

Other Names

Surname

GCE A level

1095/01

CHEMISTRY CH5

A.M. FRIDAY, 24 June 2011

13/4 hours

FOR EXAMINER'S USE ONLY			
Section	Question	Mark	
A B	1		
	2		
	3		
	4		
	5		
TOTAL MARK			

ADDITIONAL MATERIALS

In addition to this examination paper, you will need:

- a calculator
- an 8 page answer book;
- a copy of the **Periodic Table** supplied by WJEC. Refer to it for any **relative atomic masses** you require.

INSTRUCTIONS TO CANDIDATES

Use black ink or black ball-point pen.

Write your name, centre number and candidate number in the spaces at the top of this page.

Section A Answer all questions in the spaces provided.

Section B Answer both questions in Section B in a separate answer book which should then be placed inside this question-and-answer book.

Candidates are advised to allocate their time appropriately between **Section A (40 marks)** and **Section B (40 marks)**.

INFORMATION FOR CANDIDATES

The number of marks is given in brackets at the end of each question or part-question.

The maximum mark for this paper is 80.

Your answers must be relevant and must make full use of the information given to be awarded full marks for a question.

You are reminded that marking will take into account the Quality of Written Communication in all written answers.

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SECTION A

Answer all questions in the spaces provided.

 Chromium and aluminium both form amphoteric compounds. (a) State what is meant by the term <i>amphoteric</i>. 					unds.	[1]	
	(b)	Use Cr ³⁺		o show the e	electronic structu	ares of the chromit	um atom and the
		Chro	omium atom, Cr				
1s		2s	2p	3s	3p	3d	4s
		Chro	omium(III) ion, Cı	-3+			
1s	_	2s	2p	3s	3p	3d	4s
	(c)		r_2O_7 , the following	g reaction oc	curs.	ution of potassium	dichromate(VI),
		(i)	$\operatorname{Cr}_2\operatorname{O}_7^{2^{-2}}$ State the colour		\longrightarrow 2CrO ₄ ² -is seen.	+ H ₂ O	[1]
		(ii)	Use the oxidation	n states of ch	nromium to show	v that this is not a r	edox reaction. [2]
		•••••					

Examiner only

1095 010003

(d)	mag	minium chloride is a compound of the amphoteric element aluminium, whilst nesium chloride contains the non-amphoteric element magnesium. Explain how um hydroxide can be used to distinguish between solutions of these two compounds. [3]
(e)	Aluı	minium chloride, AlCl ₃ , commonly exists as the dimer Al ₂ Cl ₆ .
	(i)	Draw the structure of the dimer formed, and explain why the two AlCl ₃ monomers join together. [3]
	(ii)	Aluminium chloride monomer may combine with another chloride ion to form tetrachloroaluminate(III) ions, AlCl ₄ ⁻ . Using valence shell electron pair repulsion theory (VSEPR), state and explain the shape of this anion. [2]
	•••••	Total [14]

Total [14]

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- **2.** Fuel cells have been proposed as an alternative method of providing energy for vehicles. These use chemical reactions within electrochemical systems to generate electricity.
 - (a) A typical fuel cell uses hydrogen as a fuel and reacts this with oxygen. The two half-equations for the processes occurring at the electrodes are given in the table below.

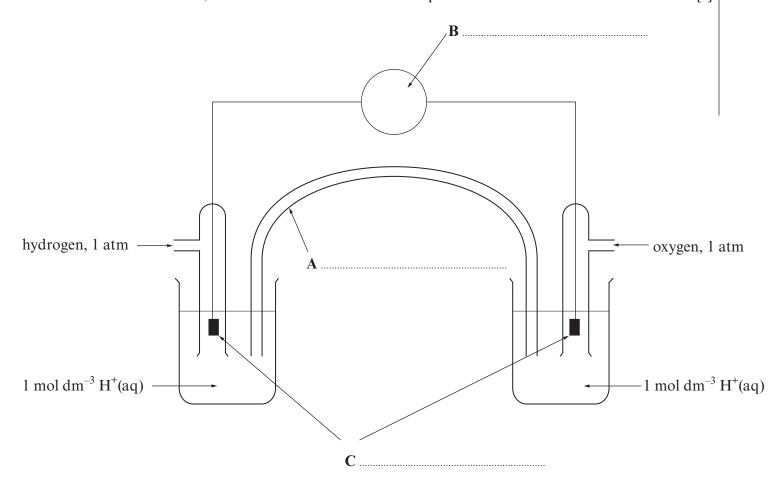
Half-equation	E ⇔ / V
$2H^+ + 2e^- \rightleftharpoons H_2$	0.00
$1/_2O_2 + 2H^+ + 2e^- \rightleftharpoons H_2O$	1.23

(i) Write an equation for the overall reaction occurring.

[1]

(ii) Give **one** benefit of the use of fuel cells as a replacement for traditional vehicle energy sources. [1]

(iii) The same reaction as above can be undertaken in a traditional electrochemical cell, such as the one below. Name the parts labelled **A-C**. [3]



(b) A different fuel for use in fuel cells is methanol, CH₃OH, which would undergo the following reaction with oxygen.

$$CH_3OH(l) \ + \ l^{1}\!/_{2}O_2(g) \ \longrightarrow \ CO_2(g) \ + \ 2H_2O(l)$$

Compound	Standard enthalpy change of formation, $\Delta H_{\mathrm{f}}^{-\Phi}/\mathrm{kJ}\mathrm{mol}^{-1}$
CH₃OH	-239
CO ₂	-394
H ₂ O	-286

(i)	Calculate the standard enthalpy change of combustion for methanol. [2	
		••
(ii)	The entropy change of this reaction is calculated as follows.	••
ΔS =	(Sum of all entropies for products) – (Sum of all entropies for reactants)	
	$\Delta S = 354 - 435$	
	$\Delta S = -81 \text{ J K}^{-1} \text{ mol}^{-1}$	
	The reaction was repeated using gaseous methanol, $CH_3OH(g)$, in place of th liquid methanol, $CH_3OH(l)$, used above. What effect, if any, would this have o the value of the entropy change ΔS given above? Explain your answer. [2]	n
(iii)	Use the values in parts (i) and (ii) of this question to calculate the value of the Gibbs free energy, ΔG , for this reaction at 298K and state what information the gives about the feasibility of the reaction.	S

Total [11]

3. Read the passage below and then answer questions (a) to (c) in the spaces provided.

The oxides of nitrogen

The atmosphere around us consists principally of two elements – nitrogen gas, N_2 , and oxygen gas, O_2 . The relative stability of this mixture of two elements hides the fact that the elements can combine to form a number of oxides of nitrogen. Their original names are shown below.

5	Dinitrogen monoxide	N_2O
	Nitrogen monoxide	NO
	Dinitrogen trioxide	N_2O_3
	Nitrogen dioxide	NO_2
	Dinitrogen tetroxide	N_2O_4
10	Dinitrogen pentoxide	N_2O_5
	Nitrogen trioxide	NO_3

Many of these oxides are useful but several can also cause environmental problems.

Dinitrogen monoxide, N2O

This gas was one of the first gaseous compounds to be identified and is probably one of the best known of the oxides of nitrogen. Commonly called 'laughing gas', due to the behaviour of those exposed to the gas, this oxide has since been used as an anaesthetic. It was initially used for the relief of pain during dental treatment and it remained one of the dentist's most useful aids for over a century. It was also commonly used to relieve the pain of childbirth due to the rarity of any adverse reactions to the gas.

Nitrogen dioxide, NO₂

15

Nitrogen dioxide is a brown gas with a notable sharp odour. It can prove toxic by inhalation. The properties of the pure material are difficult to identify due to the existence of the following equilibrium, which leads to the presence of N_2O_4 in any sample of NO_2 .

$$2NO_2(g) \rightleftharpoons N_2O_4(g)$$

Nitrogen dioxide is a key intermediate in the production of nitric acid. The nitrogen dioxide is produced by the oxidation of ammonia and this is then combined with water in a disproportionation reaction.

$$3NO_2(g) + H_2O(1) \longrightarrow 2HNO_3(aq) + NO(g)$$

Nitrogen dioxide, NO₂, along with nitrogen monoxide, NO, is considered to be a key air pollutant and these two oxides are grouped together as NO_x when air quality measurements are undertaken. Both gases are produced during combustion using air as a source of oxygen, such as in the combustion of fuel in vehicle engines. They contribute to the production of atmospheric nitric acid, a key component of acid rain.

Dinitrogen pentoxide, N₂O₅

Dinitrogen pentoxide is a colourless solid at temperatures around 0 °C, however when warmed to 32 °C the oxide sublimes to form $N_2O_5(g)$. In the gas phase the dinitrogen pentoxide is unstable and decomposes, producing nitrogen dioxide.

$$2N_2O_5(g) \longrightarrow 4NO_2(g) + O_2(g)$$

colourless gas brown gas colourless gas

Solutions of dinitrogen pentoxide dissolved in trichloromethane, CHCl₃, have been used as nitration agents to introduce the —NO₂ grouping into organic compounds. The use of this reagent requires a great deal of care as it is a strong oxidising agent and forms explosive mixtures with a range of organic materials.

- End of passage -

(1095-01)

Turn over.

Exa	m	ine	
C	nl	v	

Dinitrogen pentoxide, N ₂ O ₅ , decomposes in the gas phase according to the equation	shown in
line 38.	

(a)	Suggest two methods of studying the kinetics of this reaction.	[2]
	1	

2.

(b) The initial rates of this reaction for different concentrations of N_2O_5 were measured and are given in the table below.

Concentration of N ₂ O ₅ / mol dm ⁻³	Initial rate / mol dm ⁻³ s ⁻¹	
4.00×10^{-3}	3.00×10^{-5}	
6.00 × 10 ⁻³	4.50×10^{-5}	
8.00 × 10 ⁻³	6.00 × 10 ⁻⁵	

The rate equation for this reaction is:

Rate =
$$k[N_2O_5]^1$$

(i)	Show that the rate equation is consistent with the data above.	

(ii)	Calculate the value of the rate constant under these conditions.	Give your answer
	to three significant figures and state its units.	[3]

Units

Examiner only

(iii) Two possible mechanisms have been suggested for this reaction. These are shown below.

rogen dioxide, NO ₂ , produgen tetroxide, N ₂ O ₄ . (line 2	ced in this reaction exists in dynamic equilibrium with (24)
$2NO_2(g) \rightleftharpoons N_2O_4(g)$	$\Delta H = -57.2 \text{ kJ mol}^{-1}$
Vrite an expression for the	equilibrium constant, K_p , for this reaction.
tate and explain the effect	of increasing the temperature on the value of $K_{\rm p}$. [
At a temperature of 373 K, $.00 \times 10^5$ Pa. When the maressure of the remaining N	the partial pressure of a pure sample of NO_2 was ixture was allowed to reach equilibrium, the parti NO_2 was 2.81×10^5 Pa.
Calculate the value of $K_{\rm p}$, so	tating its units.
	gen tetroxide, N_2O_4 . (line 2 $2NO_2(g) \Rightarrow N_2O_4(g)$) Vrite an expression for the tate and explain the effect At a temperature of 373 K, $.00 \times 10^5$ Pa. When the maressure of the remaining N

Total [15]

SECTION B

Answer **both** questions in the separate answer book provided.

- 4. (a) (i) State what is meant by the term transition element. [1](ii) Explain why both iron and copper are classed as transition elements whilst zinc is
 - (ii) Explain why both iron and copper are classed as transition elements, whilst zinc is not. [1]
 - (b) Transition elements such as copper frequently form coloured complexes. Copper(II) complexes are usually blue, but the exact colour can vary, with $[Cu(H_2O)_6]^{2+}$ being pale blue and $[Cu(NH_3)_4(H_2O)_2]^{2+}$ being royal blue. Copper(I) complexes are usually colourless.

Explain why transition metal complexes are usually coloured. Your answer should include details of:

- The origin of colour in transition metal complexes;
- Why the copper(II) species above are coloured blue;
- Why the colours seen in different copper(II) complexes are different;
- Why copper(I) complexes do not form coloured compounds. [6]

(OWC) [2]

- (c) Iron is usually extracted from iron(III) oxide, Fe₂O₃, in a blast furnace using carbon monoxide, CO, as a reducing agent, releasing metallic iron and the gas carbon dioxide.
 - (i) Write the overall equation for this reaction. [1]
 - (ii) Explain in terms of oxidation states why carbon monoxide is considered to be the reducing agent in this reaction. [2]
 - (iii) Explain why carbon monoxide, CO, can be used as a reducing agent but the corresponding oxide of lead, PbO, cannot. [2]

- (d) The iron content of an alloy can be determined by a redox titration using acidified potassium dichromate(VI) solution, $K_2Cr_2O_7$. A piece of alloy of mass 1.870g was dissolved completely in acid to form Fe²⁺ ions, and the solution made up to 250.0 cm³. A 25.00 cm³ sample of this solution was titrated against acidified $K_2Cr_2O_7$. This required 23.80 cm³ of $K_2Cr_2O_7$ solution of concentration 0.0200 mol dm⁻³ for complete reaction.
 - (i) The half-equations for the processes occurring are:

$$Cr_2O_7^{2-} + 14H^+ + 6e^- \longrightarrow 2Cr^{3+} + 7H_2O$$

 $Fe^{3+} + e^- \longrightarrow Fe^{2+}$

(1095-01)

Write an **ionic** equation for the reaction between Fe^{2+} ions and $Cr_2O_7^{2-}$ ions in acid solution. [1]

- (ii) Calculate the number of moles of Fe²⁺ ions present in the 25.00 cm³ sample used in the titration. [2]
- (iii) Calculate the percentage of iron in the original alloy sample. [2]

Total [20]

5. (a) Give a current use for a named compound of chlorine.

- [1]
- (b) Chlorine gas, Cl₂, is used in the industrial preparation of bromine, Br₂. Sea water contains small amounts of bromide ions and by bubbling chlorine gas through the sea water these can be converted to Br₂.
 - (i) Write an ionic equation for the reaction occurring.

[1]

(ii) Use the standard electrode potentials, E^{\bullet} , listed below to explain why chlorine can react with bromide ions but iodine cannot react with bromide ions. [3]

Half-equation	E [⇔] /V
$I_2 + 2e^- \rightleftharpoons 2I^-$	+0.54
$Br_2 + 2e^- \rightleftharpoons 2Br^-$	+1.09
$Cl_2 + 2e^- \rightleftharpoons 2Cl^-$	+1.36

- (c) Sodium chloride and sodium iodide are both compounds which contain halide ions.
 - (i) Silver nitrate solution may be used to differentiate between solutions of sodium chloride and sodium iodide. Give the observations that would be expected in **both** cases.
 - (ii) Both sodium chloride and sodium iodide react with concentrated sulfuric acid. The observations made during both reactions are very different. Discuss the reactions occurring. Your answer should include
 - the observations made during both reactions,
 - the identities of any products,
 - the reasons for any differences in the reactions that occur.

(*QWC*) [1]

- (d) Chlorine produces a range of oxoacids, including chloric(I) acid, HOCl, and chloric(VII) acid, HClO₄. Chloric(I) acid is considered to be a weak acid whilst chloric(VII) acid is considered to be a strong acid.
 - (i) What is meant by the term *strong acid*?

[1]

[5]

- (ii) Write an expression for the acid dissociation constant, K_a , of chloric(I) acid, HOCl.
- (iii) The pH of a solution of chloric(I) acid of concentration 0.100 mol dm⁻³ was found to be 4.23. Calculate the concentration of hydrogen ions in this solution. [2]
- (iv) Using the information from part (iii), calculate the value of the acid dissociation constant, K_a , for chloric(I) acid. [2]
- (v) When the weak acid HOCl reacts with the strong base sodium hydroxide it forms the salt sodium chlorate(I), NaOCl. Suggest a pH value for a solution of NaOCl, giving a reason for your answer. [2]

Total [20]



GCE A level

1095/01-A

CHEMISTRY CH5 PERIODIC TABLE

A.M. FRIDAY, 24 June 2011

THE PERIODIC TABLE

					2			
0	4.00 He Helium		20.2 Ne Neon	40.0 Ar Argon 18	83.8 Kr Krypton 36	Xe Xenon 54	(222) Rn Radon 86	
7			19.0 F	35.5 CI Chlorine	79.9 Bromine	127 I Iodine 53	(210) At Astatine 85	
9		ock	16.0 O Oxygen 8	32.1 S Sulfur 16	Selenium	128 Te Tellurium 52	(210) Po Polonium 84	
w		p Block	14.0 N	31.0 Phosphorus	74.9 As	Sb Antimony 51	209 Bi Bismuth 83	
4			12.0 C Carbon 6	Si Silicon	72.6 Ge Germanium	Sn Tin 50	207 Pb Lead 82	
ю		•	10.8 B Boron 5	Al Aluminium	69.7 Ga Gallium	In Indium 49	204 T1 Thallium 81	
					65.4 Zn Zinc 30	Cd Cadmium 48	Hg Mercury	
					63.5 Cu Copper 29	Ag Silver	197 Au Gold 79	
		ive iic	a ca		58.7 Ni Nickel	106 Pd Palladium 46	195 Pt Platinum 78	
Group	Key	\forall	nass atomic number		58.9 Co Cobalt 27	103 Rh Rhodium 45	192 Ir Iridium 77	
Ö		Ą	Symbol Name	ock	55.8 Fe Iron 26	Ruthenium 44	190 Os Osmium 76	
				d Block	54.9 Mn Manganese 25	98.9 Tc Technetium	186 Re Rhenium	
					52.0 Cr Chromium 24	95.9 98.9 Mo Tc Molybdenum Technetium 42 43	184 W W Tungsten 74	
					50.9 V Vanadium 23	92.9 Nb Niobium 41	181 Ta Tantalum 73	
					47.9 Ti Titanium 22	91.2 Zr Zirconium 40	179 Hf Hafnium 72	
					45.0 Sc Scandium 21	88.9 Y Yttrium 39	139 ► La Lanthanum 57	(227) A Actinium 89
7	ock		9.01 Be Beryllium	24.3 Mg Magnesium	40.1 Ca Calcium	87.6 Sr Strontium 38	137 Ba Barium 56	(226) Ra Radium 88
1	s Block	Hydrogen 1	6.94 Li Lithium 3	23.0 Na Sodium	39.1 K Potassium 19	85.5 Rb Rubidium	133 Cs Caesium 55	(223) Fr Francium 87
	Period	ı	2	3	4	5	9	7

anthanoid	140 Ce	141 Pr	144 Nd	(147) Pm	150 Sm	(153) Eu	157 Gd	159 Tb	163 Dy	165 Ho	167 Er	169 Tm	173 Yb	175 Lu
l	Cerium 58	Praseodymium Neodym	Neodymium 60	Promethium 61	Samarium 62	Europium 63	Gadolinium 64	Terbium 65	Dysprosium 66	Holmium 67	Erbium 68	Thulium 69	Ytterbium 70	Lutetium 71
	232	(231)	238	(237)		(243)	(247)	(245)	(251)	(254)	(253)	(256)	(254)	(257)
Actinoid	Th	Pa	n	dN	Pu	Am	Cm	Bk	Ç	Es	Fm	Мd	No	Ľ
lements	Thorium	Protactinium	Uranium	Neptunium	щ	Americium	Curium	Berkelium	Californium	Einsteinium	Fermium	Mendelevium	Nobelium	Lawrencium
	06	91	92	93	94	95	96	76	86	66	100	101	102	103