

Cambridge International AS & A Level

CANDIDATE NAME					
CENTRE NUMBER			CANDIDATE NUMBER		

CHEMISTRY 9701/23

Paper 2 AS Level Structured Questions

October/November 2023

1 hour 15 minutes

You must answer on the question paper.

No additional materials are needed.

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 60.
- 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.

This document has 20 pages. Any blank pages are indicated.

1 The elements phosphorus, sulfur and chlorine are in Period 3 of the Periodic Table.

Table 1.1 shows some properties of the elements P to Cl.

The first ionisation energy of S is **not** shown.

Table 1.1

property	Р	S	Cl
number of electrons in 3p subshell			
total number of unpaired electrons			
first ionisation energy /kJ mol ⁻¹	1060		1260
formula of most common anion	P ³⁻	S ²⁻	Cl ⁻

(a) (Complete Table 1.1 to shoumber of unpaired electron		ectrons in the 3p subshell a and $\mathrm{C}l$.	and the tota [2]
(i	i) (Construct an equation to re	epresent the first ionis	ation energy of P.	
					[1]
(ii	i) -	Three possible values for t	the first ionisation ene	rgy of S are given.	
		1000 kJ mol ⁻¹	1160 kJ mol ⁻¹	1320 kJ mol ⁻¹	
	(Circle the correct value.			
	I	Explain your choice by cor	mparing your chosen v	value to those of P and C1.	
					L4.

(b)	P ³⁻ ,	S ²⁻ and C <i>l</i> ⁻ have the same number of electrons.
	(i)	Give the full electronic configuration of P ^{3–} .
		[1]
	(ii)	State the trend in ionic radius shown by P^{3-} , S^{2-} and Cl^{-} .
		Explain your answer.
		[2]
(c)	A st	udent does three tests on separate samples of NaCl(aq).
	Con	nplete Table 1.2 with the observations the student makes in each test.

Table 1.2

test	test	observations
1	addition of a few drops of Br ₂ (aq)	
2	addition of a few drops of concentrated H ₂ SO ₄	
3	addition of a few drops of dilute AgNO ₃ (aq)	

[3]

(d) $POCl_3$ shows similar chemical properties to PCl_5 .

(i)

(ii)

 $POCl_3$ has a melting point of 1°C and a boiling point of 106°C.

 $\mathsf{POC}\mathit{l}_{\!3}$ reacts vigorously with water, forming misty fumes and an acidic solution.

Explain how the information in (d) suggests the structure and bonding of $POCl_3$ is simple covalent.
[2
Construct an equation for the reaction of $POCl_3$ with water.
$POCl_3$ +

(iii) $POCl_3$ contains a double covalent bond between P and O.

Complete the dot-and-cross diagram, in Fig. 1.1, to show the bonding in $POCl_3$. Show outer shell electrons only.

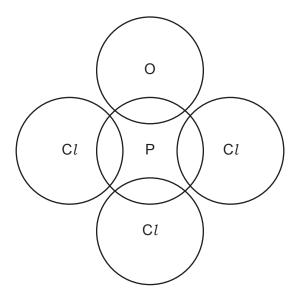


Fig. 1.1

[2]

(e) $POCl_3(g)$ forms when $PCl_3(g)$ reacts with $O_2(g)$.

$$2\mathsf{PC}\mathit{l}_{3}(\mathsf{g}) + \mathsf{O}_{2}(\mathsf{g}) \, \longrightarrow \, 2\mathsf{POC}\mathit{l}_{3}(\mathsf{g})$$

Table 1.3 gives some relevant data.

Table 1.3

process	value / kJ mol ^{−1}
enthalpy change of formation of $PCl_3(g)$	-289
enthalpy change of formation of $POCl_3(g)$	-592
$O_2(g) \rightarrow 2O(g)$	+496

(i)	Define enthalpy change of formation, $\Delta H_{\rm f}$.	
		[2
(ii)	Calculate the bond energy of P=O in ${ m POC}l_3$ using the data in Table 1.3.	
	Show your working.	

bond energy of P=O =	kJ mol ⁻¹
	[2]

[Total: 22]

2 Barium hydroxide, $Ba(OH)_2$, is a strong base used in inorganic and organic reactions.

Fig. 2.1 shows a reaction scheme involving $Ba(OH)_2$.

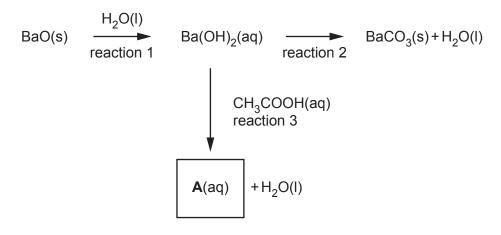


Fig. 2.1

(a)	(i)	State the variation in solubilities of group 2 hydroxides.	
			[1]
((ii)	State what is observed in reaction 1.	
			[1]
(i	iii)	Suggest a reactant for reaction 2.	
			[1]
(i	iv)	Identify A.	
			[1]
((v)	Ba(OH) ₂ is made by the reaction of Ba with water.	
		Write an equation for this reaction.	
			[1]

State which compound decomposes first when barytocalcite is heated.

(b)	The mineral barytocalcite contains both BaCO ₃ and CaCO ₃ . Both compounds decompose on
	heating.

Explain your answer.

.....

(ii) Construct an equation for the complete thermal decomposition of barytocalcite.

The formula of barytocalcite is $BaCa(CO_3)_2$.

 $BaCa(CO_3)_2$ [1]

(c) $Ba(OH)_2$ is used to hydrolyse organic compounds.

Fig. 2.2 shows the reaction of **B** with $Ba(OH)_2$, followed by acidification.

Draw the structures of the organic products of the process shown in Fig. 2.2.

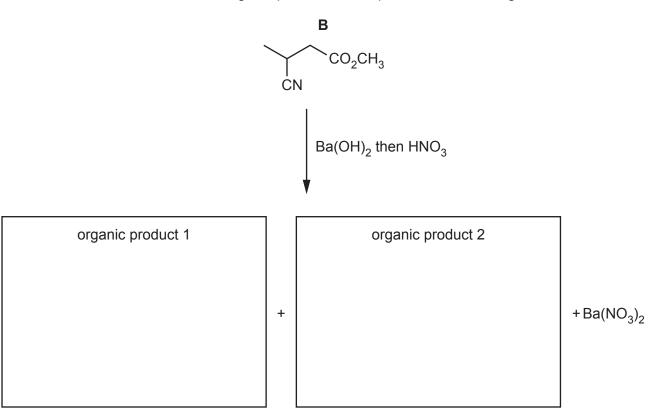


Fig. 2.2

[3]

[Total: 10]

(a)	Define oxidising ager	nt.	
			נין
(b)	KClO ₃ (s) decompose	es when heated.	
	MnO ₂ (s) catalyses th	e exothermic decomposition reaction.	
	Complete and label the of $KClO_3(s)$.	ne diagram in Fig. 3.1 to show the effect of MnO ₂ (s) on the decomposit	ion
	enthalpy		
		reactants	
		products	
	Į	progress of reaction	
		Fig. 3.1	
			[2]
(c)	When KClO ₃ is heate	ed without a catalyst, $KClO_4$ and KCl form.	
	2	$KClO_3 \rightarrow 3KClO_4 + KCl$	
	Explain why this read	tion is described as a disproportionation reaction.	

(d) Molten $KClO_3$ reacts with glucose, $C_6H_{12}O_6$.

$$4KClO_3 + C_6H_{12}O_6 \rightarrow 6CO_2 + 6H_2O + 4KCl$$

 $\mathrm{KC}l\mathrm{O}_3$ melts at 630 K. At this temperature, both CO_2 and $\mathrm{H}_2\mathrm{O}$ are gases.

(i) Use the ideal gas equation to calculate the volume, in m^3 , of one mole of gas at 630 K and 1.00×10^5 Pa.

Show your working. Give your answer to 3 significant figures.

volume of 1 mol of gas =
$$\dots$$
 m³

(ii) 5.00 g of $C_6H_{12}O_6$ reacts completely with molten $KClO_3$.

Use your answer to (d)(i) to calculate the total volume of gas released at 630 K and 1.00×10^5 Pa in this reaction.

(If you were unable to answer (d)(i), use $0.0463\,\text{m}^3$ in this question. This is **not** the correct answer to (d)(i).)

(e) The structure of glucose, $C_6H_{12}O_6$, is shown in Fig. 3.2.

glucose

Fig. 3.2

(i) Complete Table 3.1 to identify the number of primary, secondary and tertiary alcohol groups present in the structure shown in Fig. 3.2.

Table 3.1

type of alcohol group	primary	secondary	tertiary
number of groups			

[1]

(ii) Separate samples of aqueous glucose are tested with the reagents shown in Table 3.2.

Complete Table 3.2 with the observation for each reaction.

Write "no reaction" if applicable.

Table 3.2

reagent and conditions	observation with glucose
acidified KMnO ₄ (aq) and warm	
Fehling's reagent and warm	
alkaline I ₂ (aq) and warm	

	٦
٠,	

(iii) There are many structural isomers of $\mathrm{C_6H_{12}O_6}$.

Define structural isomers.

.....

[Total: 12]

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4 Compounds **C** and **D** are alkenes with the same molecular formula, C_5H_{10} .



Fig. 4.1

(a) (i)	Give the systematic name of D .	
(ii)		. [1]
(iii)	Draw the structure of a molecule that is a positional isomer of C and D .	. [1]

(iv) Give the structural formula of the compound formed when ${\bf D}$ reacts with ${\bf H_2}({\bf g})$ in the presence of a Pt catalyst.

______[1]

(v) C can form an addition polymer.

Draw the structure of **one** repeat unit of this addition polymer.

[1]

[1]

(b) The mass spectrum of $\bf C$ shows a molecular ion peak at m/e = 70. This peak has a relative intensity of 48.7.

The relative intensity of the [M+1] peak is 2.7.

Show that this information is consistent with the molecular formula of **C**.

[2]

- (c) C and D both react with HBr.
 - (i) C reacts with HBr to form E.

Complete the diagram in Fig. 4.2 to show the mechanism for this reaction.

Draw the structure of the organic intermediate.

Include charges, dipoles, lone pairs of electrons and curly arrows, as appropriate.

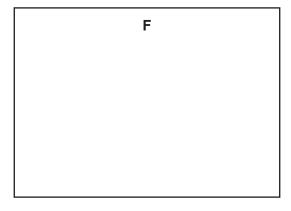


Fig. 4.2

[3]

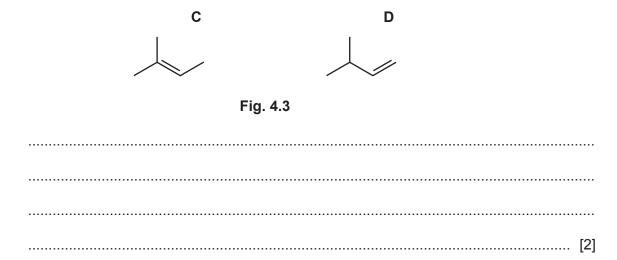
(ii) **D** reacts with HBr to produce **F**, a chiral bromoalkane.

Draw the structure of **F**.



[1]

(iii) Explain why the reaction of HBr with **C** and **D** produces different major products.

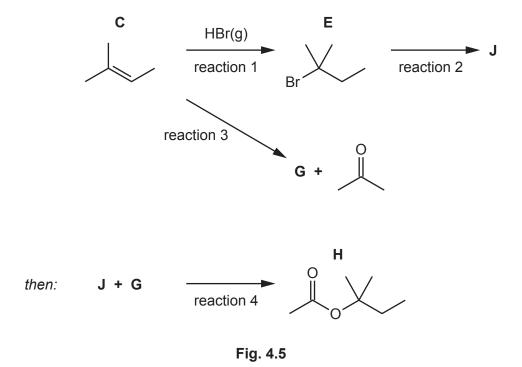


(d) C can be used to form H.

Fig. 4.4

One possible synthesis of ${\bf H}$ is shown in Fig. 4.5. Different portions of ${\bf C}$ are used in reactions 1 and 3. Some of the products are then combined to produce ${\bf H}$.

Fig. 4.5 does not show any of the inorganic products of the reactions.



Complete Table 4.1 with the reagents and conditions required for each of the reactions shown in Fig. 4.5.

Table 4.1

		reagent and conditions
reaction 1	c o e	HBr(g)
reaction 2	$ extsf{E} ightarrow extsf{J}$	
reaction 3	$c \rightarrow g + \bigcirc$	
reaction 4	$ extsf{J} + extsf{G} ightarrow extsf{H}$	

[3]

[Total: 16]

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17

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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	$L = 6.022 \times 10^{23} \text{mol}^{-1}$
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

								T						_									uo.	
	18	2	He	helium 4.0	10	Ne	neon	70.7	9	Ā	argon 39.9	36	궃	kryptor 83.8	72	Xe	xenon 131.3	98	R	radon	118	O	oganess	
	16 17				6	ш	fluorine	0.81	17	Cl	chlorine 35.5	35	Ā	bromine 79.9	53	н	iodine 126.9	85	¥	astatine -	117	<u>s</u>	tennessine -	
					80	0	oxygen	16.0	16	ഗ	sulfur 32.1	8	Se	selenium 79.0	52	<u>e</u>	tellurium 127.6	28	Ъо	polonium –	116	_	livermorium -	
	15				7	z	nitrogen	14.0	15	ட	phosphorus 31.0	33	As	arsenic 74.9	51	Sb	antimony 121.8	83	Ξ	bismuth 209.0	115	Mc	moscovium	
	14				9	ပ	carbon	12.0	4	S	silicon 28.1	32	Ge	germanium 72.6	50	Sn	tin 118.7	82	Ър	lead 207.2	114	Εl	flerovium	
	13				2	В	boron	10.8	13	Ρl	aluminium 27.0	31	Ga	gallium 69.7	49	Г	indium 114.8	18	11	thallium 204.4	113	R	nihonium	
											12	30	Zn	zinc 65.4	48	рO	cadmium 112.4	80	Hg	mercury 200.6	112	ű	copernicium	
											7	59	J	copper 63.5	47	Ag	silver 107.9	62	Αu	gold 197.0	111	Rg	roentgenium	
dn											10	28	Z	nickel 58.7	46	Pd	palladium 106.4	78	చ	platinum 195.1	110	Ds	darmstadtium -	
Group												6	27	ဝိ	cobalt 58.9	45	R	rhodium 102.9	77	'n	iridium 192.2	109	Μţ	meitnerium -
		-	I	hydrogen 1.0							80	56	Fe	iron 55.8	44	R	ruthenium 101.1	9/	SO	osmium 190.2	108	Η̈́	hassium	
				ı						7	25	Mn	manganese 54.9	43	ည	technetium -	75	Re	rhenium 186.2	107	Bh	bohrium		
						loc		SS			9	24	ဝ်	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	qN	niobium 92.9	73	<u>ra</u>	tantalum 180.9	105	o O	dubnium	
					В	atoi		rela			4	22	F	titanium 47.9	40	Zr	zirconium 91.2	72	士	hafnium 178.5	104	쪼	rutherfordium -	
								_			က	21	Sc	scandium 45.0	39	>	yttrium 88.9	57–71	lanthanoids		89–103	actinoids		
	2				4	Be	beryllium	0.0	12	Mg	magnesium 24.3	20	Ca	calcium 40.1	38	S	strontium 87.6	56	Ba	barium 137.3	88	Ra	radium	
	-				3	=	lithium	6.9	=	Na	sodium 23.0	19	×	potassium 39.1	37	윉	rubidium 85.5	55	S	caesium 132.9	87	ъ.	francium	

71	Ľ	lutetium 175.0	103	Ļ	lawrencium	I	
20	Υp	ytterbium 173.1	102	Š	nobelium	_	
69	٦	thulium 168.9	101	Md	mendelevium	Ι	
68	ш	erbium 167.3	100	Fm	fermium	_	
29	운	holmium 164.9	66	Es	einsteinium	-	
99	ò	dysprosium 162.5	86	Ç	californium	1	
65	Д	terbium 158.9	26	Ř	berkelium	_	
64	В	gadolinium 157.3	96	Cm	curium	_	
63	En	europium 152.0	92	Am	americium	_	
62	Sm	samarium 150.4	96	Pu	plutonium	_	
61	Pm	promethium	93	ď	neptunium	1	
09	ρN	neodymium 144.4	92	\supset	uranium	238.0	
69	Ą	praseodymium 140.9	91	Ра	protactinium	231.0	
58	Ce	cerium 140.1	06	드	thorium	232.0	
22	Гa	lanthanum 138.9	88	Ac	actinium	-	

lanthanoids

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