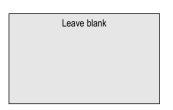
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Centre Number						Candida	ate Number		
Candidate Signat	ure								



General Certificate of Education June 2005 Advanced Level Examination



CHEMISTRY CHM5 Unit 5 Thermodynamics and Further Inorganic Chemistry (including Synoptic Assessment)

Tuesday 28 June 2005 Morning Session

In addition to this paper you will require:

a calculator.

Time allowed: 2 hours

Instructions

- Use blue or black ink or ball-point pen.
- Fill in the boxes at the top of this page.
- Answer **all** questions in **Section A** and **Section B** in the spaces provided. All working must be shown.
- Do all rough work in this book. Cross through any work you do not want marked.
- The Periodic Table/Data Sheet is provided on pages 3 and 4. Detach this perforated sheet at the start of the examination.
- **Section B** questions are provided on a perforated sheet. Detach this sheet at the start of the examination.

Information

- The maximum mark for this paper is 120.
- Mark allocations are shown in brackets.
- This paper carries 20 per cent of the total marks for Advanced Level.
- You are expected to use a calculator where appropriate.
- The following data may be required. Gas constant $R = 8.31 \text{ J K}^{-1} \text{ mol}^{-1}$
- Your answers to questions in **Section B** should be written in continuous prose, where appropriate. You will be assessed on your ability to use an appropriate form and style of writing, to organise relevant information clearly and coherently, and to use specialist vocabulary, where appropriate.

Advice

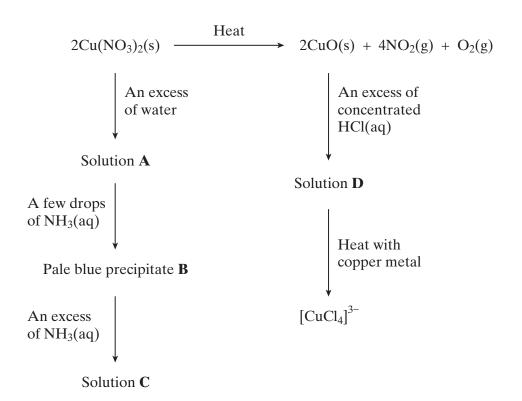
• You are advised to spend about 1 hour on **Section A** and about 1 hour on **Section B**.

For Examiner's Use									
Number	Mark	Number	er Mark						
1									
2									
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4									
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TOTAL									
Examine	r's Initials								

SECTION A

Answer all questions in the spaces provided.

1 Consider the reaction scheme below and answer the questions which follow.



(a)	A redox reaction occurs when Cu(NO ₃) ₂ is decomposed by heat. Deduce the oxidation
	state of nitrogen in Cu(NO ₃) ₂ and in NO ₂ and identify the product formed by oxidation
	in this decomposition.

Oxidation state of nitrogen in $Cu(NO_3)_2$

Oxidation state of nitrogen in NO₂

(b) Identify and state the shape of the copper-containing species present in s	olution A
--	-----------

Copper-containing species

Shape

(2 marks)

The Periodic Table of the Elements

■ The atomic numbers and approximate relative atomic masses shown in the table are for use in the examination unless stated otherwise in an individual question.

1 H H H H H H H			T	1							
1	0	4.0 He Helium 2	20.2 Ne Neon	39.9 Ar Argon 18	83.8 7.	Krypton 36	131.3 Xe	Xenon 54	222.0 Rn	Radon 86	
Frequency Freq	=		19.0 F Fluorine	35.5 C Chlorine 17	79.9 Br	Bromine 35	126.9 	lodine 3	210.0 At	Astatine 85	
H H H H H H H H H H	5		3.0 Oxygen	2.1 Sulphur	90.0 Se	selenium 4	127.6 Te	Tellurium 52	210.0 Po	Polonium 84	
H	>		4.0 N itrogen	1.0 Phosphoru 5	4.9 As	Arsenic 3	S1.8 Sb	Antin 1	209.0 Bi	Bismuth 83	
Hamman H	≥		12.0 C Carbon 6	28.1 Si Silicon	72.6 Ge	Germanium 32	118.7 Sn	Tin 50			
Hamman H	=		10.8 B Boron 5	27.0 AI Aluminium 13	69.7 Ga	Gallium 31	114.8 "	Indium 49	204.4 TI	Thallium 81	
Hearting Frequency Frequ					65.4 Zn	Zinc 30	112.4 Cd	Cadmium 48	200.6 Hg	Mercury 80	
Hearium Strontium Stront											
Hearthium Actinium Actinium					1		106.4 Pd	Palladium 46	195.1 Pt	Platinum 78	
Headium Lanthanum Hafnium Lanthanum Lanthanu					88.9 0	Cobalt 27	02.9 Rh	Rhodium 15	192.2 Ir	Iridium 77	
Headium Hernium Tantalum					55.8 Fe	Iron 26	101.1 Ru	Ruthenium 44	190.2 Os	Osmium 76	
Heryllium			6.9 Li Lithium 3		54.9 Mn	Manganese 25		Technetium 43			
Hearyllium			388 ———		52.0 Ç	Chromium 24	95.9 Mo	Molybdenum 42	183.9 W	ngsten	
## 137.3 138.9 Bardium 4			atomic ma umber —						180.9 Ta	Tantalum 73	
Beryllium 4 4 4 40.1 Calcium 12 Calcium 20 87.6 87.6 87.6 87.6 87.6 87.6 87.6 87.6		Key	relative a atomic n		47.9 Ti	anium	_	conium	178.5 H	Hafnium 72	
Beryllium 4 4 4 40.1 Calcium 12 Calcium 20 87.6 87.6 87.6 87.6 87.6 87.6 87.6 87.6						Scandium 21		Yttrium 39	138.9 La	Lanthanum 57 *	227 Actinium 89
	=		9.0 Be Beryllium 4	24.3 Mg Magnesium 12	á	Calcium 20		_		arinm	226.0 Ra Radium 88
	_	1.0 H Hydrogen				Potassium 19	Q	⊏	132.9 Cs	Caesium 55	Ę

175.0	Lutetium	(260)	ım Lawrencium
Lu	71	Ľ	103
173.0	Ytterbium	(259)	Nobeliu
Yb	70	No	102
168.9 Tm			Mendelevium 101
167.3	Erbium	257)	Fermium
Er	68	Fm	00
164.9 Ho	olmium	52) ES	ısteinium
162.5	Dysprosium	247.1 252.1 (29	Californium
Dy	66	Bk Cf	98
158.9	Terbium	247.1	Berkelium
Tb	65	BK	97
157.3	3adoliniur	247.1	Curium
Gd	34	Cm	96
52.0	uropium	243.1	Americium
Eu	3	Am	35
50.4	Samarium	239.1 2	Plutonium
Sm	32		94
144.9	Promethiur	237.0	
Pm	61	Np	
144.2	Neodymium	238.0	Uranium
Nd	60	U	92
40.1 140.9 144.2 Ce Pr Nd	Praseodymium Neodymium 59 60	232.0 231.0 238.0 Th Pa Da	Protactinium Uranium 91
40.1	Cerium 58	232.0	Thorium
Ce		Th	90

* 58 - 71 Lanthanides

† 90 - 103 Actinides

Table 1 Proton n.m.r chemical shift data

Type of proton	δ/ppm
RCH_3	0.7–1.2
R_2CH_2	1.2–1.4
R_3CH	1.4–1.6
$RCOCH_3$	2.1–2.6
$ROCH_3$	3.1–3.9
$RCOOCH_3$	3.7–4.1
ROH	0.5–5.0

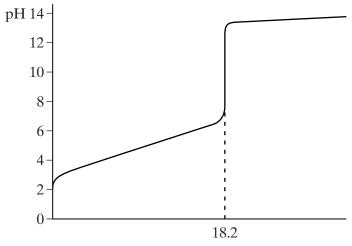
Table 2 Infra-red absorption data

Bond	Wavenumber/cm ⁻¹
С—Н	2850-3300
С—С	750–1100
C = C	1620–1680
C=O	1680–1750
С—О	1000-1300
O—H (alcohols)	3230–3550
O—H (acids)	2500–3000

(c)	(i)	Identify the pale blue precipitate \mathbf{B} and write an equation, or equations, to show how \mathbf{B} is formed from the copper-containing species in solution \mathbf{A} .
		Identity of precipitate B
		Equation(s)
	(ii)	In what way does the NH ₃ behave as a Brønsted–Lowry base?
		(3 marks)
(d)	(i)	Identify the copper-containing species present in solution \mathbf{C} . State the colour of this copper-containing species and write an equation for its formation from precipitate \mathbf{B} .
		Identity
		Colour
		Equation
	(ii)	In what way does the NH ₃ behave as a Lewis base?
		(4 marks)
(e)		tify the copper-containing species present in solution D . State the colour and shape is copper-containing species.
	Iden	tity
	Colo	ur
	Shap	e(3 marks)
(f)	The	exidation state of copper in $[CuCl_4]^{3-}$ is +1.
` ,	(i)	Give the electron arrangement of a Cu ⁺ ion.
	(ii)	Deduce the role of copper metal in the formation of $[CuCl_4]^{3-}$ from the copper-containing species in solution D .
		(2 marks)



2 The pH curve shown below was obtained when a 0.150 mol dm⁻³ solution of sodium hydroxide was added to 25.0 cm³ of an aqueous solution of a weak monoprotic acid, HA.



Volume of $0.150\,\mathrm{mol\,dm}^{-3}\,\mathrm{NaOH}$ added/cm³

(a)	Use the information given to calculate the concentration of the acid.						
	•••••						
	•••••	(2 marks)					
(b)	(i)	Write an expression for the acid dissociation constant, K_a , for HA.					
	(ii)	Write an expression for pK_a					
	(iii)	Using your answers to parts (b)(i) and (b)(ii), show that when sufficient sodium hydroxide has been added to neutralise half of the acid,					
		pH of the solution = pK_a for the acid HA					
		(4 marks)					

(c)	-	Explain why dilution with a small volume of water does not affect the pH of a buffer solution.					
	•••••						
	*****	(2 marks)					
(d)	(i)	Calculate the change in pH when $0.250\mathrm{moldm^{-3}}$ hydrochloric acid is diluted with water to produce $0.150\mathrm{moldm^{-3}}$ hydrochloric acid.					
	(ii)	Calculate the volume of water which must be added to $30.0\mathrm{cm^3}$ of $0.250\mathrm{moldm^{-3}}$ hydrochloric acid in order to reduce its concentration to $0.150\mathrm{moldm^{-3}}$.					
		(4 marks)					



TURN OVER FOR THE NEXT QUESTION

(3 marks)

3 (a) (i) The addition of aqueous silver nitrate, ammonia, can be used to distinguish between sodium bromide and sodium iodide. Record what is observed in the table below.						between separate aqueous solutions of				
				The addition of AgNO3(aq)	followed	by the addition of concentrated NH3(aq)				
			Observation with NaBr(aq)							
			Observation with NaI(aq)							
		(ii)	Explain why it is no nitrate and sodium			een separate solutions of sodium ver nitrate solution. (5 marks)				
	(b)		n aqueous sodium that colourless solution		led to solid	silver bromide a reaction occurs				
		(i)	Identify the silver-o	containing species	present in tl	ne colourless solution.				
		(ii)	Write an equation	for this reaction.						
		(iii)	Give one use of thi	s reaction.						

ethanoyl chloride.

(c) Aqueous silver nitrate can be used to distinguish between chloroethanoic acid and

	(i)	Draw the structure of ethanoyl chloride. Predict what, if anything, you would observe when ethanoyl chloride is added to aqueous silver nitrate.
		Structure of ethanoyl chloride
		Observation
	(ii)	Draw the structure of chloroethanoic acid. Predict what, if anything, you would observe when chloroethanoic acid is added to aqueous silver nitrate.
		Structure of chloroethanoic acid
		Observation
		(4 marks)
(d)	(i)	Tollens' reagent is formed by the addition of aqueous ammonia to aqueous silver nitrate. Identify the silver-containing complex present in Tollens' reagent and state its shape.
		Silver-containing complex
		Shape
	(ii)	Draw the structure of methanoic acid. By reference to this structure, suggest why a silver mirror is formed when this acid reacts with Tollens' reagent.
		Structure
		Explanation
	(iii)	Deduce the identity of a carbon-containing species formed when methanoic acid reacts with Tollens' reagent.
		(5 marks)

4 Where appropriate, use the standard electrode potential data in the table below to answer the questions which follow.

	<i>E</i> [⊕] /V
$Zn^{2+}(aq) + 2e^- \rightarrow Zn(s)$	-0.76
$V^{3+}(aq) + e^- \rightarrow V^{2+}(aq)$	-0.26
$SO_4^{2-}(aq) + 2H^+(aq) + 2e^- \rightarrow SO_3^{2-}(aq) + H_2O(1)$	+0.17
$VO^{2+}(aq) + 2H^{+}(aq) + e^{-} \rightarrow V^{3+}(aq) + H_2O(1)$	+0.34
$Fe^{3+}(aq) + e^{-} \rightarrow Fe^{2+}(aq)$	+0.77
$VO_2^+(aq) + 2H^+(aq) + e^- \rightarrow VO^{2+}(aq) + H_2O(l)$	+1.00
$Cl_2(aq) + 2e^- \rightarrow 2Cl^-(aq)$	+1.36

(a)	From the table above select the species which is the most powerful reducing agent.
	(1 mark)
(b)	From the table above select
	(i) a species which, in acidic solution, will reduce $VO_2^+(aq)$ to $VO^{2+}(aq)$ but will not reduce $VO^{2+}(aq)$ to $V^{3+}(aq)$,
	(ii) a species which, in acidic solution, will oxidise $VO^{2+}(aq)$ to $VO_2^+(aq)$.
	(2 marks)
(c)	The cell represented below was set up under standard conditions.
	$Pt Fe^{2+}(aq), Fe^{3+}(aq) Tl^{3+}(aq), Tl^{+}(aq) Pt$ Cell e.m.f. = + 0.48 V

(i) Deduce the standard electrode potential for the following half-reaction.

$$Tl^{3+}(aq) + 2e^{-} \rightarrow Tl^{+}(aq)$$

(ii) Write an equation for the spontaneous cell reaction.

(3 marks)

(d) After acidification, $25.0\,\mathrm{cm}^3$ of a solution of hydrogen peroxide reacted exactly with $16.2\,\mathrm{cm}^3$ of a $0.0200\,\mathrm{mol\,dm}^{-3}$ solution of potassium manganate(VII). The overall equation for the reaction is given below.

$$2MnO_4^- + 6H^+ + 5H_2O_2 \rightarrow 2Mn^{2+} + 8H_2O + 5O_2$$

(i)	Use the equation for this reaction to determine the concentration, in g dm ⁻³ , of the
(-)	hydrogen peroxide solution.
(ii)	Calculate the maximum volume of oxygen, measured at a pressure of 98 kPa and a temperature of 298 K, which would be evolved in this reaction.
	(8 marks)



NO QUESTIONS APPEAR ON THIS PAGE

SECTION B

Detach this perforated sheet. Answer **all** questions in the space provided on pages 15 to 20 of this booklet.

- 5 (a) State and explain the effect of a catalyst on the rate and on the equilibrium yield in a reversible reaction. (5 marks)
 - (b) Explain the terms *heterogeneous* and *active sites* as applied to a catalyst. Give **two** reasons why a ceramic support is used for the catalyst in catalytic converters in cars. Explain how lead poisons this catalyst. (7 marks)
 - (c) In aqueous solution, Fe²⁺ ions act as a homogeneous catalyst in the reaction between I^- and $S_2O_8^{2-}$ ions. Give **one** reason why the reaction is slow in the absence of a catalyst. Write equations to show how Fe²⁺ ions act as a catalyst for this reaction. (5 marks)
- 6 Most elements can be extracted from one of their naturally-occurring compounds by one of the following methods:

electrolysis of a molten chloride or oxide reduction of the oxide with carbon displacement of the element from its compound by a more reactive element.

(a) Aluminium is extracted by the electrolysis of aluminium oxide which has been dissolved in molten cryolite.

Suggest why aluminium is not extracted by electrolysis of aluminium chloride.

(2 marks)

(b) Use the information given below to calculate the minimum temperature at which the following carbon reduction process is feasible.

$$Al_2O_3(s) + 3C(s) \rightarrow 2Al(s) + 3CO(g)$$

	Al ₂ O ₃ (s)	Al(s)	CO(g)	C(s)
$\Delta H_{\rm f}^{\Theta}/{\rm kJmol}^{-1}$	-1676	0	-111	0
$S^{\Theta}/J K^{-1} \text{ mol}^{-1}$	51	28	198	6

(8 marks)

- (c) (i) State why fluorine cannot be prepared by displacement from one of its compounds by another halogen.
 - (ii) Identify a fluorine-containing product formed when solid sodium fluoride reacts with concentrated sulphuric acid. Write an equation for the reaction which occurs.

 (3 marks)

- 7 (a) By a consideration of the structure of and the bonding in each compound, explain why magnesium chloride has a high melting point and why silicon(IV) chloride has a low boiling point. (6 marks)
 - (b) State what is observed when separate samples of sodium oxide and phosphorus(V) oxide are added to water. Write equations for the reactions which occur and, in each case, state the approximate pH of the solution formed. (6 marks)
- 8 (a) Identify a reagent which reacts with ethanal by a nucleophilic addition reaction to form a racemic mixture. Write an equation for the reaction occurring. Explain why a racemic mixture is obtained. (4 marks)
 - (b) Identify a reagent which undergoes an addition reaction with propene to form a mixture of isomers. Outline a mechanism for the reaction occurring. Explain why isomers are obtained. (7 marks)
 - (c) Identify a reagent which reacts with 2-bromobutane to form a mixture of isomers by an elimination reaction. Outline a mechanism for this reaction, and draw the structures of the isomers obtained. (7 marks)

END OF QUESTIONS

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