



Cambridge International Examinations

Cambridge International Advanced Subsidiary and Advanced Level

CANDIDATE NAME			
CENTRE NUMBER		CANDIDATE NUMBER	
CHEMISTRY			9701/33
Paper 3 Advanced Practical Skills 1		Oc	tober/November 2018
			2 hours
Candidates ans	wer on the Question Paper.		

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.

As listed in the Confidential Instructions

Write in dark blue or black pen.

You may use an HB pencil for any diagrams or graphs.

Do not use staples, paper clips, glue or correction fluid.

DO **NOT** WRITE IN ANY BARCODES.

Answer all questions.

Additional Materials:

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 14 and 15. A copy of the Periodic Table is printed on page 16.

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		
3		
Total		

This document consists of 14 printed pages and 2 blank pages.



Quantitative Analysis

Read through the whole method before starting any practical work. Where appropriate, prepare a table for your results in the space provided.

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

You will determine the enthalpy change, ΔH , for the reaction between magnesium and aqueous copper(II) sulfate. To do this you will measure the change in temperature when magnesium powder reacts with aqueous copper(II) sulfate. The magnesium is in excess.

$$Mg(s) + CuSO_4(aq) \rightarrow MgSO_4(aq) + Cu(s)$$

FA 1 is a solution containing 151.9 g dm⁻³ copper(II) sulfate, $CuSO_4$. **FA 2** is magnesium powder, Mg.

(a) Method

- Weigh the container with **FA 2** and record the mass in the space below.
- Support the plastic cup in the 250 cm³ beaker.
- Use the measuring cylinder to transfer 25 cm³ of **FA 1** into the plastic cup.
- Place the thermometer in the solution and tilt the cup if necessary so that the bulb of the thermometer is fully covered. Record the temperature at time zero in the table of results.
- Start timing and do not stop the clock until the whole experiment has been completed.
- Record the temperature of the solution every half minute for 2 minutes.
- At $2\frac{1}{2}$ minutes carefully transfer **all** of **FA 2** into the solution in the cup and stir the mixture.
- Record the temperature of the mixture at 3 minutes and complete the table by recording the temperature every half minute. Stir the mixture continuously between thermometer readings.
- Weigh the container with any residual FA 2 and record the mass below.
- Calculate and record the mass of FA 2 used.

Keep FA 1 for use in Question 2.

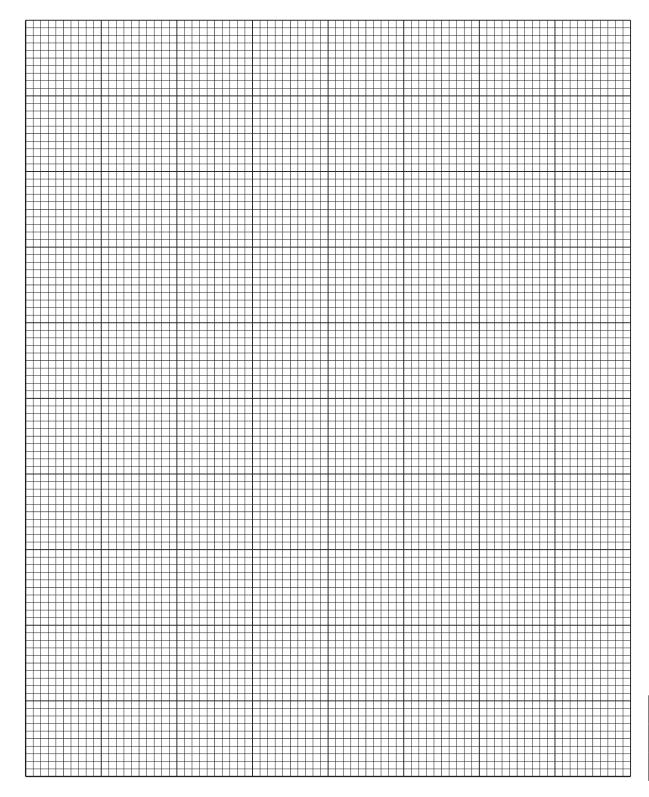
Results

time/minutes	0	1/2	1	1 ¹ / ₂	2	$2\frac{1}{2}$	3	3 ¹ / ₂	4
temperature/°C									
	Г	I	T	I			T	T	1
time/minutes	$4\frac{1}{2}$	5	5 ¹ / ₂	6	6 1 / ₂	7	$7\frac{1}{2}$	8	
temperature/°C									

(b) (i) Plot a graph of temperature on the *y*-axis against time on the *x*-axis on the grid below. The scale for temperature should extend at least 10 °C above your highest recorded temperature.

You will use the graph to determine the theoretical maximum temperature rise at $2\frac{1}{2}$ minutes.

Draw two lines of best fit through the points on your graph, the first for the temperature before adding **FA 2** and the second for the cooling of the mixture. Label any points you consider anomalous.



I III

	(11)	fit and so determine the theoretical rise in temperature, ΔT , at this time.
		$\Delta T = \dots ^{\circ}C$ [2]
(c)	Cal	culations
	(i)	Use the information given to calculate the concentration of copper(II) sulfate in $\textbf{FA 1}$ in $\text{mol}\text{dm}^{-3}.$
		concentration of FA 1 = mol dm ⁻³ [1]
	(ii)	Hence calculate the number of moles of $\operatorname{copper}(\Pi)$ sulfate you used in (a) .
		moles of CuSO ₄ = mol [1]
	(iii)	Use your answer to (b)(ii) to calculate the heat energy evolved when FA 2 is added to FA 1 . (Assume $4.2\mathrm{J}$ of heat energy changes the temperature of $1.0\mathrm{cm^3}$ of the mixture by $1.0\mathrm{^\circ C}$.)
		heat energy evolved = J [1]
	(iv)	Calculate the enthalpy change, in $kJ mol^{-1}$, when 1 mole of $CuSO_4$ reacts with magnesium.
		enthalpy change = kJ mol ⁻¹ [1] (sign) (value)

	(v)	Show by calculation that the magnesium was in excess.
		[1]
(d)	(i)	A student suggested that the calculated enthalpy change would be more accurate if magnesium turnings were used instead of magnesium powder.
		State whether you agree with the student and give a reason for your answer.
		[1]
	(ii)	The enthalpy change determined in (c)(iv) is only an approximation of the actual value.
	(,	
		Suggest and explain one improvement to the method in (a) you would make to increase the accuracy of the experiment.
		[1]
		[Total: 16]

In this experiment you will determine the enthalpy change, ΔH , for the reaction between aqueous copper(II) sulfate and zinc. The zinc is in excess. The method used in this question is slightly different from that used in **Question 1**.

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

FA 3 is zinc powder, Zn.

(a) Method

- Weigh a clean, dry plastic cup and record the mass.
- Add between 1.8 g and 2.0 g of zinc powder, **FA 3**, and record the mass.
- Place the plastic cup in the 250 cm³ beaker.
- Pour 25 cm³ of **FA 1** into the 25 cm³ measuring cylinder.
- Place the thermometer in the solution in the measuring cylinder and record the initial temperature.
- Pour the 25 cm³ of FA 1 into the plastic cup.
- Stir the contents of the cup and record the highest temperature of the mixture. Tilt the cup if necessary to ensure the thermometer bulb is fully covered.
- Calculate and record the mass of **FA 3** used and the change in temperature.

Keep the remainder of FA 3 for use in Question 3.

Results

[3]

(b)	C -		lati	
W	, vai	u	ıalı	ons

	(i)	Use your answer to $1(c)(ii)$ and the volume of FA 1 used in $2(a)$ to calculate the enthalpy change, in kJ mol ⁻¹ , when 1 mole of CuSO ₄ reacts with zinc. (Assume 4.2 J of heat energy changes the temperature of 1.0 cm ³ of the mixture by 1.0 °C.)
	(ii)	enthalpy change = kJ mol $^{-1}$ [2] (sign) (value) The maximum error in a single thermometer reading is $\pm 0.5^{\circ}$ C. Calculate the maximum percentage error in the temperature change for the experiment in (a).
(c)	solu	maximum percentage error = % [1] tudent decided to repeat this experiment using the same concentration of copper(II) sulfate ution and the same batch of zinc powder. However, 50 cm³ of FA 1 and double the mass of
	Wh The	at effect would doubling the quantities of FA 1 and FA 3 have on the temperature rise? e possible results are listed below. The temperature rise would be approximately half that for the first experiment. The temperature rise would be approximately the same as that for the first experiment. The temperature rise would be approximately double that for the first experiment.
		[1]

(d)	Use the information given in Question 1 and Question 2 and your answers to 1(c)(iv) and
	2(b)(i) to construct a Hess' cycle diagram for the reaction between magnesium and zinc sulfate
	to form magnesium sulfate and zinc.

Calculate the enthalpy change for this reaction.

(If you were unable to complete the calculations then assume the enthalpy change for 1(c)(iv) is $-320\,kJ\,\text{mol}^{-1}$ and the enthalpy change for 2(b)(i) is $-201\,kJ\,\text{mol}^{-1}$. These are not the correct values.)

enthalpy change =		$kJ mol^{-1}$	[2]
	(sign)	(value)	

[Total: 9]

Qualitative Analysis

Where reagents are selected for use in a test, the **full name** or **correct formula** of the element or compound must be given.

At each stage of any test you are to record details of the following:

- colour changes seen;
- the formation of any precipitate and its solubility in an excess of the reagent added;
- the formation of any gas and its identification by a suitable test.

You should indicate clearly at what stage in a test a change occurs.

If any solution is warmed, a **boiling tube** must be used.

Rinse and reuse test-tubes and boiling tubes where possible.

No additional tests for ions present should be attempted.

3 (a) FA 4 is a solution of a transition metal compound dissolved in acid.

Carry out the following tests on **FA 4** and record your observations.

test	observations			
To a 1 cm depth of FA 4 in a test-tube, add a 1 cm depth of potassium iodide, then				
add aqueous sodium thiosulfate.				
To a 3 cm depth of FA 4 in a boiling tube, add a spatula measure of zinc powder, FA 3 . If no reaction occurs, carefully warm the mixture. Leave the mixture for 5–15 minutes, observing the mixture occasionally.				
Keep the mixture of FA 4 and FA 3 for use in (c). The solution formed from the mixture is FA 6. Complete part (b) while FA 4 and FA 3 continue reacting				

[5]

(b) (i) FA 5 is the acid used to make FA 4. The Notes.	he anion in FA 5 is listed in the Qualitative Analysis
List the reagents you would use to ider testing for.	ntify this anion and state which acid the reagents are
	[2]
(ii) Carry out one of your tests in (i) and re	ecord your observations.
positively identify this anion.	nt in FA 5 . Write 'unknown' if your test could not
	[2]
(c) Carry out the following tests on FA 6 (the s	solution formed in (a)).
test	observations
To a 1 cm depth of FA 6 in a test-tube, add acidified potassium manganate(VII) a few drops at a time until in excess. At this stage the solution will be pink.	
To a 1 cm depth of FA 6 in a test-tube, add a 1 cm depth of FA 4 .	

(d) ((i)	State the type of reaction occurring in (c) when acidified potassium manganate(VII) was added to FA 6 . Give a reason for your answer.
		[2]
(i	ii)	The ion present in FA 4 when in acidic solution is usually written as $\mathbf{MO_2}^+$ where \mathbf{M} represents the transition metal.
		Calculate the oxidation state of \mathbf{M} in $\mathbf{MO_2}^+$.
		oxidation state of M =[1]
		[Total: 15]

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Qualitative Analysis Notes

1 Reactions of aqueous cations

ion	reaction with									
ion	NaOH(aq)	NH ₃ (aq)								
aluminium, A <i>l</i> ³+(aq)	white ppt. soluble in excess	white ppt. insoluble in excess								
ammonium, NH₄⁺(aq)	no ppt. ammonia produced on heating	_								
barium, Ba ²⁺ (aq)	faint white ppt. is nearly always observed unless reagents are pure	no ppt.								
calcium, Ca ²⁺ (aq)	white ppt. with high [Ca ²⁺ (aq)]	no ppt.								
chromium(III), Cr³+(aq)	grey-green ppt. soluble in excess	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), 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								
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								

2 Reactions of anions

ion	reaction
carbonate, CO ₃ ²⁻	CO ₂ liberated by dilute acids
chloride, C <i>l</i> ⁻ (aq)	gives white ppt. with Ag ⁺ (aq) (soluble in NH ₃ (aq))
bromide, Br ⁻ (aq)	gives cream ppt. with Ag ⁺ (aq) (partially soluble in NH ₃ (aq))
iodide, I-(aq)	gives yellow ppt. with Ag ⁺ (aq) (insoluble in NH ₃ (aq))
nitrate, NO ₃ ⁻ (aq)	NH₃ liberated on heating with OH⁻(aq) and A <i>l</i> foil
nitrite, NO ₂ -(aq)	NH ₃ liberated on heating with OH ⁻ (aq) and A <i>l</i> foil
sulfate, SO ₄ ²⁻ (aq)	gives white ppt. with Ba ²⁺ (aq) (insoluble in excess dilute strong acids)
sulfite, SO ₃ ²⁻ (aq)	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	urns 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					

The Periodic Table of Elements

	18	2	He	helium 4.0	10	Ne	neon 20.2	18	Ā	argon 39.9	36	궃	krypton 83.8	25	Xe	xenon 131.3	98	R	radon				
	17				6	ш	fluorine 19.0	17	Cl	chlorine 35.5	35	Ŗ	bromine 79.9	53	Н	iodine 126.9	85	Ą	astatine				
	16				8	0	oxygen 16.0	16	S	sulfur 32.1	34	Se	selenium 79.0	52	<u>e</u>	tellurium 127.6	84	Po	polonium -	116		livermorium —	
	15				7	z	nitrogen 14.0	15	۵	phosphorus 31.0	33	As	arsenic 74.9	51	Sp	antimony 121.8	83	ä	bismuth 209.0				
	14				9	ပ	carbon 12.0	14	S	silicon 28.1	32	Ge	germanium 72.6	50	Sn	tin 118.7	82	Ър	lead 207.2	114	Εl	flerovium —	
	13				5	В	boron 10.8	13	Ρl	aluminium 27.0	31	Ga	gallium 69.7	49	In	indium 114.8	81	lΤ	thallium 204.4				
										12	30	Zu	zinc 65.4	48	ၓ	cadmium 112.4	80	Ą	mercury 200.6	112	ပ်	copernicium -	
										7	29	Cn	copper 63.5	47	Ag	silver 107.9	62	Αn	gold 197.0	111	Rg	roentgenium -	
Group										10	28	Ż	nickel 58.7	46	Pd	palladium 106.4	78	₹	platinum 195.1	110	Ds	darmstadtium -	
Gre										0	27	රි	cobalt 58.9	45	몬	rhodium 102.9	77	٦	iridium 192.2	109	₹	meitnerium -	
		-	I	hydrogen 1.0						80	26	Ъе	iron 55.8	44	Ru	ruthenium 101.1	92	SO	osmium 190.2	108	H	hassium -	
										7	25	Mn	manganese 54.9	43	ည	technetium -	75	Re	rhenium 186.2	107	В	bohrium —	
							pol	ass			9	24	ပ်	chromium 52.0	42	Mo	molybdenum 95.9	74	≯	tungsten 183.8	106	Sg	seaborgium -
							Key	atomic number	atomic symbol	name relative atomic mass			2	23	>	vanadium 50.9	41	q	niobium 92.9	73	Та	tantalum 180.9	105
						ato	rek			4	22	F	titanium 47.9	40	Zr	zirconium 91.2	72	Ξ	hafnium 178.5	104	꿆	rutherfordium -	
										က	21	Sc	scandium 45.0	39	>	yttrium 88.9	57–71	lanthanoids		89-103	actinoids		
	2				4	Be	beryllium 9.0	12	Mg	magnesium 24.3	20	Ca	calcium 40.1	38	Š	strontium 87.6	56	Ba	barium 137.3	88	Ra	radium -	
	_				3	=	lithium 6.9	+	Na	sodium 23.0	19	¥	potassium 39.1	37	Rb	rubidium 85.5	55	S	caesium 132.9	87	ъ	francium —	

L U	175.0	103	۲	lawrencium	ı
ν Aγ	ytterblum 173.1	102	å	nobelium	ı
₆₉ T	mullum 168.9	101	Md	mendelevium	ı
₆₈ Г	erblum 167.3	100	Fm	ferminm	ı
67 Ho	nolmium 164.9	66	Es	einsteinium	1
°6 Dy	aysprosium 162.5	86	Ċ	californium	ı
65 Tb	158.9	26	益	berkelium	ı
² Od	gadolinium 157.3	96	CB	curium	ı
63 Eu	europium 152.0	92	Am	americium	ı
Sm	150.4	94	Pn	plutonium	ı
Pm	prometnium -	93	ď	neptunium	ı
09 P	neodymium 144.4	92	⊃	uranium	238.0
P.	praseodymium 140.9	91	Ра	protactinium	231.0
. Se	140.1	06	Т	thorium	232.0
57 La	138.9	88	Ac	actinium	ı

lanthanoids

actinoids

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