

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

CANDIDATE NAME			
CENTRE NUMBER		CANDIDATE NUMBER	
CHEMISTRY	·		9701/33
Paper 33 Practi	ical lest	Oc	tober/November 2009 2 hours
Candidates ans	wer on the Question Paper.		
Additional Mate	rials: As listed in the Confidential Instructions		
READ THESE I	NSTRUCTIONS 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 10 and 11.

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	
Laboratory	

For Examiner's Use	
1	
2	
3	
Total	

This document consists of 10 printed pages and 2 blank pages.



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1 Read through question 1 before starting any practical work.

You are provided with the following reagents.

- **FA 1**, hydrated iron(II) sulfate
- FA 2, aqueous iron(II) sulfate
- **FA 3**, aqueous potassium manganate(VII)
- FA 4, sulfuric acid

The formula of hydrated iron(II) sulfate is $FeSO_4$. xH_2O where x shows the number of molecules of water of crystallisation present.

The value of **x** can be found by two different methods.

Method 1 involves heating to drive off water of crystallisation while **Method 2** uses a titration to determine the concentration of $Fe^{2+}(aq)$.

- (a) Method 1
 - Weigh a crucible and record the mass.
 - Add between 1.80 g and 2.00 g of **FA 1** and record the new mass.
 - Place the crucible containing **FA 1** on a pipe clay triangle and heat gently for about four minutes with a Bunsen burner.
 - Allow the crucible to cool. You should continue with **Method 2** while the crucible is cooling.
 - Weigh the crucible and its contents.

Record all masses in the space below.

[3]

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(b) Calculate the mass of water lost and the mass of iron(II) sulfate that remained after heating.

mass of water lost = g

mass of iron(II) sulfate remaining = g

[1]

(c) Use your answer to (b) to calculate how many moles of water were lost and the moles of iron(II) sulfate, FeSO₄, remaining after heating.
 Show all of your working.
 [A_r: Fe, 55.8; H, 1.0; O, 16.0; S, 32.1]

The hydrated iron(II) sulfate contained mol of water

and mol of FeSO₄.

(d) Use your answer to (c) to determine the value of \boldsymbol{x} in the formula of hydrated iron(II) sulfate, FeSO₄. \boldsymbol{x} H₂O.

(e) Method 2

- Fill the burette with **FA 3**, aqueous potassium manganate(VII).
- Pipette 25.0 cm³ of **FA 2** into a conical flask and use a measuring cylinder to add approximately 20 cm³ of **FA 4**.
- Titrate this solution with **FA 3** from the burette until the first permanent pink colour remains in the solution.
- Perform sufficient further titrations to obtain accurate results.
- Record your titration results in the space below. Make certain that your recorded results show the precision of your working.

Summary

 25.0 cm^3 of **FA 2** reacted with cm³ of **FA 3**.

Show which results you used to obtain the value of the volume of **FA 3** by placing a tick (\checkmark) under the readings used in your results. [11]

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i ii iii iv v v vi vii viii ix x xi

For Examiner's Use

[2]

[Turn over

(f) All experimental methods contain errors, some of which are concerned with uncertainty of measurements.

For Examiner's Use

[2]

Complete the table below to show the uncertainties in measuring the volume of potassium manganate(VII) used in **Method 2**.

maximum uncertainty in a single reading with a burette	± cm ³
volume of potassium manganate(VII), FA 3 , from the summary in (e)	cm ³
maximum percentage error in the volume of potassium manganate(VII) used	%

(g) Method 1 is usually less accurate than Method 2 for finding the value of x in the formula of hydrated iron(II) sulfate, FeSO₄.xH₂O.

A group of students carried out **Method 1** correctly but calculated a value of 9 for x. The true value for x is 7.

Suggest an error in the practical procedure of the experiment that could account for this difference.

.....[1]

.....

(h) Suggest a modification that could be made to the practical procedure in **Method 1** to reduce this error.

Explain why this modification should give an answer nearer to 7.

modification

......[2]

[Total: 24]

BEFORE STARTING QUESTION 2, heat a half-full 250 cm^3 beaker of water for use as a hot water-bath in question 3.

2 The four solutions **FA 5**, **FA 6**, **FA 7** and **FA 8** each contain one of the following anions.

- chloride, Cl⁻
- iodide, I⁻
- nitrate, NO₃
- nitrite, NO_2^{-3}

Use information from the Qualitative Analysis Notes on page 11 to answer the following questions.

(a) Which single reagent could you use to identify the solution containing the nitrite ion?

.....

Which single reagent could you use to identify the solutions containing the chloride and the iodide ion?

-[1]
- (b) Use the reagents selected in (a) to test each of the solutions.

Rinse and reuse test-tubes where possible.

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

From your observations identify the solutions containing chloride, iodide and nitrite ions. In each case give evidence to support your answer.

solution contains the chloride ion.
supporting evidence
solution contains the iodide ion.
supporting evidence
solution contains the nitrite ion.
supporting evidence

i	
ii	
iii	
iv	
v	
vi	
vii	

For Examiner's Use

For	Do not carry out this test.	(c)
Examiner's Use	State another test that you could use to confirm the presence of chloride and iodide ions.	
	[1]	
	[Total: 9]	

3 (a) You are to carry out the tests given in the table below on solutions FA 9 and FA 10.You should record details of colour changes seen and the formation of any precipitate.

No additional tests should be attempted.

Reheat your water bath until the water boils. Turn off the Bunsen burner.

	test	observations
(i)	To 1 cm depth of FA 9 in a test-tube, add 1 cm depth of dilute hydrochloric acid.	
(ii)	To 1 cm depth of FA 9 in a test-tube, add 1 cm depth of dilute sulfuric acid.	
(iii)	To 1 cm depth of FA 10 in a boiling-tube, add dilute sulfuric acid until no further change occurs. Use this solution for test (iv).	
(iv)	To the solution left after test (iii) add 1 cm depth of ethanol. Place the mixture in your hot water bath and leave for approximately 3 minutes.	
(v)	To 1cm depth of FA 9 in a test-tube add 1 cm depth of FA 10 .	

For Examiner's Use [Total: 7]

Key: [ppt. = precipitate.]

1 Reactions of aqueous cations

	react	ion with
	NaOH(aq)	NH ₃ (aq)
aluminium, A <i>l</i> ³⁺ (aq)	white ppt. soluble in excess	white ppt. insoluble in excess
ammonium, NH ₄ +(aq)	no ppt. 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), Cu ²⁺ (aq)	pale blue ppt. insoluble in excess	blue ppt. soluble in excess giving dark blue solution
iron(II), Fe ²⁺ (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), Pb ²⁺ (aq)	white ppt. soluble in excess	white ppt. insoluble in excess
magnesium, Mg ²⁺ (aq)	white ppt. insoluble in excess	white ppt. insoluble in excess
manganese(II), Mn ²⁺ (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, Zn ²⁺ (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 soln turns orange with H ⁺ (aq); gives yellow ppt. with Ba ²⁺ (aq); gives bright yellow ppt. with Pb ²⁺ (aq)
chloride,	gives white ppt. with Ag ⁺ (aq) (soluble in $NH_3(aq)$);
C <i>l</i> ⁻ (aq)	gives white ppt. with $Pb^{2+}(aq)$
bromide,	gives cream ppt. with Ag ⁺ (aq) (partially soluble in $NH_3(aq)$);
Br ⁻ (aq)	gives white ppt. with $Pb^{2+}(aq)$
iodide,	gives yellow ppt. with Ag ⁺ (aq) (insoluble In NH ₃ (aq));
I (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 Al foil, NO liberated by dilute acids (colourless NO \rightarrow (pale) brown NO ₂ in air)
sulfate,	gives white ppt. with Ba ²⁺ (aq) (insoluble in excess dilute strong acid);
SO ₄ ²⁻ (aq)	gives white ppt. with Pb ²⁺ (aq)
sulfite,	SO ₂ liberated with dilute acids;
SO ₃ ^{2–} (aq)	gives white ppt. with Ba ²⁺ (aq) (soluble in excess dilute strong acid)

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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