

UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS General Certificate of Education Advanced Level

CANDIDATE NAME				
CENTRE NUMBER		CANDIDATE NUMBER		

CHEMISTRY 9701/43

Paper 4 Structured Questions

May/June 2010

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				
Total				

This document consists of 17 printed pages and 3 blank pages.



Occilon I

Answer all questions in the spaces provided.

1 Phenacyl chloride has been used as a component of some tear gases. Its lachrymatory and irritant properties are due to it reacting with water inside body tissues to produce hydrochloric acid.

It undergoes a nucleophilic substitution reaction with NaOH(aq).

(a) Write the formulae of the products of this reaction in the two boxes above.

[2]

When the rate of this reaction was measured at various concentrations of the two reagents, the following results were obtained.

experiment number	[phenacyl chloride]	[NaOH]	relative rate
1	0.020	0.10	1.0
2	0.030	0.10	1.5
3	0.025	0.20	2.5

(i)	What is meant by the term <i>order of reaction</i> ?
(ii)	Use the above data to deduce the order with respect to each reactant. Explain your reasoning.
(iii)	Write the overall rate equation for the reaction.

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(b)

	(iv)	Describe the mechanism for this reaction that is consistent with your overall rate equation. You should show all intermediates and/or transition states and partial charges, and you should represent the movements of electron pairs by curly arrows.	For Examiner's Use
		[7]	
(c)	(i)	Describe an experiment that would show that $\mathrm{CH_3COC}l$ reacts with water at a much faster rate than phenacyl chloride. Include the reagents you would use, and the observations you would make with each chloride.	
	(ii)	Suggest an explanation for this difference in reactivity.	
	(,		
		[4]	
		[Total: 13]	

2	(a)		escribe and explain how the solubilities of the sulfates of the Group II elements vary own the group.					
	4.				[3]			
	(b)		following table lists some enpounds.	enthalpy changes for ma	agnesium and strontium			
			enthalpy change	value for magnesium /kJ mol ⁻¹	value for strontium /kJ mol ⁻¹			
	lattice	entl	halpy of M (OH) ₂	-2993	-2467			
	entha	lpy c	change of hydration of M ²⁺ (g)	-1890	-1414			
	entha	lpy c	change of hydration of OH ⁻ (g)	– 550	-550			
		(i)	Use the above data to calculate Mg(OH) ₂					
				$\Delta H_{\text{solution}}^{\Theta} = \dots$	kJ mol ⁻¹			
			Sr(OH) ₂					
				$\Delta H_{\text{solution}}^{\Theta} = \dots$	kJ mol ⁻¹			
		(ii)	Use your results in (i) to sugge than is Mg(OH) ₂ . State any ass	st whether Sr(OH) ₂ is mo umptions you make.	re or less soluble in water			
	((iii)	Suggest whether Sr(OH) ₂ woul Explain your reasoning.	d be more or less soluble	in hot water than in cold.			
					[5]			

(c)	Cal	cium hydroxide, Ca(OH) ₂ , is slightly soluble in water.	For Examiner's					
	(i)							
		$K_{\rm sp}$ = units						
	(ii)	$25.0\rm cm^3$ of a saturated solution of Ca(OH) $_2$ required 21.0 cm 3 of 0.0500 mol dm $^{-3}$ HC l for complete neutralisation.						
		Calculate the [OH-(aq)] and the [Ca^2+(aq)] in the saturated solution, and hence calculate a value for $K_{\rm sp}$.						
		[OH ⁻ (aq)] =						
		[Ca ²⁺ (aq)] =						
		$K_{sp} = \dots$						
1	(iii)	How would the solubility of $Ca(OH)_2$ in $0.1\mathrm{moldm^{-3}}$ NaOH compare with that in water? Explain your answer.						
		[6]						
		[Total: 14]						

	ggest a reason	for this difference.		
				[1]
	edict whether or appropriate co	-	s will have an overall dipole. Pla	ce a tick in
	compound	molecule has an overall dipole	molecule does not have an overall dipole	
	BCl ₃			
	PCl ₃			
	CCl ₄			
1				
	SF ₆			
	SF ₆			[2]
	ron and silicon react with wate Suggest a rea	er, whereas BC $l_{ m 3}$ and SiC $l_{ m 4}$ ason for this difference in re		CCl ₄ does
not	ron and silicon react with wate Suggest a rea	er, whereas $\mathrm{BC}l_3$ and $\mathrm{SiC}l_4$ ason for this difference in re	do react. activity.	CCl ₄ does
not	ron and silicon react with water Suggest a react with water.	er, whereas $\mathrm{BC}l_3$ and $\mathrm{SiC}l_4$ ason for this difference in results as a substitution of the reaction under the reaction of the reacti	do react. activity.	CCl ₄ does

(d)	When reacted with a small quantity of water, $SiCl_4$ produces an oxychloride X , $Si_xCl_vO_2$.
	The mass spectrum of X shows peaks at mass numbers of 133, 149, 247, 263 and 396.
	(You should assume that the species responsible for all these peaks contain the ¹⁶ O,
	the ³⁵ Cl and the ²⁸ Si isotopes only.)

/:\	Use these	data to		the me	Jaculari	formula	of V
(1)	use mese	Calac) deduce	me mo	necular	юннила	()I A

molecular formula

(ii) Suggest the structures of the fragments responsible for the peaks at the following mass numbers.

mass number	structure
133	
247	
263	

(iii) Hence suggest the displayed formula of X.

[5]

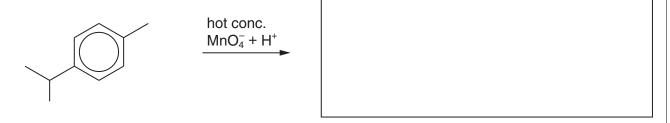
4

(a)	Cor	mplete the e	electronic structures of the Cr ³⁺ and Mn ²⁺ ions.	
		Cr ³⁺	1s ² 2s ² 2p ⁶	
		Mn ²⁺	1s ² 2s ² 2p ⁶ [2	2]
(b)	(i)	Describe slowly and a large exc	what observations you would make when dilute ${\rm KMnO_4}({\rm aq})$ is added with shaking to an acidified solution of ${\rm FeSO_4}({\rm aq})$ until the ${\rm KMnO_4}$ is increase.	d n
	(ii)	Construct	an ionic equation for the reaction that occurs.	
			[4	[]
(c)	Fe ²	+(aq) are re	elevant E^{Θ} data from the <i>Data Booklet</i> explain why acidified solutions of elatively stable to oxidation by air, whereas a freshly prepared precipitate eadily oxidised to Fe(OH) ₃ under alkaline conditions.	
	rele	vant <i>E</i> ^e val	ues and half equations	
	ехр	lanation		
			[4	<u> </u>

(d) Predict the organic products of the following reactions and draw their structures in the boxes below. You may use structural or skeletal formulae as you wish.

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$$\frac{\text{hot conc.}}{\text{MnO}_4^- + \text{H}^+}$$



[4]

(e) $\rm KMnO_4$ and $\rm K_2Cr_2O_7$ are the reagents that can be used to carry out the following transformation.

- (i) Draw the structure of intermediate **E** in the box above.
- (ii) Suggest reagents and conditions for the following.

reaction I	
reaction II	
	[3]

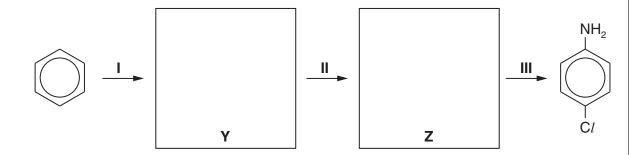
[Total: 17]

(ii) Briefly explain why all the carbon-carbon bonds in benzene are the same ler (b) Benzene can be nitrated by warming it with a mixture of concentrated sulfur nitric acids. (i) By means of an equation, illustrate the initial role of the sulfuric acid reaction. (ii) Name the type of reaction and describe the mechanism for the nitration re including curly arrows showing the movement of electrons and all charges. type of reaction	
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reaction. (ii) Name the type of reaction and describe the mechanism for the nitration re including curly arrows showing the movement of electrons and all charges. type of reaction	c and
including curly arrows showing the movement of electrons and all charges. type of reaction	n this
including curly arrows showing the movement of electrons and all charges. type of reaction	
	action,
mechanism	
(c) State the reagents and conditions needed to convert benzene into chlorobenzene	[4]

(d) Nitrobenzene undergoes further substitution considerably more slowly than chlorobenzene. In nitrobenzene the incoming group joins to the benzene ring in the 3-position, whereas in chlorobenzene the incoming group joins to the benzene ring in the 4-position.

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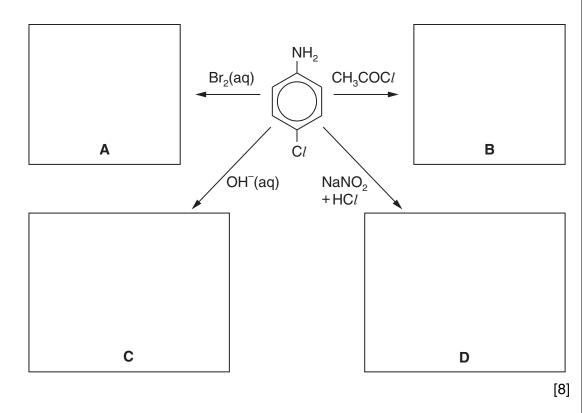
(i) Use these ideas to suggest the structures of the intermediate compounds **Y** and **Z** in the following synthesis of 4-chlorophenylamine.



(ii) Suggest the reagents and conditions needed for reaction III in the above synthesis.

.....

(iii) Suggest the structural formulae of the products **A**, **B**, **C** and **D** of the following reactions. If no reaction occurs write "no reaction" in the relevant box.



[Total: 15]

Section B

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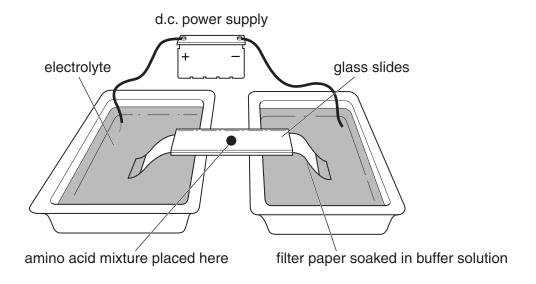
Answer all questions in the spaces provided.

6 Human hair and silk both consist of proteins. Proteins are described as having three levels of structure: primary, secondary and tertiary.					
	(a)	Outline what is meant by the terms <i>primary structure</i> and <i>tertiary structure</i> of a protein.			
		primary structure			
		tertiary structure			
		[2]			
	(b)	In hair, the secondary structure consists of α -helices which are cross-linked by disulfide bonds. The amino acid responsible for this cross-linking is cysteine, $H_2NCH(CH_2SH)CO_2H$.			
		(i) Show by means of a diagram how the disulfide cross-links are formed.			
		(ii) What type of reaction is this?			
		(ii) What type of reaction is this?			

	(iii)	State three other interactions that stabilise the tertiary structure of proteins.
		[4]
(c)		β -pleated sheet is a different form of secondary structure found in proteins, such as se in silk.
	(i)	What type of bonding is responsible for stabilising the β -pleated sheet in silk?
	(ii)	On the diagram below, draw a second polypeptide strand and show how bonds would be formed that stabilise this β -pleated sheet.
		R H O R H O C CH N C CH H O R H O R
		[3]
(d)	acid sucl	cysteine-containing protein in hair is called α -keratin. A similar sequence of amino is can produce β -keratin proteins found in the scales, claws and shells of reptiles in as tortoises. In β -keratin the secondary structure of the protein is in the form of a eated sheet.
	_	gest what makes the $\beta\text{-pleated}$ sheet in $\beta\text{-keratin}$ so much less flexible than the eated sheet in silk.
		[1]
		[Total: 10]

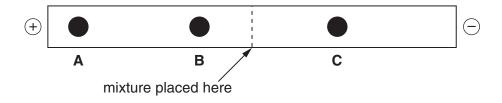
7 A mixture of amino acids may be separated using electrophoresis. A typical practical set-up is shown in the diagram.

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 Vhen the power supply is switched on, some amino acids may not move, but remain tationary. Suggest an explanation for this observation.
 [2]
The amino acid glycine has the formula $H_2NCH_2CO_2H$. Identify the species formed on the filter paper if glycine moves to the left (positive) end of the filter paper.
[4]

(c) The following result was obtained from another electrophoresis. What can be deduced about the relative sizes of, and charges on, the amino acid species A, B and C?



amino acid	relative size	charge
A		
В		
С		

[3]

(d) The sequence of amino acids in a polypeptide may be determined by partial of the chain into smaller pieces, often tripeptides.							drolysi		
	(i) Following such a partial hydrolysis, the following tripeptides were obtained given polypeptide.								
	ala-gly-asp gly-ala-gly lys-val-ser ser-ala-gly val-ser-ala								
					lysine (lys) sug I give the above	gest the amino acid se tripeptides.	quence		
	The	e structural for	mulae of the ar	mino acids	in the polypepti	de are given below.			
		abbreviation	amino ac	id	structura	al formula			
		ala	alanine	H ₂	H ₂ NCH(CH ₃)CO ₂ H				
		asp	aspartic a	cid H ₂	NCH(CH ₂ CO ₂ H	I)CO ₂ H			
		gly	glycine	H ₂	NCH ₂ CO ₂ H				
		lys	lysine	H ₂	NCH(CH ₂ CH ₂ C	CH ₂ CH ₂ NH ₂)CO ₂ H			
		ser	serine	H ₂	NCH(CH ₂ OH)C	CO ₂ H			
		val	valine	H ₂	NCH(CH(CH ₃) ₂)CO ₂ H			
	(ii)	Which of the	tripeptides in ((i) has the l	owest <i>M</i> _r ?				

[4]

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[Total: 10]

						16					
	The	e des	ign and deve	lopment of batte	eries ha	ıs been a	major re	esearch a	ırea in re	cent yea	rs.
(a) Lead-acid batteries, used in cars, are made up of a number of recharges series, and were first developed in 1860. They have the disadvantage of a re mass compared to the energy stored. During discharge, the electrode reac cells of these batteries are as follows.						relatively	high				
			I	Pb + SO_4^{2-}	→ PbS	SO ₄ + 26	e ⁻				
			II	$PbO_2 + 4H^+$	+ SO	²⁻ + 2e ⁻	→ Pb	SO ₄ + 2	H ₂ O		
				nese reactions of ining your answ		it the pos	itive ele	ctrode in	a lead-ad	cid cell du	uring
	(b)			ooklet and the ed ell under standa	quations	s I and II a					
											[2]
	(c)	incr can oxo	reasingly conneras that ne	dride batteries nmon particular ed near-consta iO(OH)) as one strode.	ly for si int sour	mall devi	ces sucl lectrical	n as mob energy. T	ile phone hese ce	es and d Ils use n	igital ickel
		One	e reaction tha	ıt takes place in	these b	oatteries	is				
				NiO(OH) +	H ₂ O +	e⁻ <==	Ni(OH) ₂	+ OH ⁻			
		(i)	State the ox	xidation state of	nickel i	n NiO(OF	H)				
		(ii)	Suggest a li	kely advantage	of thes	e batterie	es compa	ared with	lead-acid	d batterie	es.

[2]

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8

(u)	pow	rered vehicles. In these fuel cells hydrogen is oxidized to produce water, using a alyst and inert electrodes.	For Examiner's Use
	(i)	Suggest a material for the electrodes.	
	(ii)	Use your knowledge of hydrogen to suggest a disadvantage of these fuel cells in powering vehicles.	
		[2]	
(e)		ny of the world's countries are developing ways of recycling materials which are table or which require large amounts of energy to produce.	
	imp	each of the following recyclable materials, state whether recycling of this material is ortant in saving energy or in saving resources. Use your knowledge of chemistry to lain each choice.	
	glas	ss	
	stee	st	
	plas	etics	
		[3]	
		[Total: 10]	

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