



Cambridge International Examinations

Cambridge International Advanced Subsidiary and Advanced Level

CANDIDATE NAME						
CENTRE NUMBER				CANDIDATE NUMBER		

CHEMISTRY 9701/42

Paper 4 A Level Structured Questions

February/March 2018

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)	(i)	State how the solubilities of the hydroxides of the Group 2 elements vary down the group.
			[1]
		(ii)	Explain the factors that are responsible for this variation.
			[3]
	(b)	The	e solubility of $Sr(OH)_2$ is 3.37×10^{-2} mol dm ⁻³ at 0 °C.
		(i)	Write an expression for the solubility product of Sr(OH) ₂ .
			$K_{\rm sp}$ =
			[1]
		(ii)	Calculate the value of $K_{\rm sp}$ at 0 °C. Include units in your answer.
		()	ς μ · · · · · · · · · · · · · · · · · ·
			\mathcal{K}_{sp} = units = [2]

(c)	Metal peroxides contain the ¬O–O¬ ion. The peroxides of the Group 2 elements, MO₂, decompose on heating to produce a single gand the solid oxide, MO, only.								
	(i)	Write an equation for the thermal decomposition of strontium peroxide, SrO ₂ .							
		[1]							
	(ii)	Suggest how the temperature at which thermal decomposition of MO ₂ occurs varies down Group 2.							
		Explain your answer.							
		[3]							
(d)	(i)	The ethanedioates of the Group 2 elements, MC_2O_4 , decompose on heating to produce a mixture of two different gases and the solid oxide, MO, only.							
		Complete the equation for the thermal decomposition of barium ethanedioate.							
		$BaC_2O_4 \to \dots + \dots + \dots + \dots $							
	(ii)	Describe two observations you would make during the reaction when ethanedioic acid, $H_2C_2O_4$, is warmed with acidified manganate(VII) ions.							
		[2]							
		[Total: 14]							
		-							

2	(a)	Describe the trend in the reactivity of the halogens Cl_2 , Br_2 and I_2 as oxidising agents. Explain this trend using values of E° (X_2/X^-) from the <i>Data Booklet</i> .					
	(b)	(i)	Write an equa	tion for the reaction between chlorine and	d water.		
		/ii\	Llee standard		eta Booklet to calculate the E ^e for		
		(ii)	the following i	electrode potential, E° , data from the Da eaction.	ata bookiet to calculate the E cell for		
				$Cl_2 + 2OH^- \rightleftharpoons Cl^- + ClO^- + H$	₂ O		
					E _{cell} = V [2]		
		(iii)	The [OH ⁻] was	increased and the $E_{\mbox{\tiny cell}}$ was measured.			
				he value of the E_{cell} measured would comck (\checkmark) in the table.	pare to the $E_{\mathrm{cell}}^{\circ}$ calculated in (ii) by		
				$E_{\rm cell}$ becomes less positive than $E_{\rm cell}^{\circ}$.			
				E_{cell} stays the same as $E_{\text{cell}}^{\bullet}$.			
				$E_{\rm cell}$ becomes more positive than $E_{\rm cell}^{\rm e}$.			
			Explain your a	answer.			
					[2]		

(c)	A h	half-equation involving bromate(V) ions, BrO_3^- , and bromide ions is shown.							
		$BrO_3^-(aq) + 3H_2O(I) + 6e^- \iff Br^-(aq) + 6OH^-(aq) \qquad E^{\circ} = +0.58V$							
	(i)	An alkaline solution of chlorate(I), ClO^- , can be used to oxidise bromide ions to bromate(V) ions.							
		Use the Data Booklet and the half-equation shown to write an equation for this reaction.							
		[41]							
		[1]							
	(ii)	Calculate the E_{cell}^{\bullet} for the reaction in (i).							
		<i>E</i> ^o _{cell} = ∨ [1]							
((iii)	When a concentrated solution of $bromic(V)$ acid, $HBrO_3,$ is warmed, it decomposes to form bromine, oxygen and water only.							
		Write an equation for this reaction. The use of oxidation numbers may be helpful.							
		[1]							
		[Total: 10]							

3	(a)	Hydrogen	cyanide,	HCN,	is a	weak	acid	in a	queous	solution.
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$$HCN(aq) \rightleftharpoons H^{+}(aq) + CN^{-}(aq)$$
 $K_a = 6.2 \times 10^{-10} \,\text{mol dm}^{-3}$

(i) Calculate the pH of 0.10 mol dm⁻³ HCN(aq).

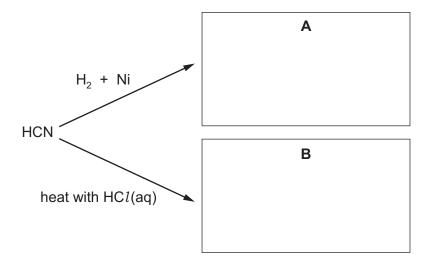
(ii) Draw a 'dot-and-cross' diagram to represent the bonding in the hydrogen cyanide molecule. Show the outer shell electrons only.

[1]

(iii) State the hybridisation of the carbon and nitrogen atoms in hydrogen cyanide, and give the H–C–N bond angle.

hybridisation of C	
hybridisation of N	
H–C–N bond angle	
	[2]

(iv) Suggest structures for the organic products **A** and **B** in the following reactions. Assume that HCN reacts in a similar way to RCN.



[2]

(b) Add	ding a m	easured quantity	y of KCN to a so	lution of NiC l_2 pr	oduces the complex	$\times [Ni(CN)_2Cl_2]^x$.
	(i)	Deduc	e the overall cha	arge, x, on this c	omplex.		
						x =	[1]
		e comple kel ion.	ex can exist as t	wo separate iso	mers with the sa	ame geometry (sha	pe) around the
	(ii)	State t	he type of isome	erism shown by t	hese isomers.		
							[1]
	(iii)	If brom	ide ions are pre	sent in the solut	ion, the complex	[Ni(CN) ₂ ClBr] ^x can	form.
		Assum of isom	ing that [Ni(CN) ners of [Ni(CN) ₂ 0	$_{2}ClBr]^{x}$ has the s $ClBr]^{x}$ that could	same geometry a exist, and draw t	as $[Ni(CN)_2Cl_2]^x$, statheir structures in the	ate the number ne box.
		• nu	mber of isomers	of [Ni(CN) ₂ ClB	r]×		
			struct	ures of the isom	ers of [Ni(CN) ₂ Cl	[Br] ^x	
							[3]
							[3]
(c)				CN is gradually s of reactions ta		tion of NiSO ₄ until	the KCN is in
Nico	k	(CN	•	more KCN	Б	KCN in excess	-
NiSO	4		C green precipitat	e	D yellow solution		E red solution
	•	None of C contact The K:	cidation state of of C, D or E contains no potassium in E is 3	ains sulfur. m. 2:1.	change during th	ese reactions.	
	Use	e the info	ormation to sug	gest the formulae	e of C , D and E .		
	С.						
	D.						
	Ε.						
							[3]
							[Total: 15]

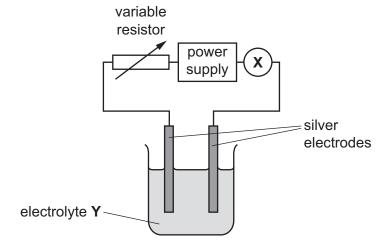
1	(a)	(i)	State what is meant by the term partition coefficient.
		An	monia is soluble in both water and organic solvents. aqueous solution of ammonia is shaken with the immiscible organic solvent trichloromethane. e mixture is left to reach equilibrium.
		Sar	mples are taken from each layer and titrated with dilute hydrochloric acid.
		•	A 25.0 cm³ sample from the trichloromethane layer requires $13.0\mathrm{cm}^3$ of $0.100\mathrm{moldm}^{-3}$ HC l to reach the end-point. A $10.0\mathrm{cm}^3$ sample from the aqueous layer requires $12.5\mathrm{cm}^3$ of $0.100\mathrm{moldm}^{-3}$ HC l to reach the end-point.
		(ii)	Calculate the partition coefficient, $K_{\rm partition}$, of ammonia between trichloromethane and water.
			$K_{\text{partition}} = \dots $ [2]
		(iii)	Butylamine, C ₄ H ₉ NH ₂ , is also soluble in both water and organic solvents.
			Suggest how the value of $K_{\rm partition}$ of butylamine between trichloromethane and water would compare to the value of $K_{\rm partition}$ calculated in (ii). Explain your answer.

.....[2]

(b) (i)	Explain why butylamine is basic.
	[1]
(ii)	Write an equation to show butylamine reacting as a base.
	[1]
(iii)	State how the basicity of butanamide, C ₃ H ₇ CONH ₂ , compares to that of butylamine.
	[1]
(iv)	State a reagent for the conversion of butanamide into butylamine.
	[1]
	[Total: 9]

(a)	(i)	Complete the e	lectronic configurati	on of a chromium a	itom.	
		1s ² 2s ² 2p ⁶ 3s ²				[1]
	(ii)	State the two h	ighest oxidation sta	tes of chromium co	mmonly seen in its	compounds.
	()				•	
				•••••		[1]
(b)	of a	aqueous NH ₃ , ad		concentrated aque	nd N , are formed weous HC <i>l</i> are sepa	
				reagent		
		solution	an excess of NH ₃ (aq)	an excess of NaOH(aq)	an excess of concentrated HCl(aq)	
		Cu ²⁺ (aq)	Н	J	K	
		Co²+(aq)	L	M	N	
	(i)	нк			lexes.	
		IVI				[2]
	(ii)	Write the formu	lae of the following	compounds or com	plexes.	
		L				
		N				
						[2]
	(iii)	State the appear	rance of compound	1 J .		
						[1]
						[Total: 7]

6 The apparatus shows a cell which can be used to determine a value of the Avogadro constant, L.



(a)	(i)	Name	com	pone	nt	X	ζ.
----	---	-----	------	-----	------	----	---	----

		[1]
(ii)	Suggest a suitable electrolyte Y.	
		[1]

(b) In an experiment, a current of 0.200 A was passed through the cell for 40.0 minutes. The mass of the silver cathode increased by 0.500 g.

The charge on the electron is -1.60×10^{-19} C.

Calculate the:

- number of moles of silver deposited on the cathode
- number of coulombs of charge passed
- number of electrons passed
- number of electrons needed to deposit 1 mol of silver at the cathode.

[3]

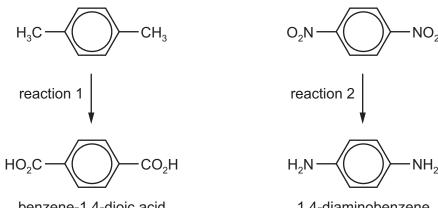
7 (a) (i)	Complete the equations to show the two types of polymerisation. Draw one repeat unit for each polymer. Include any other products.
	addition polymerisation
n CH₂=C	CHCH ₃ (g) →
	condensation polymerisation
nHO ₂ CCH ₂ + nHOCH ₂ CI	$CO_2H(s)$ \rightarrow $H_2OH(I)$
	[3]
(ii)	Suggest the sign of the entropy changes, ΔS° , for each of the two types of polymerisation. Explain your answers.
	 ΔS^e for addition polymerisation
	 ΔS° for condensation polymerisation
	[2]

(b) An amide bond forms when a carboxylic acid reacts with an amine.

	(i)	Complete the equation by writing the products in the box.
R–CO₂H	1 +	$H_2N-R' \rightarrow$
		[1]
	(ii)	Use your answer to (i) to work out the bonds that are broken and the bonds that are formed during the reaction between a carboxylic acid and an amine.
		bonds that are broken
		bonds that are formed
		[2]
((iii)	Use bond energy values from the <i>Data Booklet</i> to calculate the enthalpy change, $\Delta H^{\rm e}$, when one mole of amide bonds is formed in the reaction in (i).
		Δ <i>H</i> ° = kJ [2]
(c)	Am	de bonds can also be formed by reacting acyl chlorides with amines.
	The	enthalpy change for this process, $\Delta H^{\rm e}$, is $-6.00{\rm kJmol^{-1}}$.
	Cal 298	culate the minimum entropy change, ΔS^e , for this reaction to be spontaneous (feasible) at K.
		$\Delta S^{e} = \dots J K^{-1} \text{mol}^{-1} [2]$

The repeat unit of the polyamide *Kevlar* is shown.

(d) The monomers of Kevlar, benzene-1,4-dioic acid and 1,4-diaminobenzene, can be synthesised as follows.



benzene-1,4-dioic acid

1,4-diaminobenzene

State the reagents and conditions needed for:

O) reaction	

......[1]

reaction 2 (ii)

(e) Kevlar is both strong and rigid.

Complete the table to identify two intermolecular forces and the groups involved which are responsible for these properties of Kevlar.

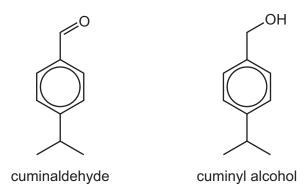
intermolecular force	group(s) involved

[2]

[Total: 17]

Question 8 starts on the next page.

8 (a) Cumin is a spice used to flavour food. Two compounds responsible for its flavour are cuminaldehyde and cuminyl alcohol.



(i)	Deduce th	he	number	of	peaks	that	would	be	present	in	the	¹³ C	NMR	spectrum	of
	cuminyl al	lcoh	nol.												

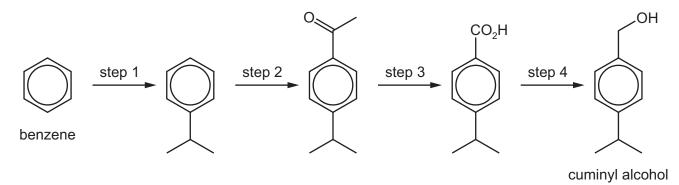
number of	peaks	[1]	1
	pound	ъ.	1

(ii) Identify **two** bonds that are responsible for the differences in the infra-red spectra of cuminaldehyde and cuminyl alcohol, and state their absorption ranges.

	absorption range in the infra-red spectrum/cm ⁻¹						
bond responsible for the difference	cuminaldehyde	cuminyl alcohol					

[2]

(b) Cuminyl alcohol can be synthesised from benzene by the following route.



(i)	Suggest reagents and conditions for steps 1–4.	
	step 1	
	step 2	
	step 3	
	step 4	 [4]
		[ד]
(ii)	Name the mechanism of step 2 and state the type of reaction in step 4.	
	mechanism of step 2	
	type of reaction in step 4	
		[2]

[Total: 9]

- **9 (a)** Two molecules of compound **F** react together under suitable conditions to form compound **G**. Some information about compounds **F** and **G** is given.
 - The mass spectrum of **F** has a peak due to the molecular ion at m/e = 106, and a peak at m/e = 107 with an abundance 8% of the 106 peak.
 - The mass spectrum of **G** has a peak due to the molecular ion at m/e = 212, and major peaks at m/e = 91 and m/e = 105.
 - Both **F** and **G** contain oxygen and are neutral compounds which are insoluble in water.
 - The ¹H NMR spectrum of **F** includes a singlet peak at δ = 10.0 due to one proton.
 - The ¹H NMR spectrum of **G** includes a singlet peak at δ = 5.1 due to two protons.
 - When **G** is heated with dilute sulfuric acid, benzoic acid, C₆H₅CO₂H, and phenylmethanol, C₆H₅CH₂OH, are produced.

Use this information to answer (i)-(vi).

(i)	Explain how the mass spectrum of F shows that a molecule of F contains seven carbon
	atoms. Show your working.

]	[1]						
(ii)	Suggest the molecular formula of the fragment of G at m/e = 91.								
		[1]						
(iii)	Suggest the molecular	formulae of F and G .							
	F								
	G								
			2]						
(iv)	Suggest structures for	compounds F and G .							
	F	G							

(v) On the structures you have drawn in (iv), circle the protons responsible for the 1H NMR peaks at $\delta = 10.0$ in **F** and $\delta = 5.1$ in **G**. [1]

[2]

	(vi)	State the type of reaction	that G undergoes when he	ated with dilute sulfuric acid.	
					[1]
(b)	Des	scribe and explain the relativ	e acidities of benzoic acid,	phenylmethanol and 4-methylpl	nenol.
	<	CO ₂ H	CH ₂ OH	H_3C OH	
		benzoic acid	phenylmethanol	4-methylphenol	
					[3]

(c) The ester 4-methylphenyl benzoate is used in the manufacture of perfumes.

4-methylphenyl benzoate

Suggest a **two**-step route for the synthesis of 4-methylphenyl benzoate from 4-methylphenol and benzoic acid.

Include reagents and conditions for each step, and the structure of the intermediate compound.

[3]

[Total: 14]

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