

UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS

General Certificate of Education Advanced Subsidiary Level and Advanced Level

CANDIDATE NAME							
CENTRE NUMBER				CANDI NUMBI			

CHEMISTRY 9701/04

Paper 4 Structured Questions

October/November 2007

1 hour 45 minutes

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 a pencil for any diagrams, graphs or rough working.

Do not use staples, paper clips, highlighters, glue or correction fluid.

DO NOT WRITE IN ANY BARCODES.

Section A

Answer all questions.

Section B

Answer all questions.

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.

For Examiner's Use				
1				
2				
3				
4				
5				
6				
7				
8				
9				
10				
Total				

This document consists of 15 printed pages and 1 blank page.



Section A

1

		a carboxylic acid, RCC	₂ H, to wri	ite equations to explain the
following terms				ded. ite equations to explain the
η ρ/ _α				[2]
The p $K_{\!a}$ values	s of four carbo	xylic acids are listed ir	the table	below.
	acid	formula of acid	p <i>K</i> _a	
	1	CH ₃ CH ₂ CO ₂ H	4.9	
	2	CH ₃ CHC <i>l</i> CO ₂ H	2.8	
	3	CH ₃ CC <i>l</i> ₂ CO ₂ H	1.4	
	4	CH ₂ ClCH ₂ CO ₂ H	4.1	
(i) Describe a	and explain the	e trend in acid strength	n shown b	y acids 1, 2 and 3.
	•••••			
	•••••			
	•••••			
		for the difference in the	17	
i) Suggest a	n explanation	for the difference in th	e pr _a vall	Jes for acids 2 and 4.

	Cl_2 + trace of P		NH ₃ (excess)	
CH ₃ CH ₂ CO ₂ H		CH ₃ CHC1CO ₂ H		CH₃CH(NH₂)CO₂H
v – –		_	II	alanine

	42
	3 M. D.
A g	ood way of making synthetic amino acids uses chloro-acids as intermediates
CH₂C	ood way of making synthetic amino acids uses chloro-acids as intermediates $CO_2H \xrightarrow{Cl_2 + \text{trace of P}} CH_3CHC_1CO_2H \xrightarrow{NH_3(\text{excess})} CH_3CH(NH_2)CO_2H$ alanine
(i)	Suggest the role that the trace of phosphorus plays in reaction I.
(ii)	Write a fully balanced equation for reaction I.
(iii)	State the type of mechanism of reaction II.
(iv)	When 10.0 g of propanoic acid was used in this 2-stage synthesis, a yield of 9.5 g of alanine was obtained. Calculate the overall percentage yield.

(d) In the solid state and in aqueous solutions, alanine exists as a zwitterion. Draw the structural formula of this zwitterion.

[2]

[5]

[Total: 15]

2

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(a)	Des eler	nents.
		[4]
(b)	tren	salts formed by the Group II elements with other divalent anions show a similar d in their solubilities, whereas most of their salts with monovalent anions are very ble.
	des	this information to predict the identities of compounds A and B in the following cription of some reactions of Group II compounds, and write balanced equations for reactions.
	mag cold	gnesium hydroxide, $\mathrm{Mg}(\mathrm{OH})_2$, is almost insoluble in water. Stirring a mixture of gnesium hydroxide and aqueous ethanedioic acid, $\mathrm{H_2C_2O_4}$, produces a clear burless solution containing A . When a solution of calcium nitrate, $\mathrm{Ca}(\mathrm{NO_3})_2$, is added, hite precipitate of B is formed.
	ider	ntity of A identity of B
	equ	ations
		[3]
(c)	The 2.0	solubility product, K_{sp} , of magnesium hydroxide has a numerical value of \times 10 ⁻¹¹ .
	(i)	Write an expression for the K_{sp} of magnesium hydroxide, stating its units.
	(ii)	Use the value of K_{sp} given to calculate the concentration of ${\rm Mg(OH)_2}$ in a saturated solution.
	(iii)	Explain whether magnesium hydroxide would be more or less soluble in 0.1 $\rm moldm^{-3}MgSO_4(aq)$ than in water.
		[5]

[Total: 12]

[Total: 9]

3 The following account describes the preparation of Péligot's salt, named after the 19th French chemist who first made it.

www.papaCambridge.com Place 6.0 g of potassium dichromate(VI) in a 100 cm³ beaker and add 8.0 g of concentrated hydrochloric acid and 1.0 cm³ water. Warm the mixture gently; if carefully done the dichromate(VI) will dissolve without the evolution of chlorine. On cooling the beaker in an ice bath the solution will deposit long orange-red crystals of Péligot's salt.

An analysis of Péligot's salt showed that it contained the following percentages by mass: K, 22.4%; Cr, 29.8%; Cl, 20.3%; O, 27.5%.

(a) Calculate the empirical formula of Péligot's salt.

		[2]
(b)	Sug	gest a balanced equation for the formation of Péligot's salt.
		[1]
(c)	The	instructions suggest that strong heating might cause chlorine to be evolved.
	(i)	What type of reaction would produce chlorine in this system?
	(ii)	Use the <i>Data Booklet</i> to identify relevant half equations and $E^{\rm e}$ values for the production of chlorine from the reaction between ${\rm K_2Cr_2O_7}$ and ${\rm HC}\it{l}.$
		Use these equations to write the overall full ionic equation for this reaction.
	(iii)	The use of ${\bf dilute}~{\rm HC}\it{l}({\rm aq})$ does not result in the production of chlorine. Suggest why this is so.
	(iv)	Use the $\it Data Booklet$ to suggest a reason why it is $\it not$ possible to prepare the bromine analogue of Péligot's salt by using HBr(aq) instead of HC $\it l(aq)$.
		[6]

(a) By choosing the chlorides of two of the Group IV elements as examples, described trend in the reactions of these chlorides with water. Suggest an explanation for differences, and write equations for any reactions that occur. (b) The standard enthalpy changes of formation of lead(II) chloride and lead(IV) chloride are given in the following table. $\Delta H_{\rm f}^{\Theta}/{\rm kJ\,mol^{-1}}$ compound PbC $l_2(s)$ -359 $PbCl_4(I)$ -329Use these data, and also bond energy data from the Data Booklet, to calculate the enthalpy changes for the following two reactions. (i) $CCl_2(g) + Cl_2(g) \longrightarrow CCl_4(g)$ $\Delta H^{\Theta} = \dots kJ \, \text{mol}^{-1}$ (ii) $PbCl_2(s) + Cl_2(g) \longrightarrow PbCl_4(l)$ $\Delta H^{\Theta} = \dots kJ \, \text{mol}^{-1}$ Make use of your answers to parts (i) and (ii) to suggest how the relative stabilities of the two oxidation states vary down the Group.

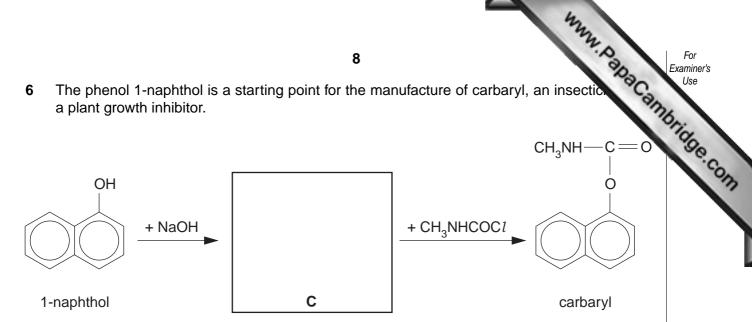
[Total: 6]

[3]

[Total: 6]

5	Pota in h the	nouse	m manganate(VII) can be used to estimate the percentage of hydrogen pehold bleach. The following unbalanced equation represents the reaction between	Cal
		MnO	$O_4^- + \dots + O_2^- + \dots + O_2^- + \dots + O_2^-$	2
	(a)	Bala	ance this equation by putting the appropriate numbers in the spaces above.	[1]
	(b)	Use	data from the <i>Data Booklet</i> to calculate the E_{cell}^{θ} for the reaction.	
				[1]
	(c)	Whe	en $0.020\mathrm{moldm^{-3}}$ KMnO ₄ (aq) was added from a burette into an acidified 25.0 cm aple of $\mathrm{H_2O_2}$, $15.0\mathrm{cm^3}$ of KMnO ₄ was required to reach the end-point.	m ³
		(i)	Describe what you would see during this titration, and also at the end-point.	
		(ii)	Calculate the concentration of H ₂ O ₂ in the sample.	
				 [4]

6 The phenol 1-naphthol is a starting point for the manufacture of carbaryl, an insection a plant growth inhibitor.



Suggest a structure for the intermediate **C** and draw it in the box above.

(ii)	Name the functional groups in carbaryl.

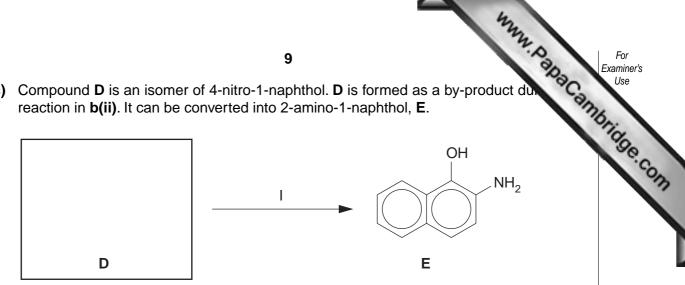
(iii) Suggest structures for the three products formed when carbaryl is hydrolysed.

- What reagents and conditions would you use for this hydrolysis? [7]
- (b) Suggest reagents and conditions for converting 1-naphthol into each of the following compounds.





(c) Compound **D** is an isomer of 4-nitro-1-naphthol. **D** is formed as a by-product du reaction in b(ii). It can be converted into 2-amino-1-naphthol, E.

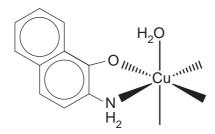


- Suggest the structural formula of the isomer **D**.
- Suggest reagents needed for reaction I. (ii)

(iii) Suggest the structural formula of the compound formed when compound E reacts with an excess of CH₃COC1.

[3]

- (d) When an alkaline solution of compound **E** is added to a solution containing Cu²⁺(aq) ions, a pale green-blue precipitate F forms. Analysis of F shows that its formula is $Cu(C_{10}H_8NO)_2(H_2O)_2$.
 - (i) Complete the following structural formula of **F**.



When an excess of concentrated NH₃(aq) is added to **F**, the precipitate dissolves to form a deep blue solution.

State the formula of the ion responsible for the deep blue colour.

(iii) What type of reaction is occurring here?

[3]

7 The nitration of benzene occurs in the following steps.

(a) What reagents and conditions are needed for this reaction?

.....[2]

(b) Write an equation showing how the electrophile NO_2^+ is formed from the reagents.

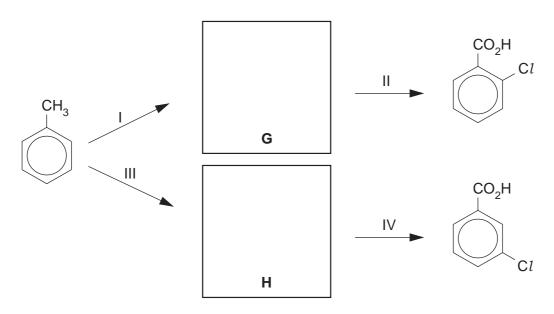
_____[1]

(c) The nitration of methylbenzene produces mainly 2-nitromethylbenzene, whereas the nitration of benzoic acid produces mainly 3-nitrobenzoic acid.

2-nitromethylbenzene

3-nitrobenzoic acid

Use this information to suggest suitable intermediates ${\bf G}$ and ${\bf H}$ in the following two 2-stage syntheses of chlorobenzoic acids, and suggest suitable reagents for reactions I to IV.



reagents:

reaction I reaction II

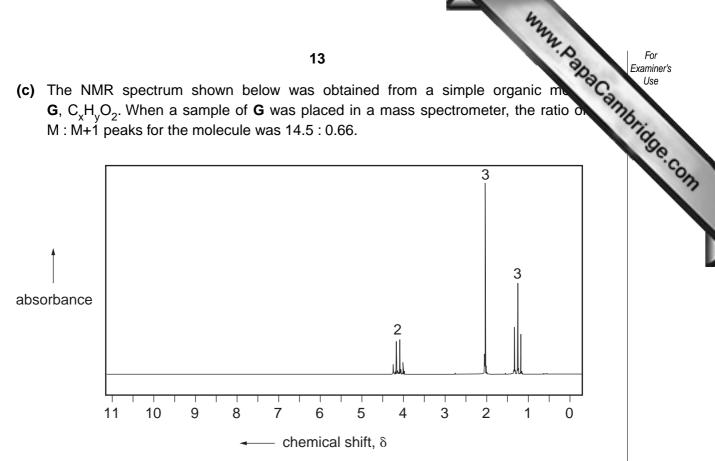
reaction III reaction IV

Section B – Applications of Chemistry

		Section B – Applications of Chemistry Answer all questions in the spaces provided. A carries the genetic code in living organisms and consists of a double helix.	Co
(-\	DNI	Answer all questions in the spaces provided.	
(a)		A carries the genetic code in living organisms and consists of a double helix.	
	(i)	Describe what is meant by a double helix.	
	(ii)	How are the strands of the double helix held together?	
			 [2]
(b)	perf	eplicating the genetic code two RNA molecules, mRNA and tRNA, are used orm functions called <i>transcription</i> and <i>translation</i> . cribe the role of the RNA molecules in these two functions.	to
	tran	scription	
		slation	
			[4]
(c)	Whe	en an egg is boiled, the protein changes from a viscous liquid to a solid.	
	(i)	Suggest what causes this change as the protein is heated.	
	(ii)	Why is there no change to the primary structure of the protein under the conditions?	

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d)	Describe in outline how energy is provided in animal cells.	dindy.
		de
	[3	 3]
	[Total: 11	1]
	Explain with reference to energy states how ¹ H NMR can supply information about th structure of molecules.	е
b)	Nuclear magnetic resonance is used in magnetic resonance imaging scanners. Thes scanners are increasingly used in hospitals to detect tumours. Suggest why magneti resonance techniques are better than X-rays.	e
b)	Nuclear magnetic resonance is used in magnetic resonance imaging scanners. Thes scanners are increasingly used in hospitals to detect tumours. Suggest why magnetic	e

(c) The NMR spectrum shown below was obtained from a simple organic management \mathbf{G} , $C_xH_vO_2$. When a sample of \mathbf{G} was placed in a mass spectrometer, the ratio \mathbf{O} M: M+1 peaks for the molecule was 14.5: 0.66.



(i) Calculate how many carbon atoms there are in the molecule.

(ii) Use the NMR spectrum and the Data Booklet to work out the structure of G.

10 Read the following article about the use of bacteria in mining, and then answer the qu that follow it.

www.PapaCambridge.com The discovery that bacteria could 'mine' metals for us was made in Spain. The Rio Tinto mine, in the southwest corner of Spain, was originally mined for copper by the Romans some 2,000 years ago. In 1752, some mining engineers looked over the mine to see if it could possibly be re-opened. They noticed streams of a blue-green liquid running from spoil heaps of the processed rock that lay around the mine. When this blue-green liquid ran over iron, it coated the iron with a brown film. The brown film was metallic copper.

There was still some copper left in the spoil heaps. At the time, everybody thought that the copper was being dissolved in the liquid through a simple chemical reaction. But in 1947, US scientists discovered that the copper was being 'mined' by a bacterium called *Thiobacillus ferrooxidans*.

The bacterium *Thiobacillus ferrooxidans* lives off the chemical energy trapped in metal sulphides. In the ore, the copper exists as copper sulphide. The bacteria gain energy by converting the copper sulphide to copper sulphate, which is then excreted. At the same time, they absorb the difference in energy in the chemical bonds. These bacteria can also obtain energy in similar reactions with ores of zinc, lead and uranium.

(a)	Use the <i>Data Booklet</i> to explain why the blue-green liquid coated the iron with copper. Write an equation for the reaction.
	[2]
(b)	Suggest two reasons why this method of extracting copper might be useful for ore containing only a small percentage of copper.
	(i)
	(ii)
	[2]
(c)	Suggest one disadvantage of using bacteria rather than traditional mining and smelting methods.
	[1]

(d) In conventional copper mining, the ore will typically contain 0.5 - 2.0% copper, gives an idea of what a valuable resource copper is.

	my my	
	15 For Examiner's	
	conventional copper mining, the ore will typically contain 0.5 – 2.0% copper copper ses an idea of what a valuable resource copper is.	
(i)	The ore from a particular mine contains 0.75% copper, and 150 000 tonnes of ore are mined each year. From this ore about 60% of the copper is extracted, and the remainder is left in the 'spoil heaps' of processed ore.	
	What mass of copper is extracted each year?	

(ii) If the use of bacteria can recover a further 17% of copper from the spoil heaps, what is the extra mass of copper produced?

(e)	Suggest why bacteria are unlikely to be used in the extraction of aluminium.
(f)	Metals like copper and zinc from abandoned mines can contaminate ground-water. Suggest one way of removing these contaminants.
	[1]

[Total: 9]

[2]

16

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