## Pearson Edexcel

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 meaning 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 <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1}$ | The only correct answer isA(decreases; gains electrons) |  |
|  | $\boldsymbol{B} \quad$ is incorrect because nitrogen gains electrons |  |
| $\boldsymbol{C} \quad$ is incorrect because the oxidation number of nitrogen decreases |  |  |
| $\boldsymbol{D} \quad$ is incorrect because the oxidation number of nitrogen decreasesand it gains electrons |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{2}$ | The only correct answer is $\mathbf{D}(100000 \mathrm{~Pa}$ ) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because the standard pressure is 1 atm not 1 Pa |  |
|  | $\boldsymbol{B} \quad$ is incorrect because the standard pressure is 100 kPa not 100 Pa |  |
| $\boldsymbol{C} \quad$ is incorrect because the standard pressure is 100000 Pa not 1000 Pa |  |  |

$\left.\begin{array}{|l|l|c|}\hline \begin{array}{l}\text { Question } \\ \text { number }\end{array} & \text { Answer } & \text { Mark } \\ \hline \text { 3(a) } & \text { The only correct answer is D(platinum and platinum) } & \text { (1) } \\ & \boldsymbol{A} \quad \text { is incorrect because neither chromium metal nor titanium metal are involved in the cell } \\ \boldsymbol{B} \quad \text { is incorrect because chromium metalis not involved in the cell } \\ \boldsymbol{C} \quad \text { is incorrect because titanium metalis not involved in the cell }\end{array}\right]$

| Question number | Answer | Mark |
| :---: | :---: | :---: |
| 3(b) | The only correct answer is $\mathbf{C}(+0.19 \mathrm{~V})$ <br> A is incorrect because the $E^{\ominus}$ cell value has been added to the electrode potential of the $\mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}, \mathrm{Cr}^{3+}$ electrode system rather than subtractedand the sign has been reversed. <br> B is incorrect because the sign has been reversed. <br> D is incorrect because the $E^{\ominus}$ cell value has been added to the electrode potential of the $\mathrm{Cr}_{2} \mathrm{O}_{7}{ }^{2-}, \mathrm{Cr}^{3+}$ electrode system rather than subtracted | (1) |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{4}$ | The only correct answer is D(more positive; unchanged) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because the Ecellvalues must be increasing and Ea values are not affected by concentrations. |  |
|  | $\boldsymbol{B} \quad$ is incorrect because the Ecellvalues must be increasing |  |
| $\boldsymbol{C} \quad$ is incorrect because Ea values are not affected by concentrations. |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :--- |
| $\mathbf{5}$ | The only correct answer is C(the only stable zinc ion has the electronic configuration $[\mathrm{Ar}] 3 \mathrm{~d}^{10}$ ) <br> A is incorrect because the electronic configuration of the atom does not determine the classification <br> B is incorrect because while the formation of more than one type of ion is a characteristic of transition metals it does <br> not determine the classification <br> is incorrect because while catalytic properties are a characteristic of transition metals, they do not determine the <br> classification | $\mathbf{( 1 )}$ |



| Question number | Answer | Mark |
| :---: | :---: | :---: |
| 7 | The only correct answer is A(structure X only) <br> B is incorrect because only the cis isomer is used <br> C is incorrect because $\operatorname{Pt}\left(\mathrm{NH}_{3}\right)_{2} \mathrm{Cl}_{2}$ is square planar <br> D is incorrect because only the cis isomer is used | (1) |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{8}$ | The only correct answer is A(bonds reversibly to an iron(II) ion) | (1) |
|  | $\boldsymbol{B} \quad$ is incorrect because the reaction is reversible |  |
| $\boldsymbol{C} \quad$ is incorrect because the iron(II) ion is not replaced |  |  |
| $\boldsymbol{D} \quad$ is incorrect because the iron(II) ion is not replaced and the reaction is reversible |  |  |


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


| Question <br> number | Answer | Mark |
| :--- | :--- | :--- |
| $\mathbf{1 0}$ | The only correct answer is C(12 ( $\sigma$ bonds) $3(\pi$ bonds)) <br> $\boldsymbol{A} \quad$ is incorrect because this does not include the $C-H$ bonds <br> $\boldsymbol{B} \quad$ is incorrect because this does not include the $C-H$ bonds and counts all the $C-C$ bonds asboth $\sigma$ bonds and <br> $\pi b o n d s$ <br> $\boldsymbol{D}$ is incorrect because this counts all the $C-C$ bonds as both $\sigma$ bonds and $\pi$ bonds | $\mathbf{( 1 )}$ |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 1}$ | The only correct answer is D(concentrated sulfuric acid containing dissolved sulfur trioxide) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because fuming sulfuric acid contains dissolved sulfur trioxide |  |
| $\boldsymbol{B} \quad$ is incorrect because fuming sulfuric acid contains dissolved sulfur trioxide |  |  |
| $\boldsymbol{C} \quad$ is incorrect because fuming sulfuric acid contains dissolved sulfur trioxide |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :--- |
| $\mathbf{1 2 ( a )}$ | The only correct answer is B(2-amino-3-methylpentanoic acid) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because the longest carbon chain has five atoms <br> $\boldsymbol{C} \quad$ is incorrect because the longest carbon chain has five atoms and the carbon chain is numbered from the acid group <br> D incorrect because the carbon chain is numbered from the acid group |  |


| Question number | Answer | Mark |
| :---: | :---: | :---: |
| 12(b) | The only correct answer is $B$ <br> A is incorrect because this structure will occur at alkaline pH <br> C is incorrect because the amine group will protonate in preference to the carboxylic acid group at this pH . <br> D is incorrect because this zwitterion will occur at neutral pH | (1) |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 3}$ | The only correct answer is $\mathbf{B}\left(\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{NH}_{3} \mathrm{Cl}\right)$ | (1) |
|  | A is incorrect because substitution does not occur in these conditions |  |
| C is incorrect because an amide does not form in this way |  |  |
| $\boldsymbol{D} \quad$ is incorrect because a carboxylic acid does not form in these conditions |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 4}$ | The only correct answer is C(butanenitrile; lithium tetrahydridoaluminate(III)) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because propanenitrile would give propylamine <br> $\boldsymbol{B} \quad$ is incorrect because propanenitrile would give propylamineand tin with HCl does not reduce nitriles <br> D is incorrect because tin with HCl does not reduce nitriles |  |


| Question <br> number | Answer | Mark |  |
| :--- | :--- | :--- | :--- |
| $\mathbf{1 5}$ |  |  | (1) |
|  | The only correct answer is A |  |  |
| B $\quad$is incorrect because the methyl group cannot form part of the polymer chain <br> is incorrect because the peptide link cannot form from an amide group <br> is incorrect because the methyl group cannot form part of the polymer chain and the peptide link cannot form from <br> an amide group |  |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 6}$ | The only correct answer is A(four) | (1) |
|  | $\boldsymbol{B} \quad$ is incorrect because onlythe ring carbon atomshave been considered and symmetry has been ignored |  |
| $\boldsymbol{C} \quad$ is incorrect because the ring symmetry has been ignored |  |  |
| $\boldsymbol{D} \quad$ is incorrect because this is just the total number of carbon atoms |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 7}$ | The only correct answer is C(cocaine less soluble;cocaine higher pH) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because carboxylic acids are more soluble than esters |  |
| $\boldsymbol{B} \quad$ is incorrect because carboxylic acids are more soluble and more acidic than esters |  |  |
| D is incorrect because carboxylic acids are more acidic than esters |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 8}$ | The only correct answer is B(soluble (at high temperature); insoluble (at low temperature) <br> Ais incorrect because the hot filtration removes the insoluble impurities <br> $\boldsymbol{C}$ is incorrect because the hot filtration removes the insoluble impurities and the cold filtration separates the soluble <br> impurities <br> $\boldsymbol{D} \quad$ is incorrect because the hot filtration removes the insoluble impurities | $\mathbf{( 1 )}$ |

## Section B

Do not penalise case errors e.g. CO for Co and $\mathrm{Pv}=$ NRT

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


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 19(b)(i) | A justification that makes reference to the following <br> - equation for reaction of thiosulfate ions and $\mathrm{VO}^{2+}$ <br> - equation for reaction of thiosulfate ions and $\mathrm{V}^{3+}$ <br> - calculation of first $E_{\text {cell }}$ value (using correct half equations) <br> and <br> (positive $E_{\text {cell }}$ value indicates) reduction to $\mathrm{V}^{3+}$ feasible <br> - calculation of second $E_{\text {cell }}$ value <br> and <br> (negative $E_{\text {cell }}$ value indicates) further reduction (to $\mathrm{V}^{2+}$ ) <br> not feasible | $\begin{equation*} 2 \mathrm{VO}^{2+}+4 \mathrm{H}^{+}+2 \mathrm{~S}_{2} \mathrm{O}_{3}{ }^{2-} \rightleftharpoons 2 \mathrm{~V}^{3+}+\mathrm{S}_{4} \mathrm{O}_{6}{ }^{2-}+2 \mathrm{H}_{2} \mathrm{O} \tag{1} \end{equation*}$ $\begin{equation*} 2 \mathrm{~V}^{3+}+2 \mathrm{~S}_{2} \mathrm{O}_{3}{ }^{2-} \rightleftharpoons 2 \mathrm{~V}^{2+}+\mathrm{S}_{4} \mathrm{O}_{6}{ }^{2-} \tag{1} \end{equation*}$ <br> Penalise incorrect balancing once only. <br> Penalise uncancelled species (including $\mathrm{e}^{-}$) once only Four correct half-equations scores (1) <br> Do not award use of $\mathrm{S}_{2} \mathrm{O}_{3}{ }^{2-} / \mathrm{S}$ half-equation $\begin{equation*} E_{\text {cell }}=0.34-0.09=(+) 0.25(\mathrm{~V}) \tag{1} \end{equation*}$ $E_{\text {cell }}=-0.26-0.09=-0.35(\mathrm{~V})$ <br> Allow 1 mark for both $E_{\text {cell }}$ values calculated without explanation <br> Ignore state symbols even if not correct <br> No TE on incorrect ionic half-equations | 4 |


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


| Question <br> number | Answer | Additional guidance | Mark |
| :--- | :--- | :--- | :---: |
| 19(c)(i) | An answer that makes reference to the following | Allow just 'pink' <br> Do not award purple / red <br> Ignore reference to the colour before the end-point | $\mathbf{1}$ |


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


| uestion number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 19(c)(iii) |  | Example of calculation | 7 |
|  | - calculation of moles of manganate(VII) in mean titre | $\begin{aligned} & \mathrm{mol} \mathrm{MnO}_{4}^{-}=\frac{22.50 \times 0.0195}{1000} \\ & =4.3875 \times 10^{-4} / 0.00043875 \end{aligned}$ |  |
|  | - calculation of moles of $\mathrm{Fe}^{2+}$ in $25 \mathrm{~cm}^{3}$ after reaction | $\begin{aligned} & \mathrm{mol} \mathrm{Fe}^{2+} \text { in } 25 \mathrm{~cm}^{3}=5 \times \mathrm{mol} \mathrm{MnO} \\ & 4 \\ &=2.1938 \times 10^{-3} / 0.0021938 \end{aligned}$ |  |
|  | - calculation of moles of $\mathrm{Fe}^{2+}$ in $25 \mathrm{~cm}^{3}$ at start | $\begin{align*} \mathrm{mol} \mathrm{Fe} \tag{1} \end{align*}{ }^{2+} \text { in } 25 \mathrm{~cm}^{3}=\frac{25.00 \times 0.250}{1000}, ~=6.25 \times 10^{-3} / 0.00625$ |  |
|  | - calculation of mol Fe ${ }^{2+}$ that reacted in $25 \mathrm{~cm}^{3}$ | $6.25 \times 10^{-3}-2.1938 \times 10^{-3}=4.0562 \times 10^{-3}$ |  |
|  | - $\mathrm{mol} \mathrm{Fe}^{2+}=\mathrm{mol} \mathrm{VO}_{3}{ }^{-}$ions (stated or implied) <br> and <br> scales $\mathrm{VO}_{3}{ }^{-}$ions to $250 \mathrm{~cm}^{3}$ | $\begin{align*} \mathrm{mol} \mathrm{VO}_{3}-\mathrm{in} 250 \mathrm{~cm}^{3} & =10 \times 4.0562 \times 10^{-3} \\ & =4.0562 \times 10^{-2} \tag{1} \end{align*}$ |  |
|  | - calculation of mass of vanadium | $\text { mass of } \begin{aligned} \mathrm{V} & =50.9 \times 4.0562 \times 10^{-2} \\ & =2.06463(\mathrm{~g}) \text { [using unrounded values] } \end{aligned}$ |  |
|  | - calculation of percentage vanadium in ferrovanadium | $\% \text { vanadium }=\frac{100 \times 2.06463}{487}$ |  |
|  | If formula mass of $\mathrm{VO}_{3}^{-}$(98.9) is used instead of $A_{\mathrm{r}}$ of vanadium, mass $=98.9 \times 4.0562 \times 10^{-2}=4.0116(\mathrm{~g})$ percentage $=100 \times 4.0116 \div 4.87=82.373(\%)$ scores ( 6 ) | $=42.395 \%$ [using unrounded values] $=42.4 \%$ TE provided some attempt to use titration data to calculate mass and $\%<100$ |  |
|  | $\text { If M3 and M4 omitted } \begin{aligned} \% & =100 \times 0.021938 \times 50.9 \div 4.87 \\ & =22.939(\%) \text { scores (5) } \end{aligned}$ | Ignore SF except 1 SF <br> Correct answer with some working scores (7) <br> Do not penalise correct rounding of intermediate |  |
|  |  | values |  |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 19(d) | An answer that makes reference to the following <br> - balanced equation showing the reduction of vanadium(V) oxide by sulfur dioxide (1) <br> - balanced equation showing the oxidation of vanadium species formed by reduction back to vanadium(V) oxide (1) | Examples of equations $\mathrm{V}_{2} \mathrm{O}_{5}+\mathrm{SO}_{2} \rightarrow \mathrm{~V}_{2} \mathrm{O}_{4}+\mathrm{SO}_{3}$ <br> $\mathrm{V}_{2} \mathrm{O}_{4}+1 / 2 \mathrm{O}_{2} \rightarrow \mathrm{~V}_{2} \mathrm{O}_{5}$ <br> Ignore uncancelled $\mathrm{SO}_{3}$ <br> Allow for 2 marks any balanced equations showing formation of a lower oxidation state oxide by reaction with $\mathrm{SO}_{2}$ then a higher oxidation state by reaction with $\mathrm{O}_{2}$ <br> Penalise error in formula of intermediate (e.g. $\mathrm{V}_{2} \mathrm{O}_{4}{ }^{-}$) once only <br> Two balanced equations showing oxidation by $\mathrm{O}_{2}$ followed by reduction by $\mathrm{SO}_{2}$ scores (1) <br> Allow multiples <br> Ignore state symbols even if incorrect | 2 |


| Question number | Answer |  |
| :---: | :---: | :---: |
| *20 | This question assesses the student's ability to show a coherent and logically structured answer with linkages and fully sustained reasoning. <br> Marks are awarded for indicative content and for how the answer is structured and shows lines of reasoning. <br> The following table shows how the marks should be awarded for indicative content. |  |
|  | Number of indicative marking points seen in answer | Number of marks awarded for indicative marking points |
|  | 6 | 4 |
|  | 5-4 | 3 |
|  | 3-2 | 2 |
|  | 1 | 1 |
|  | 0 | 0 |

The following table shows how the marks should be awarded for structure and lines of reasoning

|  | Number of marks awarded <br> for structure of answer and <br> sustained lines of reasoning |
| :--- | :---: |
| Answer shows a coherent logical <br> structure with linkages and fully <br> sustained lines of reasoning <br> demonstrated throughout | 2 |
| Answer is partially structured <br> with some linkages and lines of <br> reasoning | 1 |
| Answer has no linkages between <br> points and is unstructured | 0 |



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


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


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


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


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


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 22(a) | - equating of \% water with mass in 1 mol of viridian <br> - calculation of molar mass of viridian <br> - calculation of molar mass of metal oxide <br> - calculation of molar mass of metal <br> and deduction of its identity | Example of calculation <br> $19.15 \%$ of 1 mol of viridian weighs 36 g <br> 1 mol of viridian weighs $100 \times 36 \div 19.15$ (g) <br> and <br> molar mass $=(100 \times 36 \div 19.15)=188\left(\mathrm{~g} \mathrm{~mol}^{-1}\right)$ <br> molar mass $=188-36=152\left(\mathrm{~g} \mathrm{~mol}^{-1}\right)$ <br> molar mass of metal $=1 / 2(152-3 \times 16)=52\left(\mathrm{~g} \mathrm{~mol}^{-1}\right)$ <br> and <br> element is chromium / Cr <br> TE at each stage <br> Correct answer with some working scores (4) <br> Correct answer with no working (0) | 4 |
| Question number | Answer | Additional guidance | Mark |
| 22(b)(i) | An answer that makes reference to <br> - ligand exchange / ligand replacement / ligand substitution |  | 1 |


| Question number | Answer |  | Additional guidance | Mark |
| :---: | :---: | :---: | :---: | :---: |
| 22(b)(ii) | An answer that makes reference to <br> - Full formula of one complex <br> - completion of full balanced ionic equation | (1) <br> (1) | Allow omission of square brackets Ignore extra brackets e.g. $[\mathrm{Co}(\mathrm{Cl}) 4]^{2-}$ <br> $\left[\mathrm{Co}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}$ or $\left[\mathrm{CoCl}_{4}\right]^{2-}$ $\left[\mathrm{Co}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}+4 \mathrm{Cl}^{-} \rightarrow\left[\mathrm{CoCl}_{4}\right]^{2-}+6 \mathrm{H}_{2} \mathrm{O}$ <br> Ignore state symbols even if incorrect | 2 |


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

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

## Section C



| Question <br> number | Answer | Additional guidance | Mark |  |
| :--- | :--- | :--- | :--- | :--- |
| 23(b)(i) | An answer that makes reference to the following | (1) | Standalone marks <br> sodium dichromate(VI) $/ \mathrm{Na}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7} /$ <br> potassium dichromate(VI) $/ \mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}$ | $\mathbf{2}$ |
|  | - a suitable oxidising agent | Allow dichromate(VI) $/ \mathrm{Cr}_{2} \mathrm{O}_{7}{ }^{2-}$ <br> Ignore omission of oxidation state but, if given, <br> oxidation state must be correct |  |  |
|  | sulfuric acid $/ \mathrm{H}_{2} \mathrm{SO}_{4}$ <br> and <br> (heat under) reflux | Allow <br> acid $/$ acidified $/ \mathrm{H}^{+} / \mathrm{H}_{3} \mathrm{O}^{+}$ <br> Ignore concentration of the acid <br> Do not award other acids <br> Do not award other solvents $/$reagents <br> If name and formula are given both must be correct. |  |  |


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


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 23(c)(i) | - equation showing formation of electrophile | Allow <br> Structural formulae <br> Allow <br> $\mathrm{CO}+\mathrm{HCl}+\mathrm{AlCl}_{3} \rightarrow \mathrm{HCO}^{+}+\mathrm{AlCl}_{4}^{-}$ <br> Allow $\begin{array}{lll} \mathrm{CO}+\mathrm{HCl} \xrightarrow{\mathrm{AlCl}_{3}} \mathrm{HCO}^{+}+\mathrm{Cl}^{-} \\ \mathrm{Ignore} \\ \mathrm{CO}+\mathrm{HCl} \longrightarrow & \mathrm{HCO}^{+}+\mathrm{Cl}^{-} \end{array}$ | 1 |


| Question number | Answer | Additional guidance | Mar $\mathbf{k}$ |
| :---: | :---: | :---: | :---: |
| 23(c)(ii) | A mechanism showing the following <br> - curly arrow from on or within the circle to the positively charged carbon | Example of mechanism <br> 0 <br> Allow curly arrow from anywhere in the hexagon Allow arrow to positive charge provided the charge is on the C atom TE on any correct electrophile formed in (c)(i) <br> Allow TE on any electrophile from (c)(i) <br> Correct Kekule mechanism scores (3) <br> Ignore connectivity of undisplayed CHO in M2 and M3 | 3 |
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|  |  |  |  |
|  |  |  |  |
|  | - intermediate structure including charge with horseshoe covering at least 3 carbon atoms |  |  |
|  | and <br> facing the tetrahedral carbon <br> and <br> with some part of the positive charge within the horseshoe |  |  |
|  | - curly arrow from $\mathrm{C}-\mathrm{H}$ bond to anywhere in the benzene ring reforming delocalised structure and giving benzaldehyde |  |  |
|  | Ignore use of $\mathrm{AlCl}_{4}{ }^{-}$in mechanism |  |  |
|  |  |  |  |


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


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


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

