

Cambridge International Examinations

Cambridge International Advanced Subsidiary and Advanced Level

| CANDIDATE NAME | | | | | | |
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CHEMISTRY 9701/43

Paper 4 A Level Structured Questions

May/June 2017

2 hours

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.

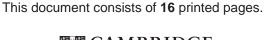
Electronic calculators may be used.

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 | (a) | Describe and explain the variation in the solubilities of the hydroxides of the Group 2 elements. |
|---|-------------|--|
| | | |
| | | |
| | | |
| | | |
| | | |
| | | [4] |
| | The ions | table lists the standard enthalpy changes of formation, $\Delta H_{\rm f}^{\rm e}$, for some compounds and aqueous s. |

| species | $\Delta H_{\rm f}^{\rm e}/{\rm kJmol^{-1}}$ | | | |
|-----------------------|---|--|--|--|
| Ba ²⁺ (aq) | -538 | | | |
| OH⁻(aq) | -230 | | | |
| CO ₂ (g) | -394 | | | |
| BaCO ₃ (s) | -1216 | | | |
| H ₂ O(I) | -286 | | | |

(b) (i) Reaction 1 occurs when $CO_2(g)$ is bubbled through an aqueous solution of $Ba(OH)_2$.

Use the data in the table to calculate the standard enthalpy change for reaction 1, $\Delta H_{\rm r1}^{\rm e}$.

$$Ba(OH)_2(aq) + CO_2(g) \rightarrow BaCO_3(s) + H_2O(l)$$
 reaction 1

 $\Delta H_{r_1}^{\bullet} = \dots kJ \, \text{mol}^{-1} \, [2]$

| If CO ₂ (g) is bubbled through an aqueous solution of Ba(OH) ₂ | for a long time, the precipitated |
|--|-----------------------------------|
| BaCO ₃ (s) dissolves, as shown in reaction 2. | |

$$BaCO_3(s) + CO_2(g) + H_2O(l) \rightarrow Ba(HCO_3)_2(aq)$$
 reaction 2

The standard enthalpy change for reaction 2, ΔH_{r2}^{e} , = -26 kJ mol⁻¹.

| (ii) | Use this information and the data in the table to calculate the standard enthalpy change of |
|------|---|
| | formation of the HCO ₃ -(aq) ion. |

$$\Delta H_{\rm f}^{\bullet} \, HCO_{3}^{-}(aq) = \dots \, kJ \, mol^{-1} \, [2]$$

(iii) The overall process is shown by reaction 3.

Use your answer to (ii), and the data given in the table, to calculate the standard enthalpy change for reaction 3, $\Delta H_{r_3}^{e}$.

$$Ba(OH)_2(aq) + 2CO_2(g) \rightarrow Ba(HCO_3)_2(aq)$$
 reaction 3

$$\Delta H_{r_3}^{\bullet} = \dots kJ \, \text{mol}^{-1} \, [1]$$

(iv) How would the value of $\Delta H_{r_3}^{e}$ compare with the value of $\Delta H_{r_4}^{e}$ for the similar reaction with $Ca(OH)_2(aq)$ as shown in reaction 4? Explain your answer.

$$Ca(OH)_2(aq) + 2CO_2(g) \rightarrow Ca(HCO_3)_2(aq)$$
 reaction 4

(c) The standard entropy change for reaction 1 is $\Delta S_{r_1}^{\bullet}$.

Suggest, with a reason, how the standard entropy change for reaction 3 might compare with $\Delta S_{r_1}^e$.

[Total: 13]

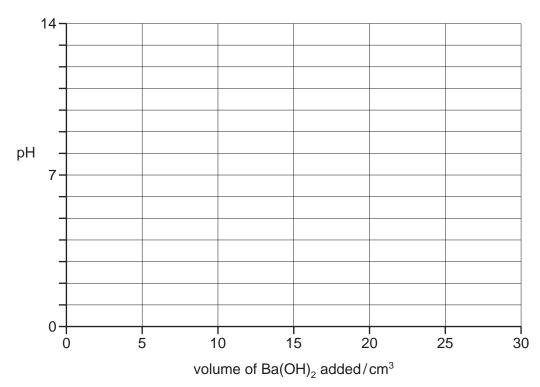
| (i) | Draw 'dot-and-cross' diagrams of these two acids, showing outer shell electrons only. |
|---------|--|
| | HOCN, cyanic acid |
| | HNCO, isocyanic acid |
| (ii) | Suggest the values of the bond angles HNC and NCO in isocyanic acid . |
| , , | HNC NCO |
| (iii) | Suggest which acid, cyanic or isocyanic, will have the shorter C–N bond length. Explain your answer. |
| (b) (i) | Isocyanic acid is a weak acid. |
| | HNCO \rightleftharpoons H ⁺ + NCO ⁻ $K_a = 1.2 \times 10^{-4} \text{ mol dm}^{-3}$ |
| | Calculate the pH of a 0.10 mol dm ⁻³ solution of isocyanic acid. |
| | |
| | pH = |
| (ii) | $pH =$ Sodium cyanate, NaNCO, is used in the production of isocyanic acid. Sodium cyanate is prepared commercially by reacting urea, $(NH_2)_2CO$, with sodicarbonate. Other products in this reaction are carbon dioxide, ammonia and steam. |

- (c) Barium hydroxide, Ba(OH)₂, is completely ionised in aqueous solutions. During the addition of 30.0 cm³ of 0.100 mol dm⁻³ Ba(OH)₂ to 20.0 cm³ of 0.100 mol dm⁻³ isocyanic acid, the pH was measured.
 - (i) Calculate the [OH-] at the end of the addition.

$$[OH^{-}] = \dots moldm^{-3}$$
 [2]

(ii) Use your value in (i) to calculate [H⁺] and the pH of the solution at the end of the addition.

(iii) On the following axes, sketch how the pH changes during the addition of a total of 30.0 cm³ of 0.100 mol dm⁻³ Ba(OH)₂ to 20.0 cm³ of 0.100 mol dm⁻³ isocyanic acid.



| (d) | The | e cyanate ion, NCO ⁻ , can act as a <i>monodentate ligand</i> . |
|-----|------|--|
| | (i) | State what is meant by the terms |
| | | monodentate, |
| | | ligand |
| | | [2] |
| | Silv | rer ions, Ag⁺, react with cyanate ions to form a linear complex. |
| | (ii) | Suggest the formula of this complex, including its charge. |
| | | [2] |
| (e) | | en heated with HC $l(aq)$, organic isocyanates, RNCO, are hydrolysed to the amine salt, H_3Cl , and CO_2 . |
| | | $RNCO \; + \; H_2O \; + \; HC\mathit{l} \; \rightarrow \; RNH_3C\mathit{l} \; + \; CO_2$ |
| | was | .00 g sample of an organic isocyanate, RNCO, was treated in this way, and the CO_2 produced a absorbed in an excess of aqueous $Ba(OH)_2$ according to the equation shown. The solid CO_3 precipitated weighed 1.66 g. |
| | | $Ba(OH)_2(aq) + CO_2(g) \rightarrow BaCO_3(s) + H_2O(l)$ |
| | (i) | Calculate the number of moles of BaCO ₃ produced. |
| | | moles of BaCO ₃ = [1] |
| | (ii) | Hence calculate the M_r of the organic isocyanate RNCO. |
| | | |
| | | $M_{\rm r}$ of RNCO =[1] |
| | | |

| The R group in RNCO a | nd RNH ₃ C <i>l</i> contains | carbon and hydrog | gen only. |
|-----------------------|---|-------------------|-----------|
|-----------------------|---|-------------------|-----------|

| (iii) | Use your M_r value isocyanate RNCO. | calculated in (| (ii) to suggest | the molecular for | mula of the organic |
|-------|---------------------------------------|------------------|----------------------------|--------------------|-----------------------------|
| | | | | | |
| | molecular formula of | RNCO | | | [1] |
| (iv) | Suggest a possible s | structure of the | amine RNH ₂ , v | which forms the an | nine salt, RNH₃C <i>l</i> . |
| | | | | | |

[Total: 23]

[1]

| (a) (i) | Determine the oxidation state of the | | • | |
|---------|---|------------------|----------------------|-------------------|
| (ii) | Name the two types of reaction uncomplex ions. | | | |
| | | | | |
| (iii) | The complex $[Co(NH_3)_4Cl_2]^+$ shows | s isomerism. | | |
| | Draw three-dimensional structures shown here. | of the two isome | ers, and suggest the | type of isomerism |
| | | | | |
| | isomer 1 | | isomer 2 | |
| | type of isomerism | | | |
| | | | | [3] |
| b) (i) | What is meant by the term co-ordi | nation number? | | |
| | | | | [41] |

(ii) Complete the table by predicting appropriate co-ordination numbers, formulae and charges for the complexes C, D, E and F.

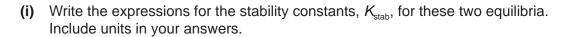
| complex | metal ion | ligand | co-ordination number | formula of complex | charge on complex |
|---------|------------------|---|-------------------------|--|----------------------|
| С | Cr³+ | CN- | | | 3- |
| D | Ni ²⁺ | H ₂ NCH ₂ CH ₂ NH ₂ | 6 | | |
| E | Pt ²⁺ | Cl- | | | 2- |
| F | Fe ³⁺ | -O ₂ C-CO ₂ - | | [Fe(O ₂ CCO ₂) ₃] | |

[6]

| (| c) | Iron(III |) forms con | nplexes in a | separate | reactions | with both | SCN- | ions and | C <i>l</i> −io | ns |
|---|----|----------|-------------|--------------|----------|-----------|-----------|------|----------|----------------|----|
| | | | | | | | | | | | |

$$Fe^{3+}(aq) + SCN^{-}(aq) \rightleftharpoons [FeSCN]^{2+}(aq)$$
 equilibrium 1

$$Fe^{3+}(aq) + 4Cl^{-}(aq) \rightleftharpoons [FeCl_4]^{-}(aq)$$
 equilibrium 2



$$K_{\text{stab1}} =$$

$$K_{\text{stab2}} =$$

(ii) An equilibrium can be set up between these two complexes as shown in equilibrium 3.

$$[FeCl_4]^-(aq) + SCN^-(aq) \rightleftharpoons [FeSCN]^{2+}(aq) + 4Cl^-(aq)$$

equilibrium 3

Write an expression for K_{eq3} in terms of K_{stab1} and K_{stab2} .

$$\mathcal{K}_{\text{eq3}} = \dots$$
 [1]

(iii) The numerical values for these stability constants are shown.

$$K_{\text{stab1}} = 1.4 \times 10^2$$
 $K_{\text{stab2}} = 8.0 \times 10^{-2}$

Calculate the value of K_{eq3} stating its units.

$$K_{\text{eq3}} = \dots$$
 units = \dots [2]

[Total: 19]

4 Carvone occurs in spearmint and a stereoisomer of carvone occurs in caraway seeds. Treating either isomer with hydrogen over a nickel catalyst produces a mixture of isomers with the structural formula **X**.

$$\begin{array}{c|c} & & & \\ \hline \\ & & \\ \hline \\ \text{carvone} \end{array}$$

(a) (i) State the type of stereoisomerism carvone can show. Explain your answer.

....

(ii) Write an equation, using molecular formulae, for this conversion of carvone to X.

.....[2]

X can be synthesised from methylbenzene by the following route.

| | •• | |
|------|---|--------|
| (i) | Name the mechanism in step 1. | |
| | | [1] |
| (ii) | What type of reaction is occuring in the following steps? | |
| | step 3 | |
| | • | |
| | | [2] |
| iii) | Suggest reagents and conditions for each of the following steps. | |
| | step 1 | |
| | step 2 | |
| | step 3 | |
| | \cdot | |
| | | [6] |
| | | Χ. |
| (i) | State the reagents and conditions needed for this reaction. | |
| | | [1] |
| (ii) | Complete the part structure to show the structure of the isomer of ${\bf X}$ that would most like be obtained during this reaction. | эly |
| | Dur The | step 3 |

X

[2]

[Total: 15]

5 Compounds **J**, **K**, **L** and **M** are isomers of each other with the molecular formula C₉H₁₁NO. All four isomers contain a benzene ring.

Two of the isomers contain a chiral centre.

The results of six tests carried out on J, K, L and M are shown in the table.

| test | | observations with each isomer | | | |
|------|---|-------------------------------|------------------------|--------------------|--|
| | | J | K | L | M |
| 1 | add cold HCl(aq) | soluble | soluble | soluble | insoluble |
| 2 | add 2,4-DNPH reagent | orange ppt. | orange ppt. | orange ppt. | no reaction |
| 3 | add NaOH(aq) + I ₂ (aq) | pale yellow ppt. | no reaction | pale yellow ppt. | no reaction |
| 4 | warm with Fehling's solution | no reaction | red ppt. | no reaction | no reaction |
| 5 | heat with NaOH(aq) | no reaction | no reaction | no reaction | $P(C_6H_7N)$ and $Q(C_3H_5O_2Na)$ produced |
| 6 | diazotization and addition of alkaline phenol | no dye produced | orange dye produced | no dye produced | no dye produced |

(a) Use the experimental results in the table above to determine the group(s), in addition to the benzene ring, present in each of the four isomers J, K, L and M.

Complete the table below, identifying the group(s) present in each isomer.

| group(s) in compound | | | | |
|----------------------|---|---|---|--|
| J | К | L | M | |
| | | | | |
| | | | | |
| | | | | |
| | | | | |
| | | | | |

[5]

| (b) (i) | Name the <i>type of reaction</i> occurring i | in test 5 that converts W into P + Q . | [1] |
|-------------------------|---|---|---------|
| (ii) | Suggest structures for compounds P | | [1] |
| | P (C ₆ H ₇ N) | Q (C ₃ H ₅ O ₂ Na) | [2] |
| (c) Iso | mers J , K , L and M all have the molec | cular formula C ₉ H ₁₁ NO. | |
| | e the information in (a) to suggest a st boxes. Draw circles around all chiral | tructure for each of these isomers and draw to centres in K and L . | hese in |
| | | | |
| | J | K | |
| | L | M | |
| | | | [5] |
| cor N c Wh | mpound N is another isomer which hatains a benzene ring. Sontains the same functional group as I sen heated with NaOH(aq), N produces aggest the structure of W . | | |
| | | W | [1] |

[Total: 14]

The rate of this reaction can be studied by measuring the amount of hydroxide ions that remain in solution at a given time. The reaction can effectively be stopped if the solution is diluted with an ice-cold solvent.

- (a) Describe a suitable method for studying the rate of this reaction at a temperature of 40°C, given the following.
 - a solution of 0.10 mol dm⁻³ 1-chloro-1-phenylethane, labelled A
 - a solution of 0.10 mol dm⁻³ sodium hydroxide, labelled B
 - 0.10 mol dm⁻³ HC*l*
 - volumetric glassware
 - ice-cold solvent
 - stopclock

| access to standard laboratory equipment and chem |
|--|
|--|

| [4 |
|----|

(b) The rate of this reaction was measured at different initial concentrations of the two reagents. The table shows the results obtained.

| experiment | [C ₆ H ₅ CHC <i>l</i> CH ₃] /moldm ⁻³ | [OH ⁻] /moldm ⁻³ | relative rate |
|------------|---|--|------------------|
| 1 | 0.05 | 0.10 | 0.5 |
| 2 | 0.10 | 0.20 | 1.0 |
| 3 | 0.15 | 0.10 | 1.5 |
| 4 | 0.20 | 0.15 | to be calculated |

| (i) | Deduce the order of reaction with respect to each of $[C_6H_5CHClCH_3]$ and $[OH^-]$. Explain your reasoning. |
|-----|--|
| | order with respect to [C ₆ H ₅ CHClCH ₃] |
| | |
| | order with respect to [OH-] |
| | order with respect to [OTT] |
| | |

[2]

| (ii) | Write the rate equation for this reaction, stating the units of the rate constant, <i>k</i> . |
|--------------|--|
| | $rate = \dots mol dm^{-3} s^{-1}$ |
| | units of $k = \dots$ |
| (iii) | [1] Calculate the relative rate for experiment 4. |
| | |
| | relative rate for experiment 4 =[1] |
| (c) (i) | Use your answers in (b)(i) to help you to draw the mechanism for the reaction of 1-chloro-1-phenylethane with hydroxide ions, including the following. |
| | all relevant lone pairs and dipoles curly arrows to show the movement of electron pairs the structures of any transition state or intermediate |
| | |
| | |
| | |
| | |
| | rol |
| /:: \ | [3] |
| (ii) | This reaction was carried out using a single optical isomer of 1-chloro-1-phenylethane. |
| | Use your mechanism in (i) to predict whether the product will be a single optical isomer or a mixture of two optical isomers. Explain your answer. |
| | |
| | [1] |

(d) The proton NMR spectrum of a sample of 1-phenylethanol shows four peaks: a multiplet for the C_6H_5 protons and three other peaks as shown in the table. When the sample is shaken with D_2O and the proton NMR spectrum recorded, **fewer** peaks are seen.

Complete the table for the proton NMR spectrum of 1-phenylethanol, $C_6H_5CH(OH)CH_3$. Use of the *Data Booklet* might be helpful.

| δ/ppm | number of ¹ H atoms responsible for the peak | group responsible for the peak | splitting pattern | result on shaking with D ₂ O |
|---------|---|--------------------------------|-------------------|---|
| 1.4 | | | | |
| 2.7 | | | | |
| 4.0 | | | | |
| 7.2-7.4 | 5 | C ₆ H ₅ | multiplet | peak remains |

[4]

[Total: 16]

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