| Surname | Centre Number | Candidate Number |
|---------------|------------------|---------------------|
| First name(s) | | 2 |



GCE A LEVEL

1410U40-1

TUESDAY, 18 JUNE 2024 – MORNING

CHEMISTRY – A2 unit 4 Organic Chemistry and Analysis

1 hour 45 minutes

Section A Section B

| For Exa | aminer's us | e only |
|----------|-----------------|-----------------|
| Question | Maximum Mark | Mark Awarded |
| 1. to 6. | 10 | |
| 7. | 14 | |
| 8. | 15 | |
| 9. | 15 | |
| 10. | 12 | |
| 11. | 14 | |
| Total | 80 | |

ADDITIONAL MATERIALS

- A calculator, pencil and ruler
- · Data Booklet supplied by WJEC

INSTRUCTIONS TO CANDIDATES

Use black ink or black ball-point pen. Do not use gel pen or correction fluid.

You may use a pencil for graphs and diagrams only.

Write your name, centre number and candidate number in the spaces at the top of this page.

Section A Answer all questions.

Section B Answer all questions.

Write your answers in the spaces provided in this booklet. If you run out of space, use the additional page(s) at the back of the booklet, taking care to number the question(s) correctly.

INFORMATION FOR CANDIDATES

The number of marks is given in brackets at the end of each question or part-question.

The maximum mark for this paper is 80.

Your answers must be relevant and must make full use of the information given to be awarded full marks for a question.

The assessment of the quality of extended response (QER) will take place in Q10(a).



[1]

SECTION A

Answer all questions.

1. Phenylmethyl ethanoate, used in perfumery, is made by reacting phenylmethanol and ethanoyl chloride.

Complete the equation for this reaction, showing the structure of the organic product. [1]

2. An azo dye has a maximum absorption of visible light at 403 nm.

(a) Calculate the frequency corresponding to this wavelength.

Frequency = _____Hz

(b) Explain why this dye is yellow in white light. [1]



$$\begin{array}{c} {\rm H_2C} = {\rm CH} - {\rm CH} - {\rm CH_2OCH_3} \\ | \\ {\rm OCH_3} \end{array}$$

compound **D**

compound E

$$\begin{array}{c} \operatorname{CH_3CH_2CH_2CH_2} - \operatorname{C} - \operatorname{CH_2} - \operatorname{OH}_2 \\ || \\ \operatorname{O} \end{array}$$

compound F

compound G

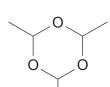
| and at 3200–3500 cm ⁻¹ but not at 1650–1750 cm ⁻¹ . Give reasons for your answer. | [2] |
|--|--|
| | ······································ |

Give the formula of the triphenylmethyl radical.

[1]



5. The structure of paraldehyde is shown below.



Explain what would be seen in the ¹³C NMR spectrum of this compound.

References to the positions of the chemical shifts are **not** required. [2]

6. Ethanal is oxidised by selenium dioxide giving ethanedial.

$$CH_3CHO + SeO_2 \longrightarrow C - C + Se + H_2O_2$$

Calculate the atom economy of this reaction to produce ethanedial.

Give your answer to an **appropriate** number of significant figures. [2]

Atom economy = %

10



| 9 | F | C | TI | 0 | Ν | R |
|---|---|---|----|---|----|---|
| J | _ | v | | U | IA | D |

Answer all questions.

7. (a) Nitrobenzene can be made by the nitration of benzene.

(i) State the reagent(s) used for this reaction.

[1]

(ii) State the **formula** of the nitrogen-containing electrophile that takes part in this reaction.

[1]

.....



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(b) Aromatic nitro-compounds are useful in synthesis as they are often the starting materials for producing other compounds.

Study the reaction sequence below and then answer the questions that follow.

stage 3 room temperature

$$\begin{array}{c|c} O & OH \\ \hline \\ Stage 4 \\ \hline \\ Cr_2O_7^{2-}/H^+ \\ \hline \\ O & OH \\ \end{array}$$

cyclohexadiene-1,4-dione

benzene-1,4-diol

| (i) | State the type of reaction occurring during stage 1. | [1] |
|-------|---|-----|
| (ii) | State the reagent(s) used in stage 2. | [1] |
| (iii) | State the reagent(s) used to produce benzene-1,4-diol in stage 3. | [1] |
| (iv) | State the colour change undergone by the reagents in stage 4. | [1] |



I. State the type of reaction mechanism that occurs when an alkene reacts with bromine.

[1]

Draw the structure of the compound obtained when 1 mol of

cyclohexadiene-1,4-dione reacts with 2 mol of bromine.

[1]

(c) Methyl orange is a water soluble acid-base indicator.

(i) Give the structure of the starting sulfur-containing compound that is coupled with N,N-dimethylphenylamine to give methyl orange. [1]

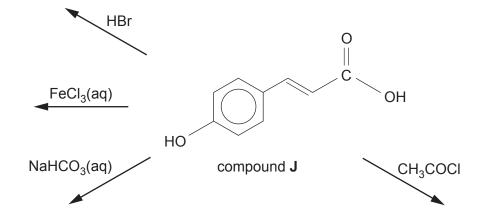
(ii) Suggest why, in basic solution, methyl orange exists as the species below. [1]

$$O = S - O - N = N - O - N(CH_3)$$



(d) Compound **J** is present in relatively large quantities in red peppers.

Study the diagram below and then answer the questions that follow.



(i) State what is seen when compound ${\bf J}$ reacts with iron(III) chloride solution. [1]

(ii) State what is seen when compound **J** reacts with sodium hydrogencarbonate solution. [1]

Give the structure of the compound formed when compound **J** reacts with

(iii) Give the structure of the compound formed when compound **J** reacts with hydrogen bromide. [1]



PMT

(iv) Give the structure of the compound formed when compound ${\bf J}$ reacts with ethanoyl chloride.

[1]

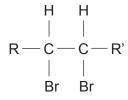
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8. (a) Compound M is an aliphatic compound, which contains two bromine atoms.



compound ${\bf M}$

(i) Compound **M** reacts with an excess of alcoholic potassium hydroxide solution to give compound **N**, which has the formula R—C≡C—R'.

Suggest the role of the alcoholic potassium hydroxide in this elimination reaction.

[1]

.....

(ii) Compound M reacts with aqueous sodium hydroxide to give compound P.



compound P

State the type of reaction mechanism occurring.

[1]



compound P

compound **S**

compound T

The mass spectrum of compound **S** showed a molecular ion at m/z 72.

I. The mass spectrum of compound **S** showed that the alkyl group R was branched.

Deduce the structure of compound **S**. Show your reasoning.

II. Compounds **S** and **T** both reacted with 2,4-dinitrophenylhydrazine to give solid derivatives.

State the colour of these derivatives.

III. The melting temperatures of the derivatives of compounds **S** and **T** in part II above are shown in the table.

| Compound | Melting temperature/°C |
|----------|------------------------|
| S | 185–187 |
| Т | 186–189 |

Describe how the melting temperature of a 50:50 mixture of the two derivatives differs from those shown in the table.

[1]

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Turn over.

[2]

[1]

(b) Pinacolone is prepared from a diol containing 6 carbon atoms.

$$(CH_3)_3C - C - CH_3$$

pinacolone

After the reaction, flammable pinacolone is distilled from the reaction mixture. The distillate consists of two layers – an aqueous layer and a less dense layer containing mainly pinacolone.

- (i) Describe how you would separate these two layers. [1]
- (ii) Describe a chemical test, giving reagent(s) and observations, to show that pinacolone contains a C group. [2]

(iii) The high resolution ¹H NMR spectrum of pinacolone consists of two signals.

$${\rm (CH_3)_3C \longrightarrow C \longrightarrow CH_3}$$

pinacolone

Complete the table below that describes these two signals.

Protons Splitting pattern Relative peak area

CH₃CO (CH₃)₃C



Examiner only

[2]

These three compounds were separated by gas chromatography.

$$\begin{array}{c} \mathsf{CH_3} \\ | \\ \mathsf{H_3C} - \mathsf{C} - \mathsf{CH_2} - \mathsf{CH_3} \\ | \\ \mathsf{CH_3} \end{array}$$

2,2-dimethylbutane

3,3-dimethylbutan-2-ol

$$\begin{array}{c} \mathsf{CH_3} \; \mathsf{O} \\ | \; \; || \\ \mathsf{H_3C} - \mathsf{C} - \mathsf{C} - \mathsf{C} - \mathsf{CH_2} \\ | \; \; \; \\ \mathsf{CH_3} \end{array}$$

pinacolone

 Mary said that the boiling temperature of 3,3-dimethylbutan-2-ol would be the highest of these three compounds and that this could be used to identify it.

Explain why she is correct. [2]

II. The $^{13}\mathrm{C}$ NMR spectra of the remaining two compounds were taken.

Suggest how the ¹³C NMR spectra of these two compounds would be similar and how they would be different.

[2]

Turn over.

Similar

Different

15

13

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| | | | | Examir |
|----|-------------|----------------|--|--------|
| 9. | (a) | Etha | nedioic acid is an important industrial chemical. | only |
| | | | COOH | |
| | | | COOH | |
| | | | COOM | |
| | | oxyg | method for its manufacture is by the reaction of butan-1-ol, carbon monoxide and gen at a pressure of 10 MPa and at a temperature of 100 °C in the liquid phase. process uses a catalyst of palladium mounted on graphite. | |
| | | | first stage gives the ester di-1-butyl ethanedioate, which is then hydrolysed to give nedioic acid and butan-1-ol. | |
| | | (i) | Balance the equation for the first stage of this process. [1] |] |
| | | C ₂ | $_{4}H_{9}OH$ + CO + O_{2} \longrightarrow $COOC_{4}H_{9}$ $COOC_{4}H_{9}$ | |
| | | (ii) | Suggest one reason why this stage of the process may be seen as relatively uneconomic. [1] | I |
| | | (iii) | Hydrolysis of the ester gives ethanedioic acid and butan-1-ol. | |
| | | | Explain why ethanedioic acid is soluble in water but di-1-butyl ethanedioate is virtually insoluble in water. Include a diagram in your answer. [3] |] |
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[3]

| (b) | An older method for manufacturing ethanedioic acid produces insoluble calcium |
|-----|---|
| | ethanedioate (M_r 128) as an intermediate. This is reacted with aqueous sulfuric acid to |
| | produce calcium sulfate and an aqueous solution of ethanedioic acid. |

$$(COO)_2Ca(s) + H_2SO_4(aq) \longrightarrow CaSO_4(s) + (COOH)_2(aq)$$

Ethanedioic acid crystallises as its dihydrate.

In a laboratory experiment 11.52 g of calcium ethanedioate was reacted with aqueous sulfuric acid of concentration 2.00 mol dm⁻³.

(i) Calculate the minimum volume of this aqueous sulfuric acid needed to just react with all the calcium ethanedioate. [2]

Minimum volume =cm³

(ii) After removal of insoluble calcium sulfate, the residue was washed with water. The solution was boiled until $50.0\,\mathrm{cm^3}$ ($50.0\,\mathrm{g}$) of solution remained. The solubility of ethanedioic acid dihydrate, (COOH)₂.2H₂O (M_r 126) is 14.3 g/100 g of solution at 20 °C.

Calculate the mass of ethanedioic acid dihydrate that crystallised from this solution at 20 °C.

Mass = _____g



(c) The removal of alkyl fluorocarbons and CFCs from the environment presents a difficult problem.

One suggestion is to heat these halogenoalkanes with sodium ethanedioate. For example, difluoromethane reacts with sodium ethanedioate, $(COO)_2Na_2$, to give hexafluorobenzene, sodium carbonate (Na_2CO_3) and water.

Write an equation for this reaction.

[2]

+ + + + +

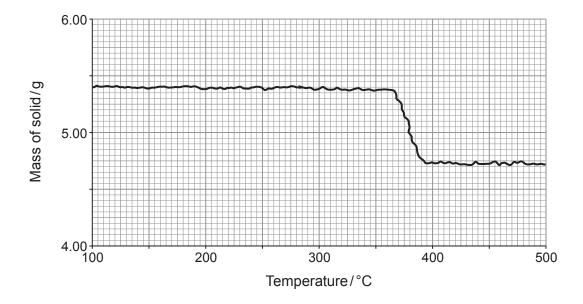
(d) On heating, barium ethanedioate undergoes decomposition.

Use the graph to show that the equation for this reaction is

$$(COO)_2Ba(s)$$
 — Ba $CO_3(s)$ + $CO(g)$

 $M_{\rm r}$ 225 $M_{\rm r}$ 197

[2]



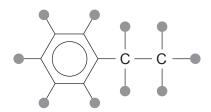


| | | Exa |
|-----|---|-----|
| (e) | Decarboxylation occurs when carboxylic acids are heated with sodalime. | 0 |
| | State the name of the organic compound produced when 4-ethylbenzenecarboxylic acid | |
| | State the name of the organic compound produced when 4-ethylbenzenecarboxylic acid is heated with sodalime. [1] | |
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[6 QER]

10. (a) Compound **B** contains only carbon, hydrogen and bromine. The hydrogen and bromine atoms could be at any of the positions indicated by a circle in the following structure.



- 5.35 g of compound **B** contains 3.74 g of bromine and 0.11 g of hydrogen
- · It rotates the plane of plane polarised light

Explain your reasoning.

• When 4.36 g of compound **B** is heated with aqueous sodium hydroxide, the bromide ions produced react with silver ions to give 4.77 g of silver bromide (M_r 187.9)

Use ${f all}$ of this information to deduce a possible structure for compound ${f B}.$

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| crose | + ethanoic ar | nhydride | sucrose oct | taethanoate + etha | noic acid |
|--------------|---|-----------------------------|--|---|------------------------|
| (i) | A 0.0700 mol s | sample c | of sucrose was used in | a preparation. | |
| | | | n volume of ethanoic ar in this sample of sucros | nhydride needed to read se. | ct with all the [2] |
| | ethanoic anhy | dride | $M_{\rm r}$ | Density/g cm ⁻³ | |
| | ethanole annyt | unue | 102 | 1.08 | |
| | | | | | |
| (;; <u>)</u> | | | | volume = | cm ³ |
| (ii) | | | ducing materials by 'gr | een' methods. | |
| (ii) | Two methods | are prop | ducing materials by 'groosed for making sucros | een' methods. se octaethanoate in the | laboratory. |
| (ii) | Two methods | are prop | ducing materials by 'groosed for making sucrosed the mixture at 139°C | een' methods. se octaethanoate in the using an electric hotpla | laboratory. |
| (ii) | Two methods Method X Method Y Apart from ten | are prop Refluxing Ultrason | ducing materials by 'groosed for making sucros g the mixture at 139°C ic irradiation at room te | een' methods. se octaethanoate in the using an electric hotpla emperature that should be consider | laboratory. ate |
| (ii) | Two methods Method X Method Y Apart from ten | are prop Refluxing Ultrason | ducing materials by 'groosed for making sucros g the mixture at 139°C ic irradiation at room te e, suggest two factors | een' methods. se octaethanoate in the using an electric hotpla emperature that should be consider | laboratory. ate |



| (c) | Lact | ose is a reducing sugar and reacts with Fehling's reagent. | |
|-----|------|---|-----|
| | (i) | State what is seen when lactose reacts with the dark blue solution of Fehling's reagent. | [1] |
| | (ii) | State which functional group must be present in a molecule of lactose for this result with Fehling's reagent. | [1] |
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[2]

11. (a) A method to produce the polyamide **W**, starting from adipamide is shown below.

$$C - (CH_2)_4 - C$$
 NH_2

adipamide

polyamide W

(i) In the first stage adipamide is hydrolysed using aqueous sodium hydroxide, giving disodium hexanedioate as one of the products.

Complete and balance the equation for this stage.

O
$$C - (CH_2)_4 - C$$
 + $NaOH$ \rightarrow $C - (CH_2)_4 - C$ + $Na^+O^ O^-Na^+$

(ii) In another stage, adipamide is reacted with bromine in the presence of alkali to give butane-1,4-diamine, $\rm H_2N(CH_2)_4NH_2$.

$$H_2NOC(CH_2)_4CONH_2$$
 $\xrightarrow{Br_2/NaOH}$ $H_2N(CH_2)_4NH_2$ + inorganic products $M_r 88$

In this stage 0.075 mol of adipamide was used. If the percentage yield of the diamine was 60%, calculate the mass of butane-1,4-diamine produced. [2]

Mass =g



(iii) The sodium salt of hexanedioic acid produced in part (i) is then acidified and the hexanedioic acid produced is reacted with butane-1,4-diamine to give polyamide **W**.

Examiner only

Polyamides contain a peptide link.

Show the peptide link present in the polyamide **on the diagram above**. [1]

(iv) An alternative method for producing hexanedioic acid is to treat adipamide with nitric(III) acid.

$$H_2NOC(CH_2)_4CONH_2 + 2HNO_2 \longrightarrow HOOC(CH_2)_4COOH + 2N_2 + 2H_2O$$

- I. Give **one** advantage of the method using nitric(III) acid. [1]
- II. Suggest how you would know that this alternative reaction using nitric(III) acid is complete. [1]



(b) A tripeptide is formed from three different amino acids.

Its formula (shown as a zwitterion) is

- (i) State why the formation of a peptide from amino acids is called a condensation reaction. [1]
- (ii) Write the formula of the species formed from phenylalanine in acid solution. [1]



Examiner The tripeptide is hydrolysed to give its amino acid fragments. A solution containing these three amino acids is separated using thin layer chromatography. (iii) Use the blank chromatogram to show the position of the 'spot' given by alanine, which has an R_f value of 0.54. Show how you arrived at your answer. solvent front start line cm



(c) A solution contains a mass of **z**g of **one** of the two enantiomers of alanine. It rotates the plane of plane polarised light in the clockwise direction (+) by 6° (+6°).

The other enantiomer rotates the plane of plane polarised light in the anticlockwise direction (–).

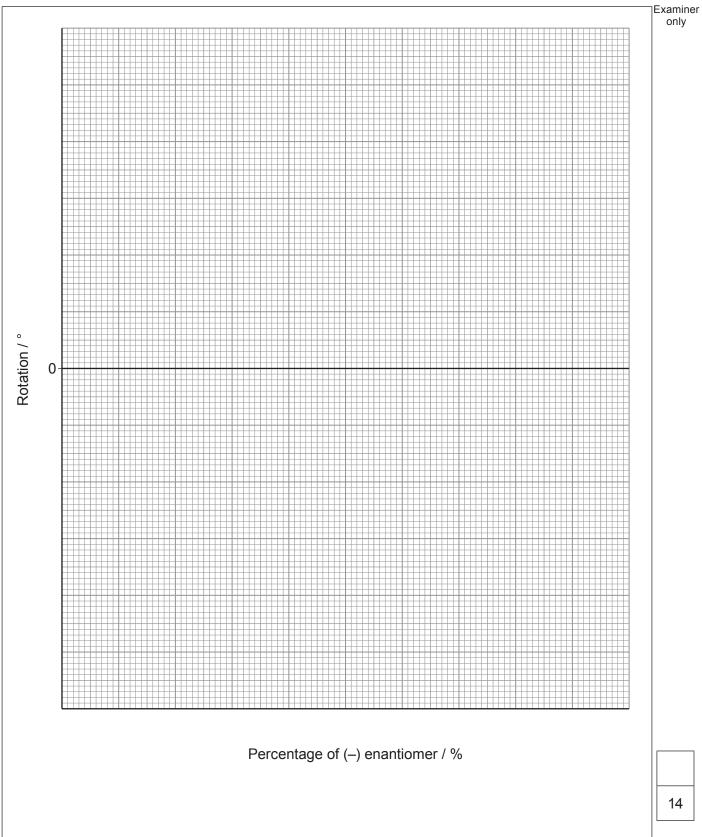
Another solution contains the same total mass zg of a **mixture** of both enantiomers. This mixture rotates the plane of plane polarised light by 1.6° in the anticlockwise direction (-1.6°).

Use this information to plot a graph to show how the rotation changes with the percentage of the (–) enantiomer. Hence determine the percentages of the (+) and (–) enantiomers in the mixture. [3]

Percentage of (+) enantiomer %

Percentage of (-) enantiomer %





END OF PAPER



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