

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

CANDIDATE NAME		
CENTRE NUMBER	CANDIDATE NUMBER	
CHEMISTRY Advanced Prac		9701/31
		er 2013
Candidates ans Additional Mate		2 hours
Candidates ans	swer on the Question Paper.	
Additional Mate	erials: As listed in the Confidential Instructions	
READ THESE		

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.

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 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

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1	
2	
3	
Total	

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



1 In this experiment you are to determine the relative formula mass of an iron(II) salt by titration with potassium manganate(VII).

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FA 1 is the iron(II) salt. FA 2 is $0.0200 \text{ mol dm}^{-3}$ potassium manganate(VII), KMnO₄. FA 3 is dilute sulfuric acid, H₂SO₄.

(a) Method

Preparing a solution of FA 1

- Weigh the 250 cm³ beaker and record the mass in the space below.
- Add all the **FA 1** provided to the beaker. Weigh the beaker with **FA 1** and record the mass.
- Calculate the mass of **FA 1** used and record this in the space below.
- Use a measuring cylinder to add approximately 100 cm³ of **FA 3** to the beaker. Stir until all the solid has dissolved.
- Transfer the solution into the 250 cm³ volumetric (graduated) flask labelled **FA 4**.
- Wash out the beaker thoroughly using distilled water and add the washings to the volumetric flask. Make the solution up to the mark using distilled water.
- Shake the flask thoroughly to mix the solution before using it for your titrations.
- This solution of the iron(II) salt is **FA 4**.

Titration

- Pipette 25.0 cm³ of **FA 4** into a conical flask.
- Use a measuring cylinder to add 20 cm³ of **FA 3** to the flask.
- Fill the burette with **FA 2**.
- Titrate FA 4 with FA 2 until the solution changes to a permanent pink colour.
- Perform a rough titration and record your burette readings in the space below.

The rough titre is cm³.

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- Carry out as many accurate titrations as you think necessary to obtain consistent results.
- Make sure 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 **FA 2** added in each accurate titration.

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[7]

(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 cm^3 of FA 4 required cm³ of FA 2 [1]

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(c) Calculations

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

(i) Calculate the number of moles of potassium manganate(VII) present in the volume of FA 2 calculated in (b).

moles of $KMnO_4$ = mol

(ii) The half-equation for the reduction of a manganate(VII) ion is:

 MnO_4^- + $8H^+$ + $5e^- \rightarrow Mn^{2+}$ + $4H_2O$

Give the half-equation for the oxidation of an iron(III) ion to an iron(III) ion.

.....

Therefore, 1 mole of manganate(VII) ions reacts with 5 moles of iron(II) ions.

(iii) Calculate the number of moles of iron(II) ions present in 25.0 cm³ of solution FA 4.

moles of Fe^{2+} in 25.0 cm³ of **FA 4** = mol

(iv) Calculate the number of moles of iron(II) ions present in 250 cm³ of solution FA 4.

moles of Fe^{2+} in 250 cm³ of **FA 4** = mol

(v) In 1 mole of the iron(II) salt, FA 1, there is 1 mole of iron(II) ions.
 Use the mass of FA 1 you weighed out to calculate the relative formula mass of the iron(II) salt.

(d) (i) A 25 cm³ pipette is accurate to ±0.06 cm³. Calculate the maximum percentage error when the pipette was used to measure solution **FA 4**.

percentage error in measuring FA 4 =%

(ii) State the maximum error in the mass of the 250 cm³ beaker that you recorded in (a).

maximum error = g

(iii) Calculate the maximum percentage error in the mass of FA 1 used in (a).

maximum percentage error =%

[Total: 15]

2 In this experiment you are to determine the formula of hydrated barium chloride, FA 5, by Examiner's heating to remove the water of crystallisation. You will heat two separate samples. The anhydrous barium chloride does not decompose when heated.

FA 5 is hydrated barium chloride, BaCl₂.**x**H₂O

(a) Method

Record **all** weighings, in an appropriate form, in the space below.

- Record the mass of the empty crucible without its lid.
- Add between 2.0 and 2.4 g of FA 5 into the crucible. Record the mass of the crucible and its contents.
- Use a pipe-clay triangle to support the crucible and contents on a tripod.
- Heat the crucible and its contents gently for about one minute with the lid off. Then heat strongly for a further four minutes.
- Put the lid on the crucible and leave to cool for approximately 10 minutes.

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

- When the crucible is cool, **remove the lid**, and weigh the crucible with the residue.
- Record the mass of anhydrous barium chloride remaining in the crucible after heating and the mass of water lost.
- To prepare for the second experiment, use a spatula to remove the residue from the crucible into the beaker labelled waste.
- Reweigh the empty crucible without its lid.
- Carry out the experiment again. This time use between 1.5 and 1.9 g of FA 5.

Ι	
II	
III	
IV	
V	
VI	

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[6]

		-	
(b)	Cal	culation	For Examiner's
Sh		ow your working in each step.	Use
	(i)	Calculate the mean number of moles of water removed from the hydrated salt in the experiments. (<i>A</i> _r : H, 1.0; O, 16.0)	
		moles of $H_2O = \dots \dots \dots \dots \dots \dots \dots \dots$	
	(ii)	Calculate the mean number of moles of anhydrous barium chloride produced in the experiments. (A_r : Ba, 137; Cl, 35.5)	
		moles of $BaCl_2 = \dots mol$	
((iii)	Calculate the value of x in the formula of hydrated barium chloride, $BaCl_2$. x H ₂ O.	
		x =[3]	
(c)	(i)	Suggest how the experimental procedure could be modified to ensure that all of the water of crystallisation had been removed by heating hydrated FA 5 .	
	(ii)	Do you think that the results from your two experiments are consistent with each other? Justify your answer by carrying out appropriate calculations.	

[3]

[Total: 12]

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) **FA 5** is hydrated barium chloride.

FA 6 is the same iron(II) salt used in Question 1. It contains **one other** cation and **one** anion.

(i) Place a small spatula measure of **FA 6** into a test-tube. Dissolve the solid in about a 5 cm depth of distilled water. Use the solution for the following tests.

test	observations
To a 1 cm depth of aqueous FA 6 in a boiling tube, add aqueous sodium hydroxide until no further change occurs, then	
heat the mixture carefully.	
Dissolve a few crystals of FA 5 in a 1 cm depth of distilled water in a test-tube. Add a 1 cm depth of FA 6 , then	
add excess dilute hydrochloric acid to the mixture.	

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(ii)	Identify the ions present in FA 6.	For Examiner's
	cations: Fe ²⁺ and anion:	Use
(iii)	Give the ionic equation for the reaction of iron(II) ions with hydroxide ions.	
(iv)	Place a small spatula measure of FA 6 into a hard-glass test-tube. Heat gently, then strongly, until no further change is observed. Record your observations in the space below.	

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(b) FA 7, FA 8 and FA 9 are aqueous solutions. Each contains one cation and one anion. **FA 3** is dilute sulfuric acid, H_2SO_4 .

Mix pairs of solutions so that you can complete the table below. For each test, use 1 cm depths of each solution in clean test-tubes. Record your observations in the table.

	FA 7	FA 8	FA 9
FA 3			
FA 7			
FA 8			

From your observations, complete the following statements.

The anion in FA 7 is	
-----------------------------	--

The cation in FA 8 is

The anion in **FA 9** is

[6]

[Total: 13]

Qualitative Analysis Notes

Key: [ppt. = precipitate]

1 Reactions of aqueous cations

inn	reaction with	
ion	NaOH(aq)	NH ₃ (aq)
aluminium, Al 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), Fe³+(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 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, NO₂⁻(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)
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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