Cambridge International AS & A Level

Cambridge International Advanced Subsidiary and Advanced Level

	CANDIDATE NAME							
	CENTRE CANDIE NUMBER NUMBE							
* 3	CHEMISTRY	9701/33						
6	Paper 3 Advanced Practical Skills 1	October/November 2015						
1 2	2 hou							
7 4	Candidates answer on the Question Paper.							
	Additional Materials: As listed in the Confidential Instructions							
7 *								
	READ THESE INSTRUCTIONS FIRST							
	 Write your Centre number, candidate number and name on all the work you hand in. Give details of the practical session and laboratory where appropriate, in the boxes provided. Write in dark blue or black pen. You may use an HB pencil for any diagrams or graphs. Do not use staples, paper clips, glue or correction fluid. DO NOT WRITE IN ANY BARCODES. Answer all questions. Electronic calculators may be used. You may lose marks if you do not show your working or if you do not use appropriate units. Use of a Data Booklet is unnecessary. 							
	Qualitative Analysis Notes are printed on pages 10 and 11.	Session						
	A Periodic Table is printed on page 12.							
	At the end of the examination, fasten all your work securely together.	Laboratory						
	The number of marks is given in brackets [] at the end of each question or part question.							
		For Examiner's Use						
		1						
		2						
		3						
		Total						

This document consists of **12** printed pages.



1 The formula of hydrated copper(II) sulfate is $CuSO_4$. xH_2O where x is the number of moles of water of crystallisation in one mole of salt. You will determine the value of x by titration.

When aqueous copper(II) ions react with aqueous iodide ions, I⁻, iodine is produced.

 $2Cu^{2+}(aq) + 4I^{-}(aq) \rightarrow I_2(aq) + 2CuI(s)$

The amount of iodine, I₂, produced can be found by titrating it with aqueous thiosulfate ions, S₂O₃²⁻.

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

FA 1 is aqueous $CuSO_4$.**x** H_2O containing 26.2 g dm⁻³. **FA 2** is 0.100 mol dm⁻³ sodium thiosulfate, $Na_2S_2O_3$. **FA 3** is aqueous potassium iodide, KI. starch indicator

(a) Method

- Pipette 25.0 cm³ of **FA 1** into a conical flask.
- Use the measuring cylinder to add 15 cm³ of **FA 3**, an excess of KI, to the conical flask. The solution will turn brown because iodine is formed.
- Fill the burette with **FA 2**.
- Add **FA 2** from the burette until the colour of the mixture changes to pale brown.
- Add 10 drops of starch indicator. The mixture will turn blue-black.
- Continue adding **FA 2** from the burette until the dark colour suddenly disappears to leave an off-white solid. This is the end point of the titration.
- Carry out a rough titration and record your burette readings in the space below.

The rough titre is cm³.

- Carry out as many accurate titrations as you think necessary to obtain consistent results.
- Make certain any recorded results show the precision of your practical work.
- Record in a suitable form below, all your burette readings and the volume of **FA 2** added in each accurate titration.

Ι	
II	
III	
IV	
V	
VI	
VII	

(b) From your accurate titration results, obtain a suitable value to be used in your calculations. Show clearly how you have obtained this value.

25.0 cm³ of **FA 1** required cm³ of **FA 2**. [1]

(c) Calculations

Show your working and appropriate significant figures in **each** step of your calculations.

(i) Calculate the number of moles of thiosulfate ions present in the volume of FA 2 you have calculated in (b).

moles of $S_2O_3^{2-}$ = mol

(ii) Use your answer to (i), and the equations for the reactions involved, to deduce the number of moles of Cu²⁺ present in 25.0 cm³ of **FA 1**.

 $2Cu^{2+}(aq) + 4I^{-}(aq) \rightarrow I_{2}(aq) + 2CuI(s)$ $2S_{2}O_{3}^{2-}(aq) + I_{2}(aq) \rightarrow S_{4}O_{6}^{2-}(aq) + 2I^{-}(aq)$

moles of Cu^{2+} = mol

(iii) Use your answer to (ii) and the mass of $CuSO_4$.**x** H_2O present in the solution, to calculate the relative molecular mass, M_r , of $CuSO_4$.**x** H_2O .

 $M_{\rm r}$ of CuSO₄. **x**H₂O =

(iv) Determine the value of *x*. (Use data from the Periodic Table on page 12.)

x =

[6]

[Total: 14]

2 FA 4 is an impure sample of hydrated calcium chloride, $CaCl_2.2H_2O$. On heating, hydrated calcium chloride loses its water of crystallisation.

 $CaCl_2.2H_2O(s) \rightarrow CaCl_2(s) + 2H_2O(g)$

You will determine the purity of **FA 4** by measuring the loss in mass that occurs when it is heated. The impurity present in **FA 4** is not decomposed on heating.

(a) Method

You should read the instructions carefully before starting any practical work and draw a table for your results in the space below.

- Weigh a crucible and record its mass.
- Add between 1.80 g and 2.00 g of **FA 4** into the crucible.
- Reweigh the crucible and its contents and record the mass.
- Place the crucible on the pipe-clay triangle and heat gently for 1 minute and then strongly for a further 2 minutes.
- Allow the crucible and its contents to cool. Reweigh the crucible and contents and record the mass.
- Heat the crucible strongly for a further 2 minutes. Allow it to cool. Reweigh the crucible and contents and record the mass.
- Repeat the heating, cooling and weighing until you are satisfied that all the water of crystallisation has been removed.
- Calculate and record the mass of **FA 4** used and the total mass of water lost.

While you are waiting for the crucible to cool, you may wish to start work on Question 3.

Ι	
II	
III	
IV	
V	
VI	

[6]

(b) Calculations

Show your working and appropriate significant figures in the final answer to **each** stage of your calculations.

(i) The percentage loss in mass on heating is defined as

 $\frac{\text{the loss in mass on heating}}{\text{the original mass}} \times 100.$

Calculate the percentage loss in mass of FA 4.

percentage loss in mass = %

(ii) Calculate the percentage loss in mass when **pure** hydrated calcium chloride, $CaCl_2.2H_2O$, is heated.

percentage loss in mass = %

(iii) Use your results to (i) and (ii) to calculate the percentage purity of FA4, impure CaC l_2 .2H₂O.

percentage purity =%
[3]

(c) A student carried out this experiment using 2.60 g of FA 4.

Suggest whether this experiment would give a more accurate result for the percentage purity of **FA 4**. Explain your answer.

.....[1]

(d) In your calculations you assumed that the impurity in FA 4 does not decompose on heating.

State how the percentage purity that you calculated in **(b)(iii)** would change if the impurity were to decompose on heating. Explain your answer.

......[1]

[Total: 11]

3 Qualitative Analysis

At each stage of any test you are to record details of the following.

- colour changes seen
- the formation of any precipitate
- the solubility of such precipitates in an excess of the reagent added

Where gases are released they should be identified by a test, **described in the appropriate place in your observations**.

You should indicate clearly at what stage in a test a change occurs. No additional tests for ions present should be attempted.

If any solution is warmed, a boiling tube MUST be used.

Rinse and reuse test-tubes and boiling tubes where possible.

Where reagents are selected for use in a test, the name or correct formula of the element or compound must be given.

(a) (i) FA 5, FA 6 and FA 7 are aqueous solutions each containing one anion and one cation.

Carry out the experiments described below and record your observations for each solution in the table.

	FA 5	FA 6	FA 7
To a 1 cm depth in a test-tube, add a 1 cm depth of aqueous sodium carbonate.			
To a 1 cm depth in a test-tube, add a 1 cm depth of aqueous copper(II) sulfate.			
To a 1 cm depth in a test-tube, add a 1 cm depth of aqueous barium chloride or aqueous barium nitrate.			

- (ii) What ion is present in **both FA 6** and **FA 7**?
- (iii) The anion in **FA 5** is one of carbonate, chloride, nitrate or sulfate.

Which anion is present in **FA 5**?

.....

.....

(iv) Write the ionic equation, including state symbols, for the reaction between **FA 5** and aqueous copper(II) sulfate.

.....

[7]

- (b) FA 8 contains two anions and two cations from the lists on pages 10 and 11.
 - To a 5 cm depth of distilled water in a boiling tube, add all the **FA 8**.
 - Shake the boiling tube thoroughly for one minute to make sure that no more of the solid will dissolve.
 - Filter the mixture into a clean boiling tube.
 - Place the filter funnel in a conical flask and wash the residue with a little distilled water.
 - Keep both filtrate and residue for tests (i) and (ii) below.

(i) Tests on the filtrate (the solution in the boiling tube)

Carry out the following tests and record your observations in the table below.

test	observations
To a 1 cm depth of the filtrate in a test-tube, add aqueous sodium hydroxide, then	
add aqueous hydrogen peroxide.	

(ii) Tests on the residue

Carry out the following tests and record your observations in the table below.

test	observations
Place the funnel containing the residue into a clean boiling tube. Pour approximately 5 cm ³ of dilute nitric acid onto the residue. Collect a 1 cm depth of solution in the boiling tube. Remove the funnel and return it to the conical flask.	
To this solution in the boiling tube, add aqueous sodium hydroxide.	

(iii)	Identify two cations present in FA 8 .
	cations present and
(iv)	Identify one anion present in FA 8 .
	anion present
(v)	Suggest what type of reaction is happening when hydrogen peroxide is added in test (b)(i).

[8]

test

[Total: 15]

Qualitative Analysis Notes

Key: [ppt. = precipitate]

1 Reactions of aqueous cations

ien	reaction with					
ion	NaOH(aq)	NH ₃ (aq)				
aluminium, A <i>l</i> ³⁺(aq)	white ppt. soluble in excess	white ppt. insoluble in excess				
ammonium, NH₄⁺(aq)	no ppt. ammonia produced on heating	_				
barium, Ba²⁺(aq)	no ppt. (if reagents are pure)	no ppt.				
calcium, Ca²⁺(aq)	white ppt. with high [Ca²+(aq)]	no ppt.				
chromium(III), Cr³⁺(aq)	grey-green ppt. soluble in excess giving dark green solution	grey-green ppt. insoluble in excess				
copper(II), Cu²⁺(aq)	pale blue ppt. insoluble in excess	blue ppt. soluble in excess giving dark blue solution				
iron(II), Fe²+(aq)	green ppt. turning brown on contact with air insoluble in excess	green ppt. turning brown on contact with air insoluble in excess				
iron(III), Fe³+(aq)	red-brown ppt. insoluble in excess	red-brown ppt. insoluble in excess				
magnesium, Mg²⁺(aq)	white ppt. insoluble in excess	white ppt. insoluble in excess				
manganese(II), Mn²⁺(aq)	off-white ppt. rapidly turning brown on contact with air insoluble in excess	off-white ppt. rapidly turning brown on contact with air insoluble in excess				
zinc, Zn²⁺(aq)	white ppt. soluble in excess	white ppt. soluble in excess				

2 Reactions of anions

ion	reaction
carbonate, CO ₃ ^{2–}	CO ₂ liberated by dilute acids
chloride, C <i>l⁻</i> (aq)	gives white ppt. with Ag ⁺ (aq) (soluble in $NH_3(aq)$)
bromide, Br⁻(aq)	gives cream ppt. with Ag ⁺ (aq) (partially soluble in $NH_3(aq)$)
iodide, I⁻(aq)	gives yellow ppt. with Ag ⁺ (aq) (insoluble in NH ₃ (aq))
nitrate, NO₃⁻(aq)	NH_3 liberated on heating with $OH^-(aq)$ and Al foil
nitrite, NO₂⁻(aq)	NH_3 liberated on heating with OH ⁻ (aq) and Al foil; NO liberated by dilute acids (colourless NO \rightarrow (pale) brown NO ₂ in air)
sulfate, SO ₄ ²⁻ (aq)	gives white ppt. with Ba ²⁺ (aq) (insoluble in excess dilute strong acids)
sulfite, SO ₃ ²-(aq)	SO ₂ liberated with dilute acids; gives white ppt. with Ba ²⁺ (aq) (soluble in excess dilute strong acids)

3 Tests for gases

gas	test and test result			
ammonia, NH ₃	turns damp red litmus paper blue			
carbon dioxide, CO ₂	gives a white ppt. with limewater (ppt. dissolves with excess CO ₂)			
chlorine, Cl_2	bleaches damp litmus paper			
hydrogen, H ₂	"pops" with a lighted splint			
oxygen, O ₂	relights a glowing splint			
sulfur dioxide, SO ₂	turns acidified aqueous potassium manganate(VII) from purple to colourless			

			- - - -		. · ·	E	a , ^c	– c	tium O		mini
		0	4.0 Helium	20.2 Neon 10	39.9 Ar Argon	83.8 Krypton 36	131 Xe Xenon 54	Radon 86	Ununoctium 118	175 Lutetium 71	Lr Lawrencium 103
		N		19.0 Fluorine	35.5 C 1 Chlorine	79.9 Br 35	127 I lodine 53	At Astatine 85		173 Yb 70	Nobellum 102
		N		16.0 Oxygen 8	32.1 S uffur 16	79.0 Se Selenium 34	128 Te ^{Tellunium} 52	Polonium 84	Ununhexium 116	169 Thulium 69	Mendelevium 101
		>		14.0 N Nitrogen 7	31.0 Phosphorus 15	74.9 AS Arsenic 33	122 Sb Antimony 51	209 Bismuth 83		167 Er 68	Fermium 100
		2		12.0 C Carbon 6	28.1 Silicon	72.6 Ge Germanium 32	119 Sn	207 Pb 82 Lead	Ununquadium 114	165 Holmium 67	Einsteinium 99
		■		10.8 Boron 5	27.0 Al Aluminium 13	69.7 Ga Gallium 31	115 Ln Indium 49	204 T1 Thallium 81		163 Dysprosium 66	Californium 98
ents						65.4 Zn 30	112 Cadmium 48	201 Hg ^{Mercury} 80	Ununbium 112	159 Tb ^{Terbium}	BK Berkelium 97
e Eleme	Group					63.5 Cu ^{Copper}	108 Ag Silver	197 Au Gold 79	Unununium 111	157 Gdd Gadolinium 64	66 Curium
ble of th						58.7 Nickel 28	106 Pd Palladium 46	195 Pt Platinum 78	Ununnilium 110	152 Europium 63	Americium 95
The Periodic Table of the Elements						58.9 Co ²⁷	103 Rhodium 45	192 Ir Iridium 77	Mt Meitnerium 109	150 Samarium 62	Putonium 94
			Hydrogen 1.0			55.8 Fe Iron 26	101 Ru Ruthenium 44	190 OS Osmium 76	Hassium 108	Promethium 61	Neptunium 93
						54.9 Manganese 25	Tc Technetium 43	186 Re Rhenium 75	Bh Bohrium 107	144 Neodymium 60	Uranium 92
						52.0 CT Chromium 24	95.9 Mo Molybdenum 42	184 X Tungsten 74	Seaborgium 106	141 Praseodymium 59	Pa Protactinium 91
						50.9 Vanadium 23		181 Ta Tantalum 73	Db Dubnium 105	140 Cerium 58	Thorium 90
						47.9 Ti Titanium 22	91.2 Zr Zirconium 40	178 Haffnium 72	Rf Rutherfordium 104	*	mic mass † nbol mic) number
						45.0 Sc Scandium 21	88.9 Yttrium 39	139 La Lanthanum 57 *	Actinium 89	S	a = relative atomic mass † X = atomic symbol b = proton (atomic) number
		=		9.0 Be Beryllium	24.3 Magnesium 12	40.1 Ca Calcium 20	87.6 Sr Strontium 38	137 Ba Barium 56	Radium 88	*58-71 Lanthanides 190-103 Actinides	
		_		6.9 Lithium 3	23.0 Na Sodium	39.1 Potassium 19	85.5 Rb Rubidium 37	133 CS Caesium 55	Francium 87	*58-71 L †90-103	۴ Кеу

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