



Cambridge International AS & A Level

CANDIDATE NAME		
CENTRE NUMBER	CANDIDATE NUMBER	

Paper 3 Advanced Practical Skills 1

May/June 2025

2 hours

9701/37

You must answer on the question paper.

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

INSTRUCTIONS

CHEMISTRY

- 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.

Session
Laboratory

For Examiner's Use		
1		
2		
3		
Total		

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

Quantitative analysis

Read through the whole method before starting any practical work. Where appropriate, prepare a table for your results in the space provided.

2

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

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

1 Hydrogen peroxide, H₂O₂, reacts rapidly with acidified potassium manganate(VII), KMnO₄.

$$5H_2O_2(aq) + 2MnO_4^-(aq) + 6H^+(aq) \rightarrow 5O_2(g) + 8H_2O(I) + 2Mn^{2+}(aq)$$

You will determine the concentration of a solution of hydrogen peroxide. You will first dilute the solution and then carry out a titration with acidified potassium manganate(VII).

FA 1 is aqueous hydrogen peroxide, H₂O₂.

FA 2 is 0.0300 mol dm⁻³ potassium manganate(VII), KMnO₄.

FA 3 is $1.0 \,\mathrm{mol}\,\mathrm{dm}^{-3}$ sulfuric acid, $\mathrm{H}_2\mathrm{SO}_4$.

(a) Method

Dilution of FA 1

- Pipette 25.0 cm³ of FA 1 into the 250 cm³ volumetric flask.
- Add distilled water to make 250 cm³ of solution.
- Shake the flask thoroughly.
- Label this diluted solution of hydrogen peroxide FA 4.

Titration

- Fill the burette with FA 2.
- Rinse the 25.0 cm³ pipette with distilled water and then with **FA 4**.
- Pipette 25.0 cm³ of FA 4 into a conical flask.
- Use the 25 cm³ measuring cylinder to add 10 cm³ of **FA 3** to the conical flask.
- Perform a rough titration and record your burette readings in the space below.

The rough titre is = \dots cm³.

- Carry out as many accurate titrations as you think necessary to obtain consistent results.
- Make sure your recorded results show the precision of your practical work.
- Record, in a suitable form in the space for results, all your burette readings and the volume of FA 2 added in each accurate titration.

Keep FA 1 for use in Questions 2 and 3. Keep FA 2 for use in Question 3.





Results

I	
II	
III	
IV	
V	
VI	
VII	

[7]

(b) From your accurate titration results, calculate a suitable mean value to use in your calculations. Show clearly how you obtain the mean value.

3

- (c) Calculations
 - (i) Give your answers to (c)(ii), (c)(iii) and (c)(iv) to an appropriate number of significant figures.
 - (ii) Calculate the amount, in mol, of manganate(VII) ions in the volume of FA 2 in (b).

amount of
$$MnO_4^- = \dots mol [1]$$

(iii) Calculate the amount, in mol, of hydrogen peroxide in $25.0\,\mathrm{cm}^3$ of FA 4.

amount of
$$H_2O_2$$
 = mol [1]

(iv) Calculate the concentration, in mol dm⁻³, of hydrogen peroxide in **FA 1**.

concentration of
$$H_2O_2$$
 in **FA 1** = mol dm⁻³ [1]

(d) A student suggests that the experiment would be more accurate if a 10 cm³ pipette is used to measure **FA 3** in place of the measuring cylinder.

State whether the student's suggestion is correct. Explain your answer. Include a calculation as part of your explanation.

		[2]

[Total: 14]



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2 You will now determine the concentration of a solution of hydrogen peroxide by a different method.

Hydrogen peroxide decomposes slowly into water and oxygen at room temperature. This reaction is exothermic. When a catalyst is added, the decomposition is fast and there is a measurable temperature rise.

$$H_2O_2(aq) \rightarrow H_2O(l) + \frac{1}{2}O_2(g)$$
 $\Delta H = -98.2 \text{ kJ mol}^{-1}$

FA 1 is aqueous hydrogen peroxide, H_2O_2 . **FA 5** is manganese(IV) oxide, MnO_2 .

(a) Method

Experiment 1

- Support one of the cups in the 250 cm³ beaker.
- Use the 50 cm³ measuring cylinder to add 25.0 cm³ of **FA 1** to the cup.
- Place the thermometer in the **FA 1** and tilt the cup, if necessary, so that the bulb of the thermometer is fully covered. Record the temperature in the space for results.
- Add a heaped spatula measure of FA 5 to the solution in the cup.
- Stir constantly until the maximum temperature is reached. Record this temperature.
- Calculate and record the temperature rise.
- · Rinse and dry the thermometer.

Experiment 2

- Support the second cup in the 250 cm³ beaker.
- Use the 50 cm³ measuring cylinder to add 40.0 cm³ of **FA 1** to the second cup.
- Measure and record the initial temperature of the solution.
- Add a heaped spatula measure of **FA 5** to the solution in the second cup.
- Stir constantly until the maximum temperature is reached. Record this temperature.
- Calculate and record the temperature rise.

Keep FA 5 for use in Question 3.

Results

I	
II	
III	
IV	
V	

[5]





(b) Calculations

(i) Calculate the energy change, in J, in Experiment 2.

(ii) Use the information given and your answer to (b)(i) to calculate the concentration, in mol dm⁻³, of hydrogen peroxide in FA 1.

7

concentration of
$$H_2O_2$$
 in **FA 1** = mol dm⁻³ [2]

(c) The concentration of hydrogen peroxide in **FA 1** calculated using the method given for Question 1 is more accurate than that using the method given for Question 2.

(i)	Heat loss is a large source of error when carrying out the method for Question 2. Describe and explain the effect of heat loss on the value of the concentration of hydrogen peroxide calculated.

[1]

(ii) A student suggests that calculating the concentration of hydrogen peroxide using the method in Question 2 would be less accurate when the concentration is lower.

Suggest whether the student is correct. Explain your answer.

ouggest mistrer the	z etagent le contect. Explain year anomen	
		[1]



[Total: 10]



[2]



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
- 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.

J	(a)	(1)	aqueous sodium hydroxide followed by a small spatula measure of FA 5 , MnO ₂ . Stir for approximately 30 seconds then filter the mixture into a second test-tube.
			observations
			Add dilute sulfuric acid to the filtrate until no further change.
			observations
			[2]
		(ii)	Transfer a 1 cm depth of aqueous iron(II) sulfate into a boiling tube. Add the same depth of dilute sulfuric acid followed by a very small spatula measure of FA 5 , MnO_2 . Carefully warm the mixture using a Bunsen burner for about 20 seconds. Filter the warm mixture into a test-tube.
			observations
			Add aqueous sodium hydroxide dropwise to the filtrate until no further change.
			observations



(111)	(a)(ii).	s in
	FA 5 is acting as	[1]
(iv)	Write an ionic equation for the reaction between aqueous sodium hydroxide and filtrate in (a)(ii). Include state symbols.	the
		[1]

Question 3 continues on page 10.



(b) FA 6 and FA 7 are both aqueous solutions of salts. Neither solution includes an ion that contains sulfur.

FA 6 contains two cations and one anion. Two of the ions are listed in the Qualitative analysis notes.

FA 7 contains one cation and one anion. One of the ions is listed in the Qualitative analysis notes.

(i) Carry out the following tests and record your observations in Table 3.1.

Use a 1 cm depth of **FA 6** or **FA 7** in a boiling tube for Test 1. Use a 1 cm depth of **FA 6** or **FA 7** in a test-tube for Tests 2, 3 and 4.

Table 3.1

toot	observ	vations
test	FA 6	FA 7
Test 1 Add aqueous sodium hydroxide, then		
warm gently.		
Test 2 Add acidified aqueous KMnO ₄ dropwise until no further change.		
Test 3 Add an equal volume of FA 1 , H_2O_2 , then		
add a few drops of starch solution.		
Test 4 Add aqueous sodium carbonate.		

* 0000800000011 *

i) The anion in **FA** 6 does **not** contain nitrogen. Select **one** further reagent to identify the anion present in **FA** 6.

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Carry out a test with this reagent and record your observations in Table 3.2.

Table 3.2

reagent	observations

[1]

(iii) Give the formulae of the ions present in **FA 6** and **FA 7**. If you are unable to identify an ion from your tests, write 'unknown.'

The ions present in FA 6 are and and and

The ions present in **FA 7** are and and

[3]

[Total: 16]

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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

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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	$L = 6.02 \times 10^{23} \mathrm{mol}^{-1}$
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

	18	2	D C	helium 4.0	10	Ne	neon 20.2	18	Ā	argon 39.9	36	궃	krypton 83.8	25	Xe	xenon 131.3	98	R	radon	118	Og	oganesson	
	17				6	ш	fluorine 19.0	17	Cl	chlorine 35.5	35	ъ	bromine 79.9	53	П	iodine 126.9	82	Ąŧ	astatine	117	<u>s</u>	tennessine -	
	16				80	0	oxygen 16.0	16	S	sulfur 32.1	34	Se	selenium 79.0	52	<u>е</u>	tellurium 127.6	84	Ъо	moloulum	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>.</u>	bismuth 209.0	115	Mc	moscovium	
	41				9	O	carbon 12.0	14	S	silicon 28.1	32	Ge	germanium 72.6	20	S	tin 118.7	82	Ъ	lead 207.2	114	Εl	flerovium	
	13				2	Ф	boron 10.8	13	Αl	aluminium 27.0	31	Ga	gallium 69.7	49	In	indium 114.8	84	11	thallium 204.4	113	R	nihonium	
										12	30	Zn	zinc 65.4	48	පි	cadmium 112.4	80	£	mercury 200.6	112	ပ်	copernicium	
										7	59	Cn	copper 63.5	47	Ag	silver 107.9	62	Αu	gold 197.0	111	Rg	roentgenium	
dno										10	28	z	nickel 58.7	46	Pd	palladium 106.4	78	చ	platinum 195.1	110	Ds	darmstadtium -	
Group										6	27	ပိ	cobalt 58.9	45	뫈	rhodium 102.9	77	'n	iridium 192.2	109	₩	meitnerium -	
		-]	С	hydrogen 1.0						œ	26	Ъе	iron 55.8	44	Ru	ruthenium 101.1	9/	Os	osmium 190.2	108	Ϋ́	hassium -	
										_	25	Mn	manganese 54.9	43	ည	technetium -	75	Re	rhenium 186.2	107	Bh	bohrium –	
					_	pol	SS			9	24	ပ်	chromium 52.0	42	Mo	molybdenum 95.9	74	≥	tungsten 183.8	106	Sg	seaborgium	
				Key	atomic number	atomic symbol	name relative atomic mass			2	23	>	vanadium 50.9	14	g	niobium 92.9	73	<u>⊾</u>	tantalum 180.9	105	9	dubnium –	
						atc	2			4	22	F	titanium 47.9	40	Zr	zirconium 91.2	72	Ξ	hafnium 178.5	104	¥	rutherfordium -	
										က	21	Sc	scandium 45.0	39	>	yttrium 88.9	57-71	lanthanoids		89–103	actinoids		
	2				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	
	~				3	:=	lithium 6.9	=	Na	sodium 23.0	19	\prec	potassium 39.1	37	Вb	rubidium 85.5	55	S	caesium 132.9	87	<u>ن</u>	francium -	

1.1	lutetium 175.0	103	۲	lawrencium	ı
07 X	ytterbium 173.1	102	No	nobelium	ı
69 F	thulium	101	Md	mendelevium	1
68 г	erbium	100	Fm	ferminm	I
⁶⁷	holmium 254 9	66	Es	einsteinium	ı
99	dysprosium	98	Ç	californium	ı
65 Th	terbium	97	Ř	berkelium	ı
م را م	gadolinium 157.3	96	Cm	curium	ı
63 H-1	europium	95	Am	americium	ı
62 0	samarium 150.4	94	Pu	plutonium	ı
61	promethium	93	ď	neptunium	ı
09	neodymium 144.2	92	\supset	uranium	238.0
88 9	praseodymium	91	Ра	protactinium	231.0
85 C	Cerium 140 1	06	H	thorium	232.0
57	anthanum	89	Ac	actinium	ı

lanthanoids

actinoids

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