Write your name here		
Surname	Other	rnames
Pearson Edexcel International Advanced Level	Centre Number	Candidate Number
Chemistry	/	
Advanced Unit 6: Chemistry Lal		II
Friday 15 May 2015 – Morn Time: 1 hour 15 minutes	ing	Paper Reference WCH06/01
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.

Turn over ▶



(1)

Answer ALL the questions. Write your answers in the spaces provided.	
A white solid, $\bf A$, contains one cation and one anion. When water is added slowly, the solid turns blue and then dissolves to form a blue solution, $\bf B$.	
(a) When aqueous barium chloride is added to an acidified portion of solution B , a white precipitate forms.	
(i) Give the formula of the anion in B .	(1)

When aqueous ammonia is added to another portion of solution B , a blue precipitate forms. When more aqueous ammonia is added, this precipitate
dissolves to form a deep blue solution, C .

(ii) Name a suitable acid for acidifying solution **B** in this test.

(i) Identify, by name or formula, the blue precipitate. (1)

(ii) Give the **formula** of the ion responsible for the deep blue colour of solution **C**. (1)

(c) Give the **formula** of the complex ion which gives the blue colour to solution **B**. Include the ligands in your answer.

(1)

(d) Give the **formula** of the white solid **A**. (1)

(e) Why is solid **A** white and not coloured blue? Justify your answer. (2)

(Total for Question 1 = 8 marks)

(b)

2	 A salt, P, contains carbon, hydrogen, oxygen and one metallic element. (a) When a flame test is carried out on P, a yellow flame results. Give the formula of the metal ion in the salt. 	
		(1)
	 (b) P dissolves in water to form a weakly alkaline solution, Q, with pH 8.1. The pH of Q can be measured by using a calibrated pH meter or an indicator. (i) Describe how to calibrate a pH meter. 	
	(i) Describe now to camprate a primeter.	(2)
	(ii) Name a suitable indicator you could use and state the colour you would expect to observe.	
		(2)
	(iii) Which of the two methods will give the more accurate value for the pH of Q ? Justify your answer.	(1)



(c)	Some of the solution Q is acidified with concentrated hydrochloric acid.	
	An organic compound, R , forms in the solution.	
	Methanol is added and the mixture warmed, forming a new organic compound S .	
	This mixture is added to sodium carbonate solution in an evaporating basin. A fruity smell is detected.	
	(i) Describe and explain what you would see as the mixture is added to the	
	sodium carbonate solution.	(2)
	(ii) What type of compound is S ?	
		(1)



	(d) The high resolution proton nuclear magnetic resonance (nmr) spectrum of S has only two peaks which are both singlets and have the same area.	
	Deduce the structural formulae of S , R , and P .	
		(3)
S		
R		
Р		
-		
	(Total for Question 2 = 12 mar	ks)
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3	Cupron	ickel is an alloy of copper and nickel. It is used to make 'silver' coins.	
	A coin i	s analysed by the following method.	
	Step 1	It is weighed on a balance which reads to two decimal places and found to have mass 4.00 g.	
	Step 2	Water is added to the coin in a beaker. Concentrated nitric and sulfuric acids are added and the coin dissolves.	
	Step 3	When the coin is completely dissolved, the solution is neutralized.	
	Step 4	The neutral solution is transferred, with the washings, to a 100 cm ³ volumetric flask, made up to the mark with water and mixed thoroughly.	
	Step 5	10 cm ³ samples of the solution are taken and an excess of potassium iodide is added, producing iodine.	
	Step 6	The iodine is titrated with 0.200 mol dm ⁻³ sodium thiosulfate solution.	
	(a) Why	, in Step 2 , is water added before, rather than after, the acids?	(1)
	(b) Wha	at is the colour of an aqueous iodine solution?	(1)
		To make the end point of the titration more obvious, an indicator is added just before the colour of the iodine disappears.	
		Name this indicator.	
			(1)
		Suggest why the indicator is not added to the iodine solution earlier in the titration.	
			(1)
		Give the colour change at the end point when the indicator is used in this titration.	
			(1)



(d) The results for the titrations are shown below.

Titration number	1	2	3	4
Burette reading (final) / cm³	24.10	47.90	23.55	47.00
Burette reading (initial) / cm³	0.00	24.10	0.00	23.55
Titre / cm³				

	(i)	Compl	ete	the	table
--	-----	-------	-----	-----	-------

(1)

(11)	Which titres should be used to calculate the mear	n? Explain your choice.	
			(1

(iii)	Calculate	the	mean	titre
١	(III <i>)</i>	Calculate	uic	mean	uuc.

(1)



(iv) Calculate the percentage by mass of copper in the coin.

Use the equations below.

$$2Cu^{2+}(aq) \ + \ 4I^{-}(aq) \ \rightarrow \ 2Cul(s) \ + \ I_{2}(aq)$$

$$2S_2O_3^{2-}(aq) + I_2(aq) \rightarrow S_4O_6^{2-}(aq) + 2I^{-}(aq)$$

(5)

(v) The uncertainty in each burette reading is ± 0.05 cm³ and the uncertainty in each reading of the balance is ± 0.005 g.

Calculate the percentage uncertainty in the third titre value and in the mass measurement. Use your results to decide whether using a balance that weighs to three decimal places would significantly improve the accuracy of the result.

(2)

(Total for Question 3 = 15 marks)



- **4** Bromobenzene can be prepared from benzene by the following steps.
 - **Step 1** Reflux 20.0 cm³ of benzene with 6.0 cm³ of bromine and about 10 g of iron filings, by heating on a water bath at 50 °C.
 - **Step 2** After the reaction has finished, remove the water bath and heat to boiling until no bromine vapour can be seen.
 - **Step 3** Cool the mixture and add 25 cm³ of ethoxyethane (diethyl ether) to extract the bromobenzene.
 - **Step 4** Wash the ethoxyethane layer with aqueous sodium hydroxide. Separate the ethoxyethane layer.
 - **Step 5** Wash the ethoxyethane layer with water and repeat the separation.
 - **Step 6** Dry the ethoxyethane layer with a suitable drying agent.
 - **Step 7** Decant the dried solution.
 - **Step 8** Distil the separated solution, collecting the fraction boiling around the boiling temperature of bromobenzene, 156°C.
 - (a) Calculate the number of moles of bromine, Br₂, used in the experiment.

[Density of bromine 3.1 g cm⁻³]

(2)

(b) Bromine reacts with iron to form iron(III) bromide, which reacts with bromine to produce the attacking electrophile in **Step 1**.

Write the chemical equations for these reactions.

(2)



(c)	Why is the ethoxyethane layer washed with sodium hydroxide solution in Step 4 ?	(1)
(d)	Draw a diagram of the apparatus used to separate the ethoxyethane layer from the aqueous layer in Step 5 . Clearly label the ethoxyethane layer.	
	[Densities: water 1.0 g cm ⁻³ , ethoxyethane 0.7 g cm ⁻³]	(2)
(e)	The bromobenzene formed in this reaction can be nitrated to make 2,4-dinitrobromobenzene.	
(e)		(1)
(e)	2,4-dinitrobromobenzene.	(1)



(f) 2,4-dinitrobromobenzene reacts with hydrazine hydrate to make 2,4-dinitrophenylhydrazine crystals.

The percentage yields for the reactions are:

75% for the formation of bromobenzene from benzene

70% for the formation of 2,4-dinitrobromobenzene from bromobenzene

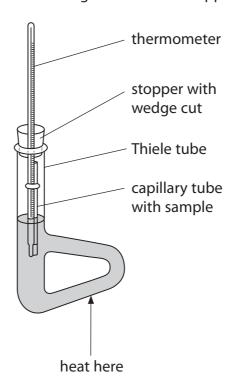
70% for the formation of 2,4-dinitrophenylhydrazine from 2,4-dinitrobromobenzene

Calculate the overall percentage yield of 2,4-dinitrophenylhydrazine from benzene, for this series of reactions.

(1)



(g) The purity of the 2,4-dinitrophenylhydrazine crystals can be checked by carrying out a melting temperature determination using the Thiele tube apparatus shown below.



(i)	The capillary tube must be sealed at one end
	Describe how this is done.

(1)

(ii) When crystals are placed in the capillary tube they often stick in the top. Describe how to ensure the crystals reach the bottom of the capillary tube.

(1)

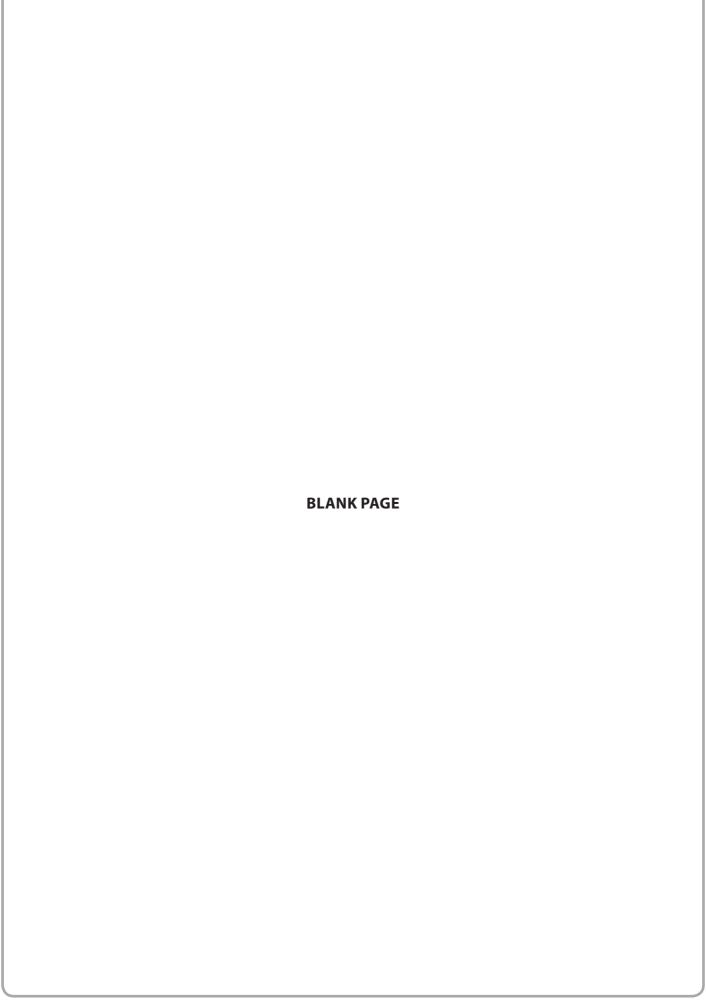
(iii) Dibutyl phthalate is often used as the liquid in the Thiele tube.

Suggest **two** properties of dibutyl phthalate that make it a suitable liquid for this purpose.

(2)



TOTAL FOR PAPER = 50 MA	DIVE					
(Total for Question 4 = 15 ma	marks)					
After recrystallization						
Before recrystallization						
	(2)					
Suggest the temperature range over which you would expect the crystals to melt before and after purification by recrystallization.						
(iv) The melting temperature for crystals of 2,4-dinitrophenylhydrazine is 201°C.						





2	(1) (2)	6,9 9.0 Li Be Utthlum beryllium 3 4	23.0 24.3 Mg sodium magnesium (3) (1)	39.1 40.1 45.0 4	um calcium scandium	87.6 88.9	Rb Sr Y zironbidium strontium yttrium ziron 37 38 39	137.3 138.9	Cs Ba La* P caesium barium tanthanum haf 55 57	[223] [226] [227] [2 Fr Ra Ac* I	nm radium actinium 88 89		* Actinide series
		aton atonic ((4)	47.9	E	91.2	Zr zirconium 40	178.5	Hf hafinium 72	[261] Rf	5		cerium pr
	Key	relative atomic mass atomic symbol name atomic (proton) number	(5)	6,05	En .	92.9	Mobium #	180.9	Ta tantalum 73	[262] Ob	dubnium seaborgium 105 106	141 Pr	praseodymium ne 59
		nass iol	(9)	52.0	Ę	6.56	Mo molybdenum 42	183.8	W tungsten 74	[266]	eaborgium 106	441 Nd	60
			(7)	54.9 Mr	manganese 25	[86]	Tc technetium 43	186.2	Re rhenium 75	[264] Bh	bohrium 107	[147] Pm	odymium promethium 60 61
	1.0 X hydrogen		(8)	55.8	iron 26	101.1	Ru ruthenium 44	190.2	OS osmium 76	[277] Hs	E	150 Sm	samarrum 62
			(6)	58.9	cobalt 27	102.9	Rh rhodium 45	192.2	iridium 77	[268] Mt	meitnerium 109	152 Eu	europium 63
			(10)	58.7	nicket 28	106.4	Pd palladium 46	195.1	Pt platinum 78	[1771] Ds	5		gadolinium 64
			(11)	63.5	copper 29	107.9	Ag silver 47	197.0	Au gold 79	[272] Rø	roentgenium 111		terbium 65
			(12)	65.4	Zinc 30	112.4	Cd cadmium 48	9.002	Hg mercuny 80			163 Dy	dysprosium 66
m	(13)	10.8 B boron 5	27.0 AI atuminium 13	69.7	gallium 31	114.8	Indium 49	2	Th thallium 81	Elements with atomic numbers 112-116 have been reported			holmium 67
4	(14)	12.0 C carbon 6	Si Sificon 14	72.6	germanium 32	118.7	S # 8	207.2	Pb lead 82	atomic nu	but not f	167 Er	erbium 68
Ŋ	(15)	14.0 N nitrogen 7	31.0 P phosphorus 15	74.9	arsenic 33	121.8	Sb antimony 51	209.0	Bi bismuth 83	mbers 112	but not fully authenticated	169 Tm	thulium 69
9	(16)	16.0 O oxygen 8	32.1 S sulfur 16	79.0	selenium 34	127.6	Te tellurium 52	[209]	Po polonium 84	-116 have	nticated	173 Yb	ytterbium 70
1	(17)	19,0 F fluorine 9	35.5 CL chlorine 17	79.9 Rr	bromine 35	126.9	iodine 53	[210]	At astatine 85	been repor		175 Lu	tutetium 71
0 (8)	(18) 4,0 He helium 2	20.2 Ne neon	39.9 Ar argon 18	83.8 Kr	krypton 36	131,3	Xenon xenon 54	[222]	Rn radon 86	ted			