## $A Q A=$

Please write clearly in block capitals.
Centre number

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Candidate number

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Surname
Forename(s)
Candidate signature
I declare this is my own work.

## A-level CHEMISTRY

## Paper 1 Inorganic and Physical Chemistry

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.

| For Examiner's Use |  |
| :---: | :---: |
| Question | Mark |
| 1 |  |
| 2 |  |
| 3 |  |
| 4 |  |
| 5 |  |
| 6 |  |
| 7 |  |
| 8 |  |
| TOTAL |  |

- 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.




## Question 1 continues on the next page

$$
\mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g})
$$

0.430 mol of carbon monoxide is mixed with 0.860 mol of hydrogen.

At equilibrium, the total pressure in the flask is 250 kPa and the mixture contains 0.110 mol of methanol.

Calculate the amount, in moles, of carbon monoxide present at equilibrium.
Calculate the partial pressure, in kPa , of carbon monoxide in this equilibrium mixture.

Amount of carbon monoxide $\qquad$ mol

Partial pressure $\qquad$ kPa

| 0 | 1 | 4 | Give an expression for the equilibrium constant $\left(K_{p}\right)$ for this reaction. |
| :--- | :--- | :--- | :--- |

$$
\mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g})
$$

$K_{p}$

| 0 | 1 | $\mathbf{5}$ | A different mixture of carbon monoxide and hydrogen is left to reach equilibrium at a |
| :--- | :--- | :--- | :--- | temperature $T$.

Some data for this equilibrium are shown in Table 1.

## Table 1

| Partial pressure of $\mathbf{C O}$ | 125 kPa |
| :--- | :---: |
| Partial pressure of $\mathrm{CH}_{3} \mathbf{O H}$ | 5.45 kPa |
| $\boldsymbol{K}_{\mathrm{p}}$ | $1.15 \times 10^{-6} \mathrm{kPa}^{-2}$ |

$$
\mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g})
$$

Calculate the partial pressure, in kPa , of hydrogen in this equilibrium mixture.
$\qquad$ kPa

| 0 | 1 | 6 |
| :--- | :--- | :--- | temperature $T$.

$$
\mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g}) \rightleftharpoons \mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g})
$$

Give the units for $K_{p}$


| $\mathbf{0}$ | $\mathbf{2}$ Rhenium has an atomic number of 75 |
| :--- | :--- | :--- |


| $\mathbf{0}$ | $\mathbf{2} .1$ | Define the term relative atomic mass. |
| :--- | :--- | :--- |

$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{2}$ | $\mathbf{2}$ The relative atomic mass of a sample of rhenium is 186.3 |
| :--- | :--- | :--- |

Table 2 shows information about the two isotopes of rhenium in this sample.

## Table 2

| Relative isotopic mass | Relative abundance |
| :---: | :---: |
| 185 | 10 |
| To be calculated | 17 |

Calculate the relative isotopic mass of the other rhenium isotope. Show your working.

Relative isotopic mass $\qquad$
$\begin{array}{llll}0 & 2 & 3 & \text { State why the isotopes of rhenium have the same chemical properties. }\end{array}$
$\qquad$
$\qquad$

A sample of rhenium is ionised by electron impact in a time of flight (TOF) mass spectrometer.
 1.450 m flight tube.

The kinetic energy of the ion is given by the equation $K E=\frac{1}{2} m v^{2}$
where
$m=$ mass / kg
$v=$ speed $/ \mathrm{m} \mathrm{s}^{-1}$
$K E=$ kinetic energy $/ \mathrm{J}$
Calculate the time, in seconds, for the ion to reach the detector.
The Avogadro constant, $L=6.022 \times 10^{23} \mathrm{~mol}^{-1}$

| $\mathbf{0}$ | $\mathbf{2}$ | $\mathbf{5}$ State how the relative abundance of ${ }^{185} \mathrm{Re}^{+}$is determined in a TOF mass |
| :--- | :--- | :--- | :--- | spectrometer.

$\qquad$
$\qquad$
$\qquad$
Turn over for the next question

| $\mathbf{0}$ | $\mathbf{3}$ This question is about hydrogen peroxide, $\mathrm{H}_{2} \mathrm{O}_{2}$ |
| :--- | :--- |

The half-equation for the oxidation of hydrogen peroxide is

$$
\mathrm{H}_{2} \mathrm{O}_{2} \rightarrow \mathrm{O}_{2}+2 \mathrm{H}^{+}+2 \mathrm{e}^{-}
$$

Hair bleach solution contains hydrogen peroxide.
A sample of hair bleach solution is diluted with water.
The concentration of hydrogen peroxide in the diluted solution is $5.00 \%$ of that in the original solution.
A $25.0 \mathrm{~cm}^{3}$ sample of the diluted hair bleach solution is acidified with dilute sulfuric acid.
This acidified sample is titrated with $0.0200 \mathrm{~mol} \mathrm{dm}^{-3}$ potassium manganate(VII) solution.
The reaction is complete when $35.85 \mathrm{~cm}^{3}$ of the potassium manganate(VII) solution are added.

| $\mathbf{0}$ | $\mathbf{3}$. | $\mathbf{1}$ Give an ionic equation for the reaction between potassium manganate(VII) and |
| :--- | :--- | :--- | acidified hydrogen peroxide.

Calculate the concentration, in mol $\mathrm{dm}^{-3}$, of hydrogen peroxide in the original hair bleach solution.
(If you were unable to write an equation for the reaction you may assume that the mole ratio of potassium manganate(VII) to hydrogen peroxide is $3: 4$ This is not the correct mole ratio.)
$\qquad$ $\mathrm{mol} \mathrm{dm}^{-3}$

| $\mathbf{0}$ | $\mathbf{3} .2$ | $\mathbf{2}$ State why an indicator is not added in this titration. |
| :--- | :--- | :--- |

$\qquad$
$\qquad$

| 0 | $\mathbf{3}$. | $\mathbf{3}$ Give the oxidation state of oxygen in hydrogen peroxide. |
| :--- | :--- | :--- |


| 0 | $\mathbf{3} .4$ | $H y d r o g e n ~ p e r o x i d e ~ d e c o m p o s e s ~ t o ~ f o r m ~ w a t e r ~ a n d ~ o x y g e n . ~$ |
| :--- | :--- | :--- |

Give an equation for this reaction.
Calculate the amount, in moles, of hydrogen peroxide that would be needed to produce $185 \mathrm{~cm}^{3}$ of oxygen gas at 100 kPa and 298 K

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

Equation

Amount $\qquad$ mol

| 0 | 3 | 5 | $\mathbf{5}$ Hydrazine $\left(\mathrm{N}_{2} \mathrm{H}_{4}\right)$ is used as a rocket fuel that is oxidised by hydrogen peroxide. |
| :--- | :--- | :--- | :--- |

The equation for this reaction in the gas phase is


H
The enthalpy change for this reaction, $\Delta H=-789 \mathrm{~kJ} \mathrm{~mol}^{-1}$
Table 3 shows some mean bond enthalpy values.
Table 3

|  | N-H | N-N | NミN | O-H |
| :--- | :---: | :---: | :---: | :---: |
| Mean bond <br> enthalpy $/ \mathrm{kJ} \mathrm{mol}^{-1}$ | 388 | 163 | 944 | 463 |

Define the term mean bond enthalpy.
Use the equation and the data in Table 3 to calculate a value for the $\mathrm{O}-\mathrm{O}$ bond enthalpy in hydrogen peroxide.

Definition $\qquad$

Bond enthalpy $\qquad$ $\mathrm{kJ} \mathrm{mol}^{-1}$
Turn over for the next question


H

| 0 | 4 | 2 | 2 |
| :--- | :--- | :--- | :--- |

Calculate the pH of a $0.0100 \mathrm{~mol} \mathrm{dm}^{-3}$ solution of strontium hydroxide at $10^{\circ} \mathrm{C}$ You may assume that strontium hydroxide is completely dissociated in this solution.

At $10^{\circ} \mathrm{C}$ the ionic product of water, $K_{w}=2.93 \times 10^{-15} \mathrm{~mol}^{2} \mathrm{dm}^{-6}$
$\qquad$

| $\mathbf{0}$ | $\mathbf{4}$ | $\mathbf{3}$ The pH of a barium hydroxide solution is lower at $50^{\circ} \mathrm{C}$ than at $10^{\circ} \mathrm{C} \mathrm{C}$ |
| :--- | :--- | :--- | :--- |

At $50^{\circ} \mathrm{C}$ a $25 \mathrm{~cm}^{3}$ sample of this barium hydroxide solution was neutralised by $22.45 \mathrm{~cm}^{3}$ of hydrochloric acid added from a burette.

Deduce the volume of this hydrochloric acid that should be added from a burette to neutralise another $25 \mathrm{~cm}^{3}$ sample of this barium hydroxide solution at $10^{\circ} \mathrm{C}$

Circle ( $\bigcirc$ ) the correct answer.

$$
>22.45 \mathrm{~cm}^{3} \quad=22.45 \mathrm{~cm}^{3} \quad<22.45 \mathrm{~cm}^{3}
$$

Explain your answer $\qquad$
$\qquad$
$\qquad$
$\qquad$

| 0 | 4 | 4 | State how a buffer solution can be made from solutions of potassium hydroxide and |
| :--- | :--- | :--- | :--- | ethanoic acid.

Give an equation for the reaction between potassium hydroxide and ethanoic acid.
State how this buffer solution resists changes in pH when a small amount of acid is added.

How buffer solution is made $\qquad$
$\qquad$
$\qquad$
$\qquad$
Equation

How buffer solution resists pH change $\qquad$
$\qquad$
$\qquad$
$\qquad$

| 0 | 4 | 5 |
| :--- | :--- | :--- |
| 5 |  |  | $500 \mathrm{~cm}^{3}$ of $1.00 \mathrm{~mol} \mathrm{dm}^{-3}$ ethanoic acid solution.

Calculate the pH of this buffer solution at $25^{\circ} \mathrm{C}$
Give your answer to 2 decimal places.
For ethanoic acid, $K_{\mathrm{a}}=1.74 \times 10^{-5} \mathrm{~mol} \mathrm{dm}^{-3}$ at $25^{\circ} \mathrm{C}$


| $\mathbf{0}$ | $\mathbf{5}$ | $\mathbf{1}$ Which is not a correct statement about magnesium hydroxide? |
| :--- | :--- | :--- |

Tick ( $\checkmark$ ) one box.

It is used to neutralise stomach acid


It forms a solution with $\mathrm{pH}=14$ at $25^{\circ} \mathrm{C}$


It has the empirical formula $\mathrm{H}_{2} \mathrm{MgO}_{2}$


| 0 | 5 | $\mathbf{2}$ Give an equation for the reaction of aluminium oxide with sulfuric acid. |
| :--- | :--- | :--- |


| $\mathbf{0}$ | $\mathbf{5}$ | $\mathbf{3}$ | Identify a reagent or test that could be used to distinguish between aqueous solutions |
| :--- | :--- | :--- | :--- | of sulfur dioxide and sulfur trioxide with the same concentrations.

State the observation in each case.

Reagent or test $\qquad$
Observation with sulfur dioxide solution $\qquad$
$\qquad$
Observation with sulfur trioxide solution $\qquad$
$\qquad$

Question 5 continues on the next page

| $\mathbf{0}$ | $\mathbf{5}$ | $\mathbf{4}$ The mass spectrum of the element phosphorus has a peak at $\frac{\mathrm{m}}{\mathrm{z}}=124$. |
| :--- | :--- | :--- | :--- |

Give the formula of the species responsible for this peak.
$\qquad$

| $\mathbf{0}$ | $\mathbf{5}$ | $\mathbf{5}$ | Give an equation for the reaction of phosphorus $(\mathrm{V})$ oxide with sodium hydroxide |
| :--- | :--- | :--- | :--- | solution.


| $\mathbf{0}$ | $\mathbf{5} .6$ | Draw the displayed formula of the molecule formed when phosphorus $(\mathrm{V})$ oxide reacts |
| :--- | :--- | :--- | :--- | with water.



| 0 | 5 | $\mathbf{7}$ | Table 4 shows the melting points of three substances. |
| :--- | :--- | :--- | :--- |

## Table 4

| Substance | Melting point / K |
| :--- | :---: |
| sodium chloride | 1074 |
| chlorine | 172 |
| hydrogen chloride | 158 |

Explain why the melting points of these substances are different.
You should refer to the structure of and bonding in each substance.
[6 marks]
You should refer to the structure of and bonding in each substance.
$\qquad$
$\qquad$
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$\qquad$


| 0 | 6 | This question is about some elements in Group 7 and their compounds. |
| :--- | :--- | :--- |


| $\mathbf{0}$ | $\mathbf{6}$. | $\mathbf{1}$ Chlorine is added to some drinking water supplies to decrease the risk of people |
| :--- | :--- | :--- | :--- | suffering from diseases such as cholera.

State why the amount of chlorine added must be controlled.
$\qquad$
$\qquad$

| 0 | 6 | 2 |
| :--- | :--- | :--- | two acids.

Explain, with reference to electrons, why this is a redox reaction.

Equation
$\qquad$
Explanation $\qquad$
$\qquad$
$\qquad$
$\qquad$

| 0 | 6 | 3 | A student bubbles chlorine gas through a solution of sodium iodide. |
| :--- | :--- | :--- | :--- |

State the observation the student would make.
Give an ionic equation for the reaction.

Observation $\qquad$
Ionic equation
$\qquad$

| $\mathbf{0}$ | $\mathbf{6}$. | $\mathbf{4}$ The student adds a few drops of concentrated sulfuric acid to a small amount of |
| :--- | :--- | :--- | :--- | solid sodium iodide.

Two gaseous sulfur-containing products are formed.
Give an equation for the formation of each of these sulfur-containing products.
State the role of sulfuric acid in the formation of these products.

Equation 1

## Equation 2

Role

| 0 | 6 | $\mathbf{5}$ | The student adds a few drops of acidified silver nitrate solution to a solution of an |
| :--- | :--- | :--- | :--- | unknown impure sodium halide.

The student observes bubbles of gas and a colourless solution.
The student bubbles the gas through calcium hydroxide solution and a white precipitate forms.

Deduce the identity of the sodium halide.
Suggest the identity of the gas.
Give an ionic equation for the formation of this gas from the impurity.

Identity of sodium halide $\qquad$
Identity of gas $\qquad$
Ionic equation

| $\mathbf{0}$ | $\mathbf{6}$ | .6 |
| :--- | :--- | :--- |
| $\mathbf{T}$ |  |  | $\mathrm{The}_{\mathrm{Cl}}^{2}{ }^{+}$ion contains two different Group 7 elements.

Use your understanding of the electron pair repulsion theory to draw the shape of this ion.

Include any lone pairs of electrons that influence the shape.
Explain why the ion has the shape you have drawn.
Suggest a value for the bond angle in the ion.

Shape

Explanation $\qquad$
$\qquad$
$\qquad$
Bond angle

| 0 | 6 | .7 |
| :--- | :--- | :--- | Magnesium is used in the extraction of titanium from titanium(IV) chloride.

Give an equation for this reaction.
Explan

## [3

| $\mathbf{0}$ | $\mathbf{7}$ | Copper(II) complexes are coloured. |
| :--- | :--- | :--- |

The colour is caused by the d electrons of copper moving from their ground state to an excited state.

| $\mathbf{0}$ | $\mathbf{7}$. | $\mathbf{1}$ Explain why aqueous solutions containing $\left[\mathrm{CuCl}_{4}\right]^{2-}$ ions are yellow. ${ }^{2}$. |
| :--- | :--- | :--- |

$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{7} .2$ | $\mathbf{2}$ When a d electron moves from the ground state to the excited state in a |
| :--- | :--- | :--- | copper complex, the energy change is $3.98 \times 10^{-19} \mathrm{~J}$

The Planck constant, $h=6.63 \times 10^{-34} \mathrm{~J} \mathrm{~s}$
Calculate the frequency, in $\mathrm{s}^{-1}$, of the light absorbed.
$\qquad$

| 0 | 7 | 3 | State three ways in which a transition metal complex can be changed to alter its |
| :--- | :--- | :--- | :--- | colour.

1

2

3 $\qquad$

Question 7 continues on the next page


| 0 | 7 | 4 | Name the shape of the $\left[\mathrm{CuCl}_{4}\right]^{2-}$ ion. |
| :--- | :--- | :--- | :--- |

$\qquad$


| $\mathbf{0}$ | $\mathbf{7}$. | $\mathbf{6}$ | State the colour of the solution containing the complex ion $\mathbf{Q}$. |
| :--- | :--- | :--- | :--- |

Give an ionic equation for the conversion of $\left[\mathrm{CuCl}_{4}\right]^{2-}$ to $\mathbf{Q}$.

Colour
Equation

| 0 | $\mathbf{7}$ | $\mathbf{7}$ | Identify complex ion $\mathbf{R}$. |
| :--- | :--- | :--- | :--- |


| 0 | 8 |
| :--- | :--- |


| $\mathbf{0}$ | $\mathbf{8}$. | $\mathbf{1}$ The half-equations for two electrodes that combine to make a non-rechargeable cell |
| :--- | :--- | :--- | are

$\mathrm{Zn}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Zn}(\mathrm{s})$
$2 \mathrm{MnO}_{2}(\mathrm{~s})+2 \mathrm{NH}_{4}{ }^{+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Mn}_{2} \mathrm{O}_{3}(\mathrm{~s})+2 \mathrm{NH}_{3}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\mathrm{I}) \quad E^{\ominus}=+0.52 \mathrm{~V}$
Identify the oxidising agent in this cell.

Figure 1 shows a cross-section through a rechargeable silver-zinc cell.
Figure 1

$\begin{array}{llll}0 & 8 & 2 & \text { Suggest the function of the porous separator in Figure } 1 .\end{array}$
$\qquad$

| 0 | 8 | .3 | The standard electrode potentials for two half-equations for the silver-zinc cell are |
| :--- | :--- | :--- | :--- |

$$
\begin{array}{ll}
\mathrm{Ag}_{2} \mathrm{O}(\mathrm{~s})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l})+2 \mathrm{e}^{-} \rightarrow 2 \mathrm{Ag}(\mathrm{~s})+2 \mathrm{OH}^{-}(\mathrm{aq}) & E^{\ominus}=+0.34 \mathrm{~V} \\
\mathrm{ZnO}(\mathrm{~s})+\mathrm{H}_{2} \mathrm{O}(\mathrm{I})+2 \mathrm{e}^{-} \rightarrow \mathrm{Zn}(\mathrm{~s})+2 \mathrm{OH}^{-}(\mathrm{aq}) & E^{\circ}=-1.26 \mathrm{~V}
\end{array}
$$

Give an equation for the overall reaction that occurs when the cell is recharging.
$\qquad$

The EMF of an alkaline hydrogen-oxygen fuel cell is +1.23 V The standard electrode potential for one of the electrodes in the alkaline hydrogen-oxygen fuel cell is

$$
2 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})+2 \mathrm{e}^{-} \rightarrow 2 \mathrm{OH}^{-}(\mathrm{aq})+\mathrm{H}_{2}(\mathrm{~g}) \quad E^{\ominus}=-0.83 \mathrm{~V}
$$

| 0 | 8 | 4 |
| :--- | :--- | :--- | Give the half-equation for the other electrode and calculate its standard electrode potential.

Equation
$\qquad$
$E^{\ominus}$ $\qquad$

| $\mathbf{0}$ | $\mathbf{8}$ | $\mathbf{5}$ Suggest why the EMF values of the acidic and alkaline hydrogen-oxygen fuel cells |
| :--- | :--- | :--- | :--- | are the same.

There are no questions printed on this page

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