



#### **Cambridge International Examinations**

Cambridge International Advanced Subsidiary and Advanced Level

CANDIDATE NAME									
CENTRE NUMBER						CANDIDATE NUMBER			
CHEMISTRY								970	)1/41
Paper 4 A Leve	l Structu	red Ques	stions				May/	June	2016
								2 h	ours
Candidates ans	wer on th	ne Quest	ion Pape	r.					
Additional Mater	rials:	Data Bo	ooklet						

#### **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)		ewater is a saturated solution of $\mathrm{Ca}(\mathrm{OH})_2$ in water. It is used to test for the presence of $\mathrm{CO}_2$ gaseous mixture.
		(i)	Write an equation for the reaction between limewater and CO <sub>2</sub> .
			[1
			aturated solution of Ba(OH) <sub>2</sub> can be used instead of Ca(OH) <sub>2</sub> to test for CO <sub>2</sub> . A saturated ution of Mg(OH) <sub>2</sub> cannot be used for this test.
		(ii)	Explain why a saturated solution of Ba(OH) <sub>2</sub> can be used to test for CO <sub>2</sub> .
			[1
		(iii)	Explain why a saturated solution of $\mathrm{Mg}(\mathrm{OH})_2$ cannot be used to test for $\mathrm{CO}_2$ .
			[2
	(b)		scribe and explain the trend in the thermal stabilities of the carbonates down Group 2.
			[3
	(c)	Nic	kel carbonate, NiCO <sub>3</sub> , decomposes on heating.
			e the Data Booklet to explain whether $NiCO_3$ will decompose more or less readily than $CO_3$ .
			[2
			[Z

2

(a) (i	) Complete the foll	owing electror	nic config	urations.		
	the cobalt at	om, Co	1s <sup>2</sup> 2s <sup>2</sup> 2p <sup>6</sup>	······		
	• the cobalt(II)	) ion, Co <sup>2+</sup>	1s²2s²2p <sup>6</sup>	3		[1]
(ii)	State the colours solution of cobalt Give the formulae	(II) nitrate, Co	$(NO_3)_2$ .		HCl(aq) is added to d.	an aqueous
						[5]
С	$So(R_3P)_2I_2$ . (R = alkyl	l, R₃P is a mon	odentate	e ligand)	one structure with voisomers of Ni(R₃F	
	Ni(R <sub>3</sub>	.P).T.		Ni <i>i</i>	$(R_3P)_2I_2$	
	111(13)	3. 72-2			1	_
			Co(R <sub>3</sub> F	$\left( I_{1}\right) _{2}I_{2}$		
					•	[3]

[Total: 9]

3	One method of producing hydrogen from natural gas is the reaction between hydrogen sulfide and
	methane.

$$2H_2S(g) + CH_4(g) \rightleftharpoons CS_2(g) + 4H_2(g)$$

	(2)	Write the	ovnroccion	for K	for	thic	roaction	and	ctata	ito	unito
1	(a)	vviile liie	expression	IOI N	ຸເບເ	แแร	reaction,	anu	State	เเร	units

$$K_p =$$

(b) The initial partial pressures of the two gases in a mixture at 1000 K are recorded.

$$H_2S(g)$$
 200 atm

The mixture is left to reach equilibrium.

It is found that the equilibrium partial pressure of  $CS_2(g)$  is 2atm and that of the remaining  $CH_4(g)$  is 98 atm.

(i) Calculate the equilibrium partial pressures of  $H_2S(g)$  and  $H_2(g)$ .

$$p(H_2S) = \dots$$
 atm

$$p(H_2) = \dots$$
 atm [2]

(ii) Calculate the value of  $K_{\rm p}$  at this temperature.

$$K_p$$
 = .....[1]

(c)	(i)	Predict the sign of $\Delta S^e$ for this reaction. Explain your answer.
		$2H_2S(g) + CH_4(g) \rightleftharpoons CS_2(g) + 4H_2(g)$ $\Delta H^{\circ} = +241 \text{ kJ mol}^{-1}$
		[1]
	The	free energy change, $\Delta G^{\circ}$ , for this reaction at 1000 K is +51 kJ mol <sup>-1</sup> .
	(ii)	Calculate the value of $\Delta S^{\circ}$ for this reaction, stating its units.
		$\Delta S^{\circ}$ = units [2]
(d)		v would the value of $\Delta G^{\circ}$ , and hence the spontaneity (feasibility) of this reaction change as temperature increases? Explain your answer.
		[2]
		[Total: 10]

			6	
4	(a)	(i)	Define the term standard cell potential, $E_{\text{cell}}^{\bullet}$ .	
			[	1]
		The for	e following incomplete diagram shows the apparatus that can be used to measure the $E_5^6$ a cell composed of the Fe <sup>3+</sup> /Fe <sup>2+</sup> and Ag <sup>+</sup> /Ag half-cells.	∍ cell
			A C	
			ט	
		(ii)	Complete the diagram, labelling the components you add.	1]
		(iii)	Identify the components <b>A-D</b> .	
			A	
			В	
			C	
			D	
				3]
	(b)	(i)	Use $E^{\circ}$ values to write an equation for the cell reaction that takes place if the two electrode in <b>(a)</b> are connected by a wire and the circuit is completed.	
			г	
		(ii)	Another electrochemical cell was set up using 0.31 mol dm <sup>-3</sup> Ag <sup>+</sup> (aq) instead of th standard Ag solution.	е
			Use the Nernst equation, $E = E^{\circ} + 0.059 \log [Ag^{+}(aq)]$ , and the relevant $E^{\circ}$ values to calculate the new $E_{cell}$ in this experiment.	to

[Total: 8]

5 Dicarboxylic acids dissociate in stages.

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$$\mathsf{HO_2C}(\mathsf{CH_2})_\mathsf{n}\mathsf{CO_2H} \quad \overset{\mathsf{stage 1}}{\longleftarrow} \quad \mathsf{HO_2C}(\mathsf{CH_2})_\mathsf{n}\mathsf{CO_2}^- \, + \, \mathsf{H}^+ \quad \overset{\mathsf{stage 2}}{\longleftarrow} \quad {}^-\mathsf{O_2C}(\mathsf{CH_2})_\mathsf{n}\mathsf{CO_2}^- \, + \, \mathsf{2H}^+$$

(a) The  $pK_a$  values for stage 1 and stage 2 for some dicarboxylic acids are listed below.

n in HO <sub>2</sub> C(CH <sub>2</sub> ) <sub>n</sub> CO <sub>2</sub> H	pK <sub>a</sub> (1) for stage 1	pK <sub>a</sub> (2) for stage 2		
1	2.83	5.69		
2	4.16	5.61		
3	4.31	5.41		

For comparison, the  $pK_a$  of ethanoic acid,  $CH_3CO_2H$ , is 4.76.

(i)	State the mathematical relationship between $pK_a$ and the acid dissociation constant $K_a$ .
	[1]
(ii)	With reference to the table above, suggest why the $pK_a(1)$ values
	• are all smaller than the $pK_a$ of ethanoic acid,
	become larger as n increases.
	[3]
	Suggest why all the $pK_a(2)$ values in the table above are larger than the $pK_a$ of ethanoic acid.
	[1]
(b) The	monosodium salts of edible dicarboxylic acids are added to some foodstuffs as buffers.
(i)	Explain what is meant by the term buffer solution.
	[2]
	Write two equations to show how monosodium butanedioate, $HO_2CCH_2CH_2CO_2Na$ , acts as a buffer.
	[2]

[Total: 9]

6	Nitrobenzene,	C <sub>6</sub> H <sub>5</sub> NO <sub>2</sub> ,	can be	reduced to	phenylamine,	C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub> ,	in acid	solution	in a	two	step
	process.										

(a)	(i)	Balance the half-equation for this reaction to work out how many moles of electrons are
		needed to reduce one mole of nitrobenzene

$$C_{6}H_{5}NO_{2} + \dots H^{+} \rightarrow C_{6}H_{5}NH_{2} + \dots H_{2}O$$
 [1]

(ii) The reducing agent normally used is granulated tin and concentrated hydrochloric acid. In the first step, the reduction of nitrobenzene to phenylammonium chloride can be represented by the equation shown.

Use oxidation numbers or electrons transferred to balance this equation. You might find your answer to (i) useful.

.....
$$C_6H_5NO_2 + .....HCl + .....Sn \rightarrow .....C_6H_5NH_3Cl + .....SnCl_4 + .....H_2O$$
 [2]

(b) When 5.0 g of nitrobenzene was reduced in this reaction, 4.2 g of phenylammonium chloride, C<sub>6</sub>H<sub>5</sub>NH<sub>3</sub>C*l*, was produced.

Calculate the percentage yield.

- (c) Following the reaction in (b), an excess of NaOH(aq) was added to liberate phenylamine from phenylammonium chloride.
  - (i) Calculate the mass of phenylamine, C<sub>6</sub>H<sub>5</sub>NH<sub>2</sub>, produced when 4.20 g of phenylammonium chloride reacts with an excess of NaOH(aq).

The final volume of the alkaline solution of phenylamine in (i) was 25.0 cm<sup>3</sup>. The phenylamine was extracted by addition of 50 cm<sup>3</sup> of dichloromethane. After the extraction, the dichloromethane layer contained 2.68 g of phenylamine.

(ii) Use the data to calculate the partition coefficient,  $K_{\text{partition}}$ , of phenylamine between dichloromethane and water.

$$K_{\text{partition}} = \dots$$
 [2]

(d)	How does the basicity of	phenylamine compare to the	nat of ethylamin	e? Explain your answer.
				[2]
(e)	Phenol can be synthesise	ed from phenylamine in two	steps.	
	NH <sub>2</sub> step 1		step 2	OH + N <sub>2</sub>
		E		

(i) State the reagents and conditions for steps 1 and 2.

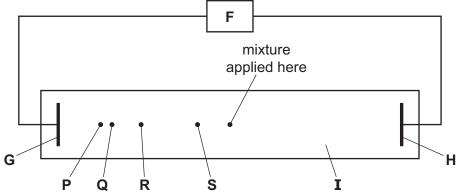
step 1	 	 
step 2		
		[2]

(ii) Draw the structure of the intermediate compound **E** in the box above. [1]

[Total: 13]

7	(a)	(i)	Use the <i>Data Booklet</i> to draw the structure of the tripeptide Ala-Ser-Gly showing its peptide bonds in full.
			Ala-Ser-Gly [2]
		(ii)	Calculate the relative molecular mass, $M_r$ , of Ala-Ser-Gly.
			$M_{\rm r} = \dots $ [1]
	(b)	Αn	ctrophoresis can be used to separate mixtures of amino acids and peptides.  nixture of the tripeptide Ala-Ser-Gly and its three constituent amino acids was subjected to ctrophoresis in a buffer at pH 11.
		(i)	Draw the structure of serine at pH11.
			[1]

At the end of the experiment the following results were seen.



(ii)	Identify the components labelled <b>F-I</b> in the above diagram.
	F
	G
	Н
	<b>I</b> [4
(iii)	Suggest the identities of the species responsible for
	spot <b>P</b> ,
	spot <b>S</b> .
	Explain your answers.
	[3
(c) (i)	State the reagents and conditions needed for converting the tripeptide into its three constituent amino acids.
	[1
(ii)	Name the type of reaction in (i).
	[1
	[Total: 13

8 (a) Chromyl chloride,  $CrO_2Cl_2$ , can be prepared by heating a mixture of potassium dichromate(VI) and potassium chloride with concentrated sulfuric acid.

$$\mathsf{K_2Cr_2O_7} \ + \ \mathsf{4KC}\mathit{l} \ + \ \mathsf{3H_2SO_4} \ \rightarrow \ \mathsf{2CrO_2C}\mathit{l_2} \ + \ \mathsf{3K_2SO_4} \ + \ \mathsf{3H_2O}$$

Use the following data to complete the Hess' Law cycle and calculate the enthalpy change of the reaction,  $\Delta H_{\rm r}$ .

compound	enthalpy change of formation, $\Delta H_{\rm f}^{\rm e}/{\rm kJmol^{-1}}$
K <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub>	-2061
KC1	-437
H <sub>2</sub> SO <sub>4</sub>	-814
CrO <sub>2</sub> Cl <sub>2</sub>	-580
K <sub>2</sub> SO <sub>4</sub>	-1438
H <sub>2</sub> O	-286

$$K_2Cr_2O_7 + 4KCl + 3H_2SO_4$$

$$\longrightarrow$$

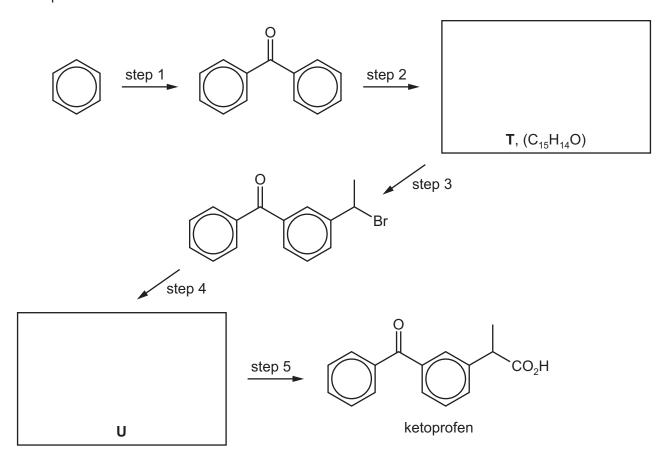
$$2CrO_2Cl_2 + 3K_2SO_4 + 3H_2O$$

elements

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

(b)		ere are two isomeric complex ions with the formula $[Cr(NH_3)_4Cl_2]^+$ . One is green and the er is violet.
	(i)	Suggest the type of isomerism shown by these complex ions.
		[1]
	(ii)	Explain why these two complex ions
		are coloured,
		have different colours.
		[4]
		[Total: 7]

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



(a) Suggest the structures of compounds **T** and **U** and draw them in the boxes above.

[2]

(b) Suggest reagents and conditions for steps 1-5.

step 1	
step 2	
step 3	
step 4	
step 5	[5]

(c) What types of reaction are steps 1 and 5?

step 1	
step 5	
	[2]

[Total: 9]

- **10** (a) Ethanedioic acid, C<sub>2</sub>O<sub>4</sub>H<sub>2</sub>, occurs in many vegetables. The amount that occurs in spinach can be estimated as follows.
  - 40.0 g of spinach leaves are crushed and mixed with distilled water, using a mortar and pestle.
  - The mixture is filtered, and the leaves are washed with a little more water.
  - The combined filtrate and washings are made up to 100.0 cm<sup>3</sup> with water.
  - A 25.0 cm<sup>3</sup> portion of the resulting solution is added to a conical flask, along with an excess of dilute sulfuric acid.
  - The acidified solution is warmed, and then titrated with 0.0200 mol dm<sup>-3</sup> KMnO<sub>4</sub>.

The equation for the reaction between ethanedioic acid and acidified manganate(VII) ions is shown.

$$2MnO_4^- + 6H^+ + 5C_2O_4H_2 \rightarrow 2Mn^{2+} + 10CO_2 + 8H_2O$$

In the titration, 15.20 cm<sup>3</sup> of KMnO<sub>4</sub> was required to reach the end-point.

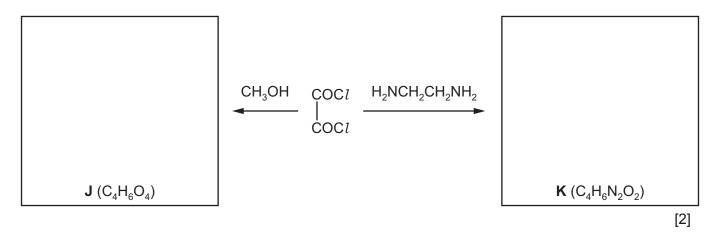
Calculate the percentage by mass of ethanedioic acid in the spinach leaves.

percentage of ethanedioic acid = ..... % [3]

(b) Ethanedioic acid can be converted into ethanedioyl chloride:

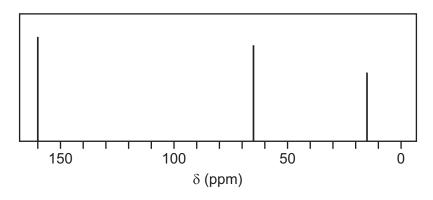
(i) State a suitable reagent for this reaction.

(ii) For the reactions of ethanedioyl chloride below, suggest the structures of compounds **J** and **K** and draw them in the boxes.

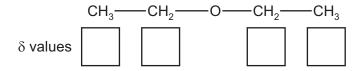


(c) When ethanedioyl chloride is reacted with silver ethanedioate,  $AgO_2CCO_2Ag$ , in ethoxyethane at  $-30\,^{\circ}C$ , an oxide of carbon, **L**, is formed. The molecule of **L** has no overall dipole and has molecular formula  $C_4O_6$ .

The carbon-13 NMR spectrum of a solution of **L** in ethoxyethane, CH<sub>3</sub>CH<sub>2</sub>OCH<sub>2</sub>CH<sub>3</sub>, is shown below.



(i) Use the *Data Booklet* to state in the boxes below the  $\delta$  values for the peaks in the spectrum which are due to the carbon atoms in ethoxyethane.



[2]

(ii) Explain what the rest of the carbon-13 NMR spectrum indicates about the structure of  ${\bf L}$ .

.....[1]

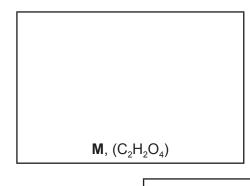
(	d)	When pure L	is reacted	with an excess	s of CH <sub>a</sub> Ol	l a mixture o	f three com	pounds is fo	rmed
•	u,	VVIICII PUIC L		WILLI GIT CACCO		i, a illixtale o		pourius is io	/I I I I I C C

 ${\bf M}$  is formed as one of the products when either  ${\bf N}$  or  ${\bf O}$  is heated with aqueous acid.

The table gives information of the peaks recorded in the carbon-13 NMR spectra of M, N and O.

compound peaks recorded in carbon-13 NMR spectr	
M	δ 162
N	δ 53 δ 160 δ 162
0	δ 53 δ 160

(i) Suggest the structures of M, N and O.



**N**, (C<sub>3</sub>H<sub>4</sub>O<sub>4</sub>)

**O**, (C<sub>4</sub>H<sub>6</sub>O<sub>4</sub>)

[3]

(ii) Suggest a structure for L that fits all the data given in (c) and (d).

L, (C<sub>4</sub>O<sub>6</sub>)

[1]

[Total: 13]

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