

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

CANDIDATE NAME					
CENTRE NUMBER			CANDIDATE NUMBER		

CHEMISTRY 9701/31

Paper 31 Advanced Practical Skills

May/June 2008

2 hours

Candidates answer on the Question Paper.

Additional Materials: As listed in the Instructions to Supervisors

#### **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 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 are advised to show all working in calculations.

Use of a Data Booklet is unnecessary.

Qualitative Analysis Notes are printed on pages 11 and 12.

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.

Session	
Laboratory	

For Examiner's Use		
1		
2		
Total		

This document consists of 11 printed pages and 1 blank page.

SPA (NF/KN) T58586/1 © UCLES 2008



[Turn over



## 1 Read through question 1 before starting any practical work.

You are provided with the following reagents.

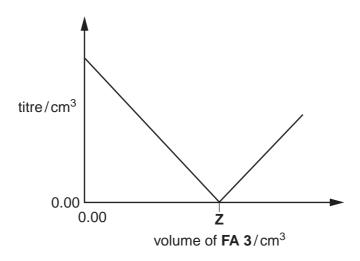
- **FA 1** containing 15.68 g dm<sup>-3</sup> of hydrated ammonium iron(II) sulphate (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>.FeSO<sub>4</sub>.6H<sub>2</sub>O
- FA 2, 0.015 mol dm<sup>-3</sup> potassium manganate(VII), KMnO<sub>4</sub>
- FA 3 containing 0.025 mol dm<sup>-3</sup> of a reagent X
- 1.0 mol dm<sup>-3</sup> sulphuric acid, H<sub>2</sub>SO<sub>4</sub>

Iron(II) ions, Fe<sup>2+</sup>, are oxidised by acidified manganate(VII) ions.

$$MnO_4^-(aq) + 8H^+(aq) + 5e^- \rightarrow Mn^{2+}(aq) + 4H_2O(l)$$
  
 $Fe^{2+}(aq) \rightarrow Fe^{3+}(aq) + e^-$ 

Reagent **X** oxidises  $Fe^{2+}$  to  $Fe^{3+}$  and is also oxidised by acidified  $MnO_4^-$ .

If varying volumes of **FA 3**, containing reagent **X**, are added to  $25.0\,\mathrm{cm}^3$  of **FA 1** in the presence of  $\mathrm{H_2SO_4}$  and the mixtures are titrated against **FA 2**, a graph of the results can be drawn as shown.



You are to determine experimentally

- Z, the exact volume of FA 3 which reacts with 25.0 cm<sup>3</sup> of FA 1,
- the mole ratio for the reaction of **FA 1** with reagent **X**.

## (a) Method

- Fill a burette with FA 2.
- Pipette 25.0 cm<sup>3</sup> of **FA 1** into a conical flask.
- Use a measuring cylinder to add approximately 10 cm<sup>3</sup> of 1.0 mol dm<sup>-3</sup> H<sub>2</sub>SO<sub>4</sub> to the solution in the flask.
- Titrate the **FA 1** in the flask with **FA 2** until the first permanent pink colour remains in the solution.

The end-point should be found after the addition of approximately 13 cm<sup>3</sup> of **FA 2**. **One** titration, performed accurately, will be sufficient.

You are reminded that just before the end-point the pink colour from a single drop of **FA 2** spreads through the whole of the solution before disappearing.



Use

For Examiner's Record your titration results in the space below.

For Examiner's Use

[1]

#### (b) Method

- Fill the second burette with FA 3.
- Empty and rinse the conical flask used in part (a).
- Pipette 25.0 cm<sup>3</sup> of FA 1 into the conical flask and add 10 cm<sup>3</sup> of H<sub>2</sub>SO<sub>4</sub> using a measuring cylinder.
- Run 12.00 cm<sup>3</sup> of FA 3 from the second burette into the flask.
- Titrate against **FA 2** until the first permanent pink colour remains in the solution.

The end-point should be found after the addition of approximately 5 cm<sup>3</sup> of **FA 2**. **One** titration, performed accurately, will be sufficient.

You are reminded that just before the end-point the pink colour from a single drop of **FA 2** spreads through the whole of the solution before disappearing.

Record your titration results in the space below.

[1]

(c) The volume of FA 3 added to the flask in (b) reacts with some but not all of the FA 1 present. Calculate the difference between the titres obtained in parts (a) and (b). Use this difference and the volume of FA 3 added to the flask in (b) to calculate the volume of FA 3 that you would expect to react with all of the Fe<sup>2+</sup> ions in 25.0 cm<sup>3</sup> of FA 1.

volume of **FA 3** = .....  $cm^3$ 

[1]



(d) The value you have obtained in (c) is an approximate value of **Z**. You are to perform four more titrations, **each** with a **different** volume of **FA 3** added to 25.0 cm<sup>3</sup> of **FA 1**, in order to plot a graph of the form shown on page 2 and to obtain an exact value for **Z**.

For Examiner's Use

One titration, performed accurately, will be sufficient for each volume of FA 3 added.

The volume of **FA 3** you have obtained in **(c)** will help you to choose suitable volumes of **FA 3** to be added for each titration.

(If you were unable to calculate the volume of **FA 3** in **(c)** assume that the value lies in the range 19.0 cm<sup>3</sup> to 21.0 cm<sup>3</sup>.)

#### Remember

- you should not use more than 40.0 cm<sup>3</sup> of FA 3 for any single titration,
- you already have data for titrations with no FA 3 added and with 12.00 cm<sup>3</sup> added.

Prepare a table in the space below and use it to record the titration results for each volume of **FA 3** added. Include in your table the titre values from parts **(a)** and **(b)**.

[6]

(e) Use the grid on the opposite page to plot a graph of titre against volume of FA 3 added.

Draw **two** straight lines through the plotted points to find **Z**, the volume of **FA 3** that just reacts with the  $Fe^{2+}$  ions in  $25.0 \, \text{cm}^3$  of **FA 1**.



For Examiner's Use

[7]

[Turn over

Calculations  Show your working and appropriate significant figures in all of your calculations. [2]  (g) Calculate how many moles of Fe <sup>2+</sup> ions were pipetted into the flask.  [A <sub>r</sub> : Fe, 55.8; H, 1.0; N, 14.0; O, 16.0; S, 32.1]	(f)	Circle on the graph <b>one</b> point where a repeat titration might be appropriate and justify you decision. If you do not think that any titration needs to be repeated, explain why you have come to that conclusion.	For Examiner's Use
Show your working and appropriate significant figures in <b>all</b> of your calculations. [2]  (g) Calculate how many moles of Fe <sup>2+</sup> ions were pipetted into the flask. [A <sub>i</sub> : Fe, 55.8; H, 1.0; N, 14.0; O, 16.0; S, 32.1]		[1]	
(g) Calculate how many moles of Fe <sup>2+</sup> ions were pipetted into the flask.  [A <sub>i</sub> : Fe, 55.8; H, 1.0; N, 14.0; O, 16.0; S, 32.1]	Cal	culations	
[A <sub>r</sub> : Fe, 55.8; H, 1.0; N, 14.0; O, 16.0; S, 32.1]	Sho	w your working and appropriate significant figures in <b>all</b> of your calculations. [2]	
(h) Calculate how many moles of X are present in Z, the volume of FA 3 read from your graph.	(g)		
graph.		mol of Fe <sup>2+</sup> were pipetted into the flask. [2]	
(i) Calculate, to 3 significant figures, the number of moles of Fe <sup>2+</sup> ions that react with 1 mol of <b>X</b> .	(h)	· · · · · · · · · · · · · · · · · · ·	
1 mol of <b>X</b> .		mol of <b>X</b> were present in cm <sup>3</sup> of <b>FA 3</b> . [1]	
	(i)		
		[2]	
		[Total: 24]	

studyguide.pk

7

**BLANK PAGE** 



2 The three solutions **FA 4**, **FA 5**, and **FA 6 each** contain **one** of the following. aluminium sulphate,  $Al_2(SO_4)_3$  ammonium iodide,  $NH_4I$  zinc nitrate,  $Zn(NO_3)_2$ 

For Examiner's Use

(a) Use the information on page 12 to select two suitable reagents to use to discover which solution contains iodide ions.

Record, in the space below, the reagents used and the observations made.

From these tests, solution FA ...... contains iodide ions. [5]

You are to perform the tests given in the table opposite on each of **FA 4**, **FA 5** and **FA 6** to identify, where possible, the cation and anion present in each solution.

Record details of colour changes seen, the formation of any precipitate and the solubility of any such precipitate 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 table.

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.



For Examiner's Use

	test	observations with <b>FA 4</b>	observations with FA 5	observations with FA 6
(b)	To 1 cm depth of solution in a test-tube, add aqueous sodium hydroxide drop-by-drop until it is in excess.			
(c)	To 1 cm depth of solution in a test-tube, add aqueous ammonia drop-by-drop until it is in excess.			
(d)	To 1 cm depth of solution in a test-tube, add aqueous barium chloride,			
	add dilute hydrochloric acid.			
(e)	To 1 cm depth of solution in a boiling-tube add 2 cm depth of water and 1 cm depth of aqueous lead(II) nitrate,			
	if a precipitate has formed, cautiously warm until the solution boils,			
	cool the tube by standing it in a beaker of cold water.			

[6]



For Examiner's Use

(T)	performed to identify the cations and anions present.  State clearly where a cation or an anion has <b>not</b> been specifically identified.					
	FA 4 contains					
	supporting evidence					
	FA 5 contains					
	supporting evidence					
	FA 6 contains					
	supporting evidence					
	[4]					
(g)	When testing a solution containing both $NH_4I$ and $Zn(NO_3)_2$ , suggest why a student should identify the $NH_4^+$ ion before attempting to identify the $NO_3^-$ ion. The Qualitative Analysis Notes on pages 11 and 12 should help you to answer this.					
	[1]					
	[Total: 16]					



**(f)** 

# **Qualitative Analysis Notes**

Key: [ppt. = precipitate]

# 1 Reactions of aqueous cations

ion	reaction with					
	NaOH(aq)	NH <sub>3</sub> (aq)				
aluminium, Al <sup>3+</sup> (aq)	white ppt. soluble in excess	white ppt. insoluble in excess				
ammonium, NH <sub>4</sub> <sup>+</sup> (aq)	ammonia produced on heating					
barium, Ba <sup>2+</sup> (aq)	no ppt. (if reagents are pure)	no ppt.				
calcium, Ca <sup>2+</sup> (aq)	white ppt. with high [Ca <sup>2+</sup> (aq)]	no ppt.				
chromium(III), Cr <sup>3+</sup> (aq)	grey-green ppt. soluble in excess giving dark green solution	grey-green ppt. insoluble in excess				
copper(II), Cu <sup>2+</sup> (aq)	pale blue ppt. insoluble in excess	blue ppt. soluble in excess giving dark blue solution				
iron(II), Fe <sup>2+</sup> (aq)	green ppt. insoluble in excess	green ppt. insoluble in excess				
iron(III), Fe <sup>3+</sup> (aq)	red-brown ppt. insoluble in excess	red-brown ppt. insoluble in excess				
lead(II), Pb <sup>2+</sup> (aq)	white ppt. soluble in excess	white ppt. insoluble in excess				
magnesium, Mg <sup>2+</sup> (aq)	white ppt. insoluble in excess	white ppt. insoluble in excess				
manganese(II), Mn <sup>2+</sup> (aq)	off-white ppt. insoluble in excess	off-white ppt. insoluble in excess				
zinc, Zn <sup>2+</sup> (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 <sub>2</sub> liberated by dilute acids
chromate(VI), CrO <sub>4</sub> <sup>2-</sup> (aq)	yellow solution turns orange with H <sup>+</sup> (aq); gives yellow ppt. with Ba <sup>2+</sup> (aq); gives bright yellow ppt. with Pb <sup>2+</sup> (aq)
chloride, Cl <sup>-</sup> (aq)	gives white ppt. with Ag <sup>+</sup> (aq) (soluble in NH <sub>3</sub> (aq)); gives white ppt. with Pb <sup>2+</sup> (aq)
bromide, Br <sup>-</sup> (aq)	gives pale cream ppt. with Ag <sup>+</sup> (aq) (partially soluble in NH <sub>3</sub> (aq)); gives white ppt. with Pb <sup>2+</sup> (aq)
iodide, I <sup>-</sup> (aq)	gives yellow ppt. with Ag <sup>+</sup> (aq) (insoluble in NH <sub>3</sub> (aq)); gives yellow ppt. with Pb <sup>2+</sup> (aq)
nitrate, NO <sub>3</sub> (aq)	NH <sub>3</sub> liberated on heating with OH <sup>-</sup> (aq) and A <i>l</i> foil
nitrite, NO <sub>2</sub> (aq)	NH <sub>3</sub> liberated on heating with OH⁻(aq) and A <i>l</i> foil; NO liberated by dilute acids (colourless NO → (pale) brown NO <sub>2</sub> in air)
sulphate, SO <sub>4</sub> <sup>2-</sup> (aq)	gives white ppt. with $\mathrm{Ba^{2+}}(\mathrm{aq})$ or with $\mathrm{Pb^{2+}}(\mathrm{aq})$ (insoluble in excess dilute strong acids)
sulphite, SO <sub>3</sub> <sup>2-</sup> (aq)	SO <sub>2</sub> liberated with dilute acids; gives white ppt. with Ba <sup>2+</sup> (aq) (soluble in excess dilute strong acids)

## 3 Tests for gases

gas	test and test result
ammonia, NH <sub>3</sub>	turns damp red litmus paper blue
carbon dioxide, CO <sub>2</sub>	gives a white ppt. with limewater (ppt. dissolves with excess CO <sub>2</sub> )
chlorine, Cl <sub>2</sub>	bleaches damp litmus paper
hydrogen, H <sub>2</sub>	'pops' with a lighted splint
oxygen, O <sub>2</sub>	relights a glowing splint
sulphur dioxide, SO <sub>2</sub>	turns potassium dichromate(VI) (aq) from orange to green

Permission to reproduce items where third-party owned material protected by copyright is included has been sought and cleared where possible. Every reasonable effort has been made by the publisher (UCLES) to trace copyright holders, but if any items requiring clearance have unwittingly been included, the publisher will be pleased to make amends at the earliest possible opportunity.

University of Cambridge International Examinations is part of the Cambridge Assessment Group. Cambridge Assessment is the brand name of University of Cambridge Local Examinations Syndicate (UCLES), which is itself a department of the University of Cambridge.

