

UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS General Certificate of Education Advanced Level

CHEMISTRY		9701/42
CENTRE NUMBER	CANDIDATE NUMBER	
CANDIDATE NAME		

Paper 4 Structured Questions

October/November 2013

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 a soft 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.

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.

For Exam	iner's Use
1	
2	
3	
4	
5	
6	
7	
8	
Total	

This document consists of 16 printed pages and 4 blank pages.



Section A

Answer all the questions in the spaces provided.

1 (a) Gaseous ammonia reacts with gaseous hydrogen chloride to form solid ammonium chloride.

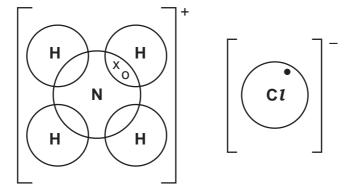
$$NH_3 + HCl \rightarrow NH_4Cl$$

The bonding in ammonium chloride includes ionic, covalent and co-ordinate (dative covalent) bonds.

Complete the following 'dot-and-cross' diagram of the bonding in ammonium chloride. For **each** of the six atoms show **all** the electrons in its outer shell. Three electrons have already been included.

Use the following code for your electrons.

- electrons from chlorine
- x electrons from hydrogen
- o electrons from nitrogen



[3]

- **(b)** When a sample of dry ammonia is needed in the laboratory, the gas is passed through a tower containing lumps of solid calcium oxide, CaO.
 - (i) Suggest why the usual drying agent for gases, concentrated H₂SO₄, is **not** used for ammonia.

(ii) Write an equation for the reaction between CaO and H₂O.

.....

(iii) Suggest why CaO rather than MgO is used to dry ammonia.

[3]

(c)	(i)	Write an equation showing the thermal decomposition of calcium nitrate, $Ca(NO_3)_2$.	For Examiner's Use
	(ii)	State and explain how the thermal stabilities of the nitrates vary down Group II.	
		[4]	
		[Total: 10]	

element	melting point/K
С	3925
Si	1683
Ge	1210
Sn	505

Suggest an explanation for each of the following.

2

(i)	The melting point of silicon is less than that of carbon.
(ii)	The melting point of tin is less than that of germanium.
	[2]

(b) Using data from the *Data Booklet* where appropriate, write equations for the following reactions of compounds of Group IV elements.

(i)	$SiCl_4(I) + H_2O(I)$				

(ii) the action of heat on $PbCl_4(I)$

|--|--|

(iv) SnO₂(s) + NaOH(aq)

[4]

[Total: 6]

3

For

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(a)	(i)	-	to represent a Brønsted-Lowry acid, write equations which stances acting as Brønsted-Lowry bases.
		NH ₃ +	\rightarrow
		CH ₃ OH +	\rightarrow
	(ii)	-	o represent a Brønsted-Lowry base, write equations which stances acting as Brønsted-Lowry acids.
		NH ₃ +	\rightarrow
		CH ₃ OH +	\rightarrow
			[4]
(b)	Sta	te briefly what is meant l	by the following terms.
	(i)	reversible reaction	
	(ii)	dynamic equilibrium	
			[2]
(c)	(i)	Explain what is meant to	by a buffer solution.
	(ii)		g of a buffer solution relies on a reversible reaction involving such as HZ and a Brønsted-Lowry base such as Z
			[4]

- (d) Propanoic acid, $CH_3CH_2CO_2H$, is a weak acid with $K_a = 1.34 \times 10^{-5} \, \text{mol dm}^{-3}$.
 - (i) Calculate the pH of a 0.500 mol dm⁻³ solution of propanoic acid.

Buffer solution **F** was prepared by adding 0.0300 mol of sodium hydroxide to 100 cm³ of a 0.500 mol dm⁻³ solution of propanoic acid.

(ii) Write an equation for the reaction between sodium hydroxide and propanoic acid.

(iii) Calculate the concentrations of propanoic acid and sodium propanoate in buffer solution **F**.

[propanoic acid] = $mol dm^{-3}$

[sodium propanoate] = mol dm⁻³

(iv) Calculate the pH of buffer solution F.

pH =[6]

(e) Phenyl propanoate cannot be made directly from propanoic acid and phenol. Suggest the identities of the intermediate **G**, the reagent **H** and the by-product **J** in the following reaction scheme.

$$CH_3CH_2CO_2H$$
 \longrightarrow G \longrightarrow ONa \longleftrightarrow O \longleftrightarrow O \longleftrightarrow O \longleftrightarrow O

G is

H is

J is

[2]

[Total: 18]

4	(a)	Exp	lain what is mean	t by the term bon	d energy.	
	(b)	(i)	Describe and exp where X = F, C <i>l</i> , I		ond energies	[2] s of the C–X bond in halogenoalkanes,
		(ii)	Describe the rela	=	the reactiv	ity of halogenoalkanes, RX, and the
	(c)	mu		the ozone layer	•	[3] as to why CFCs such as $\mathrm{CF_2C}l_2$ are arbons such as $\mathrm{CF_4}$ or hydrocarbons
						[3]
	(d)	Pre belo	dict the products ow. The molecular	of the following r formula of each p	reactions an oroduct is given	d draw their structures in the boxes ven, where $X = Cl$, Br or I.
		H ₂ C	+ Cl_	CI		$C_3H_5O_2X$
		H ₂ C	+ 1	✓ C1		C ₃ H ₇ OX

[3]

 C_7H_7OX

H₂O +

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(e) Ethane reacts with chlorine according to the following equation.

$$C_2H_6 + Cl_2 \rightarrow C_2H_5Cl + HCl$$

(i) State the conditions needed for this reaction.

(ii) State the type of reaction occurring here.

.....

One of the steps during this reaction is the following process.

$$Cl^{\bullet} + CH_3CH_3 \rightarrow HCl + CH_3CH_2^{\bullet}$$

(iii) Use the *Data Booklet* to calculate the enthalpy change, ΔH , of this step.

 $\Delta H = \dots kJ \text{ mol}^{-1}$

(iv) Use the *Data Booklet* to calculate the enthalpy change, ΔH , of the similar reaction:

 $\Delta H = \dots kJ \, \text{mol}^{-1}$

(v) Hence suggest why it is **not** possible to make iodoethane by reacting together iodine and ethane.

(vi) Complete the following equations of some possible steps in the formation of chloroethane.

$$Cl_2 \rightarrow \dots$$

$$Cl^{\bullet} + CH_3CH_3 \rightarrow HCl + CH_3CH_2^{\bullet}$$

$$CH_3CH_2^{\bullet}$$
 + \rightarrow +

$$..... + \rightarrow \mathsf{CH_3CH_2C}\mathit{l}$$

[8]

[Total: 19]

5 Super-absorbent polymers have the ability to absorb 200-300 times their own mass of water. They are classified as hydrogels and they are widely used in personal disposable hygiene products such as babies' nappies (diapers).

These polymers are commonly made by the polymerisation of compound ${\bf K}$ mixed with sodium hydroxide in the presence of an initiator.

$$CH_2 = C$$
 CO_2H

(a)	(i)	Explain what is meant by the term <i>polymerisation</i> .
	(ii)	What type of polymerisation is involved in the formation of hydrogels?
	(iii)	Describe the changes in chemical bonding that occur during the polymerisation of K .
		[3]
(b)		ylic acid is the common name for compound K . ggest the systematic (chemical) name of K .
		[1]
(c)	(i)	Draw the structure of at least two repeat units of the polymer formed by the above method from acrylic acid, K , when mixed with NaOH.
	(ii)	The C–C–C bond angle in compound ${\bf K}$ changes when the polymer is formed. State and explain how the C–C–C bond angle differs between a molecule of ${\bf K}$ and the polymer.
		angle changes from to
		explanation
		[4]

(d)	(i)	Draw a detailed diagram of a portion of the polymer you have drawn in (c)(i) to explain how it can absorb a large volume of water.
	/ 110	
	(11)	A student added 0.10 g of the polymer to 10 cm ³ of aqueous copper(II) sulfate solution. Predict, with a reason, what you expect to observe.
		[4]
(e)		mpound \mathbf{L} , $\mathrm{CH_2}$ = $\mathrm{CHCONH_2}$, can also be polymerised to form a super-absorbent ymer.
	(i)	Name the two functional groups in compound L .
	Cor	mpound K can be converted into compound L by the following two-step route.
H ₂ C	H	CO_2H $\frac{\text{step 1}}{\text{H}_2C}$ $\frac{\text{H}}{\text{C}}$ $\frac{\text{Step 2}}{\text{CO}_2^-\text{NH}_4^+}$ $\frac{\text{step 2}}{\text{H}_2C}$ $\frac{\text{H}}{\text{C}}$ $\frac{\text{CONH}_2}{\text{CONH}_2}$
	K	L.
	(ii)	Suggest a reagent for step 1.
((iii)	What other product is formed in step 2?

[Total: 17]

[5]

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(iv) State the reagents and conditions necessary to re-form **K** from **L**.

Answer all the questions in the spaces provided.

6 (a) Protein molecules are formed by the polymerisation of amino acids in the body. The structures of three amino acids are given.

$$H_2N$$
 OH OH_2 OH_2 OH_3 OH_4 OH_4 OH_4 OH_5 OH_4 OH_5 OH_5 OH_6 OH_6 OH_6 OH_6 OH_6 OH_7 OH_8 OH_8 OH_8 OH_8 OH_8 OH_8 OH_9 OH_9

(i)	How many	different	tripeptides	can	be	made	using	one	molecule	of	each	of	the
	amino acids	s shown?											

(ii) Draw the tripeptide ser-gly-val, showing the peptide bonds in displayed form.

iii)	Within the tripeptide, which amino acid provides a hydrophobic side chain?

(iv) Polypeptide chains can form bonds giving proteins their secondary and tertiary structures.

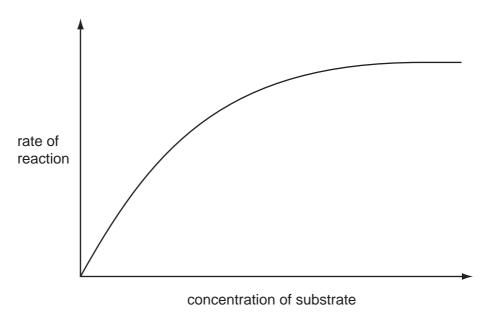
Using the tripeptide in (ii), state two types of bonding that can be formed and the groups in the tripeptide that are involved in this bonding.

bond	groups
bond	groups[6]

(b) Enzymes are particular types of proteins that catalyse chemical reactions. The efficiency of enzymes can be reduced by the presence of other molecules known as inhibitors. Explain how both *competitive* and *non-competitive* inhibitors prevent enzymes from working efficiently.

(i)	competitive inhibitors
()	
(ii)	non-competitive inhibitors
` ,	

(iii) The graph shows the rate of an enzyme-catalysed reaction against the substrate concentration in the absence of an inhibitor.



On the same axes, sketch a graph showing the rate of this reaction if a non-competitive inhibitor was present.

[4]

[Total: 10]

Electrophoresis is a technique which can be used fragments present in a mixture.	d to separate amino acids or peptide
fragments present in a mixture. (a) Draw a diagram to show the apparatus used to label each of the relevant parts of the apparatus.	
	[4]
(b) How far an amino acid will travel during electrophore For a given potential difference, state two other amino acid travels in a fixed time during electrophore.	r factors that will affect how far a given
1	
2	
	[2]
(c) A number of analytical and separation technique partition coefficients.	es rely on substances having different
State what is meant by the term partition coeffici	ent.
	[1]

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For Examiner's Use (d) The partition coefficient of X between ethoxyethane and water is 40.0.
 A solution contains 4.00 g of X dissolved in 0.500 dm³ of water.

 Calculate the mass of X that can be extracted from this aqueous solution by shaking it with

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(i) 0.050 dm³ of ethoxyethane,

(ii) two successive portions of 0.025 dm³ of ethoxyethane.

[4]

[Total: 11]

- In a world with a rapidly increasing population, access to clean drinking water is critical. For many countries, groundwater sources, rather than stored rainwater or river-water, are vital. *Groundwater* is water that exists in the pore spaces and fractures in rock and sediment beneath the Earth's surface. The World Health Organisation (WHO) provides maximum recommended concentrations for different ions present in drinking water.
 - (a) The geological nature of the soil determines the chemical composition of the groundwater. The table shows some ions which may contaminate groundwater.

ion present	WHO maximum permitted concentration/mg dm ⁻³
Ba ²⁺	0.30
Cl-	250.00
NO ₃ -	50.00
Pb ²⁺	0.01
Na⁺	20.00
SO ₄ ²⁻	500.00

	(i)	Nitrate, NO ₃ ⁻ , ions are difficult to remove from groundwater. What is the reason for this?
	(ii)	State which ions in the table above are likely to be removed from the water by treatment with powdered limestone, CaCO ₃ , giving reasons for each of your answers.
		[4]
b)		ates and phosphates can enter water courses such as rivers or streams as a result luman activity. Both of these ions are nutrients for algae.
	(i)	What is the origin of these nitrates?

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(ii) S	uggest an origin for the phosphates found in water courses.
(iii) W	/hat effect do nitrates and phosphates have on water courses?
	ro1
	[3]
• •	ain can have a major impact on natural waters, particularly lakes. In recent years has been a worldwide effort to reduce the amount of acid rain produced.
(i) W	/rite equations to show the production of acid rain from sulfur dioxide, SO ₂ .
	he use of fossil fuels is one major source of sulfur dioxide. ame another major industrial source.
	[2]
	[Total: 9]

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