

New
Specification



Rewarding Learning

ADVANCED SUBSIDIARY (AS)
General Certificate of Education
2017

Centre Number

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Candidate Number

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Chemistry

Assessment Unit AS 1

assessing

Basic Concepts in Physical
and Inorganic Chemistry



[SCH12]

SCH12

FRIDAY 26 MAY, MORNING

TIME

1 hour 30 minutes.

INSTRUCTIONS TO CANDIDATES

Write your Centre Number and Candidate Number in the spaces provided at the top of this page.

Answer **all fifteen** questions.

Answer **all ten** questions in **Section A**. Record your answers by marking the appropriate letter on the answer sheet provided. Use only the spaces numbered 1 to 10. Keep in sequence when answering.

Answer **all five** questions in **Section B**. **You must answer the questions in the spaces provided.**

Do not write outside the boxed area on each page or on blank pages.

Complete in black ink only. **Do not write with a gel pen.**

INFORMATION FOR CANDIDATES

The total mark for this paper is 90.

Quality of written communication will be assessed in Question **13(c)**.

In Section A all questions carry equal marks, i.e. **one** mark for each question.

In Section B the figures in brackets printed down the right-hand side of pages indicate the marks awarded to each question or part question.

A Periodic Table of Elements, containing some data, is included with this question paper.

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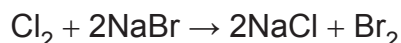
20SCH1201

Section A – Multiple Choice

Select the correct response in each case and mark its code letter by connecting the dots as illustrated on the answer sheet.

Each multiple choice question is worth 1 mark.

- 1 Bromine is formed in the reaction below.



Which statement about the reaction is correct?

- A Bromide ions lose electrons
- B Bromine is reduced by chlorine
- C Chloride ions are reduced
- D Chlorine is a weaker oxidising agent than bromide
- 2 Which trend in the Periodic Table is correct?
- A Boiling point decreases from fluorine to bromine
- B First ionisation energy decreases from lithium to caesium
- C First ionisation energy increases from nitrogen to oxygen
- D Melting point decreases from sodium to silicon
- 3 Which of the following is the structure of $^{55}\text{Mn}^{2+}$?

	protons	neutrons	electrons
A	25	30	23
B	25	30	27
C	27	30	25
D	30	25	28



- 4 Potassium iodide is formed when potassium is warmed in iodine vapour. Which of the following shows the bonding in the three species?

	potassium	iodine	potassium iodide
A	ionic	covalent	ionic
B	metallic	ionic	covalent
C	covalent	covalent	ionic
D	metallic	covalent	ionic

- 5 The element astatine lies below iodine in the Periodic Table and is likely to

- A be black.
- B be a volatile liquid at room temperature and pressure.
- C form an astatide ion, At^{2-} .
- D oxidise iodide ions to iodine.

- 6 Which molecule is non-polar?

- A H_2S
- B NH_3
- C PF_3
- D SF_6

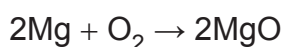
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- 7 The element boron has a relative atomic mass of 10.8. In this sample, boron exists as two isotopes, ^{10}B and ^{11}B . The percentage abundance of ^{10}B in this sample of boron is

A 10.8%.
B 20.0%.
C 80.0%.
D 89.2%.

- 8 When burned in oxygen magnesium forms magnesium oxide.



What is the number of molecules of oxygen required for the complete oxidation of 1.2g of magnesium?

- A 1.5×10^{22}
B 3.0×10^{22}
C 3.0×10^{23}
D 6.0×10^{23}
- 9 Which statement describes the trends in electronegativity values in the Periodic Table?
- A Decrease across a Period and increase down a Group
B Decrease across a Period and decrease down a Group
C Increase across a Period and increase down a Group
D Increase across a Period and decrease down a Group



10 Which of the following would exactly neutralise 10.0 cm³ of 1.00 mol dm⁻³ NaOH(aq)?

- A 2.50 cm³ of 1.00 mol dm⁻³ CH₃COOH
- B 5.00 cm³ of 1.00 mol dm⁻³ HCl
- C 5.00 cm³ of 1.00 mol dm⁻³ H₂SO₄
- D 3.00 cm³ of 1.00 mol dm⁻³ H₃PO₄

[Turn over



Section B

Answer **all five** questions in the spaces provided.

- 11** Sulfate, hydrogensulfate and thiosulfate ions are formed when sulfuric and thiosulfuric acids ionise.

(a) (i) Write the equation for the complete ionisation of thiosulfuric acid.

_____ [2]

(ii) Write the formula for the hydrogensulfate ion.

_____ [1]

(b) (i) Write the formula for ammonium sulfate.

_____ [1]

(ii) Describe the bonding in ammonium sulfate.

_____ [2]

(c) Describe how you could use chemical tests on an aqueous solution of ammonium sulfate to prove that it contains ammonium ions and sulfate ions.

_____ [4]



12 Some properties of the metals sodium and aluminium are shown in the table below.

metal	charge on metal ion	electronic structure of the atom	melting point /°C
sodium	1+	$1s^2 2s^2 2p^6 3s^1$	98
aluminium	3+	$1s^2 2s^2 2p^6 3s^2 3p^1$	660

(a) Describe, without using a diagram, the bonding in sodium metal.

[2]

(b) Explain why aluminium has a higher melting point than sodium.

[2]

(c) (i) Write the equation, including state symbols, for the first ionisation energy of sodium.

[2]

(ii) The first six ionisation energies, in kJ mol^{-1} , of sodium are 496, 4563, 6913, 9544, 13352 and 16611. Explain which of these values can be used to identify sodium as belonging to Group I of the Periodic Table.

[2]

[Turn over]



- (iii) The outer electron in the sodium atom is located in the 3s orbital.
Explain what is meant by the term **orbital**.

[2]

(d) Aluminium forms covalent bonds with chlorine.

- (i) Explain what is meant by the term **covalent bond**.

[2]

- (ii) Write the equation for the reaction of aluminium with chlorine to form aluminium chloride, AlCl_3 .

[1]

- (iii) State the octet rule and explain whether the atoms in aluminium chloride obey the rule.

[3]



- 13 (a)** Zinc reacts with chlorine to form the ionic compound zinc chloride. Draw a dot and cross diagram, using outer electrons only, to show how zinc chloride, ZnCl_2 , is formed from zinc and chlorine atoms.

[2]

- (b)** Zinc is an essential trace element. People who have a zinc deficiency can take hydrated zinc sulfate, $\text{ZnSO}_4 \cdot x\text{H}_2\text{O}$, as a dietary supplement.

The value of x can be determined by heating hydrated zinc sulfate to constant mass.

A student heated 5.65 g of hydrated zinc sulfate and obtained 3.85 g of anhydrous zinc sulfate.

- (i)** Calculate the number of moles of anhydrous zinc sulfate obtained.

[1]

- (ii)** Calculate the mass of water present in the hydrated zinc sulfate.

[1]

- (iii)** Calculate the number of moles of water present in the hydrated zinc sulfate.

[1]

- (iv)** Calculate the value of x in $\text{ZnSO}_4 \cdot x\text{H}_2\text{O}$

[1]

[Turn over]



- (c) Describe how you would prepare 250.0 cm³ of a 28.7 g dm⁻³ zinc sulfate solution from the anhydrous solid.

In this question you will be assessed on using your written communication skills including the use of specialist scientific terms.

[6]



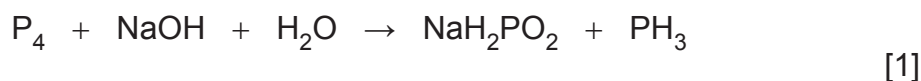
14 Nitrogen and phosphorus are Group V elements. They form the toxic hydrides ammonia and phosphine.

- (a)** Ammonia is formed by the reversible reaction of nitrogen with hydrogen. Write the equation for this reaction.

_____ [2]

- (b)** Phosphine is formed by the reaction of phosphorus with aqueous sodium hydroxide.

- (i)** Balance the equation for the formation of phosphine.



- (ii)** Deduce the oxidation number of phosphorus in:

P_4 _____

NaH_2PO_2 _____

PH_3 _____ [3]

- (iii)** Explain, using the oxidation numbers of phosphorus, why the reaction is described as disproportionation.

_____ [3]

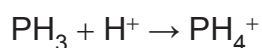
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- (c) The boiling point of ammonia is -33°C while that of phosphine is -88°C . Explain why the boiling point of ammonia is higher than that of phosphine.

[3]

- (d) Both ammonia and phosphine molecules react with H^+ ions.



- (i) Name the type of bond formed between a phosphine molecule and the H^+ ion.

[1]

- (ii) Draw and name the shapes of the molecule PH_3 and the ion PH_4^+ .



Shape _____



Shape _____

[4]

- (iii) Explain why the bond angle in PH_3 is different from the bond angle in PH_4^+ .

[3]



- (e) Ammonia is very soluble in water. Draw diagrams to show the two ways in which a molecule of ammonia can be attracted to a molecule of water. Include all partial charges and lone pairs in your diagram.

[4]

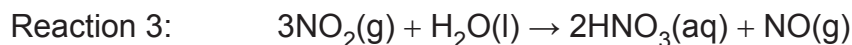
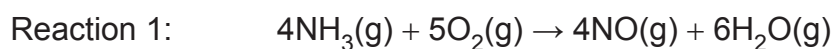
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20SCH1213

15 Ammonia is used to make nitric acid by the Ostwald Process outlined below.



- (a) (i)** Calculate the number of moles of oxygen needed to react with 6.8 kg of ammonia.

[3]

- (ii)** Calculate the number of moles of nitrogen(IV) oxide which can be obtained from 6.8 kg of ammonia.

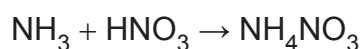
[2]

- (iii)** Calculate the concentration of nitric acid, in g dm^{-3} , produced on reacting the nitrogen(IV) oxide obtained in part **(ii)** with 50 dm^3 of water.

[3]



(b) Ammonia reacts with nitric acid according to the equation below.



The following results were obtained by diluting 25.0 cm^3 of a concentrated ammonia solution to 250.0 cm^3 in a volumetric flask and then titrating 25.0 cm^3 portions of the diluted ammonia solution using 0.100 mol dm^{-3} nitric acid.

titration	initial burette reading / cm^3	final burette reading / cm^3	titre / cm^3
rough	0.00	22.00	22.00
first accurate	0.10	21.40	21.30
second accurate	0.20	21.60	21.40

(i) Name a suitable indicator for the titration and state the colour change at the end point.

_____ [3]

(ii) Calculate the mean titre.

_____ [1]

[Turn over]



- (iii) A burette has an uncertainty of $\pm 0.05 \text{ cm}^3$. Calculate the uncertainty when two burette readings are used to calculate a titre value.

[1]

- (iv) Calculate the concentration of the concentrated ammonia solution in mol dm^{-3} .

[5]





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20SCH1219

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Question Number	Marks
Section A	
1–10	
Section B	
11	
12	
13	
14	
15	
Total Marks	

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General Information

1 tonne = 10^6 g

1 metre = 10^9 nm

One mole of any gas at 293 K and a pressure of 1 atmosphere (10^5 Pa) occupies a volume of 24 dm³

Avogadro Constant = 6.02×10^{23} mol⁻¹

Planck Constant = 6.63×10^{-34} J s

Specific Heat Capacity of water = $4.2 \text{ J g}^{-1} \text{ K}^{-1}$

Speed of Light = $3 \times 10^8 \text{ m s}^{-1}$

Characteristic absorptions in IR spectroscopy

Wavenumber/cm ⁻¹	Bond	Compound
550–850	C–X (X = Cl, Br, I)	Haloalkanes
750–1100	C–C	Alkanes, alkyl groups
1000–1300	C–O	Alcohols, esters, carboxylic acids
1450–1650	C=C	Arenes
1600–1700	C=C	Alkenes
1650–1800	C=O	Carboxylic acids, esters, aldehydes, ketones, amides, acyl chlorides
2200–2300	C≡N	Nitriles
2500–3200	O–H	Carboxylic acids
2750–2850	C–H	Aldehydes
2850–3000	C–H	Alkanes, alkyl groups, alkenes, arenes
3200–3600	O–H	Alcohols
3300–3500	N–H	Amines, amides

Proton Chemical Shifts in Nuclear Magnetic Resonance Spectroscopy (relative to TMS)

Chemical Shift	Structure	
0.5–2.0	–CH	Saturated alkanes
0.5–5.5	–OH	Alcohols
1.0–3.0	–NH	Amines
2.0–3.0	–CO–CH	Ketones
	–N–CH	Amines
	C ₆ H ₅ –CH	Arene (aliphatic on ring)
2.0–4.0	X–CH	X = Cl or Br (3.0–4.0) X = I (2.0–3.0)
4.5–6.0	–C=CH	Alkenes
5.5–8.5	RCONH	Amides
6.0–8.0	–C ₆ H ₅	Arenes (on ring)
9.0–10.0	–CHO	Aldehydes
10.0–12.0	–COOH	Carboxylic acids

These chemical shifts are concentration and temperature dependent and may be outside the ranges indicated above.

New
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GCE

CHEMISTRY DATA SHEET

GCE A/AS EXAMINATIONS

CHEMISTRY

Including the Periodic Table of the Elements

For the use of candidates taking
Advanced Subsidiary and Advanced Level
Chemistry Examinations

Copies must be free from notes or additions of any kind.
No other type of data booklet or information sheet is
authorised for use in the examinations.

I II

THE PERIODIC TABLE OF ELEMENTS

Group

III IV V VI VII 0

13 14 15 16 17 18

1 H Hydrogen 1																	4 He Helium 2
7 Li Lithium 3	9 Be Beryllium 4											11 B Boron 5	12 C Carbon 6	14 N Nitrogen 7	16 O Oxygen 8	19 F Fluorine 9	20 Ne Neon 10
23 Na Sodium 11	24 Mg Magnesium 12											27 Al Aluminium 13	28 Si Silicon 14	31 P Phosphorus 15	32 S Sulfur 16	35.5 Cl Chlorine 17	40 Ar Argon 18
39 K Potassium 19	40 Ca Calcium 20	45 Sc Scandium 21	48 Ti Titanium 22	51 V Vanadium 23	52 Cr Chromium 24	55 Mn Manganese 25	56 Fe Iron 26	59 Co Cobalt 27	59 Ni Nickel 28	64 Cu Copper 29	65 Zn Zinc 30	70 Ga Gallium 31	73 Ge Germanium 32	75 As Arsenic 33	79 Se Selenium 34	80 Br Bromine 35	84 Kr Krypton 36
85 Rb Rubidium 37	88 Sr Strontium 38	89 Y Yttrium 39	91 Zr Zirconium 40	93 Nb Niobium 41	96 Mo Molybdenum 42	98 Tc Technetium 43	101 Ru Ruthenium 44	103 Rh Rhodium 45	106 Pd Palladium 46	108 Ag Silver 47	112 Cd Cadmium 48	115 In Indium 49	119 Sn Tin 50	122 Sb Antimony 51	128 Te Tellurium 52	127 I Iodine 53	131 Xe Xenon 54
133 Cs Caesium 55	137 Ba Barium 56	139 La [*] Lanthanum 57	178 Hf Hafnium 72	181 Ta Tantalum 73	184 W Tungsten 74	186 Re Rhenium 75	190 Os Osmium 76	192 Ir Iridium 77	195 Pt Platinum 78	197 Au Gold 79	201 Hg Mercury 80	204 Tl Thallium 81	207 Pb Lead 82	209 Bi Bismuth 83	210 Po Polonium 84	210 At Astatine 85	222 Rn Radon 86
223 Fr Francium 87	226 Ra Radium 88	227 Ac [†] Actinium 89	261 Rf Rutherfordium 104	262 Db Dubnium 105	266 Sg Seaborgium 106	264 Bh Bohrium 107	277 Hs Hassium 108	268 Mt Meitnerium 109	271 Ds Darmstadtium 110	272 Rg Roentgenium 111	285 Cn Copernicium 112						

* 58–71 Lanthanum series
† 90–103 Actinium series

a^x_b

a = relative atomic mass (approx)
x = atomic symbol
b = atomic number

140 Ce Cerium 58	141 Pr Praseodymium 59	144 Nd Neodymium 60	145 Pm Promethium 61	150 Sm Samarium 62	152 Eu Europium 63	157 Gd Gadolinium 64	159 Tb Terbium 65	162 Dy Dysprosium 66	165 Ho Holmium 67	167 Er Erbium 68	169 Tm Thulium 69	173 Yb Ytterbium 70	175 Lu Lutetium 71
232 Th Thorium 90	231 Pa Protactinium 91	238 U Uranium 92	237 Np Neptunium 93	242 Pu Plutonium 94	243 Am Americium 95	247 Cm Curium 96	245 Bk Berkelium 97	251 Cf Californium 98	254 Es Einsteinium 99	253 Fm Fermium 100	256 Md Mendelevium 101	254 No Nobelium 102	257 Lr Lawrencium 103