Cambridge International AS & A Level

CHEMISTRY

Paper 4 A Level Structured Questions MARK SCHEME Maximum Mark: 100 9701/42 May/June 2021

Published

This mark scheme is published as an aid to teachers and candidates, to indicate the requirements of the examination. It shows the basis on which Examiners were instructed to award marks. It does not indicate the details of the discussions that took place at an Examiners' meeting before marking began, which would have considered the acceptability of alternative answers.

Mark schemes should be read in conjunction with the question paper and the Principal Examiner Report for Teachers.

Cambridge International will not enter into discussions about these mark schemes.

Cambridge International is publishing the mark schemes for the May/June 2021 series for most Cambridge IGCSE[™], Cambridge International A and AS Level components and some Cambridge O Level components.

Generic Marking Principles

These general marking principles must be applied by all examiners when marking candidate answers. They should be applied alongside the specific content of the mark scheme or generic level descriptors for a question. Each question paper and mark scheme will also comply with these marking principles.

GENERIC MARKING PRINCIPLE 1:

Marks must be awarded in line with:

- the specific content of the mark scheme or the generic level descriptors for the question
- the specific skills defined in the mark scheme or in the generic level descriptors for the question
- the standard of response required by a candidate as exemplified by the standardisation scripts.

GENERIC MARKING PRINCIPLE 2:

Marks awarded are always **whole marks** (not half marks, or other fractions).

GENERIC MARKING PRINCIPLE 3:

Marks must be awarded **positively**:

- marks are awarded for correct/valid answers, as defined in the mark scheme. However, credit is given for valid answers which go beyond the scope of the syllabus and mark scheme, referring to your Team Leader as appropriate
- marks are awarded when candidates clearly demonstrate what they know and can do
- marks are not deducted for errors
- marks are not deducted for omissions
- answers should only be judged on the quality of spelling, punctuation and grammar when these features are specifically assessed by the question as indicated by the mark scheme. The meaning, however, should be unambiguous.

GENERIC MARKING PRINCIPLE 4:

Rules must be applied consistently, e.g. in situations where candidates have not followed instructions or in the application of generic level descriptors.

GENERIC MARKING PRINCIPLE 5:

Marks should be awarded using the full range of marks defined in the mark scheme for the question (however; the use of the full mark range may be limited according to the quality of the candidate responses seen).

GENERIC MARKING PRINCIPLE 6:

Marks awarded are based solely on the requirements as defined in the mark scheme. Marks should not be awarded with grade thresholds or grade descriptors in mind.

Science-Specific Marking Principles

- 1 Examiners should consider the context and scientific use of any keywords when awarding marks. Although keywords may be present, marks should not be awarded if the keywords are used incorrectly.
- 2 The examiner should not choose between contradictory statements given in the same question part, and credit should not be awarded for any correct statement that is contradicted within the same question part. Wrong science that is irrelevant to the question should be ignored.
- 3 Although spellings do not have to be correct, spellings of syllabus terms must allow for clear and unambiguous separation from other syllabus terms with which they may be confused (e.g. ethane / ethene, glucagon / glycogen, refraction / reflection).
- 4 The error carried forward (ecf) principle should be applied, where appropriate. If an incorrect answer is subsequently used in a scientifically correct way, the candidate should be awarded these subsequent marking points. Further guidance will be included in the mark scheme where necessary and any exceptions to this general principle will be noted.

5 <u>'List rule' guidance</u>

For questions that require *n* responses (e.g. State **two** reasons ...):

- The response should be read as continuous prose, even when numbered answer spaces are provided.
- Any response marked *ignore* in the mark scheme should not count towards *n*.
- Incorrect responses should not be awarded credit but will still count towards *n*.
- Read the entire response to check for any responses that contradict those that would otherwise be credited. Credit should **not** be awarded for any responses that are contradicted within the rest of the response. Where two responses contradict one another, this should be treated as a single incorrect response.
- Non-contradictory responses after the first *n* responses may be ignored even if they include incorrect science.

6 <u>Calculation specific guidance</u>

Correct answers to calculations should be given full credit even if there is no working or incorrect working, **unless** the question states 'show your working'.

For questions in which the number of significant figures required is not stated, credit should be awarded for correct answers when rounded by the examiner to the number of significant figures given in the mark scheme. This may not apply to measured values.

For answers given in standard form (e.g. $a \times 10^n$) in which the convention of restricting the value of the coefficient (a) to a value between 1 and 10 is not followed, credit may still be awarded if the answer can be converted to the answer given in the mark scheme.

Unless a separate mark is given for a unit, a missing or incorrect unit will normally mean that the final calculation mark is not awarded. Exceptions to this general principle will be noted in the mark scheme.

7 <u>Guidance for chemical equations</u>

Multiples / fractions of coefficients used in chemical equations are acceptable unless stated otherwise in the mark scheme.

State symbols given in an equation should be ignored unless asked for in the question or stated otherwise in the mark scheme.

Question		Answer			Marks
1(a)(i)	[1s ²] 2s ² 2p ⁶ 3s ² 3p ⁶ 3d ³				1
1(a)(ii)	(a molecule or ion forn	ned by a central) metal atom/ion surrounded	by / bonded to one or mo	re ligands	1
1(b)	[Cr(H₂O) ₆]³+(aq)	formula of chromium species formed	type of reaction		5
	+ NaOH(aq)	Cr(OH) ₃ or Cr(OH) ₃ (H ₂ O) ₃	precipitation		
	+ H ₂ O ₂ (aq)	Cr ₂ O ₇ ²⁻ / CrO ₄ ²⁻	redox / oxidation		
	+ excess NH ₃ (aq)	Cr(NH ₃) ₆ ³⁺	ligand substitution		
	chromium species: one type of reaction: two ce	e mark for each correct species prrect for one mark and three correct for two	marks		
1(c)	M1: ∆ <i>E</i> is different OR	energy gap between d-orbitals is different			2
	M2: different frequenc OR different energy / li	y / wavelength is absorbed ght in visible region is absorbed			
1(d)(i)	ethanoate ions are bid OR ethanoate ions for OR ethanoate ions do	entate whereas as H ₂ O are monodentate m two dative bonds whereas H ₂ O forms one nate two lone pairs whereas H ₂ O donates on	(dative) bond e (lone) pair		1
1(d)(ii)	(coordination number)	six AND (geometry around Cr) octahedral			1
1(d)(iii)	coordinate / (dative) co	valent			1
1(e)(i)	$4\mathrm{Cr}^{2+} + \mathrm{O}_2 + 4\mathrm{H}^+ \rightarrow 4\mathrm{C}$	$Cr^{3+} + 2H_2O \mathbf{OR} 2Cr^{2+} + O_2 + 2H^+ \rightarrow 2Cr^{3+} + Cr^{3+}$	H ₂ O ₂		2
	M1: correct species M2: balancing				
1(e)(ii)	<i>E</i> ^e _{cell} = 1.23 − (−0.41) = value linked to (e)(i)	$= (+)1.64 \vee \mathbf{OR} \ E^{\circ}_{\text{cell}} = 0.68 - (-0.41) = (+)1$.09 V		1

Question	Answer	Marks
2(a)	M1: increases down the group	3
	M2: radius / size of cation / M^{2+} increases OR charge density of cation / M^{2+} decreases	
	M3: less polarisation / less distortion of anion / NO ₃ ⁻ ion OR less weakening of NO bond	
2(b)(i)	$Pb(NO_3)_2 \rightarrow PbO + 2NO_2 + \frac{1}{2}O_2$	1
2(b)(ii)	lead nitrate / Pb(NO ₃) ₂ would decompose more / easier AND as Pb^{2+} is smaller / Pb ²⁺ has larger charge density (so more polarising)	1
2(c)(i)	$BaC_2O_4 \rightarrow BaO + CO_2 + CO OR BaC_2O_4 \rightarrow BaO + 2CO + \frac{1}{2}O_2$	1
2(c)(ii)	M1: [a] initial moles $MnO_{4^{-}} = 0.0200 \times 0.050 = 1.00 \times 10^{-3}$ [b] moles $Fe^{2+} = 0.050 \times 0.0304 = 1.52 \times 10^{-3}$	4
	[b] moles MnO ₄ ⁻ reacted = $1.00 \times 10^{-3} - 3.04 \times 10^{-4} = 6.96 \times 10^{-4}$	
	M3: moles $C_2O_4^{2-}$ reacted = $6.96 \times 10^{-4} \times 5/2 = 1.74 \times 10^{-3}$	
	M4: mass of BaC₂O₄ = 225.3 × 1.74 × 10 ⁻³ = 0.392 g % Purity of BaC₂O₄ = 100 × 0.392/0.50 = 78.4	
2(d)	M1: $[OH^{-}] = 2 \times 0.12 = 0.24 \text{ (mol dm}^{-3})$ $[H^{+}] = 1 \times 10^{-14}/0.24 = 4.17 \times 10^{-14} \text{ / pOH} = -\log(0.24) \text{ OR } 0.62$	2
	M2: pH = -log[H ⁺] = 13.4 OR pH = 14 - 0.6 = 13.4	

Question	Answer	Marks
3(a)(i)	M1: voltage of an electrode / a half-cell compared to / connected to (standard) hydrogen electrode / half-cell	2
	M2: (at concentration of) 1 mol / dm³ AND (pressure of) 1 atm / 101 kPa (or in Pa) AND 298 K / 25°C	
3(a)(ii)	E + redox system	1
	Most B negative	
	↑ C	
	Least A A	
3(a)(iii)	$M_{N_{N_{N_{N_{N_{N_{N_{N_{N_{N_{N_{N_{N_$	3
3(b)(i)	E_{cell}^{Θ} = 1.07 – 0.80 = (+)0.27 V AND direction of electron flow = Ag ⁺ / Ag to Br ₂ / Br ⁻	1
3(b)(ii)	M1: <i>E</i> ^e _{cell} 3rd box ticked	2
	M2: [Ag ⁺] decreases AND so (Ag ⁺ /Ag) equilibrium shifts to the left OR [Ag ⁺] decreases AND <i>E</i> for (Ag ⁺ /Ag) becomes less positive / more negative	

Question	Answer	Marks
3(c)(i)	(a species) that uses / shares a lone pair of electrons to form a coordinate bond to a metal atom / ion	1
3(c)(ii)	$K_{\rm c} = [Ag(S_2O_3)_2^3] [Br] / [S_2O_3^2]^2$	1
3(c)(iii)	M1: $K_c = K_{stab} \times K_{sp} = 15.7$ M2: 1 OR none / no units	2
3(d)	M1: highest $[Ag(CN)_2]^- [Ag(S_2O_3)_2]^{3-} [Ag(NH_3)_2]^+$ lowest	2
	M2: K_{stab} [Ag(CN) ₂] ⁻ is highest / [Ag(CN) ₂] ⁻ is the most stable OR higher K_{stab} forms the more stable complex	

Question	Answer	Marks
4(a)(i)	M1: energy change when 1 mole of a ionic compound is formed M2: from its gaseous ions under standard conditions	2
4(a)(ii)	$\Delta H_{sol} = (-2099) + (2 \times -378) - (-2824)$ $\Delta H_{sol} = -31 \text{ kJ mol}^{-1}$	2
	M1: use of ×2 as only multiplier M2: correct signs and evaluation	
4(a)(iii)	M1: Cu ²⁺ is smaller OR Cu ²⁺ has a higher charge density	2
	M2: Cu ²⁺ attracts water molecules more / stronger OR (Cu ²⁺) forms stronger ion-dipole forces to water molecules	
4(b)(i)	anode: chlorine / C <i>l</i> ₂ cathode: hydrogen / H ₂	1
4(b)(ii)	M1: Q = 0.75 × 60 × 60 = 2700 C AND 96 500 or 193 000 used	2
	M2: [a] moles of Ca = 2700 / 193 000 = 0.0140 [b] mass = 0.0140 × 40.1 = 0.56 g	

Question	Answer	Marks
4(c)(i)	measure / degree of (dis)order / randomness (of a system) OR the number of possible arrangements of the particles and their energy (in a given system)	
4(c)(ii)	ΔS is negative ΔS is zero ΔS is positive	1
	solid dissolving in water	
	water solidifying to ice 🗸	
4(c)(iii)	$\frac{positive}{\Delta G} = \frac{positive}{25 + 50 + 75 + 100} temperature °C$ $\frac{requestive}{1000} temperature °C$ $\frac{two correct for 1 mark, three correct for two marks:}{10000} temperature temp$	
4(d)	M1: ΔH negative / – , ΔS negative / – M2: as temperature increase, ΔG becomes (more) positive / less negative ora OR at low(er) T, (ΔH more negative than T ΔS) so ΔG is negative OR at high(er) T, (ΔH less negative than T ΔS) so ΔG is positive	2

Question	Answer	Marks
5(a)	M1: ethylamine phenylamine 4-nitrophenylamine most basic least basic	4
	 M2 / 3 / 4: explanation two correct for one mark, three correct for two marks, four correct for three marks (basicity linked to) lone pair / p orbital on N AND being able accept / donate to / coordinate to a proton / H⁺ 	
	 ethyl / alkyl group is electron donating / has a positive inductive effect (so lone pair on N is more able to accept a proton) 	
	 (phenylamines are less basic than ethylamine as) p orbital / lone pair on N is delocalised (into the ring so less able to accept a proton) 	
	 (4-nitrophenylamine is less basic than phenylamine as) nitro / NO₂ group is electron withdrawing (so lone pair on N is less able to accept a proton) 	
5(b)(i)	O_2N \longrightarrow N	1
5(b)(ii)	M1: step 1: HNO ₂ , ≤ 10°C OR NaNO ₂ , HC <i>l</i> (aq), ≤ 10°C	2
	M2: step 2: NaOH / alkaline AND 1-naphthol / α-naphthol / structure below	
5(c)(i)	4-bromo-2-nitrobenzoic acid OR 4-bromo-2-nitro(-1-)benzenecarboxylic acid	1

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Question	Answer	Marks
5(c)(ii)	$E = \bigcup_{Br}^{CH_3} OR \qquad \bigcup_{Br}^{CH_3}$ $F = \bigcup_{Br}^{CH_3} NO_2$	2
5(c)(iii)	M1: step 1 conc. H_2SO_4 and conc. HNO_3	3
	M2: step 2 Br_2 and $AlBr_3$	
	M3: step 3 hot (alkaline / acidified) MnO ₄ ⁻ / KMnO ₄	

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Question	Answer	Marks
6(a)	three peaks four peaks	4
	five peaks six peaks	
	correct isomers and correct assignment to peaks: mark as $\cdot \checkmark \cdot \checkmark \cdot \checkmark \cdot \checkmark$	
6(b)(i)	$RCl + AlCl_3 \rightarrow R^+ + AlCl_4^-$	1
	OR $Cl(CH_2)_3COOH + AlCl_3 \rightarrow {}^+(CH_2)_3COOH + AlCl_4^-$	
6(b)(ii)	$W \longrightarrow V \longrightarrow V = V = V = V = V = V = V = V = V$	2

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Question	Answer	Marks
6(b)(iii)	SOC <i>l</i> ₂ OR PC <i>l</i> ₅ ALLOW PC <i>l</i> ₃ AND heat	1
6(b)(iv)	$ \overbrace{R^{+}}^{\text{intermediate}} \qquad $	3

Question	Answer	Marks
7(a)(i)	the power to which a concentration of a reactant is raised in the rate equation / law	1
7(a)(ii)	M1: (using expt 1 and 3) as $[ClO_2] \times 2.5$ rate $\times 6.25$ so 2nd order	2
	M2: (using expt 1 and 2) as $[OH^-] \times 4$ rate $\times 4$ so 1st order	
7(a)(iii)	rate = k[C <i>l</i> O ₂] ² [OH ⁻]	1
7(a)(iv)	M1: $k = rate / [ClO_2]^2[OH^-]$ $k = 7.20 \times 10^{-4} / (0.02)^2 (0.03)$ k = 60 M2: mol ⁻² dm ⁶ min ⁻¹	2

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Question	Answer	Marks
7(b)(i)	structure of phenol: C ₆ H ₅ OH OR	1
7(b)(ii)	tangent drawn correctly AND rate = $0.015/260 = 5.8 \times 10^{-5}$ ALLOW values consistent with tangent drawn at 100 sec	1
7(c)	AND half-life decreases (1st box)	1

Question	Answer				
8(a)	M1: acyl chlorides > alkyl chlorides > aryl chlorides fastest slowest				
	 M2 / 3: Any two from: acyl chlorides carbon in C-Cl bond is more electron deficient since it is also attached to an oxygen atom OR C-Cl bond is weakest / weakened in acyl chlorides since it is also attached to an oxygen atom / two electronegative atoms aryl chlorides (no hydrolysis) C-Cl bond is part of delocalised system / partially double bond character (so C-Cl bond is 				
	 stronger) OR lone pair / p-orbital on Cl delocalises with π ring (so C-Cl bond is stronger) alkyl chlorides C-Cl bond strengthened by electron donating effect / positive inductive effect of alkyl / R group (as compared to acyl chlorides) OR carbon atom has a smaller δ+ and the C-Cl bond is stronger (than the C-Cl bond in COCl) due to (the carbon) being only attached to one electronegative atom 				
8(b)	HO HO HO HO HO HO HO HO	3			
	ALLOW amine salt for the third structure – mono or di ion				
8(c)(i)	ОН	1			

Question	Answer				Marks	
8(c)(ii)	chemical shift (δ)	environment of proton	splitting pattern (words required)	number of ¹ H atoms responsible for the peak		3
	0.95	alkane / CH3	doublet	6		
	1.90	alkane / CH ALLOW alkyne	multiplet	1		
	2.20	R / alkyl / CH ₂ next to C=O / COOH	doublet	2		
	mark as • • •	/ •• √ •• √	1	l		