

MINISTRY OF EDUCATION, SINGAPORE in collaboration with CAMBRIDGE INTERNATIONAL EDUCATION General Certificate of Education Advanced Level

CANDIDATE NAME						
CENTRE NUMBER	S			INDEX NUMBER		

012345678

CHEMISTRY 9476/03

Paper 3 Structured Questions

For examination from 2026

SPECIMEN PAPER 2 hours

You must answer on the question paper.

You will need: Data booklet

INSTRUCTIONS

- Section A: answer all questions.
- Section B: answer one question.
- Use a black or dark blue pen. You may use an HB pencil for any diagrams or graphs.
- Write your name, centre number and index number in the boxes at the top of the page.
- Write your answer to each question in the space provided.
- Do not use an erasable pen. Do not use correction fluid or tape.
- Do not write on any bar codes.
- You may use an approved calculator.

INFORMATION

- The total mark for this paper is 75.
- The number of marks for each question or part question is shown in brackets [].

This document has **24** pages. Any blank pages are indicated.



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Section A

Answer all the questions in this section in the spaces provided.

1

(a)	Sta	ate what is meant by dynamic equilibrium.	
At h	nigh t	temperatures, steam and carbon undergo the reaction shown.	
		$H_2O(g) + C(s) \rightleftharpoons H_2(g) + CO(g)$ $\Delta H^{\ominus} = +131 \text{ kJ mol}^{-1}$	
(b)		an experiment, steam at a pressure of 2.00 atm and a temperature of 1000 K is introto a vessel containing an excess of powdered carbon.	duced
	At e	equilibrium, the partial pressure of hydrogen is 1.40 atm.	
	(i)	Write an expression for $K_{\rm p}$ for this reaction. State its units.	
	(ii)	Calculate the partial pressures of $\rm H_2O(g)$ and $\rm CO(g)$ at equilibrium. Hence, calculate total pressure at equilibrium.	ate the

[3]

(iii) Use your answer from 1(b)(ii) to calculate the value of $K_{\rm p}$.

[1]

(c)		separate experiment, the same amount of carbon was used, but in the form of lumps er than powder. All other conditions were kept the same.
	(i)	State how this change would affect the time taken to reach equilibrium. Explain your answer.
		[1
	(ii)	State how this change would affect both the position of equilibrium and the numerical value of $K_{\rm p}$.
		[1
(d)	Car	bon monoxide is poisonous.
	-	lain how carbon monoxide affects the function of haemoglobin in the transport of oxygen ne human body.
		[2

- (e) Carbon monoxide can be 'inserted' into some organic molecules by different types of addition reactions.
 - 1 the reaction of an alcohol with CO, for example the synthesis of ethanoic acid from methanol

2 the reaction of an alkene with CO and H₂O, for example the synthesis of propanoic acid from ethene

$$C_2H_4 + CO + H_2O \rightarrow CH_3CH_2CO_2H$$

3 the reaction of an aldehyde with CO and H₂O, for example the synthesis of hydroxyethanoic acid from methanal

$$CH_2O + CO + H_2O \rightarrow CH_2(OH)CO_2H$$

(i) A, B and C can be synthesised by the addition reactions described.

(CH ₃) ₂ CHCH ₂ CO ₂ H	C ₆ H ₅ CH(OH)CO ₂ H	(CH ₃) ₃ CCO ₂ H
Α	В	С
55	naterials for the synthesis of A , be, 1, 2 or 3, for the synthesis of	
		[3]

(ii) Figure 1.1 shows the conversion of **B** into the diacid **F**, $C_6H_5C(OH)(CO_2H)_2$, in three steps.

D forms an orange precipitate with 2,4-DNPH.

Figure 1.1

State the reagents and conditions you would use for steps 1, 2 and 3.

Suggest the structures of compounds **D** and **E**.

	re:
 	 [5]

[Total: 20]

2 (a) An electrochemical cell is set up as shown in Figure 2.1.

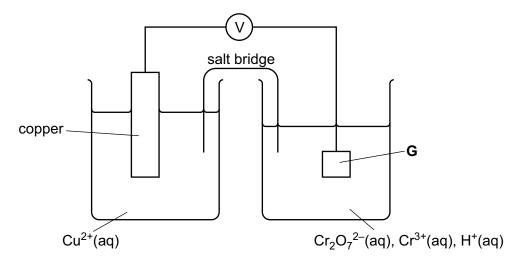


Figure 2.1

Table 2.1 lists relevant electrode potentials for some electrode reactions.

Table 2.1

electrode reaction	E [⊕] /V
$Cr_2O_7^{2-} + 14H^+ + 6e^- \Rightarrow 2Cr^{3+} + 7H_2O$	+1.33
Cu ²⁺ + 2e ⁻ ⇌ Cu	+0.34

(i) Suggest a suitable material for the electrode labelled **G** in Figure 2.1.

.....[1]

(ii) Describe **one** condition specified by the symbol $^{\circ}$ when the electrode potentials shown in Table 2.1 are measured at 298 K.

.....

.....[1]

(iii) For the cell drawn in Figure 2.1, use the data in Table 2.1 to calculate the $E_{\rm cell}^{\ominus}$ for this electrochemical cell.

(iv) Construct the overall equation for the reaction that occurs in the cell when a current is allowed to flow.

.....[1]

(v) Calculate a value of ΔG^{\ominus} for the cell reaction, using the $E_{\text{cell}}^{\ominus}$ you calculated in **2(a)(iii)**, and your overall equation in **2(a)(iv)**.

[2]

(b) A sample of finely ground copper is contaminated with zinc powder.

An excess of H⁺(aq) is added to this sample and all the Zn is oxidised to Zn²⁺.

The reaction forms 126 cm 3 of hydrogen gas, measured at a temperature of 300 K and a pressure of 1.00 \times 10 5 Pa.

$$Zn(s) + 2H^{+}(aq) \rightarrow Zn^{2+}(aq) + H_{2}(g)$$

The mixture is filtered to remove the copper. Copper reacts with acidified manganate(VII) ions, MnO_4^- (aq).

 4.88×10^{-3} mol of acidified $\mathrm{MnO_4}^{-}(\mathrm{aq})$ is required for all the Cu to be oxidised to Cu^{2^+} .

$$5Cu(s) + 2MnO_4^{-}(aq) + 16H^{+}(aq) \rightarrow 5Cu^{2+}(aq) + 2Mn^{2+}(aq) + 8H_2O(I)$$

Calculate the masses of zinc and copper present in the sample.

			8	
(c)		iqueous solution, dichi wn in equilibrium 2.1.	omate(VI) ions exist in equ	ilibrium with chromate(VI) ions, as
		equilibrium 2.1	$\text{Cr}_2\text{O}_7^{\ 2-}(\text{aq}) + \text{H}_2\text{O}(\text{I}) \rightleftharpoons \text{orange}$	2CrO ₄ ²⁻ (aq) + 2H ⁺ (aq) yellow
	(i)	When aqueous sodic yellow. Explain why.	ım hydroxide is added, the	solution changes colour from orange to
				[1]

to orange.

Use Le Chatelier's Principle to state and explain what you can conclude about the enthalpy change of the forward reaction for equilibrium 2.1.

The resulting solution from 2(c)(i) is cooled. The colour changes from yellow back

[2]

(iii) Barium chromate(VI), BaCrO $_4$, is sparingly soluble, while barium dichromate(VI), BaCr $_2$ O $_7$, is soluble.

Aqueous barium nitrate, $Ba(NO_3)_2(aq)$, is added to an equilibrium mixture consisting of $Cr_2O_7^{\ 2-}(aq)$ and $CrO_4^{\ 2-}(aq)$.

State the effect on the equilibrium position of equilibrium 2.1 and the pH of the resulting solution. Explain your answer.

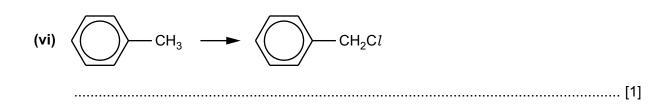
.....[2]

[Total: 15]

3	(a)	Halogeris can behave as Lewis acids of Lewis bases.
		Use the reactions of chlorine with aluminium chloride and iodine with iodide ions to form
		triiodide ions, ${\rm I_3}^-$, to illustrate this behaviour. Write equations for the reactions that occur.

• • • •
\sim

- **(b)** There are several ways of introducing chlorine atoms into organic molecules. For each of the following reactions, state the reagents and conditions necessary.
 - (i) $C_2H_4 \rightarrow C_2H_5Cl$
 - (ii) $C_2H_4 \rightarrow C_2H_4Cl_2$
 - (iii) $C_2H_5OH \rightarrow C_2H_5Cl$ [1]



(c)		anes are generally considered to be unreactive compounds, showing an inertness to muon reagents such as NaOH, $\rm H_2SO_4$ and $\rm K_2Cr_2O_7$.
	(i)	Suggest a reason why these reagents do not react with an alkane such as CH ₄ .
		[1]
	(ii)	Ethane can be converted into chloroethane according to reaction 3.1.
		reaction 3.1 $C_2H_6 + Cl_2 \rightarrow C_2H_5Cl + HCl$
		Name the type of mechanism in reaction 3.1.
		[1]
	(iii)	Use equations to describe the mechanism for reaction 3.1. Name each step involved.
		[3]

(d) It is found by experiment that during reactions of alkanes with chlorine, hydrogen atoms of the primary, secondary and tertiary carbons are replaced by chlorine atoms at different rates, as shown in Table 3.1.

Table 3.1

reaction	relative rate
$RCH_3 \rightarrow RCH_2Cl$	1
$R_2CH_2 \rightarrow R_2CHCl$	7
$R_3CH \rightarrow R_3CCl$	21

When 2-methylpropane is chlorinated, a mixture of two organic products, **H** and **I**, is formed.

Use the information in Table 3.1 and the number of hydrogen atoms of each type (primary, secondary or tertiary) within the molecule to predict the relative ratio of **H** and **I** formed.

......[1]

(e) Draw the skeletal formulae of **four** different structural isomers of C₅H₁₁C*l* that could be obtained from the chlorination of 2-methylbutane.

Indicate any chiral centres in your structures by an asterisk (*).

[3]

(f) (i) Figure 3.1 shows the polymer poly(ethene), which can be used in food plastic wrapping.

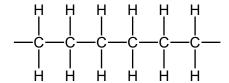


Figure 3.1

This polymer macromolecule has a relative molecular mass of between 15 000 and 20 000.

Calculate the minimum number of repeating units of ethene needed to be present on one macromolecule of poly(ethene).

[1]		
	Suggest why people are encouraged to recycle poly(alkenes).	(ii)
[2]		
[Total: 20]		

Section B

Answer **one** question from this section in the spaces provided.

4

(a)	Exp	plain what is meant by a transition element.	
			 [1]
(b)	Sug	ggest a reagent that could be used to convert $[Cu(H_2O)_6]^{2+}$ into $[CuCl_4]^{2-}$ in the laborate	
(c)	The	e complex [Cu(H ₂ O) ₆] ²⁺ is octahedral.	[1]
		an octahedral complex, the d subshell of a transition metal ion is split into two sets of ergy levels.	
	(i)	On Figure 4.1, draw one of the d orbitals at the lower energy level in an octahedral complex.	[1]
	(ii)	On Figure 4.2, draw one of the d orbitals at the upper energy level in an octahedral complex.	
		$\frac{z}{y}$	
		Figure 4.1 Figure 4.2	[1]
	(iii)	Use your d orbital diagrams in 4(c)(i) and 4(c)(ii) to explain why d orbital splitting occuin an octahedral complex.	
			 [1]

(d)	Copper can form stable compounds containing $copper(II)$ ions or $copper(I)$ ions.				
	$\label{lem:copper} Copper(II) \ salts \ are \ usually \ coloured, \ whereas \ copper(I) \ salts \ are \ usually \ white \ or \ colourless.$				
		complex ion $[Cu(H_2O)_6]^{2^+}$ is blue, whereas the complex ion $[Mn(H_2O)_6]^{3^+}$ is red. Both of se complexes are octahedral.			
	(i)	State the electronic configuration of a copper(I) ion.			
	(ii)	Explain why copper(II) salts are usually coloured, whereas copper(I) salts are usually white or colourless.			
		[3			
	(iii)	Suggest why $[Cu(H_2O)_6]^{2+}$ and $[Mn(H_2O)_6]^{3+}$ have different colours.			
		[1			
	(iv)	Draw the three-dimensional diagram for [Cu(H ₂ O) ₆] ²⁺ .			

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

Table 4.1 lists the $\Delta H_{\rm f}^{\ominus}$ values for some compounds.

Table 4.1

compound	∆H _f [⊕] /kJ mol ⁻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, as shown in reaction 4.1.

reaction 4.1
$$Cu(NO_3)_2(s) \rightarrow CuO(s) + 2NO_2(g) + \frac{1}{2}O_2(g)$$

Calculate ΔH^{\oplus} for reaction 4.1, using suitable $\Delta H^{\oplus}_{\rm f}$ values from Table 4.1.

[1]

(ii) Copper(I) oxide can be formed by heating copper(II) oxide, as shown in reaction 4.2.

reaction 4.2
$$2\text{CuO}(s) \neq \text{Cu}_2\text{O}(s) + \frac{1}{2}\text{O}_2(g)$$
 $\Delta H^{\circ} = +146 \text{ kJ mol}^{-1}$

Suggest which of the reactions, reaction 4.1 or reaction 4.2, would have the higher entropy change per mole of reactant. Explain your answer.

(f) Some copper compounds are used as reagents for organic reactions.

A mixture of copper(II) ethanoate and lead(IV) ethanoate is used for the conversion of carboxylic acids into alkenes containing one less carbon atom.

When carboxylic acid $\bf J$, $\rm C_8H_{14}O_2$, is treated in this way, alkene $\bf K$, $\rm C_7H_{12}$, is formed. Treatment of $\bf K$ with a hot acidified solution of manganate(VII) ions produces compound $\bf L$, $\rm C_7H_{12}O_3$.

A series of tests are carried out on \mathbf{L} . The tests and the results are recorded in Table 4.2.

Table 4.2

test on L	observations / results	type of reaction	functional groups identified in L
addition of Na ₂ CO ₃ (aq)	effervescence	acid-base	carboxylic acid
addition of 2,4-DNPH	orange precipitate formed		
addition of alkaline aqueous iodine, followed by acid	yellow precipitate and organic compound formed, HO ₂ CCH ₂ CH(CH ₃)CH ₂ CO ₂ H		

(i) Complete Table 4.2 to give the type of reaction described and what the information tells you about the functional groups present in **L**. [4]

(ii) Suggest structures for compounds $\boldsymbol{J},\,\boldsymbol{K}$ and $\boldsymbol{L}.$

[3]

[Total: 20]

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(a) (i)	State the Arrhenius theory of acids and bases.	
	[2]	
(ii)	Separate samples of $\rm Na_2O$ and MgO are shaken with water. The pH value of the solution formed with $\rm Na_2O$ is greater than the pH value of the solution formed with MgO.	
	The pH of the solution formed when SO_3 is shaken with water is less than both of these.	
	Explain these observations using the Arrhenius theory. Write equations for all the reactions described.	
	[3]	

- (b) The ozone, ${\rm O_3}$, molecule contains a dative covalent bond.
 - (i) Draw a 'dot-and-cross' diagram to show the bonding in a molecule of ${\rm O_3}$. Include all outer shell electrons in your diagram.

	(11)	Suggest a value for the bond angle in O_3 . Explain your answer.
		[3]
(c)		compound FO_2 does not exist, but CIO_2 does. In the molecule of CIO_2 the central atom nlorine.
	Sug	gest reasons for this difference, by considering the possible types of bonding in each of two compounds.
		·
		[2]

(d) Ozone can be made by passing oxygen gas through a tube between two highly charged electrical plates. This forms an equilibrium mixture of the two gases O₂ and O₃.

$$3O_2(g) \rightleftharpoons 2O_3(g)$$

The concentration of O₃ in the mixture can be determined by its reaction with aqueous KI.

$$O_3$$
 + 2KI + $H_2O \rightarrow I_2$ + O_2 + 2KOH

The iodine formed can be estimated by its reaction with sodium thiosulfate.

$$2Na_2S_2O_3 + I_2 \rightarrow Na_2S_4O_6 + 2NaI$$

In an experiment, 500 cm³ of an oxygen / ozone gaseous mixture at a temperature of 300 K is passed into an excess of aqueous KI.

The iodine formed required 15.0 $\rm \,cm^3$ of 0.100 $\rm \,mol\,\,dm^{-3}\,\,Na_2S_2O_3$ for complete reaction.

(i) Calculate the number of moles of iodine produced.

[1]

(ii) Calculate the concentration of ozone gas.

[1]

(e) (i) Compounds $\bf M$ and $\bf N$ each have the formula $\bf C_9 \bf H_{10}$. Compound $\bf P$ has the formula $C_9H_{12}O$. Compound **M** reacts with $Br_2(aq)$, but compound **N** does **not**.

Compound **P** reacts with alkaline aqueous iodine.

In separate reactions, M, N and P are all oxidised by hot concentrated alkaline KMnO₄ and then acidified. The product of each of these oxidation and acidification reactions is benzene-1,2-dicarboxylic acid, Q.

$$CO_2H$$
 CO_2H

Suggest structures for compounds M, N and P.

(ii) Compound **S** can be synthesised from compound **Q** in two steps as shown in Figure 5.1.

$$CO_2H$$
 $Step 1$ R $Step 2$ NH R S

Figure 5.1

Suggest reagents for step 1 and for step 2.

Suggest the structure of the intermediate **R**.

13

(f) Figure 5.2 shows the structure of 2-hydroxypropanoic acid.

Figure 5.2

Draw the structure of a polyester that can be formed from 2-hydroxypropanoic acid. Show at least two repeat units.

[1]

[Total: 20]