

Cambridge	
International	
AS & A Level	

Cambridge International Examinations

Cambridge International Advanced Subsidiary and Advanced Level

CHEMISTRY	ced Practical Skills 1		9701/35 May/June 2016
OUEMOTOV			0704/05
CENTRE NUMBER		CANDIDATE NUMBER	
CANDIDATE NAME			

2 hours

Candidates answer on the Question Paper.

As listed in the Confidential Instructions Additional Materials:

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 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.

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 13 printed pages and 3 blank pages.



1 In this experiment you will determine the concentration of a solution of sulfuric acid by titration.

FA 1 is sulfuric acid, H₂SO₄.

FA 2 is aqueous sodium hydroxide, containing 4.20 g NaOH dissolved in 1.00 dm³ of water. thymolphthalein indicator

(a) Method

Dilution of FA 1

- Pipette 10.0 cm³ of FA 1 into the 250 cm³ volumetric flask.
- Make the solution up to the mark using distilled water.
- Shake the flask thoroughly.
- This diluted solution of sulfuric acid is FA 3. Label the flask FA 3.

Titration

- Fill the burette with **FA 2**.
- Pipette **25.0 cm³** of **FA 3** into a conical flask.
- Add a few drops of thymolphthalein indicator.
- Perform a rough titration and record your burette readings in the space below. The end point is reached when the solution turns a permanent pale blue colour.

The rough titre is	cm ³
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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.

Keep solution FA 1 for use in Questions 2 and 3.

I	
II	
III	
IV	
V	
VI	
VII	

[7]

(b)	in y	m your accurate titration results, obtain a suitable value for the volume of FA 2 to be used our calculations. ow clearly how you obtained this value.
		25.0 cm ³ of FA 3 required cm ³ of FA 2 . [1]
(c)	Cal	culations
		ow your working and appropriate significant figures in the final answer to each step of your culations.
	(i)	Calculate the number of moles of sodium hydroxide present in the volume of FA 2 calculated in (b)
		in (b) . Use the data in the Periodic Table on page 16.
		moles of NaOH = mol
	(ii)	Complete the equation for the reaction of sulfuric acid with sodium hydroxide. State symbols are required.
		$+$ \rightarrow $Na_2SO_4(aq) +$
((iii)	Use your answers to (i) and (ii) to calculate the number of moles of sulfuric acid used in each titration.
		moles of H ₂ SO ₄ = mol
((iv)	Calculate the concentration, in mol dm ⁻³ , of sulfuric acid in FA 3 .
		concentration of H_2SO_4 in FA 3 = mol dm ⁻³
	(v)	Calculate the concentration, in mol dm ⁻³ , of sulfuric acid in FA 1 .
	-	
		concentration of H_2SO_4 in FA 1 = mol dm ⁻³ [5]
		[Total: 13]

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2 In this experiment you will determine the enthalpy change, ΔH , for the decomposition of magnesium carbonate to magnesium oxide.

$$MgCO_3(s) \rightarrow MgO(s) + CO_2(g)$$

In order to do this, you will determine the enthalpy changes for the reactions of magnesium carbonate and magnesium oxide with sulfuric acid. Excess of the two magnesium compounds will be used in each experiment.

Then you will use Hess' Law to calculate the enthalpy change for the reaction above.

FA 1 is sulfuric acid, H₂SO₄.

FA 4 is magnesium carbonate, MgCO₃.

FA 5 is magnesium oxide, MgO.

(a) Determination of the enthalpy change for the reaction of magnesium carbonate, FA 4, with sulfuric acid, FA 1

(i) Method

- Support the plastic cup inside the 250 cm³ beaker.
- Use a measuring cylinder to transfer 25 cm³ of **FA 1** into the plastic cup.
- Measure and record the initial temperature of the **FA 1** in the space below.
- Add all the FA 4 from the container to the FA 1 in the plastic cup.
- Stir constantly until the maximum temperature is reached.
- Measure and record the maximum temperature of the contents of the cup.
- Rinse out the plastic cup and shake to dry for use in (b).
- Calculate and record the temperature rise.

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

(ii) Calculate the energy produced during this reaction.
[Assume that 4.2 J are needed to raise the temperature of 1.0 cm³ of solution by 1.0 °C.]

energy produced = J

(iii) Use your answer to 1(c)(v) to calculate the number of moles of sulfuric acid in 25 cm³ of FA 1.

(If you were unable to calculate the concentration of sulfuric acid in **FA 1**, assume that it is $1.27 \, \text{mol dm}^{-3}$. This is not the true value.)

moles of $H_2SO_4 = \dots mol$

(iv) Calculate the enthalpy change, in kJmol⁻¹, for the reaction below.

$$\label{eq:mgco3} MgCO_3(s) \ + \ H_2SO_4(aq) \ \rightarrow \ MgSO_4(aq) \ + \ CO_2(g) \ + \ H_2O(l)$$

enthalpy change = $kJ \, mol^{-1}$ (sign) (value)

[6]

(b) Determination of the enthalpy change for the reaction of magnesium oxide, FA 5, with sulfuric acid, FA 1

(i) Method

- Use the measuring cylinder to transfer approximately 40 cm³ of **FA 1** into the **100 cm³** beaker.
- Place the beaker on a tripod and gauze.
- Heat FA 1 in the beaker until the temperature is between 40 °C and 50 °C.
- Support the plastic cup in the **250 cm**³ beaker.
- Use the measuring cylinder to transfer 25 cm³ of hot FA 1 into the plastic cup. CARE.
- Measure and record, in the space below, the initial temperature of FA 1 in the plastic cup.
- Immediately, add all the **FA 5** from the container to the **FA 1** in the plastic cup.
- Stir constantly until the maximum temperature is reached.
- Measure and record the maximum temperature.
- Calculate and record the temperature rise.

Calculations

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

(ii) Calculate the energy produced during this reaction. [Assume that 4.2J are needed to raise the temperature of 1.0 cm³ of solution by 1.0 °C.]

(iii) Use your answer to (a)(iii) to calculate the enthalpy change, in kJ mol⁻¹, for the reaction below.

$$MgO(s) + H_2SO_4(aq) \rightarrow MgSO_4(aq) + H_2O(l)$$

enthalpy change = $kJ \, mol^{-1}$ (sign) (value)

[4]

	7
(c)	Use your values for the enthalpy changes calculated in (a)(iv) and (b)(iii) to calculate the enthalpy change for the reaction below.
	Show clearly how you obtained your answer by drawing a Hess' Law energy cycle.
	(If you were unable to calculate the enthalpy changes, assume that the value of the enthalpy change in (a)(iv) is -58.7 kJ mol ⁻¹ and the value in (b)(iii) is -140.3 kJ mol ⁻¹ . Note: these are not the correct values.)

 $\mathsf{MgCO_3}(\mathsf{s}) \ \to \ \mathsf{MgO}(\mathsf{s}) \ + \ \mathsf{CO_2}(\mathsf{g})$

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

(d) (i) Calculate the maximum percentage error in the temperature rise in (b)(i).

	percentage error = %
(ii)	The magnesium oxide, FA 5 , was weighed with a balance measuring to one decimal place A student suggested that the accuracy of the experiment in (b)(i) would be improved by weighing FA 5 using a balance measuring to two decimal places. State and explain whether or not the student is correct.

[Total: 14]

[2]

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)	(i)	FA 6 is a salt containing one cation and one anion from those listed on pages 14 and 15 Transfer a small spatula measure of FA 6 into a hard-glass test-tube. Heat gently at first, then heat strongly until no further change occurs.
		Record all your observations below.

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(ii) Dissolve the remainder of **FA 6** in an approximate depth of 5 cm of distilled water in a boiling tube for use in the following tests. Record your observations in the table below.

observations
A 6 with cold sodium hydroxide.

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(b) (i) FA 7 is a solution containing one cation and one anion from the list on pages 14 and 15.Carry out the following tests and record your observations in the table below.

	test	observations								
•	To a 1 cm depth of FA 7 in a test-tube, add aqueous sodium hydroxide.									
-	To a 1 cm depth of FA 7 in a test-tube, add aqueous ammonia.									
<u>-</u>	To a 1 cm depth of FA 7 in a test-tube, add a few drops of acidified potassium manganate(VII), followed by a few drops of aqueous starch.									
(i	i) Identify FA 7.									
	FA 7 is									
(ii	i) Carry out one further test of your cho	Carry out one further test of your choice to confirm the identity of the anion in FA 7 .								
	reagent(s) used									
	observation(s)									

[Total: 13]

[5]

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Qualitative Analysis Notes

Key: [ppt. = precipitate]

1 Reactions of aqueous cations

	reaction with									
ion	NaOH(aq)	NH ₃ (aq)								
aluminium, Al³+(aq)	white ppt. soluble in excess	white ppt. insoluble in excess								
ammonium, NH ₄ +(aq)	no ppt. ammonia produced on heating	_								
barium, Ba²+(aq)	faint white ppt. is nearly always observed unless reagents are pure	no ppt.								
calcium, Ca ²⁺ (aq)	white ppt. with high [Ca2+(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), 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, 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, 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)	NH ₃ 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 Al 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) (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	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

The Periodic Table of Elements

	18	2	e L	helium 4.0	10	Ne	neon 20.2	18	Ā	argon 39.9	38	궃	krypton 83.8	25 25	Xe	xenon 131.3	98	Ru	radon			
	17				6	ш	fluorine 19.0	17	Cl	chlorine 35.5	35	ğ	bromine 79.9	53	Ι	iodine 126.9	85	At	astatine -			
	16				80	0	oxygen 16.0	16	ഗ	sulfur 32.1	34	Se	selenium 79.0	52	<u>e</u>	tellurium 127.6	84	Ъ	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	Ξ	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	Pp	lead 207.2	114	Εl	flerovium —
	13				2	В	boron 10.8	13	Αl	aluminium 27.0	31	Ga	gallium 69.7	49	In	indium 114.8	18	11	thallium 204.4			
										12	30	Zu	zinc 65.4	48	g	cadmium 112.4	80	Нg	mercury 200.6	112	ပ်	copernicium —
										1	29	J.	copper 63.5	47	Ag	silver 107.9	79	Αn	gold 197.0	111	Rg	roentgenium -
dn										10	28	z	nickel 58.7	46	Pd	palladium 106.4	78	₫	platinum 195.1	110	Ds	darmstadtium -
Group										6	27	ပိ	cobalt 58 9	45	R	rhodium 102.9	77	Ϊ́	iridium 192.2	109	¥	meitnerium -
		-]	I	hydrogen 1.0						80	26	Pe	iron 8 77	44	Ru	ruthenium 101.1	9/	SO	osmium 190.2	108	£	hassium -
					ı					7	25	Mn	manganese	43	ပ	technetium -	75	Re	rhenium 186.2	107	B	bohrium —
						loc	SS			9	24	ပ်	chromium 52 0	42	Mo	molybdenum 95.9	74	≥	tungsten 183.8	106	Sg	seaborgium -
				Key	atomic number	atomic symbo	name relative atomic mass			2	23	>	vanadium 50 Q	41	g	niobium 92.9	73	<u>n</u>	tantalum 180.9	105	<u>а</u>	dubnium —
					10	ato	rela			4	22	F	titanium 47 o	40	Zr	zirconium 91.2	72	Ξ	hafnium 178.5	104	쪼	rutherfordium -
										က	21	လွ	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	88	Š	strontium 87.6	56	Ba	barium 137.3	88	Ra	radium _
	-				3	:-	lithium 6.9	#	Na	sodium 23.0	19	×	potassium	37	Rb	rubidium 85.5	55	S	caesium 132.9	87	ь	francium -

71	P	lutetium 175.0	103	۲	lawrencium	ı	
		ytterbium 173.1				ı	
69	T	thulium 168.9	101	Md	mendelevium	ı	
89	Ē	erbium 167.3	100	Fm	ferminm	I	
29	웃	holmium 164.9	66	Es	einsteinium	I	
99	۵	dysprosium 162.5	86	Ç	californium	ı	
65	Д	terbium 158.9	26	益	berkelium	I	
64	gq	gadolinium 157.3	96	CB	curium	I	
63	En	europium 152.0	98	Am	americium	I	
62	Sm	samarium 150.4	94	Pu	plutonium	ı	
19	Pm	promethium —	93	ď	neptunium	ı	
09	PZ	neodymium 144.4	92	\supset	uranium	238.0	
59	Ą	praseodymium 140.9	91	Ра	protactinium	231.0	
58	Ce	cerium 140.1	06	Ч	thorium	232.0	
22	Га	lanthanum 138.9	88	Ac	actinium	ı	

lanthanoids

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