

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

	CANDIDATE NAME		
	CENTRE CAN NUMBER NUM	NDIDATE MBER	
* 59470	CHEMISTRY Paper 31 Advanced Practical Skills		9701/31 May/June 2008 2 hours
	Candidates answer on the Question Paper.		
4	Additional Materials: As listed in the Instructions to Supervisors		
0 *	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.



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1 Read through question 1 before starting any practical work.

You are provided with the following reagents.

- FA 1 containing 15.68 g dm⁻³ of hydrated ammonium iron(II) sulphate (NH₄)₂SO₄.FeSO₄.6H₂O
- **FA 2**, 0.015 mol dm⁻³ potassium manganate(VII), KMnO₄
- FA 3 containing 0.025 mol dm⁻³ of a reagent X
- 1.0 mol dm⁻³ sulphuric acid, H_2SO_4

Iron(II) ions, Fe²⁺, are oxidised by acidified manganate(VII) ions.

$$MnO_{4}^{-}(aq) + 8H^{+}(aq) + 5e^{-} \rightarrow Mn^{2+}(aq) + 4H_{2}O(I)$$

Fe²⁺(aq) → Fe³⁺(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 cm^3 of **FA 1** in the presence of 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 cm³ 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 cm³ of **FA 1** into a conical flask.
- Use a measuring cylinder to add approximately 10 cm³ of 1.0 mol dm⁻³ H₂SO₄ 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^3 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.

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Record your titration results in the space below.

(b) Method

- Fill the second burette with **FA 3**.
- Empty and rinse the conical flask used in part (a).
- Pipette 25.0 cm³ of FA 1 into the conical flask and add 10 cm³ of H₂SO₄ using a measuring cylinder.
- Run 12.00 cm³ 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^3 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 Fe²⁺ ions in 25.0 cm³ of FA 1.

volume of **FA 3** = \dots cm³

[1]

[1]

For Examiner's Use (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 cm³ of FA 1, in order to plot a graph of the form shown on page 2 and to obtain an exact value for Z.

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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^3 to 21.0 cm^3 .)

Remember

- you should not use more than 40.0 cm³ of **FA 3** for any single titration,
- you already have data for titrations with no **FA 3** added and with 12.00 cm³ 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 cm³ of **FA 1**.

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- 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, $Zn(NO_3)_2$
 - (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

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	test	observations with FA 4	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, then			
	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, then			
	if a precipitate has formed, cautiously warm until the solution boils, then			
	cool the tube by standing it in a beaker of cold water.			

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Qualitative Analysis Notes

Key: [ppt. = precipitate]

1 Reactions of aqueous cations

ion	reaction with		
	NaOH(aq)	NH ₃ (aq)	
aluminium,	white ppt.	white ppt.	
Al ³⁺ (aq)	soluble in excess	insoluble in excess	
ammonium, NH ₄ (aq)	ammonia produced on heating		
barium, Ba ²⁺ (aq)	no ppt. (if reagents are pure)	no ppt.	
calcium, Ca ²⁺ (aq)	white ppt. with high [Ca ²⁺ (aq)]	no ppt.	
chromium(III), Cr ³⁺ (aq)	grey-green ppt. soluble in excess giving dark green solution	grey-green ppt. insoluble in excess	
copper(II),	pale blue ppt.	blue ppt. soluble in excess	
Cu ²⁺ (aq)	insoluble in excess	giving dark blue solution	
iron(II),	green ppt.	green ppt.	
Fe ²⁺ (aq)	insoluble in excess	insoluble in excess	
iron(III),	red-brown ppt.	red-brown ppt.	
Fe ³⁺ (aq)	insoluble in excess	insoluble in excess	
lead(II),	white ppt.	white ppt.	
Pb ²⁺ (aq)	soluble in excess	insoluble in excess	
magnesium,	white ppt.	white ppt.	
Mg ²⁺ (aq)	insoluble in excess	insoluble in excess	
manganese(II),	off-white ppt.	off-white ppt.	
Mn ²⁺ (aq)	insoluble in excess	insoluble in excess	
zinc,	white ppt.	white ppt.	
Zn ²⁺ (aq)	soluble in excess	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 pale cream ppt. with Ag ⁺ (aq) (partially soluble in $NH_3(aq)$); gives white ppt. with $Pb^{2+}(aq)$
iodide, I ⁻ (aq)	gives yellow ppt. with Ag ⁺ (aq) (insoluble in NH ₃ (aq)); gives yellow ppt. with Pb ²⁺ (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 A <i>l</i> foil; NO liberated by dilute acids (colourless NO \rightarrow (pale) brown NO ₂ in air)
sulphate, SO ₄ ²⁻ (aq)	gives white ppt. with $Ba^{2+}(aq)$ or with $Pb^{2+}(aq)$ (insoluble in excess dilute strong acids)
sulphite, SO ₃ ^{2−} (aq)	SO ₂ liberated with dilute acids; gives white ppt. with Ba ²⁺ (aq) (soluble in excess dilute strong acids)

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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
sulphur dioxide, SO ₂	turns potassium dichromate(VI) (aq) from orange to green

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