## AQA

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 2 Organic and Physical Chemistry

Monday 19 June 2023
Afternoon

## 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.

Time allowed: 2 hours

## Information

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

| Answer all questions in the spaces provided. |  |
| :---: | :---: |
| 0 0 1 | This question is about rates of reaction. |
|  | Figure 1 shows apparatus used to measure the rate of reaction when an acid reacts with an excess of solid sodium hydrogencarbonate, $\mathrm{NaHCO}_{3}$ |
|  | When different monoprotic organic acids are used, the rates at which gas escapes can be used to compare the strengths of the acids. |
|  | A timer is started when the $\mathrm{NaHCO}_{3}$ is added to the acid and the mass of $\mathrm{CO}_{2}$ gas lost is recorded at regular intervals. (It is assumed that any change in mass is due to the loss of $\mathrm{CO}_{2}$ ) |

Figure 1


| $\mathbf{0}$ | $\mathbf{1}$ | $\mathbf{1}$ | Suggest a reason why using a conical flask instead of a beaker would give more |
| :--- | :--- | :--- | :--- | accurate results in this experiment.

$\qquad$
$\qquad$

Figure 2 shows the results of this experiment when $25.0 \mathrm{~cm}^{3}$ of a $2.23 \mathrm{~mol} \mathrm{dm}^{-3}$ solution of ethanoic acid reacts with an excess of $\mathrm{NaHCO}_{3}$

Figure 2


| 0 | 1 | $\mathbf{2}$ | Use Figure 2 to calculate the rate of reaction at 2 minutes. |
| :--- | :--- | :--- | :--- |

Deduce the units of your calculated rate.

Rate $\qquad$ Units $\qquad$

| 0 | 1 | 3 |
| :--- | :--- | :--- |

Sketch, on Figure 2, the curve you would expect when $25.0 \mathrm{~cm}^{3}$ of a $2.23 \mathrm{~mol} \mathrm{dm}^{-3}$ solution of chloroethanoic acid reacts with an excess of $\mathrm{NaHCO}_{3}$

Suggest why chloroethanoic acid is a stronger acid than ethanoic acid.
$\qquad$
$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{2} \quad \mathrm{A}$ and B react together in the presence of an acid catalyst..$~$ |
| :--- | :--- |

$$
A(a q)+2 B(a q) \rightarrow C(a q)+D(a q)
$$

The rate equation for this reaction is

$$
\text { rate }=k[\mathrm{~B}]^{2}\left[\mathrm{H}^{+}\right]
$$

Table 1 shows how the values of the relative initial rate vary with different concentrations of each reagent at the same temperature.

## Table 1

| Experiment | $[\mathbf{A}]$ <br> $/ \mathrm{mol} \mathrm{dm}^{-3}$ | $[\mathrm{B}]$ <br> $/ \mathrm{mol} \mathrm{dm}^{-3}$ | $\left[\mathrm{H}^{+}\right]$ <br> $/ \mathrm{mol} \mathrm{dm}^{-3}$ | Relative initial <br> rate |
| :--- | :---: | :---: | :---: | :---: |
| 1 | 0.40 | 0.20 | 0.10 | 1.00 |
| 2 | 0.50 | 0.20 | 0.10 |  |
| 3 | 0.40 |  | 0.10 | 0.64 |
| 4 | 0.50 | 0.30 | 0.06 |  |


| 0 | 2 | 1 |
| :--- | :--- | :--- |


| $\mathbf{0}$ | $\mathbf{2} .2$ | $\mathbf{2}$ A suggested mechanism for the reaction is shown. |
| :--- | :--- | :--- |

Step $1 \quad \mathrm{~B}+\mathrm{H}^{+} \rightarrow \mathrm{BH}^{+}$
Step $2 \mathrm{BH}^{+}+\mathrm{B} \rightarrow \mathrm{B}_{2} \mathrm{H}^{+}$
Step $3 \quad \mathrm{~B}_{2} \mathrm{H}^{+}+\mathrm{A} \rightarrow \mathrm{C}+\mathrm{D}$
Deduce the rate-determining step for this reaction.
Give a reason for your answer.

Rate-determining step $\qquad$
Reason $\qquad$
$\qquad$
Turn over for the next question Turn over

| $\mathbf{0}$ | $\mathbf{3} \quad$ This question is about intermediates in reaction mechanisms. |
| :--- | :--- |


| 0 | 3 | $\mathbf{1}$ Figure 3 shows an intermediate formed in the first step of a |
| :--- | :--- | :--- | nucleophilic addition-elimination mechanism.

Figure 3


Complete Figure 3 to show the structures of the two reactant species with curly arrows and relevant lone pairs of electrons involved in the formation of the intermediate.

Draw curly arrows and relevant lone pairs of electrons on the intermediate to show how the final products are formed.

| $\mathbf{0}$ | $\mathbf{3} .2$ | Figure $\mathbf{4}$ shows an intermediate formed in the first step of a reaction mechanism of |
| :--- | :--- | :--- | :--- | methylbenzene.

Figure 4


Complete Figure 4 to show the reactant species and any curly arrows involved in the formation of the intermediate.

Draw a curly arrow on the intermediate to show how the product is formed.
Give the name of the reaction mechanism.

Name of mechanism $\qquad$

Figure 5


Complete Figure 5 to show the structure of the intermediate formed with curly arrows involved in its formation.

Give the name of the reaction mechanism.

Name of mechanism




Give the name of the reaction mechanism.

| 0 | 4 | Acrylonitrile, $\mathrm{H}_{2} \mathrm{C}=\mathrm{CHCN}$, can be used as a starting material for the synthesis of |
| :--- | :--- | :--- | butane-1,4-diamine, as shown in this reaction scheme.



Isomer W

$\mathrm{H}_{2} \mathrm{~N}\left(\mathrm{CH}_{2}\right)_{4} \mathrm{NH}_{2}$

| $\mathbf{0}$ | $\mathbf{4} .1$ |
| :--- | :--- |
| $\mathbf{1}$ | Use IUPAC rules to name isomer $\mathbf{W}$. |


| 0 | $\mathbf{4}$ | $\mathbf{2}$ Reaction 1 produces a mixture of $\mathbf{W}$ and two other isomers. |
| :--- | :--- | :--- |

Draw the structures of the two other isomers.
Explain, by considering the mechanism of this reaction, why all three isomers are formed.


The reaction scheme is repeated here.


| $\mathbf{0}$ | $\mathbf{4}$. | $\mathbf{3}$ Identify the reagent that is warmed with isomer $\mathbf{W}$ in reaction 2. |
| :--- | :--- | :--- |

State the other reaction condition needed.

Reagent $\qquad$
Condition $\qquad$

| $\mathbf{0}$ | $\mathbf{4}$. | $\mathbf{4}$ | State the reagent and reaction conditions needed for reaction 3. |
| :--- | :--- | :--- | :--- |

Give an equation for reaction 3.

Reagent and conditions $\qquad$
Equation
$\qquad$

| 0 | $\mathbf{4}$ | $\mathbf{5}$ An incomplete equation for the formation of nylon 4,6 from five molecules of |
| :--- | :--- | :--- | :--- | butane-1,4-diamine and five molecules of hexanedioic acid is shown.



Deduce the values of $x$ and $y$ in this equation.

## Deduce that of $x$ and in

$x$ $\qquad$ $y$ $\qquad$

| $\mathbf{0}$ | $\mathbf{4} .6$ | Figure $\mathbf{6}$ shows a section of the nylon 4,6 polymer molecule. |
| :--- | :--- | :--- | :--- |

Figure 6


Draw, on Figure 6, another section of nylon 4,6 polymer showing two hydrogen bonds between the two sections.


Figure 7 shows the infrared spectrum of $\mathbf{Z}$.
Figure 7

$\begin{array}{llll}\mathbf{0} & \mathbf{5} . & \mathbf{1} \text { Identify the bond that causes the absorption at } 1725 \mathrm{~cm}^{-1} 1\end{array}$

Figure 8 shows the ${ }^{13} \mathrm{C}$ NMR spectrum of $\mathbf{Z}$.
Figure 8

$\begin{array}{lll}\mathbf{0} & \mathbf{5} .2 & \mathbf{2} \text { How many different carbon environments are there in a molecule of } \mathbf{Z} \text { ? }\end{array}$

|  | 5 | 6 | 7 | 8 |
| :--- | :--- | :--- | :--- | :--- |
| Tick $(\checkmark)$ one box |  |  |  |  |


| 0 | 5 | 3 |
| :--- | :--- | :--- | Use Table C in the Data Booklet to help you answer this question.



## Table 2

| Chemical shift ס/ppm | 4.10 | 2.60 | 2.56 | 2.19 | 1.26 |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Integration ratio | 2 | 2 | 2 | 3 | 3 |
| Splitting pattern | quartet | triplet | triplet | singlet | triplet |

Explain what can be deduced from the splitting patterns and chemical shift values for the peaks at $\delta=4.10 \mathrm{ppm}$ and at $\delta=1.26 \mathrm{ppm}$

Deduce the part of the structure of $\mathbf{Z}$ that causes the peaks at $\delta=4.10 \mathrm{ppm}$ and $\delta=1.26 \mathrm{ppm}$

Use Table B in the Data Booklet to help you answer this question.

Peak at $\delta=4.10 \mathrm{ppm}$ $\qquad$
$\qquad$
$\qquad$
Peak at $\delta=1.26 \mathrm{ppm}$ $\qquad$
$\qquad$
$\qquad$

Part of structure

| $\mathbf{0}$ | $\mathbf{5} .5$ | Deduce the part of the structure of $\mathbf{Z}$ that causes the peak at $\delta=2.19 \mathrm{ppm}, ~$ |
| :--- | :--- | :--- |

Part of structure

Figure 9 shows the ${ }^{1} \mathrm{H}$ NMR spectrum of compound $\mathbf{Z}$.
Figure 9


| 0 | $\mathbf{5} .6$ | 6 |
| :--- | :--- | :--- | spectrum in Figure 9, without the information in Table 2 on page 13.

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

| 0 | $\mathbf{5}$. | $\mathbf{7}$ |
| :--- | :--- | :--- |

Do not write outside the box

## Turn over for the next question

| 0 | 6 | $\begin{array}{l}\text { A student plans a series of chemical t } \\ \text { liquids. }\end{array}$ |
| :--- | :--- | :--- |

Liquid J


Liquid $\mathbf{L}$




Liquid $\mathbf{K}$


Liquid $\mathbf{M}$

This is the student's method.
To separate test tubes containing samples of each liquid:
Test 1 add potassium dichromate(VI) solution and warm gently
Test 2 add Fehling's solution and cool in iced water
Test 3 add sodium hydrogencarbonate solution and test any gas produced with a lighted splint

Test 4 add bromine water and shake at room temperature.

| 0 | 6 | 1 | 1 |
| :--- | :--- | :--- | :--- |


| 0 | 6 | 2 | In addition to the missing reagent in Test 1, there is a mistake in the method for two |
| :--- | :--- | :--- | :--- | of the other tests.

State the two mistakes.
Suggest how each of the mistakes should be corrected.

Mistake 1
Suggestion $\qquad$
$\qquad$
Mistake 2
Suggestion $\qquad$
$\qquad$

| 0 | 6 | 3 |
| :--- | :--- | :--- | The missing reagent is added and the mistakes are corrected.

Identify the liquid(s), J, K, L and $\mathbf{M}$, that would react in each test.
State the expected observation for each reaction.

Liquid(s) that react in Test 1
Expected observation $\qquad$
$\qquad$
Liquid(s) that react in Test 2
Expected observation $\qquad$
$\qquad$
Liquid(s) that react in Test 3
Expected observation $\qquad$
$\qquad$
Liquid(s) that react in Test 4
Expected observation $\qquad$
$\qquad$

| 0 | 6 | 4 |
| :--- | :--- | :--- | liquids $\mathbf{K}$ and $\mathbf{M}$ using fractional distillation.

Suggest labels that should be added to positions S, T and U in Figure 10.
Explain why fractional distillation is preferred to simple distillation to separate liquids $\mathbf{K}$ and $\mathbf{M}$.

Figure 10


Label S $\qquad$
Label T $\qquad$
Label U
Explanation $\qquad$
Turn over for the next question

| $\mathbf{0}$ | $\mathbf{7}$ | A gas syringe that does not have any graduations is calibrated using a known mass of |
| :--- | :--- | :--- | box propanone (boiling point $=56.2^{\circ} \mathrm{C}$ ).

The sealed gas syringe contains 0.146 g of propanone $\left(M_{\mathrm{r}}=58.0\right)$ at a temperature of $95^{\circ} \mathrm{C}$ and a pressure of 103 kPa

| 0 | $\mathbf{7}$. | 1 |
| :--- | :--- | :--- |
| Calculate the volume, in $\mathrm{cm}^{3}$, of propanone in the gas syringe. |  |  |

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

Volume of propanone $\qquad$ $\mathrm{cm}^{3}$

| $\mathbf{0}$ | $\mathbf{7} .2$ |
| :--- | :--- | :--- | The gas syringe is then cooled to $75^{\circ} \mathrm{C}$, without changing the pressure.

Calculate the decrease in volume.
(If you were unable to calculate the volume in Question 07.1, you should use the volume $89 \mathrm{~cm}^{3}$. This is not the correct answer.)

| $\mathbf{0}$ | $\mathbf{7}$ | $\mathbf{3}$ The total uncertainty in using the balance to measure the mass of propanone in |
| :--- | :--- | :--- | :--- | Question 07.1 is $\pm 0.001 \mathrm{~g}$

Calculate the uncertainty that this causes in the volume, in $\mathrm{cm}^{3}$, of propanone calculated in Question 07.1.
(If you were unable to calculate the volume in Question 07.1, you should use the volume $89 \mathrm{~cm}^{3}$. This is not the correct answer.)

## Uncertainty

$\qquad$ $\mathrm{cm}^{3}$

| 0 | 7 | 4 | A $600 \mathrm{~cm}^{3}$ sample of propanone is mixed with $2800 \mathrm{~cm}^{3}$ of oxygen in a container at |
| :--- | :--- | :--- | :--- | $60^{\circ} \mathrm{C}$ and 100 kPa . The mixture is ignited.

When the reaction is complete, the remaining mixture of gases is cooled to $60^{\circ} \mathrm{C}$ at 100 kPa

$$
\mathrm{CH}_{3} \mathrm{COCH}_{3}(\mathrm{~g})+4 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow 3 \mathrm{CO}_{2}(\mathrm{~g})+3 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})
$$

Calculate the total volume of the remaining gas mixture.
$\qquad$ $\mathrm{cm}^{3}$

| 0 | 8 | This question is about biofuels. |
| :--- | :--- | :--- |

Palmitic acid, $\mathrm{CH}_{3}\left(\mathrm{CH}_{2}\right)_{14} \mathrm{COOH}$, can be made by hydrolysis of the triester in palm oil under acidic conditions.
Palmitic acid can be used as a biofuel.

| $\mathbf{0}$ | $\mathbf{8} .1$ | $\mathbf{1}$ |
| :--- | :--- | :--- | acidic conditions.



| 0 | $\mathbf{8}$. | $\mathbf{2}$ Palmitic acid burns in air. |
| :--- | :--- | :--- |

In a calorimetry experiment, combustion of 387 mg of palmitic acid increases the temperature of 0.150 kg of water from $23.9^{\circ} \mathrm{C}$ to $37.5^{\circ} \mathrm{C}$

Calculate a value, in $\mathrm{kJ} \mathrm{mol}^{-1}$, for the enthalpy of combustion of palmitic acid in this experiment.
Give your answer to the appropriate number of significant figures.
The specific heat capacity of water is $4.18 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~g}^{-1}$

| $\mathbf{0}$ | $\mathbf{8}$. | $\mathbf{3}$ | State how the value calculated in Question $\mathbf{0 8 . 2}$ is likely to differ from |
| :--- | :--- | :--- | :--- | data book values.

Give one reason, other than heat loss, for this difference.

Difference $\qquad$
Reason $\qquad$
$\qquad$

| 0 | 8 |
| :--- | :--- | $\mathbf{4}$ A sample of a different biofuel, made from sewage sludge, is found to contain $37.08 \%$ carbon, $5.15 \%$ hydrogen and $24.72 \%$ oxygen by mass. The rest of the sample is sulfur.

Calculate the empirical formula of this biofuel.
$\qquad$

| 0 | 8 | 5 | Complete combustion of the biofuel made from sewage sludge produces the |
| :--- | :--- | :--- | :--- | greenhouse gas carbon dioxide.

Suggest one other possible environmental problem with the complete combustion of this biofuel.

State the formula of the pollutant responsible for this problem.

Environmental problem $\qquad$
$\qquad$
Formula $\qquad$


$$
\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6} \rightarrow 2 \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}+2 \mathrm{CO}_{2}
$$

Glucose has the structural formula shown.


Table 3 shows some mean bond enthalpy values.

## Table 3

|  | C-H | C-C | C-O | C=O | O-H |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Mean bond enthalpy <br> $/$ kJ mol $^{-1}$ | 412 | 348 | 360 | 805 | 463 |

Use the equation and the data in Table 3 to calculate an approximate value of $\Delta H$ for the fermentation of glucose. For this calculation you should assume that all the substances are in the gaseous state.
$\qquad$ $\mathrm{kJ} \mathrm{mol}{ }^{-1}$

| $\mathbf{0}$ | $\mathbf{8} .7$ | The carbon dioxide produced from fermentation can be reacted with steam to make |
| :--- | :--- | :--- | :--- | more ethanol.

The equation for this reaction is

$$
2 \mathrm{CO}_{2}(\mathrm{~g})+3 \mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \rightarrow \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}(\mathrm{~g})+3 \mathrm{O}_{2}(\mathrm{~g})
$$

Table 4 shows some standard enthalpies of formation.

## Table 4

|  | $\mathbf{C O}_{2}(\mathbf{g})$ | $\mathbf{O}_{\mathbf{2}}(\mathbf{g})$ | $\mathbf{C}_{\mathbf{2}} \mathbf{H}_{\mathbf{5}} \mathbf{O H}(\mathrm{g})$ | $\mathbf{H}_{\mathbf{2}} \mathbf{O}(\mathrm{g})$ |
| :--- | :---: | :---: | :---: | :---: |
| $\Delta_{\mathrm{f}} \mathrm{H}^{\ominus} / \mathrm{kJ} \mathrm{mol}^{-1}$ | -394 | 0 | -235 | -242 |

Use the data in Table 4 to calculate a standard enthalpy change value for this reaction.
$\qquad$ $\mathrm{kJ} \mathrm{mol}^{-1}$

## Turn over for the next question

| 0 | 9 | This question is about ethanoic anhydride. |
| :--- | :--- | :--- |

In the gas phase, ethanoic anhydride $\left(\mathrm{CH}_{3} \mathrm{CO}\right)_{2} \mathrm{O}$ decomposes to form ethenone.
The equation is


| $\mathbf{0}$ | $\mathbf{9}$. | $\mathbf{1}$ Ethenone is the simplest member of the ketene homologous series. |
| :--- | :--- | :--- |

Ketenes all contain one $\mathrm{C}=\mathrm{C}$ double bond and one $\mathrm{C}=\mathrm{O}$ double bond.
Deduce the general formula for the ketene homologous series.

0 . 9 . 2 Figure 11 shows an incomplete suggested mechanism for the decomposition of ethanoic anhydride.

Figure 11


Complete the mechanism in Figure 11 by adding three curly arrows and any relevant lone pairs of electrons.
[3 marks]

| 0 | 9 | 3 | $F o r ~ a ~ c h e m i c a l ~ r e a c t i o n ~ t h e ~ r e l a t i o n s h i p ~ b e t w e e n ~ t h e ~ r a t e ~ c o n s t a n t, ~$ |
| :--- | :--- | :--- | :--- | , and the temperature, $T$, is shown by the Arrhenius equation.

$$
k=A e^{\frac{-E_{\mathrm{e}}}{R T}}
$$

For the decomposition of gaseous ethanoic anhydride
the activation energy, $E_{\mathrm{a}}=34.5 \mathrm{~kJ} \mathrm{~mol}^{-1}$
the Arrhenius constant, $\mathrm{A}=1.00 \times 10^{12} \mathrm{~s}^{-1}$
At temperature $T_{1}$ the rate constant, $k=2.48 \times 10^{8} \mathrm{~s}^{-1}$
Calculate $T_{1}$
The gas constant, $R=8.31 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}$
$T_{1}$ $\qquad$ K

Question 9 continues on the next page

| 0 | 9 | $\mathbf{4}$ Sketch the Maxwell-Boltzmann distribution of molecular energies for |
| :--- | :--- | :--- | gaseous ethanoic anhydride at temperature $T_{1}$ and at a higher temperature $T_{2}$

Include a label for each axis, and mark on the appropriate axis a typical position for the activation energy.

Explain why the rate of reaction is faster at $T_{2}$
$\qquad$

Explanation
$\qquad$
$\qquad$
There are no questions printed on this page

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