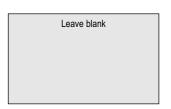
Surname				Other	Names			
Centre Nu	mber				Candida	ate Number		
Candidate	Signat	ure						



General Certificate of Education January 2003 Advanced Subsidiary Examination



CHEMISTRY CHM2 Unit 2 Foundation Physical and Inorganic Chemistry

Friday 10 January 2003 Morning Session

In addition to this paper you will require:
a calculator.

Time allowed: 1 hour 30 minutes

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.

Information

- The maximum mark for this paper is 90.
- Mark allocations are shown in brackets.
- This paper carries 30 per cent of the total marks for AS. For Advanced Level this paper carries 15 per cent of the total marks.
- You are expected to use a calculator where appropriate.
- The following data may be required. Gas constant $R = 8.31 \,\mathrm{J \, K^{-1} \, mol^{-1}}$
- 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 30 minutes on **Section B**.

	For Exam	iner's Use	
Number	Mark	Number	Mark
1			
2			
3			
4			
5			
6			
7			
Total (Column	1)	→	
Total (Column	2)	→	
TOTAL			
Examine	r's Initials		

SECTION A

Answer all questions in the spaces provided.

1 The table below contains some standard enthalpy of formation data.

Substance	C(s)	$N_2(g)$	H ₂ O(g)	CO ₂ (g)	NH ₄ NO ₃ (s)
$\Delta H_{\rm f}^{\Theta}/{\rm kJmol}^{-1}$	0	0	-242	-394	-365

(a)	Why are the values of the standard enthalpy of formation for carbon and nitrogen zero?
	(1 mark)
(b)	State Hess's Law.
	(2 marks)

(c) Use $\Delta H_{\rm f}^{\Theta}$ data from the table to calculate a value for the enthalpy change for the following reaction.

$$NH_4NO_3(s) + \frac{1}{2}C(s) \rightarrow N_2(g) + 2H_2O(g) + \frac{1}{2}CO_2(g)$$

$$(3 marks)$$



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.

_	=											≡	≥	>	5	=	0
T drogen			Key														4.0 He Helium 2
Li ithium	9.0 Be Beryllium 4		relative atomic mass atomic number ——	atomic ma umber —		6.9 Li Lithium 3						10.8 B Boron 5	12.0 C Carbon 6	14.0 N Nitrogen 7	16.0 Oxygen 8	9.0 F Fluorine	20.2 Ne Neon
0 Na odium					-							27.0 Al Aluminium 13	28.1 Si Silicon	AI Silcon Phosphorus Sulphur Aluminium Silicon Phosphorus Sulphur 13 14 15 16	32.1 S Sulphur 16	55.5 CI Chlorine 7	39.9 Ar Argon 18
1 K assium	40.1 Ca Calcium 20	Scandium 21	47.9 Ti Ti Titanium 22	50.9 Vana 23	L 52.0 54.9 55.8 58.9 58.7 Indiget Cromium Manganese Iron Cobatt Nicks 24 25 26 27 28	Mn Manganese 25	55.8 Fe Iron 26	58.9 Co Cobalt 27	<u></u>	63.5 Cu Copper 29	65.4 Zn Zinc 30	69.7 Ga Gallium 31	72.6 Ge Germanium 32	74.9 As As Arsenic 33	79.0 Se Selenium 34	79.9 Br Bromine	83.8 Kr Krypton 36
5 Rb bidium	87.6 Sr Strontium 38	88.9 Y Yttrium 39	91.2 Zr Zirconium 40	92.9 Niok	95.9 Mo Molybdenum 42	98.9 Tc Technetium 43	Ruthenium 44	102.9 Rh Rhodium 45	_	107.9 Ag Silver 47	112.4 Cd Cadmium 48	114.8 In Indium 49	Sn Tin	121.8 127.6 1 Sb Te Antimony 52 52 5	127.6 Te Tellurium 52	26.9 – lodine i3	131.3 Xe Xenon 54
Cs Cs Resium	137.3 Ba Barium 56	بد⊇ا	178.5 Hf Hafnium 72	180.9 T Tant 73	183.9 W Tungsten 74	186.2 Re Rhenium 75	190.2 Os Osmium 76	192.2 r r Iridium 77	Ę	197.0 Au Gold 79	200.6 Hg Mercury 80	204.4 Tl Thallium Thallium 81	207.2 Pb Lead 32	209.0 Bi Bismuth 83	210.0 Po Polonium 84	.10.0 At Astatine 55	222.0 Rn Radon 86
www.teea	226.0 Ra Radium 88	227 Ac Actinium 89 †															
llpapers.co	seded 2008	ınides		140.1 Ce Cerium 58	140.9 Pr Praseodymium 59	144.2 Nd Neodymium 60	Pm Pm Promethium 61	150.4 Sm Samarium 62	140.9 144.2 144.9 150.4 152.0 157.3 158.9 162.5 164.9 167.3 168.9 173.0 175.0 Praseodymium Promethium Promethiu	157.3 Gd Gadolinium 64	158.9 Tb Terbium 65	162.5 Dy Dysprosium 66	164.9 Ho Holmium 67	167.3 Er Erbium 68	168.9 Tm Thulium 69	173.0 Yb Ytterbium 70	175.0 Lu Lutetium 71
om <u>0</u>	03 Actini	ges		232.0 Th Thorium	232.0 231.0 238.0 Th Pa U Thorium Protactinium Uranium	238.0 U Uranium	237.0 Np Neptunium	239.1 Pu Plutonium	243.1 Am Americium	247.1 Cm Curium	247.1 BK Berkelium	252.1 Cf Californium	(252) ES Einsteinium	(257) Fm Fermium	(258) Md Mendelevium	(259) Nobelium	(260) Lr Lawrencium

140.1 Ce	140.9 Pr	144.2 Nd	٦٤	150.4 Sm	152.0 Eu	157.3 Gd	158.9 Tb	162.5 Dy	164.9 Ho		168.9 Tm		175.0 Lu
Cerium F	Cerium Praseodymium Neodymium Prome 58 59 60 61	Neodymium 60	Ę	٤		٦			Holmium 67	Erbium 68		Ytterbium 70	Lutetium 71
232.0 Th	232.0 231.0 238.0 237.0 Th Pa U Np	238.0 U		239.1 Pu	- E	247.1 Cm	u	252.1 Cf	ွှ		_	l	(260) Lr
Thorium 90	Protactinium 91	Protactinium Uranium 91 92	Ę	_	Americium 95		Berkelium 97	Californium 98	E	Fermium 100	.E		Lawrencium 103

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_3 CH	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

2 The table below contains some mean bond enthalpy data.

Bond	Н—Н	С-С	C=C	N≡N	N-H
Mean bond enthalpy/kJ mol ⁻¹	436	348	612	944	388

(a)	Expl	ain the term mean bond enthalpy.
	•••••	
	•••••	
	•••••	
	•••••	(2 marks)
(b)	(i)	Write an equation for the formation of one mole of ammonia, NH ₃ , from its elements.
	(ii)	Use data from the table above to calculate a value for the enthalpy of formation of ammonia.
		(4 marks)

(c) Use the following equation and data from the table above to calculate a value for the C–H bond enthalpy in ethane.

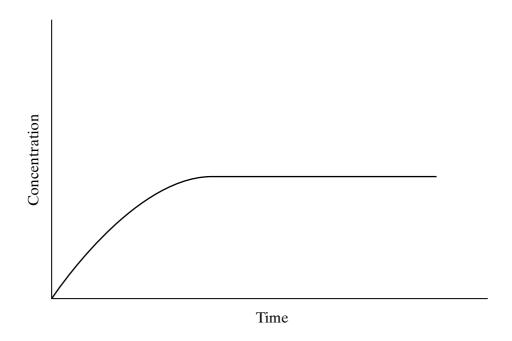
	H H	НН
(3 marks)		

3	(a)	Define the term activation energy for a reaction.
		(2 marks)
	(b)	Give the meaning of the term <i>catalyst</i> .
		(2 marks)
	(c)	Explain in general terms how a catalyst works.
		(2 marks)

(d) In an experiment, two moles of gas \mathbf{W} reacted completely with solid \mathbf{Y} to form one mole of gas \mathbf{Z} as shown in the equation below.

$$2W(g) + Y(s) \rightarrow Z(g)$$

The graph below shows how the concentration of \mathbf{Z} varied with time at constant temperature.



- (i) On the axes above, sketch a curve to show how the concentration of **W** would change with time in the same experiment. Label this curve **W**.
- (ii) On the axes above, sketch a curve to show how the concentration of \mathbf{Z} would change with time if the reaction were to be repeated under the same conditions but in the presence of a catalyst. Label this curve \mathbf{Z} .
- (iii) In terms of the behaviour of particles, explain why the rate of this reaction decreases with time.

••••••	•••••	•••••	(6 marks)

4	Methanol can be synthesised from carbon monoxide by the reversible reaction shown below.
	$CO(g) + 2H_2(g) \rightleftharpoons CH_3OH(g) \Delta H = -91 \text{ kJ mol}^{-1}$

The process operates at a pressure of $5\,\mathrm{MPa}$ and a temperature of $700\,\mathrm{K}$ in the presence of a copper-containing catalyst. This reaction can reach dynamic equilibrium.

(a) By reference to rates and concentrations, explain the meaning of the terr dynamic equilibrium.
(2 marks
(b) Explain why a high yield of methanol is favoured by high pressure.
(2 marks
(c) Suggest two reasons why the operation of this process at a pressure much higher tha 5 MPa would be very expensive.
Reason 1
Reason 2(2 marks
(d) State the effect of an increase in temperature on the equilibrium yield of methanol an explain your answer.
Effect
Explanation
(3 marks
(e) If a catalyst were not used in this process, the operating temperature would have to b greater than 700 K. Suggest why an increased temperature would be required.
(1 mark



5 (a)	Stat	State and explain the trend in electronegativity down Group VII from fluorine to iodine.				
	Tren	Trend				
	Exp	lanation				
	•••••	(3 marks)				
(b)	(i)	Describe what you would observe when an aqueous solution of bromine is added to an aqueous solution containing iodide ions. Write an equation for the reaction occurring.				
		Observation				
		Equation				
	(ii)	Explain why bromine does not react with aqueous chloride ions.				
		(3 marks)				
(c)		cribe what you would observe when aqueous silver nitrate is added to separate eous solutions of potassium fluoride and potassium bromide.				
	Obs	ervation with KF(aq)				
	Obs	ervation with KBr(aq)				
		(2 marks)				
(d)		te an equation to show how solid potassium fluoride reacts with concentrated huric acid.				
	•••••	(1 mark)				
(e)	Writ acid	te an equation for the redox reaction of sodium bromide with concentrated sulphuric.				
	•••••	(2 marks)				



6 (a)		In acidic conditions, hydrogen peroxide, H_2O_2 , oxidises iodide ions to iodine. The hydrogen peroxide is reduced to water. In H_2O_2 , oxygen has an oxidation state of -1 .		
		(i)	Construct a half-equation for the reduction of hydrogen peroxide to water in acidic conditions.	
		(ii)	Construct a half-equation for the oxidation of ${\rm I}^-$ ions to iodine.	
		(iii)	Construct an equation for the overall reaction.	
			(3 marks)	
aqueous sodium thiosulphate. In a titration reacted with exactly 19.5 cm ³ of a 0.120 molds		aque	concentration of an aqueous iodine solution can be determined by titration with sous sodium thiosulphate. In a titration, 25.0 cm ³ of an aqueous iodine solution ted with exactly 19.5 cm ³ of a 0.120 mol dm ⁻³ solution of sodium thiosulphate. Write an equation for the reaction between iodine and thiosulphate ions.	
		(-)		
		(ii)	Calculate the concentration of the iodine solution.	
			(If you are unable to answer part $(b)(i)$, assume that one mole of iodine reacts with three moles of thiosulphate ions. This is not the correct ratio.)	
			(5 marks)	

(c) Chlorine reacts with water as shown in the following equation.

$$Cl_2 + H_2O \implies Cl^- + HClO + H^+$$

In this reaction, chlorine acts both as an oxidising agent and as a reducing agent.

Construct a half-equation for the reduction of chlorine to chloride ions.

-
- (ii) Deduce the oxidation state of chlorine in HClO.
- (iii) Construct a half-equation for the oxidation of chlorine, in reaction with water, to form HClO and H⁺ ions.
- (iv) Give **one** reason why chlorine is used in the water industry.

(4 marks)



TURN OVER FOR THE NEXT QUESTION

SECTION B

Answer the question below in the space provided on pages 12 to 16 of this booklet.

- 7 In this question, credit will be given for appropriate equations.
 - (a) Describe the process by which titanium metal is extracted from titanium(IV) oxide. Explain why this extraction process causes titanium to be an expensive metal.

 (11 marks)

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- (b) Explain how the impurity silicon(IV) oxide is removed during the extraction of iron in the Blast Furnace.
 - Discuss the chemical reactions by which impurities are removed from iron in the basic oxygen converter. (13 marks)
- (c) Describe how scrap iron is recycled. Discuss the social and environmental benefits of this recycling. (6 marks)

END OF QUESTIONS