Surname	Centre Number	Candidate Number
First name(s)		2



GCE AS





B410U10-1

TUESDAY, 17 MAY 2022 - MORNING

CHEMISTRY - AS component 1

The Language of Chemistry, Structure of Matter and Simple Reactions

1 hour 30 minutes

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In addition to this examination paper, you will need a:

- calculator;
- Data Booklet supplied by WJEC.

Section B

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Maximum Mark	Mark Awarded			
10				
8				
16				
16				
13				
17				
80				
	Mark 10 8 16 16 13			

For Examiner's use only

INSTRUCTIONS TO CANDIDATES

Use black ink or black ball-point pen. Do not use gel pen or correction fluid. You may use a pencil for graphs and diagrams only.

Write your name, centre number and candidate number in the spaces at the top of this page.

Section A Answer **all** questions. **Section B** Answer **all** questions.

Write your answers in the spaces provided in this booklet. If you run out of space, use the additional page(s) at the back of the booklet, taking care to number the question(s) correctly.

Candidates are advised to allocate their time appropriately between **Section A (10 marks)** and **Section B (70 marks)**.

INFORMATION FOR CANDIDATES

The number of marks is given in brackets at the end of each question or part-question.

The maximum mark for this paper is 80.

Your answers must be relevant and must make full use of the information given to be awarded full marks for a question.

The assessment of the quality of extended response (QER) will take place in **Q.9**(a).



SECTION A	Ex
	[4]
(a) State what is meant by a polar covalent bond.	[1]
(b) On the diagram below mark any permanent dipole.	[1]
H — F	
lodine-131 decays by beta emission.	
Identify the element formed.	[1]
A mass of 2.750g is weighed by difference.	
Calculate the percentage error in this value when a 3 decimal place balance is used.	[2]
Percentage error =	%
Complete the electron arrangement of iron.	[1]
1s ² 2s ² 2p ⁶	
Write an equation for the thermal decomposition of calcium hydroxide.	[1]
	Identify the element formed. A mass of 2.750 g is weighed by difference. Calculate the percentage error in this value when a 3 decimal place balance is used. Percentage error = Complete the electron arrangement of iron. 1s ² 2s ² 2p ⁶



				Evamia
6.	(a)	State why magnesium and barium are described as s-block elements.	[1]	Examin only
	(b)	Describe how the results of a flame test would differentiate between samples of magnesium sulfate and barium sulfate.	[1]	
	(c)	Barium sulfate can be made using a precipitation reaction between solutions of two metal compounds. One of the compounds is barium chloride.		
		Identify a metal compound that could be used for the other solution.	[1]	
				10



SECTION B

Answer all questions.

7. When heated above 100 °C, nitrosyl chloride (NOCI) partially decomposes to form nitrogen monoxide and chlorine.

 $2NOCl(g) \rightleftharpoons 2NO(g) + Cl_2(g)$

(a) Write an expression for the equilibrium constant (K_c) for this reaction.

Give the unit of K_c .

[2]

Unit

(b) At a fixed temperature, a mixture of NOCI, NO and Cl₂ reached equilibrium in a sealed container.

The equilibrium mixture formed contained NOCI, at a concentration of 0.126 mol dm $^{-3}$, and NO, at a concentration of 5.73×10^{-2} mol dm $^{-3}$.

The value of K_c for the equilibrium at this temperature was 7.40×10^{-3} .

Calculate the concentration of Cl_2 in this equilibrium mixture.

[3]

Concentration of $\operatorname{Cl}_2 = \dots \mod \operatorname{dm}^{-3}$



(c)	State and explain, with reference to Le Chatelier's principle, how the amount of chlorine would change if the reaction were carried out at a higher pressure. [3]	Exam on
		8



)			
Xenon is found in Group 0 of the Periodic Table. (a) Xenon has a first ionisation energy of 1170 kJ mol ⁻¹ .				
	(i)	Explain why the first ionisation energy of xenon is lower than that of krypton.	[2]	
	(ii)	Explain why the first ionisation energy of xenon is higher than that of iodine.	[2]	
(b)		on is an unreactive element but can be made to react with the very electronega nent fluorine. State the meaning of the term electronegativity.	tive [1]	
(b)	elem	nent fluorine.	[1]	



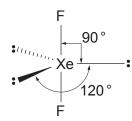
(iv) Calculate the volume of xenon gas required, at a temperature of 400 °C and a pressure of 1.00×10^5 Pa, to make 5.00 g of xenon difluoride.

Give your answer in dm³.

[3]

Volume of xenon = dm³

(v) Xenon difluoride molecules are linear.



Using your knowledge and understanding of VSEPR theory, suggest why the electron pairs arrange themselves in this way.

.....

[2]

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(vi)	Xenon difluoride is a strong oxidising agent.	Exa o
	An example of such a reaction is given below.	
	$2CrO_2F_2 + XeF_2 \longrightarrow 2CrOF_3 + Xe + O_2$	
	Use the oxidation states of xenon and oxygen to show why xenon difluoride is an oxidising agent in this reaction. [2]	
•••••		
•••••		-
•••••		-
•••••		•
(vii)	Xenon difluoride reacts with water to produce hydrogen fluoride and two elements.	
	Write an equation to represent this reaction. [2]	I
		. _

16



9.	Wate this th	r authorities provide millions of customers with a safe and reliable water supply. To do ney operate a comprehensive monitoring system.
	(a)	Mass spectrometry is just one of the many techniques that can be used to analyse water samples. Both chlorine and chlorine-containing compounds can be analysed in this way.
		Outline how the mass spectrometer works and describe the mass spectrum of chlorine. [6 QER]
	•••••	



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(b) There are regulations to govern limits of certain chemicals in drinking water.

Species	Level not to be exceeded
arsenic	10 μg/l
copper	2 mg/l
chromium	50 mg/l
fluoride	1.5 mg/l

(i)	The volume of water needed to fill a glass is 330 cm ³	•
-----	---	---

Calculate the mass of the water in the glass.

[1]

(ii) Calculate the number of hydrogen atoms present in this mass of water. [2]

Number of hydrogen atoms =



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	(iii)	A student has two different water samples contained in glasses identical to the one in (b)(i).
		One sample is a full glass containing water contaminated with arsenic and the other sample is a glass one-third full, containing water contaminated with copper.
		Given that both these water samples contain the maximum mass of these contaminants, according to the limits in the table, determine which sample contains the greater number of moles of contaminant.
		You must show your working. [3
(c)		e water authorities add calcium fluoride to their drinking water and other water orities do not.
(c)	autho	orities do not. ain your own view on the fluoridation of drinking water, including arguments for and
	autho Expla agair	orities do not. ain your own view on the fluoridation of drinking water, including arguments for and
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(d)	Water from the Peak District has percolated through rocks containing calcium fluoride in the form of the mineral fluorspar.

Calcium fluoride, CaF_2 , has a solubility of $1.7 \times 10^{-3} \, \text{g} / 100 \, \text{g}$ of water.

Calculate the number of moles of calcium fluoride dissolved in 100 g of this water. [1]

Number of moles = mol

16



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Galliu tempe	im is an unusual metal because it melts if you hold it in your hand. It has a melting erature of 29.8 °C.	
(a)	Aluminium is above gallium in the Periodic Table. It melts at 660 °C.	
	Describe the solid structure of aluminium using a labelled diagram to support your answer.	[3]
• • • • • • • • • • • • • • • • • • • •		
•••••		
(b)	In gallium the bonding between two neighbouring particles is covalent and the struction is built up from Ga ₂ dimers, similar to the structure of iodine.	ture
(b)	In gallium the bonding between two neighbouring particles is covalent and the structise built up from Ga_2 dimers, similar to the structure of iodine. Use this information to explain the unusual melting temperature of gallium compared aluminium.	
(b)	is built up from Ga ₂ dimers, similar to the structure of iodine. Use this information to explain the unusual melting temperature of gallium compared	d to
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 (c) Gallium has 31 known isotopes but only two of them are stable and occur naturally – gallium-69 and gallium-71. Given that the relative atomic mass of natural gallium is 69.798, determine the percentage abundance of these two isotopes. 	[3]
	[3]
Abundance of Ga-69 =	%
Abundance of Ga-71 =	%
(d) (i) Write an equation for the first ionisation energy of gallium.	[1]
(ii) Ionisation energy can be measured in electron volts where	
$1 \text{eV} = 1.602 \times 10^{-19} \text{J}$	
The first ionisation energy of gallium has a value of 5.9993 eV.	
Calculate the wavelength of radiation, in m, that will ionise an atom of gallium form a Ga^+ ion.	to [3]
Wavelength =	m



Turn over.

13

[1]

11. (a) In a titration experiment, good technique is essential for an accurate result to be obtained.

titration.

(i) Suggest a reason for removing the funnel after it has been used for filling the burette. [1]

(ii) Suggest **one** other potential source of error in **using** the burette to carry out a

(b) Sodium hydroxide solutions are often contaminated with sodium carbonate owing to the ease with which sodium hydroxide reacts with carbon dioxide in the air.

The technique of double titration allows us to determine the extent to which a solution has been contaminated.

(i) Write an equation for the reaction between sodium hydroxide and carbon dioxide.

(ii) The method of double titration uses two indicators to determine two end-points. During the titration, hydroxide ions and carbonate ions react according to the following equations.

hydroxide: $OH^- + H^+ \longrightarrow H_2O$ carbonate: $CO_3^{2-} + H^+ \longrightarrow HCO_3^-$ both complete at phenolphthalein stage

 $\mathrm{HCO_3}^-$ + H^+ \longrightarrow $\mathrm{CO_2}$ + $\mathrm{H_2O}$ complete at methyl orange stage

By recording the volumes of hydrochloric acid added at each end-point it is possible to calculate the concentrations of the hydroxide ions and carbonate ions in the original solution and thus determine the extent of contamination.

Assume that hydroxide ions and carbonate ions are the only anions in the initial solution.

A 1.00 dm³ flask containing sodium hydroxide solution had been left open to the air in a laboratory for a period of time.

A student performed a double titration to determine the volume of carbon dioxide that had reacted with the sodium hydroxide solution.

He titrated a 25.0 cm³ sample of the solution with 1.00 mol dm⁻³ hydrochloric acid. He recorded the volume used at the phenolphthalein end-point.

He then added methyl orange and continued titrating to its end-point and recorded the total volume of hydrochloric acid added.

Results

Volume of HCl at the phenolphthalein end-point/cm ³	22.50
Total volume of HCl added at the methyl orange end-point/cm ³	25.00

l.	Calculate the number of moles of hydrochloric acid used in the first stage)
	of the titration. This gives the total number of moles of hydroxide ions plu	S
	carbonate ions in the 25.0 cm ³ sample.	[1]

Number of moles =	=	mol
-------------------	---	-----

II.	Calculate the number of moles of hydrogencarbonate ions reacting in the)
	second stage of the reaction.	[2]



Examine	
I. and II., determine the number of moles e number of moles of hydroxide ions in the [2]	III. Using yo of carbor 25.0 cm ³
arbonate ions = mol	
nydroxide ions = mol	
arbonate ions determined in part III., dioxide absorbed into the original solution at [3]	IV. Using the calculate 298 K an
arbon dioxide = dm ³	



			19	
(c)	(i)	Expl	ain what is meant by a strong acid and write an equation to show how ochloric acid behaves as a strong acid.	[2]
	(ii)	I.	Calculate the pH of 500 cm ³ of 0.50 mol dm ⁻³ hydrochloric acid.	[1]
			pH =	
		II.	Calculate the volume of water that must be added to this solution to increase its pH to a value of 1.0. Show your working.	[3]
			Volume of water added =	dm³ -
			END OF PAPER	



Question number	Additional page, if required. Write the question number(s) in the left-hand margin.	Examiner only
		1
		•





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CHEMISTRY – AS component 1 Data Booklet

Avogadro constant molar gas constant molar gas volume at 273 K and 1 atm molar gas volume at 298 K and 1 atm Planck constant speed of light density of water specific heat capacity of water ionic product of water at 298 K fundamental electronic charge

 $N_A = 6.02 \times 10^{23} \,\mathrm{mol}^{-1}$ R^{-} = 8.31 Jmol⁻¹ K⁻¹ $V_m = 22.4 \, \text{dm}^3 \, \text{mol}^{-1}$ $V_m = 24.5 \text{ dm}^3 \text{ mol}^{-1}$ $h = 6.63 \times 10^{-34} \text{ Js}$ $c = 3.00 \times 10^8 \,\mathrm{m\,s^{-1}}$ $d = 1.00 \,\mathrm{g}\,\mathrm{cm}^{-3}$ $c = 4.18 \text{ Jg}^{-1} \text{K}^{-1}$ $K_w = 1.00 \times 10^{-14} \text{ mol}^2 \text{dm}^{-6}$

 $= 1.60 \times 10^{-19} \text{ C}$

temperature (K) = temperature (°C) + 273

$$1 \,dm^3 = 1000 \,cm^3$$

 $1 \,m^3 = 1000 \,dm^3$
 $1 \,tonne = 1000 \,kg$
 $1 \,atm = 1.01 \times 10^5 \,Pa$

Multiple	Prefix	Symbol
10-9	nano	n
10-6	micro	μ
10-3	milli	m

Multiple	Prefix	Symbol	
10³	kilo	k	
10 ⁶	mega	M	
10 ⁹	giga	G	

Infrared absorption values

Bond	Wavenumber/cm ⁻¹
C-Br	500 to 600
C-CI	650 to 800
C - O	1000 to 1300
C = C	1620 to 1670
C = O	1650 to 1750
$C \equiv N$	2100 to 2250
C-H	2800 to 3100
O — H (carboxylic acid)	2500 to 3200 (very broad)
O — H (alcohol / phenol)	3200 to 3550 (broad)
N-H	3300 to 3500

13 C NMR chemical shifts relative to TMS = 0

Type of carbon	Chemical shift, δ (ppm)			
$-\overset{\mid}{\operatorname{c}}-\overset{\mid}{\operatorname{c}}-$	5 to 40			
R — C — Cl or Br	10 to 70			
R-c-c- 0	20 to 50			
R-C-N	25 to 60			
-c-o-	50 to 90			
c = c	90 to 150			
$R-C \equiv N$	110 to 125			
	110 to 160			
R — C — (carboxylic acid / es O	ter) 160 to 185			
R — C — (aldehyde / ketone) O	190 to 220			

¹H NMR chemical shifts relative to TMS = 0

Type of proton	Chemical shift, δ (ppm)					
$-CH_3$	0.1 to 2.0 0.9 1.3 2.0 2.0 to 2.5 2.0 to 3.0 2.2 to 2.3					
R-CH ₃						
R-CH ₂ -R						
$CH_3-C \equiv N$						
CH ₃ -C						
$-CH_2-C$						
\bigcirc —CH $_3$						
HC-Cl or HC-Br	3.1 to 4.3 3.3 to 4.3					
HC-O						
R-OH	4.5 *					
-C = CH	4.5 to 6.3					
-C = CH - CO	5.8 to 6.5					
\leftarrow CH=C	6.5 to 7.5					
\leftarrow H	6.5 to 8.0 7.0 *					
ОН						
R-C H	9.8 *					
R-COH	11.0 *					

^{*}variable figure dependent on concentration and solvent

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THE PERIODIC TABLE

	0 2	Helium	19.0 20.2 F Ne Neon 9 10	35.5 40.0 Cl Chlorine Argon 17 18	79.9 83.8 Br Kr m Bromine Krypton 35	127 131 Xe Xe Nenon 53 54	(210) (222) At Rn n Astatine Radon 85		175 Lu Lutetium 71	(257) Lr Lawrendum 103
	9	p bld q Nitrogen	0	32.1 S Sulfur 16	79.0 Se ic Selenium	128 Te Tellurium 52	(210) Po Polonium 84		173 Yb Ytterbium 70	(254) No Nobelium 102
	3		Ž	31.0 Phosphorus	AS AS Arsenic	Sb Antimony 51	209 Bi Bismuth 83		169 Tm Thulium 69	(256) Md Mendelevium 101
	4		12.0 C C Carbon	Silicon 14	72.6 Ge m Germanium 32	Sn Tin 50	207 Pb m Lead 82		167 Er n Erbium 68	(253) Fm Fermium 100
	က	,	10.8 B Boron 5	Aluminium 13	69.7 Ga Gallium	115 In Indium 49	204 TI Thallium		165 Ho m Holmium 67	(254) ES m Einsteinium 99
Щ					65.4 Zn Sinc 30	Cd Cd Cdmium 48	201 Hg Mercury 80	f block	163 Dy Dysprosium 66	(251) Cf Cf Californium 98
HE PERIODIC TABI					63.5 Cu Copper	Ag Ag Silver	197 Au Gold 79		159 Tb n Terbium 65	(245) BK Bk 97
					Nickel Nickel 28	106 Pd m Palladium 46	195 Pt n Platinum 78		157 Gd n Gadolinium 64	(247) Cm Curium 96
ERIC			58.9 Co Cobalt 27	103 Rh Rhodium 45	192 Ir Iridium		(153) Eu Europium 63	(243) Am Americium 95		
出	roup	y	mass mass atomic number	d block	55.8 Fe Iron 26	101 Ru Ruthenium 44	190 Os n Osmium 76		Sm Samarium 62	(242) Pu Plutonium 94
F	Ō	, A	Symbol Name Z	d d	Manganese 25	98.9 TC Technetium 43	186 Re Rhenium		(147) Pm n Promethium 61	(237) Np Neptunium 93
					52.0 Cr Chromium 24	95.9 Molybdenum 42	184 W Tungsten 74		144 Nd Neodymium 60	238 U Uranium 92
				50.9 Vanadium 23	92.9 Nbbium 41	181 Ta Tantalum 73	-	141 Pr Praseodymium 59	(231) Pa Protactinium 91	
					47.9 Ti Titanium 22	91.2 Zr Zirconium 40	179 Hafnium		Cerium 58	232 Th Thorium 90
		A			Scandium 21	88.9 Y Yttrium 39	139 La La La Lanthanum	(227) AC AC Actinium 89	► Lanthanoid elements	▶ Actinoid elements
	7	OCK C	9.01 Be Beryllium	Mg Magnesium	40.1 Ca Calcium 20	87.6 Sr Strontium 38	137 Ba Barium 56	(226) Ra Radium 88	<u> </u>	A
	_	s block 1.01 Hydrogen	6.94 Li Lithium	23.0 Na Sodium	39.1 K Potassium 19	85.5 Rb Rubidium 37	133 Cs Caesium 55	(223) Fr Francium 87		
		Period 1	7	က	4	Ŋ	9			
		Q	© WJEC C	BAC Ltd.	(B410U10	0-1A)				