## Cambridge International Examinations

Cambridge International Advanced Subsidiary and Advanced Level

## International AS \& A Level



## CENTRE NUMBER

$\square$ | $\begin{array}{l}\text { CANDIDATE } \\ \text { NUMBER }\end{array}$ |  |  |  |  |
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## CHEMISTRY

Paper 4 A Level Structured Questions
Candidates answer on the Question Paper.
Additional Materials: Data Booklet

## READ THESE INSTRUCTIONS FIRST

Write your Centre number, candidate number and name on all the work you hand in.
Write in dark blue or black pen.
You may use an HB pencil for any diagrams or graphs.
Do not use staples, paper clips, glue or correction fluid.
DO NOT WRITE IN ANY BARCODES.
Answer all questions.

Electronic calculators may be used.
You may lose marks if you do not show your working or if you do not use appropriate units.
A Data Booklet is provided.
At the end of the examination, fasten all your work securely together.
The number of marks is given in brackets [ ] at the end of each question or part question.

Answer all the questions in the spaces provided.

1 Copper is a transition element and has atomic number 29.
(a) Complete the electronic configuration for the copper atom and the copper ion in the +2 oxidation state.

- copper atom
[Ar] $\qquad$
- copper ion in the +2 oxidation state
[Ar] $\qquad$
(b) The following equilibrium exists between two complex ions of copper in the +2 oxidation state.

$$
\left[\mathrm{Cu}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}+4 \mathrm{Cl}^{-} \rightleftharpoons\left[\mathrm{CuCl}_{4}\right]^{2-}+6 \mathrm{H}_{2} \mathrm{O}
$$

(i) Name the type of reaction occurring here.
$\qquad$
(ii) State the colours of these two complex ions.
$\left[\mathrm{Cu}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}$
$\left[\mathrm{CuCl}_{4}\right]^{2-}$
(iii) State the shape of the $\left.[\mathrm{CuCl}]_{4}\right]^{2-}$ ion.
$\qquad$
(iv) Write the expression for the stability constant, $K_{\text {stab }}$, for this equilibrium.

$$
\begin{equation*}
K_{\text {stab }}= \tag{1}
\end{equation*}
$$

(c) Copper also forms the complex ions $\left[\mathrm{Cu}\left(\mathrm{NH}_{3}\right)_{2}\left(\mathrm{H}_{2} \mathrm{O}\right)_{4}\right]^{2+}$ and $\left[\mathrm{Cu}(e n)\left(\mathrm{H}_{2} \mathrm{O}\right)_{4}\right]^{2+}$ where en is the bidentate ligand ethane-1,2-diamine, $\mathrm{H}_{2} \mathrm{NCH}_{2} \mathrm{CH}_{2} \mathrm{NH}_{2}$.

$$
\begin{array}{rll}
{\left[\mathrm{Cu}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}+2 \mathrm{NH}_{3}} & \rightleftharpoons\left[\mathrm{Cu}\left(\mathrm{NH}_{3}\right)_{2}\left(\mathrm{H}_{2} \mathrm{O}\right)_{4}\right]^{2+}+2 \mathrm{H}_{2} \mathrm{O} & \text { equilibrium } 1 \\
{\left[\mathrm{Cu}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}+\text { en }} & \rightleftharpoons\left[\mathrm{Cu}(e n)\left(\mathrm{H}_{2} \mathrm{O}\right)_{4}\right]^{2+}+2 \mathrm{H}_{2} \mathrm{O} & \text { equilibrium } 2
\end{array}
$$

(i) What is meant by the term bidentate ligand?
$\qquad$
$\qquad$
(ii) The table lists the values of stability constants for these two complexes.

|  | stability constant, $K_{\text {stab }}$ |
| :---: | :---: |
| $\left[\mathrm{Cu}\left(\mathrm{NH}_{3}\right)_{2}\left(\mathrm{H}_{2} \mathrm{O}\right)_{4}\right]^{2+}$ | $7.94 \times 10^{7}$ |
| $\left[\mathrm{Cu}(e n)\left(\mathrm{H}_{2} \mathrm{O}\right)_{4}\right]^{2+}$ | $3.98 \times 10^{10}$ |

What do these $K_{\text {staa }}$ values tell us about the relative positions of equilibria 1 and 2 ?
$\qquad$
$\qquad$
(d) Nickel forms the complex ion $\left[\mathrm{Ni}(e n)_{3}\right]^{2+}$ in which it is surrounded octahedrally by six nitrogen atoms.
(i) Name the type of stereoisomerism displayed by $\left[\mathrm{Ni}(e n)_{3}\right]^{2+}$.
$\qquad$
(ii) Draw three-dimensional diagrams to show the two stereoisomers of $\left[\mathrm{Ni}(e n)_{3}\right]^{2+}$.
(e) Ethane-1,2-diamine is a useful reagent in organic chemistry.
(i) Explain how the amino groups in ethane-1,2-diamine allow the molecule to act as a Brønsted-Lowry base.
$\qquad$
$\qquad$
(ii) Write an equation for the reaction of ethane-1,2-diamine with an excess of hydrochloric acid.
$\qquad$
(f) (i) Under certain conditions, ethane-1,2-diamine reacts with ethanedioic acid, $\mathrm{HO}_{2} \mathrm{CCO}_{2} \mathrm{H}$, to form the polymer $\mathbf{Z}$.

Draw the structure of this polymer, $\mathbf{Z}$, showing two repeat units.
(ii) Name the type of reaction occurring during this polymerisation.
$\qquad$
(iii) Polymer $\mathbf{Z}$ is an example of a biodegradable polymer.

Name a polymer that is non-biodegradable.

2 (a) When copper(II) carbonate is heated strongly, it decomposes in a similar way to Group 2 carbonates.

Predict what would be observed when anhydrous copper(II) carbonate is heated.
$\qquad$
$\qquad$
(b) Describe and explain how the thermal stability of the Group 2 carbonates varies down the group.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(c) Calcium cyanamide, $\mathrm{CaCN}_{2}$, can be used as a fertiliser.
(i) Complete the 'dot-and-cross' diagram for the cyanamide ion, $\mathrm{CN}_{2}{ }^{2-}$.

Use the following key for the electrons.

- electrons from carbon
$\times$ electrons from nitrogen
- added electron(s) responsible for the overall negative charge

(ii) $\mathrm{CaCN}_{2}$ decomposes readily on contact with water forming an insoluble white solid and ammonia only.

Suggest an equation for this reaction.

3 The spontaneity (feasibility) of a chemical reaction depends on the standard Gibbs free energy change, $\Delta G^{\ominus}$. This is related to the standard enthalpy and entropy changes by the equation shown.

$$
\Delta G^{\ominus}=\Delta H^{\ominus}-\mathrm{T} \Delta S^{\ominus}
$$

(a) State and explain whether the following processes will lead to an increase or decrease in entropy.
(i) the reaction of magnesium with hydrochloric acid
entropy change $\qquad$
explanation $\qquad$
(ii) solid potassium chloride dissolving in water
entropy change $\qquad$
explanation $\qquad$
(iii) steam condensing to water
entropy change $\qquad$
explanation $\qquad$
(b) Magnesium carbonate can be decomposed.

$$
\mathrm{MgCO}_{3}(\mathrm{~s}) \rightarrow \mathrm{MgO}(\mathrm{~s})+\mathrm{CO}_{2}(\mathrm{~g}) \quad \Delta H^{\ominus}=+117 \mathrm{kJmol}^{-1}
$$

Standard entropies are shown in the table.

| substance | $\mathrm{MgCO}_{3}(\mathrm{~s})$ | $\mathrm{MgO}(\mathrm{s})$ | $\mathrm{CO}_{2}(\mathrm{~g})$ |
| :---: | :---: | :---: | :---: |
| $\mathrm{S}^{\ominus} / \mathrm{Jmol}^{-1} \mathrm{~K}^{-1}$ | +65.7 | +26.9 | +214 |

(i) Calculate $\Delta G^{\ominus}$ for this reaction at 298 K . Include a relevant sign and give your answer to three significant figures.

$$
\Delta G^{\ominus}=
$$

$\qquad$ $\mathrm{kJmol}^{-1}$
(ii) Explain, with reference to $\Delta G^{\ominus}$, why this reaction becomes more feasible at higher temperatures.
$\qquad$
$\qquad$
(c) On heating, sodium hydrogencarbonate decomposes into sodium carbonate as shown.

$$
\begin{array}{ll}
2 \mathrm{NaHCO}_{3}(\mathrm{~s}) \rightarrow \mathrm{Na}_{2} \mathrm{CO}_{3}(\mathrm{~s})+\mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \quad \begin{array}{l}
\Delta H^{\ominus}=+130 \mathrm{kJmol}^{-1} \\
\Delta S^{\ominus}=+316 \mathrm{Jmol}^{-1} \mathrm{~K}^{-1}
\end{array}
\end{array}
$$

Calculate the minimum temperature at which this reaction becomes spontaneous (feasible). Show your working.
temperature =
(d) The solubility of Group 2 sulfates decreases down the Group.

Explain this trend.
$\qquad$
$\qquad$
$\qquad$
$\qquad$

4 (a) Cobalt is a transition element that forms complex ions with oxidation states +2 and +3 .
Explain what is meant by the term transition element.
$\qquad$
$\qquad$
(b) The following scheme shows some reactions of $\left[\mathrm{Co}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}$.

(i) State the formula of each of the following.

A

B $\qquad$
C
(ii) State the colour of the following solutions.
$\left[\mathrm{Co}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}$ $\qquad$
solution of B $\qquad$
solution of C $\qquad$
(c) Define the term standard electrode potential.
$\qquad$
$\qquad$
$\qquad$
(d) An electrochemical cell was set up to measure the standard electrode potential, $E_{\text {cell, }}^{\ominus}$, of a cell made of a $\mathrm{Co}^{2+} / \mathrm{Co}$ half-cell and a $\mathrm{Fe}^{3+} / \mathrm{Fe}^{2+}$ half-cell.
(i) Complete the table with the substance used to make the electrode in each of these half-cells.

| half-cell | electrode |
| :---: | :---: |
| $\mathrm{Co}^{2+} / \mathrm{Co}$ |  |
| $\mathrm{Fe}^{3+} / \mathrm{Fe}^{2+}$ |  |

(ii) Write the equation for the overall cell reaction.
$\qquad$
(iii) Use the Data Booklet to calculate the $E_{\text {cell }}^{\ominus}$.

$$
\begin{equation*}
E_{\text {cell }}^{\ominus}= \tag{1}
\end{equation*}
$$

(e) The electrochemical cell in (d) was set up again but this time the concentration of $\mathrm{Co}^{2+}(\mathrm{aq})$ was $0.050 \mathrm{~mol} \mathrm{dm}^{-3}$.

The Nernst equation can be used to calculate the value of an electrode potential at different concentrations.

$$
E=E^{\ominus}+(0.059 / \mathrm{z}) \log \left[\mathrm{Co}^{2+}(\mathrm{aq})\right] \quad \text { Nernst equation }
$$

(i) Use the Data Booklet and the Nernst equation to calculate the value of $E$ for the $\mathrm{Co}^{2+} / \mathrm{Co}$ half-cell in this experiment.

$$
E \text { for } \mathrm{Co}^{2+} / \mathrm{Co}=
$$

(ii) Suggest how this change will affect the overall cell potential, $E_{\text {cell }}$, compared to $E_{\text {cell }}^{\ominus}$ in (d)(iii).

Circle your answer.
less positive no change more positive
(f) Iron(III) ions can oxidise vanadium metal.

Construct an equation for the reaction of an excess of iron(III) ions with vanadium metal.
Use of the Data Booklet will be helpful.

5 Compound $\mathbf{F}$ contains the elements carbon, hydrogen and oxygen only. All carbon-carbon bonds in $\mathbf{F}$ are single bonds. The structure of $\mathbf{F}$ was analysed by mass spectrometry and infra-red and NMR spectroscopy.
(a) The mass spectrum shows that the $m / e$ value for the $M$ peak is 90 .

The ratio of the heights of the $M$ and $M+1$ peaks is $22.1: 0.7$.
(i) Use the ratio of the heights of the M and $\mathrm{M}+1$ peaks to calculate the number of carbon atoms in a molecule of $\mathbf{F}$.
number of carbon atoms $=$
(ii) Suggest the molecular formula of $\mathbf{F}$.
molecular formula $=\mathrm{CHO}$
(b) The infra-red spectrum of $\mathbf{F}$ was obtained.

Use the Data Booklet and your knowledge of infra-red spectroscopy to identify the type of bond and the functional group responsible for these three absorptions.

| absorption $/ \mathrm{cm}^{-1}$ | appearance of the peak | type of bond | functional group |
| :---: | :---: | :---: | :---: |
| 3350 | broad and strong |  |  |
| 2680 | very broad and strong |  |  |
| 1725 | strong |  |  |

(c) $\mathbf{F}$ was dissolved in deuterated trichloromethane, $\mathrm{CDCl}_{3}$, and the proton NMR spectrum of this solution obtained.

(i) Use the Data Booklet and your answer to (a)(ii) to complete Table 1 for the proton NMR spectrum of $F$.
The actual chemical shifts for the four absorptions in $\mathbf{F}$ have been added for you.
Table 1

| $\delta / \mathrm{ppm}$ | type of proton | relative peak area |
| :---: | :---: | :---: |
| 1.4 |  |  |
| 3.9 |  |  |
| 4.7 |  |  |
| 12.9 |  |  |

(ii) Describe and explain the splitting pattern for the absorption at $\delta=1.4$.
$\qquad$
$\qquad$
(iii) F was dissolved in $\mathrm{D}_{2} \mathrm{O}$ and the proton NMR spectrum of this new solution obtained. Two of the absorptions in Table 1 were not present in this spectrum.

Which absorptions were not present?
$\qquad$
and
(iv) Suggest the structure of F .
(d) Molecules of cycloheptadiene, $\mathrm{C}_{7} \mathrm{H}_{10}$, consist of a seven-membered ring with two carbon-carbon double bonds.
(i) Complete the skeletal formulae of two isomers of cycloheptadiene.


The isomers $\mathbf{P}$ and $\mathbf{Q}$ were analysed using carbon-13 NMR spectroscopy.
(ii) Predict the number of peaks that will be seen in the carbon-13 NMR spectra of $\mathbf{P}$ and $\mathbf{Q}$.

| isomer | number of peaks |
| :---: | :---: |
| $\mathbf{P}$ |  |
| $\mathbf{Q}$ |  |

Question 6 starts on the next page.

6 Ibuprofen and paracetamol are pain-relief drugs.

ibuprofen

paracetamol
(a) Ibuprofen and paracetamol both contain the aryl (benzene) functional group.

Name the other functional groups present in each molecule.
ibuprofen $\qquad$
paracetamol $\qquad$
(b) Ibuprofen contains a chiral centre and shows stereoisomerism.
(i) State what is meant by the term chiral centre.
$\qquad$
$\qquad$
(ii) Draw the two stereoisomers of ibuprofen.


(c) Draw the structures of the organic products when ibuprofen and paracetamol react separately with $\mathrm{LiAlH}_{4}$.
$\square$

| product with paracetamol |
| :---: |

(d) A student carried out some reactions with solutions of ibuprofen and paracetamol using reagents $\mathbf{D}$ and $\mathbf{E}$ and the following results were obtained.
( $\checkmark$ means a reaction took place.)

| reagent | ibuprofen | paracetamol |
| :---: | :---: | :---: |
| D | $\checkmark$ | $\boldsymbol{x}$ |
| E | $\boldsymbol{x}$ | $\checkmark$ |

(i) Suggest a possible identity for each reagent $\mathbf{D}$ and $\mathbf{E}$.

D $\qquad$
E $\qquad$
(ii) Give the structure of the organic product formed when reagent $\mathbf{D}$ reacted with ibuprofen.

(iii) Give the structure of the organic product formed when reagent E reacted with paracetamol.

(e) One of the steps in the manufacture of ibuprofen is shown.

(i) Write an equation for the reaction between $\mathrm{CH}_{3} \mathrm{COCl}$ and $\mathrm{AlCl}_{3}$.
$\qquad$
(ii) Complete the mechanism for the conversion of $\mathbf{X}$ into $\mathbf{Y}$. Include all necessary curly arrows, any relevant dipoles and charges.


X
(iii) Name the mechanism in (ii).
$\qquad$

Question 7 starts on the next page.

7 (a) Sodium chlorate(I), NaClO , is the active ingredient in commercial bleach. The concentration of chlorate(I) ions was determined by titration.

- $10.0 \mathrm{~cm}^{3}$ of a bleach solution was diluted to $250 \mathrm{~cm}^{3}$ in a volumetric flask using distilled water.
- Dilute sulfuric acid and an excess of potassium iodide solution were added to a $25.0 \mathrm{~cm}^{3}$ portion of this solution to liberate iodine.
- The resulting solution required $20.80 \mathrm{~cm}^{3}$ of $0.100 \mathrm{moldm}^{-3}$ aqueous sodium thiosulfate solution to react with the iodine produced.

The titration reactions are shown.

$$
\begin{gathered}
\mathrm{ClO}^{-}+2 \mathrm{I}^{-}+2 \mathrm{H}^{+} \rightarrow \mathrm{I}_{2}+\mathrm{Cl}^{-}+\mathrm{H}_{2} \mathrm{O} \\
\mathrm{I}_{2}+2 \mathrm{~S}_{2} \mathrm{O}_{3}{ }^{2-} \rightarrow 2 \mathrm{I}^{-}+\mathrm{S}_{4} \mathrm{O}_{6}{ }^{2-}
\end{gathered}
$$

Calculate the concentration, in $\mathrm{moldm}^{-3}$, of $\mathrm{ClO}^{-}$ions in the bleach solution.

$$
\text { concentration of } \mathrm{ClO}^{-}=
$$

$\qquad$ $\mathrm{moldm}^{-3}$
(b) An indicator was used in the thiosulfate-iodine titration.
(i) Name a suitable indicator for this titration.
(ii) State the expected colour change you would observe at the end-point in this titration.
$\qquad$
(iii) State when in the procedure you would add the indicator.
$\qquad$
$\qquad$
(c) The concentration of chlorate(I) ions can also be determined by adding an excess of hydrogen peroxide to the sample of bleach and measuring the volume of oxygen gas produced.

$$
\mathrm{H}_{2} \mathrm{O}_{2}+\mathrm{NaClO} \rightarrow \mathrm{NaCl}+\mathrm{O}_{2}+\mathrm{H}_{2} \mathrm{O}
$$



When an excess of aqueous hydrogen peroxide was added to $5.0 \mathrm{~cm}^{3}$ of a different bleach solution, $82 \mathrm{~cm}^{3}$ of oxygen was produced at room temperature and pressure.

Calculate the concentration of ClO - ions in this bleach solution.

$$
\begin{equation*}
\text { concentration of } \mathrm{ClO}^{-}= \tag{2}
\end{equation*}
$$

$\qquad$ $\mathrm{moldm}^{-3}$
(d) Trichlorocyanuric acid, $\mathrm{C}_{3} \mathrm{Cl}_{3} \mathrm{~N}_{3} \mathrm{O}_{3}$, acts as a chlorine buffer and disinfectant for swimming pools. It reacts with water to give chloric(I) acid, HClO .

$$
\mathrm{C}_{3} \mathrm{Cl}_{3} \mathrm{~N}_{3} \mathrm{O}_{3}+3 \mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{C}_{3} \mathrm{H}_{3} \mathrm{~N}_{3} \mathrm{O}_{3}+3 \mathrm{HCl} \mathrm{O}
$$

(i) Write the expression for $K_{\mathrm{c}}$ for this equilibrium.

$$
K_{\mathrm{c}}=
$$

(ii) In outdoor swimming pools, the HClO is decomposed by sunlight. The decomposition of HClO is a redox reaction which forms a gas that relights a glowing splint.

Describe and explain the effect of the decomposition of HClO on the equilibrium in (d). State the effect on $K_{\mathrm{c}}$.
$\qquad$
$\qquad$
effect on $K_{c}$ $\qquad$
(iii) The decomposition of HClO is a redox reaction.

## Suggest an equation for this reaction.

(e) The buffer solution in blood is a mixture of carbonic acid, $\mathrm{H}_{2} \mathrm{CO}_{3}$, and hydrogencarbonate ions, $\mathrm{HCO}_{3}^{-}$. Healthy blood has a pH of 7.40.

$$
\mathrm{H}_{2} \mathrm{CO}_{3}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{HCO}_{3}^{-}+\mathrm{H}_{3} \mathrm{O}^{+} \quad K_{\mathrm{a}}=7.94 \times 10^{-7} \mathrm{moldm}^{-3}
$$

(i) Explain how this buffer system acts to control the blood pH. Include equations in your answer.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) A patient's blood has a $\left[\mathrm{HCO}_{3}^{-}\right]:\left[\mathrm{H}_{2} \mathrm{CO}_{3}\right]$ ratio of 9.5:1.

Calculate the pH of the patient's blood.

$$
\mathrm{pH}=
$$

[Total: 16]

[^0]
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