



## **Cambridge International Examinations**

Cambridge International Advanced Subsidiary and Advanced Level

CANDIDATE NAME			
CENTRE NUMBER		CANDIDATE NUMBER	
CHEMISTRY			9701/42
Paper 4 A Level Struc	ctured Questions		May/June 2016
			2 hours
Candidates answer or	n 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** questions in the spaces provided.

1	ma In t	gnesium nitrate, $Mg(NO_3)_2$ , is very soluble in water. When a hot saturated solution of gnesium nitrate is cooled, crystals of the hydrate, $Mg(NO_3)_2$ .6H <sub>2</sub> O, are formed. he crystals, six water molecules bond to each $Mg^{2+}$ ion, and some of these water molecules <b>also</b> bonded to the nitrate ions.
	(i)	Suggest the type of bonding that occurs between
		H <sub>2</sub> O and Mg <sup>2+</sup> ,
		H <sub>2</sub> O and NO <sub>3</sub> <sup>-</sup> .
	(ii)	Describe the arrangement of the water molecules around the Mg <sup>2+</sup> ion.
		[1]
	(iii)	Describe in detail what you would observe when crystals of $Mg(NO_3)_2$ .6H <sub>2</sub> O are heated in a boiling tube, gently at first and then more strongly. Write equations for any reactions that occur.
		[4]
	(iv)	Calculate the percentage <b>loss</b> in mass when $Mg(NO_3)_2.6H_2O$ is heated strongly to constant mass.

percentage loss = ..... % [2]

(b)	Explain why the Group 2 nitrates become more stable to heat down the group.
	[2]
(c)	Magnesium nitrate and silver nitrate, $AgNO_3$ , decompose on heating to produce the same gases. Silver nitrate also produces silver metal during decomposition.
	Write an equation for the decomposition of AgNO <sub>3</sub> .
	[1]
	[Total: 12]

(a) ⊏x	plain what is mea	ant by the	e term <i>weak</i> acid.		
<b>(b)</b> Th	e p $K_a$ values of f	our acids	are listed below.		
		acid	structural formula	p <i>K</i> <sub>a</sub>	
		1	CH₃CO₂H	4.8	
		2	CH <sub>3</sub> CH <sub>2</sub> CO <sub>2</sub> H	4.9	
		3	CH <sub>3</sub> CHC <i>l</i> CO <sub>2</sub> H	2.8	
		4	CH <sub>2</sub> ClCH <sub>2</sub> CO <sub>2</sub> H	4.0	
(i) (ii)			y, explain the difference		cid dissociation constant
	<ul><li>acid 1 and</li></ul>	acid 2,			
	acid 2 and	acid 3,			

[3]

(c)	(i)	Draw a fully labelled diagram of the equipment needed to measure the voltage of an electrochemical cell consisting of the standard hydrogen electrode and the standard $\text{Cu/Cu}^{2+}$ electrode.
		[4]
	(ii)	For the cell drawn in (i), calculate the $E_{\text{cell}}^{\text{e}}$ and state which electrode is positive.
		$E_{\text{cell}}^{\text{e}}$ = identity of the positive electrode
(d)	A m	nonobasic acid, <b>D</b> , has $K_a = 1.23 \times 10^{-5}  \mathrm{mol  dm^{-3}}$ .
	(i)	Calculate the pH of a 0.100 mol dm <sup>-3</sup> solution of <b>D</b> .
		pH =[2]
	(ii)	An electrochemical cell similar to the one you have drawn in <b>(c)(i)</b> was set up using a $0.100\mathrm{moldm^{-3}}$ solution of <b>D</b> in the hydrogen electrode instead of the standard solution.
		Use the data and the Nernst equation, $E = E^{\circ} + 0.059 \log [H^{+}(aq)]$ , to calculate the new $E_{cell}$ in this experiment.
		E <sub>cell</sub> = V [2]
		[Total: 14]

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(a) 2-bromopropane can be used to synthesise methylethylamine and 2-methylpropylamine.

	$\rightarrow$ NH <sub>2</sub>	
	reaction 1 methylethylamine  Br	
2-br	omopropane	
	reaction 2  reaction 3  NH <sub>2</sub> 2-methylpropylamine	
(i)	Draw the structure of the intermediate <b>X</b> in the box above.	[1]
(ii)	Suggest reagents and conditions for	
	reaction 1,	
	reaction 2,	
	reaction 3.	
		[3]
(b) (i)	Write an equation showing why aqueous solutions of ethylamine are alkaline.	
(ii)	Compare the basicities of ethylamine and ammonia. Explain your answer.	[1]

(c)	Sol	Solutions containing mixtures of amines and their salts are buffer solutions.				
	(i)	Explain what is meant by the term <i>buffer solution</i> .				
	(ii)	Write two equations to show how a solution containing a mixture of CH <sub>3</sub> NH <sub>2</sub> and CH <sub>3</sub> NH <sub>3</sub> C <i>l</i>				
	(11)	acts as a buffer.				
		[2]				
		[Total: 10]				

(i)	Draw diagram	s to show the three-c	limensional strud	ctures of the two is	omers.
	is	somer 1		isomer 2	
					<u> </u>
(ii)	Comment on	the polarity of the two	isomers of Pt(N	$(\mathrm{IH_3})_2\mathrm{C}l_2$ . Explain yo	our answer.
Ox	xaloplatin is ano		cancer drug in v		
Ox	xaloplatin is ano	other successful anti- ne same as that in Pt	cancer drug in v		
Ox	xaloplatin is ano atinum atom is th	other successful anti- ne same as that in Pt	cancer drug in v $(NH_3)_2Cl_2$ . $H_2$ $O$ $C$ $H_2$ $O$ $C$		

b)	Onl	y one structure of the	e complex [Ni( $R_3P$ ) <sub>2</sub> C $l_2$ ] is known. (R = C	CH <sub>3</sub> , R <sub>3</sub> P is a monodentate ligand)
	(i)	What does this ind	icate about the stereochemistry around	d the nickel atom?
				[1]
	(ii)	Draw a three-dime	nsional diagram showing the structure	of this complex.
				[1]

[Total: 6]

Cadmium ions form complexes with primary amines and with 1,2-diaminoethane.

5

$Cd^{2+}(aq) + 4CH_3NH_2(aq) \rightleftharpoons [Cd(CH_3NH_2)_4]^{2+}(aq)$	$K_{\mathrm{stab}}$ = 3.6 $ imes$ 10 <sup>6</sup>	equilibrium I
$Cd^{2+}(aq) + 2H_2NCH_2CH_2NH_2(aq) \rightleftharpoons [Cd(H_2NCH_2CH_2NH_2)_2]^{2+}(aq)$	$K_{\rm stab} = 4.2 \times 10^{10}$	equilibrium II

(a) (i) Write an expression for the stability constant,  $K_{\text{stab}}$ , for equilibrium I, and state its units.

Cadmium ions are poisonous and need to be removed from some water supplies. This is often done by adding a complexing agent.

(ii) In a sample of ground water the concentration of  $Cd^{2+}(aq)$  is  $1.00 \times 10^{-4}$  mol dm<sup>-3</sup>.

Calculate the concentration of  $CH_3NH_2(aq)$  needed to reduce the concentration of  $Cd^{2+}(aq)$  in this dilute solution by a factor of one thousand.

concentration of  $CH_3NH_2(aq) = \dots mol dm^{-3}$  [2]

[2]

(b) Values for  $\Delta H^{\circ}$  and  $\Delta G^{\circ}$  for equilibria I and II, and the value of  $\Delta S^{\circ}$  for equilibrium I, are given in the table below. All values are at a temperature of 298 K.

equilibrium	ΔH <sup>o</sup> / kJ mol <sup>-1</sup>	∆G° /kJ mol⁻¹	ΔS <sup>e</sup> /JK <sup>-1</sup> mol <sup>-1</sup>
I	-57.3	-37.4	-66.8
II	-56.5	-60.7	to be calculated

(i)	Suggest a reason why the $\Delta H^{e}$ values for the two equilibria are very similar.
(ii)	
(iii)	$\Delta S^{\circ} = J K^{-1}  \text{mol}^{-1}   [1]$ Suggest a reason for the difference between the $\Delta S^{\circ}$ you have calculated for equilibrium II and that for equilibrium I given in the table.
(iv)	
	[1]

**6** Esterases are enzymes that hydrolyse esters.

$$R - C + H_2O \rightarrow R - C + HO-R'$$

Enzymes can be quite specific in the structures of the substrates they act upon. For example, an esterase isolated from the mould *Aspergillus niger* will hydrolyse phenyl ethanoate,  $CH_3CO_2C_6H_5$ , but not its isomer methyl benzoate,  $C_6H_5CO_2CH_3$ .

(a)	Outline how enzymes catalyse reactions, and explain their specificity
	Use diagrams in your answer where appropriate.

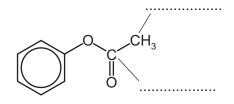
.....

- **(b)** Sample bottles of each of the isomers phenyl ethanoate and methyl benzoate have lost their labels and so have been named isomer **A** and isomer **B**.
  - (i) The carbon-13 NMR spectra of isomers **A** and **B** contain the following peaks.

isomer A	isomer B
δ 52	δ 26
δ 128	δ 122
δ 129	δ 126
δ 130	δ 129
δ 133	δ 151
δ 167	δ 169

The identity of the compound responsible for each spectrum can be deduced by studying the chemical shifts ( $\delta$ ) of the peaks in the spectra.

Use the *Data Booklet* to assign the correct peaks to the labelled carbon atoms in the structures of the isomers below. Write each value next to the relevant carbon atom and hence deduce the identity of each isomer.



C CH<sub>3</sub>

phenyl ethanoate is isomer .....

methyl benzoate is isomer .....

[2]

(ii) These two isomers are difficult to distinguish chemically.

Describe a method of converting them to suitable products in step 1 which can then be tested in step 2.

You should state the reagents and conditions for each step, and any observations you would make.

tep 1	
tep 2	
	[3]

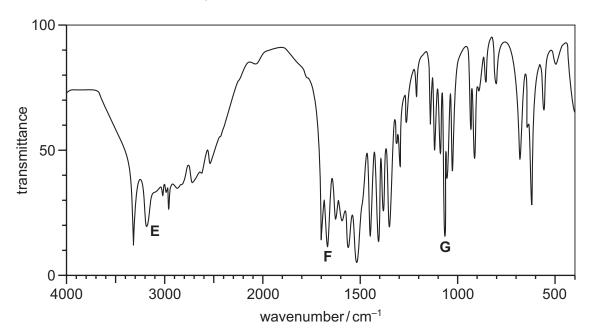
[Total: 8]

(a)	Am	ino acids can be separated by electrophoresis.									
	(i)	Draw a labelled diagram of the apparatus used to separate a mixture by electrophoresis.									
	(ii)	[3] Explain the principles of the separation of amino acids by electrophoresis.									
		[2]									
(b)	Ele	ctrophoresis is usually carried out in a buffer solution.									
	wou You	Given three buffers, with pH values of 2.0, 7.0 and 12.0, suggest, with a reason, which buffer would be the most suitable for the separation of the following amino acid mixtures. Your reasons should refer to the structure of each molecule. (The structures of these amino acids are given in the <i>Data Booklet</i> .)									
	(i)	Asp and Val									
		buffer pH									
		reason									
	(ii)	Lys and Ser									
		reason									
	(iii)	Tyr and Phe									
		buffer pH									
		reason									
		[3]									

(c) (i) Draw the structure of the dipeptide Gly-Ser, showing the peptide bond in full.

[2]

The infra-red spectrum of Gly-Ser is shown below.



(ii) Use the *Data Booklet* to identify the bond in the molecule of Gly-Ser that is responsible for each of the peaks indicated on the above infra-red spectrum.


F .....

•							

[2]

[Total: 12]

(a)	De	scribe and explain the trend in the solubility of the hydroxides down Group 2.
		[3]
(b)	Cal	cium reacts vigorously with HC $l(aq)$ producing $H_2(g)$ .
		$Ca(s) + 2HCl(aq) \rightarrow CaCl_2(aq) + H_2(g)$
	(i)	How would you expect the enthalpy change for this reaction to compare with the enthalpy change for the reaction where $HNO_3(aq)$ is used in place of $HCl$ but all other conditions are the same? Explain your answer.
		743

(	ii)	) The	ionic	equation	for	this	reaction	is	shown
۸		,		09444					00

Ca(s) + 
$$2H^+(aq) \rightarrow Ca^{2+}(aq) + H_2(g)$$
  $\Delta H^{\circ} = \mathbf{x} \text{ kJ mol}^{-1}$ 

Construct a **fully labelled** Hess' Law cycle to connect each side of this equation to the relevant gas phase ions.

Use your cycle, the following data, **and** data from the *Data Booklet*, to calculate a value for **x**.

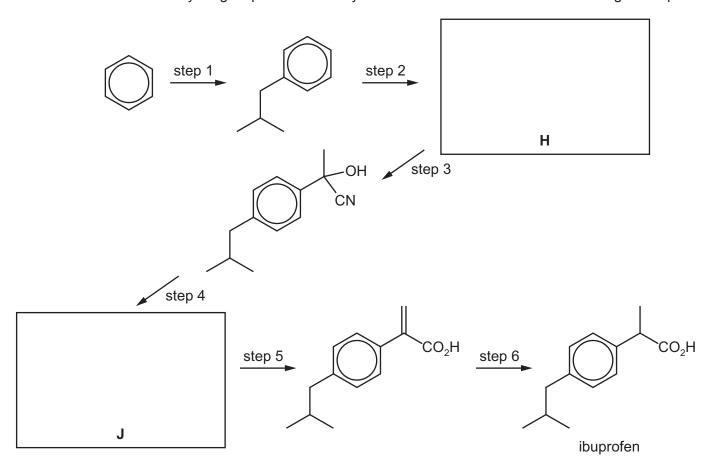
standard enthalpy of atomisation of Ca(s), $\Delta H_{\rm at}^{\rm e}({\rm Ca})$	+178 kJ mol <sup>-1</sup>
standard enthalpy of hydration of Ca <sup>2+</sup> (g), $\Delta H_{\text{hyd}}^{\bullet}(\text{Ca}^{2+})$	–1576 kJ mol <sup>–1</sup>
standard enthalpy of hydration of $H^{+}(g)$ , $\Delta H^{e}_{hyd}(H^{+})$	–1090 kJ mol <sup>–1</sup>

c)	The standard enthalpy change for the reaction between $Ca(s)$ and $CH_3CO_2H(aq)$ is less negative than <b>x</b> by $2  kJ  mol^{-1}$ .	>
	Suggest an explanation for this.	

[Total: 10]

 $x = ..... kJ mol^{-1} [4]$ 

**9** The anti-inflammatory drug ibuprofen can be synthesised from benzene via the following six steps.



- (a) Draw circles around any chiral carbon atoms in the above five formulae. [1]
- (b) Suggest the structures of compounds **H** and **J** and draw them in the boxes above. [2]
- (c) Suggest reagents and conditions for steps 1-6.

step 1	
step 2	
step 3	
step 4	
step 5	
step 6	 61

(d) Name the mechanism of step 1 and state the *type of reaction* for step 6.

step 1	 	 	 	 	 	
step 6	 	 	 	 	 	

[Total: 11]

[2]

				19	
10	(a)	(i)	Complete the electron	ic configuration of the iron	atom.
			Fe 1s <sup>2</sup> 2s <sup>2</sup> 2p <sup>6</sup>		[1]
		(ii)	In some of its complex	kes, the Fe³+ ion has <b>only o</b>	ne unpaired electron in its d orbitals.
				nd ↓ to represent electrons I orbital electronic configura	of opposite spins, complete the following tion of <b>this</b> Fe <sup>3+</sup> ion.
			3d {		energy
			(		[1]
	<ul> <li>(b) A solution containing a mixture of Sn²+(aq) and Sn⁴+(aq) is added to a solution containing mixture of Fe²+(aq) and Fe³+(aq).</li> <li>Use E⁰ data from the Data Booklet to predict the reaction that might take place when the solutions are mixed, and write an equation for the reaction.</li> </ul>				(aq) is added to a solution containing a
					•

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(c)	Hexaaquairon(III) ions are pale violet. They form a colourless complex with fluoride ions, F-	.,
	equilibrium 1, and a deep-red complex with thiocyanate ions, SCN-, equilibrium 2.	

$$[\text{Fe}(\text{H}_2\text{O})_6]^{3^+} + \text{F}^- \rightleftharpoons [\text{Fe}(\text{H}_2\text{O})_5\text{F}]^{2^+} + \text{H}_2\text{O}$$
 equilibrium **1**  $K_{\text{stab}} = 2.0 \times 10^5 \, \text{mol}^{-1} \, \text{dm}^3$  violet 
$$[\text{Fe}(\text{H}_2\text{O})_6]^{3^+} + \text{SCN}^- \rightleftharpoons [\text{Fe}(\text{H}_2\text{O})_5\text{SCN}]^{2^+} + \text{H}_2\text{O}$$
 equilibrium **2**  $K_{\text{stab}} = 1.0 \times 10^3 \, \text{mol}^{-1} \, \text{dm}^3$  violet 
$$\text{deep-red}$$

(i) Predict and explain the **sequence** of colour changes you would observe in each of the following experiments.

•	KF(aq).
•	A few drops of KF(aq) are added to 5 cm <sup>3</sup> of Fe <sup>3+</sup> (aq), followed by a few drops o KSCN(aq).

(ii)	What type of reaction is occurring during the experiments in (i)?	
		[1]

[Total: 9]

[4]

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