

Mark Scheme (Results)

October 2022

Pearson Edexcel International Advanced Level in Chemistry (WCH15) Paper 01:Transition Metals and Organic Nitrogen 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.

Section A (Multiple Choice)

Question number	Answer	Mark
1	The only correct answer isA(decreases; gains electrons)	(1)
	<i>B</i> is incorrect because nitrogen gains electrons	
	<i>C</i> is incorrect because the oxidation number of nitrogen decreases	
	D is incorrect because the oxidation number of nitrogen decreases and it gains electrons	

Question	Answer	Mark
number		
2	The only correct answer is D(100000 Pa)	(1)
	<i>A</i> is incorrect because the standard pressure is 1 atm not 1 Pa	
	<i>B</i> is incorrect because the standard pressure is 100 kPa not 100 Pa	
	<i>C</i> is incorrect because the standard pressure is 100000 Pa not 1000 Pa	

Question number	Answer	Mark
3(a)	The only correct answer is D(platinum and platinum)	(1)
	<i>A</i> is incorrect because neither chromium metal nor titanium metal are involved in the cell	
	<i>B</i> is incorrect because chromium metalis not involved in the cell	
	<i>C</i> is incorrect because titanium metalis not involved in the cell	

Question number	Ans	swer	Mark
3(b)	The	e only correct answer is C(+0.19 V)	(1)
	A	is incorrect because the E°_{cell} value has been added to the electrode potential of the Cr_2O7^{2-} , Cr^{3+} electrode system rather than subtracted and the sign has been reversed.	
	B	is incorrect because the sign has been reversed.	
	D	is incorrect because the E°_{cell} value has been added to the electrode potential of the $Cr_2O_7^{2-}$, Cr^{3+} electrode system rather than subtracted	

Question number	Answer	Mark
4	The only correct answer is D(more positive; unchanged)	(1)
	<i>A</i> is incorrect because the E_{cell} values must be increasing and E_a values are not affected by concentrations.	
	B is incorrect because the E _{cell} values must be increasing	
	C is incorrect because E_a values are not affected by concentrations.	

Question number	Ans	swer	Mark
5	The	e only correct answer is C(the only stable zinc ion has the electronic configuration [Ar] 3d ¹⁰)	(1)
	A	is incorrect because the electronic configuration of the atom does not determine the classification	
	B	is incorrect because while the formation of more than one type of ion is a characteristic of transition metals it does not determine the classification	
	D	is incorrect because while catalytic properties are a characteristic of transition metals, they do not determine the classification	

Question	Answer	Mark
number		
6	The only correct answer is B	(1)
	$[Ar] \uparrow \downarrow \uparrow \uparrow \uparrow \uparrow \uparrow$	
	<i>A</i> is incorrect because both 4s electrons are removed when the Fe2+ ion is formed from the Fe atom	
	C is incorrect because both 4s electrons are removed when the Fe2+ ion is formed from the Fe atom	
	D is incorrect because electrons occupy a subshell with the maximum possible number of unpaired electrons	

Question number	Answer	Mark
7	The only correct answer is A(structure X only) $\begin{pmatrix} C & N \\ P & 3 \\ C & N \\ C & N \\ M \\ 3 \end{pmatrix}$	(1)
	<i>B</i> is incorrect because only the cis isomer is used	
	C is incorrect because $Pt(NH_3)_2Cl_2$ is square planar	
	<i>D</i> is incorrect because only the cis isomer is used	

Question number	Answer	Mark
8	The only correct answer is A(bonds reversibly to an iron(II) ion)	(1)
	B is incorrect because the reaction is reversible	
	<i>C</i> is incorrect because the iron(II) ion is not replaced	
	<i>D</i> is incorrect because the iron(II) ion is not replaced and the reaction is reversible	

Question number	Answer	Mark
9	The only correct answer is C(the oxidation of Ti ⁺ is catalysed by Ag ⁺ ions)	(1)
	<i>A</i> is incorrect because Ag^+ is unchanged at the end of the sequence while the Ce^{4+} ions are reduced	
	B is incorrect because Ag^+ is unchanged at the end of the sequence while the Ti^{2+} ions are formed from Ti^+ then further oxidised to Ti^{3+}	
	D is incorrect because Ag^{2+} ions are formed and removed in the reaction	

Question number	Answer	Mark
10	The only correct answer is C(12 (σ bonds) 3 (π bonds))	(1)
	<i>A</i> is incorrect because this does not include the C–H bonds	
	B is incorrect because this does not include the C–H bonds and counts all the C–C bonds as both σ bonds and π bonds	
	D is incorrect because this counts all the C–C bonds as both σ bonds and π bonds	

Question number	Answer	Mark
11	The only correct answer is D(concentrated sulfuric acid containing dissolved sulfur trioxide)	(1)
	A is incorrect because fuming sulfuric acid contains dissolved sulfur trioxide	
	B is incorrect because fuming sulfuric acid contains dissolved sulfur trioxide	
	<i>C</i> is incorrect because fuming sulfuric acid contains dissolved sulfur trioxide	

Question number	Answer	Mark
12(a)	12(a) The only correct answer is B(2-amino-3-methylpentanoic acid)	
	<i>A</i> is incorrect because the longest carbon chain has five atoms	
	<i>C</i> is incorrect because the longest carbon chain has five atoms and the carbon chain is numbered from the acid group	
	<i>D</i> is incorrect because the carbon chain is numbered from the acid group	

Question number	Answer	Mark
12(b)	The only correct answer is B $\left(\begin{array}{c} & \circ \\ & &$	(1)
	<i>A</i> is incorrect because this structure will occur at alkaline pH	
	<i>C</i> is incorrect because the amine group will protonate in preference to the carboxylic acid group at this pH.	
	<i>D</i> is incorrect because this zwitterion will occur at neutral pH	

Question	Answer	Mark
number		
13	13The only correct answer is B(CH ₃ CH ₂ CH ₂ CH ₂ CH ₂ NH ₃ Cl)	
	<i>A</i> is incorrect because substitution does not occur in these conditions	
	<i>C</i> is incorrect because an amide does not form in this way	
	<i>D</i> is incorrect because a carboxylic acid does not form in these conditions	

Question number	Answer	Mark
14		(1)
	The only correct answer is C(butanenitrile; lithium tetrahydridoaluminate(III))	
	<i>A</i> is incorrect because propanenitrile would give propylamine	
	B is incorrect because propanenitrile would give propylamineand tin with HCl does not reduce nitriles	
	D is incorrect because tin with HCl does not reduce nitriles	

Question number	Answer	Mark
15	The only correct answer is A $\begin{array}{c} H & d = 0 \\ d & d \\ d + 3 \end{array}$	(1)
	B is incorrect because the methyl group cannot form part of the polymer chain	
	<i>C</i> is incorrect because the peptide link cannot form from an amide group	
	<i>D</i> is incorrect because the methyl group cannot form part of the polymer chain and the peptide link cannot form from an amide group	

Question	Answer	Mark
number		
16	The only correct answer is A(four)	(1)
	<i>B</i> is incorrect because onlythe ring carbon atomshave been considered and symmetry has been ignored	
	<i>C</i> is incorrect because the ring symmetry has been ignored	
	D is incorrect because this is just the total number of carbon atoms	

Question number	Answer	Mark
17	The only correct answer is C(cocaine less soluble;cocaine higher pH)	
	<i>A</i> is incorrect because carboxylic acids are more soluble than esters	
	B is incorrect because carboxylic acids are more soluble and more acidic than esters	
	<i>D</i> is incorrect because carboxylic acids are more acidic than esters	

Question number	Answer	Mark
18	The only correct answer is B(soluble (at high temperature); insoluble (at low temperature)	(1)
	Ais incorrect because the hot filtration removes the insoluble impurities	
	<i>C</i> is incorrect because the hot filtration removes the insoluble impurities and the cold filtration separates the soluble impurities	
	<i>D</i> is incorrect because the hot filtration removes the insoluble impurities	

Total for Section A = 20 marks

Section B

Do not penalise case errors e.g. CO for Co and Pv = NRT

Question number	Answer	Additional guidance	Mark
19(a)	An explanation that makes reference to the following		3
	• +5 (1)	Allow 5+ / 5 / +V / V / V ⁵⁺	
	• vanadium has 3 (unpaired) electrons in the (3)d subshell and 2 electrons in the 4s (orbital / subshell) (1)	Accept electronic configuration of V is $[Ar]3d^3 4s^2 / [Ar]4s^2 3d^3 /$ 'electrons in boxes' Ignore errors in $[Ar]$ configuration Do not award incorrect electronic configurations of 3d and 4s	
	• the total number of 3d and 4s electrons gives the highest (stable) oxidation state (1)	Allow electronic configuration of V ⁵⁺ shown if electronic configuration of V given for M2	
		Allow vanadium has 5 electrons in its valence shell Allow vanadium has 5 outer electrons	
		Allow loss of 5 electrons from electronic configuration [Ar] 3d ⁵	
		Do not award 'orbit' for orbital / shell / subshell	
		No TE on other incorrect electronic configurations Ignore reference to stability of V ⁵⁺ Ignore just 'loses 5 electrons'	

Question number	Answer	Additional guidance	Mark
19(b)(i)	A justification that makes reference to the following		4
	• equation for reaction of thiosulfate ions and VO^{2+} (1)	$2VO^{2+} + 4H^+ + 2S_2O_3^{2-} \rightleftharpoons 2V^{3+} + S_4O_6^{2-} + 2H_2O$	
	• equation for reaction of thiosulfate ions and V^{3+} (1)	$2V^{3+} + 2S_2O_3^{2-} \rightleftharpoons 2V^{2+} + S_4O_6^{2-}$	
		Penalise incorrect balancing once only. Penalise uncancelled species (including e ⁻) once only Four correct half-equations scores (1)	
	• calculation of first <i>E</i> _{cell} value (using correct half – equations)	Do not award use of $S_2O_3^{2-} / S$ half-equation	
	and (positive E_{cell} value indicates) reduction to V ³⁺ feasible (1)	$E_{\text{cell}} = 0.34 - 0.09 = (+)0.25(\text{V})$	
	 calculation of second <i>E</i>_{cell} value and (negative <i>E</i>_{cell} value indicates) further reduction (to V²⁺) not feasible (1) 	$E_{\text{cell}} = -0.26 - 0.09 = -0.35(\text{V})$	
		Allow 1 mark for both E_{cell} values calculated without explanation	
		Ignore state symbols even if not correct No TE on incorrect ionic half-equations	

Question number	Answer	Additional guidance	Mark
19(b)(ii)	An explanation that makes reference to the following		2
	• Electrode potential for Ni $ Ni^{2+}$ is very close to $V^{2+} V^{3+}$ (1)	Accept $E_{cell} = -0.26 - (-0.25) = -0.01(V)$ Allow E_{cell} close to zero /(only) 0.01 V difference	
	• Some reduction of V^{3+} will occur and although E_{cell} is negative (1)	Allow Some reduction of V^{3+} will occur and $Ni + 2V^{3+} \Rightarrow Ni^{2+} + 2V^{2+}$ / equilibrium formed	
		Allow Some reduction of V ³⁺ will occur and if conditions are not standard /conditions are changed	
		Ignore just 'V ³⁺ reduced' Ignore just 'nickel is a strong(er) reducing agent'	

Question number	Answer	Additional guidance	Mark
19(c)(i)	An answer that makes reference to the following		1
	• pale pink	Allow just 'pink' Do not award purple / red Ignore reference to the colour before the end-point	

Question	Answer	Additional guidance	Mark
number			
19(c)(ii)	An answer that makes reference to the following		2
	 Because MnO₄⁻ ions have more positive standard electrode potential than VO²⁺, VO²⁺ should be oxidised using up MnO₄⁻ (1) 	Accept $E_{cell} = 1.51 - 1.00 = +0.51(V)$	
	• so reaction rate must be slow or	Allow conditions not standard	
	activation energy is high (1)		
		Allow MnO ₄ ⁻ can oxidise VO ²⁺ to VO ₃ ⁻ (reaction ratio 1:5) (1)	
		Then VO_3^- will oxidise Fe^{2+} to Fe^{3+} and	
		Overall this is equivalent to MnO_4^- oxidising Fe^{2+} to Fe^{3+} (1)	
		Ignore reference to catalysis	

uestion number	Answer		Additional guidance	Mark
19(c)(iii)	• calculation of moles of manganate(VII) in mean titre	(1)	Example of calculation mol MnO ₄ ⁻ = $\frac{22.50 \times 0.0195}{1000}$ = 4.3875 x 10 ⁻⁴ / 0.00043875	7
	• calculation of moles of Fe ²⁺ in 25 cm ³ after reaction	(1)	mol Fe ²⁺ in 25 cm ³ = 5 x mol MnO ₄ ⁻ = $2.1938 \times 10^{-3} / 0.0021938$	
	• calculation of moles of Fe ²⁺ in 25 cm ³ at start	(1)	mol Fe ²⁺ in 25 cm ³ = $\frac{25.00 \times 0.250}{1000}$ = 6.25 x 10 ⁻³ / 0.00625	
	• calculation of mol Fe ²⁺ that reacted in 25 cm ³	(1)	$6.25 \times 10^{-3} - 2.1938 \times 10^{-3} = 4.0562 \times 10^{-3}$	
	 mol Fe²⁺ = mol VO₃⁻ ions (stated or implied) and scales VO₃⁻ ions to 250 cm³ 	(1)	mol VO ₃ ⁻ⁱⁿ 250 cm ³ = 10 x 4.0562 x 10 ⁻³ = 4.0562 x 10 ⁻²	
	• calculation of mass of vanadium	(1)	mass of V = 50.9 x 4.0562 x 10^{-2} = 2.06463(g) [using unrounded values]	
	• calculation of percentage vanadium in ferrovanadium	(1)	% vanadium = $\frac{100 \times 2.06463}{4.87}$	
	If formula mass of VO ₃ ⁻ (98.9) is used instead of A_r of vanadium, mass = 98.9 × 4.0562 x 10 ⁻² = 4.0116 (g) percentage = 100 × 4.0116 ÷ 4.87 = 82.373(%) scores (6)		= 42.395% [using unrounded values] = 42.4% TE provided some attempt to use titration data to calculate mass and $\% < 100$	
	If M3 and M4 omitted % = $100 \ge 0.021938 \ge 50.9 \div 4.87$ = $22.939(\%) = (5)$		Ignore SF except 1 SF Correct answer with some working scores (7) Do not penalise correct rounding of intermediate values	

Question number	Answer	Additional guidance	Mark
19(d)	An answer that makes reference to the following	Examples of equations	2
	• balanced equation showing the reduction of vanadium(V) oxide by sulfur dioxide (1)	$V_2O_5 + SO_2 \rightarrow V_2O_4 + SO_3$	
	• balanced equation showing the oxidation of vanadium species formed by reduction back to vanadium(V) oxide (1)	$V_2O_4 + \frac{1}{2}O_2 \rightarrow V_2O_5$ Ignore uncancelled SO ₃	
		Allow for 2 marks any balanced equations showing formation of a lower oxidation state oxide by reaction with SO ₂ then a higher oxidation state by reaction with O ₂	
		Penalise error in formula of intermediate (e.g. $V_2O_4^-$) once only	
		Two balanced equations showing oxidation by O ₂ followed by reduction by SO ₂ scores (1)	
		Allow multiples Ignore state symbols even if incorrect	

Total for Question 19 = 21marks

nswer		Additional guidance	Mark
8	kages and fully sustained ontent and for how the answer is ning.	 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, a response with five indicative marking points that 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). If there were no linkages between the points, then the same 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 	
ne following table shows how the ructure and lines of reasoning	marks should be awarded for		
Answer shows a coherent logical tructure with linkages and fully ustained lines of reasoning emonstrated throughout Answer is partially structured with some linkages and lines of easoning Answer has no linkages between	Number of marks awarded for structure of answer and sustained lines of reasoning 2 1 0	 if there is incorrect chemistry, deduct mark(s) from the reasoning. If no reasoning mark(s) awarded do not deduct mark(s). Comment: Look for the indicative marking points first, then consider the mark for the structure of the answer and sustained line of reasoning 	
Answer vith son easonin Answer	is partially structured ne linkages and lines of	is partially structured 1 ne linkages and lines of g has no linkages between 0	is partially structured1Comment: Look for the indicative marking pointsne linkages and lines of1first, then consider the mark for the structure of the answer and sustained line of reasoninghas no linkages between0

Question number	Answe	er	Additional guidance	Mark
20 continued	Indica	tive content	Accept structures for names throughout Do not penalise unbalanced / incomplete / incorrect equations If name and formula given both must be correct	6
	IP1	(the π electrons of) both benzene and cyclohexene attract / react with electrophiles (similarity)	Allow both benzene and cyclohexene form carbocation (intermediates) Do not award nucleophilic reactions	
	IP2	delocalised (π) electron ring in benzene is (very)stable	Allow delocalised (π) electrons make benzene stable If neither IP1 or IP2 given, allow an IP for benzene has delocalised electrons but cyclohexene does not.	
	IP3	so benzene undergoes (electrophilic) substitution whereas cyclohexene undergoes (electrophilic) addition	Allow benzene forms bromobenzene and cyclohexene forms 1,2-dibromocyclohexane. Do not award nucleophilic reactions	
	IP4	both benzene and phenol undergo (electrophilic) substitution (similarity)	Allow benzene forms bromobenzene and phenol forms 2,4,6-tribromophenol	
	IP5	the lone pair of electrons on the oxygen in phenol interacts with / overlaps the delocalised (π) electron system of the benzene ring	Allow the lone pair of electrons on the oxygen in phenol increases the electron density of the (benzene) ring	
	IP6	phenol reacts (with electrophiles) much faster / under much milder conditions / does not require a catalyst / has a lower activation energy / E_a	Accept benzene requires a halogen carrier / catalyst Allow AlBr ₃ / AlCl ₃ / FeBr ₃ / Fe & Br ₂ Ignore just 'bromine reacts more easily with phenol than with benzene' / 'phenol is more susceptible to electrophilic attack'	

Question number	Answer	Additional guidance	Mark
21(a)(i)		Example of calculation	4
	• rearrangement of the ideal gas equation (IGE) (1)	$pV = nRT = \frac{m \times R \times T}{M} M = \frac{m \times R \times T}{p \times V}$	
	• conversion of temperature to Kelvin (1)	T = 185 + 273 (= 458 K)	
	• conversion of volume of vapour to m^3 (1)	$V = 67.1 \div 1 \ge 10^6 = 6.71 \ge 10^{-5} \text{ (m}^3)$	
	• solve IGE for M (1)	$M = \frac{0.235 \text{ x } 8.31 \text{ x } 458}{105000 \text{ x } 6.71 \text{ x } 10^{-5}} = 126.95 / 127 \text{ (g mol^{-1})}$	
	Alternative method	TE on incorrect conversions for M4	
	• rearrangement of the ideal gas equation (IGE) (1)	$pV = nRT n = \frac{pV}{RT}$	
	• conversion of temperature to Kelvin (1)	T = 185 + 273 (= 458 K)	
	• conversion of volume of vapour to m^3 (1)	$V = 67.1 \div 1 \ge 10^6 = 6.71 \ge 10^{-5}$	
	• solve IGE for M (1)	$n = \frac{105000 \ge 6.71 \ge 10^{-5}}{8.31 \ge 458} = 1.8512 \ge 10^{-3}$	
		$M = \frac{0.235}{1.8512 \text{ x } 10^{-3}} = 126.947 / 126.9 / 127 \text{ (g mol}^{-1}\text{)}$	
		Use of 1.85 gives 127.03 Allow units of g or no units Allow conversion of pressure to kPa & volume to dm ³ Correct answer with no working scores zero	

Question number	Answer	Additional guidance	Mark
21(a)(ii)	An answer that makes reference to the following points:		2
	• calculation of the molar mass of the ring structure of coniine (1)	the molar mass of ring = 5 x $12 + 14 + 10 x 1$ = 84 g mol ⁻¹	
	• calculation of the molar mass of R and	the molar mass of $R = 127 - 84 = 43 \text{ (g mol}^{-1}\text{)}$	
	deduction of the molecular formula of R (1)	R is C ₃ H ₇ (only possible alkyl group formula for molar mass = $43 \text{ (g mol}^{-1}\text{)}$)	
		Allow structural formulae e.g. –CH ₂ CH ₂ CH ₃ Allow this mark if the formula is given in (b)(i)	
		TE on incorrect molar mass of ring provided R	
		contains C and H only	

Question number	Answer	Additional guidance	Mark
21(b)(i)	An answer that makes reference to the following points:		3
	• identification of the two possible R structures (1)	-CH ₂ CH ₂ CH ₃ and –CH(CH ₃) ₂ Do not award if additional structures are given	
	• -CH ₂ CH ₂ CH ₃ has three proton environments (as all three are different) (1)	Accept the peak areas show that the environments have 2, 2 and 3 protons as in -CH ₂ CH ₂ CH ₃	
	• -CH(CH ₃) ₂ has two proton environments (because the methyl groups are equivalent) (1)	Accept –CH(CH ₃) ₂ would have peak areas 1 and 6 If no other mark is scored identifying R as - CH ₂ CH ₂ CH ₃ with any valid reason based on the NMR data, scores (1)	
		No TE if R not given as an alkyl group e.g. CH ₃ CO	

Question number	Answer	Additional guidance	Mark
21(b)(ii)	 An answer that makes reference to the following points: proton environment 1 (has 3 protons and) will be a triplet (1) proton environment 3 (has 2 protons and) will be a quartet (1) 	May be shown on a diagram	3
	 Either proton environment 1(is a triplet because it) is adjacent to a CH₂ in the chain or as proton environment 3(is a quartet because it) is adjacent to a CH in the ring and to a CH₂ in the chain (1) 	Allow there are 2 adjacent protons Allow there are 3 adjacent protons M3 dependent on correct splitting pattern(s)	

Question number	Answer	Additional guidance	Mark
21(c)	An explanation that makes reference to the following points: EITHER • identification of the asymmetric carbon atom in coniine (1)	asymmetric carbon H	2
	• the molecule is non-superimposable on its mirror image and hence coniine shows optical isomerism (1)	Ignore incorrect alkyl groups Allow carbon atom is attached to four different groups and hence coniine shows optical isomerism OR Allow carbon atom is chiral and coniine shows optical isomerism Do not award four different 'molecules'	
	OR	Diagram showing the two enantiomers and a mirror plane scores M1 and M2 (dots and wedge not required)	
	 There is not free rotation around the N-C(-R) bond (because of the ring) (1) Coniine shows cis-trans / E-Z / geometrical isomerism and because the lone pair on N leads to the H having two possible orientations (1) 	This diagram including the mirror plane but without H	
		shown (and chiral carbon unlabelled) scores (1) Total for Ouestion 21 =	14

Total for Question 21 = 14 marks

Question number	Answer	Additional guidance	Mark
22(a)		Example of calculation	4
	• equating of % water with mass in 1 mol of viridian (1)	19.15% of 1 mol of viridian weighs 36 g	
	• calculation of molar mass of viridian (1)	1 mol of viridian weighs 100 x $36 \div 19.15$ (g) and molar mass = (100 x $36 \div 19.15$) = 188 (g mol ⁻¹)	
	• calculation of molar mass of metal oxide (1)	molar mass = $188 - 36 = 152$ (g mol ⁻¹)	
	 calculation of molar mass of metal and deduction of its identity (1) 	molar mass of metal = $\frac{1}{2}(152 - 3 \times 16) = 52 \text{ (g mol}^{-1})$ and element is chromium / Cr	
		TE at each stage Correct answer with some working scores (4) Correct answer with no working (0)	

Question number	Answer	Additional guidance	Mark
22(b)(i)	 An answer that makes reference to ligand exchange / ligand replacement / ligand substitution 		1

Question number	Answer	Additional guidance	Mark
22(b)(ii)	An answer that makes reference to	Allow omission of square brackets Ignore extra brackets e.g. [Co(Cl)4] ^{2–}	2
	• Full formula of one complex (1	$[Co(H_2O)_6]^{2+}$ or $[CoCl_4]^{2-}$	
	• completion of full balanced ionic equation (1	$[Co(H_2O)_6]^{2+} + 4Cl^- \rightarrow [CoCl_4]^{2-} + 6H_2O$ Ignore state symbols even if incorrect	

Question number	Answer	Additional guidance	Mark
number 22(b)(iii)	 An explanation that makes reference to the following points (the aqua complex is octahedral and) chloro complex of cobalt / tetrachlorocobaltate(II) / CoCl4²⁻ is tetrahedral (1) because chloride ions / Cl⁻ are too large for six ions to coordinate around the cobalt(II) ion / Co²⁺ (1) 	Allow octahedral (complex) changes to tetrahedral (complex) Allow the complex formed is tetrahedral Do not award Cl / chlorine atoms / molecules If no other mark is scored, chloride ions / Cl ⁻ are	2
		larger than H ₂ O scores (1) [Both species needed] If no other mark is scored, coordination number changes from 6 to 4 scores (1) Ignore just 'chloride ions / Cl ⁻ are larger'	

Total for Question 22 = 9 marks Total for Section B = 50 marks

Section C

Question number	A	nswer						Additional guidance	Mark
23(a)								Example of calculation	4
	•	working ou	ıt molec	ular formul	a		(1)	$C_{13}H_{18}O_7$ (Accept numbers of C, H and O indicated in calculation)	
	•	calculation	of mola	r mass			(1)	286	
	•	calculation	of perce	entage by n	nass of one	e element	(1)	% by mass of carbon = $100 \ge 13 \ge 12 \div 286$ = 54.545	
	•	calculation	of perce	entage by n	nass of oth	er elements	(1)	% by mass of hydrogen = $100 \times 18 \times 1 \div 286$ = 6.294 and	
		Some comr	non inco	orrect answ	ers (score	(3))		% by mass of oxygen = $100 \times 7 \times 16 \div 286$ or	
		Formula	Mr	%C	%Н	%O		% by mass of oxygen = $100 - (54.545 + 6.2937)$	
		C13H10O7	278	56.12	3.60	40.29		= 39.161	
		C13H11O7	279	55.91	3.94	40.14	1		
		C13H13O7	281	55.52	4.63	39.86		TE at each stage	
		C13H17O7	285	54.74	5.96	39.30			
		C13H20O7	288	54.17	6.94	38.89		Ignore SF except 1 SF	
		C13H23O7	291	53.61	7.90	38.49			
		C13H24O7	292	53.42	8.22	38.36			

Question number	Answer		Additional guidance	Mark
23(b)(i)	An answer that makes reference to the following		Standalone marks	2
	• a suitable oxidising agent	(1)	sodium dichromate(VI) / Na2Cr2O7 / potassium dichromate(VI) / K2Cr2O7	
	• sulfuric acid / H ₂ SO ₄		Allow dichromate(VI) / Cr ₂ O ₇ ^{2–} Ignore omission of oxidation state but, if given, oxidation state must be correct	
	and	(1)	Allow acid / acidified / H ⁺ / H ₃ O ⁺ Ignore concentration of the acid Do not award other acids Do not award other solvents / reagents If name and formula are given both must be correct.	

Question	Answer	Additional guidance	Mark
number			
23(b)(ii)	An answer that makes reference to the following		2
	• salicylaldehyde needs to be distilled / separated (from the reaction mixture to prevent its further oxidation) (1)		
	EITHER		
	• (because) at high temperatures salicylaldehyde will be oxidised (to salicylic acid / carboxylic acid) (1)	Allow salicylaldehyde is more easily oxidised than salicyl alcohol Allow oxidation of salicylaldehyde is fast	
	 OR (because) its high boiling temperature means that salicylaldehyde cannot be easily be separated as the other components of the mixture have lower / similar 	Allow salicylaldehyde is hard to separate from the mixture	
	boiling temperatures (1)	If no other mark is scored, salicylaldehyde / salicyl alcohol will be oxidised to salicylic acid / carboxylic acid scores (1)	

Question number	Answer	Additional guidance	Mark
23(c)(i)	equation showing formation of electrophile	Example of equation $ \begin{array}{ccccccccccccccccccccccccccccccccccc$	1

Question number	Answer	Additional guidance	Mar k
23(c)(ii)	 A mechanism showing the following curly arrow from on or within the circle to the positively charged carbon (1) 	Example of mechanism 0 H H H H H H H H	3
	 intermediate structure including charge with horseshoe covering at least 3 carbon atoms and facing the tetrahedral carbon and with some part of the positive charge within the horseshoe (1) 		
	 curly arrow from C—H bond to anywhere in the benzene ring reforming delocalised structure and giving benzaldehyde (1) Ignore use of AlCl4⁻ in mechanism 	Allow TE on any electrophile from (c)(i) Correct Kekule mechanism scores (3) Ignore connectivity of undisplayed CHO in M2 and M3	

Question number	Answer		Additional guidance	Mark
23(d)	An answer that makes reference to the following points:		Penalise use of Cl or I in Grignard once only	4
	• reagent and conditions for the formation of the Grignard reagent	(1)	Reaction of bromoethane with magnesium (powder) in (dry) ether (under reflux) Ignore heat	
	• identification of Grignard reagent	(1)	CH ₃ CH ₂ MgBr / ethyl magnesium bromide	
	• addition product of reaction between salicylaldehyde and Grignard reagent	(1)	ОН	
	• identification of a suitable dehydrating reagent	(1)	Ignore reference to hydrolysis of the Grignard intermediate (conc) phosphoric((V)) acid / H ₃ PO ₄ / (conc) sulfuric acid / H ₂ SO ₄ Do not award dilute acids Allow conversion of OH to halogen using a suitable reagent (e.g. PCl ₅) and Dehydrohalogenation using alcoholic KOH / NaOH	

Question number	Answer	Additional guidance	Mark
23(e)(i)	A diagram showing	Example of diagram	2
	• nickel(II) ion coordinated to the two oxygen and two nitrogen atoms of the salen ligand (1)		
	• at least one dative covalent bond from an oxygen or a nitrogen to the nickel (II) ion in any structure involving one salen ligand (1)	Accept charge shown over whole structure Ignore omission of C=N double bond Allow 'dative /coordinate (covalent) bond' stated provided complex structure is involves a salen ligand and at least one O—Ni ²⁺ or N—Ni ²⁺ bond is shown Penalise incorrect connectivity of OH with Ni ²⁺ only in M2. If two OH groups are shown, both must be correctly connected to the Ni ²⁺	

23(e)(ii)An answer that makes reference to the following points:Standalone marks2• converting the aqua complex to the salen ligand complex results in an increase in the number of particles (in solution)Allow 'molecules' for 'particles'2• (thus) the entropy of the system increases / ΔS_{system} is positive(1)Accept ΔS_{system} increases Ignore just 'entropy increases' If no other mark is scored balanced equation (1)2	Question number	Answer	Additional guidance	Mark
an increase in the number of particles (in solution) (1) • (thus) the entropy of the system increases / ΔS_{system} is positive (1) Accept ΔS_{system} increases Ignore just 'entropy increases'	23(e)(ii)	An answer that makes reference to the following points:	Standalone marks	2
Ignore just 'entropy increases'			Allow 'molecules' for 'particles'	
		• (thus) the entropy of the system increases / ΔS_{system} is positive (1)	Ignore just 'entropy increases'	

TOTAL FOR SECTION C = 20 MARKS

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