Vrite your name here Surname	Oth	ner names
Pearson Edexcel GCE	Centre Number	Candidate Number
Chemist		
Advanced Subsid Unit 2: Applicatio	liary	ples of Chemistry
Advanced Subsid	iary n of Core Princi Afternoon	ples of Chemistry Paper Reference 6CH02/01

Instructions

- Use **black** ink or ball-point pen.
- Fill in the boxes at the top of this page with your name, centre number and candidate number.
- Answer **all** questions.
- Answer the questions in the spaces provided there may be more space than you need.

Information

- The total mark for this paper is 80.
- The marks for **each** question are shown in brackets – use this as a guide as to how much time to spend on each question.
- Questions labelled with an asterisk (*) are ones where the quality of your written communication will be assessed
 you should take particular care with your spelling, punctuation and grammar, as well as the clarity of expression, on these questions.
- A Periodic Table is printed on the back cover of this paper.

Advice

- Read each question carefully before you start to answer it.
- Keep an eye on the time.
- Try to answer every question.
- Check your answers if you have time at the end.

Turn over 🕨







CECTION

	SECTION A			
Answer ALL the questions in this section. You should aim to spend no more than 20 minutes on this section. For each question, select one answer from A to D and put a cross in the box ⊠. If you change your mind, put a line through the box ⊠ and then mark your new answer with a cross ⊠.				
1	The bo	ond angle in beryllium chloride, BeCl ₂ , is most likely to be		
	🖾 A	90°		
	B	104.5°		
	C	120°		
	D	180°		
		(Total for Question 1 = 1 mark)		
2	•	ite and buckminsterfullerene are forms of carbon. Buckminsterfullerene ves in octane but graphite does not. This is because		
	A	the bonds between carbon atoms in buckminsterfullerene are weaker than those in graphite.		
	B	buckminsterfullerene is molecular whereas graphite is a giant structure.		
	🖾 C	graphite has delocalised electrons but buckminsterfullerene does not.		
	D	graphite has covalent bonds and London forces but buckminsterfullerene has just London forces.		
		(Total for Question 2 = 1 mark)		
3	Which	of the following molecules is polar?		
	A 🛛	Carbon dioxide, CO ₂		
	B	Silicon tetrachloride, SiCl ₄		
	🖾 C	Ammonia, NH ₃		
	D	Boron trifluoride, BF ₃		
		(Total for Question 3 = 1 mark)		

P 4 4 8 8 1 A 0 2 2 4

4	ln whi to righ	ch series of compounds does covalent character increase when going from left	
	⊠ A	KI, KBr, KCI	
	B	Nal, Kl, Rbl	
	🖾 C	NaCl, MgCl ₂ , AlCl ₃	
	🛛 D	SO ₂ , P ₄ O ₁₀ , SiO ₂	
		(Total for Question 4 = 1 ma	ark)
5	Hydro becau	gen bromide has a lower boiling temperature than hydrogen iodide. This is se	
	🖾 A	hydrogen bromide has a smaller permanent dipole than hydrogen iodide.	
	B	hydrogen bromide has weaker London forces than hydrogen iodide.	
	🖾 C	hydrogen iodide forms hydrogen bonds but hydrogen bromide does not.	
	D 🛛	the H—I bond is stronger than the H—Br bond.	
		(Total for Question 5 = 1 ma	ark)
		$\begin{bmatrix} 1 & 1 & 1 & 1 \\ P & 4 & 4 & 8 & 8 & 1 & A & 0 & 3 & 2 & 4 \end{bmatrix}$	3 Turn ove

6	Consider the following compounds, E , F , G and H .
	$\begin{array}{cccc} CH_2 & CH_3 \\ CH_2 & CH_2 \\ CH_3 & CH_3 \\ E & F \end{array}$
	$ \begin{array}{cccc} I - CH_2 & CH_3 \\ CH_2 - CH_2 & I - C - CH_3 \\ CH_3 & CH_3 \\ G & H \end{array} $
	The boiling temperature of these compounds increases in the order
	A HGFE
	B GHEF
	C EFGH
	D FEHG
	(Total for Question 6 = 1 mark)
7	In which of the following reactions is sulfuric(IV) acid, H_2SO_3 , acting as an oxidizing agent?
7	In which of the following reactions is sulfuric(IV) acid, H ₂ SO ₃ , acting as an oxidizing
7	In which of the following reactions is sulfuric(IV) acid, H_2SO_3 , acting as an oxidizing agent?
7	In which of the following reactions is sulfuric(IV) acid, H ₂ SO ₃ , acting as an oxidizing agent? \square A H ₂ SO ₃ + H ₂ O \rightarrow H ₃ O ⁺ + HSO ₃ ⁻
7	In which of the following reactions is sulfuric(IV) acid, H ₂ SO ₃ , acting as an oxidizing agent? $\square \ \mathbf{A} \ H_2SO_3 + H_2O \rightarrow H_3O^+ + HSO_3^-$ $\square \ \mathbf{B} \ H_2SO_3 \rightarrow SO_2 + H_2O$
7	In which of the following reactions is sulfuric(IV) acid, H_2SO_3 , acting as an oxidizing agent? A $H_2SO_3 + H_2O \rightarrow H_3O^+ + HSO_3^-$ B $H_2SO_3 \rightarrow SO_2 + H_2O$ C $H_2SO_3 + 2FeCI_3 + H_2O \rightarrow 2FeCI_2 + H_2SO_4 + 2HCI$
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8		of the following properties decreases on descending Group 2 of the lic Table?
	🖾 A	Solubility of the sulfates.
	B	Solubility of the hydroxides.
	🖾 C	Reactivity of the elements.
	🖾 D	lonic character of the oxides.
		(Total for Question 8 = 1 mark)
9	white	ound X is an anhydrous, white solid which decomposes on heating to form a solid residue, a colourless gas, and a colourless vapour which condenses to a less liquid.
	Comp	ound X is
	🖾 A	sodium carbonate.
	B	sodium hydrogencarbonate.
	🖾 C	sodium nitrate.
	🖾 D	sodium sulfate.
		(Total for Question 9 = 1 mark)



10 These questions concern the Maxwell-Boltzmann energy distribution shown below.



(a) What is the best way to describe the activation energy, E_{a} , of a reaction?

(1)

- A The average energy of the particles that react.
- **B** The minimum energy required for a reaction to occur.
- **C** The energy difference between the reactants and products.
- **D** The energy produced by the particles that react.
- (b) How does the curve above change when the temperature is **increased**?

(1)

- A The peak increases in height and moves to the left.
- **B** The peak increases in height and moves to the right.
- C The peak decreases in height and moves to the left.
- **D** The peak decreases in height and moves to the right.



	(Total for Question 11 = 1 mark)	
_ 2	a mass spectrum.	
🖾 D	spectrum. The compounds have molecular ion peaks at different mass to charge ratios in	
🖾 C	The compounds produce different fragmentation patterns in a mass	
B		
A	The compounds produce different patterns in the fingerprint region of the IR spectrum.	
these	compounds are compared using physical methods of analysis, which of the	
	(Total for Question 10 = 4 marks)	
🛛 D	move to the left and the peak would be unchanged.	
TI	he activation energy would (1)	
(d) W	/hat would be the effect on the diagram if a catalyst was added?	
🛛 D		
Δ		
	creased?	
	in A B C D C D C A B C D Propathese follov A C C C C C C C C C C C C C	 (d) What would be the effect on the diagram if a catalyst was added? The activation energy would (1) A be unchanged and the peak would move to the right. B move to the left and the peak would move to the right. C move to the left and the peak would move to the left. D move to the left and the peak would be unchanged. (Total for Question 10 = 4 marks) Propanal, CH ₃ CH ₂ CHO, and propanone, CH ₃ COCH ₃ , are carbonyl compounds. When these compounds are compared using physical methods of analysis, which of the following statements is not correct? A The compounds produce different patterns in the fingerprint region of the IR spectrum. B The carbonyl groups absorb at frequencies in the same region of the IR spectrum. C The compounds produce different fragmentation patterns in a mass spectrum. D The compounds have molecular ion peaks at different mass to charge ratios in



12	12 A sample of propanone, CH ₃ COCH ₃ , was heated under reflux with potassium dichromate(VI) acidified with sulfuric acid, and then the mixture was distilled. Apart from the peaks due to the C—C and C—H bonds, what peak(s) would be present in the IR spectrum of the distillate?					
	\times	Α	A peak due to the C=C) only.		
	\times	В	A peak due to the O—H	t only.		
	X	C	Peaks due to C=O and	О—Н.		
	X	D	Peaks due to C—O, C=	O and O—H.		
				ſ	Total for Question 12 = 1	mark)
13			of the following describe ions?	es the appearance of iodin	e under the stated	
			Solid	Dissolved in aqueous potassium iodide	Dissolved in a liquid hydrocarbon	
	\times	A	purple	brown	purple	
	X	В	brown	blue-black	yellow	
	X	C	shiny grey	brown	purple	
	\times	D	shiny grey	brown	brown	
				('	Total for Question 13 = 1	mark)
14			o calculations of global v phere?	varming exclude the effect		
	X	Α	Water vapour is not a gi	reenhouse gas.		
	X	B	Water vapour is a much	less potent greenhouse g	as than carbon dioxide.	
	X	C	The average concentrat	ion of water vapour in the	atmosphere is fairly const	ant.
	\mathbf{X}	D The concentration of water vapour in the atmosphere is much lower than that of carbon dioxide.				nat
				(*	Total for Question 14 = 1	mark)

15 A com	pound of nitrogen and hydrogen only is analyzed and found to contain 97.7%			
by mass of nitrogen. What is the empirical formula of the compound?				
	Molar masses /g mol ⁻¹ : $H = 1$; $N = 14$			
Α ⊠	NH_3			
B	NH ₂			
⊠ C	N_3H_5			
D 🛛	N₃H			
	(Total for Question 15 = 1 mark)			
16 Chloro involv	ofluorocarbons, CFCs, damage the ozone layer. The mechanism of the process es			
🖾 A	homolytic fission.			
B	heterolytic fission.			
🖾 C	electrophilic addition.			
D	nucleophilic substitution.			
	(Total for Question 16 = 1 mark)			
17 Floctr	ophiles are			
	electron pair donors that are attracted to regions of high electron density.			
B	electron pair donors that are attracted to regions of low electron density.			
	electron pair acceptors that are attracted to regions of high electron density.			
⊡ C	electron pair acceptors that are attracted to regions of low electron density.			
	(Total for Question 17 = 1 mark)			
	(Total for Question 17 = 1 mark)			
	TOTAL FOR SECTION A = 20 MARKS			



SECTION B

Answer ALL the questions. Write your answers in the spaces provided.

18 Halogenoalkanes react slowly with water to form alcohols. The equation for the reaction is

 $\mathsf{RX} \ + \ \mathsf{H}_2\mathsf{O} \ \rightarrow \ \mathsf{ROH} \ + \ \mathsf{H}^+ \ + \ \mathsf{X}^-$

(a) The rate of this reaction for different halogenoalkanes was investigated using the apparatus below.



In one experiment, equal amounts of 1-chlorobutane, 1-bromobutane and 1-iodobutane were placed in separate test tubes with 5 cm³ of ethanol. These test tubes were placed in the water bath together with a test tube containing aqueous silver nitrate. After about 5 minutes, 1 cm³ of the silver nitrate solution was added to each test tube containing a halogenoalkane and the time taken for a precipitate to form in each test tube was noted. The temperature of the water bath was maintained at 50 °C.

(i) Why is ethanol used as a solvent in this experiment?

(1)

(ii) Explain why the apparatus was left for 5 minutes before the silver nitrate was added.

(1)



1
I



		(3)
	Show the lone pair involved in the mechanism and any relevant dipoles and curly arrows.	(3)
(iii)	Give the mechanism for the reaction of 1-bromobutane with aqueous alkali.	
		(1)

(iv)	One student suggested that the final reaction presence of an alcohol using phosphorus(V) c using potassium dichromate(VI) with sulfuric a Describe the result of a positive test for alcoho	hloride while another suggested acid.				
	Explain which test is better for the final reaction	on mixture.	(3)			
Observatio	on with PCI₅					
Observatio	on with acidified $K_2Cr_2O_7$					
Explanatio	n					
	romobutane is classified as a primary halogen uctural isomers with a molecular formula C ₄ H ₉ B					
	e the skeletal formula of the three isomers, ot	her than 1-bromobutane,				
cla	ssifying the halogenoalkane in each case.		(3)			
	Skeletal formula Classification					
			_			
		(Total for Question 18 = 18 ma	rks)			



19 Two white powders, A and B , known to be Group 2 carbonates, are investigated.	
(a) (i) The presence of the carbonate ion is usually confirmed using a simple test carried out in two stages at room temperature. Describe this test and its results.	
	(2)
Test	
Results	
(ii) Using barium carbonate as your example, write the equation for each of the stages of the carbonate test. Include state symbols in both equations.	(3)
First equation (test)	
Second equation (result)	
Second equation (result)	

(i) Describe how a flame test is carried out.	
·,	(3)
(ii) Give the formulae of the metal ions in A and B .	
	(2)
А В	
*(iii) Explain the origin of the flame colour.	
	(3)
(iv) Suggest why compound A produces no flame colour.	(1)
(iv) Suggest why compound A produces no flame colour.	(1)



-	1 • • • • •	arbon dioxide		1	.1	
Exj is c	plain why the thern lescended.	hal stability of	the metal ca	rbonates incre	ases as the grou	р
						(3)
				(Total for Ou	lestion 19 = 17	marks)
						111a1K5/

P 4 4 8 8 1 A 0 1 6 2 4

20 Induced hydraulic fracturing, commonly known as fracking, which was developed in 1947, is a technique for extracting natural gas (mainly methane) from shale deposits. While natural gas is a much cleaner fuel than coal, it is difficult to carry out fracking without leakage. Because methane is a far more potent greenhouse gas than carbon dioxide, it has been calculated that leakage rates of around 2% are sufficient to increase global warming. (a) Suggest what is meant by 'natural gas is a much cleaner fuel than coal'. (1) (b) Explain how greenhouse gases cause global warming. (2) (c) Suggest why methane is a far more potent greenhouse gas than carbon dioxide. (1) (Total for Question 20 = 4 marks) **TOTAL FOR SECTION B = 39 MARKS** 17

881

A 0

SECTION C

Answer ALL the questions. Write your answers in the spaces provided.



Swimming Pool Chemistry

The circulation of water in swimming pools is much slower than that in most natural water courses, but the number of people using a given volume of water is often far greater. If steps are not taken to keep microorganisms and other contaminants under control, the water will become hazardous. Filters are used to remove solid material and chemicals are added to disinfect the water.

The most common method of disinfection involves the use of chlorine compounds, but systems using bromine have advantages. These systems depend on their oxidizing properties. With chlorine systems, the key species is the chlorate(I) ion (OCI⁻) which kills bacteria by damaging the structure of their cell walls and disrupting enzyme activity.

A simple way of adding chlorate(I) ions to water is by using chlorine. The weak acid, chloric(I) acid (HOCI), is formed and this dissociates producing the chlorate(I) ion.

(a) The equation for the reaction of chlorine with water is:

 $Cl_2(aq) + H_2O(l) \implies HOCI(aq) + HCI(aq)$

By referring to the relevant oxidation numbers, explain why this is a disproportionation reaction.

(3)





4 4 8 8 1 A 0 1 9 2

(c) If the concentration of chlorate(I) ions in a swimming pool is too low, bacteria will flourish. If it is too high, swimmers will be harmed. In the United Kingdom, it is recommended that the concentration of chlorine be in the range 1–2 mg dm⁻³. The total amount of chlorine, Cl₂, present can be measured by titration.

In such a determination, excess sulfuric acid and potassium iodide were added to a 1.00 dm³ sample of swimming pool water and the resulting solution required 9.65 cm³ of 0.00550 mol dm⁻³ sodium thiosulfate for complete reaction.

The equations for the reactions are:

$$\begin{array}{rcl} \mathsf{CI}_2 \ + \ 2\mathsf{I}^{\scriptscriptstyle -} \ \rightarrow \ \mathsf{I}_2 \ + \ 2\mathsf{CI}^{\scriptscriptstyle -} \\ \\ \mathsf{I}_2 \ + \ 2\mathsf{S}_2\mathsf{O}_3^{2^{\scriptscriptstyle -}} \ \rightarrow \ 2\mathsf{I}^{\scriptscriptstyle -} \ + \ \mathsf{S}_4\mathsf{O}_6^{2^{\scriptscriptstyle -}} \end{array}$$

(i) Show that the concentration of chlorine, Cl₂, in this sample meets the UK recommended requirements.

All steps in your calculation must be shown.

(3)



(ii) Analysing a single sample of swimming pool water in this way is likely to give unreliable results because it is not possible to repeat the titration. Suggest another way in which this method is unreliable.	
	(1)
(d) One advantage of the bromine disinfection system is that, while chloric(I) acid decomposes in sunlight forming hydrogen chloride and oxygen, bromic(I) acid (HOBr) is stable. Another advantage is that bromine is less volatile than chlorine.	
(i) Write an equation for the decomposition of chloric(I) acid in sunlight. State symbols are not required.	
	(1)
(ii) Explain why bromine is less volatile than chlorine.	(2)
	21
	Turn over

(e) The pH of the swimming pool affects the performance of the disinfectant. The data below show how the concentrations of bromine species vary with pH.

рН	% bromine as HOBr	% bromine as OBr-
6.0	100	0.0
7.0	98.0	2.0
8.0	83.0	17.0
8.5	57.0	43.0

The equation for the reaction of bromic(I) acid in water is

 $HOBr(aq) \rightleftharpoons H^+(aq) + OBr^-(aq)$

(i) Assume the pH of the swimming pool is neutral. Explain how any changes in the pH of the swimming pool affect the concentration of the bromine species.

(3)

(ii) The higher the concentration of bromate(I) ions, the more effective the disinfectant.

Suggest a disadvantage of too high a pH.

(1)

(Total for Question 21 = 21 marks)

TOTAL FOR SECTION C = 21 MARKS TOTAL FOR PAPER = 80 MARKS





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	0 (8)	(18)	4.0	He helium	2	20.2	Ne	neon	10	39.9	Ar	argon 18	83.8	Кr	krypton 36	131.3	Xe	xenon	54	[222]	Rn	radon 86		ted								_		
	7				(17)	19.0	Ŀ	fluorine	9	۲. ۲۵	บ	cniorine 17	79.9	Br	bromine 35	126.9	_	iodine	53	[210]	At	astatine 85		Elements with atomic numbers 112-116 have been reported but not fully authenticated		175	Lu	lutetium 71	[257]	Lr Internetium	103			
	9				(16)	16.0	0	oxygen	8	32.1	S	sultur 16	79.0	Se	selenium 34	127.6	Te	tellurium	52	[209]	Po	polonium 84		-116 have h nticated		173	Υb	ytterbium 70	[254]	No	102			
	2				(15)	14.0	z	nitrogen		31.0	م -	pnospnorus 15	74.9	As	a	121.8	Sb	antimony	51	209.0	Bi	bismuth 83		tomic numbers 112-116 hav but not fully authenticated		169	Tm	thulium 69	[256]	Md	101			
	4				(14)	12.0	υ	carbon	9	28.1		silicon 14	72.6	e B	germanium 37	118.7	Sn	tin	50	207.2	Pb	lead 82		atomic nu but not i		167	E	erbium 68	[253]		100			
	ε				(13)	10.8	8	boron	2 2	27.0	A	aluminium 13	69.7	Ga	gallium 31	114.8	<u>_</u>	indium	49	204.4	F	thallium 81		nents with		165	Ю		[254]	Cf Es	66			
SULLA												(12)	65.4	Zn	zinc	112.4	DQ	cadmium	48	200.6	Hg	mercury 80			_	163	D	dysprosium 66	[251]	Cf	98			
Llell														(11)	63.5	Cu	copper	107.9		° v	47	197.0	Au	gold 79	[272]	Rg roentgenium	111	159		terbium 65	[245]	BK hertalium	97	
												(10)	58.7	Ż	nickel 28	106.4	Ρd	palladium	46	195.1	Ł	platinum 78	[271]	DS damstadtium	110	157		gadolinium 64	[247]		96			
		_						(6)					58.9	ပိ	cobalt 77	102.9	Rh	£	45	192.2	۔	iridium 77	[268]	Mt meitnerium	109	152		europium 63	[243]	Am	95			
		•		h ydrogen	-							(8)	55.8	Fe		101.1	Ru	rut	44	190.2	S	osmium 76	[277]	Hs hassium	108	150		samarium 62		Pu	94			
lle re												(2)	54.9	Wn	ma	[98]	μ	molybdenum technetium	43	186.2	Re	rhenium 75	[264]	Bh bohrium	107	[147]	Pm	promethium 61	[237]	Np	93			
_						mass	bol	rodani	Initiber			(9)	52.0	J	chr	95.9	Wo	molybdenum	42	183.8	3	tungsten 74	[266]	Sg seaborgium	106	144	PN	præcodymium neodymium promethium 59 60 61		U	92			
					Key	relative atomic mass	atomic symbol	name	מנטווור (הנטרטוו) העוווטפו			(5)	50.9	>	vanadium	92.9	qN	Ē	41	180.9	Ta	tantalum 73	[262]	Db dubnium	105	141	Pr	praseodymium 59	[231]	Pa	91			
							relat	atc	tomic	qroillic			(4)	47.9	ï	titanium 22	91.2	Zr	zirconium	40	178.5		hafnium 72	[261]	Rf rutherfordium	104	140	Ce	cerium 58	232	H	60		
						_						(3)	45.0	Sc	scandium 21	88.9	7	yttrium	39	138.9	La*	lanthanum 57	[227]	Ac* actinium	89		es							
	2				(2)	0.6	Be	beryllium	4	24.3	Mg	magnesium 12	40.1	Ca	calcium	87.6		str	38	137.3	Ba	barium 56	[226]	Ra radium	88		* Lanthanide series	* Actinide series						
	-				(1)	6.9	:-	lithium		23.0	Na	sodium 11	39.1	¥	potassium 10	85.5	Rb	rubidium	37	132.9	S	caesium 55	[223]	Fr francium	87		* Lanth	* Actin						

The Periodic Table of Elements

