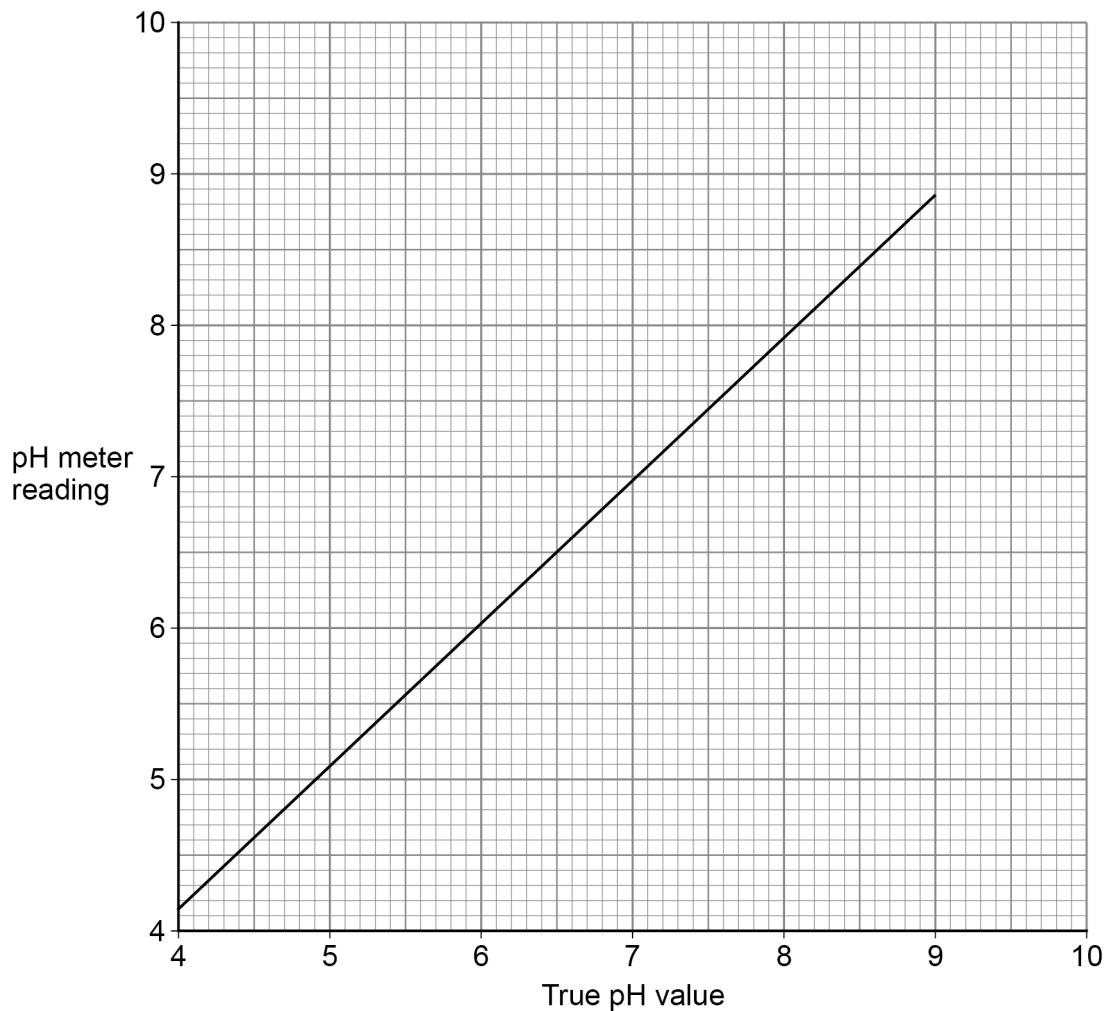


A pH meter is calibrated using a calibration graph.

To create the calibration, the pH meter is used to measure the pH of separate solutions, each with a known, accurate pH.

**Figure 3** shows the calibration graph.

**Figure 3**



- 0 6 . 4** Use **Figure 3** to give the true pH value when the pH meter reading is 5.6

[1 mark]

- 
- 0 6 . 5** Suggest why the pH probe is washed with distilled water between each of the calibration measurements.

[1 mark]

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**0 6 . 6**

The calibrated pH meter is used to monitor the pH during a titration of hydrochloric acid with sodium hydroxide.

Explain why the volume of sodium hydroxide solution added between each pH measurement is smaller as the end point of the titration is approached.

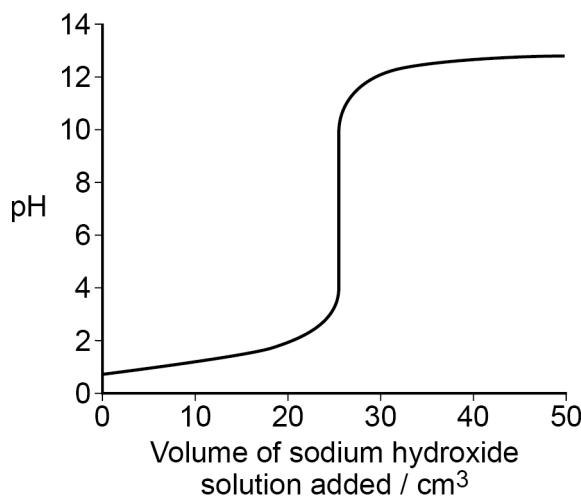
[1 mark]

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**Figure 4** shows the pH curve for a titration of hydrochloric acid with sodium hydroxide solution.

**Figure 4**

**Table 6** shows data about some indicators.

**Table 6**

Indicator	pH range	Colour at low pH	Colour at high pH
Bromocresol green	3.8 – 5.4	yellow	blue
Phenol red	6.8 – 8.4	yellow	red
Thymolphthalein	9.3 – 10.5	colourless	blue

The student plans to do the titration again using one of the indicators in **Table 6** to determine the end point.

**0 6 . 7**

State why all three of the indicators in **Table 6** are suitable for this titration.

[1 mark]

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**Turn over ►**

**0 6 . 8**

36.25 cm<sup>3</sup> of 0.200 mol dm<sup>-3</sup> sodium hydroxide solution are added to 25.00 cm<sup>3</sup> of 0.150 mol dm<sup>-3</sup> hydrochloric acid.

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Calculate the pH of the final solution at 25 °C

$$K_w = 1.00 \times 10^{-14} \text{ mol}^2 \text{ dm}^{-6} \text{ at } 25^\circ\text{C}$$

**[5 marks]**

pH \_\_\_\_\_

**16**



2 2

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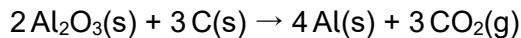


2 3

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**0 7**

This question is about thermodynamics.  
Consider the reaction shown.



**Table 7** shows some thermodynamic data.

**Table 7**

Substance	$\text{Al}_2\text{O}_3(\text{s})$	$\text{Al}(\text{s})$	$\text{C}(\text{s})$	$\text{CO}_2(\text{g})$
$\Delta_f H^\ominus / \text{kJ mol}^{-1}$	-1669	0	0	-394
$S^\ominus / \text{J K}^{-1} \text{mol}^{-1}$	51	28	6	214

**0 7 . 1**

Explain why the standard entropy value for carbon dioxide is greater than that for carbon.

[1 mark]

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**0 7 . 2**

State the temperature at which the standard entropy of aluminium is  $0 \text{ J K}^{-1} \text{ mol}^{-1}$

[1 mark]

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**0 7 . 3**

Use the equation and the data in **Table 7** to calculate the minimum temperature, in K, at which this reaction becomes feasible.

**[7 marks]**

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Minimum temperature \_\_\_\_\_ K

**9****Turn over ►**

2 5

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**0 8**

This question is about electrode potentials and electrochemical cells.

**0 8 . 1**

State the meaning of the term electrochemical series.

**[1 mark]**


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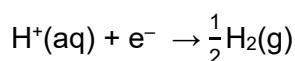
**Table 8** shows some electrode potentials.

**Table 8**

	$E^\ominus / V$
$[\text{Fe}(\text{H}_2\text{O})_6]^{2+}(\text{aq}) + 2 \text{e}^- \rightarrow \text{Fe(s)} + 6 \text{H}_2\text{O(l)}$	-0.44
$\text{H}^+(\text{aq}) + \text{e}^- \rightarrow \frac{1}{2} \text{H}_2(\text{g})$	0.00
$[\text{Co}(\text{NH}_3)_6]^{3+}(\text{aq}) + \text{e}^- \rightarrow [\text{Co}(\text{NH}_3)_6]^{2+}(\text{aq})$	+0.11
$[\text{Fe}(\text{H}_2\text{O})_6]^{3+}(\text{aq}) + \text{e}^- \rightarrow [\text{Fe}(\text{H}_2\text{O})_6]^{2+}(\text{aq})$	+0.77
$\text{VO}_2^+(\text{aq}) + 2 \text{H}^+(\text{aq}) + \text{e}^- \rightarrow \text{VO}^{2+}(\text{aq}) + \text{H}_2\text{O(l)}$	+1.00
$[\text{Co}(\text{H}_2\text{O})_6]^{3+}(\text{aq}) + \text{e}^- \rightarrow [\text{Co}(\text{H}_2\text{O})_6]^{2+}(\text{aq})$	+1.81

**0 8 . 2**

State **two** conditions needed for the following half-cell to have  $E^\ominus = 0.00 \text{ V}$

**[1 mark]**


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**0 8 . 3**

Identify the weakest reducing agent in **Table 8**.

**[1 mark]**


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2 6

- 0 8 . 4** Use half-equations from **Table 8** to deduce an equation for the reduction of  $\text{VO}_2^+$  to form  $\text{VO}^{2+}$  in aqueous solution by iron.

[2 marks]

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- 0 8 . 5** Use data from **Table 8** to explain why  $[\text{Co}(\text{H}_2\text{O})_6]^{3+}(\text{aq})$  will undergo a redox reaction with  $[\text{Fe}(\text{H}_2\text{O})_6]^{2+}(\text{aq})$

Give an equation for this reaction.

[2 marks]

Explanation

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Equation

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- 0 8 . 6** Suggest why the **two** cobalt(III) complex ions in **Table 8** have different electrode potentials.

[1 mark]

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8

**Turn over for the next question**

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**0 9**

This question is about the development of lithium cells.

The value of  $E^\circ$  for lithium suggests that a lithium cell could have a large EMF.

**Table 9** shows some electrode potential data.

**Table 9**

	$E^\circ / \text{V}$
$\text{Li}^+(\text{aq}) + \text{e}^- \rightarrow \text{Li}(\text{s})$	-3.04
$2\text{H}_2\text{O}(\text{l}) + 2\text{e}^- \rightarrow \text{H}_2(\text{g}) + 2\text{OH}^-(\text{aq})$	-0.83
$\frac{1}{2}\text{I}_2(\text{s}) + \text{e}^- \rightarrow \text{I}^-(\text{aq})$	+0.54

**0 9 . 1**

Use data in **Table 9** to explain why an aqueous electrolyte is **not** used for a lithium cell.

[2 marks]

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**0 9 . 2**

In the 1970s lithium-iodine cells became a common power source for heart pacemakers. Lithium iodide is the final product of the cell reaction.

Use the data in **Table 9** to calculate the cell EMF of a standard lithium-iodine cell.

[1 mark]

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**0 9 . 3**

An EMF value for a commercial lithium-iodine cell is 2.80 V

Suggest why this value is different from the value calculated in Question **09.2**.

[1 mark]

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**0 9 . 4** In some lithium cells, lithium perchlorate ( $\text{LiClO}_4$ ) is used as the electrolyte.

Deduce the oxidation state of chlorine in  $\text{LiClO}_4$

[1 mark]

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In other lithium cells, lithium cobalt oxide electrodes **and** lithium electrodes are used.

**0 9 . 5** Give an equation for the reaction that occurs at the positive lithium cobalt oxide electrode.

[1 mark]

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**0 9 . 6** Give an equation for the reaction that occurs at the negative lithium electrode.

[1 mark]

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**7**

**END OF QUESTIONS**



2 9

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