

# 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

October/November 2009

2 hours

Candidates answer on the Question Paper.

Additional Materials: As listed in the Confidential Instructions

#### **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.



# **BLANK PAGE**

1 You are provided with the following reagents.

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[1]

- two weighing bottles labelled FA 1, each containing between 2.90g and 3.00g of zinc powder
- FA 2, 0.80 mol dm<sup>-3</sup> copper sulfate, CuSO<sub>4</sub>

You are to determine the enthalpy change,  $\Delta H$ , for the following reaction.

$$Zn(s) + CuSO_4(aq) \rightarrow Cu(s) + ZnSO_4(aq)$$

You will carry out the experimental procedure twice.

#### Read through the instructions below before starting the experiment.

(a) You will weigh each bottle and later in the experiment weigh it again after the zinc powder has been tipped into copper sulfate solution. In the space below prepare a table to record the weighings and the mass of zinc powder used in each experiment.

Weigh accurately, to at least one decimal place, one of the weighing bottles labelled **FA 1**.

Record this mass in the table you have prepared.

(b) Procedure

- Support the plastic cup in the 250 cm<sup>3</sup> beaker and, using a pipette, place 25.0 cm<sup>3</sup> of **FA 2** into the plastic cup.
- Stir gently, taking a temperature reading every ½ minute until a steady temperature has been obtained for a period of at least 2 minutes. You may need to tilt the beaker in order to cover the bulb of the thermometer with solution.
- On a precise minute reading tip the zinc powder from the weighing bottle into the plastic cup.

#### Do not read the temperature at this time or at the following ½ minute.

- Continue to stir the mixture thoroughly. Starting 1 minute after the addition of the zinc powder, record the temperature every ½ minute until the temperature has reached a maximum value and then decreased steadily for at least 5 minutes.
- Reweigh the empty weighing bottle. Record the mass of the bottle + any residual zinc powder and the mass of zinc powder used in the experiment in the table you prepared in (a).
- Record your results in an appropriate form in the space on the following page.

Repeat the experiment using the contents of the second weighing bottle and 25.0 cm<sup>3</sup> copper sulfate solution pipetted into a clean plastic cup.

### (b) continued

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**Results** Make certain your readings of temperature display the precision of the apparatus used.

[11]

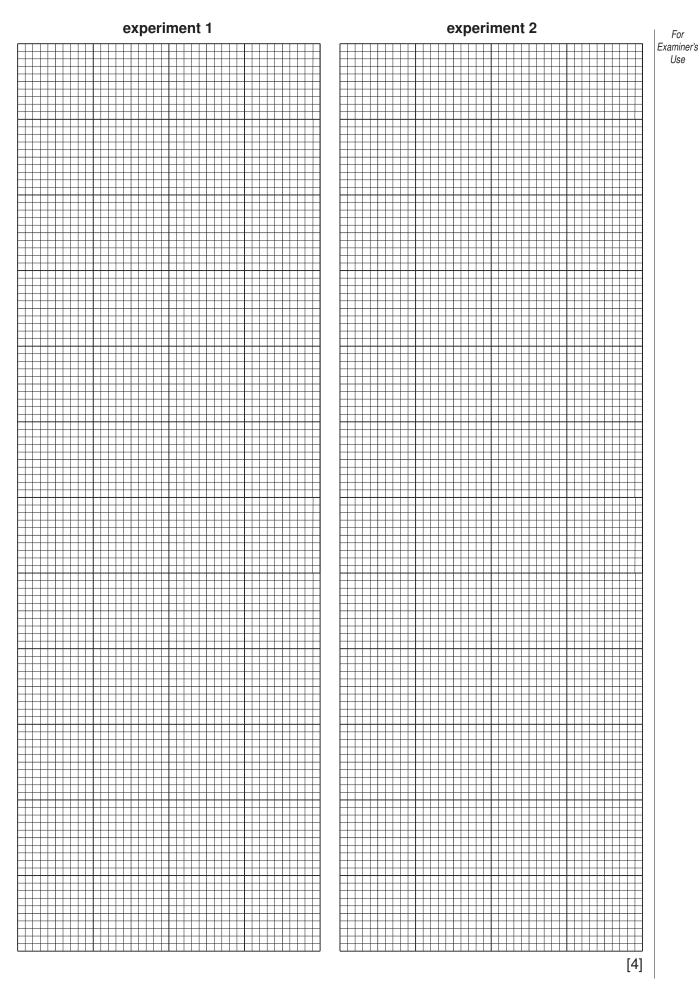
(c) Plot your temperature and time readings separately for each experiment on the grids on the next page. Your temperature axis should extend 10°C **above** the highest temperature you recorded.

Draw lines as instructed below.

On each graph draw a horizontal straight line through the steady initial temperature.

Extrapolate the cooling section of each graph back to the time when you added the zinc powder.

Draw construction lines on the graphs to deduce the "theoretical" **temperature rise** at the moment of mixing the reagents.



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(d)	The "theoretical" temperature rises are	e°C	and	. °C.
	The mean "theoretical" temperature ris	se is	. °C.	[1]
	Calculations			
	Show working and appropriate significa	nt figures in <b>all</b> of	your calculations.	[2]
(e)	Calculate how many moles of copper cup.	sulfate, CuSO <sub>4</sub> ,	were pipetted into	the plastic
		mol of Cus	SO <sub>4</sub> were pipetted	into the cup
	For each experiment calculate how many plastic cup. [A <sub>r</sub> : Zn, 65.4]	nany moles of zir	nc powder were ac	dded to the
<b>1</b> <sup>st</sup> (	experiment	2 <sup>nd</sup> experiment		
	In the 1 <sup>st</sup> experiment mol			
	тт ито д тохротипоне	1 01 21110 9011401 11	roro addod to trio p	[1]
(f)	Use your answers to (e) and the equawas in excess and which was the limiting			nich reagent
	$Zn(s) + CuSO_4(aq) -$	→ Cu(s) + ZnSO	<sub>4</sub> (aq)	
				[1]

(g)	From your mean "theoretical" temperature rise at the time of mixing, calculate the heat energy released in the plastic cup by the reaction of zinc powder with copper sulfate solution.  [You may assume that 4.3 J are required to raise the temperature of 1 cm <sup>3</sup> of any solution by 1 °C and that the mass of any solid may be ignored.]	For Examiner's Use
	of heat energy are released. [1]	
(h)	Calculate, correct to 3 significant figures, the enthalpy change in $kJ  mol^{-1}$ for the following reaction.	
	$Zn(s) + CuSO_4(aq) \longrightarrow Cu(s) + ZnSO_4(aq)$	
	$\Delta H = \dots kJ  \text{mol}^{-1}$ [2]	
(i)	Identify and explain one source of error in the experiment you have carried out.	
	[1]	
(j)	Suggest a way in which the experimental method you used could be improved in a school or college laboratory in order to minimise this error.	
	[1]	
	[Total: 26]	

2 The three boiling-tubes, labelled **FA 3**, **FA 4**, and **FA 5**, each contain a solid with one cation and one anion from those listed on pages 11 and 12.

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You will carry out specified tests to deduce the cations and anions present in **FA 3**, **FA 4** and **FA 5**.

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.

No additional tests for ions present should be attempted. If any solution is warmed a boiling-tube MUST be used.

(a) Heat the boiling-tube containing **FA 5** gently at first then more strongly. Record your observations in the space below.

[2]

(b) In their boiling-tubes, dissolve FA 3, FA 4 and the cold residue after heating FA 5 in a minimum of dilute nitric acid and then add distilled water so that each boiling-tube is approximately ½ full. Warm to dissolve if necessary. Record your observations in the space below.
Use these solutions for tests (d), (e) and (f).

[1]

(c)	Which anion can be identified from your observations in <b>(a)</b> and <b>(b)</b> ? Explain your answer.	For Examiner's
		Use
	[1]	
(d)	The cations present in <b>FA 3</b> , <b>FA 4</b> and <b>FA 5</b> can be identified by reaction of each solution with aqueous sodium hydroxide and with aqueous ammonia.  React 1 cm depth of each of the solutions prepared in <b>(b)</b> with each of these two reagents.	
	Record, in an appropriate form in the space below, your observations for these reactions.	
		i
		ii
		iii
		iv
		vi
	Conclusions	
	Using your observations you should be able to identify the cation present in two of the solutions. For the remaining solution you should be able to identify two possible cations.	
	FA 3 contains the cation(s)	
	FA 4 contains the cation(s)	
	<b>FA 5</b> contains the cation(s)	

[6]

		11 and 12 to select a as present in one of the	reagent to distinguish bene solutions in (d).	etween the
Carry out	the test with the sele	ected reagent.		
reagent				
observati	on			
conclusio	n			
(f) Carry out	the following tests.			[2]
test		observ	ations	
lesi	FA 3	FA	4 FA	5
To 1 cm depth of solution in a test-tube, add 1 cm de of aqueous barium nitrate,	epth			
add 2 cm depth o dilute nitric acid.	f			
To 1 cm depth of solution in a test-tube, add 1 cm de of aqueous silver nitrate,	epth			
allow any precipit formed to settle, pour off the soluti and add aqueous ammonia to the precipitate.	ion			
What con	clusions can be mad	de from the observation	ns above?	
				[2]

[Total: 14]

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# **Qualitative Analysis Notes**

*Key:* [ppt. = precipitate]

# 1 Reactions of aqueous cations

	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> (aq)	no ppt. 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. turning brown on contact with air insoluble in excess	green ppt. turning brown on contact with air 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. 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 <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 <sub>3</sub> <sup>2-</sup>	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, C <i>l</i> <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 $Ag^+(aq)$ (partially soluble in $NH_3(aq)$ ); gives white ppt. with $Pb^{2+}(aq)$
iodide, I <sup>-</sup> (aq)	gives yellow ppt. with $Ag^+(aq)$ (insoluble in $NH_3(aq)$ ); gives yellow ppt. with $Pb^{2+}(aq)$
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)	$NH_3$ 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 <sub>4</sub> <sup>2-</sup> (aq)	gives white ppt. with Ba <sup>2+</sup> (aq) or with Pb <sup>2+</sup> (aq) (insoluble in excess dilute strong acid)
sulfite, 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 acid)

## 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
sulfur dioxide, SO <sub>2</sub>	turns acidified aqueous potassium dichromate(VI) (aq) from orange to green

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