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Pearson Edexcel International Advanced Level	Centre Number	Candidate Number
Chemistry Advanced Subsidiar Unit 3: Chemistry Lal	ry	
Thursday 21 January 2016 Time: 1 hour 15 minutes	– Morning	Paper Reference WCH03/01
Candidates may use a calcula	tor.	Total Marks

Instructions

- Use **black** ink or ball-point pen.
- **Fill in the boxes** at the top of this page with your name, centre number and candidate number.
- Answer **all** questions.
- Answer the questions in the spaces provided
 - there may be more space than you need.

Information

- The total mark for this paper is 50.
- The marks for each question are shown in brackets
 use this as a guide as to how much time to spend on each question.
- You will be assessed on your ability to organise and present information, ideas, descriptions and arguments clearly and logically, including your use of grammar, punctuation and spelling.
- A Periodic Table is printed on the back cover of this paper.

Advice

- Read each question carefully before you start to answer it.
- Keep an eye on the time.
- Try to answer every question.
- Check your answers if you have time at the end.

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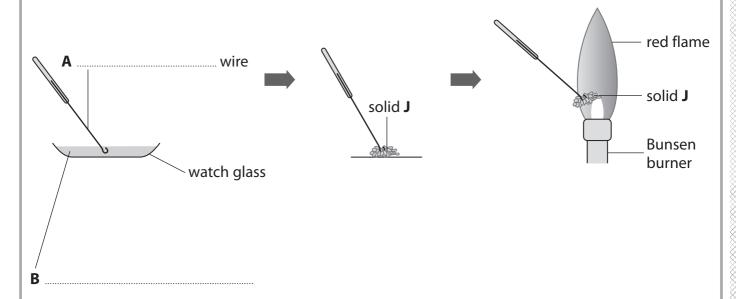
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Answer ALL the questions. Write your answers in the spaces provided.

- 1 A series of tests is carried out on a white solid, **J**, which is a mixture of two compounds. One compound contains a Group 1 cation, and the other a Group 2 cation. The two compounds contain the same anion.
 - (a) Complete the labels **A** and **B** in the diagram below, which shows the procedure and result for a flame test on solid **J**.





(b) It is known that the Group 2 cation in **J** gives no colour in a flame test, so the red colour seen must be due to the Group 1 cation.

Give the name or formula of the Group 1 cation, which is responsible for the red colour observed in the flame test, and give the name or formula of the Group 2 cation.

(2)

Group 1 ca	tion	 	 	 	

Group 2 cation





(c)	Dilute hydrochloric acid is added to a sample of J. J dissolves in the acid but there is no other change.	
	(i) If J contained a carbonate anion, what would be the observation on the addition of dilute hydrochloric acid?	(1)
	(ii) Give the name or formula of another anion which would produce the same observation with dilute hydrochloric acid as the carbonate anion.	(1)
	(iii) Aqueous barium chloride is then added to the solution of J in hydrochloric ac A white precipitate forms. Give the formulae for the two salts present in J .	id. (2)
	(Total for Question 1 = 8 m	arks)



2 This question is about the following two dicarboxylic acids. Both acids are solid at room temperature.

(a) (i) Phosphorus(V) chloride is often used to confirm the presence of an —OH group in a compound.

Suggest a practical problem if solid phosphorus(V) chloride is used with these two dicarboxylic acids.

(1)

(ii) Suggest a reagent that could be used to confirm the presence of an **acid** group in either of the two compounds above, and the positive observation that would be made.

(2)

Reagent

Observation

(b) Bromine dissolved in an organic solvent reacts rapidly with butenedioic acid.

Complete the equation for the reaction of butenedioic acid with bromine.

(c) Propanedioic acid can be produced by the oxidation of propane-1,3-diol.	
(i) Draw the skeletal formula of propane-1,3-diol.	(1)
(ii) Identify one way in which the infrared spectrum of propanedioic acid would be different from that of the infrared spectrum of propane-1,3-diol.	
Wavenumber data are not required.	(1)
(Total for Question 2 = 6 m	arks)



3 Washing soda is hydrated sodium carbonate, Na₂CO₃.xH₂O, where the number of moles of water of crystallization, x, can vary.

A sample of washing soda is analysed. Two methods are used to determine the value of x in the sample.

Method 1: Heating

2.50 g of the washing soda is placed in a crucible. The crucible is gently heated for three minutes and then heated strongly for five minutes. The mass of the solid after heating is 1.06 g.

$$Na_2CO_3.xH_2O(s) \rightarrow Na_2CO_3(s) + xH_2O(g)$$
 Equation 1

(a) Suggest why the crucible is heated gently for the first three minutes.

(1)

(b) What additional step after heating strongly for five minutes is needed to make sure that all of the water of crystallization has been removed?

(1)

(c) What is the correct chemical term for sodium carbonate without water of crystallization?

(1)

(d) (i) Calculate the number of moles of sodium carbonate that remain after heating the sample, assuming that all of the water of crystallization has been removed.

(2)

(ii) Calculate the number of moles of water lost from the sample of washing soda on heating.

(iii) Hence deduce the value of x in the sample of washing soda, $Na_2CO_3.xH_2O$, obtained using Method 1.

(1)

Y =

Method 2: Titration

Sodium carbonate reacts with hydrochloric acid as follows:

$$Na_2CO_3 + 2HCI \rightarrow 2NaCI + CO_2 + H_2O$$
 Equation 2

A 2.50 g sample of the washing soda is placed in a beaker and dissolved in deionized water. This solution is poured into a 250 cm³ volumetric flask, made up to the mark and mixed thoroughly.

A pipette is then used to transfer 25.00 cm³ of the washing soda solution to each of three conical flasks. A burette is filled with hydrochloric acid, of concentration 0.100 mol dm⁻³, and titrations are carried out. The results are shown in the table.

Titration numbers	1	2	3
Burette reading (final) / cm ³	17.00	33.55	16.45
Burette reading (initial) / cm ³	0.00	17.00	0.00
Titre / cm ³	17.00	16.55	16.45

(e)	What should be done to make sure that all of the washing soda is transferred to
	the volumetric flask?

(1)

(f) Explain why only titrations 2 and 3 are used to calculate the mean titre.





(g) (i) Calculate the mean titre, and then calculate the number of moles of hydrochloric acid in the mean titre.

(1)

(ii) Using your answer to part (g)(i) and **Equation 2**, calculate the number of moles of sodium carbonate present in 25 cm³ of the washing soda solution.

(1)

(iii) Hence calculate the total number of moles of sodium carbonate present in 250 cm³ of the washing soda solution.

(1)

(iv) Calculate the molar mass of the hydrated washing soda, Na₂CO₃.xH₂O. Hence deduce the value of x in the sample of washing soda from the data in Method 2.

(2)

x =

(h) A student carrying out Method 2 overshot the end-point of each titration.	
Explain how this would affect the calculated value of x.	(2)
(Total for Question 3 = 16 ma	rks)

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4 The thermal decomposition of copper(II) carbonate is

$$CuCO_3(s) \rightarrow CuO(s) + CO_2(g)$$

Equation 3

The enthalpy change for this reaction, ΔH_3 , cannot be determined directly. However, it can be calculated using Hess's law, from the enthalpy changes for the reaction of sulfuric acid with copper(II) carbonate and with copper(II) oxide.

(a) A student carried out an experiment to determine the enthalpy change, ΔH_4 , for the reaction

$$CuCO_3(s) + H_2SO_4(aq) \rightarrow CuSO_4(aq) + H_2O(l) + CO_2(g)$$
 Equation 4

In the experiment, a known mass of copper(II) carbonate was mixed with a known volume of sulfuric acid in a polystyrene cup, and the temperature change measured. The results of the experiment are shown in the table.

Measurement	Reading
Mass of copper(II) carbonate added to sulfuric acid	2.54 g
Volume of sulfuric acid, 1 mol dm ⁻³	50.0 cm ³
Initial temperature of sulfuric acid before addition of copper(II) carbonate	24.3°C
Maximum temperature of sulfuric acid after the addition of copper(II) carbonate	29.0℃

(i) Calculate the energy transferred, in joules, for this reaction using the expression Energy transferred (J) = $50.0 \times 4.18 \times \text{temperature change}$

(1)

(ii) Calculate the number of moles of copper(II) carbonate used.

Molar mass of copper(II) carbonate = 123.5 g mol^{-1}



(iii) Use your answers to (a)(i) and (a)(ii) to calculate, in kJ mol ⁻¹ , the enthalpy change, ΔH_4 , for the reaction shown in Equation 4 . Include a sign for ΔH_4 and give your answer to three significant figures.	(2)
(iv) Why does the sulfuric acid need to be in excess?	(1)

(v) The enthalpy change obtained from this experiment is much less negative than the Data Booklet value.

Suggest one likely reason for this difference, other than a measurement error.

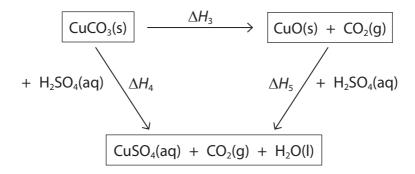


(b) The student then carried out a similar experiment to determine the enthalpy change, ΔH_5 , for the reaction between copper(II) oxide and sulfuric acid.

$$CuO(s) \hspace{0.3cm} + \hspace{0.3cm} H_2SO_4(aq) \hspace{0.3cm} \rightarrow \hspace{0.3cm} CuSO_4(aq) \hspace{0.3cm} + \hspace{0.3cm} H_2O(I) \hspace{0.3cm} \textbf{Equation 5}$$

From the results of this experiment, ΔH_5 was calculated to be -56.1 kJ mol⁻¹.

The values of ΔH_4 and ΔH_5 can be used to determine the enthalpy change for the thermal decomposition of copper(II) carbonate using the Hess cycle shown below.



Use Hess's law to calculate the value of ΔH_3 , in kJ mol⁻¹. Include a sign in your answer. (2)

(c) Suggest why it is not possible to determine directly the enthalpy change for the thermal decomposition of copper(II) carbonate.

(1)

(Total for Question 4 = 9 marks)

5 A method for the preparation of iodoethane is given in outline below.

Procedure

- **Step 1** Suitable quantities of red phosphorus and iodine are placed in a round-bottom flask. The flask is fitted with a reflux condenser and immersed in cold water.
- **Step 2** Using a dropping pipette, a suitable volume of ethanol is added, in 1 cm³ portions, down the condenser.
- **Step 3** After the addition of the ethanol is complete, and a further 15 minutes have passed, the cold water bath is removed and the mixture in the flask is heated under reflux for 45 minutes.
- **Step 4** The apparatus is allowed to cool and the condenser is rearranged for distillation.
- **Step 5** The crude iodoethane is distilled off.
- **Step 6** The distillate is washed with dilute sodium carbonate solution.
- **Step 7** The washed iodoethane is separated from the aqueous solution.
- **Step 8** Anhydrous calcium chloride is added to the washed iodoethane.

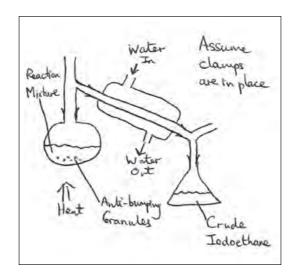
The equations for the reactions are

$$2P + 3I_2 \rightarrow 2PI_3$$

 $3CH_3CH_2OH + PI_3 \rightarrow 3CH_3CH_2I + H_3PO_3$

(a) What does the way in which ethanol is added in **step 2** suggest about the nature of the reaction?

(b) A student drew a diagram of the apparatus used in **step 5**. There are a number of errors in the diagram.



(i) One of the errors is that the flow of water in the condenser is the wrong way round. Explain the effect of this error.

(1)

(ii) Identify the most significant error in the diagram and explain the effect of this.

(1)

Effect

(c) Suggest why, in step 6 , the crude iodoethane is washed with dilute sodium carbon solution.					
Solution.	(1)				
(d) Draw a diagram of the apparatus that would be used in step 7 to separate the iodoethane. Name the apparatus and label its contents.					
The density of iodoethane is 1.5 g cm $^{-3}$ and the density of the aqueous solution is about 1.0 g cm $^{-3}$.					
Name of apparatus	(3)				
Diagram					
(e) How will the appearance of the iodoethane be changed by the addition of anhydrous calcium chloride in step 8 ?					
, , , , , , , , , , , , , , , , , , ,	(1)				
(f) How would the iodoethane be separated from the calcium chloride after step 8 ?	(1)				



(g) To obtain pure iodoethane, one further step in the preparation is needed. What is this step?

 (1)

 (h) It is not possible to effectively produce iodoethane by reacting ethanol with a mixture of sodium iodide and 50% sulfuric acid. This is because the sulfur in the sulfuric acid can be reduced to form substances such as hydrogen sulfide and sulfur.
 State what happens to the iodide ions in the sodium iodide when this occurs.

(Total for Question 5 = 11 marks)

TOTAL FOR PAPER = 50 MARKS

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[257] ᆣ 103

[254] 2 102

[256] PW 101

Cm curium 9%

uranium 238 U

[231]
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protactinium

92

9

90

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Table
2
<u>10</u>
Periodic
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_	, E	2	0 5	80 . Vo	m e	73		
0 (8)	4.0 He helium 2	20.2 Ne neon	39.9 Ar argon 18	83.8 Kr krypton 36	Xe xenon 54	[222] Rn radon 86	ted	
1	(77)	19.0 F fluorine	35.5 CI chlorine 17	79.9 Br bromine 35	126.9 	[210] At astatine 85	sen repor	175 Lu lutetium
0	(16)	16.0 O oxygen 8	32.1 S sulfur 16	79.0 Se selenium	127.6 Te tellurium 52	Po Polanium 84	Elements with atomic numbers 112-116 have been reported but not fully authenticated	173 Yb ytterbium
n	(15)	14.0 Nitrogen	31.0 P phosphorus 15	As As arsenic 33	121.8 Sb antimony 51	Bi bismuth 83	tomic numbers 112-116 hav but not fully authenticated	169 Tm thu(ium
4	(14)	12.0 C carbon 6	Si Silicon 14	72.6 Ge germanium	118.7 Sn tin 50	207.2 Pb tead 82	atomic nun but not fu	167 Er erbíum
7	(13)	10.8 B boron 5	27.0 Al aluminium 13	Ga gallium 31	114.8 Indium 49	204.4 Tl thallium 81	ents with	165 Ho holmium
			(21)	65.4 Zn zinc 30	Cd cadmium 48	200.6 Hg mercury 80	Elem	163 Dy dysprosium
			(11)	63.5 Cu copper	107.9 Ag silver 47	197.0 Au gold 79	Rg reentgenium 111	159 Tb terbium
			(01)	58.7 Nicket	Pd Pd pattadium 46	195.1 Pt platinum 78	Ds darmstadtlum n 110	157 Gd gadolinium
			6)	S8.9 Co cobalt	6 E	192.2 Ir rridium 77	[268] Mt meitnerium 109	152 Eu europium
	1.0 H hydrogen		(8)	55.8 Fe iron 26	Ru ruthenium 44	190.2 Os osmium 76	[277] Hs hassium 108	
			0	Mn Manganese		Re rhenium 75	[264] Bh bohrium 107	141 144 [147] 150 15
		mass sol umber	(9)	52.0 54.9 Cr Mn chromium manganese 24 25	95.9 [98] Mo Tc molybdenum technetium 42 43	183.8 W tungsten 74	[266] Sg seaborgium 106	Nd Nd neodymium
	Key	relative atomic mass atomic symbol name atomic (proton) number	(5)	50.9 V vanadium	E	180.9 Ta tantalum 73	[262] Db dubnium 105	Pr Pr raseotymium
		relati atoric	(4)	47.9 Ti titanium	91.2 Zr zirconium 40	178.5 Hf hafnium 72	[261] Rf rutherfordium 104	Cerium
			(3)	Sc scandium	88.9 Y yttrium 39	138.9 La* tanthanum 57	[227] Ac* actinium 89	85
7	(2)	9.0 Be beryllium 4	Mg magnesium 12	Calcium	87.6 Sr strontium 38	137.3 Ba barium 56	[226] Ra radium 88	*Lanthanide series *Actinide series
-	(1)	6.9 Li lithium	Na sodium 11	39.1 K potassium	85.5 Rb rubidium 37	132.9 Cs caesium 55	[223] Fr francium 87	*Lanth

58 232 **Th** thorium * Actinide series