

Mark Scheme (Results)

Summer 2025

Pearson Edexcel GCE A Level In Chemistry (9CH0)

Paper 01: Advanced Inorganic and Physical

Chemistry

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General Marking Guidance

- All candidates must receive the same treatment. Examiners must mark the first candidate in exactly the same way as they mark the last.
- Mark schemes should be applied positively. Candidates must be rewarded for what they have shown they can do rather than penalised for omissions.
- Examiners should mark according to the mark scheme not according to their perception of where the grade boundaries may lie.
- There is no ceiling on achievement. All marks on the mark scheme should be used appropriately.
- All the marks on the mark scheme are designed to be awarded. Examiners should always award full marks if deserved, i.e. if the answer matches the mark scheme. Examiners should also be prepared to award zero marks if the candidate's response is not worthy of credit according to the mark scheme.
- Where some judgement is required, mark schemes will provide the principles by which marks will be awarded and exemplification may be limited.
- When examiners are in doubt regarding the application of the mark scheme to a candidate's response, the team leader must be consulted.
- Crossed out work should be marked UNLESS the candidate has replaced it with an alternative response.

Using the Mark Scheme

Examiners should look for qualities to reward rather than faults to penalise. This does NOT mean giving credit for incorrect or inadequate answers, but it does mean allowing candidates to be rewarded for answers showing correct application of principles and knowledge. Examiners should therefore read carefully and consider every response: even if it is not what is expected it may be worthy of credit.

The mark scheme gives examiners:

- an idea of the types of response expected
- how individual marks are to be awarded
- the total mark for each question
- examples of responses that should NOT receive credit.

/ means that the responses are alternatives and either answer should receive full credit.

() means that a phrase/word is not essential for the award of the mark, but helps the examiner to get the sense of the expected answer.

Phrases/words in **bold** indicate that the <u>meaning</u> of the phrase or the actual word is **essential** to the answer. ecf/TE/cq (error carried forward) means that a wrong answer given in an earlier part of a question is used correctly in answer to a later part of the same question.

Candidates must make their meaning clear to the examiner to gain the mark. Make sure that the answer makes sense. Do not give credit for correct words/phrases which are put together in a meaningless manner. Answers must be in the correct context.

Quality of Written Communication

Questions which involve the writing of continuous prose will expect candidates to:

- write legibly, with accurate use of spelling, grammar and punctuation in order to make the meaning clear
- select and use a form and style of writing appropriate to purpose and to complex subject matter
- organise information clearly and coherently, using specialist vocabulary when appropriate.

Full marks will be awarded if the candidate has demonstrated the above abilities.

Questions where QWC is likely to be particularly important are indicated (QWC) in the mark scheme, but this does not preclude others.

Question Number	Answer		Additional Guidance	Mark
1(a)	An answer that makes reference to the following points:			(2)
	• (relative isotopic mass) is the mass of an atom (of the isotope / element)	(1)	Allow mass of an isotope of an atom	
			Do not award any reference to average / mean / weighted or moles in M1 or M2, but penalise only once if in both	
	• relative to 1/12 th (the mass of an atom) of carbon-12	(1)	Allow for two marks Mass of one atom of an isotope × 12 Mass of one atom of carbon-12	
			COMMENT M1 can be awarded if atom mentioned somewhere in response	

Question Number	Answer		A	Additional Guidance			Mark
1(b)	An answer that makes reference to the following points:						(2)
	• all isotopes have 18 protons (and 18 electrons)	(1)	Do not award inc	correct number	r of electrons		
	• the isotopes have different numbers of neutrons, 18,	(1)	Isotope	Protons	Neutrons		
	20 and 22	(1)	³⁶ Ar	18	18		
			³⁸ Ar	18	20		
			⁴⁰ Ar	18	22		
			If no other mark same number of neutrons for 1 m Ignore same ator number	protons but a cark	different number	of	

Question Number	Answer	Additional Guidance	Mark
1(c)		Example of calculation	(2)
	• correct expression for calculation of $A_{\rm r}$ (1)	$A_{\rm r} = \underbrace{((35.97 \times 0.337) + (37.96 \times 0.0630) + (39.96 \times 99.6))}_{100}$ (= 39.945)	
	• value given to 2 dp (1)	= 39.95	
		TE on correct calculation from incorrect transposed numbers or use of whole numbers for relative isotopic mass	
		Ignore g mol ⁻¹ but penalise other units	
		Correct answer with no working scores 2	

(Total for Question 1 = 6 marks)

Question Number	Answer		Additional Guidance	Mark
2(a)	oxygen below both nitrogen and fluorine and above carbon	(1) (1) (1)	Example of graph 2500 2000 First ionisation 1500 energy / kJ mol ⁻¹ 1000 H He Li Be B C N O F Ne Na Element Accept just points	(3)
			Penalise plots on x-axis between elements once only	

Question Number	Answer		Additional Guidance	Mark
2(b)	An explanation that makes reference to the following points:		Allow reverse argument throughout Penalise omission of 'outer' once only	(4)
			Ignore electron configurations unless incorrect	
	(the value for) beryllium is higher than lithium • as beryllium has one more proton / a higher nuclear charge	(1)	Allow greater proton number	
	 shielding is the same / similar in beryllium and lithium or electron repulsion is the same / similar in beryllium and lithium or the outer electron is a same /similar distance from the nucleus or the outer electron is in the same / 2s orbital 	(1)	Ignore reference to atomic or ionic radius	
	 (the value for) beryllium is higher than boron as the outer electron in boron is in the (2)p orbital / subshell (rather than the 2s orbital in beryllium) 	(1)	Allow outer electron in B is in a higher energy level	
	 the effect of the additional proton / increased nuclear charge in boron is outweighed by increased shielding / repulsion (from inner subshell) 	(1)	Allow by increased distance from the nucleus (of the outer electron in B)	

Question Number	Answer	Additional Guidance	Mark
2(c)	An answer that makes reference to the following point:		(1)
	correct equation with state symbols	$Be^{+}(g) \rightarrow Be^{2+}(g) + e^{(-)}$	
		Allow Be ⁺ (g) - $e^{(-)} \rightarrow Be^{2+}(g)$	
		Ignore state symbol for the electron Do not allow	

Question Number	Answer	Mark
2(d)	The only correct answer is C (4s)	(1)
	A is not correct because this would be Mg	
	B is not correct because this could relate to a p-block element in period 3	
	D is not correct because this could relate to a p-block element in period 4	

Question Number	Answer	Mark
2(e)	The only correct answer is A (S ²⁻)	(1)
	B is not correct because the ionic radius of Cl^- is smaller than S^{2-} as it has one more proton	
	C is not correct because the ionic radius of K^+ is smaller than both S^{2-} and Cl^- as it has more protons	
	$m{D}$ is not correct because Ca^{2+} has the smallest ionic radius since it has the most protons	

Question Number	Answer		Additional Guidance	Mark
2(f)	An answer that makes reference to the following points:		Penalise reference to covalent bonds breaking once only	(6)
	hydrogen bonding is the strongest intermolecular force	(1)	Allow H bonds take more energy to break than other types of intermolecular force	
	• there is hydrogen bonding between H ₂ O (molecules) and between HF (molecules)	(1)		
	• H ₂ O has two hydrogen bonds (per molecule, as it has two lone pairs on the oxygen) and so its boiling temperature is	(1)	Allow water can form twice any many hydrogen bonds (than HF)	
	higher than HF		Ignore H ₂ O has more hydrogen bonds	
			Do not award H ₂ O forms 3 hydrogen bonds, HF forms 2 hydrogen bonds	
	there are permanent dipoles between the HCl molecules	(1)	Allow HCl has dipole-dipole (forces) Ignore HI has a permanent dipole	
	there are stronger / more London forces between HI (molecules) than HCl (molecules) because there are more	(1)	Accept van der Waals / temporary dipole – induced dipole	
	electrons (in a molecule of HI)		Allow London forces require more energy	
	• (the increased) London forces between HI (molecules) outweigh the permanent dipole forces (between HCl	(1)	to break / be overcome than permanent dipole (forces)	
	(molecules)		Allow London forces outweigh the (greater) electronegativity difference between H and Cl	

(Total for Question 2 = 16 marks)

Question Number	Answer	Additional Guidance	Mark
3(a)	An answer that makes reference to the following point:		(1)
	heat energy must be supplied so the energy change is not just due to the thermal decomposition reaction or it is difficult to determine when the (decomposition) reaction is complete	Allow it is impossible to measure ΔT when heat energy is supplied / cannot measure temperature (easily) whilst heating Ignore the reaction is not complete	
	or it's difficult to (accurately) measure the temperature of a solid (whilst heating)	Allow you cannot measure the temperature of a solid (easily)	

Question Number	Answer	Additional Guidance		
3(b)(i)	. 1 11 . 6			(1)
Clip all of	 correct values in table for mass and temperature 	Substance	Value	
(b)	•	Mass of weighing boat with calcium carbonate / g	(4.30)	
		Mass of weighing boat after emptying out calcium carbonate / g	(0.20)	
		Mass of calcium carbonate used / g	4.10	
		Start temperature / °C	(19.0)	
		Highest temperature / °C	24.5	
		Temperature change / °C	(5.5)	

Question Number	Answer	Additional Guidance	Mark
3(b)(ii)	• mol of HCl (1)	Example of calculation $= 3.00 \times \frac{30}{1000} = 0.090$	(2)
	mol of HCl required and more mol HCl present than required (1)	= $0.041 \times 2 = 0.082$ and 0.09 > 0.082 (mol)	
		Allow $0.090 \div 2 = 0.045$ (mol CaCO ₃ that you would need to react with 0.09 mol HCl and) $0.045 > 0.041$ (mol)	
		Ignore SF Allow calculation of excess HCl by subtraction from 0.090	

Question Number	Answer		Additional Guidance	Mark
3(b)(iii)	• energy change (Q) for Reaction 2	(1)	Example of calculation energy change = $30 \times 4.18 \times 5.5$ = $689.7 \text{ (J)} / 0.6897 \text{ (kJ)}$	(3)
	• enthalpy change $(\Delta_r H_2)$ for Reaction 2	(1)	Ignore any signs in M1 enthalpy change = $\frac{0.6897}{0.041}$ (kJ) / $\frac{689.7}{0.041}$ (J)	
	answer with negative sign	(1)	$= -16.822 \text{ (kJ mol}^{-1}) / -16822 \text{ J mol}^{-1} / -17.0 \text{ (kJ mol}^{-1}) / -17000 \text{ J mol}^{-1}$	
			Accept correct answer with no working Allow TE throughout Ignore SF except 1SF	

Question Number	Answer		Additional Guidance	Mark
3(b)(iv)	An answer that makes reference to the following points: • correct species with state symbols in the bottom box • arrow on left hand side in correct direction and labelled with 2HCl((aq))	(1)		(2)

Question Number	Answer	Additional Guidance	Mark
3(b)(v)	calculation of enthalpy change of thermal decomposition of calcium carbonate	Example of calculation $\Delta_{\rm r}H_I = -16.8176.3 = (+) \ 159.5 \ \rm kJ \ mol^{-1}$ TE on their incorrect value from 3(b)(iii) No TE on incorrect cycle	(1)

Question Number	Answer	Additional Guidance	Mark
3(c)(i)	An explanation that makes reference to the following points:	Example of calculation	(3)
	• calculation of $\Delta S^{\Theta}_{\text{system}}$ (1)	= $(39.7 + 213.6) - (92.9)$ = $(+) 160.4 (J K^{-1} mol^{-1}) / (+) 0.1604 (kJ K^{-1} mol^{-1})$	
	• calculation of ΔG^{Θ} (1)	$(\Delta G = \Delta H - T\Delta S_{\text{system}})$ = $(+178.2 \times 1000) - (298 \times 160.4)$ = $(+) 130 400 \text{ (J mol}^{-1})/(+) 130.4 \text{ (kJ mol}^{-1})$ Allow TE from M1 to M2	
	 reason why not feasible Alternative for M2 and 3 using ΔS_{total} 	ΔG positive / > 0 so not feasible No TE on negative value of ΔG	
	• $\Delta S_{\text{surroundings}} = -\frac{178\ 200}{298} = -597.99 \text{ (J K}^{-1} \text{ mol}^{-1}\text{)}$ (1)		
	• $\Delta S_{\text{total}} = 160.4 - 597.99 = -437.59 \text{ (J K}^{-1} \text{ mol}^{-1}\text{)} \text{ and}$ Value is negative and so not feasible (1)	Ignore minor rounding error in decimal places of ΔS_{total}	
		Ignore SF except 1SF	

Question Number	Answer		Additional Guidance	Mark
3(c)(ii)	• rearrangement of ΔG expression and calculation of feasible temperature	(1)	Example of calculation $T = \Delta H / \Delta S_{\text{system}}$ $T = 178.2 / 0.1604 = 1111 \text{ (K)}$ Allow 1110 (K) for M1	(2)
	conversion to degrees Celsius	(1)	1111 – 273 = 838 (°C) Allow 1110 – 273 = 837 (°C)	
			Accept method that uses $\Delta S_{\text{total}} = 0$ $\Delta S_{\text{total}} = 0 = 160.4 - (178\ 200\ /\ T)$ $T = -178\ 200\ /\ -160.4 = 1111(K)$ 1111 - 273 = 838	
			838 / 837 with or without working scores both marks	
			Do not award 840°C	
			Allow TE from M1 to M2 for values > - 273 °C	

Question Number	Answer	Additional Guidance	Mark
3(d)	An explanation that makes reference to the following points:	Accept reverse argument Penalise reference to just Mg or Ca once only	(3)
	• Mg ²⁺ is a smaller cation / ion / has a smaller ionic radius / (1) has a greater charge density	Ignore 'more covalent character' / comments related to atomic radius unless they contradict the relative size of the ions	
	• Mg ²⁺ polarises the carbonate / CO ₃ ²⁻ / anion more (1)	Allow Mg ²⁺ has greater polarising power / Mg ²⁺ distorts the electron cloud more Ignore Mg ²⁺ polarises the C-O bond more	
	• (so in MgCO ₃) the bond between C and O is weakened more (1)	Allow bond(s) within the anion are weakened more Allow C-O (bond) / C=O (bond) is weakened more	

(Total for Question 3 = 18 marks)

Question Number	Answer		Additional Guidance							
4(a)			Example of calcu	Example of calculation						
				C ₂ H ₄ (g)	H ₂ O(g)	C ₂ H ₅ OH(g)				
			Initial moles	90.0	54.0	0.0				
			Change in moles	-4.5	-4.5	+4.5				
	calculation of moles at equilibrium	(1)	Equilibrium moles	85.5	49.5	4.5				
	calculation of mole fractions	(1)	Mole fraction	$85.5 \div 139.5$ = 0.6129	$49.5 \div 139.5 \\ = 0.3548$	$4.5 \div 139.5 \\ = 0.0323$				
	calculation of partial pressures	(1)	Partial pressure	0.6129×60 = 36.77	0.3548×60 = 21.29	0.0323 × 60 = 1.94				
	• expression for K_p	(1)	$K_p = p(C_2H_5OH_2)$ $p(C_2H_4) p(H_2A)$ Allow p inside by Allow p Allow without by Do not award square	eO) racket rackets						
	• value of K_p and units	(1)	$K_p = \frac{1.94}{(36.77 \times 21)}$ = 0.00248 / 2. Allow TE throug Ignore SF except Correct answer v K_p with no early	$0.48 \times 10^{-3} / 0.00$ chout for 1 SF with no working		3 and M5				

Question Number	Answer		Additional Guidance	Mark
4(b)(i)	An answer that makes reference to the following points: • (so increased pressure) will increase the yield of ethanol	(1)	Mark points independently	(2)
	because the position of equilibrium will move to the RHS as there are less moles / molecules / particles (of gas) on the RHS	(1)	Allow because the position of equilibrium will move to the products side as there are less moles / molecules (of gas) on the RHS Allow forward direction for RHS	

Question Number	Answer	Mark
4(b)(ii)	The only correct answer is C (decreasing the temperature)	(1)
	A is not correct as changing the catalyst might affect the rate at which equilibrium was established, but would have no effect on the value of K_p	
	${\it B}$ is not correct because although increasing the pressure would allow equilibrium to be established faster, this would not affect the value of $K_{\rm p}$	
	$m{D}$ is not correct because increasing the volume would mean that equilibrium would be established more slowly, but this would not affect the value of $K_{p.}$	

(Total for Question 4 = 8 marks)

Question Number	Answer	Mark
5(a)	The only correct answer is C (acid H ₃ O ⁺ , conjugate base H ₂ O)	(1)
	A is not correct as H_2O is not the conjugate base of HNO_3	
	B is not correct because the NO_3^- is the conjugate base and HNO_3 is the acid	
	${m D}$ is not correct because the H_2O is the conjugate base and H_3O^+ is the acid	

Question Number	Answer	Additional Guidance	Mark
5(b)	An answer that makes reference to the following point:		(1)
	$ \bullet (pH =) - \log_{10} [H^{+}(aq)] $ or	Allow log / lg for log ₁₀	
	$(pH =) -log_{10} [H_3O^+(aq)]$	Ignore missing (aq)	
		Do not award —log conc H ⁺	
		Do not award round brackets / no brackets for concentration, but allow round brackets around the square brackets e.g. $-\log_{10}\left([H^+(aq)]\right)$	

Question Number	Answer	Mark
5(c)	The only correct answer is B (3.16×10^{-5})	(1)
	$m{A}$ is not correct because this is the value for the $m{H}^+$ concentration	
	C is not correct as the value for the concentration has been doubled	
	$m{D}$ is not correct because $10^{4.5}$ has been used instead of $10^{-4.5}$	

(Total for Question 5 = 3 marks)

Question Number	Answer	Mark
6(a)(i)	The only correct answer is D ([Ar] 3d ⁵ 4s ¹ , [Ar] 3d ³)	(1)
	A is not correct because neither of the electronic configurations is correct	
	B is not correct because the Cr atom is correct, but the Cr^{3+} ion is not correct	
	C is not correct because the Cr atom is not correct, but the Cr^{3+} ion is correct	

Question Number	Answer	Additional Guidance	Mark
6(a)(ii)	An answer that makes reference to the following point:		(1)
	• (Cr) is a transition metal because it forms (one or more stable) ion(s) with incompletely filled <i>d</i> -orbital(s) / <i>d</i> -subshell	Ignore d-shell	

Question Number	Answer		Additional Guidance	Mark
6(b)(i)	 An answer that makes reference to the following points: six water ligands around chromium ion with 3D structure indicated (ignore connectivity for M1) dative bonds between water ligands and chromium indicated by arrows from the oxygen of the water ligands to the chromium ion or bonds to Cr from lone pairs shown on the oxygen atoms 	(1)	Example of structure 3+ Ignore missing square brackets and charge Ignore incorrect charge Ignore comments linked to shape names / bond angles Allow bonds labelled as dative / co-ordinate	(2)

Question Number	Answer	Additional Guidance	Mark
6(b)(ii)	An answer that makes reference to the following point:		(1)
	• octahedral	Allow octahedron Do not award octagonal / octagon	

Question Number	Answer		Additional Guidance	Mark
6(c)	An answer that makes reference to the following points:			(3)
	• equation for [Cr(OH) ₃ (H ₂ O) ₃] with H ⁺	(1)		
	• equation for [Cr(OH) ₃ (H ₂ O) ₃] with OH	(1)		
	all state symbols correct	(1)	M3 dependent on M1 and M2 No TE for state symbol mark	
			Allow reversible arrows Penalise full equations once only Ignore missing square brackets	

Question Number	Answer	Additional Guidance	Mark
6(d)	An answer that makes reference to the following point:		(1)
	ligand exchange / ligand substitution	Allow ligand replacement	

Question Number	Answer		Additional Guidance	Mark
6(e)(i)	An explanation that makes reference to the following points:			(4)
	• (when a ligand approaches) d-orbitals split or d-subshell splits (into higher and lower energy levels)	(1)	Do not award the splitting of a single d –orbital Ignore d-shell	
	• electron is promoted / d-d transition / excited	(1)	Ignore electrons returning to ground state	
	• absorbing (visible) light (energy) / photons (of specific wavelength / frequency)	(1)		
	• complementary colour / seen / transmitted / reflected	(1)	Allow remaining light seen / transmitted / reflected	
			Do not award any reference to emission (of light)	

Question Number	Answer	Additional Guidance	Mark
6(e)(ii)	 An answer that makes reference to the following point: (colours are different because) with different ligands the energy gap / splitting / ΔE between the d-orbitals is different or (colours are different because) with different coordination numbers as the energy gap / splitting / ΔE between the d-orbitals is different 	Allow different ligands / coordination number so energy needed to promote electron(s) is different Allow different ligand / coordination number changes the wavelength / frequency of light absorbed Ignore different oxidation states (of metal ion)	(1)

Question Number	Answer		Additional Guidance	Mark
6(f)(i)	An answer that makes reference to the following points:			(2)
	 multiplication of equation (a) by 2 and multiplication of equation (b) by 3 	(1)	$2\text{CrO}_4^{2^-}(\text{aq}) + 8\text{H}_2\text{O}(1) + 6\text{e}^- \rightleftharpoons 2\text{Cr}(\text{OH})_3(\text{s}) + 10\text{OH}^-(\text{aq})$ Accept half equation for the chromium written as an oxidation $3\text{H}_2\text{O}_2(\text{aq}) + 6\text{e}^- \rightleftharpoons 6\text{OH}^-(\text{aq})$ M1 can be shown by use of $2\text{Cr}(\text{OH})_3$: $3\text{H}_2\text{O}_2$ in attempted equation or via equations in stem of question	
	• overall equation	(1)	2Cr(OH) ₃ (s) + 3H ₂ O ₂ (aq) + 4OH ⁻ (aq) → 2CrO ₄ ²⁻ (aq) + 8H ₂ O(l) Allow multiples Ignore state symbols even if incorrect Allow reversible arrow for full equation as long as reaction is written in the forward direction Award 2 marks for correct overall equation with no working	

Question Number	Answer	Additional Guidance	Mark
6(f)(ii)	• calculation of $E^{\bullet}_{\text{cell}}$ value (1)	Example of calculation $E^{\circ}_{\text{cell}} = 0.13 + 0.88 = (+) 1.01 \text{ (V)}$ Allow correct answer without calculation	(2)
	 E^e_{cell} / answer is positive / > 0 and the reaction is (thermodynamically) feasible (1) 	Allow as TE if negative $E^{\rm e}_{\rm cell}$ calculated Answer is negative / < 0 and the reaction is not (thermodynamically) feasible	

Question Number	Answer	Mark
6(f)(iii)	The only correct answer is A (E°_{cell}) is directly proportional to $\ln K$)	(1)
	B is not correct because E^{Θ}_{cell} is directly proportional to ΔS_{total} , and not ΔS_{system}	
	${m C}$ is not correct because $E^{m{e}}_{ m cell}$ is directly proportional to ΔS_{total} , and not $\ln \Delta S_{total}$	
	$m{D}$ is not correct because $E^{\bullet}_{\text{cell}}$ is not directly proportional to K but to $\ln K$	

(Total for Question 6 = 19 marks)

Question Number	Answer	
7(a)	The only correct answer is C (the energy change when 1 mol of an ionic substance is formed from its gaseous ions)	(1)
	A is not correct because it is the energy change not the energy absorbed	
	B is not correct because it is gaseous ions and not the elements in their standard states	
	D is not correct because it is the energy change and gaseous ions	

Question Number	Answer	Mark
7(b)	The only correct answer is A (NaCl)	(1)
	B is not correct because the I^- ion is larger than the Cl^- ion and so the lattice energy is less exothermic than that of NaCl	
	C is not correct because the K^+ ion is larger than Na^+ and so the lattice energy is less exothermic than that of $NaCl$	
	${\bf D}$ is not correct because K^+ and I^- are both larger than Na^+ and Cl^- respectively and so the lattice energy is less exothermic than that of $NaCl$	

Question Number	Answer	Mark
7(c)	The only correct answer is B (NaI)	(1)
	A is not correct because NaCl has a smaller difference between the experimental and theoretical lattice energies because the Cl ⁻ ion is not polarised as much as I ⁻ by the Na ⁺ ion as it is small compared to I ⁻ C is not correct because KCl has a smaller difference between the experimental and theoretical lattice energies because the Cl ⁻ ion is not polarised as much as I ⁻ by the K ⁺ ion as it is small compared to I ⁻ D is not correct because the difference in polarisation between K ⁺ and I ⁻ is not as great as between Na ⁺ and I ⁻ because K ⁺ is a larger ion than Na ⁺	

(Total for Question 7 = 3 marks)

Question Number	Acceptable Answer Additional Guidance		Additional Guidance	Mark
*8	coherent and logically structured answer with linkages and fully-sustained reasoning. Marks are awarded for indicative content and for how the answer is structured and shows lines of reasoning. The following table shows how the marks should be awarded for indicative content.		Guidance on how the mark scheme should be applied: The mark for indicative content should be added to the mark for lines of reasoning. For example, an answer with five indicative marking points, which is partially structured with some linkages and lines of reasoning, scores 4 marks (3 marks for indicative content and 1 mark for partial structure and some linkages and lines of reasoning).	(6)
	points seen in answer 6 5-4 3-2 1 0 The following table shows how the structure and lines of reasoning.	indicative marking points 4 3 2 1 0 marks should be awarded for	If there are no linkages between points, the same five indicative marking points would yield an overall score of 3 marks (3 marks for indicative content and no marks for linkages). In general it would be expected that 5 or 6 indicative points would get 2 reasoning marks, and 3 or 4 indicative points would get 1 mark for reasoning, and 0, 1 or 2 indicative points would score zero marks for reasoning.	
	Answer shows a coherent and logica structure with linkages and fully sustained lines of reasoning demonstrated throughout.	Number of marks awarded for structure of answer and sustained line of reasoning	If there is any incorrect chemistry, deduct mark(s) from the reasoning. If no reasoning mark(s) awarded do not deduct mark(s).	
	Answer is partially structured with some linkages and lines of reasoning			
	Answer has no linkages between points and is unstructured.	0		

 Indicative content IP1 (similarity: mechanism of action) both (increase the rate of a reaction) by lowering the activation energy by providing an alternative pathway / route for the reaction 	Ignore any references to colours of species in reactions	(6)
IP2 (similarity: oxidation number changes) both involve the change of oxidation numbers and then the return to the original oxidation number	Allow via illustration of correct changes in oxidation state Allow V ₂ O ₅ is reduced to V ₂ O ₄ , which is then oxidised to V ₂ O ₅ and Fe ²⁺ is oxidised to Fe ³⁺ which is then reduced to Fe ²⁺ Ignore catalyst is reformed / regenerated Ignore sulfur/ iodine oxidation numbers	
• IP3 (V ₂ O ₅ mechanism) reactants / SO ₂ and O ₂ adsorb(s), (reactant bonds weaken / break) and product / SO ₃ desorb(s) or	Do not award autocatalysis	
(Fe ²⁺ mechanism) (the) positive Fe ²⁺ ions attract the negative S ₂ O ₈ ²⁻ ions	Allow $S_2O_8^{2-}$ and I^- repel (so the reaction starts slowly)	
• IP4 (difference: homogeneous / heterogeneous) V ₂ O ₅ is heterogeneous / different phase / state (as reactants) and Fe ²⁺ is homogeneous / same phase / state (as reactants)	Allow specific states mentioned $-V_2O_5$ is a solid and the reactants are gases and Fe^{2+} and the reactants are all in aqueous solution	
• IP5 (Equation 1) $V_2O_5(s) + SO_2(g) \rightarrow V_2O_4(s) + SO_3(g)$ and $V_2O_4(s) + \frac{1}{2}O_2(g) \rightarrow V_2O_5(s)$	Allow multiples Ignore state symbols for IP5 and IP6	
• IP6 (Equation 2) $S_2O_8^{2-}(aq) + 2Fe^{2+}(aq) \rightarrow 2Fe^{3+}(aq) + 2SO_4^{2-}(aq)$ and $2Fe^{3+}(aq) + 2I^{-}(aq) \rightarrow I_2(aq) + 2Fe^{2+}(aq)$	If IP5 and IP6 not given allow 1IP for any two correct equations	
	 IP1 (similarity: mechanism of action) both (increase the rate of a reaction) by lowering the activation energy by providing an alternative pathway / route for the reaction IP2 (similarity: oxidation number changes) both involve the change of oxidation numbers and then the return to the original oxidation number IP3 (V₂O₅ mechanism) reactants / SO₂ and O₂ adsorb(s), (reactant bonds weaken / break) and product / SO₃ desorb(s) or (Fe²⁺ mechanism) (the) positive Fe²⁺ ions attract the negative S₂O₈²⁻ ions IP4 (difference: homogeneous / heterogeneous) V₂O₅ is heterogeneous / different phase / state (as reactants) and Fe²⁺ is homogeneous / same phase / state (as reactants) IP5 (Equation 1) V₂O₅(s) + SO₂(g) → V₂O₄(s) + SO₃(g) and V₂O₄(s) + ½₂O₂(g) → V₂O₅(s) IP6 (Equation 2) S₂O₈²⁻(aq) + 2Fe²⁺(aq) → 2Fe³⁺(aq) + 2SO₄²⁻(aq) and 	 IP1 (similarity: mechanism of action) both (increase the rate of a reaction) by lowering the activation energy by providing an alternative pathway / route for the reaction IP2 (similarity: oxidation number changes) both involve the change of oxidation numbers and then the return to the original oxidation number IP3 (V₂O₅ mechanism) reactants / SO₂ and O₂ adsorb(s), (reactant bonds weaken / break) and product / SO₃ desorb(s) or (Fe²+ mechanism) (reactants / SO₂ and O₂ adsorb(s), (reactant bonds weaken / break) and product / SO₃ desorb(s) or (Fe²+ mechanism) (the) positive Fe²+ ions attract the negative S₂O₅²- ions IP4 (difference: homogeneous / heterogeneous) V₂O₅ is heterogeneous / different phase / state (as reactants) and Fe²+ is homogeneous / same phase / state (as reactants) and V₂O₄(s) + SO₂(g) → V₂O₄(s) + SO₃(g) and V₂O₄(s) + ½O₂(g) → V₂O₃(s) IP5 (Equation 1) V₂O₅(s) + SO₂(g) → V₂O₃(s) IP6 (Equation 2) S₂O₃²-(aq) + 2Fe²⁺(aq) → 2Fe³⁺(aq) + 2SO₄²-(aq) and

(Total for Question 8 = 6 marks)

Question Number	Answer		Mark				
9(a)	• titre 3 correct						
	and initial burette reading 4 correct	Titration number	1	2	3	4	
		Final burette reading / cm ³	(15.90)	(24.00)	(15.30)	(25.00)	
		Initial burette reading / cm ³	(0.00)	(8.45)	(0.05)	9.35	
		Titre / cm ³	(15.90)	(15.55)	15.25	(15.65)	
			•	•	•	·	

Question Number	Answer	Additional Guidance	Mark
9(b)	An answer that makes reference to the following point:		(1)
	• the mean / average of the concordant titres	Allow titres 2 and 4 / 15.55 and 15.65 to identify the relevant titres	
		Do not award all four titrations added together and divided by four	

Question Number	Answer		Additional Guidance	Mark
9(c)	An answer that makes reference to the following points:			(2)
	• (from) colourless	(1)		
	• (to) (pale) pink	(1)	Do not award pink – purple / dark pink / purple / red Allow one mark for correct colours reversed	

Question Number	Answer		Additional Guidance	Mark
9(d)	• moles of KMnO ₄ (aq) / MnO ₄ ⁻ (aq) used in titration	(1)	Example of calculation = $0.0200 \times 15.60 = 3.12 \times 10^{-4} / 0.000312$ (mol) 1000	(5)
	• moles H ₂ O ₂ in 25.0 cm ³ of the diluted sanitiser	(1)	= $(3.12 \times 10^{-4} / 2) \times 5 = 7.80 \times 10^{-4} / 0.000780 \text{ (mol)}$	
	• moles H_2O_2 in the 250 cm ³ of solution = moles in 10.0 cm ³ of hand sanitiser	(1)	= $7.80 \times 10^{-4} \times 10 = 7.80 \times 10^{-3} / 0.00780$ (mol) M3 could be carried out before M2	
	• mass H ₂ O ₂ in 10 cm ³ hand sanitiser	(1)	$=34 \times 7.80 \times 10^{-3} = 0.2652 \text{ (g)}$	
	% by mass of H ₂ O ₂ in hand sanitiser and this is within the (WHO) guidelines	(1)	$= \underbrace{0.2652}_{8.2} \times 100 = 3.23 \text{ (\%)}$ Ignore SF except 1 SF Allow TE throughout unless greater than 100%	
			Allow alternative method for M4 and M5 using concentrations of H ₂ O ₂ in mol dm ⁻³	

Question Number	Answer	Additional Guidance	Mark
9(e)	• pipette uncertainty (1)	Example of calculation $= \underline{0.04} \times 100 = (\pm) \ 0.16\%$ 25	(2)
	• burette uncertainty (1)	= 0.05 × 2 × 100 = (±) 0.643 / 0.64% 15.55 Ignore absence of labels 'pipette' and 'burette' If labels given but wrong way round allow 1 mark for 2 correct numerical values Allow any SF except 1 SF Accept correct answers without working Penalise 1 SF once only	

(Total for Question 9 = 11 marks)

TOTAL FOR PAPER = 90 MARKS

