

#### UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS General Certificate of Education Advanced Level

CANDIDATE NAME					
CENTRE NUMBER			CANDIDATE NUMBER		



CHEMISTRY 9701/43

Paper 4 Structured Questions

October/November 2011

2 hours

Candidates answer on the Question Paper.

Additional Materials: Data Booklet

#### READ THESE INSTRUCTIONS FIRST

Write your name, Centre number and candidate number 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 ON 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.



# Section A

For Examiner's Use

Answer all questions in the spaces provided.

1

(a)	Con	Complete the electronic configurations of the following ions.							
	Cr <sup>3</sup>	<b>+</b> :	1s <sup>2</sup> 2s <sup>2</sup> 2p <sup>6</sup>						
	Mn <sup>2+</sup> :		1s <sup>2</sup> 2s <sup>2</sup> 2p <sup>6</sup>					[2]	
(b)	) Both k		KMnO <sub>4</sub> and K <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub> are used as oxidising agents, usually in acidic solution.						
	(i)			om the <i>Data Boo</i> the solution incr		in why th	eir oxidising	power increases	
	(ii)		nat colour chan mpletely reduce	-	observe who	en each	of these oxi	dising agents is	
		•	KMnO <sub>4</sub>	from			to		
		•	K <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub>	from			to	[4]	
(c)	Pas	sing	a stream of S		a suspension			and dilute acids. does, however,	
	(i)			oklet to suggestidation states of				nd explain what the reaction.	
	(ii)			pension of MnC t, if any, this wo			nt of this rea	ction.	
								[4]	

(d) The main ore of manganese, pyrolusite, is mainly  $MnO_2$ . A solution of  $SnCl_2$  can be used to estimate the percentage of  $MnO_2$  in a sample of pyrolusite, using the following method.

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- A known mass of pyrolusite is warmed with an acidified solution containing a known amount of SnCl<sub>2</sub>.
- The excess Sn<sup>2+</sup>(aq) ions are titrated with a standard solution of KMnO<sub>4</sub>.

In one such experiment,  $0.100\,\mathrm{g}$  of pyrolusite was warmed with an acidified solution containing  $2.00\times10^{-3}\,\mathrm{mol}\,\mathrm{Sn^{2+}}$ . After the reaction was complete, the mixture was titrated with  $0.0200\,\mathrm{mol}\,\mathrm{dm^{-3}}\,\mathrm{KMnO_4}$ , and required  $18.1\,\mathrm{cm^3}$  of this solution to reach the end point.

The equation for the reaction between  $Sn^{2+}(aq)$  and  $MnO_{\Delta}^{-}(aq)$  is as follows.

$$2\mathsf{MnO_4}^- + 5\mathsf{Sn^{2+}} + 16\mathsf{H^+} \longrightarrow 2\mathsf{Mn^{2+}} + 5\mathsf{Sn^{4+}} + 8\mathsf{H_2O}$$

(i) Use the Data Booklet to construct an equation for the reaction between  ${\rm MnO_2}$  and  ${\rm Sn^{2+}}$  ions in acidic solution.

\_\_\_\_\_\_\_

- (ii) Calculate the percentage of MnO<sub>2</sub> in this sample of pyrolusite by the following steps.
  - number of moles of MnO<sub>4</sub><sup>-</sup> used in the titration
  - number of moles of Sn<sup>2+</sup> this MnO<sub>4</sub><sup>-</sup> reacted with
  - number of moles of Sn<sup>2+</sup> that reacted with the 0.100 g sample of pyrolusite
  - number of moles of MnO<sub>2</sub> in 0.100 g pyrolusite. Use your equation in (i).
  - mass of MnO<sub>2</sub> in 0.100 g pyrolusite
  - percentage of MnO<sub>2</sub> in pyrolusite

percentage = .....%

ſΟÌ

[Total: 16]

2	(a)	(i)	What is meant b	y the term	ligand as	applied to	o the	chemistry	of the	transition
			elements?							

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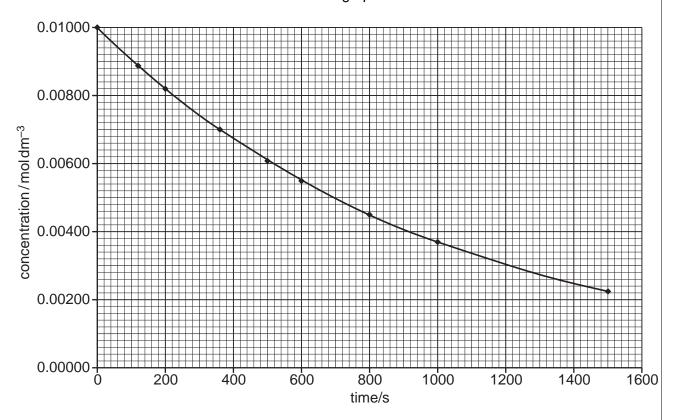
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(ii) Describe the type of bonding that occurs between a ligand and a transition element.

(b) Chromium hexacarbonyl undergoes the following ligand replacement reaction.

$$\operatorname{Cr(CO)}_6 + \operatorname{PR}_3 \rightarrow \operatorname{Cr(CO)}_5 \operatorname{PR}_3 + \operatorname{CO}$$

Two separate experiments were carried out to study the rate of this reaction. In the first experiment, the ligand  $PR_3$  was in a large excess and  $[Cr(CO)_6]$  was measured with time. The results are shown on the graph below.



In the second experiment,  $Cr(CO)_6$  was in a large excess, and  $[PR_3]$  was measured with time. The following results were obtained.

time/s	[PR <sub>3</sub> ]/moldm <sup>-3</sup>
0	0.0100
120	0.0076
200	0.0060
360	0.0028

(i) Plot the data in the table on the graph above, using the same axis scales, and draw the best-fit line through your points.

In e	each case explain how you arrived at yo	
Cr(	CO) <sub>6</sub>	
PR <sub>:</sub>	3	
	ite the rate equation for the reaction, anng the method of initial rates, or any oth	
 Sta	Ite the units of the rate constant.	
 Sta		
 Fou		n are given below. Draw a <b>circle</b> ar
Fouthe	ute the units of the rate constant.  ur possible mechanisms for this reactio letter next to the <b>one</b> mechanism which	n are given below. Draw a <b>circle</b> ar
Fou the hav	ate the units of the rate constant.  our possible mechanisms for this reaction letter next to the <b>one</b> mechanism which we written in <b>(iii)</b> . $\operatorname{Cr(CO)}_6 \rightarrow \operatorname{Cr(CO)}_5 + \operatorname{CO}$	n are given below. Draw a <b>circle</b> ar n is consistent with the rate equation fast
Fou the hav <b>A</b>	are the units of the rate constant.  our possible mechanisms for this reaction letter next to the <b>one</b> mechanism which we written in <b>(iii)</b> . $Cr(CO)_6 \rightarrow Cr(CO)_5 + CO$ $Cr(CO)_5 + PR_3 \rightarrow Cr(CO)_5 PR_3$ $Cr(CO)_6 \rightarrow Cr(CO)_5 + CO$	n are given below. Draw a <b>circle</b> are is consistent with the rate equation fast slow  slow fast fastPR <sub>3</sub> ] → Cr(CO) <sub>5</sub> PR <sub>3</sub> + CO
Fou the hav <b>A</b>	ar possible mechanisms for this reaction letter next to the <b>one</b> mechanism which we written in <b>(iii)</b> . $Cr(CO)_{6} \rightarrow Cr(CO)_{5} + CO$ $Cr(CO)_{5} + PR_{3} \rightarrow Cr(CO)_{5}PR_{3}$ $Cr(CO)_{6} \rightarrow Cr(CO)_{5} + CO$	n are given below. Draw a <b>circle</b> are is consistent with the rate equation fast slow  slow fast fastPR <sub>3</sub> ] → Cr(CO) <sub>5</sub> PR <sub>3</sub> + CO
Fou the hav A B	ar possible mechanisms for this reaction letter next to the <b>one</b> mechanism which we written in <b>(iii)</b> . $Cr(CO)_{6} \rightarrow Cr(CO)_{5} + CO$ $Cr(CO)_{5} + PR_{3} \rightarrow Cr(CO)_{5}PR_{3}$ $Cr(CO)_{6} \rightarrow Cr(CO)_{5} + CO$ $Cr(CO)_{6} \rightarrow PR_{3} \rightarrow Cr(CO)_{5}PR_{3}$ $Cr(CO)_{6} + PR_{3} \rightarrow Cr(CO)_{6}PR_{3}$ $Cr(CO)_{6} + PR_{3} \rightarrow Cr(CO)_{6}PR_{3}$	n are given below. Draw a <b>circle</b> are is consistent with the rate equation fast slow  slow fastPR <sub>3</sub> ] → Cr(CO) <sub>5</sub> PR <sub>3</sub> + CO ate) slow
Fou the hav A B	ar possible mechanisms for this reaction letter next to the <b>one</b> mechanism which we written in <b>(iii)</b> . $Cr(CO)_{6} \rightarrow Cr(CO)_{5} + CO \\ Cr(CO)_{5} + PR_{3} \rightarrow Cr(CO)_{5}PR_{3}$ $Cr(CO)_{6} \rightarrow Cr(CO)_{5} + CO \\ Cr(CO)_{5} + PR_{3} \rightarrow Cr(CO)_{5}PR_{3}$ $Cr(CO)_{6} \rightarrow PR_{3} \rightarrow Cr(CO)_{5}PR_{3}$ $Cr(CO)_{6} + PR_{3} \rightarrow Cr(CO)_{6}PR_{3}$ $Cr(CO)_{6} + PR_{3} \rightarrow Cr(CO)_{6}PR_{3}$ $Cr(CO)_{6} + PR_{3} \rightarrow Cr(CO)_{5}PR_{3} + CO$	n are given below. Draw a <b>circle</b> are is consistent with the rate equation fast slow  slow fastPR <sub>3</sub> ] → Cr(CO) <sub>5</sub> PR <sub>3</sub> + CO ate) slow

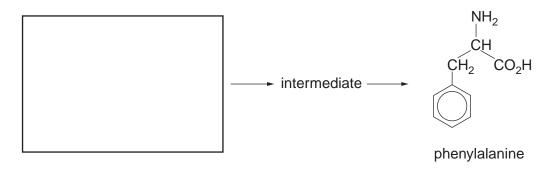
[Total: 11]

For Examiner's Use **3 (a)** Amino acids such as alanine are essential building blocks for making proteins. They can be synthesised by a general reaction of which the following is an example.

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$$\begin{array}{c|c} \text{CH}_3\text{CHO} & & \\ \hline & & \\ \hline & & \\ \text{CH}_3\text{CHO} & \\ \hline & & \\$$

- (i) Suggest the structure of the intermediate compound **E** by drawing its structural formula in the box above.
- (ii) Suggest, in the box below, the structural formula of the starting material needed to synthesise phenylalanine by the above general reaction.



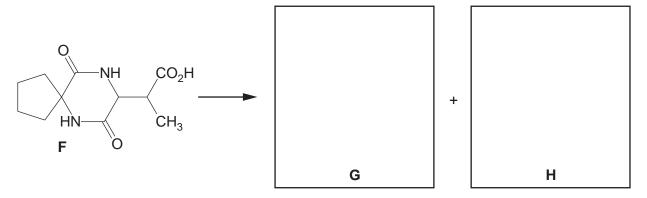
[2]

- (b) (i) What is a protein?
  - (ii) Using alanine as an example, draw a diagram to show how proteins are formed from amino acids. Show two repeat units in your answer.

[3]

(c) The hydrolysis of compound F produces two compounds G and H.





(i) State the reagents and conditions needed for this hydrolysis.

(ii) Draw the structures of the two products **G** and **H** in the boxes above.

[3]

(d) (i) Draw the zwitterionic structure of alanine.

(ii) Suggest the structural formulae of the zwitterions that could be formed from the following compounds.

compound	zwitterion
$H_2N$ — $CO_2H$	
OH NHCH <sub>3</sub>	
HO NH <sub>2</sub>	

[4]

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(e)	Sol	lutions of amino acids are good buffers.							
	(i)	What is meant by the term buffer?							
	(ii)	(ii) Write an equation to show how a solution of alanine, $CH_3CH(NH_2)CO_2H$ , behav as a buffer in the presence of an acid such as $HCl(aq)$ .							
	(iii)	Briefly describe how the pH of blood is controlled.							
	(iv) Calculate the pH of the buffer formed when $10.0\mathrm{cm^3}$ of $0.100\mathrm{moldm^{-3}}$ NaOH added to $10.0\mathrm{cm^3}$ of $0.250\mathrm{moldm^{-3}}$ CH <sub>3</sub> CO <sub>2</sub> H, whose p $K_\mathrm{a}$ = 4.76.								
		pH =[7]							
		[Total: 19]							

4

(a)	Writ	e an equation representing the action of heat on calcium nitrate, $\text{Ca(NO}_3)_2$ .
		[1]
(b)		cribe and explain the trend in the thermal stabilities of the nitrates of the Group II nents.
		[3]
(c)	Sod CO <sub>2</sub>	ium carbonate is stable to heat, but heating lithium carbonate readily produces $\varrho(g)$ .
	(i)	Suggest an equation for the action of heat on lithium carbonate.
	(ii)	Suggest a reason for the difference in reactivity of these two carbonates.
ı	(iii)	Predict what you would see if a sample of lithium nitrate was heated. Explain your answer.
		[4]
		[Total: 8]

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5	Alkanes are generally considered to be unreactive compounds, showing an inertness t common reagents such as NaOH, $\rm H_2SO_4$ , and $\rm K_2Cr_2O_7$ .						
	(a)	Sug	gest a reason why these reagents	do not attack an alkane such as CH <sub>4</sub> .			
				[1]			
	(b) When a mixture of chlorine and ethane gas is exposed to strong sunlight, an explorant occur due to the fast exothermic reaction. Under more controlled conditions, however, the following reaction occurs.						
			$C_2H_6 + Cl_2 \rightarrow$	$C_2H_5Cl + HCl$			
		(i)	What is the name of this type of re	eaction?			
		(ii)	Use equations to describe the involved.	mechanism of this reaction, naming the steps			
		(iii)					
	(	struc	tural formula of by-product	formed by			

(iv) It is found by experiment that, during this type of reaction, primary, secondary and tertiary hydrogen atoms are replaced by chlorine atoms at different rates, as shown in the following table.

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reaction	relative rate
$RCH_3 \rightarrow RCH_2Cl$	1
$R_2CH_2 \rightarrow R_2CHCl$	7
$R_3CH \rightarrow R_3CCl$	21

Using this information, and considering the number of hydrogen atoms of each type (primary, secondary or tertiary) within the molecule, predict the relative ratio of the two possible products **J** and **K** from the chlorination of 2-methylpropane. Explain your answer.

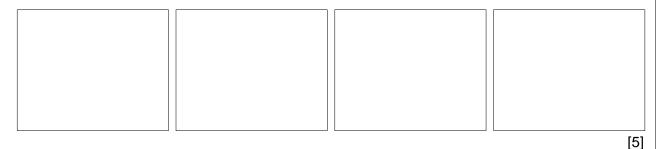
ratio **J**/**K** = .....

explanation:

		[10]

(c) In the boxes below draw the skeletal formulae of four different structural isomers of  $C_5H_{11}Cl$  that could be obtained from the chlorination of 2-methylbutane. Indicate any chiral centres in your structures by an asterisk (\*).

2-methylbutane



[3]

### Answer all questions in the spaces provided.

- **6** The formation of proteins is a key process in the growth and repair of tissues in living organisms.
  - (a) (i) Study the structures of the three molecules below. One of the molecules could be a building block for a protein while the other two could be building blocks for other biological polymers.

НО	ОН		CH <sub>2</sub> OH
	H H H OH OH	OH N H	HOH H
	J	K	L
	Which of the three c	ould be a building block for a p	protein? Explain your answer.
(ii)	For which biologica block?	I polymer could <b>one</b> of the c	other molecules form a building
	molecule	polymer	[2]
(b) Pro		four levels of structure as th	e long molecules fold and take
(i)		re is the sequence of amino at the sequence of amino acids in	acids in the protein chain. What this chain?
(ii)	What type of bondin	g can exist in <b>all</b> of the other ty	ypes of structure?
(iii)	Name one type of boot of the protein.	nding that does <b>not</b> occur in th	e primary or secondary structure

(c)		y proteins play an important role in catalysing chemical reactions in living nisms.	For Examiner's Use
	(i)	What name is given to these catalysts?	
	(ii)	Give <b>two</b> changes in conditions under which these catalysts may be inactivated, explaining the chemical reason for this in each case.	
		[4]	
		[Total: 9]	

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7 Different analytical techniques are used to build up a picture of complex molecules. Each technique on its own provides different information about complex molecules but together the techniques can give valuable structural information.

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(a) Complete the table, identifying the technique which can provide the appropriate structural information.

structural information	analytical technique
three-dimensional arrangement of atoms and bonds in a molecule	
chemical environment of protons in a molecule	
identity of amino acids present in a polypeptide	

[3]

(b)	One the	fly explain	
	(i)	paper chromatography	
	(ii)	thin-layer chromatography	
			[2]

(c) A combination of mass spectrometry and NMR spectroscopy is often enough to determine the structure of a simple organic compound. The organic compound N produced a mass spectrum in which the ratio of the M:M+1

peaks was 5.9:0.20, and which had an M+2 peak of similar height to the M peak.

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(i) Calculate how many carbon atoms are present in one molecule of N.

(ii) Deduce which element, other than carbon and hydrogen, is present in **N**.

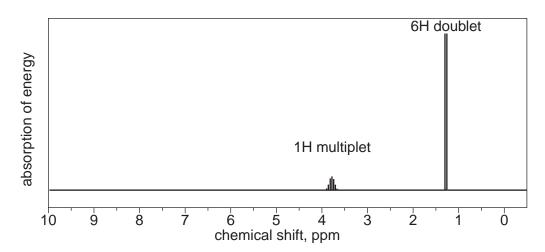
.....

(iii) Explain how many atoms of this element are present in one molecule of N.

.....

.....

The NMR spectrum of **N** is shown.



(iv) State the empirical formula of **N** and, using the NMR data, suggest the structural formula of **N**, explaining your reasons.

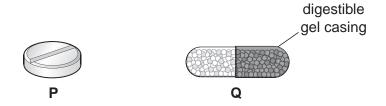
[6]

[Total: 11]

**8** Drugs can be delivered in a number of ways. The method chosen depends both on the nature of the drug, and the problem it is being used to treat.

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(a) Many common drugs are taken by mouth in forms similar to those shown.



(i)	Some drugs are available in solution. How would the speed of action of this form compare with <b>P</b> and <b>Q</b> ? Explain your answer.
(ii)	Explain which of the two forms, ${\bf P}$ or ${\bf Q}$ , would act the most rapidly when taken by mouth.
(iii)	Some drugs are broken down before they can be absorbed by the intestine. Suggest how the design of ${\bf Q}$ prevents this.
	[3]
into	er an abdominal operation drugs are often delivered by means of a 'drip' inserted a blood vessel in the patient's arm. Explain why this is more effective than taking akillers by mouth.
	[2]

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(b)

(c)	One of the molecules that has found a variety of uses in drug delivery is poly(ethylene glycol) or PEG. It is formed from dihydroxyethane, HOCH <sub>2</sub> CH <sub>2</sub> OH.			
	2n	$\mathrm{HOCH_2CH_2OH} \ \ \ \rightarrow \ \ \ \mathrm{H-(OCH_2CH_2OCH_2CH_2)_n-OH} \ + \ (2n-1)\ \mathrm{H_2O}$	Use	
	(i)	What type of reaction is this?		
	brok this:	ching a PEG molecule to a drug increases the time that it takes for the drug to be sen down and flushed from the body. There are thought to be two major reasons for firstly the PEG can form bonds to slow the passage of the drug around the body; andly it may reduce the efficiency of breakdown of the drug by enzymes.		
	(ii)	What type of bonds would the PEG part of the molecule form with molecules in the body?		
	(iii)	Suggest why attaching a PEG molecule to a drug molecule would reduce the rate of the drug's decomposition by enzymes.		
	(iv)	Drugs are often protein or polypeptide molecules. What type of reaction might occur in the breakdown of such a drug?		
		[5]		
		[Total: 10]		

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