International AS & A Level	Cambridge International Ac	dvanced Subsidiary and Advanced Lo	evel
CANDIDATE NAME			
CENTRE NUMBER		CANDIDATE NUMBER	
CHEMISTRY			9701/35
Paper 3 Advar	nced Practical Skills 1		May/June 2017
Candidates an	swer on the Question Paper.		2 hours

Cambridge International Examinations

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.

As listed in the Confidential Instructions

Answer **all** questions.

Additional Materials:

Cambridge

89062

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 10 and 11. A copy of the Periodic Table is printed on page 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.



2

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1 Sulfur forms the peroxodisulfate anion, $S_2O_8^{2-}$. This ion can oxidise iodide ions, I⁻, to iodine, I₂, as shown in the equation.

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

You will carry out a series of experiments to investigate how the rate of this reaction is affected by changing the concentration of the solutions.

The rate can be measured by adding thiosulfate ions, $S_2O_3^{2-}$, and starch indicator. As the reaction between $S_2O_8^{2-}$ and I^- occurs iodine is produced, but it reacts immediately with the thiosulfate.

 $I_2(aq) + 2S_2O_3^{2-}(aq) \rightarrow 2I^{-}(aq) + S_4O_6^{2-}(aq)$

When all the thiosulfate has reacted, the iodine will remain in the mixture and cause the starch indicator to turn blue-black. The rate of reaction may be determined by timing how long it takes the reaction mixture to turn blue-black.

FA 1 is 0.0200 mol dm⁻³ potassium peroxodisulfate, $K_2S_2O_8$. **FA 2** is 1.00 mol dm⁻³ potassium iodide, K1. **FA 3** is 0.00500 mol dm⁻³ sodium thiosulfate, $Na_2S_2O_3$. starch indicator

Read through the instructions carefully and prepare a table for your results on page 4 before starting any practical work.

(a) Method

Experiment 1

- Fill the burette labelled **FA 1** with **FA 1**.
- Use the pen to label one of the 100 cm³ beakers 'A' and the other 100 cm³ beaker 'B'.
- Run 20.00 cm³ of **FA 1** from the burette into beaker **A**.
- Use the measuring cylinder to add 20.0 cm³ of FA 2 into beaker B.
- Use the measuring cylinder to add 10.0 cm³ of **FA 3** to beaker **B**.
- Add 10 drops of starch indicator 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 a white tile.
- Stop timing as soon as the solution turns blue-black.
- Record this reaction time to the nearest second in your results table.
- Wash out both beakers and shake to remove excess water.

Experiment 2

- Fill a second burette with distilled water.
- Run 10.00 cm³ of **FA 1** into beaker **A**.
- Run 10.00 cm³ of distilled water into beaker **A**.
- Use the measuring cylinder to add 20.0 cm³ of **FA 2** into beaker **B**.
- Use the measuring cylinder to add 10.0 cm³ of **FA 3** to beaker **B**.
- Add 10 drops of starch indicator 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 a white tile.
- Stop timing as soon as the solution turns blue-black.
- Record this reaction time to the nearest second in your results table.
- Wash out both beakers and shake to remove excess water.

Experiments 3-5

Carry out three further experiments to investigate how the reaction time changes with different volumes of potassium peroxodisulfate, FA 1.
Note that the combined volume of FA 1 and distilled water must always be 20.00 cm³.
Do not use a volume of FA 1 that is less than 6.00 cm³.

Keep FA 1, FA 2, FA 3 and the starch indicator for use in (e).

Calculating the rate of the reaction

The rate of the reaction can be represented by the formula shown.

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

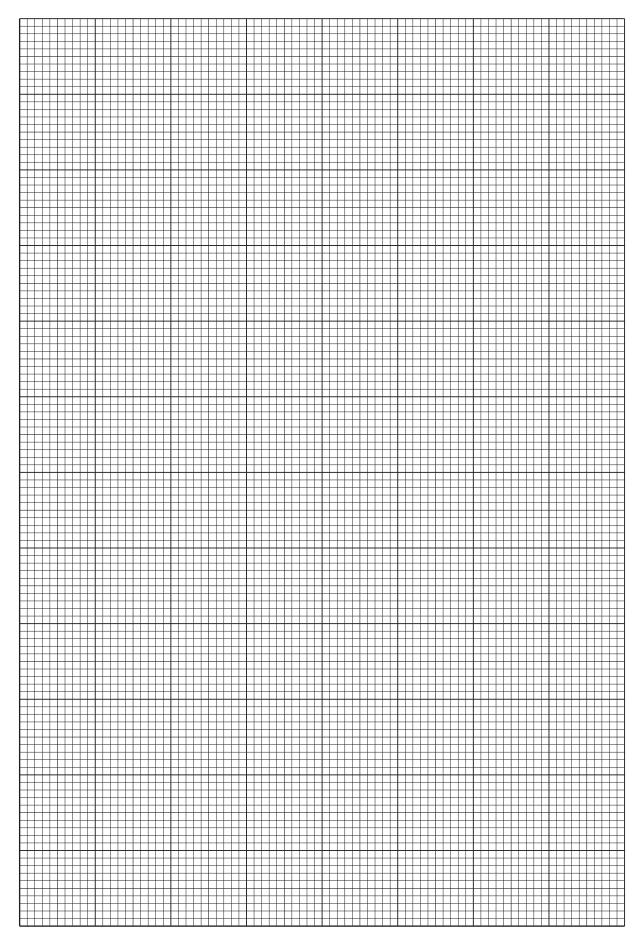
Use this formula to calculate the rate for each of your five experiments.

Record all your results in a single 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.

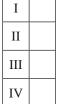
Ι	
II	
III	
IV	
V	
VI	
VII	
VIII	
IX	
X	

[10]

(b) On the grid on page 5, plot the rate (*y*-axis) against the volume of **FA 1** (*x*-axis). Include the origin in your plot. Draw a straight line of best fit and circle any clearly anomalous points.



5





[4]

[Turn over

(c) The volume of FA 1 is directly related to the concentration of potassium peroxodisulfate.

From your results, what can be stated about the relationship between the rate of reaction and the concentration of potassium peroxodisulfate?

......[1]

(d) (i) Use your graph to calculate the reaction time you would expect to measure if you carried out an experiment using 5.00 cm³ of FA 1.
Show your working.

reaction time = s

 (ii) Assume that the error in the time measured for each reaction was ±0.5s in total. Calculate the maximum percentage error in the reaction time you measured in Experiment 1. Show your working.

maximum percentage error = %

(iii) A student suggested that this error could be reduced if 0.0100 mol dm⁻³ sodium thiosulfate were used in place of **FA 3**.

Do you agree with this student? Explain your answer.

.....

(iv) A student repeated **Experiment 1** but used 0.100 mol dm⁻³ sodium thiosulfate in place of **FA 3**. The student found that the reaction mixture never turned blue-black.

Explain why.

[5]

- (e) (i) Using the same method as in (a), carry out an additional experiment to record the reaction time to the nearest second when the following solutions are mixed together.
 - 10.00 cm³ of **FA 1**
 - 20.0 cm³ of **FA 2**
 - 5.0 cm³ of **FA 3**
 - 15.00 cm³ of distilled water
 - 10 drops of starch indicator

reaction time =

(ii) Use your answer to (i) to **estimate** the reaction time that would be measured if the following solutions were mixed together.

DO NOT CARRY OUT THIS EXPERIMENT

- 10.00 cm³ of **FA 1**
- 20.0 cm³ of **FA 2**
- 20.0 cm³ of FA 3
- 10 drops of starch indicator

Explain your answer.

estimated reaction time =

[4]

[Total: 24]

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

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.

- (a) FA 4 and FA 5 are aqueous solutions. Each solution contains two different cations and the sulfate anion.
 - (i) Carry out the following tests and record your observations.

fact	obser	vations								
test	FA 4	FA 5								
To a 1 cm depth of solution in a boiling tube, add aqueous sodium hydroxide, then										
warm the tube carefully.										
To a 1 cm depth of solution in a test-tube, add aqueous ammonia.										

(ii) Identify as many as possible of the cations in FA 4 and FA 5.

FA 4 contains the cation(s)

FA 5 contains the cation(s)

- (b) FA 6 is a salt containing either the sulfate anion or sulfite anion. You will first make a solution of FA 6.
 - Rinse one of the 100 cm³ beakers with distilled water.
 - Place all the sample of **FA 6** into the beaker and add approximately 40 cm³ of distilled water.
 - Stir the mixture until the solid has dissolved.

Select reagent(s) and carry out tests to identify the anion in **FA 6**. Record your results in the space below.

(c) FA 7 is a solution containing one cation and the sulfate anion.
FA 8 is a solution containing the sodium cation and one of the anions from those listed in the Qualitative Analysis Notes.

(i) Carry out the following tests to determine the formulae of FA 7 and FA 8.

test	observations
To a 2 cm depth of FA 7 in a test-tube, add a 2 cm strip of magnesium.	
To a 1 cm depth of FA 7 in a test-tube add a 1 cm depth of FA 8 and shake the tube.	

(ii) The formula of FA 7 is

The formula of **FA 8** is

(iii) Give the ionic equation for the reaction that takes place when magnesium is added to **FA 7**. Include state symbols.

.....

[6]

Qualitative Analysis Notes

1 Reactions of aqueous cations

ian	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 NH₃(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 A <i>l</i> 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

		17 18	2	He	helium 4.0	9 10		fluorine neon 19.0 20.2			chlorine argon 35.5 39.9			bromine krypton 79.9 83.8			iodine xenon 126.9 131.3		At Rn					4		LU	Iutetium 175.0	103	Ľ	lawrencium -
		16				∞	0	oxygen 16.0	16	თ	sulfur 32.1	34	Se	selenium 79.0	52	Te	tellurium 127.6	84	Ро	polonium –	116	2	livermorium -	40		γD	ytterbium 173.1	102	No	nobelium -
		15				7	z	nitrogen 14.0	15	٩	phosphorus 31.0	33	As	arsenic 74.9	51	Sb	antimony 121.8	83	Bi	bismuth 209.0				C ^y	B H	E	thulium 168.9	101	Md	mendelevium -
		14				9	U	carbon 12.0	14	<u>S</u>	silicon 28.1	32	Ge	germanium 72.6	50	Sn	tin 118.7	82	Pb	lead 207.2	114	Fl	flerovium -	00	° L	L U	erbium 167.3	100	Еm	fermium I
		13				5	В	boron 10.8	13	Al	aluminium 27.0	31	Ga	gallium 69.7	49	In	indium 114.8	81	Τl	thallium 204.4				57	/o	0H	holmium 164.9	66	Es	einsteinium -
											12	30	Zn	zinc 65.4	48	S	cadmium 112.4	80	Нg	mercury 200.6		ű	copernicium -	55	⁸ (h	dysprosium 162.5	98	ç	californium -
ements										11	29	Cu	copper 63.5	47	Ag	silver 107.9	79	Au	gold 197.0	111	Rg	roentgenium -	5 C	6 F	a	terbium 158.9	67	凝	berkelium -	
ble of Ele	Group										10	28	ïZ	nickel 58.7	46	Ъd	palladium 106.4	78	Ŧ	platinum 195.1	110	Ds	darmstadtium -	2	- 5 (<u>9</u>	gadolinium 157.3	96	СЗ	curium I
The Periodic Table of Elements	Gro										0	27	ပိ	cobalt 58.9	45	Rh	rhodium 102.9	77	Ir	iridium 192.2	109	Mt	meitnerium -	52	2° L	П	europium 152.0	95	Am	americium -
The Pe			-	I	hydrogen 1.0						8			iron 55.8		Ru	ruthenium 101.1	76	Os	osmium 190.2	108	Чs	hassium -				samarium 150.4		Pu	plutonium -
						1					7	25	Mn	manganese 54.9	43	Ц	technetium -	75	Re	rhenium 186.2	107	Bh	bohrium –	Č	ō (ЕЧ	promethium -	93	dN	neptunium -
							loc	SS			9	24	ŗ	chromium 52.0	42	Мо	molybdenum 95.9	74	$^{>}$	tungsten 183.8	106	Sg	seaborgium -	eo.	no	Na	neodymium 144.4	92	⊃	uranium 238.0
					Key	atomic number	atomic symbol	name relative atomic mass			5	23	>	vanadium 50.9	41	ЧN	niobium 92.9	73	Та	tantalum 180.9	105	Db	dubnium –	CD	20 (ŗ	praseodymium 140.9	91	Ра	protactinium 231.0
							ato	rela			4	22	F	titanium 47.9	40	Zr	zirconium 91.2	72	Ηf	hafnium 178.5	104	Rf	rutherfordium -	E0	^ი (cerium 140.1	06	Ч	thorium 232.0
									1		С	21	Sc	scandium 45.0	39	≻	yttrium 88.9	57-71	lanthanoids		89-103	actinoids		57		Га	lanthanum 138.9	89	Ac	actinium -
		2				4	Be	beryllium 9.0	12	Mg	magnesium 24.3	20	Ca	calcium 40.1	38	ي ا	strontium 87.6	56	Ba	barium 137.3	88	Ra	radium –	_	-	Ids				
		1				e	:	lithium 6.9	1	Na	sodium 23.0	19	¥	potassium 39.1	37	Rb	rubidium 85.5	55	Cs	caesium 132.9	87	Ļ	francium -		;	lanthanoids			actinoids	

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