Cambridge
International
AS & A Level

Cambridge Assessment International Education Cambridge International Advanced Subsidiary and Advanced Level

	CANDIDATE NAME	
	CENTRE NUMBER CANE NUMB	DIDATE BER
* 5 9	CHEMISTRY	9701/33
8 2	Paper 3 Advanced Practical Skills 1	May/June 2019
5 2		2 hours
° *	Candidates answer on the Question Paper.	
5 9	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 h Give details of the practical session and laboratory where appropriate, in the 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 app Use of a Data Booklet is unnecessary.	ropriate units.
	·	Session
	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.	Laboratory
	The number of marks is given in brackets [] at the end of each question or part question.	

For Examiner's Use							
1							
2							
3							
Total							

This document consists of **12** printed pages, **4** blank pages and **1** Insert.

2

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.

1 The thiosulfate ion, $S_2O_3^{2-}$, reacts in acidic conditions as shown.

$$S_2O_3^{2-}(aq) + 2H^+(aq) \rightarrow S(s) + SO_2(g) + H_2O(I)$$

You will investigate how the concentration of the thiosulfate ions affects the rate of this reaction. The rate can be measured by timing how long it takes for the solid sulfur that is formed to make the solution too cloudy to see through.

Small amounts of SO_2 gas may be produced during this reaction. Care must be taken to avoid inhaling this SO_2 gas.

It is very important that as soon as each experiment is complete the beaker containing the reaction mixture is emptied into the quenching bath.

FA 1 is 0.100 mol dm⁻³ sodium thiosulfate, $Na_2S_2O_3$. FA 2 is 2.00 mol dm⁻³ hydrochloric acid, HC*l*. distilled water

(a) Method

Experiment 1

- Fill the burette labelled **FA 1** with **FA 1**.
- Run 45.00 cm³ of **FA 1** from the burette into the 100 cm³ beaker.
- Use the measuring cylinder to measure 10.0 cm³ of **FA 2**.
- Add the FA 2 to the FA 1 in the beaker and start timing immediately.
- Stir the mixture once and place the beaker on the printed insert.
- Look down through the solution in the beaker at the print on the insert.
- Stop timing as soon as the precipitate of sulfur makes the print on the insert just invisible.
- Record this reaction time to the nearest second in your results table.
- Empty the contents of the beaker into the quenching bath.
- Wash out the beaker thoroughly.
- Shake the beaker to remove any excess water.

Experiment 2

- Fill a second burette with distilled water.
- Refill the burette labelled **FA 1** with **FA 1**.
- Run 20.00 cm³ of **FA 1** into the 100 cm³ beaker.
- Run 25.00 cm³ of distilled water into the same beaker.
- Use the measuring cylinder to measure 10.0 cm³ of **FA 2**.
- Add the FA 2 to the FA 1 in the beaker and start timing immediately.
- Stir the mixture once and place the beaker on the printed insert.
- Look down through the solution in the beaker at the print on the insert.
- Stop timing as soon as the precipitate of sulfur makes the print on the insert **just** invisible.
- Record this reaction time to the nearest second in your results table.
- Empty the contents of the beaker into the quenching bath.
- Wash out the beaker thoroughly.
- Shake the beaker to remove any excess water.

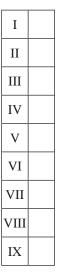
Experiments 3–5

Carry out three further experiments to investigate how the reaction time changes with different volumes of **FA 1**.

Note that the combined volume of **FA 1** and distilled water must always be 45.00 cm^3 . Do not use a volume of **FA 1** that is less than 20.00 cm^3 .

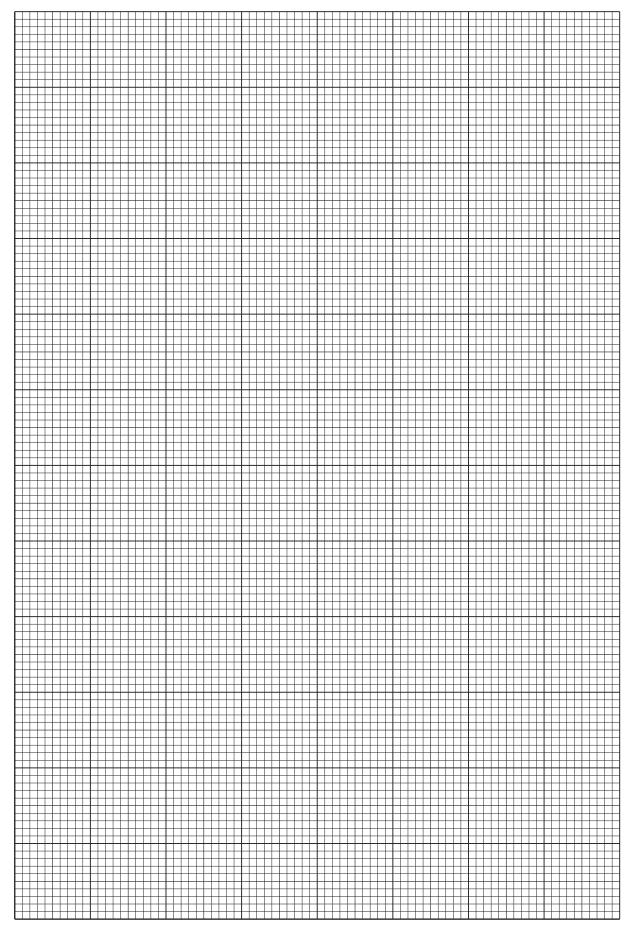
Record all your results in a table. You should include the volume of **FA 1**, the volume of distilled water, the reaction time and the reaction rate for each of your five experiments. The rate of reaction can be calculated using the following expression.

rate = $\frac{500}{\text{reaction time}}$

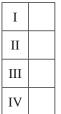


[9]

(b) On the grid, plot a graph of the rate (*y*-axis) against the volume of **FA 1** (*x*-axis). Label any anomalous points. Draw a line of best fit.



5



[4]

(c) In these experiments, the volume of **FA 1** is related to the concentration of the thiosulfate ions. From your graph state the relationship between the rate of reaction and the concentration of the thiosulfate ions.

.....[1]

(d) Assume that the error in the time measured for each experiment was $\pm 2s$.

Calculate the minimum value for the reaction rate you observed in **Experiment 2**. Show your working.

(e) (i) A student suggested that, using a 250 cm³ beaker, the time recorded for **Experiment 1** would be the same.

Discuss whether the student is correct.

(ii) A student carried out a further experiment using the same procedure as in (a). The student used 5.00 cm³ of **FA 1**, 40.00 cm³ of distilled water and 10.0 cm³ of **FA 2**. The print on the insert never became invisible. Explain why.

[Total: 18]

2 In this experiment you will determine the enthalpy change of solution for hydrated sodium thiosulfate.

FA 3 is hydrated sodium thiosulfate, $Na_2S_2O_3.5H_2O$.

(a) Method

- Support the plastic cup in the 250 cm³ beaker.
- Rinse the measuring cylinder.
- Using the measuring cylinder, pour 25.0 cm³ of distilled water into the plastic cup.
- Measure the temperature of the water in the cup.
- Weigh the container with FA 3.
- Add all the **FA 3** to the distilled water in the cup.
- Use the thermometer to stir the mixture gently until all the solid has dissolved.
- Measure the lowest temperature that is reached.
- Reweigh the container with any remaining FA 3.
- Record all your measurements.
- Calculate and record the mass of **FA 3** added and the change in temperature.

Ι	
II	
III	
IV	

[4]

- (b) The enthalpy change of solution for **FA 3** is the enthalpy change when 1 mole of **FA 3** is dissolved in 1 dm³ of solution.
 - (i) Calculate how many moles of **FA 3** were added to the water.

moles of **FA 3** = mol [1]

(ii) Calculate the energy change when the sample of **FA 3** was added to the distilled water. [Assume that 4.2J of heat energy changes the temperature of 1.0 cm^3 of solution by 1.0 °C.]

	energy change = .		J	[1]
Calculate the enthalpy change of solution of F	A 3.			
enthalpy change of	solution =			
	sign	value	units	[1]

(c) One way to improve this experiment would be to use a balance that reads to more decimal places.

Suggest two other ways in which this experiment could be altered to give a more accurate value for the enthalpy change. Explain how each would improve the accuracy.

Suggestion 1 Suggestion 2 [2]

(d) A student carrying out the experiment in Question 1 used all the FA 1. The student made up a fresh sample of FA 1 of the correct concentration by dissolving some FA 3 in water. This solution was then used immediately to repeat one of the experiments in Question 1 but the time was then much greater than had been measured previously.

Explain why the time was greater.

[1] [Total: 10]

(iii)

Qualitative Analysis

Where reagents are selected for use in a test, the **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 an aqueous solution containing a single cation and a single anion. The anion is either the sulfate ion, SO_4^{2-} , or the sulfite ion, SO_3^{2-} .
 - (i) To an approximately 1 cm depth of **FA 4** in a test-tube, add aqueous sodium carbonate. Record your observations.

		[2]
(ii)	Select reagents to identify the anion present in FA 4 . Carry out a test with these reagents and record your observations.	
	reagents	
	observations	
		[2]
(iii)	Identify FA 4.	
	The formula of FA 4 is	[1]

(b) (i) FA 5 contains one cation and two anions. Two of these ions are listed in the Qualitative Analysis Notes.

Carry out the following tests and record your observations.

test	observations
Add a small spatula measure of FA 5 to a hard-glass test-tube.	
Heat the sample gently at first and then more strongly.	
Pour a 4 cm depth of dilute sulfuric acid into a boiling tube. Carefully add the remaining FA 5 . Leave to stand until the reaction is complete. The solution produced is FA 6 .	
Keep FA 6 for use in the following tests.	
To a 1 cm depth of FA 6 in a test-tube add aqueous sodium hydroxide.	
To a 1 cm depth of FA 6 in a test-tube add aqueous ammonia.	
	[5]

(ii)	State the type of reaction observed when	FA 5 was heated.	
			[1]
(iii)	Give the formula of the cation and one of	he anions present in FA 5 .	
	cation:	anion:	[1]
		[Total:	12]

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12

13

Qualitative Analysis Notes

1 Reactions of aqueous cations

in a	reac	tion 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)	faint white ppt. is nearly always observed unless 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	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							
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, NO₃⁻(aq)	NH_3 liberated on heating with $OH^-(aq)$ and Al foil
nitrite, NO₂⁻(aq)	NH_3 liberated on heating with $OH^-(aq)$ and Al 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	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	Group	13 14 15 16 17 18			hydrogen 1.0		B C N	boron carbon nitrogen oxygen fluorine neon 10.8 12.0 14.0 16.0 19.0 20.2	14 15 16	Si P S C <i>l</i>	8 9 10 11 12 aluminium silicon phosphous sulfur o 22.0 28.1 31.0 32.1	25 26 27 28 29 30 31 32 33 34 35	Cr Mn Fe Co Ni Cu Zn Ga Ge As Se Br	n dromium manganese iron cobalt nickel copper zinc gallium germanium arsenic selenium bromine krypton 52.0 54.9 55.8 58.9 58.7 63.5 65.4 69.7 72.6 74.9 79.0 79.9 83.8	42 43 44 45 46 47 48 49 50 51 52 53	Mo Tc Ru Rh Pd Ag Cd In Sn Sb Te I	m molydehum technetium ruthenium thodium palladium silver cadmium indium tin antimony tellurium iodine xenon 95.9 – 101.1 102.9 106.4 107.9 112.4 114.8 118.7 121.8 127.6 126.9 131.3	75 76 77 78 79 80 81 82 83 84 85	Re Os Ir Pt Au Hg T <i>I</i> Pb Bi Po At	tungsten rhenium osmium iridium platinum 183.8 186.2 190.2 192.2 195.1	106 107 108 109 110 111 112 114	Sg Bh Hs Mt Ds Rg Cn	seaborgium bohrium hassium meitnerium darmstadtium – – – – – – – –		60 61 62 63 64 65 66 67 68 69 70	Nd Pm Sm Eu Gd Tb Dy Ho Er Tm Yb	ium neodymium promethium samarium europium gadolinium terbium dysprosium holmium erbium truikum yrterbium lutetium 144.4 – 150.4 152.0 157.3 158.9 162.5 164.9 167.3 168.9 173.1 175.0	92 93 94 95 96 97 98 99 100	U Np Pu Am Cm Bk Cf Es Fm Md No	uranium neptunium plutonium americium curium berkelium californium einsteinium fermium mendelevium r			
The Periodic Tab	Grou				~ =	E	nydrogen 1.0						80	26	Fe	iron 55.8	44	Ru	ruthenium 101.1	76	Os	osmium 190.2	108	Hs	hassium meitnerium –	-	2.9	Sm	samarium 150.4	94	Pu	plutonium	
						Key	atomic number	atomic symbol	name relative atomic mass				24	ŗ		42	Mo		74	8		106		_	:	60	Nd	praseodymium neodymium promethiu 140.9 144.4 –	92	⊃			
									÷	atom	atomic	relative			3 4		F	scandium titanium va 45.0 47.9	40	Zr	yttrium zirconium n 88.9 91.2	72	Hf	hafnium ta 178.5	104	actinoids Rf	rutherfordium du	_	28	0e	lanthanum cerium prase 138.9 140.1 1	06	Th
		1 2				3 4	Li Be	lithium beryllium 6.9 9.0	11 12		sodium magnesium 23.0 24.3			potassium calcium 39.1 40.1			rubidium strontium 85.5 87.6	55 56	Cs Ba	caesium barium 132.9 137.3	87 88	Fr Ra	francium radium -	L		lanthanoids			actinoids				

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