## $A Q A B$

Please write clearly in block capitals.
Centre number

|  |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- |

Candidate number


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

## A-level CHEMISTRY

## Paper 3

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

| For Examiner's Use |  |
| :---: | :---: |
| Question | Mark |
| 1 |  |
| 2 |  |
| 3 |  |
| 4 |  |
| 5 |  |
| Section B |  |
| TOTAL |  | 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 90 .


## Advice

- You are advised to spend 70 minutes on Section A and 50 minutes on Section B.


Section A $\quad |$| Do not write |
| :---: |
| outsidid the |
| box |

Answer all questions in this section.

| 0 | 1 |
| :--- | :--- | A value for enthalpy of solution can be determined in two ways:

- from a cycle, using lattice enthalpy and enthalpies of hydration
- from the results of a calorimetry experiment.

| 0 | 1 | 1 |
| :--- | :--- | :--- | Define the term enthalpy of lattice dissociation.

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

| $\mathbf{0}$ | $\mathbf{1} .2$ | The enthalpy of solution for ammonium nitrate is the enthalpy change for the reaction |
| :--- | :--- | :--- | shown.

$$
\mathrm{NH}_{4} \mathrm{NO}_{3}(\mathrm{~s})+\mathrm{aq} \rightarrow \mathrm{NH}_{4}^{+}(\mathrm{aq})+\mathrm{NO}_{3}^{-}(\mathrm{aq}) \quad \Delta H=+26 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

## Table 1

|  | $\mathrm{NH}_{4}{ }^{+}(\mathrm{g})$ | $\mathrm{NO}_{3}-(\mathrm{g})$ |
| :--- | :---: | :---: |
| Enthalpy of hydration $\Delta_{\mathrm{hyd}} \mathrm{H} / \mathrm{kJ} \mathrm{mol}^{-1}$ | -307 | -314 |

Draw a suitably labelled cycle and use it, with data from Table 1, to calculate the enthalpy of lattice dissociation for ammonium nitrate.
$\qquad$ $\mathrm{kJ} \mathrm{mol}^{-1}$

## Turn over

| $\mathbf{0}$ | $\mathbf{1}$ | .3 | $\mathbf{3}$ |
| :--- | :--- | :--- | :--- |
| A student does an experiment to determine a value for the enthalpy of solution for |  |  |  | ammonium nitrate.

The student uses this method.

- Measure $25.0 \mathrm{~cm}^{3}$ of distilled water in a measuring cylinder.
- Pour the water into a beaker.
- Record the temperature of the water in the beaker.
- Add 4.00 g of solid $\mathrm{NH}_{4} \mathrm{NO}_{3}$ to the water in the beaker.
- Stir the solution and record the lowest temperature reached.

Table 2 shows the student's results.
Table 2

| Initial temperature $/{ }^{\circ} \mathrm{C}$ | 20.2 |
| :--- | :--- |
| Lowest temperature $/{ }^{\circ} \mathrm{C}$ | 12.2 |

Calculate the enthalpy of solution, in $\mathrm{kJ} \mathrm{mol}^{-1}$, for ammonium nitrate in this experiment.

Assume that the specific heat capacity of the solution, $c=4.18 \mathrm{JK}^{-1} \mathrm{~g}^{-1}$
Assume that the density of the solution $=1.00 \mathrm{~g} \mathrm{~cm}^{-3}$
$\qquad$ $\mathrm{kJ} \mathrm{mol}^{-1}$

| 0 | 1 | .4 |
| :--- | :--- | :--- | The uncertainty in each of the temperature readings from the thermometer used in this experiment is $\pm 0.1^{\circ} \mathrm{C}$

Calculate the percentage uncertainty in the temperature change in this experiment.
$\qquad$
$\begin{array}{llll}0 & 1 & \mathbf{5} \text { Suggest a change to the student's method, using the same apparatus, that would }\end{array}$ reduce the percentage uncertainty in the temperature change.

Give a reason for your answer.

Change $\qquad$
$\qquad$
Reason $\qquad$
$\qquad$

Suggest the main reason for the difference between this experimental value for the enthalpy of solution and the correct value of $+26 \mathrm{~kJ} \mathrm{~mol}^{-1}$
$\qquad$
$\qquad$

Question 1 continues on the next page

| 0 | 1 | $\mathbf{7}$ | Table 3 shows some entropy data at 298 K |
| :--- | :--- | :--- | :--- |

Table 3

|  | Entropy $\mathbf{S} / \mathbf{J K}^{-1}$ mol $^{-1}$ |
| :---: | :---: |
| $\mathrm{NH}_{4} \mathrm{NO}_{3}(\mathrm{~s})$ | 151 |
| $\mathrm{NH}_{4}{ }^{+}(\mathrm{aq})$ | 113 |
| $\mathrm{NO}_{3}{ }^{-}(\mathrm{aq})$ | 146 |

Calculate a value for the Gibbs free-energy change ( $\Delta G$ ), at 298 K , for the reaction when ammonium nitrate dissolves in water.

$$
\mathrm{NH}_{4} \mathrm{NO}_{3}(\mathrm{~s})+\mathrm{aq} \rightarrow \mathrm{NH}_{4}^{+}(\mathrm{aq})+\mathrm{NO}_{3}^{-}(\mathrm{aq}) \quad \Delta H=+26 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Use data from Table 3 and the value of $\Delta H$ from the equation.
Assume for the solvent, water, that the entropy change, $\Delta S=0$
Explain what the calculated value of $\Delta G$ indicates about the feasibility of this reaction at 298 K

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

| $\mathbf{0}$ | $\mathbf{1}$ | $\mathbf{8}$ | Ammonium nitrate decomposes as shown. |
| :--- | :--- | :--- | :--- |

$$
\mathrm{NH}_{4} \mathrm{NO}_{3}(\mathrm{~s}) \rightarrow \mathrm{N}_{2}(\mathrm{~g})+\frac{1}{2} \mathrm{O}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \quad \Delta H=+123 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

The entropy change $(\Delta S)$ for this reaction is $+144 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}$
Calculate the temperature at which this reaction becomes feasible.
$\qquad$ K

| $\mathbf{0}$ | $\mathbf{2}$ Tschermigite is a hydrated, water-soluble mineral, with relative formula mass of 453.2 |
| :--- | :--- | :--- |

The formula of tschermigite can be represented as $\mathrm{M} . \mathrm{xH}_{2} \mathrm{O}$, where M represents all the ions present.

Table 4 shows its composition by mass.
Table 4

| Element | \% by mass |
| :---: | :---: |
| N | 3.09 |
| H | 6.18 |
| Al | 5.96 |
| S | 14.16 |
| O | 70.61 |

In an analysis, it is found that the mineral contains the ions $\mathrm{NH}_{4}{ }^{+}, \mathrm{Al}^{3+}$ and $\mathrm{SO}_{4}{ }^{2-}$
Calculate the empirical formula of tschermigite and the value of $x$ in M. $x \mathrm{H}_{2} \mathrm{O}$
Describe the tests, with their results, including ionic equations, that would confirm the identities of the ions present.
[6 marks]
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
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$\qquad$
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$\qquad$
$\qquad$


| $\mathbf{0}$ | $\mathbf{3}$ Under suitable conditions, 2-bromobutane reacts with sodium hydroxide to produce a |
| :--- | :--- | :--- | mixture of five products, A, B, C, D and E.

Products A, B and C are alkenes.
$\mathbf{A}$ is a structural isomer of $\mathbf{B}$ and $\mathbf{C}$.
A does not exhibit stereoisomerism.
B and C are a pair of stereoisomers.
Products D and E are alcohols.
D and $\mathbf{E}$ are a pair of enantiomers.

| $\mathbf{0}$ | $\mathbf{3}$. | $\mathbf{1}$ Give the names of the two concurrent mechanisms responsible for the formation of |
| :--- | :--- | :--- | the alkenes and the alcohols.

Mechanism to form alkenes
Mechanism to form alcohols $\qquad$

| $\mathbf{0}$ | $\mathbf{3} .2$ | Define the term stereoisomers. |
| :--- | :--- | :--- |

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

| $\mathbf{0}$ | $\mathbf{3} .3$ | $\mathbf{3}$ Deduce the name of isomer $\mathbf{A}$.....$~$ |
| :--- | :--- | :--- |

Explain why A does not exhibit stereoisomerism.

Name $\qquad$
Explanation $\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{3} .4$ | Outline the mechanism for the reaction of 2-bromobutane with sodium hydroxide to |
| :--- | :--- | :--- | form alkene $\mathbf{A}$


| $\mathbf{0}$ | $\mathbf{3}$. | $\mathbf{5}$ Deduce the name of isomer $\mathbf{B}$ and the name of isomer $\mathbf{C}$. |
| :--- | :--- | :--- |

Explain the origin of the stereoisomerism in $\mathbf{B}$ and $\mathbf{C}$.

Names $\qquad$
$\qquad$
Explanation $\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{3}$ | $\mathbf{6}$ Draw 3D representations of enantiomers $\mathbf{D}$ and $\mathbf{E}$ to show how their structures are |
| :--- | :--- | :--- | :--- | related.


| $\mathbf{0}$ | $\mathbf{3}$. | $\mathbf{7}$ | A student compares the rates of hydrolysis of 1-chlorobutane, 1-bromobutane and |
| :--- | :--- | :--- | :--- | 1-iodobutane.

The suggested method is:

- add equal volumes of the three halogenoalkanes to separate test tubes
- add equal volumes of aqueous silver nitrate to each test tube
- record the time taken for a precipitate to appear in each test tube.

State and explain the order in which precipitates appear.

Order in which precipitates appear $\qquad$
Order in which preciptates appear $\qquad$
$\qquad$
$\qquad$
Explanation $\qquad$
$\qquad$
$\qquad$
$\qquad$


| 0 | $\mathbf{4}$ | Hydrogen peroxide solution decomposes to form water and oxygen. |
| :--- | :--- | :--- |

$$
2 \mathrm{H}_{2} \mathrm{O}_{2}(\mathrm{aq}) \rightarrow 2 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})+\mathrm{O}_{2}(\mathrm{~g})
$$

The reaction is catalysed by manganese(IV) oxide.
A student determines the order of this reaction with respect to hydrogen peroxide. The student uses a continuous monitoring method in the experiment.

The student places hydrogen peroxide solution in a conical flask with the catalyst and uses a gas syringe to collect the oxygen formed. The student records the volume of oxygen every 10 seconds for 100 seconds.

| 0 | $\mathbf{4}$ | $\mathbf{1}$ Explain why the reaction is fastest at the start. |
| :--- | :--- | :--- |

$\qquad$
$\qquad$
$\qquad$
Explain whe reacion fat
$\qquad$

| $\mathbf{0}$ | $\mathbf{4} .2$ The graph in Figure 1 shows how the concentration of hydrogen peroxide changes |
| :--- | :--- | :--- | :--- | with time in this experiment.

Figure 1


Tangents to the curve in Figure 1 can be used to determine rates of reaction.
Draw a tangent to the curve when the concentration of hydrogen peroxide solution is $0.05 \mathrm{~mol} \mathrm{dm}^{-3}$

Use your tangent to calculate the gradient of the curve at this point.
$\qquad$ $\mathrm{mol} \mathrm{dm}{ }^{-3} \mathrm{~s}^{-1}$

## Question 4 continues on the next page

| $\mathbf{0}$ | $\mathbf{4}$ | $\mathbf{3}$ The concentration of hydrogen peroxide solution at time $t$ during the experiment can |
| :--- | :--- | :--- | :--- | be calculated using this expression.

$$
\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]_{t}=\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]_{\text {initial }}\left(\frac{V_{\max }-V_{t}}{V_{\max }}\right)
$$

$\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]_{t}=$ concentration of hydrogen peroxide solution at time $t / \mathrm{mol} \mathrm{dm}^{-3}$
$\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]_{\text {initial }}=$ concentration of hydrogen peroxide solution at the start $/ \mathrm{mol} \mathrm{dm}^{-3}$ $V_{\max }=$ total volume of oxygen gas collected during the whole experiment / $\mathrm{cm}^{3}$ $V_{t}=$ volume of oxygen gas collected at time $t / \mathrm{cm}^{3}$

In this experiment, $V_{\max }=100 \mathrm{~cm}^{3}$
Use Figure 1 and the expression to calculate $\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]_{t}$ when $20 \mathrm{~cm}^{3}$ of oxygen has been collected.
$\qquad$ $\mathrm{mol} \mathrm{dm}^{-3}$

Table 5 shows data from a similar experiment.

## Table 5

| $\left[\mathrm{H}_{2} \mathrm{O}_{2}\right] / \mathrm{mol} \mathrm{dm}^{-3}$ | 0.02 | 0.03 | 0.05 | 0.07 | 0.09 |
| :--- | :--- | :--- | :--- | :--- | :--- |
| Rate $/ \mathrm{mol} \mathrm{dm}^{-3} \mathbf{s}^{-1}$ | 0.00049 | 0.00073 | 0.00124 | 0.00168 | 0.00219 |


| 0 | 4 | 4 | Plot the data from Table 5 on the grid in Figure 2. |
| :--- | :--- | :--- | :--- | Draw a line of best fit.

Figure 2


| 0 | 4 | 5 |
| :--- | :--- | :--- |
| 5 |  |  |

State how the graph shows this order.

Order $\qquad$
How the graph shows this order $\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{5}$ | This question is about catalysis. |
| :--- | :--- | :--- |


| 0 | 5 | 1 |
| :--- | :--- | :--- |

Tetradecane $\left(\mathrm{C}_{14} \mathrm{H}_{30}\right)$ can be cracked to form octane and a cycloalkane.
Give an equation for this reaction.
State the meaning of the term heterogeneous.

Equation

Heterogeneous $\qquad$
$\qquad$
$\qquad$

| 0 | 5 | 2 |
| :--- | :--- | :--- | A student determines the concentration of ethanedioate ions in an acidified solution by titration with potassium manganate(VII) solution.

$$
2 \mathrm{MnO}_{4}^{-}+5 \mathrm{C}_{2} \mathrm{O}_{4}{ }^{2-}+16 \mathrm{H}^{+} \rightarrow 2 \mathrm{Mn}^{2+}+10 \mathrm{CO}_{2}+8 \mathrm{H}_{2} \mathrm{O}
$$

The mixture is warmed before the addition of potassium manganate(VII) solution because the reaction is slow at first. When more potassium manganate(VII) solution is added, the mixture goes colourless quickly due to the presence of an autocatalyst.

Explain the meaning of the term autocatalyst.
Explain, using equations where appropriate, why the reaction is slow at first and then goes quickly.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$


## Section B

Answer all questions in this section.

Only one answer per question is allowed.
For each answer completely fill in the circle alongside the appropriate answer.
CORRECT METHOD WRONG METHODS $\quad \infty \quad \odot \quad \notin$

If you want to change your answer you must cross out your original answer as shown.


If you wish to return to an answer previously crossed out, ring the answer you now wish to select as shown.


You may do your working in the blank space around each question but this will not be marked. Do not use additional sheets for this working.

06 Which atom in the ground state contains at least one unpaired $p$ electron?

A Na
0
B Ne
C O $\square$
D Sc

```
O
```

| 0 | 7 | $C o m p l e t e ~ c o m b u s t i o n ~ o f ~$ |
| :--- | :--- | :--- |
| 0.0100 mol of an organic acid produced 0.0200 mol of |  |  | carbon dioxide.

The same amount of the acid required $20 \mathrm{~cm}^{3}$ of $1.00 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{NaOH}$ (aq) for neutralisation.

Which could be the formula of the acid?

A HCOOH

B $\mathrm{CH}_{3} \mathrm{COOH}$


C HOOCCOOH


D $\mathrm{HOOCCH}_{2} \mathrm{COOH}$ $\square$


A graphite $\square$

B methylbenzene
$\square$
D sodium $\square$

A There are more molecules with any given energy.

C There are more molecules with the most probable energy.

A Chlorine in $\mathrm{Cl}_{2}$ is -1 $\square$

A the concentration of the hydrogen ions
B the pressure of the hydrogen
C the surface area of the platinum electrode
D the temperature of the acid

| 1 | 2 |
| :--- | :--- | Some electrode potential data are shown.

$$
\begin{array}{ll}
\mathrm{Zn}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Zn}(\mathrm{~s}) & E^{\ominus}=-0.76 \mathrm{~V} \\
\mathrm{~Pb}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Pb}(\mathrm{~s}) & E^{\ominus}=-0.13 \mathrm{~V}
\end{array}
$$

Which is a correct statement about this cell?

$$
\mathrm{Zn}(\mathrm{~s})\left|\mathrm{Zn}^{2+}(\mathrm{aq}) \| \mathrm{Pb}^{2+}(\mathrm{aq})\right| \mathrm{Pb}(\mathrm{~s})
$$

A Electrons travel in the external circuit from zinc to lead.

B The concentration of lead(II) ions increases.
C The maximum EMF of the cell is 0.89 V
D Zinc is deposited. $\square$

| 1 | 3 |
| :--- | :--- | Which can not function as a Brønsted-Lowry acid?

A $\mathrm{CH}_{3} \mathrm{COO}^{-}$ $\square$
B $\mathrm{HCO}_{3}{ }^{-}$ $\square$
C $\mathrm{H}_{3} \mathrm{O}^{+}$ $\square$
D $\mathrm{NH}_{4}{ }^{+}$ $\square$

| 1 | 4 |
| :--- | :--- |$\quad$ A strong acid $\mathrm{H}_{2} \mathrm{X}$ dissociates in aqueous solution.

$$
\mathrm{H}_{2} \mathrm{X}(\mathrm{aq}) \rightarrow 2 \mathrm{H}^{+}(\mathrm{aq})+\mathrm{X}^{2-}(\mathrm{aq})
$$

What is the pH of a $0.020 \mathrm{~mol} \mathrm{dm}^{-3}$ solution of this acid?

A 1.00


B 1.40
C 1.70
D 2.00

| 1 | 5 |
| :--- | :--- | Equal volumes of two solutions, each with the same concentration, are mixed together at 298 K

Which two solutions, when mixed, form a solution with a $\mathrm{pH}>7$ ?

A HCOOH and HCOOK


B KOH and $\mathrm{CH}_{3} \mathrm{COOH}$ $\square$
C $\mathrm{NH}_{3}$ and HCl
D $\mathrm{NH}_{4} \mathrm{Cl}$ and KCl $\square$

| 1 | 6 |
| :--- | :--- | Which ionisation needs less energy than this process?

$$
\mathrm{Mg}(\mathrm{~g}) \rightarrow \mathrm{Mg}^{+}(\mathrm{g})+\mathrm{e}^{-}
$$

A $\mathrm{Al}(\mathrm{g}) \rightarrow \mathrm{Al}^{+}(\mathrm{g})+\mathrm{e}^{-}$ $\square$
B $\operatorname{Ar}(\mathrm{g}) \rightarrow \mathrm{Ar}^{+}(\mathrm{g})+\mathrm{e}^{-}$ $\square$
C $\mathrm{Be}(\mathrm{g}) \rightarrow \mathrm{Be}^{+}(\mathrm{g})+\mathrm{e}^{-}$ $\square$
D $\mathrm{Mg}^{+}(\mathrm{g}) \rightarrow \mathrm{Mg}^{2+}(\mathrm{g})+\mathrm{e}^{-}$ $\square$

| $\mathbf{1}$ | $\mathbf{7}$ | Which statement is correct about the Group 1 elements? |
| :--- | :--- | :--- |

A The $\mathrm{Cs}^{+}$ion has a more negative enthalpy of hydration than the $\mathrm{Rb}^{+}$ion.
B The enthalpy of atomisation for potassium is greater than the enthalpy of atomisation for sodium.

C The melting point of potassium is higher than the melting point of $\square$ sodium.
Dhe second ionisation energy of rubidium is lower than the second ionisation energy of lithium.
$\begin{array}{lll}1 & 8 & \text { A test for chloride ions in aqueous solution involves adding dilute nitric acid followed }\end{array}$ by aqueous silver nitrate.

What is the reason for adding the nitric acid?

A To convert $\mathrm{AgNO}_{3}$ into $\left[\mathrm{Ag}\left(\mathrm{NO}_{3}\right)_{2}\right]^{-}$ $\square$
B To decrease the solubility of silver chloride
C To increase the pH of the solution
D To prevent the precipitation of other silver compounds $\square$

| 1 | 9 |
| :--- | :--- | Which pair of reagents reacts to form a tetrahedral complex?

A $\mathrm{CoCl}_{2}(\mathrm{aq})$ and concentrated $\mathrm{NH}_{3}(\mathrm{aq})$ $\square$
B CuSO
C $\mathrm{CuSO}_{4}(\mathrm{aq})$ and sodium ethanedioate(aq) $\square$
D $\mathrm{FeCl}_{3}(\mathrm{aq})$ and concentrated $\mathrm{HCl}(\mathrm{aq})$

| 20 | Cobalt(II) chloride solution changes colour when an excess of concentrated hydrochloric acid is added. |
| :---: | :---: |
|  | What type of reaction takes place? |
|  | A hydrolysis |
|  | B ligand substitution 0 |
|  | C precipitation |
|  | D redox |

A hydrolysis


D redox $\square$

| 2 | 1 |
| :--- | :--- | The reaction between vanadium(IV) ions and manganate(VII) ions in acidic solution can be represented by the equation

$$
5 \mathrm{~V}^{4+}+\mathrm{MnO}_{4}^{-}+8 \mathrm{H}^{+} \rightarrow 5 \mathrm{~V}^{5+}+\mathrm{Mn}^{2+}+4 \mathrm{H}_{2} \mathrm{O}
$$

What volume, in $\mathrm{dm}^{3}$, of $0.020 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{KMnO}_{4}$ is needed to oxidise 0.10 mol of vanadium(IV) ions completely?

A 0.10 $\square$
B 0.50

C 1.0 $\square$
D 5.0 $\square$

22 2-Bromopropane reacts with bromine to form 2,2-dibromopropane.
What is the name of the mechanism of this reaction?

A Electrophilic addition
B Elimination $\square$
C Free-radical substitution
D Nucleophilic substitution

```
O
```



A $\mathrm{CH}_{3} \mathrm{CN}$ $\square$
B $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{NH}_{2}$ $\square$
$\square$


A They are easily oxidised to carboxylic acids by

D They show hydrogen bonding in the liquid state.

A butyl methanoate


C methyl butanoate
$\square$

$\left.\begin{array}{|ccc|}\hline \mathbf{2} \mathbf{9} & \text { Which is the main species present in an aqueous solution of aspartic acid at } \mathrm{pH}=14 \text { ? } \\ \text { [1 mark] }\end{array}\right]$


A $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{Br}$ and $\mathrm{NaOH}(\mathrm{aq})$
$\square$

```
    O
```

A covalent bonds $\square$

B hydrogen bonds
○
$\square$
treacts with hydrochloric acid $\square$


## END OF QUESTIONS







## There are no questions printed on this page

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