

UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS General Certificate of Education Advanced Subsidiary Level and Advanced Level

CANDIDATE NAME				
CENTRE NUMBER		CANDIDATE NUMBER		
CHEMISTRY			9701/36	
Advanced Pract	Oc	October/November 2012		
			2 hours	
Candidates ans	wer on the Question Paper.			
Additional Mate	rials: As listed in the Confide	ential Instructions		
READ THESE	INSTRUCTIONS FIRST			
Advanced Pract Candidates ans Additional Mate	wer on the Question Paper. rials: As listed in the Confide INSTRUCTIONS FIRST		tober/November 20	

Give details of the practical session and laboratory where appropriate, in the boxes provided. Write in dark blue or black pen. You may use a soft pencil for any diagrams, graphs or rough working. Do not use staples, paper clips, highlighters, glue or correction fluid. DO **NOT** WRITE IN ANY BARCODES.

Answer all questions.

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 15 and 16.

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At the end of the examination, fasten all your work securely together. The number of marks is given in brackets [] at the end of each question or part question.

For Exam	iner's Use
1	
2	
3	
Total	

Laboratory

This document consists of **13** printed pages and **3** blank pages.



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1 You are to determine the concentration of aqueous copper(II) sulfate by titration. The concentration of Cu^{2+} ions in a solution can be found by reaction with an excess of aqueous iodide ions to produce iodine. The amount of iodine formed can be found by titration with thiosulfate ions, $S_2O_3^{2-}$.

FB 1 is aqueous copper(II) sulfate, $CuSO_4$. **FB 2** is 0.100 mol dm⁻³ sodium thiosulfate, $Na_2S_2O_3$. **FB 3** is aqueous potassium iodide, KI. starch indicator

Read through the instructions carefully before starting any practical work.

(a) Method

- Fill the burette with **FB 2**.
- Pipette 25.0 cm³ of **FB 1** into a conical flask.
- Use a measuring cylinder to add 10 cm³ of **FB 3** into the conical flask.
- Titrate this mixture with **FB 2** until the colour of the mixture changes from brown to yellow-brown. An off-white precipitate will also be present in the flask throughout the titration.
- Add approximately 1 cm³ of starch indicator.
- Continue the titration until the blue-black colour of the starch-iodine complex just disappears leaving the off-white precipitate.
- Perform 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 of your burette readings and the volume of **FB 2** added in each accurate titration.



[6]

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25.0 cm³ of **FB 1** required cm³ of **FB 2** [1]

(c) Calculations

The equations for the formation of iodine and its reaction with thiosulfate ions are given below.

 $\begin{array}{rcl} 2 C u^{2 +} \ + \ 4 I^{-} \ \rightarrow \ 2 C u I \ + \ I_{2} \\ \\ I_{2} \ + \ 2 S_{2} O_{3}^{\ 2 -} \ \rightarrow \ S_{4} O_{6}^{\ 2 -} \ + \ 2 I^{-} \end{array}$

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

(i) Calculate the number of moles of thiosulfate ions, S₂O₃²⁻, present in the volume of FB 2 in (b).

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

(ii) Using the equations above, deduce the number of moles of Cu²⁺ ions present in each 25.0 cm³.

moles of Cu²⁺ = mol

(iii) Calculate the concentration, in mol dm⁻³, of copper(II) sulfate in **FB 1**.

concentration of $CuSO_4 = \dots mol dm^{-3}$ [3]

Show clearly how you obtained this value.

(b) From your accurate titration results, obtain a suitable value to be used in your calculations.

(d) Three students repeated the experiment but each obtained different values for the concentration of $CuSO_4$.

The students each suggested possible improvements.

Student 1 suggested that a larger quantity of starch should be added. Student 2 suggested that a larger volume of potassium iodide, **FB 3**, should be added. Student 3 suggested that the contents of the conical flask should be filtered before titration.

Comment on the effectiveness of **each** of these possible improvements. Explain your answers.

Student 1

Student 2			
Student 3			
	 	 	[2]
			[Total: 12]

2 You are to determine the enthalpy change for the reaction between aqueous copper(II) sulfate and zinc. The enthalpy change of reaction can be found by measuring the temperature change when powdered zinc is added to aqueous copper(II) sulfate.

FB 4 is 1.10 mol dm^{-3} aqueous copper(II) sulfate, CuSO₄. powdered zinc

(a) Method

- Weigh a 100 cm³ beaker.
- In the beaker weigh out between 2.1 g and 2.3 g of powdered zinc.
- Record the weighings and the mass of zinc in the space below.

mass of zinc used = g

- Support the plastic cup in a 250 cm³ beaker.
- Use a measuring cylinder to transfer 50 cm³ of **FB 4** into the plastic cup.
- Measure and record in the table below, the initial temperature of **FB 4** in the cup.
- Start the stop watch. Measure and record the temperature of **FB 4** in the cup after 1 minute, 2 minutes and 3 minutes.
- At time 3¹/₂ minutes, add the weighed zinc to **FB 4** in the cup and stir the mixture.
- From time 4 minutes, continue to stir the mixture and measure the temperature of the contents of the cup to complete the table.

Results

time/min	0	1	2	3	4	5	6	7	8	9	10	11	12
temperature/°C													

[2]

(b) (i) On the axes opposite, plot the temperature (*y*-axis) against time (*x*-axis). The temperature axis should allow you to include a point at least 5 °C greater than the maximum temperature recorded.

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- (ii) Complete the graph to show how the temperature of the contents of the cup varies with time.
 - Draw one straight line through the points between time 0 minutes and 3 minutes.
 - Draw one straight line through the points after the maximum was reached.
 - Extrapolate these two lines and draw a vertical line at time 3¹/₂ minutes.

[4]

 $Zn(s) + Cu^{2+}(aq) \rightarrow Cu(s) + Zn^{2+}(aq)$

From the mass of zinc added and the concentration of FB 4, show that the copper(II) sulfate was in excess in your reaction. [A,: Zn, 65.4]

(iv) Assuming that the copper(II) sulfate was in excess, use your answer to (ii) to calculate the enthalpy change of the reaction between Zn(s) and Cu²⁺(aq).

Give you answer in kJ mol⁻¹ and include the relevant sign.

enthalpy change of reaction	า =		kJ mol ⁻¹
	sign	value	
	-		[6]

(c) Calculation

(d) One source of error in this experiment is due to the accuracy to which the thermometer can be read.

9

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What is the maximum error in a single temperature reading on a thermometer with graduations at $1 \,^{\circ}$ C?

maximum error =°C

Calculate the maximum percentage error when measuring a temperature rise of 12.0 °C.

maximum percentage error = % [2]

[Total: 14]

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. Marks are **not** given for chemical equations. **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.

You are provided with solutions **FB 5**, **FB 6**, **FB 7** and **FB 8**. **FB 5** and **FB 6** each contain a compound of a transition element.

Half fill a 250 cm³ beaker with water. Heat to approximately 80 °C, then stop heating and switch off the Bunsen burner. You will need this as a hot water bath on (b)(i). Continue work on (a) while the water heats.

test	observations
To 1 cm depth of FB 5 in a test-tube, add aqueous ammonia.	
To 1 cm depth of FB 5 in a test-tube, add aqueous sodium hydroxide.	
To 1 cm depth of FB 5 in a test-tube, add aqueous barium chloride or aqueous barium nitrate then,	
add an excess of either hydrochloric acid or nitric acid.	

(a) (i) Carry out the following tests on FB 5.

	(ii) From these tests, what conclusions, if any, can you reach about the identity of FB 5 ? [4]	For Examiner's Use
(b)	(i) Carry out the following tests on FB 6 .	
. ,	test observations	
	To 1 cm depth of FB 6 in a boiling tube, add 1 cm depth of FB 7 then,	
	add 1 cm depth of ethanol. Place the boiling tube in the warm water bath and leave for a few minutes.	
	To 1 cm depth of FB 6 in a test-tube, add 1 cm depth of FB 8 .	
	(ii) From these tests suggest identities for the following.	
	The anion in FB 6 is	
	The cation in FB 7 is	
	The cation in FB 8 could be or	
	 (iii) Suggest a test to determine which of the two possible cations is present in FB 8. Do not carry out this test. 	
	[7]	
(c)	Using your conclusions about the possible identities of FB 5 and FB 8 , predict the result of mixing solutions of each. Do not carry out this test.	
	Prediction	
	[1]	

- (d) Suggest what happened to the ethanol when it was warmed with the mixture of FB 6 and For Examiner's Use
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 [1]
 [1]
- (e) You are to devise and carry out a test to confirm the identity of the cation in FB 7.Record the test you use and the results of the test in the space below.

[1]

[Total: 14]

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Qualitative Analysis Notes

Key: [*ppt.* = *precipitate*]

1 Reactions of aqueous cations

	reac	tion 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 [Ca ²⁺ (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), Fe ³⁺ (aq)	red-brown ppt. insoluble in excess	red-brown ppt. insoluble in excess
lead(II), Pb²+(aq)	white ppt. soluble in excess	white 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

[Lead(II) ions can be distinguished from aluminium ions by the insolubility of lead(II) chloride.]

2 Reactions of anions

ion	reaction
carbonate, CO ₃ ²⁻	CO ₂ liberated by dilute acids
chromate(VI), CrO ₄ ^{2–} (aq)	yellow solution turns orange with H ⁺ (aq); gives yellow ppt. with Ba ²⁺ (aq); gives bright yellow ppt. with Pb ²⁺ (aq)
chloride, C <i>l</i> ⁻(aq)	gives white ppt. with Ag ⁺ (aq) (soluble in NH ₃ (aq)); gives white ppt. with Pb ²⁺ (aq)
bromide, Br ⁻ (aq)	gives cream ppt. with Ag⁺(aq) (partially soluble in NH ₃ (aq)); gives white ppt. with Pb²⁺(aq)
iodide, I⁻(aq)	gives yellow ppt. with Ag⁺(aq) (insoluble in NH₃(aq)); gives yellow ppt. with Pb²⁺(aq)
nitrate, NO ₃ ⁻(aq)	NH_3 liberated on heating with $OH^-(aq)$ and Al foil
nitrite, NO ₂ -(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) or with Pb 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 dichromate(VI) from orange to green

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