

Cambridge International AS & A Level

CANDIDATE NAME						
CENTRE NUMBER				CANDIDATE NUMBER		

CHEMISTRY 9701/31

Paper 3 Advanced Practical Skills 1

May/June 2023

2 hours

You must answer on the question paper.

You will need: The materials and apparatus listed in the confidential instructions

Insert (enclosed)

INSTRUCTIONS

- Answer all questions.
- Use a black or dark blue pen. You may use an HB pencil for any diagrams or graphs.
- Write your name, centre number and candidate number in the boxes at the top of the page.
- 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.

INFORMATION

- 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.
- Important values, constants and standards are printed in the question paper.
- Notes for use in qualitative analysis are provided in the question paper.
- The insert contains additional resources referred to in the questions.

Session
Laboratory

For Examiner's Use						
1						
2						
3						
Total						

This document has 16 pages. Any blank pages are indicated.

BLANK PAGE

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 The thiosulfate ion, S₂O₃²⁻, decomposes when an acid is added.

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

The rate of this reaction can be investigated by measuring how long it takes the solution to produce enough sulfur so that it cannot be seen through.

You will investigate how the concentration of the thiosulfate ion affects the rate of the reaction.

Note: A small amount of sulfur dioxide gas may be formed in the experiment. It is very important that you avoid inhaling any fumes. As soon as each experiment is complete, empty the reaction mixture into the quenching bath and rinse the beaker thoroughly.

FA 1 is $0.100\,\mathrm{mol\,dm^{-3}}$ sodium thiosulfate, $\mathrm{Na_2S_2O_3}$. **FA 2** is $2.00\,\mathrm{mol\,dm^{-3}}$ hydrochloric acid, HC *l*.

(a) Method

Experiment 1

- Fill a burette with **FA 1**.
- Run 40.00 cm³ of **FA 1** into the 100 cm³ beaker.
- Use the 25 cm³ 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.
- View the printing on the insert from above, through the solution.
- Stop timing when the print on the insert becomes obscured.
- Record this reaction time to the nearest second in the space for results on page 4.
- Empty the contents of the beaker into the guenching bath.
- Rinse and dry the beaker and glass rod so they are ready to use in Experiment 2.

Experiment 2

- Refill the burette with FA 1.
- Fill the other burette with distilled water.
- Run 20.00 cm³ of **FA 1** into the 100 cm³ beaker.
- Run 20.00 cm³ of distilled water into the same beaker.
- Use the 25 cm³ measuring cylinder to measure 10.0 cm³ of **FA 2**.
- Add the **FA 2** to the solution in the beaker and start timing **immediately**.
- Stir the mixture once and place the beaker on the printed insert.
- View the printing on the insert from above, through the solution.
- Stop timing when the print on the insert becomes obscured.
- Record this reaction time to the nearest second.
- Empty the contents of the beaker into the quenching bath.
- Rinse and dry the beaker and glass rod so they are ready to use in the next experiment.

Experiments 3-5

Carry out three further experiments to investigate how using different volumes of FA 1
affects the reaction time.

Note that the combined volumes of **FA 1** and distilled water must always be 40.00 cm³.

Do **not** use a volume of **FA 1** that is less than 15.00 cm³.

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 formula.

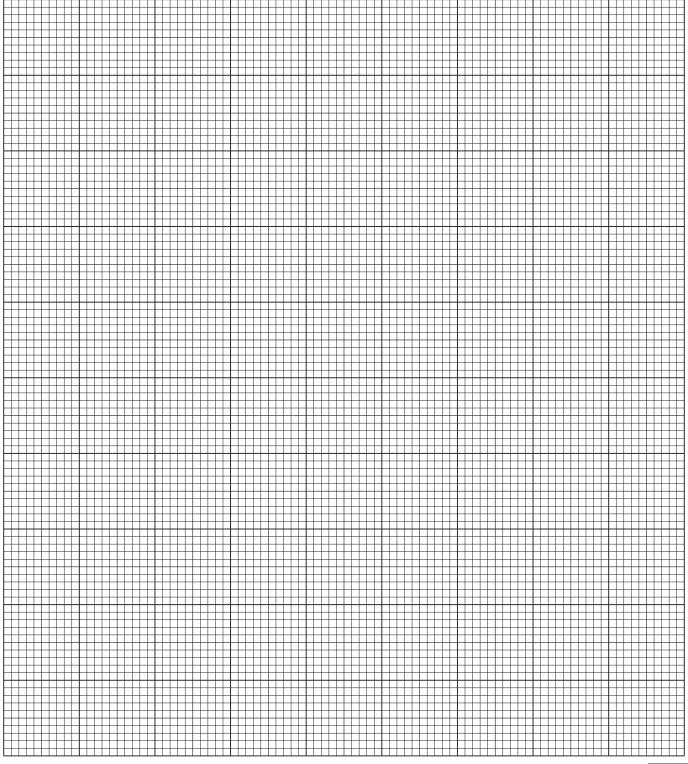
rate =
$$\frac{1000}{\text{reaction time}}$$

Results

[8]

(b) On the grid, plot the rate (*y*-axis) against the volume of **FA 1** (*x*-axis). Start each axis at the origin (0,0).

Ring any anomalous points. Draw a line of best fit.



I II III IV

[4]

(c)	In these experiments, the volume of FA 1 is a measure of the concentration of thiosulfate ions.
	A student suggested that the graph shows that the rate of reaction is directly proportional to the concentration of thiosulfate ions.
	Explain, using your graph, whether you agree with this student.
	[1]
(d)	Use your graph to calculate the time you would expect to record if you had used 12.50 cm ³ or FA 1 and followed the same method.
	Show clearly on the graph how you calculated this time.
	time = s [2]
(e)	Another student broke the beaker and decided to use a Petri dish instead.
	beaker Petri dish
	Fig. 1.1
	What effect, if any, would this have on the times measured in the experiment in (a) ? Explain your answer.
	[2]

[Total: 17]

7

BLANK PAGE

2 Many salts contain water of crystallisation which can be removed by heating to form the anhydrous salt. You will determine the enthalpy change of dehydration for hydrated magnesium sulfate.

$$MgSO_4 \cdot 7H_2O(s) \rightarrow MgSO_4(s) + H_2O(l)$$

You will determine this enthalpy change by measuring the changes in temperature when samples of hydrated magnesium sulfate and anhydrous magnesium sulfate are dissolved separately in excess water.

FA 3 is hydrated magnesium sulfate, MgSO₄•7H₂O. **FA 4** is anhydrous magnesium sulfate, MgSO₄.

(a) Method

Experiment 1

- Weigh the container with FA 3. Record the mass in the space below.
- Support the cup in the 250 cm³ beaker.
- Rinse the 25 cm³ measuring cylinder with distilled water. 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.
- Tip all of the **FA 3** into the water in the cup. Stir the mixture thoroughly.
- Record the highest or lowest temperature of the mixture.
- Calculate the change in temperature. Record this change.
- Weigh the container with any remaining FA 3. Record the mass.
- Calculate the mass of FA 3 used. Record this mass.

Experiment 2

Repeat the method using FA 4 in place of FA 3. Use the second cup.

Results

I II III IV V

[5]

(b) Calcu	lations
----	---------	---------

(i) Calculate the heat energy transferred, in J, in each experiment.

Assume that 4.18 J of heat energy changes the temperature of $1.0\,\mathrm{cm}^3$ of solution by $1.0\,^\circ\mathrm{C}$.

Experiment 1 with FA 3	Experiment 2 with FA 4
heat energy = J	heat energy = J

(ii) Calculate the enthalpy change, ΔH , in kJ mol⁻¹, when 1.0 mol of solid dissolves in water in each experiment.

Experiment 1 with FA 3	Experiment 2 with FA 4
$\Delta H_1 = \frac{1}{\text{sign}} \text{ kJ mol}^{-1}$	$\Delta H_2 = \frac{1}{\text{sign}} \text{ kJ mol}^{-1}$

[3]

[1]

(iii) Use your answers to **(b)(ii)** to calculate the enthalpy change when 1.0 mol of hydrated magnesium sulfate is dehydrated to form 1.0 mol of anhydrous magnesium sulfate.

$$MgSO_4 \cdot 7H_2O(s) \rightarrow MgSO_4(s) + 7H_2O(l)$$

Show clearly, by a Hess's energy cycle or any other suitable means, how you calculated your answer.

If you were unable to complete the calculation in **(b)(ii)** then assume that the enthalpy change in Experiment 1, ΔH_1 , is +8.7 kJ mol⁻¹ and that the enthalpy change in Experiment 2, ΔH_2 , is -15.5 kJ mol⁻¹. These may **not** be the correct values.

enthalpy change of dehydration of
$$MgSO_4 \cdot 7H_2O = \dots kJmol^{-1}$$
 [2]

[Total: 11]

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
- 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.

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

No additional tests should be attempted.

3 (a) (i) A student finds a container of a compound which is thought to be FA 3. The student labels the container FA 5.

Carry out tests to determine whether your sample of **FA 5** contains magnesium ions, sulfate ions and water of crystallisation.

Record your tests and observations in a suitable form in the space below.

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

(ii) Complete Table 3.1 to indicate the contents of FA 5.Put a tick (✓) in one box in each row.

Table 3.1

	yes	no
FA 5 contains magnesium ions		
FA 5 contains sulfate ions		
FA 5 contains water of crystallisation		

[1]

(b)	FA 6	contains	one	cation	and	one	anion,	both	of	which	are	listed	in	the	Qualitative	analy	sis
	notes	S.															

(i)	In a hard-glass test-tube heat a spatula measure of FA 6 . Heat gently at first and then heat strongly. Record all your observations.
	[2]

(ii) Carry out the following tests and record your observations.

Table 3.2

test	observations
Test 1 To a small spatula measure of FA 6 in a test-tube, add 5 cm depth of dilute hydrochloric acid.	
Keep the solution for Test 2.	
Test 2 Put 1 cm depth of the solution from Test 1 in a test-tube. Add aqueous ammonia dropwise at first and then until it is in excess.	

[2]

(iii)	Use your observations from (b)(i) and (ii) to identify the ions present in FA 6.									
	cation	anion	[1]							
(iv)	Write the ionic equation for added. Include state symb	or the reaction that occurs in Test 2 before excess arols.	mmonia is							
	If you were unable to ident	tify the cation in the solution then use the symbol M^2	+.							
			[2]							
			[Total: 12]							

Qualitative analysis notes

1 Reactions of cations

cation	reaction 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, Cl ⁻ (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)	${ m NH_3}$ liberated on heating with ${ m OH^-}({ m aq})$ and ${ m A}l$ foil; decolourises acidified aqueous ${ m KMnO_4}$
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	$F = 9.65 \times 10^4 \mathrm{C}\mathrm{mol}^{-1}$
Avogadro constant	$L = 6.022 \times 10^{23} \mathrm{mol}^{-1}$
electronic charge	$e = -1.60 \times 10^{-19} \mathrm{C}$
molar volume of gas	$V_{\rm m} = 22.4 {\rm dm^3 mol^{-1}}$ at s.t.p. (101 kPa and 273 K) $V_{\rm m} = 24.0 {\rm dm^3 mol^{-1}}$ at room conditions
ionic product of water	$K_{\rm w} = 1.00 \times 10^{-14} \rm mol^2 dm^{-6} (at 298 K (25 ^{\circ}C))$
specific heat capacity of water	$c = 4.18 \mathrm{kJ kg^{-1} K^{-1}} (4.18 \mathrm{J g^{-1} K^{-1}})$

The Periodic Table of Elements

			<i>a</i>	E		<i>a</i>	- ^·		_	- C					~	چ ₀		_	c	_		nos
	18	2	Ĭ	heliu 4.0	10	ž	neo.	18	Ā	argon 39.9	36	Ž	kryptt 83.8	54	×	xenc 131.	86	쬬	rado	118	ŏ	oganes
	17				6	ш	fluorine 19.0	17	Cl	chlorine 35.5	35	Ā	bromine 79.9	53	Н	iodine 126.9	85	Ą	astatine -	117	<u>~</u>	tennessine -
	16				80	0	oxygen 16.0	16	S	sulfur 32.1	34	Se	selenium 79.0	52	<u>e</u>	tellurium 127.6	84	Ъо	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	<u>B</u>	bismuth 209.0	115	Mc	moscovium
	4				9	ပ	carbon 12.0	14	S	silicon 28.1	32	Ge	germanium 72.6	20	S	tin 118.7	82	Ър	lead 207.2	114	Εl	flerovium
	13				2	Ф	boron 10.8	13	Ρl	aluminium 27.0	31	Ga	gallium 69.7	49	In	indium 114.8	84	<i>1</i> L	thallium 204.4	113	R	nihonium
										12	30	Zu	zinc 65.4	48	පි	cadmium 112.4	80	£	mercury 200.6	112	ပ်	copernicium
										7	29	Cn	copper 63.5	47	Ag	silver 107.9	62	Au	gold 197.0	111	Rg	roentgenium
dn										10	28	Z	nickel 58.7	46	Pd	palladium 106.4	78	귙	platinum 195.1	110	Ds	darmstadtium -
Group										6	27	ဝိ	cobalt 58.9	45	格	rhodium 102.9	77	ŗ	iridium 192.2	109	¥	meitnerium -
		-	I	hydrogen 1.0						œ	26	Pe	iron 55.8	44	Ru	ruthenium 101.1	9/	SO	osmium 190.2	108	Ϋ́	hassium
					_					7	25	Mn	manganese 54.9	43	ည	technetium -	75	Re	rhenium 186.2	107	Bh	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 50.9	41	g	niobium 92.9	73	<u>⊾</u>	tantalum 180.9	105	Op	dubnium
					o l	atol	relai			4	22	F	titanium 47.9	40	Zr	zirconium 91.2	72	Ξ	hafnium 178.5	104	꿒	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	26	Ba	barium 137.3	88	Ra	radium
	~				3	:=	lithium 6.9	#	Na	sodium 23.0	19	¥	potassium 39.1	37	Rb	rubidium 85.5	55	S	caesium 132.9	87	Ļ	francium

7.1	ŋ	lutetium 175.0	103	۲	lawrencium	-
20	Υp	ytterbium 173.1	102	%	nobelium	Ι
69	H	thulium 168.9	101	Md	mendelevium	ı
89	щ	erbium 167.3	100	Fm	ferminm	I
29	우	holmium 164.9	66	Es	einsteinium	I
99	۵	dysprosium 162.5	86	ŭ	californium	ı
65	Д	terbium 158.9	26	益	berkelium	I
49	В	gadolinium 157.3	96	CB	curium	-
63	En	europium 152.0	98	Am	americium	I
62	Sm	samarium 150.4	26	Pn	plutonium	_
61	Pm	promethium —	93	ď	neptunium	ı
09	PN	neodymium 144.4	92	\supset	uranium	238.0
59	Ą	praseodymium 140.9	91	Ра	protactinium	231.0
58	Ce	cerium 140.1	06	Т	thorium	232.0
22	Га	anthanum 138.9	89	Ac	actinium	-

lanthanoids actinoids

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.