

A

**AQA** 

**Surname** \_\_\_\_\_

**Other Names** \_\_\_\_\_

**Centre Number** \_\_\_\_\_

**Candidate Number** \_\_\_\_\_

**Candidate Signature** \_\_\_\_\_

**I declare this is my own work.**

**A-level**

**CHEMISTRY**

**Paper 1 Inorganic and Physical Chemistry**

**7405/1**

**Time allowed: 2 hours**

**At the top of the page, write your surname and other names, your centre number, your candidate number and add your signature.**

**[Turn over]**



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

**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.**
- **Answer ALL questions.**
- **You must answer the questions in the spaces provided. Do NOT write 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.**

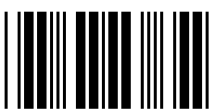


- **Do all rough work in this book. Cross through any work you do not want to be marked.**

## **INFORMATION**

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

**DO NOT TURN OVER UNTIL TOLD TO DO SO**



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**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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**[Turn over]**

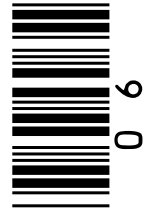


**FIGURE 1, on the opposite page, shows an incomplete Born–Haber cycle for the formation of calcium chloride.**

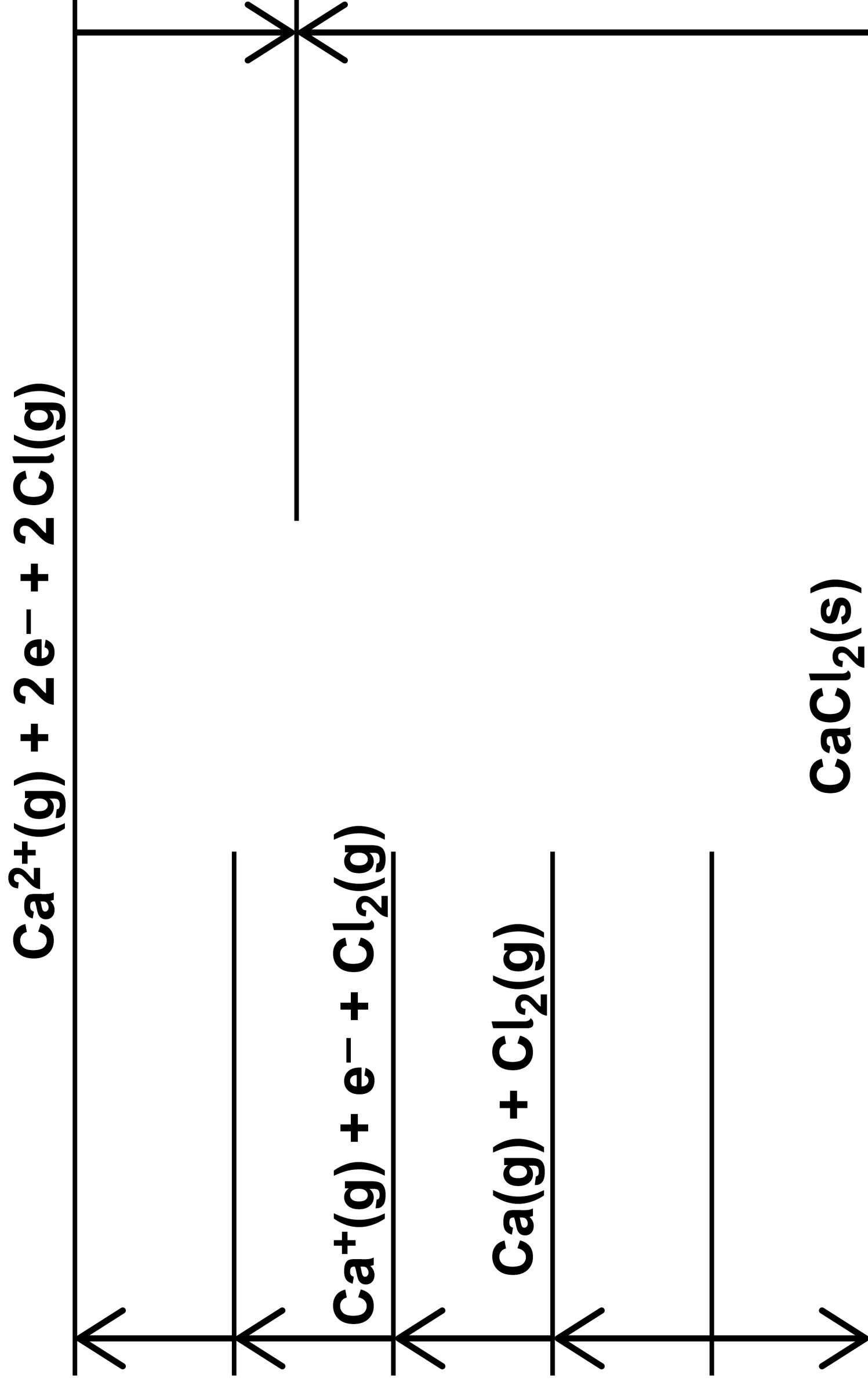
**0 1 . 2**

**Complete FIGURE 1 by writing the formulas, including state symbols, of the appropriate species on each of the three blank lines. [3 marks]**

**6**



**FIGURE 1**



07

**[Turn over]**

**TABLE 1 shows some enthalpy data.**

**TABLE 1**

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

**∞**





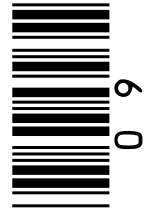
**Use FIGURE 1, on page 7, and the data in TABLE 1 to calculate a value for the enthalpy of lattice dissociation of calcium chloride. [2 marks]**

**9**

**Enthalpy of lattice dissociation**

**\_\_\_\_\_ kJ mol<sup>-1</sup>**

**[Turn over]**



**01.4**

**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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**[Turn over]**



**01.5****TABLE 2 shows some enthalpy data.****TABLE 2**

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



**Use your answer to Question 01.4, on page 10, 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>**

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**[Turn over]**



0	1	.	6
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The enthalpy of hydration of  $\text{Ca}^{2+}(\text{g})$  is  $-1650 \text{ kJ mol}^{-1}$

Suggest why this value is less exothermic than that of  $\text{Mg}^{2+}(\text{g})$  [2 marks]

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11



02

**This question is about atomic structure.**

02.1

**Define the mass number of an atom.**

**[1 mark]**

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**[Turn over]**



0 2 . 2

Complete TABLE 3 to show the numbers of neutrons and electrons in the species shown. [2 marks]

TABLE 3

	Number of protons	Number of neutrons	Number of electrons
$^{46}\text{Ti}$	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  $^{46}\text{Ti}$  \_\_\_\_\_ %

[Turn over]

6



**03**

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

**03.1**

**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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---

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**[Turn over]**



0	3	.	2
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**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** \_\_\_\_\_

0	3	.	3
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**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**

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0	3	.	5
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**Give the formula of a hydroxide of an element in Period 3 used in medicine. [1 mark]**

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0	3	.	6
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**Identify the element in Period 3, from sodium to chlorine, that has the largest atomic radius. [1 mark]**

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**[Turn over]**

12



0	4
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**This question is about iron and its ions.**

0	4	.	1
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**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]**













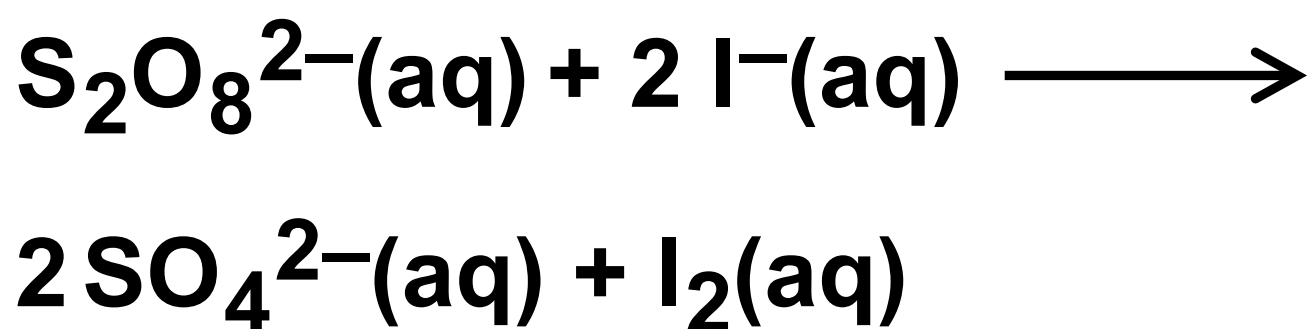
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**[Turn over]**



**04.2**

**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  $\text{Zn}^{2+}$  ions do NOT catalyse the reaction in Question 04.2. [1 mark]**

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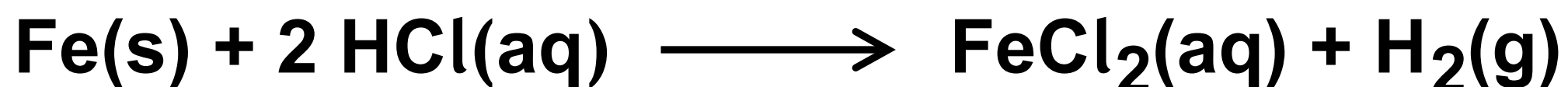
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**[Turn over]**



**04.4**

**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

$\text{m}^3$

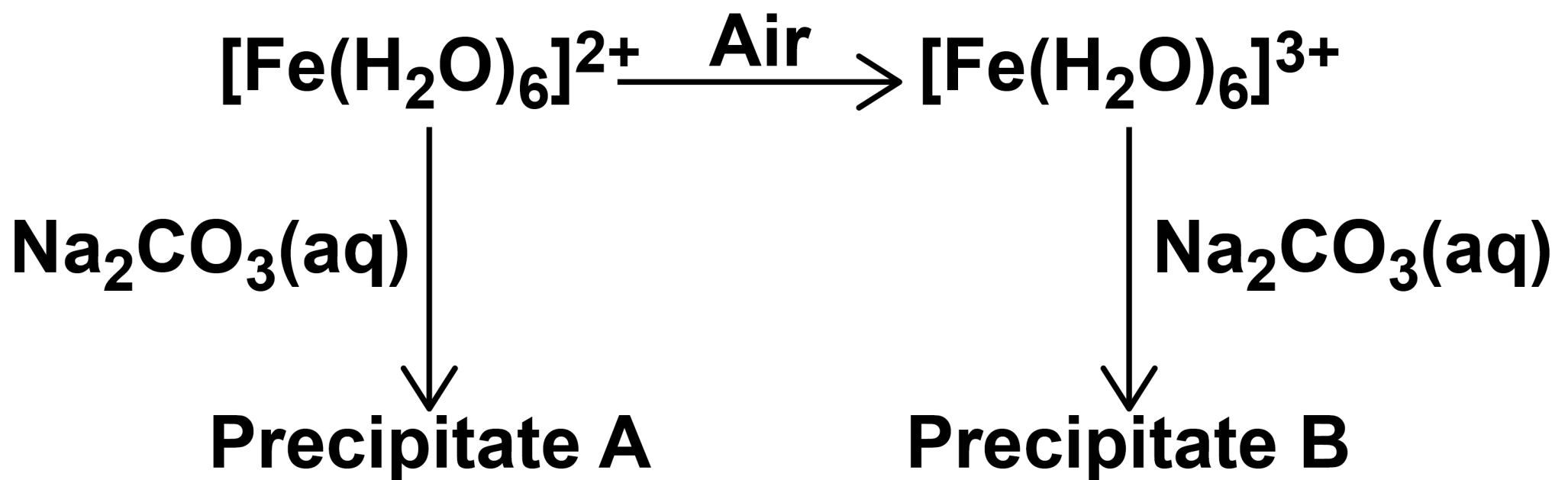


3 3

[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
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**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**

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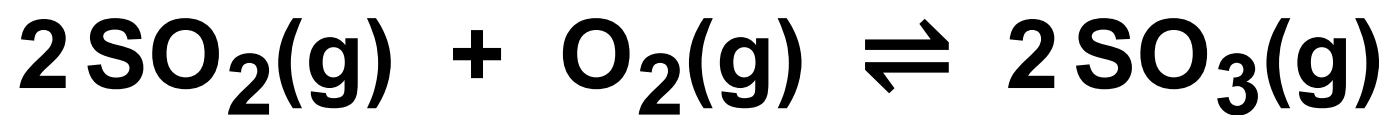
**[Turn over]**





0	5
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**This question is about the equilibrium**



0	5	.	1
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**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** \_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

**[Turn over]**



**05.2**

**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 SO<sub>2</sub>**

**kPa**

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**[Turn over]**



**05.3**

A different mixture of  $\text{SO}_2$  and  $\text{O}_2$  reaches equilibrium at a different temperature.

TABLE 4 shows the partial pressures of the gases at equilibrium.

TABLE 4

Gas	Partial pressure / kPa
$\text{SO}_2$	$1.67 \times 10^2$
$\text{O}_2$	$1.02 \times 10^2$
$\text{SO}_3$	$1.85 \times 10^2$

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]



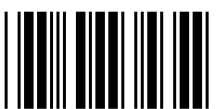


$K_p$

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

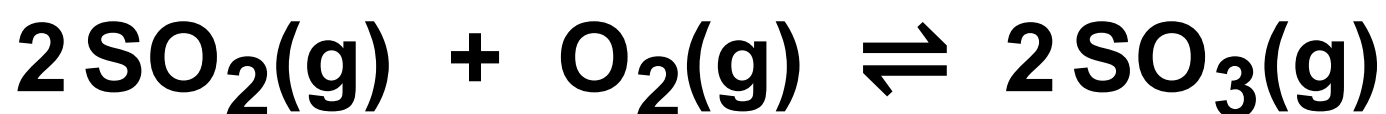
Units \_\_\_\_\_

[Turn over]



0	5	.	4
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**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 (✓) ONE box.**

**The value of  $K_p$**

**increases.**

**stays the same.**

**decreases.**

<b>11</b>



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**[Turn over]**



06

**This question is about pH.**

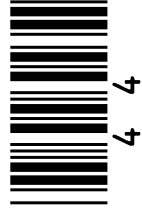
**Pure water dissociates slightly.**



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

**44**

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



06.1

**Explain why  $[H_2O]$  is not shown in the  $K_w$  expression.**

**[1 mark]**

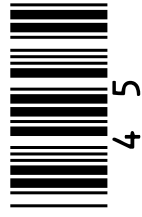
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**45**

**[Turn over]**



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

**TABLE 5**

<b>Temperature / °C</b>	<b><math>K_w</math> / mol<sup>2</sup> dm<sup>-6</sup></b>
<b>10</b>	<b><math>2.93 \times 10^{-15}</math></b>
<b>20</b>	<b><math>6.81 \times 10^{-15}</math></b>
<b>25</b>	<b><math>1.00 \times 10^{-14}</math></b>
<b>30</b>	<b><math>1.47 \times 10^{-14}</math></b>
<b>50</b>	<b><math>5.48 \times 10^{-14}</math></b>



**06.2**

**Explain why the value of  $K_w$  increases as the temperature increases. [2 marks]**

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**[Turn over]**



**06.3****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** \_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

**[Turn over]**



**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, on the opposite page, shows the calibration graph.**

**0 6 . 4**

**Use FIGURE 3 to give the true pH value when the pH meter reading is 5.6 [1 mark]**

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**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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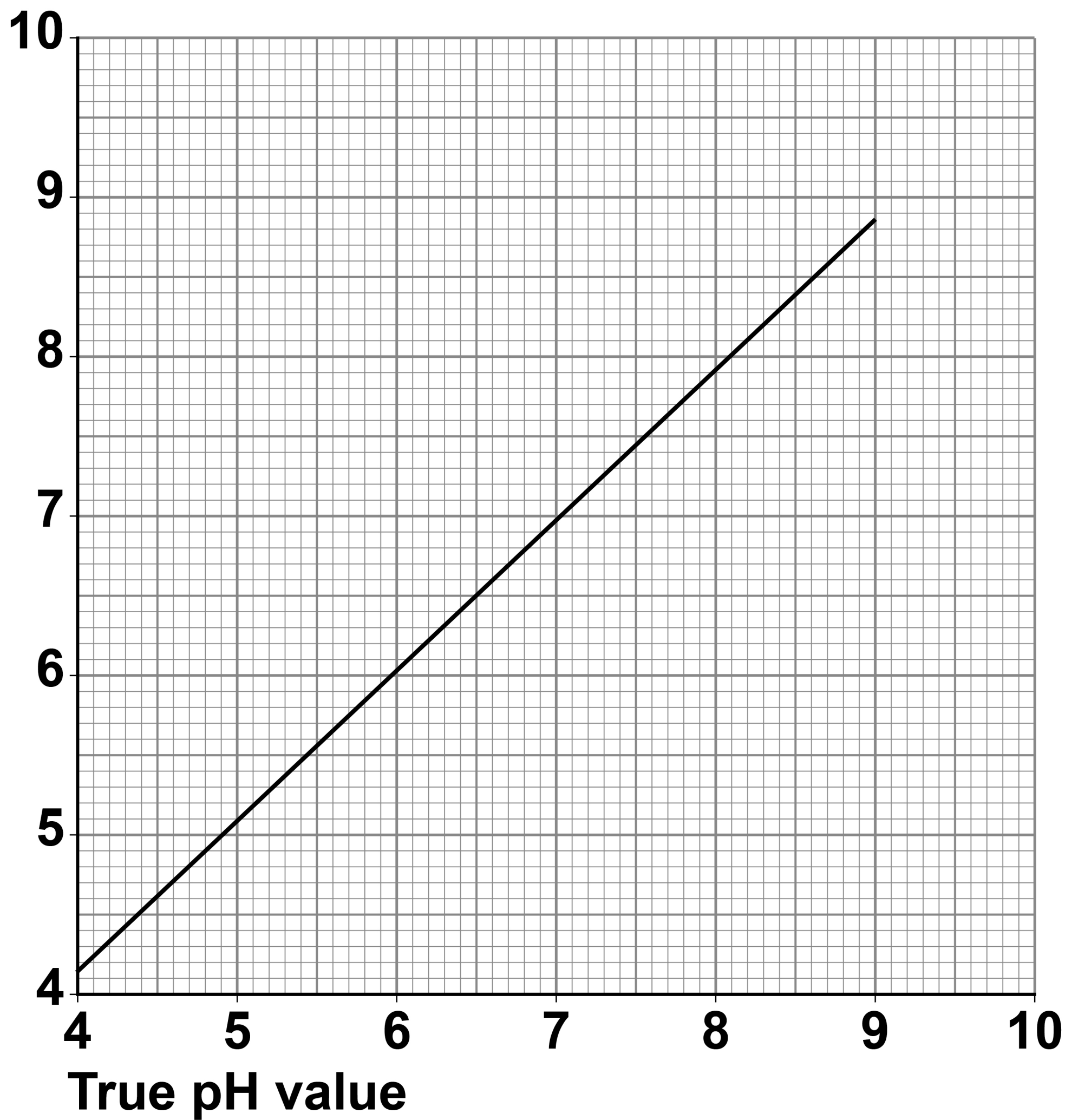
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**FIGURE 3**

**pH meter  
reading**



**[Turn over]**



**06.6**

**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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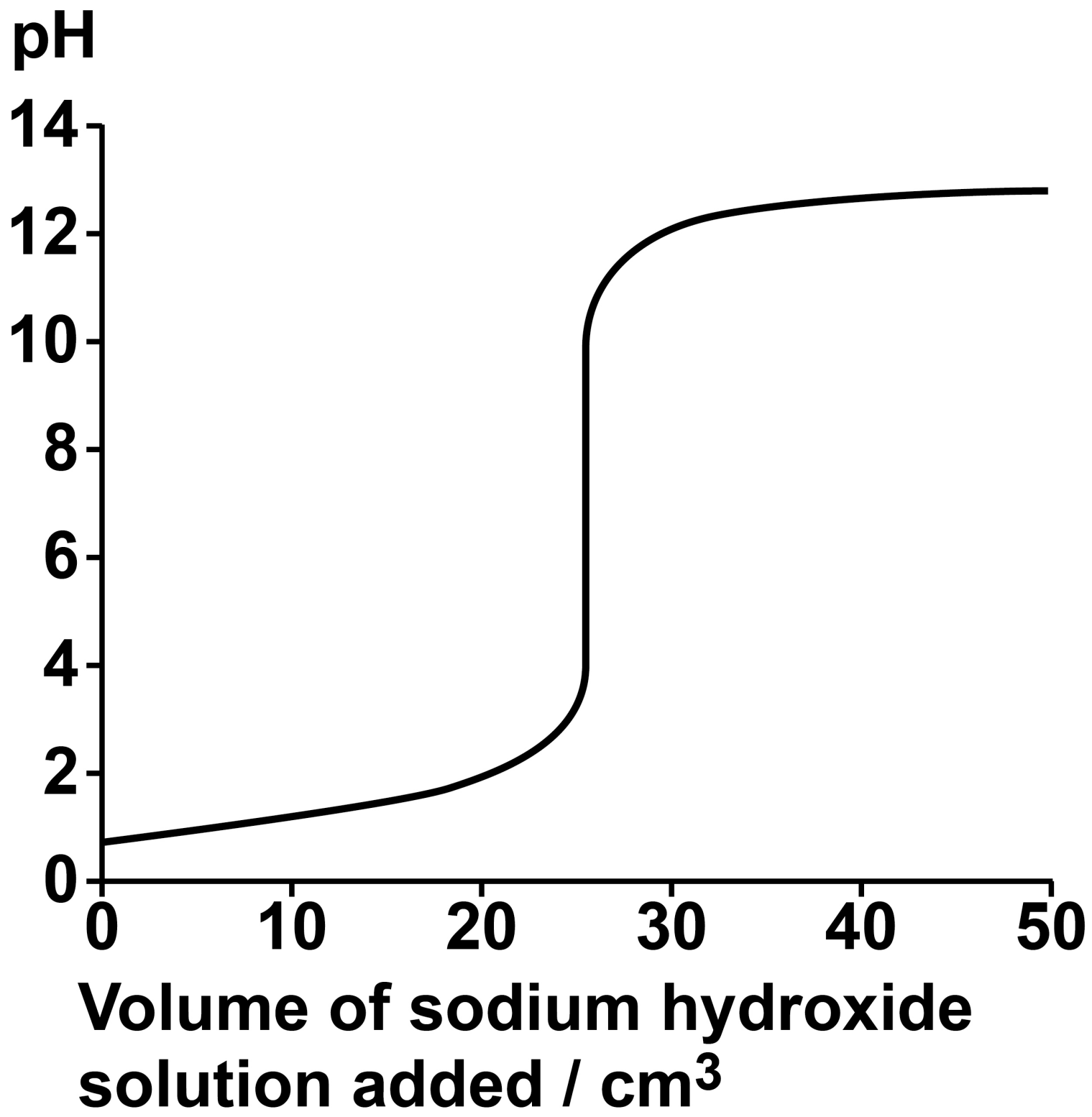
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**FIGURE 4, on the opposite page, shows the pH curve for a titration of hydrochloric acid with sodium hydroxide solution.**



FIGURE 4



[Turn over]



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

**TABLE 6**

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

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



06.7

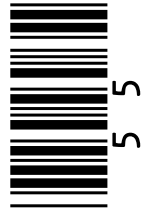
**State why all three of the indicators in TABLE 6 are suitable for this titration. [1 mark]**

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**[Turn over]**

**55**



0	6	.	8
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**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.**

**Calculate the pH of the final solution at 25 °C**

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

**[5 marks]**





pH \_\_\_\_\_

16

[Turn over]



07

This question is about thermodynamics.

Consider the reaction shown.

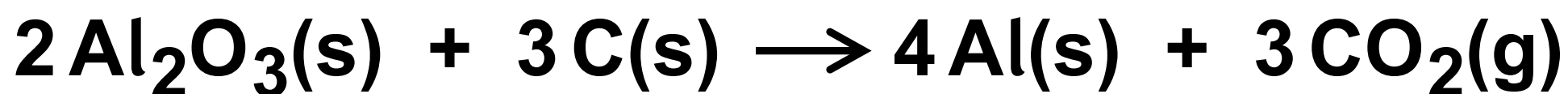


TABLE 7 shows some thermodynamic data.

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



**07.1**

**Explain why the standard entropy value for carbon dioxide is greater than that for carbon. [1 mark]**

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**07.2**

**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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**[Turn over]**



## REPEAT OF 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 . 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]



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

[Turn over]

9



0	8
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**This question is about electrode potentials and electrochemical cells.**

0	8	.	1
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**State the meaning of the term electrochemical series. [1 mark]**

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**[Turn over]**



**TABLE 8 shows some electrode potentials.**

**TABLE 8**

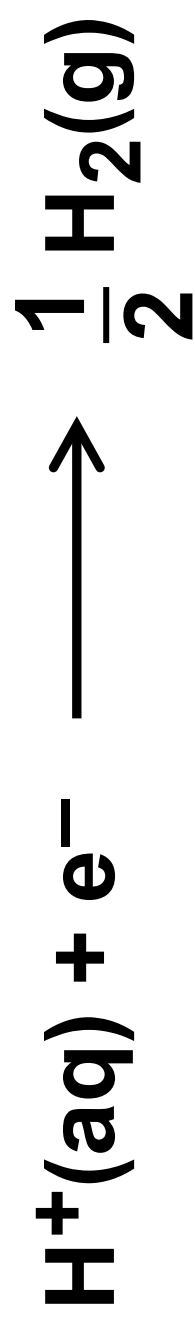
	$E^\ominus / \text{V}$
$[\text{Fe}(\text{H}_2\text{O})_6]^{2+}(\text{aq}) + 2 \text{e}^- \longrightarrow \text{Fe}(\text{s}) + 6 \text{H}_2\text{O}(\text{l})$	-0.44
$\text{H}^+(\text{aq}) + \text{e}^- \longrightarrow \frac{1}{2} \text{H}_2(\text{g})$	0.00
$[\text{Co}(\text{NH}_3)_6]^{3+}(\text{aq}) +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





08.2

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



[1 mark]

65

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08.3

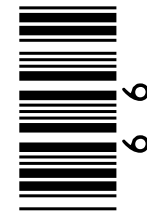
Identify the weakest reducing agent in TABLE 8. [1 mark]

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 [Turn over]

# REPEAT OF TABLE 8

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



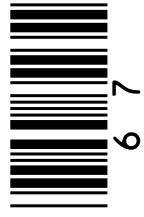
08.4

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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67

[Turn over]



# REPEAT OF TABLE 8

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



08.5

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  $[\text{Fe}(\text{H}_2\text{O})_6]^{2+}(\text{aq})$

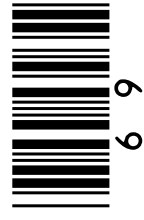
Give an equation for this reaction. [2 marks]

Explanation

69

Equation

[Turn over]



# REPEAT OF TABLE 8

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



08.6

**Suggest why the TWO cobalt(III) complex ions in TABLE 8 have different electrode potentials. [1 mark]**

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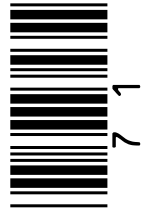
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71

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**[Turn over]**



09

This question is about the development of lithium cells.

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

TABLE 9 shows some electrode potential data.

TABLE 9

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





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**Use data in TABLE 9 to explain why an aqueous electrolyte is NOT used for a lithium cell. [2 marks]**

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**[Turn over]**



**09.2**

**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, on page 72, to calculate the cell EMF of a standard lithium-iodine cell. [1 mark]**

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**09.3**

**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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**09.4**

**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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**[Turn over]**



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

**09.5**

**Give an equation for the reaction that occurs at the positive lithium cobalt oxide electrode. [1 mark]**

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**09.6**

**Give an equation for the reaction that occurs at the negative lithium electrode. [1 mark]**

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**END OF QUESTIONS**

**7**



**Additional page, if required.**

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