

Cambridge International AS & A Level

CANDIDATE NAME			
CENTRE NUMBER		CANDIDATE NUMBER	
CHEMICTRY			0704/2

CHEMISTRY

9701/33

Paper 3 Advanced Practical Skills 1

February/March 2020

2 hours

You must answer on the question paper.

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

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, use appropriate units and use an appropriate number of significant figures.
- Give details of the practical session and laboratory, where appropriate, in the boxes provided.

Session		
Laborat	ory	

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.
- Notes for use in qualitative analysis are provided in the question paper.

For Examiner's Use				
1				
2				
3				
Total				

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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 The concentrations of solutions of hydrogen peroxide are often represented in terms of 'volume strength'. In this experiment you will determine the 'volume strength' of a solution of hydrogen peroxide by titration with acidified potassium manganate(VII).

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

FA 1 is 0.0300 mol dm⁻³ potassium manganate(VII), KMnO₄.

FA 2 is dilute sulfuric acid, H₂SO₄.

FA 3 is aqueous hydrogen peroxide, H_2O_2 .

(a) Method

Dilution of FA 3

- Pipette 25.0 cm³ of FA 3 into the 250 cm³ volumetric flask.
- Make the solution up to the mark using distilled water.
- Shake the flask thoroughly.
- Label this diluted solution of hydrogen peroxide FA 4.

Titration

- Fill the burette with FA 1.
- Rinse the pipette thoroughly with distilled water and then with a little FA 4.
- Pipette 25.0 cm³ of FA 4 into a conical flask.
- Use the 25 cm³ measuring cylinder to add 20 cm³ of **FA 2** into the same conical flask.
- Perform a rough titration and record your burette readings in the space below.

				_
The	rough	titre is	S	 cm ³

- Carry out as many accurate titrations as you think necessary to obtain consistent results.
- Make sure any recorded results show the precision of your practical work.
- Record all of your burette readings and the volume of FA 1 added in each accurate titration.

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

I	
II	
III	
IV	
V	
VI	
VII	

[7]







(b)	in	om your accurate titration results, obtain a suitable value for the volume of FA 1 to be used your calculations. ow clearly how you obtained this value.
		25.0 cm ³ of FA 4 required cm ³ of FA 1 . [1]
(c)	Ca	lculations
	(i)	Give your answers to (ii), (iii), (iv) and (v) to the appropriate number of significant figures. [1]
	(ii)	Calculate the number of moles of potassium manganate (VII) present in the volume calculated in (b).
		moles of KMnO ₄ = mol [1]
	(iii)	The equation for the reaction of potassium manganate(VII) with hydrogen peroxide is shown.
		$2MnO_4^-(aq) + 5H_2O_2(aq) + 6H^+(aq) \rightarrow 2Mn^{2+}(aq) + 8H_2O(I) + 5O_2(g)$
		Use your answer to (c)(ii) to calculate the number of moles of hydrogen peroxide used in each titration.
		moles of $H_2O_2 = \dots mol$
		Hence calculate the concentration of H ₂ O ₂ in FA 4 , in mol dm ⁻³ .
		concentration of H_2O_2 in FA 4 = mol dm ⁻³ [1]

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(iv)	Calculate	the concent	ration o	of H ₂ O ₂ i	in FA 3 .	in moldm ⁻³

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

(v) When hydrogen peroxide decomposes in the presence of a catalyst, oxygen is produced.

$$H_2O_2(aq) \rightarrow H_2O(1) + \frac{1}{2}O_2(g)$$

The 'volume strength' of hydrogen peroxide is equal to the volume of oxygen, in dm³, produced under room conditions, when 1.00 dm³ of the solution decomposes.

Use your answer to **(c)(iv)** and the equation above to calculate the volume, in dm³, of oxygen produced when 1.00 dm³ of **FA 3** decomposes. This is the 'volume strength', in vol, of **FA 3**.

(Under room conditions 1.00 mol of gas occupies a volume of 24.0 dm³. If you were unable to calculate the concentration of H_2O_2 in **FA 3**, assume that it is 1.02 mol dm⁻³. This may **not** be the correct value.)

(d) The maximum error in reading a 25.0 cm³ pipette is ±0.06 cm³.

Show by calculation that the pipette is more accurate than a burette for measuring $25.0\,\mathrm{cm^3}$ of solution.

[1]

[Total: 15]







2 In this experiment you will determine the enthalpy change, ΔH , for the catalytic decomposition of hydrogen peroxide into water and oxygen.

$$H_2O_2(aq) \rightarrow H_2O(I) + \frac{1}{2}O_2(g)$$

FA 3 is aqueous hydrogen peroxide, H_2O_2 . **FA 5** is manganese(IV) oxide, MnO₂.

(a) Method

Experiment 1

- Support one of the plastic cups inside the 250 cm³ beaker.
- Use the 50 cm³ measuring cylinder to add 30 cm³ of **FA 3** into the plastic cup.
- Measure and record the initial temperature of the solution.
- Add a heaped spatula measure of **FA 5** to the solution in the plastic cup.
- Stir constantly until the maximum temperature is reached and record this temperature.
- Calculate and record the temperature rise.
- Rinse and dry the thermometer.

Experiment 2

- Support the second plastic cup inside the 250 cm³ beaker.
- Use the 50 cm³ measuring cylinder to add 40 cm³ of **FA 3** into the plastic cup.
- Measure and record the initial temperature of the solution.
- Add a heaped spatula measure of **FA 5** to the solution in the plastic cup.
- Stir constantly until the maximum temperature is reached and record this temperature.
- Calculate and record the temperature rise.



[5]

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(i)	Calculate the energy released in Experiment 1 . [Assume that 4.2 J changes the temperature of 1.0 cm³ of solution by 1.0 °C.]
(ii)	energy released =
(iii)	moles of $\rm H_2O_2$ =
	enthalpy change = kJ mol ⁻¹ [1] sign value
(c) (i)	A student suggested that the experiment would be more accurate if the same mass of ${\bf FA}$ 5, manganese(IV) oxide, had been weighed out for each experiment.
(ii)	State and explain whether you agree with the student's suggestion. [2] The student also suggested that Experiments 1 and 2 should give the same temperature rise, even though a greater volume of FA 3 was used in Experiment 2. State and explain whether you agree with the student's suggestion.
	State and explain whether you agree with the student's suggestion.
	[1] [Total: 11]
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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

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- 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.

- (a) FA 3 is aqueous hydrogen peroxide, H₂O₂.
 FA 6 is a solution containing two cations and one anion from those listed in the Qualitative analysis notes.
 - (i) To a 1 cm depth of **FA 6** in a boiling tube, add aqueous sodium hydroxide until it is in excess. Then heat the tube, gently and carefully. Keep the mixture obtained in the boiling tube for the test in (a)(ii).

	Record all your observations. Identify the cations in FA 6 .	
	observations	
	FA 6: cations are and	[4]
(ii)	To the mixture obtained from (a)(i), carefully add a 1 cm depth of FA 3. Record your observations.	
		[1]
iii)	One reaction taking place in (a)(ii) involves oxidation of one of the cations in FA 6.	
	Give the half-equation to show this oxidation reaction. State symbols are not required.	
		[1]

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(b) FA 1 is aqueous potassium manganate(VII).

FA 2 is dilute sulfuric acid.

FA 7 and FA 8 are solutions, each containing one cation and one anion.

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

test	observations			
iesi	FA 7	FA 8		
Test 1 To a 1 cm depth of solution in a test-tube, add a small spatula measure of solid sodium carbonate.				
Test 2 To a 1 cm depth of solution in a test-tube, add an equal volume of FA 2 and a few drops of FA 1, then				
add a few drops of aqueous starch.				
Test 3 To a 1 cm depth of solution in a test-tube, add a few drops of aqueous silver nitrate, then				
add aqueous ammonia.				

		[5]
(ii)	Identify the anion in FA 7 .	
	anion	. [1]
(iii)	Identify FA 8.	
	FA 8 is	. [1]
(iv)	Carry out one further test to confirm the identity of the cation in FA 8 . State the name of the reagent you used and record the observation(s) you made.	
	reagent	
	observation(s)	
		[1]
	[Tota	: 14]







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Qualitative analysis notes

1 Reactions of aqueous cations

inn	reaction with								
ion	NaOH(aq)	NH ₃ (aq)							
aluminium, Al³+(aq)	white ppt. soluble in excess	white ppt. insoluble in excess							
ammonium, NH₄⁺(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 [Ca²+(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	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), 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					



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	18	2 H	helium 4.0	10	Ne	neon 20.2	18	Ā	argon 39.9	36	호	krypton	54	Xe	xenon 131.3	86	R	radon			
	17			6	ш	fluorine 19.0	17	Cl	chlorine 35.5	35	Ŗ	bromine	53	Ι	iodine 126.9	85	¥	astatine			
	16			80	0	oxygen 16.0	16		sulfur 32.1		Se	selenium	79.0	Те	tellurium 127.6	8	Ро	polonium	116	^	live morium –
	15			7	z	nitrogen 14.0	15	Д	phosphorus 31.0	33	As	arsenic 74.0	51	Sb	antimony 121.8	83	ï	bismuth 209.0			
	14			9	ပ	carbon 12.0	14	Si	silicon 28.1	32	Ge	germanium	50	Sn	tin 118.7	82	Ъ	lead 207.2	114	Εl	flerovium -
	13			5	В	boron 10.8	13	Αl	aluminium 27.0	31	Ga	gallium	49	In	indium 114.8	81	11	thallium 204.4			
									12	30	Zu	zinc	4.8	g	cadmium 112.4	80	Η̈́	mercury 200.6	112	ပ်	copernicium
									7	29	ರ	copper	47	Ag	silver 107.9	62	Au	gold 197.0	111	ß	roentgenium -
Group									10	28	Z	nickel	30.7	Pd	palladium 106.4	78	Ŧ	platinum 195.1	110	Ds	darmstadtium -
Gre									6	27	ပိ	cobalt	45	R	modium 102.9	11	'n	iridium 192.2	109	Mŧ	meitnerium -
		- I	hydrogen 1.0						80	56	Ьe	iron	0.00	Ru	ruthenium 101.1	9/	Os	osmium 190.2	108	Η̈́	hassium -
									7	25	Mn	manganese	94.9		technetium -		Re	rhenium 186.2	107	B	pohrium –
					pol	ass			9	24	ပ်	chromium	92.0	Mo	molybdenum 95.9	74	≥	tungsten 183.8	106	Sg	seaborgium -
			Key	atomic number	tomic symbol	name elative atomic mass			2	23	>	vanadium	90.9	qN	midolin 92.9	73	<u>ra</u>	tantalum 180.9	105	Ор	dubnium -
					ato	rela			4	22	i=	titanium	40	Zr	zirconium 91.2	72	Ξ	hafnium 178.5	104	짪	rutherfordium -
							-		က	21	လွ	scandium	39	>	yttrium 88.9	57-71	lanthanoids		89-103	actinoids	
	2			4	Be	beryllium 9.0	12	Mg	magnesium 24.3	20	Sa	calcium 40.4	38	ഗ്	strontium 87.6	56	Ba	barium 137.3	88	Ra	radium
	_			3	:=	lithium 6.9	11	Na	sodium 23.0	19	¥	potassium	37	Rb	rubidium 85,5	55	Cs	caesium 132.9	87	ъ́	francium -

71	3	lutetium 175.0	103	۲	rencium	1	
		= `			law		
20	Υb	ytterbium 173.1	102	Š	nobelium	ı	
69	Tm	thulium 168.9	101	Md	mendelevium	ı	
89	ш	erbium 167.3	100	Fm	fermium	ı	
29	웃	holmium 164.9	66	Es	einsteinium	ı	
99	ò	dysprosium 162.5	86	ర	californium	ı	
65	Д	terbium 158.9	26	益	berkelium	ı	
64	В	gadolinium 157.3	96	Cm	curium	ı	
63	En	europium 152.0	92	Am	americium	ı	
62	Sm	samarium 150.4	94	Pu	plutonium	ı	
61	Pm	promethium -	93	dN	neptunium	ı	
		neodymium 144.4					
59	Ā	praseodymium 140.9	91	Ра	protactinium	231.0	
58	Ce	cerium 140.1	06	드	thorium	232.0	
22	Га	anthanum 138.9	68	Ac	actinium	ı	

lanthanoids

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