

Cambridge International AS & A Level

	CANDIDATE NAME CENTRE NUMBER	IDATE ER	
×	CHEMISTRY		9701/35
	Paper 3 Advanced Practical Skills 1	Ν	lay/June 2024
6	•		2 hours
2690705*	You must answer on the question paper. You will need: The materials and apparatus listed in the confidential instruct	ons	
	 Answer all questions. Use a black or dark blue pen. You may use an HB pencil for any diagram Write your name, centre number and candidate number in the boxes at the Write your answer to each question in the space provided. Do not use an erasable pen or correction fluid. Do not write on any bar codes. You may use a calculator. You should show all your working and use appropriate units. 		e.
	INFORMATION	Ses	sion
	The total mark for this paper is 40.The number of marks for each question or part question is shown in		
	 brackets []. The Periodic Table is printed in the question paper. 	Labo	ratory
	 Important values, constants and standards are printed in the 		
	question paper.Notes for use in qualitative analysis are provided in the		
	question paper.	For Exam	iner's Use
		1	
		2	
		3	
		L	

Total

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 the precision of the apparatus you used in the data you record.

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

1 lodide ions in aqueous solution are oxidised to iodine by a variety of oxidising agents. One of these is the peroxodisulfate ion, $S_2O_8^{2-}$, which reacts as shown.

$$2I^{-}(aq) + S_2O_8^{2^-}(aq) \rightarrow I_2(aq) + 2SO_4^{2^-}(aq)$$

Sodium thiosulfate is added to the reaction mixture to react with iodine as it is produced. When all of the thiosulfate has reacted, further iodine produced reacts with starch indicator to give a dark colour.

You will carry out two experiments to investigate how the rate of this reaction is affected by changing the concentration of the peroxodisulfate ion.

FA 1 is $0.0200 \text{ mol dm}^{-3}$ potassium peroxodisulfate, $K_2S_2O_8$.

FA 2 is $0.00500 \text{ mol dm}^{-3}$ sodium thiosulfate, $Na_2S_2O_3$.

FA 3 is 1.00 mol dm^{-3} potassium iodide, KI.

FA 4 is starch indicator.

(a) Method

Experiment 1

- Label one of the 100 cm³ beakers **A** and the other 100 cm³ beaker **B**.
- Fill one burette with FA 1. Label this burette FA 1.
- Run 20.00 cm³ of **FA 1** from the burette into beaker **A**.
- Fill the second burette with FA 2. Label this burette FA 2.
- Run $10.00 \,\mathrm{cm^3}$ of **FA 2** from the burette into beaker **B**.
- Use the measuring cylinder to add 20.0 cm³ of **FA 3** to beaker **B**.
- Add 10 drops of **FA 4** to beaker **B**.
- Add the contents of beaker **A** to beaker **B** and start timing **immediately**.
- Stir the mixture once and place the beaker on the white tile.
- Stop timing as soon as the solution turns a dark colour.
- Record this time to the nearest second in the space for results.
- Wash out both beakers and dry them using paper towel.

Experiment 2

- Run 10.00 cm³ of **FA 1** from the burette into beaker **A**.
- Run 10.00 cm³ of **FA 2** from the burette into beaker **B**.
- Use the measuring cylinder to add 20.0 cm³ of **FA 3** into beaker **B**.
- Use the same measuring cylinder to add 10.0 cm³ of distilled water to beaker B.
- Add 10 drops of **FA 4** to beaker **B**.
- Add the contents of beaker **A** to beaker **B** and start timing **immediately**.
- Stir the mixture once and place the beaker on the white tile.
- Stop timing as soon as the solution turns a dark colour.
- Record this time to the nearest second.

Record all your data in a table. You should include the volume of **FA 1**, the volume of distilled water, the reaction time and the rate of reaction for both experiments.

Use the following formula to calculate the rate of reaction.

rate = $\frac{1000}{\text{reaction time}}$

Results

Ι	
II	
III	
IV	
V	
VI	

[6]

(b) (i) Explain why the concentration of potassium peroxodisulfate used in each experiment is proportional to the volume of **FA 1** used.

......[1]

(ii) A student thinks that the rate of reaction is proportional to the concentration of FA 1. Complete Table 1.1 to suggest volumes of reactants that could be used in a further experiment to confirm whether the student is correct. Do not carry out this experiment.

	volum	ie/cm ³		volume
FA 1	FA 2	FA 3	distilled water	FA 4
				10 drops

[2]

(c) A student correctly carried out the method in (a) but had been given a more concentrated solution of sodium thiosulfate.
 State how you would expect the student's times to differ from yours. Explain your answer.

State how you would expect the student's times to differ from yours. Explain your answer.

......[1]

(d) The potassium iodide is in a large excess in Experiments 1 and 2. Suggest why a large excess of iodide ions is needed in these experiments.

......[1]

[Total: 11] [Turn over **2** You will carry out an experiment to determine the enthalpy change, ΔH , when one mole of ammonium chloride dissolves in water.

FA 5 is ammonium chloride, NH_4Cl .

- (a) Method
 - Weigh the container with **FA 5**. Record the mass in the space for results.
 - Support the cup in the 250 cm³ beaker.
 - Use the measuring cylinder to transfer 25.0 cm³ of distilled water into the cup.
 - Place the thermometer in the water and tilt the cup, if necessary, so that the bulb of the thermometer is fully covered. Record the temperature of the water at time *t* = 0.
 - Start the stop-clock and leave it running for the whole experiment.
 - Measure and record the temperature of the water in the cup every half minute for 2 minutes.
 - At $t = 2\frac{1}{2}$ minutes, tip all the **FA 5** into the cup. Stir the contents of the cup.
 - Measure and record the temperature of the contents of the cup at t = 3 minutes and then every half minute up to and including t = 8 minutes.
 - Weigh the container with any residual **FA 5**. Record the mass.
 - Calculate and record the mass of **FA 5** added.

Results

Ι	
II	
III	
IV	
V	

[5]

(b) Plot a graph of temperature (y-axis) against time (x-axis) on the grid. You should choose a scale that allows you to plot 2 °C below the minimum temperature reached. Label any points you consider to be anomalous.

Draw **two** straight lines of best fit. One line is for the temperature before adding **FA 5** and the other line is for the warming of the solution once the minimum temperature has been reached.

Extrapolate both these lines to $t = 2\frac{1}{2}$ minutes.



5



[4]

(c) (i) Use your graph to determine the temperature change, ΔT , at $t = 2\frac{1}{2}$ minutes.

 ΔT at $t = 2\frac{1}{2}$ minutes =°C [1]

(ii) Calculate the energy change, in J, in the reaction.

energy change = J [1]

(iii) Calculate the amount, in mol, of ammonium chloride used.

amount of NH_4Cl = mol [1]

(iv) Calculate the enthalpy change, ΔH , in kJ mol⁻¹, when one mole of ammonium chloride dissolves in water.

 $\Delta H = \dots kJ \operatorname{mol}^{-1} [1]$

(d) Use your results in (a) to calculate the maximum percentage error for the temperature change from 0 to 4 minutes.

Assume that the maximum uncertainty in a single thermometer reading is ± 0.5 °C.

Show your working.

maximum percentage error = % [1]

[Total: 14]

Qualitative analysis

For each test you should record all your observations in the spaces provided.

Examples of observations include:

- colour changes seen
- the formation of any precipitate and its solubility (where appropriate) in an excess of the reagent added

7

• the formation of any gas and its identification (where appropriate) by a suitable test.

You should record clearly at what stage in a test an observation is made.

Where no change is observed, you should write 'no change'.

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

If any solution is warmed, a boiling tube must be used. If a solid is heated, a hard-glass test-tube must be used.

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

No additional tests should be attempted.

- 3 (a) FA 6 is a salt containing a Group 1 ion and an anion that consists of a transition metal and a non-metal element.
 - (i) Transfer FA 6 into a hard-glass test-tube. Heat the tube gently at first and then strongly. Record all your observations and identify the gas produced. The residue is FA 7. You will use FA 7 in (a)(ii).

observations

.....

gas produced

[3]

Allow FA 7 to cool before starting (a)(ii). While FA 7 is cooling you may wish to continue with (b)(i).

(ii) Put a 2cm depth of acidified aqueous potassium manganate(VII) in a test-tube. Add the same depth of aqueous sodium hydroxide. Then add FA 7 and stir using the glass rod for about 30 seconds. Filter the mixture and collect the filtrate. Record your observations.

.....

Put a 1 cm depth of the filtrate in a test-tube. Add sulfuric acid until in excess. Record your observations.

.....

[2]

(b) (i) FA 8 and FA 9 are both aqueous solutions of salts. FA 8 contains one cation and one anion. FA 9 contains two cations and one anion. One of the cations and both anions are listed in the Qualitative analysis notes.

Carry out the following tests and record your observations in Table 3.1. For each test use a 1 cm depth of **FA 8** or **FA 9** in a test-tube.

test	observ	vations
lesi	FA 8	FA 9
Test 1		
Add sulfuric acid.		
Test 2		
Add aqueous sodium hydroxide, then		
transfer the mixture into a boiling tube and warm.		
Test 3		
Add a few drops of aqueous barium chloride or aqueous barium nitrate, then		
add nitric acid.		
Test 4		
Add FA 8 with shaking until in excess.		

Table 3.1

Qualitative analysis notes

1 Reactions of cations

cation	reaction	on with
	NaOH(aq)	NH ₃ (aq)
aluminium, Al ³⁺ (aq)	white ppt. soluble in excess	white ppt. insoluble in excess
ammonium, NH ₄ +(aq)	no ppt. ammonia produced on warming	_
barium, Ba ²⁺ (aq)	faint white ppt. is observed unless [Ba ²⁺ (aq)] is very low	no ppt.
calcium, Ca ²⁺ (aq)	white ppt. unless [Ca ²⁺ (aq)] is very low	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	pale 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
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

2 Reactions of anions

anion	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/off-white ppt. with Ag ⁺ (aq) (partially soluble in NH ₃ (aq))
iodide, I [_] (aq)	gives pale 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 A <i>l</i> foil; decolourises acidified aqueous KMnO ₄
sulfate, SO ₄ ^{2–} (aq)	gives white ppt. with $Ba^{2+}(aq)$ (insoluble in excess dilute strong acids); gives white ppt. with high $[Ca^{2+}(aq)]$
sulfite, SO ₃ ^{2–} (aq)	gives white ppt. with Ba ²⁺ (aq) (soluble in excess dilute strong acids); decolourises acidified aqueous KMnO ₄
thiosulfate, S ₂ O ₃ ^{2–} (aq)	gives off-white/pale yellow ppt. slowly with H ⁺

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
hydrogen, H ₂	'pops' with a lighted splint
oxygen, O ₂	relights a glowing splint

4 Tests for elements

element	test and test result
iodine, I ₂	gives blue-black colour on addition of starch solution

Important values, constants and standards

molar gas constant	$R = 8.31 \mathrm{J}\mathrm{K}^{-1}\mathrm{mol}^{-1}$
Faraday constant	$F = 9.65 \times 10^4 \mathrm{C mol^{-1}}$
Avogadro constant	$L = 6.022 \times 10^{23} \text{mol}^{-1}$
electronic charge	$e = -1.60 \times 10^{-19} \mathrm{C}$
molar volume of gas	$V_{\rm m}$ = 22.4 dm ³ mol ⁻¹ at s.t.p. (101 kPa and 273 K) $V_{\rm m}$ = 24.0 dm ³ mol ⁻¹ at room conditions
ionic product of water	$K_{\rm w}$ = 1.00 × 10 ⁻¹⁴ mol ² dm ⁻⁶ (at 298K (25 °C))
specific heat capacity of water	$c = 4.18 \mathrm{kJ kg^{-1} K^{-1}} (4.18 \mathrm{J g^{-1} K^{-1}})$

							The Pe	riodic Ta	The Periodic Table of Elements	ments							
								Grc	Group								
~	2											13	14	15	16	17	18
							-										2
							т										He
				Key			hydrogen 1.0										helium 4.0
3	4	_		atomic number		_		_				5	9	7	80	6	10
	Be		ato	atomic symbol	bol							Ш	U	z	0	ш	Ne
lithium 6.9	beryllium 9.0		rels	name relative atomic mass	ISS							boron 10.8	carbon 12.0	nitrogen 14.0	oxygen 16.0	fluorine 19.0	neon 20.2
	12	_									-	13	14	15	16	17	18
	Mg											Al	Si.	٩	ა	Cl	Ar
sodium 23.0	magnesium 24.3	ი	4	5	9	7	8	6	10	11	12	aluminium 27.0	silicon 28.1	phosphorus 31.0	sulfur 32.1	chlorine 35.5	argon 39.9
	20		22	23	24	25	26	27	28	29	30	31	32	33	34	35	36
¥	Ca	Sc	F	>	ŗ	Mn	Ъе	ပိ	ïZ	Cu	Zn	Ga	Ge	As	Se	Br	Кr
potassium 39.1	calcium 40.1	scandium 45.0	titanium 47.9	vanadium 50.9	chromium 52.0	manganese 54.9	iron 55.8	cobalt 58.9	nickel 58.7	copper 63.5	zinc 65.4	gallium 69.7	germanium 72.6	arsenic 74.9	selenium 79.0	bromine 79.9	krypton 83.8
37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54
Rb	S	≻	Zr	qN	Mo	р	Ru	Rh	Ъd	Ag	рС	In	Sn	Sb	Те	Ι	Xe
rubidium 85.5	strontium 87.6	yttrium 88.9	zirconium 91.2	niobium 92.9	molybdenum 95.9	technetium -	ruthenium 101.1	rhodium 102.9	palladium 106.4	silver 107.9	cadmium 112.4	indium 114.8	tin 118.7	antimony 121.8	tellurium 127.6	iodine 126.9	xenon 131.3
55	56	57-71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86
Cs	Ba	lanthanoids	Ηf	Та	≥	Re	Os	Ir	Ę	Au	Hg	1T	Pb	Ξ	Ро	At	Rn
caesium 132.9	barium 137.3		hafnium 178.5	tantalum 180.9	tungsten 183.8	rhenium 186.2	osmium 190.2	iridium 192.2	platinum 195.1	gold 197.0	mercury 200.6	thallium 204.4	lead 207.2	bismuth 209.0	polonium –	astatine -	radon -
87	88	89-103	104	105	106	107	108	109	110	111	112	113	114	115	116	117	118
г Ц	Ra	actinoids	Rf	Db	Sg	Bh	Hs	Mt	Ds	Rg	ő	ЧN	ĿΙ	Mc	۲	Ч s	Og
francium -	radium -		rutherfordium 	dubnium –	seaborgium -	bohrium	hassium -	meitnerium -	darmstadtium -	roentgenium -	copernicium -	nihonium I	flerovium -	moscovium -	livermorium –	tennessine -	oganesson
		57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	
lanthanoids		La	Ŭ	Pr	ΡN	Рш	Sm	Eu	рд	Tb	D	Ч	ц	Tm	γb	Lu	
		lanthanum 138.9	0.	praseodymium 140.9	ēu	promethium -	samarium 150.4	europium 152.0	gadolinium 157.3	terbium 158.9	dysprosium 162.5	holmium 164.9	erbium 167.3	thulium 168.9	ytterbium 173.1	lutetium 175.0	
		89	06	91		93	94	95	96	97	98	66	100	101	102	103	
actinoids		Ac		Ра		dN	Pu	Am	CB	¥	Ç	Es	ШШ	Md	No	<u>_</u>	
		actinium -	thorium 232.0	protactinium 231.0	uranium 238.0	neptunium –	plutonium –	americium -	curium I	berkelium –	californium -	einsteinium –	fermium -	mendelevium -	nobelium 	lawrencium -	

To avoid the issue of disclosure of answer-related information to candidates, all copyright acknowledgements are reproduced online in the Cambridge Assessment International Education Copyright Acknowledgements Booklet. This is produced for each series of examinations and is freely available to download at www.cambridgeinternational.org after the live examination series.