



Cambridge International Examinations

Cambridge International Advanced Subsidiary and Advanced Level

CANDIDATE NAME										
CENTRE NUMBER						CANDIDATE NUMBER				
CHEMISTRY									970	1/33
Paper 3 Advance	ced Prac	tical Skil	s 1				Febru	Jary/N	/larch	2018
									2 h	ours
Candidates ans	swer on t	he Quest	ion Pap	er.						
Additional Mate	rials:	As liste	d in the	Confide	ntial Instructions					

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		
Total		

This document consists of 14 printed pages, 2 blank pages and 1 Insert.



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 You will investigate how increasing temperature affects the rate of a reaction.

Sodium thiosulfate reacts with acid to form a pale yellow precipitate of sulfur. The ionic equation for the reaction is given.

$$S_2O_3^{2-}(aq) + 2H^+(aq) \rightarrow S(s) + SO_2(q) + H_2O(1)$$

You will measure the time it takes for the sulfur formed in the reaction to obscure the print on the Insert supplied.

Record your results in a table on page 4. Your table should include the rate of reaction for each experiment.

FA 1 is an $18.1 \,\mathrm{g}\,\mathrm{dm}^{-3}$ solution of hydrated sodium thiosulfate, $\mathrm{Na_2S_2O_3.5H_2O}$. **FA 2** is a $0.050\,\mathrm{mol}\,\mathrm{dm}^{-3}$ solution of a strong monoprotic acid, H**Z**.

(a) Method

- Approximately half fill the 250 cm³ beaker with tap water and place it on the tripod and gauze over the Bunsen burner.
- Heat the water in the beaker to about 55 °C and then switch off the Bunsen burner. This will be your hot water bath.
- Use the 25 cm³ measuring cylinder to transfer 10 cm³ of **FA 1** into boiling tube **1**. Place boiling tube **1** into your hot water bath.
- Use the 50 cm³ measuring cylinder to transfer 20 cm³ of **FA 2** into boiling tube **2**. Place boiling tube **2** into your hot water bath.
- Leave boiling tubes 1 and 2 in the hot water bath to heat up for use in **Experiment 2**.
- Start Experiment 1.

Experiment 1

- Use the 50 cm³ measuring cylinder to transfer 20 cm³ of **FA 2** into the 100 cm³ beaker.
- Measure and record the temperature of FA 2.
- Use the 25 cm³ measuring cylinder to transfer 10 cm³ of **FA 1** into the same beaker and start timing **immediately**.
- Swirl the beaker once to mix the solutions and place the beaker on the Insert.
- Look down through the beaker and contents onto the Insert.
- Stop timing as soon as the precipitate of sulfur obscures the print on the Insert.
- Record the reaction time to the nearest second.
- Empty the contents of the beaker into the quenching bath.
- Rinse and dry the beaker so it is ready for use in **Experiment 2**.

Experiment 2

- Measure and record the temperature of FA 2 in boiling tube 2.
- Carefully transfer the hot contents of boiling tube 2 into the 100 cm³ beaker.
- Carefully transfer the hot contents of boiling tube 1 into the same beaker and start timing immediately.
- Swirl the beaker once to mix the solutions and place the beaker on the Insert.
- Look down through the beaker and contents onto the Insert.
- Stop timing as soon as the precipitate of sulfur obscures the print on the Insert.
- Record the reaction time to the nearest second.
- Empty the contents of the beaker into the guenching bath.
- Rinse and dry the beaker so it is ready for use in Experiment 3.

Experiment 3

- Use the 25 cm³ measuring cylinder to transfer 10 cm³ of **FA 1** into boiling tube **1**. Place boiling tube **1** into your hot water bath.
- Use the 50 cm³ measuring cylinder to transfer 20 cm³ of **FA 2** into boiling tube **2**. Place boiling tube **2** into your hot water bath.
- Place the thermometer in boiling tube 2. When the temperature of FA 2 is about 8 °C lower than that for Experiment 2 record the temperature. Remove the thermometer and transfer the contents of boiling tube 2 into the 100 cm³ beaker.
- Transfer the contents of boiling tube 1 into the same beaker and start timing immediately.
- Swirl the beaker once to mix the solutions and place the beaker on the Insert.
- Look down through the beaker and contents onto the Insert.
- Stop timing as soon as the precipitate of sulfur obscures the print on the Insert.
- Record the reaction time to the nearest second.
- Empty the contents of the beaker into the quenching bath.
- Rinse and dry the beaker so it is ready for use in Experiments 4 and 5.

Experiments 4 and 5

- Repeat the method for Experiment 3 but at two different temperatures.
- Keep the temperature of FA 2 between room temperature and 55 °C. Do not exceed 55 °C.

Record all your results in your table on page 4.

Results

The rate of reaction can be calculated as shown.

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

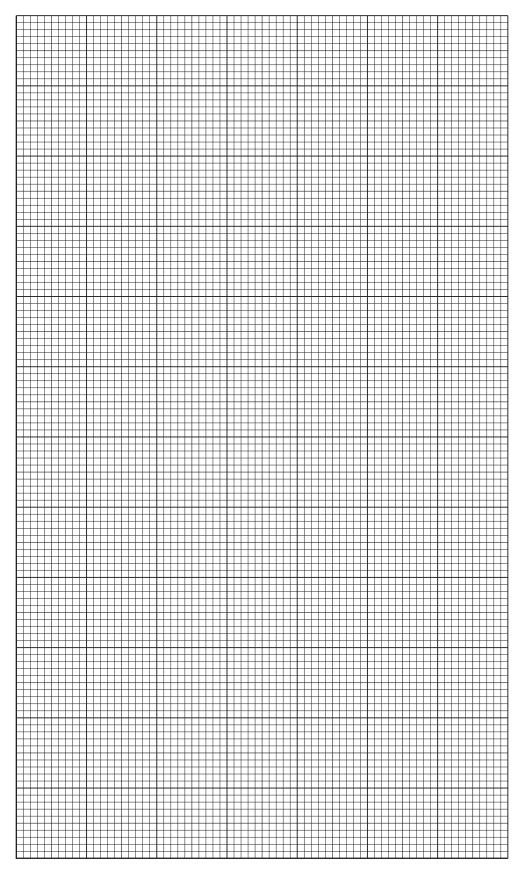
Calculate the rate of reaction for each of your **five** experiments. Record these rates in your table.

I	
II	
III	
IV	
V	
VI	
VII	
VIII	

[8]

(b) On the grid plot a graph of rate of reaction on the *y*-axis, starting at zero, against temperature on the *x*-axis. Select a scale for the *x*-axis which includes a temperature of 15.0 °C. Label your axes and any points you consider anomalous.

Draw a line of best fit and extrapolate it to 15.0 °C.



Ι	
II	
III	
IV	

[4]

(c)		e your graph to calculate the time to the nearest second that the reaction would have taken ou had carried it out at 17.5°C. Show on the grid how you obtained your answer.
		time = s [2]
(d)		plain, by referring to your graph or your table of results, how the rate of reaction is affected ncreasing temperature.
		[2]
(e)	Cal	culations
	(i)	Calculate the concentration of hydrated sodium thiosulfate, ${\rm Na_2S_2O_3.5H_2O}$, in FA 1 in mol dm ⁻³ .
		concentration of $Na_2S_2O_3.5H_2O$ in FA 1 = mol dm ⁻³ [1]
	(ii)	Calculate the concentration of the strong monoprotic acid, HZ, in the solution immediately after FA 1 was added to FA 2 in the beaker.
		concentration of HZ = mol dm ⁻³ [1]
((iii)	Use the equation on page 2 to determine which reagent, FA 1 or FA 2, was in excess.
		The reagent in excess was [2]

(f)	(i)	Calculate the maximum percentage error in measuring the reaction time you recorded for Experiment 2 . Assume that the maximum error of the timer is ± 0.5 s.
		maximum percentage error in the reaction time = % [1]
	(ii)	A student suggested that the error in measuring the reaction time in Experiment 1 was greater than for Experiment 2 .
		Give one reason why the student could be correct.
		[1]
(g)	Su	ggest two ways to improve the accuracy of the results of these experiments.
	1	
	2	
		[2]
		[Total: 24]

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.

2 (a) FA 3 is a more concentrated solution of the strong monoprotic acid, HZ, used for Question 1.

Select **two** sets of reagents and suitable apparatus to use in **two** separate tests, **Test 1** and **Test 2**, to investigate the identity of the anion, **Z**⁻, present in **FA 3**. The anion is one of those listed in the Qualitative Analysis Notes.

Complete the 'test' boxes in the table **before** starting any practical work by circling whether you would use a test-tube or a boiling tube, and stating which reagents you would use.

Carry out your tests and record your observations. You must carry out both Test 1 and Test 2.

test	observations
Test 1	
To a 1 cm depth of FA 3 in a	
test-tube/boiling tube	
add	
(reagent(s))	
Test 2	
To a 1 cm depth of FA 3 in a	
test-tube/boiling tube	
add	
(reagent(s))	
	[4]
(b) Identify the anion present in HZ from your	observations in (a).
Z - is	[1]

(c) FA 4 and FA 5 both contain one cation and one anion. The ions present in FA 4 are different from the ions present in FA 5. All four ions are listed in the Qualitative Analysis Notes. You are to identify the four different ions.

Carry out the following tests and record your observations.

test	observations
To a small spatula measure of FA 4 in a boiling tube, add a 4 cm depth of FA 3 and shake the tube well. Leave the tube to stand for at least five minutes. Label the solution formed FA 6 .	
To a 1 cm depth of FA 5 in a test-tube, add aqueous sodium carbonate.	
To a 1 cm depth of FA 5 in a test-tube, add aqueous sodium hydroxide.	
To a 1 cm depth of FA 5 in a test-tube, add aqueous ammonia.	
To a 1 cm depth of FA 5 in a test-tube, add a few drops of aqueous silver nitrate.	
To a 1 cm depth of FA 5 in a test-tube, add a few drops of aqueous barium chloride or aqueous barium nitrate, then	
add a 1 cm depth of a suitable acid.	
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.	
To a 1 cm depth of FA 6 in a test-tube, add dilute sulfuric acid.	
To a 1 cm depth of FA 6 in a test-tube, add a 1 cm depth of FA 5 .	

(a)	aqueous barium nitrate in (c) .	or
	The acid added was	[1]
(e)	Identify the ions present in FA 4 and FA 5 from your observations in (c) .	

	cation	anion
FA 4		
FA 5		

[2]

[Total: 16]

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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, NH ₄ +(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, 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), 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), 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)	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
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	He	helium 4.0	10	Ne	neon 20.2	18	Ā	argon 39.9	36	궃	kryptor 83.8	22	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				80	0	oxygen 16.0	16	S	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	Sb	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	Ър	lead 207.2	114	Εl	flerovium
13				5	В	boron 10.8	13	Ν	aluminium 27.0	31	Ga	gallium 69.7	49	In	indium 114.8	81	11	thallium 204.4			
									12	30	Zu	zinc 65.4	48	පි	cadmium 112.4	80	Нg	mercury 200.6	112	ပ်	copernicium
									7	29	Cn	copper 63.5	47	Ag	silver 107.9	79	Au	gold 197.0	111	Rg	roentgenium -
									10	28	Z	nickel 58.7	46	Pd	palladium 106.4	78	귙	platinum 195.1	110	Ds	darmstadtium -
									6	27	ပိ	cobalt 58.9	45	뫈	rhodium 102.9	77	'n	iridium 192.2	109	¥	meitnerium -
	-	I	hydrogen 1.0						80	26	Pe	iron 55.8	44	Ru	ruthenium 101.1	92	SO	osmium 190.2	108	Ϋ́	hassium
				_					7	25	Mn	manganese 54.9	43	ည	technetium -	75	Re	rhenium 186.2	107	В	bohrium
					loc	18S			9	24	ပ်	chromium 52.0	42	Mo	molybdenum 95.9	74	>	tungsten 183.8	106	Sg	seaborgium -
			Key	atomic number	mic sym	name ative atomic ma			2	23	>	vanadium 50.9	41	qN	niobium 92.9	73	Б	tantalum 180.9	105	Ср	dubnium –
					ato	rela			4	22	F	titanium 47.9	40	Zr	zirconium 91.2	72	Ξ	hafnium 178.5	104	Ŗ	rutherfordium -
									က	21	Sc	scandium 45.0	39	>	yttrium 88.9	57–71	lanthanoids		89–103	actinoids	
7				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
_				8	=	lithium 6.9	11	Na	sodium 23.0	19	\prec	potassium 39.1	37	Rb	rubidium 85.5	55	S	caesium 132.9	87	ቷ	francium
	13 14 15 16 17	13 14 15 16 17	13 14 15 16 17 H	13 14 15 16 17	13 14 15 16 17 17 18 18 17 17 18 18	2 13 14 15 16 17 17 17 18 18 19 17 17 18 19 19 19 19 19 19 19	2 13 14 15 16 17 17 18 18 19 17 18 18 19 19 19 19 19 19	13 14 15 16 17 17 18 18 19 17 18 18 19 19 19 19 19 19	1	2 13 14 15 16 17 17 18 19 19 19 19 19 19 19	1	2 13 14 15 16 17 18 18 19 19 19 19 19 19	2 13 14 15 15 16 17 18 19 19 19 19 19 19 19	2 1 1 1 1 1 1 1 1 1	2 1.2	1	1	1	1	The continue of the continue	The control of the

71 Lu	175.0	103	ځ	lawrencium	ı
70 Yb	173.1	102	9 N	nobelium	ı
E9 L	168.9	101	Md	mendelevium	ı
68 Fr	167.3	100	Fm	ferminm	ı
67 Holmium	164.9	66	Es	einsteinium	1
66 Dy	162.5	86	ర్	californium	ı
65 Tb	158.9	26	Ř	berkelium	1
Gd gadolinium	157.3	96	Cm	curium	1
63 Eu	152.0	98	Am	americium	ı
62 Sm	150.4	94	Pn	plutonium	1
61 Pm	ı	93	ď	neptunium	1
Nd Nd	144.4	92	\supset	uranium	238.0
Pr Pr		91	Pa	protactinium	231.0
S8 Ce	140.1	06	T	thorium	232.0
57 La	138.9	68	Ac	actinium	

lanthanoids

actinoids

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