Candidate Name	Centre Number	Candidate Number

WELSH JOINT EDUCATION COMMITTEE General Certificate of Education Advanced



CYD-BWYLLGOR ADDYSG CYMRU Tystysgrif Addysg Gyffredinol Uwch

336/01

CHEMISTRY CH6a

A.M. MONDAY, 25 June 2007 (1 hour 10 minutes)

_	EXAMIN JSE ONL	
Section	Question	Mark
A	1	
В	2	
С	3	
	4	
TOTAL		

ADDITIONAL MATERIALS

In addition to this examination paper, you will need a:

- calculator:
- **Data Sheet** which contains a **Periodic Table** supplied by WJEC. Refer to it for any **relative atomic masses** you require.

INSTRUCTIONS TO CANDIDATES

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

- **Section A** Answer the question in the spaces provided.
- **Section B** Answer the question in the spaces provided.
- **Section C** Answer **both** questions in the spaces provided.

Candidates are advised to allocate their time appropriately between **Section A** (10 marks), **Section B** (15 marks) and **Section C** (25 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 50.

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 used in all written answers.

Page 11 may be used for rough work.

No certificate will be awarded to a candidate detected in any unfair practice during the examination.

SECTION A

Answer the questions in the spaces provided.

1.	• This question concerns the following six solids.					
	Aminoethanoic acid (glycine)			NH ₂ CH ₂ COOH		
	Amn	noniu	m ethanoate	CH ₃ COONH ₄		
	Ethanamide			CH ₃ CONH ₂		
	Lead	(II) ic	odide	PbI_2		
	Sodi	um io	dide	NaI		
	Triio	dome	thane (iodoform)	CHI_3		
	(a)	State the two solids which are yellow in colour. [1]				
				and		
	(b)	State	e the two solids which do	o not have ionic bonding in their structures.	[1]	
				and		
	(c)	Give the structure of the organic group which can be identified on warming with alkaline iodine solution due to the formation of triiodomethane. [1]				
	(d)	(i)	Outline how a pure sa iodide.	ample of lead(II) iodide can be prepared using	sodium [2]	
		(ii)	Outline how ammoniu	m ethanoate can be converted to ethanamide.	[1]	

Section A Total [10]

(e)	Choose two solids from the list which will react with sodium hydroxide solute. For each of the compounds you choose, give the observations and the product(sthe reaction.	
	First solid	
	Observations	
	Product(s)	[2]
	Second solid	
	Observations	
	Product(s)	
		[2]

(336-01) **Turn over.**

SECTION B

2. Read the passage below and then answer the questions in the spaces provided.

Hydrogen Peroxide

Hydrogen peroxide, H₂O₂, is a widespread compound. It can be detected in drinking water, rain water, sea water and polluted air such as smog. Discovered by Louis Jacques Thénard in 1818, it is a colourless liquid that resembles water, with similar physical properties.

5 Compared to the more stable water molecule, an H₂O₂ molecule contains one extra oxygen atom, with single covalent bonds H-O-O-H. The bond between the two oxygen atoms is the so-called "peroxide" bond. There are many uses for hydrogen peroxide, which rely on two reactions:

$$2H_2O_2 \longrightarrow 2H_2O + O_2$$
 Reaction **A**

10 $H_2O_2 \longrightarrow 2HO^{\bullet}$ Reaction **B**

Hydrogen peroxide is a strong oxidising agent, more powerful than chlorine, Cl₂, or potassium manganate(VII), KMnO₄. The oxidising behaviour means it is an irritant of eyes, mucous membranes, and skin. One advantage of hydrogen peroxide is that the decomposition products, water and oxygen, are not harmful.

15 Hydrogen peroxide has been used for bleaching straw hats, pulp and paper, but perhaps its best known use is the bleaching of hair. H₂O₂ can oxidise double bonds in large organic molecules. Since the double bonds often cause the molecules to absorb light, and therefore give the molecules their colour, removal of them destroys the pigments and so removes the colour. The term "peroxide blondes" was made famous by movie stars such as Marilyn Monroe and Jean Harlow.

Another widespread application of hydrogen peroxide is as a mild disinfectant for treating cuts and scratches. What makes hydrogen peroxide unique is the foaming action that occurs when it is placed on a cut. People used to believe this indicated the presence of infection, and the H₂O₂ was foaming as it destroyed the disease. In reality, though Reaction A normally occurs very slowly at room temperature, hydrogen peroxide foams when it comes into contact with blood because enzymes in blood, such as catalases, catalyse Reaction A. H₂O₂ is present in trace amounts in honey, which, before the advent of modern preparations, was used for dressing wounds.

Hydrogen peroxide is commercially available in various grades based on the percentage of H₂O₂. For example, Pharmaceutical Grade (3%) contains 3.00 g H₂O₂ made up to 100 cm³ solution. Beautician Grade, used for hair colourings, is typically 6% strength whilst 35% Food Grade is used in producing cheese and egg-containing foods, and is sprayed onto the foil lining of drink cartons.

<i>(a)</i>	Draw a dot and cross	diagram to sho	ow the covalent	bonding in	hydrogen	peroxide
	H_2O_2 (line 6).					[1]

(b) (i) Calculate the enthalpy change, ΔH^{\oplus} , for Reaction A:

$$2H_2O_2(l) \longrightarrow 2H_2O(l) + O_2(g)$$
 [2]

Use the enthalpy changes of formation below.

$$\Delta H_{\rm f}^{-\Theta} H_2 O_2(1) = -187.8 \text{ kJ mol}^{-1}$$

$$\Delta H_{\rm f}^{\ominus}$$
 H₂O(l) = -285·8 kJ mol⁻¹

$$\Delta H_{\rm f} \stackrel{\Leftrightarrow}{\circ} {\rm O}_2({\rm g}) = 0 \quad {\rm kJ \ mol}^{-1}$$

(ii) Explain why hydrogen peroxide foams when it comes into contact with blood (*lines 9, 25 to 27*). [1]

(c) (i) Reaction A can also be carried out in the gas phase

$$2H_2O_2(g)$$
 \longrightarrow $2H_2O(g)$ + $O_2(g)$ $\Delta H^{\oplus} = -210 \text{ kJ mol}^{-1}$

Given the bond energies

$$O - H ext{ } 464 \text{ kJ mol}^{-1} ext{ } and ext{ } O = O ext{ } 498 \text{ kJ mol}^{-1}$$

use the
$$\Delta H^{\odot}$$
 value to calculate the O – O bond energy. [2]

(ii) Explain why the O – O bond is normally the one involved when H_2O_2 reacts (line 10). [1]

(d) The following table shows some standard electrode potentials.

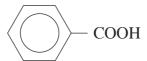
Half Reaction	Electrode Potential / V
$I_2 + 2e^- \Longrightarrow 2I^-$	+ 0.54
$Cl_2 + 2e^- \Longrightarrow 2Cl^-$	+ 1.36
$MnO_4^- + 8H^+ + 5e^- \iff Mn^{2+} + 4H_2O$	+ 1.70
$H_2O_2 + 2H^+ + 2e^- \Longrightarrow 2H_2O$	+ 1.77

(i)	Using the standard electrode potentials, explain why H_2O_2 is the strongest oxidising agent of those shown in the table (<i>lines 11, 12</i>). [2]
(ii)	Calculate the EMF for the oxidation of iodide ions, I^- , by hydrogen peroxide, H_2O_2 , in acid solution and write a balanced equation for the reaction. [2] <i>EMF</i>
	Equation
(i)	State the general name given to a group of atoms which cause a molecule to be coloured (<i>lines 17, 18</i>). [1]
(ii)	Explain why hydrogen peroxide bleaches many organic materials (lines 16 – 19). [1]
	culate the concentration of Pharmaceutical Grade (3%) H_2O_2 in mol dm ⁻³ , giving r answer to three significant figures (lines 30, 31). [2]
	Section B Total [15]

SECTION C

Answer both questions in the spaces provided.

3. The white crystalline solid benzenecarboxylic acid (benzoic acid) and its salts are widely used in industry as antimicrobial agents in products such as food, inks and hair spray. It has the following structure.



benzenecarboxylic acid

(a)	(i)	Benzenecarboxylic acid can be prepared from a hydrocarbon, compound A . Compound A contains 91·23% carbon and 8·77% hydrogen by mass, with a relative molecular mass of 92·08. Its proton NMR spectrum shows two main peaks at shifts of 2·2 ppm and 7·3 ppm with peak areas in the ratio 3:5. Showing your reasoning, determine the structure of compound A and state the reagent(s) and conditions needed to convert it to benzenecarboxylic acid. [4]
	(ii)	Given that benzenecarboxylic acid is soluble in hot water but almost insoluble in cold water, describe an experimental technique you could use to purify the benzenecarboxylic acid produced. [3]

(336-01) **Turn over.**

(b)	State the organic product, and name the type of reaction occurring, when benzenecarboxylic acid is				
	(i)	heated with sodalime,	[2]		
		Organic product			
		Type of reaction			
	(ii)	treated with lithium tetrahydridoaluminate(III), LiAlH ₄ , in dry ether.	[2]		
		Organic product			
		Type of reaction			
(c)	Benz	zenecarboxylic acid is a weak acid ($K_a = 6.3 \times 10^{-5} \text{ mol dm}^{-3}$).			
		\leftarrow COOH \rightleftharpoons H ⁺ + \leftarrow COO ⁻			
	A so Calc	plution of benzenecarboxylic acid has a concentration of 2.79×10^{-2} mol dm ⁻³ . culate the pH of this solution of benzenecarboxylic acid.	[2]		

4.	(a)	of th	minium chloride, AlCl ₃ , is an <i>electron-deficient molecule</i> . Explain the meannis term and why it applies to AlCl ₃ .	[2]
	(b)	AlC (i)	Cl ₃ , under appropriate conditions, will react with a chloride ion to form AlCl ₄ . Describe the bonding change(s) which occur during this reaction.	. [1]
		(ii)	State the shapes of the $AlCl_3$ molecule and the $AlCl_4^-$ ion. $AlCl_3$ shape	[2]
	(c)	deve poly temp	Cl ₄ is one anion found in <i>ionic liquids</i> , which in recent years have beloped as catalysts and solvents for a range of processes such as alk ymerisation. <i>Ionic liquids</i> are organic salts that generally have mely peratures below 100 °C and are liquid over a wide temperature range. React thus be performed under mild conditions.	ting
		(i)	State two advantages of being able to carry out industrial reactions under a conditions only just above room temperature.	mild [2]
		(ii)	State the reaction type involved in the polymerisation of alkenes such ethene.	h as
		(iii)	State one organic preparation, other than polymerisation, in which alumin chloride is used as a catalyst.	iium [1]
		•••••		

	d) A	أسينا	ninm	ablarida	can exist	00.0	dimor
1	u	<i>)</i> Λ	lullill	IIIuIII	Cilionae	call exist	as a	unner

$$2AlCl_3(g) \iff Al_2Cl_6(g)$$

Giving your reasons in each case, state whether

	Total	[12]
(iii) 	dimerisation in aluminium chloride would be favoured by high or pressure.	low [1]
(ii)	the proportion of dimer present at equilibrium is greater at high or temperatures,	low [1]
(i) 	dimerisation, the forward reaction above, will be exothermic or endotherm	[1]

Section C Total [25]

Rough Work