



Cambridge International Examinations

Cambridge International Advanced Level

| CANDIDATE NAME | | | |
|-------------------|----------------------------|---------------------|---------------|
| CENTRE NUMBER | | CANDIDATE NUMBER | |
| CHEMISTRY | | | 9701/41 |
| Paper 4 Structu | red Questions | | May/June 2014 |
| | | | 2 hours |
| Candidates ansv | wer on the Question Paper. | | |

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.

Data Booklet

Do not use staples, paper clips, glue or correction fluid.

DO **NOT** WRITE IN ANY BARCODES.

Section A

Answer all questions.

Additional Materials:

Section B

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.

| For Examiner's Use | | |
|--------------------|--|--|
| 1 | | |
| 2 | | |
| 3 | | |
| 4 | | |
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| 7 | | |
| 8 | | |
| Total | | |
| | | |

This document consists of 19 printed pages and 1 blank page.

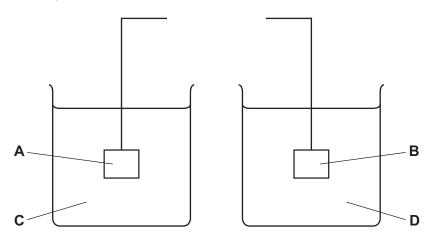


Section A

Answer all the questions in the spaces provided.

| 1 | (a) | (i) | State how the melting point and density of iron compare to those of calcium. | |
|---|-----|------|--|-----|
| | | | melting point of iron: | |
| | | | density of iron: | |
| | | (ii) | Explain why these differences occur. | |
| | | | melting point: | |
| | | | | |
| | | | density: | |
| | | | | |
| | | | | [4] |

- **(b)** The following diagram shows the apparatus used to measure the standard electrode potential, E° , of a cell composed of a Cu(II)/Cu electrode and an Fe(II)/Fe electrode.
 - (i) Finish the diagram by adding components to show the complete circuit. Label the components you add.



(ii) In the spaces below, identify or describe what the four letters **A-D** represent.

| A | |
|---|--|
| В | |
| _ | |
| С | |
| D | |

| | (iii) | Use the <i>Data Booklet</i> to calculate the E° for this cell. |
|-----|----------------------|--|
| | (iv) | Predict how the size of the overall cell potential would change, if at all, as the concentration of solution C is increased. Explain your reasoning. |
| | | |
| | | [8] |
| (c) | forr A ca with | e iron(II) complex ferrous bisglycinate hydrochloride is sometimes prescribed, in capsule m, to treat iron deficiency or anaemia. apsule containing 500 mg of this iron(II) complex was dissolved in dilute $\rm H_2SO_4$ and titrated n 0.0200 mol dm ⁻³ KMnO ₄ . 1 cm ³ of KMnO ₄ solution were required to reach the end point. |
| | The | e equation for the titration reaction is as follows. |
| | | $5Fe^{2+} + MnO_4^- + 8H^+ \rightarrow 5Fe^{3+} + Mn^{2+} + 4H_2O$ |
| | (i) | Describe how you would recognise the end point of this titration. |
| | | |
| | (ii) | Calculate |
| | | • the number of moles of Fe ²⁺ in the capsule, |
| | | the mass of iron in the capsule, |
| | | |
| | | the molar mass of the iron(II) complex, assuming 1 mol of the complex contains 1 mol of iron. |
| | | |

[4]

[Total: 16]

| (1) | State what is meant by the terms: | | | | | | | |
|-------|--|--------|--|--|--|--|--|--|
| | complex, | | | | | | | |
| | | | | | | | | |
| | ligand | | | | | | | |
| (ii) | Two of the complexes formed by coppe Draw three-dimensional diagrams of the | | | | | | | |
| | [Cu(H ₂ O) ₆] ²⁺ | | CuCl ₄ ²⁻ | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| | shape: | | shape: | | | | | |
| (iii) | Platinum forms square-planar complexe | es, in | which all four ligands lie in the same pla | | | | | |
| | the Pt atom. There are two isomeric complexes with Suggest the structures of the two isomerism in organic chemistry, suggestions. | mers | , and, by comparison with a similar ty | | | | | |
| | Structures of isomers: | | type of learners and the first | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| | isomer 1 | | isomer 2 | | | | | |
| | 13011161 1 | 1 | ISOITICI Z | | | | | |

| 0) | - | oper forms two series of compounds, one containing copper(Π) ions and the other containing oper(Π) ions. | | | | | | | |
|----|------|--|---|--------|--|--|--|--|--|
| | (i) | Complete t | he electronic structures of these ions. | | | | | | |
| | | Cu(II) | [Ar] | | | | | | |
| | | Cu(I) | [Ar] | | | | | | |
| | (ii) | Use these | electronic structures to explain why | | | | | | |
| | | copper(II) | salts are usually coloured, | | | | | | |
| | | | | | | | | | |
| | | | | | | | | | |
| | | | | | | | | | |
| | | | | | | | | | |
| | | copper(I) s | alts are usually white or colourless. | | | | | | |
| | | | | | | | | | |
| | | | [t | 5] | | | | | |

(c) Copper(I) oxide and copper(II) oxide can both be used in the ceramic industry to give blue, green or red tints to glasses, glazes and enamels.

The table lists the $\Delta H_{\rm f}^{\rm e}$ values for some compounds.

| compound | $\Delta H_{\rm f}^{\rm e}/{\rm kJmol^{-1}}$ |
|----------------------|---|
| Cu ₂ O(s) | -168.6 |
| CuO(s) | -157.3 |
| $Cu(NO_3)_2(s)$ | -302.9 |
| NO ₂ (g) | +33.2 |

(i) Copper(II) oxide can be produced in a pure form by heating copper(II) nitrate. Use suitable ΔH_f^{\bullet} values from the table to calculate the ΔH^{\bullet} for this reaction.

$$Cu(NO_3)_2(s) \rightarrow CuO(s) + 2NO_2(g) + \frac{1}{2}O_2(g)$$

 $\Delta H^{\circ} = \dots kJ \, \text{mol}^{-1}$

- (ii) Copper(I) oxide can be produced from copper(II) oxide.
 - Use suitable ΔH_f° values from the table to calculate ΔH° for the reaction.

$$2CuO(s) \rightleftharpoons Cu_2O(s) + \frac{1}{2}O_2(g)$$

 $\Delta H^{\circ} = \dots kJ \text{ mol}^{-1}$

• Hence suggest whether a low or a high temperature of oxidation would favour the production of copper(I) oxide. Explain your reasoning.

.....

[4]

[Total: 16]

3 Piperine is the compound responsible for the hot taste of black pepper.

piperine

Piperine is an amide and can be broken down as follows:

(a) Suggest reagents and conditions for this reaction.

.....[1]

(b) (i) How many stereoisomers are there with the same structural formula as piperic acid (including piperic acid itself)?

.....

(ii) Draw the skeletal structure of a stereoisomer of piperic acid, different to the one shown above.

(iii) Suggest structures for the compounds that would be formed when piperic acid is treated with an **excess** of hot concentrated acidified KMnO₄.

| (c) | (i) | Write the expression for $K_{\rm w}$. | |
|-----|-------|--|----------|
| | (ii) | Use your expression and the value of $K_{\rm w}$ in the <i>Data Booklet</i> to calculate the ph 0.150 mol dm ⁻³ NaOH(aq). | H of |
| | | | |
| | (iii) | The pH of a 0.150 mol dm ⁻³ solution of piperidine is 11.9. | |
| | | piperidine | |
| | | Suggest why this answer differs from your answer in (c)(ii). | |
| | (iv) | How would you expect the basicity of piperidine to compare to that of ammonia? Explain your reasoning. | |
| | | | |

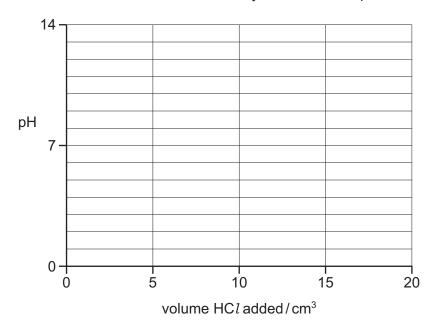
| (d) | 20.0 cm ³ of 0.10 | 0 mol dm ⁻³ | HCl was | slowly | added | to a | $10.0cm^3$ | sample of | of 0.150 r | nol dm ⁻³ |
|-----|------------------------------|------------------------|------------|--------|----------|--------|------------|-----------|------------|----------------------|
| | piperidine. The p | H was mea | sured thro | oughou | t the ad | dition | | | | |

| (i) | Calculate the number | of moles of HC | 1 remaining at | t the end of the | addition |
|-------------|------------------------|----------------|------------------|------------------|----------|
| \! <i>!</i> | Calculate the Hulliber | | t i Cilianing at | | addition |

| | - 4 1 | 101- | | |
|-------|-------|--------|------|------|
| moles | OT I | H(./.= | | |

(ii) Hence calculate the $[H^+]$ and the pH at the end of the addition.

(iii) On the following axes, sketch how the pH will change during the addition of a total of 20.0 cm³ of 0.100 mol dm⁻³ HC *l*. Mark clearly where the end point occurs.



(iv) From the following list of indicators, put a tick in the box by the side of the indicator most suitable for this titration.

| indicator | pH at which colour changes | place one tick only in this column |
|-----------|----------------------------|---|
| А | 0-1 | |
| В | 3-4 | |
| С | 11 - 12 | |
| D | 13-14 | |

[6]

[Total: 16]

4 Noradrenaline is a hormone and neurotransmitter, which is released during stress to stimulate the heart and increase blood pressure.

noradrenaline

| (a) | State the names of three functional groups in the noradrenaline molecule. | | | | | | |
|-----|---|----|--|--|--|--|--|
| | | | | | | | |
| | | | | | | | |
| | r | 3. | | | | | |

(b) (i) Consider the following two-stage synthesis of noradrenaline from dihydroxybenzaldehyde.

- Draw the structure of the intermediate **Z** in the box.
- Suggest reagents for steps 1 and 2.

| step 1 | | | | |
|--------|------|------|------|------|
| sten 2 | | | | |

| (ii) | Dihydroxybenzaldehyde reacts with Br ₂ (aq). | |
|------------------|---|-------------------|
| | Describe what you would see during this reaction. | |
| | Draw the structure of the product. | |
| | | |
| | | [5] |
| (c) Dra | aw the structures of the products when noradrenaline is reacted with dilute NaOH(aq), | |
| (-) | | |
| | | |
| (ii) | dilute HC $l(aq)$, | |
| | | |
| | | |
| <i>,</i> ,,,, | | |
| (iii) | an excess of ethanoyl chloride, CH ₃ COC <i>l</i> . | |
| | | |
| | | [4] |
| (d) Nai | me the new functional groups formed in the reaction in (c)(iii) . | נידן |
| | | |
| | | [2] Total: 14] |
| | L | iotai. 14] |

- 5 The two compounds **V** and **W** are isomers with the molecular formula C₄H₈O, and show the following properties and reactions.
 - Both compounds react with sodium metal, and both decolourise bromine water.
 - Compound V forms a yellow precipitate with alkaline aqueous iodine, whereas compound W does not.
 - When reacted with cold KMnO₄(aq), both V and W produce the same neutral compound X, C₄H₁₀O₃.
 - Both V and W exist as pairs of stereoisomers.

| (a) | Sug | ggest wh | ich functio | nal groups are re | espons | ible for the r | eactions w | ith | |
|-----|-------|-----------|---------------------|-------------------|--------|----------------|-------------|-----|-----|
| | (i) | sodium | , | | | | | | |
| | (ii) | bromine | e water, | | | | | | |
| | (iii) | alkaline | aqueous | iodine. | | | | | |
| (b) | Sug | gest stru | ctures for V | and W . | | | | | [3] |
| | | | | | | | | | |
| | | | | V | | | NA / | | |
| | | | | V | | | W | | [2] |

| (c) | State the type of stereoisostereoisomers. | omerism shown b | y compound V and draw | w the structures of the |
|-----|---|-------------------|------------------------------|-------------------------|
| | type of stereoisomerism | | | |
| | structures of stereoisomers | | | |
| | isomer | 1 | isomer 2 | [2] |
| (d) | Suggest the structure of the | e neutral compour | nd X . | [~] |
| | | | | |
| | | X | | [1] |

[Total: 8]

Section B

Answer all the questions in the spaces provided.

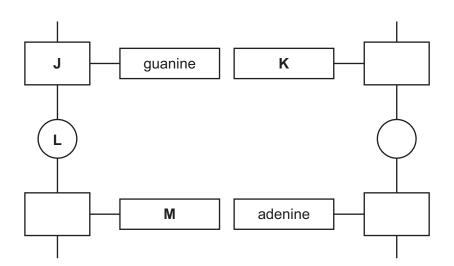
- **6** Proteins and deoxyribonucleic acid, DNA, are two important polymers that occur within living organisms.
 - (a) Proteins have a number of 'levels' of bonding: primary, secondary and tertiary.

 Complete the table to indicate the level of bonding responsible for the features described.

| feature | level of bonding |
|------------------------------|------------------|
| formation of α -helix | |
| formation of disulfide bonds | |
| formation of ionic bonds | |
| linking amino acids | |

[3]

(b) The diagram shows part of a DNA molecule. Study the diagram and give the correct names for the blocks labelled **J**, **K**, **L** and **M**.



| block letter | name |
|--------------|------|
| J | |
| K | |
| L | |
| M | |

[4]

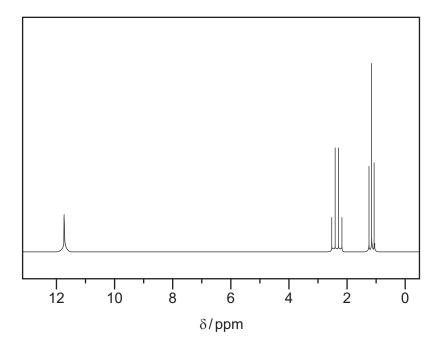
| (c) | | DNA molecule is formed from two polymer strands which are held together until DNA ication occurs. |
|-----|-------|--|
| | (i) | What type of bonding holds the strands together? |
| | | |
| | (ii) | Explain why this type of bonding allows the base pairs within the strands to separate during replication at normal body temperature. |
| | | |
| | | |
| | | [2] |
| (d) | In th | ne polymer RNA, the identities of two of the blocks, J , K , L or M , are different. |
| | For | one of these blocks that are different, give its correct name in DNA and in RNA. |
| | DN | Α: |
| | RN | Δ: |
| | | [1] |
| | | [Total: 10] |

- **7** The combination of mass spectroscopy and NMR spectroscopy provides a powerful method of analysis for organic compounds.
 - (a) The mass spectrum of a compound **G** contains M and M+1 peaks in the ratio of their heights of 74:2.5.

Use these data to calculate the number of carbon atoms present in **G**. Show your working.

[2]

(b) The NMR spectrum of compound **G** is shown.



(i) Use the *Data Booklet* and your knowledge of NMR spectroscopy to identify the type of proton responsible for each of the three absorptions.

| δ/ppm | type of proton |
|-------|----------------|
| 1.1 | |
| 2.2 | |
| 11.8 | |

| (ii) | The addition of D ₂ O causes one of these absorptions to disappear | |
|------|---|--|
| | Explain why this happens and state which absorption is affected. | |

.....

| (iii) | Draw t | he stru | ctural fo | ormula | of (| G |
|-------|--------|---------|-----------|--------|------|---|
|-------|--------|---------|-----------|--------|------|---|

[6]

- (c) Several structural isomers of **G** exist.
 - (i) Draw the structural formula of an isomer of ${\bf G}$ with only two absorptions in its NMR spectrum.

(ii) Use the Data Booklet to suggest where these absorptions would occur.

| peak | δ/ppm |
|------|-------|
| 1 | |
| 2 | |

[3]

[Total: 11]

8 (a) Many common drugs are taken orally, but some medications, such as those based on protein molecules, are injected to prevent them being broken down in the digestive system.

| the stomach. |
|--------------|
| |
| |
| |
| |

Name a functional group present in drug molecules that might be broken down by acid in

(ii) State the *type of reaction* that would cause such a breakdown.

(iii) Which one of the following compounds would not be suitable to be taken orally?

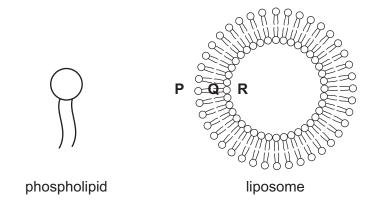
compound

(iv) On the structure of your chosen compound in (iii), circle all the functional groups that might be broken down by acid.

[5]

[Total: 9]

(b) One way of protecting drug molecules that are taken orally is to enclose them in liposomes. These are artificially created spheres made from phospholipids which have an ionic phosphate 'head' and two hydrocarbon 'tails'.



| (i) | State and explain in which location, P , Q or R , a hydrophobic drug could be carried. | |
|------|---|---------|
| | | |
| | | |
| (ii) | By considering the nature of the functional groups in $\bf A$, $\bf B$ and $\bf C$, explain why these drucan be carried at position $\bf R$ in the liposome. | ugs |
| | | |
| | | [2] |
| they | other method of protecting drug molecules is to 'trap' them inside gold nano-cages. Why reach the site where they are needed, such as a tumour, the drug is released by expossite to infra-red radiation. | |
| (i) | Suggest the size of the nano-cages in metres. | |
| | | |
| (ii) | Suggest why infra-red, rather than higher frequency radiation is used. | |
| | | |
| | | |

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(c)

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