# Mark Scheme (Provisional) 

Summer 2021

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.


## Section A (Multiple Choice)

| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1}$ | The only correct answer is $\mathbf{D}\left(\mathrm{VO}_{2}{ }^{+}\right.$and $\left.\mathrm{VO}^{2+}\right)$ |  |
| $\boldsymbol{A}$ is not correct because both $C r$ have the oxidation number +6 |  |  |
| $\boldsymbol{B}$ is not correct because both $C r$ have the oxidation number +6 |  |  |
| $\boldsymbol{C}$ is not correct because both $V$ have the oxidation number +5 |  |  |$\quad \mathbf{1}$.


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{2 ( a )}$ | The only correct answer is B $(-1.63 \mathrm{~V})$ |  |
| A $\quad$ is not correct because the electrode potential for the $\mathrm{Fe}^{3+} / \mathrm{Fe}^{2+}$ electrode system has been added to the $E^{o}$ cell <br> value instead of being subtracted <br> $\boldsymbol{C} \quad$ is not correct because the value should have a negative sign <br> $\boldsymbol{D} \quad$ is not correct because the value should have a negative sign and the electrode potential for the $\mathrm{Fe}^{3+} / \mathrm{Fe}^{2+}$ <br> electrode system has been added to the $E^{o}$ cell value instead of being subtracted |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| 2(b) | The only correct answer is C (platinum and titanium) <br> $\boldsymbol{A} \quad$ is not correct because the $F e^{3+} \mid F e^{2+}$ electrode system requires a platinum electrode <br> $\boldsymbol{B} \quad$ is not correct because the $F e^{3+} \mid F e^{2+}$ electrode system requires a platinum electrode and the $T i^{2+} \mid$ Ti electrode <br> system requires a titanium electrode. <br> $\boldsymbol{D \quad i s ~ n o t ~ c o r r e c t ~ b e c a u s e ~ t h e ~} T i^{2+} \mid$ Ti electrode system requires a titanium electrode. | $\mathbf{1}$ |


| Question number | Answer | Mark |
| :---: | :---: | :---: |
| 2(c) | The only correct answer is $\mathbf{C}\left(2 \mathrm{~mol} \mathrm{dm}^{-3}\right.$ and $\left.1 \mathrm{~mol} \mathrm{dm}^{-3}\right)$ <br> $\boldsymbol{A} \quad$ is not correct because the cell solution must be $1 \mathrm{~mol} \mathrm{dm}{ }^{-3}$ with respect to both $\mathrm{Fe}^{2+}$ and $\mathrm{Fe}^{3+}$ therefore the dilution on mixing and the two $\mathrm{Fe}^{3+}$ ions in each iron(III) sulfate must be taken into account <br> $\boldsymbol{B}$ is not correct because the cell solution must be $1 \mathrm{~mol} \mathrm{dm}{ }^{-3}$ with respect to both $\mathrm{Fe}^{2+}$ and $\mathrm{Fe}^{3+}$ therefore the dilution on mixing and the two $\mathrm{Fe}^{3+}$ ions in each iron(III) sulfate must be taken into account <br> $\boldsymbol{D}$ is not correct because the cell solution must be 1 mol $\mathrm{dm}^{-3}$ with respect to both $\mathrm{Fe}^{2+}$ and $\mathrm{Fe}^{3+}$ therefore the dilution on mixing and the two $\mathrm{Fe}^{3+}$ ions in each iron(III) sulfate must be taken into account | 1 |



| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{4}$ | The only correct answer is A (electron-pair donor) | $\mathbf{1}$ |
|  | B is not correct because ligands do not need to be negatively charged <br> C is not correct because ligands are not electron pair acceptors <br> $\boldsymbol{D}$ is not correct because ligands are not electron pair acceptors and do not need to be negatively charged |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{5}$ | The only correct answer is C (d-d transitions are not possible because the d orbitals are fully occupied) | $\mathbf{1}$ |
|  | $\boldsymbol{A} \quad$ is not correct because the d orbitals in copper(I) can be split <br> $\boldsymbol{B} \quad$ is not correct because no d-d transitions occur <br> $\boldsymbol{D} \quad$ is not correct because the ease of oxidation of an ion does not affect the colour of its complex |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{6}$ | The only correct answer is B(type of ligand is bidentate; coordination number of copper(II) is 6) | $\mathbf{1}$ |
|  | $\boldsymbol{A} \quad$ is not correct because3 is the number of ligands not the coordination number |  |
| $\boldsymbol{C} \quad$ is not correct because the ligand is bidentate and 3 is the number of ligands |  |  |
| $\boldsymbol{D} \quad$ is not correct because the ligand is bidentate |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{7}$ | The only correct answer is $\mathbf{D}($ green precipitate and precipitate turns brown $)$ | $\mathbf{1}$ |
|  | $\boldsymbol{A} \quad$ is not correct because $\mathrm{Fe}(\mathrm{OH})_{2}$ is green (initially) |  |
| $\boldsymbol{B} \quad$ is not correct because the precipitate turns brown on standing |  |  |
| $\boldsymbol{C} \quad$ is not correct because $\mathrm{Fe}(\mathrm{OH})_{2}$ is green (initially) |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{8}$ | The only correct answer is A (deprotonation and ligand exchange) | $\mathbf{1}$ |
| $\boldsymbol{B} \quad$ is not correct because the formation of the ammine complex involves ligand exchange |  |  |
| $\boldsymbol{C} \quad$ is not correct because the formation of the precipitate involves deprotonation of water ligands and the formation of |  |  |
| the ammine complex involves ligand exchange |  |  |
| $\boldsymbol{D} \quad$ is not correct because the formation of the precipitate involves deprotonation of water ligands |  |  |$\quad$.


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{9}$ | The only correct answer is C (four) | $\mathbf{1}$ |
| $\boldsymbol{A} \quad$ is not correct because this omits two isomers |  |  |
| $\boldsymbol{B} \quad$ is not correct because in the Kekulé structure1,2-dichlorobenzene has two isomers, one with the carbon atoms |  |  |
| carrying the chlorines joined by a single bond and the other with them joined by a double bond |  |  |
| $\boldsymbol{D} \quad$ is not correct because this overlooks the fact that the 1,3 and 1,5 structures are identical |  |  |\(\quad\left\{\begin{array}{l} <br>

\hline\end{array}\right.\)

| Question <br> number | Answer | Mark |  |
| :--- | :--- | :--- | :---: |
| $\mathbf{1 0}$ | The only correct answer is $\mathbf{D}$ | $\mathbf{1}$ |  |
|  |  |  |  |
|  |  |  |  |


| Question number | Answer | Mark |
| :---: | :---: | :---: |
| 11 | The only correct answer is B <br> A is not correct because the $\mathrm{N}---\mathrm{H}-\mathrm{O}$ bond angle should be $180^{\circ}$ <br> $\boldsymbol{C}$ is not correct because hydrogen atoms bonded to carbon atoms cannot form hydrogen bonds <br> D is not correct because hydrogen atoms bonded to carbon atoms cannot form hydrogen bonds | 1 |

$\left.\begin{array}{|l|l|c|}\hline \begin{array}{l}\text { Question } \\ \text { number }\end{array} & \text { Answer } & \text { Mark } \\ \hline \mathbf{1 2} & \text { The only correct answer is A(amides) } & \mathbf{1} \\ & \boldsymbol{B} \quad \text { is not correct because amino acids combine to form polypeptides and proteins, which are polyamides } \\ \boldsymbol{C} \quad \text { is not correct because diacyl chlorides combine with diamines to form polyamides } \\ \boldsymbol{D} \quad \text { is not correct because diamines combine with diacyl chlorides to form polyamides }\end{array}\right]$

| Question number | Answer | Mark |
| :---: | :---: | :---: |
| 13(a) | The only correct answer is B <br> A is not correct because this structure is only possible near neutral pH <br> C is not correct because this structure is formed at low pH <br> D is not correct because this structure dominates at neutral pH | 1 |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 3 ( b )}$ | The only correct answer is D (ionic bonds) | $\mathbf{1}$ |
|  | $\boldsymbol{A} \quad$ is not correct because van der Waals forces are the weakest forces broken |  |
| $\boldsymbol{B} \quad$ is not correct because hydrogen bonds are weaker than ionic bonds |  |  |
| $\boldsymbol{C} \quad$ is not correct because covalent bonds are not broken when amino acids melt |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 4}$ | The only correct answer is D (P is due to C-H aldehyde and Q is due to C=O aldehyde) | $\mathbf{1}$ |
|  | $\boldsymbol{A} \quad$ is not correct because $P$ is too sharp to be an OH stretch and $Q$ is outside the region of $1725-1700 \mathrm{~cm}^{-1}$ for a |  |
| $\boldsymbol{B} \quad$ is not correct because $P$ is too sharp to be an $O H$ stretch |  |  |
| $\boldsymbol{C} \quad$ is not correct because $Q$ is outside the region of $1725-1700 \mathrm{~cm}^{-1}$ for a carboxylic acid $C=O$ stretch |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 5}$ | The only correct answer is C (five) | $\mathbf{1}$ |
|  | A is not correct because the C atoms in methylcyclohexane are not all equivalent <br> $\boldsymbol{B} \quad$ is not correct because there are five types of $C$ atom in methylcyclohexane not three <br> $\boldsymbol{D}$ is not correct because the C atoms in methylcyclohexane are not all different |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 6}$ | The only correct answer is A (one singlet, one doublet and a heptet) | $\mathbf{1}$ |
|  | B is not correct because this ignores the fact that the proton environments on C1 and C3 are the same <br> $\boldsymbol{C}$ is not correct because this is the number of protons in each environment, not the splitting pattern <br> $\boldsymbol{D}$ is not correct because this ignores all the splitting except the effect of one methyl on the C2 proton |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 7}$ | The only correct answer is B $\left(8.4 \mathrm{dm}^{3}\right)$ | $\mathbf{1}$ |
| $\boldsymbol{A} \quad$ is not correct because the number of oxygen atoms in the compound has been doubled |  |  |
| $\boldsymbol{C} \quad$ is not correct because the oxygen atoms in the compound have been omitted from the calculation |  |  |
| $\boldsymbol{D}$ is not correct because the oxygen atoms in the compound have been omitted from the calculation and one oxygen |  |  |
| molecule has been allowed for the combustion of each pair of hydrogen atoms |  |  |$\quad$.

## Section B

| Question number | Answer |  | Additional guidance | Mark |
| :---: | :---: | :---: | :---: | :---: |
| 18(a)(i) | - selection of the correct half-equations from the table <br> - writing the balanced equation | (1) <br> (1) | [1] $\mathrm{MnO}_{4}^{-}+\mathrm{e}^{-} \rightleftharpoons \mathrm{MnO}_{4}{ }^{2-}$ and <br> [7] $\mathrm{MnO}_{4}{ }^{2-}+4 \mathrm{H}^{+}+2 \mathrm{e}^{-}=\mathrm{MnO}_{2}+2 \mathrm{H}_{2} \mathrm{O}$ $3 \mathrm{MnO}_{4}{ }^{2-}+4 \mathrm{H}^{+} \rightleftharpoons 2 \mathrm{MnO}_{4}^{-}+\mathrm{MnO}_{2}+2 \mathrm{H}_{2} \mathrm{O}$ <br> Allow <br> [1] $\mathrm{MnO}_{4}^{-}+\mathrm{e}^{-} \rightleftharpoons \mathrm{MnO}_{4}{ }^{2-}$ and $[5] \mathrm{MnO}_{4}^{-}+8 \mathrm{H}^{+}+5 \mathrm{e}^{-} \rightleftharpoons \mathrm{Mn}^{2+}+4 \mathrm{H}_{2} \mathrm{O}$ <br> which gives $5 \mathrm{MnO}_{4}{ }^{2-}+8 \mathrm{H}^{+} \rightleftharpoons 4 \mathrm{MnO}_{4}^{-}+\mathrm{Mn}^{2+}+4 \mathrm{H}_{2} \mathrm{O}$ <br> Accept e for $\mathrm{e}^{-}$in half equations <br> Correct equation scores (2) marks <br> Unbalanced equation with all species correct scores (1) <br> Use of alkaline half-equation to give $3 \mathrm{MnO}_{4}{ }^{2-}+2 \mathrm{H}_{2} \mathrm{O} \rightleftharpoons 2 \mathrm{MnO}_{4}^{-}+\mathrm{MnO}_{2}+4 \mathrm{OH}^{-}$ scores (1) mark <br> Ignore state symbols even if incorrect | 2 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 18(a)(ii) | - selection of appropriate values and insertion in the correct equation (1) <br> - completion of calculation <br> and <br> evaluation of the feasibility of the reaction | $\begin{aligned} & \begin{aligned} E_{\text {cell }}^{\mathrm{o}} & =2.26-0.56 \\ & =(+) 1.70(\mathrm{~V}) \text { and (positive so } \\ & \text { reaction is) feasible } \end{aligned} \\ & \text { Accept (if second equation given) } \\ & \begin{aligned} & E_{\text {cell }}^{\mathrm{o}}=1.51-0.56 \\ &=(+) 0.95(\mathrm{~V}) \text { and (positive so } \\ & \text { reaction is) feasible } \end{aligned} \end{aligned}$ <br> TE on alkaline disproportionation $E^{\circ}{ }_{\text {cell }}=0.59-0.56=(+) 0.03(\mathrm{~V})$ and (positive so reaction is) feasible scores (2) <br> TE on equations used the wrong way round $-1.70 /-0.95(\mathrm{~V})$ and not feasible scores (1) <br> Just (+)1.70 (V) / (+)0.95 (V) scores (1) <br> Do not award a non-disproportionation reaction | 2 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 18(a)(iii) | For M1 and M2 Either <br> - equation for the reaction <br> - calculation of negative $E^{0}$ cell for the equation <br> Or <br> - clear identification of the appropriate half-equations from the table (1) <br> - use of anticlockwise rule or similar to show that required reaction is not favoured <br> For M3 and M4 <br> - statement that the standard electrode potential values are close <br> - reaction may be shifted in the required direction using concentrated alkali | $\begin{align*} & 2 \mathrm{MnO}_{4}^{-}+\mathrm{MnO}_{2}+4 \mathrm{OH}^{-} \rightleftharpoons 3 \mathrm{MnO}_{4}{ }^{2-}+2 \mathrm{H}_{2} \mathrm{O}  \tag{1}\\ & \quad \text { Allow } \rightarrow \text { for } \rightleftharpoons \\ & \text { or } E_{\text {cell }}^{\mathrm{o}}=0.56-0.59 \\ & E_{\text {cell }}^{\mathrm{o}}=-0.03(\mathrm{~V}) \tag{1} \end{align*}$ <br> Reverse reaction and $E_{\text {cell }}^{0}=(+) 0.03(\mathrm{~V})$ (1) <br> Equations 1 and 2 <br> Desired reaction moves 'clockwise' so not (thermodynamically) feasible Allow calculation of $E^{\circ}$ cell as above <br> $(-) 0.03(\mathrm{~V}) / E^{\mathrm{o}}$ cell is a (very) small Or equilibrium has significant concentrations of the reactants and products <br> Allow if $E_{\text {cell }}^{\mathrm{o}}=(+) 0.03(\mathrm{~V})$ in M2 Standalone mark <br> Allow by increasing alkali concentration No TE on incorrect system Ignore just 'by changing concentrations' references to rate and temperature Ignore state symbols even if incorrect | 4 |


| Question <br> number | Answer | Additional guidance | Mark |
| :--- | :--- | :--- | :---: |
| $\mathbf{1 8 ( b ) ( i )}$ | - colourless |  |  |
|  | and <br> to pale pink | Allow pale green for colourless | $\mathbf{1}$ |
|  |  | Do not award just 'green' for the start colour <br> Accept pink |  |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 18(b)(ii) | - calculation of moles of manganate(VII) in $27.35 \mathrm{~cm}^{3}$ <br> - use of 1:5 ratio to calculate $\mathrm{mol} \mathrm{Fe}^{2+}$ in $25.0 \mathrm{~cm}^{3}$ <br> - scaling to $250.0 \mathrm{~cm}^{3}$ to give $\mathrm{mol} \mathrm{Fe}^{2+}$ in $250.0 \mathrm{~cm}^{3}$ <br> - conversion of mol to g of iron <br> - calculation of percentage of iron in the wire and gives the final value to 3 SF | Example of calculation $\begin{align*} & 27.35 \times 0.0195 \times 10^{-3}  \tag{1}\\ & =5.33325 \times 10^{-4} / 0.000533325(\mathrm{~mol}) \\ & 5 \times 5.33325 \times 10^{-4}  \tag{1}\\ & =2.666625 \times 10^{-3} / 0.002666625(\mathrm{~mol}) \\ & 10 \times 2.666625 \times 10^{-3}  \tag{1}\\ & \left.=2.666625 \times 10^{-2} / 0.02666625 \mathrm{~mol}\right) \\ & 2.666625 \times 10^{-2} \times 55.8=1.48798(\mathrm{~g})  \tag{1}\\ & 100 \times 1.48798 / 1.53=97.25338 \\ & =97.3 \% \end{align*}$ <br> If $A_{\mathrm{r}}(\mathrm{Fe})=56$ is used mass $=1.49331(\mathrm{~g})$ $\% \text { iron = 97.6\% }$ <br> Allow $100 \times 1.49 / 1.53=97.4 \%$ <br> Correct answer to 3 SF and some working scores (5) <br> TE at each stage Ignore premature rounding if final answer correct otherwise allow rounding to at least 3SF | 5 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 18(b)(iii) | An explanation that makes reference to the following <br> - brown suspension formed is manganese(IV) oxide / $\mathrm{MnO}_{2}$ <br> - $\mathrm{Mn}(\mathrm{VII})$ to $\mathrm{Mn}(\mathrm{II})$ provides 5 electrons per $\mathrm{MnO}_{4}{ }^{-}$but $\mathrm{Mn}(\mathrm{VII})$ to $\mathrm{Mn}(\mathrm{IV})$ only provides 3 electrons <br> (1) <br> - so more $\mathrm{MnO}_{4}^{-}$is needed / titre is greater | Reference to half-equations 6 \& 7 in the table <br> Allow <br> manganese(IV) oxide / $\mathrm{MnO}_{2}$ formed (in alkaline conditions) <br> Accept explanation in terms of oxidation numbers <br> Standalone mark <br> If no other mark is scored, allow one mark for the titration is no longer quantitative as another reaction is (also) taking place. | 3 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 18(c)(i) | (Balance: $\frac{100 \times 0.005 \times 2}{1.53}=0.65$ ) <br> - burette: $\frac{100 \times 0.05 \times 2}{27.35}=0.37(\%)$ <br> - pipette: $\frac{100 \times 0.06}{25}=0.24(\%)$ <br> - volumetric flask: $\frac{100 \times 0.3}{250}=0.12(\%)$ <br> $($ Total $=1.38)$ | All three percentages correct scores (2) Any two percentages correct scores (1) IGNORE SF | 2 |
| Question number | Answer | Additional guidance | Mark |
| 18(c)(ii) | An explanation that makes reference to <br> - total percentage uncertainty is (approximately) $1.38 \%$ (1) <br> - because the total percentage uncertainty is much bigger than 0.863 , the answer should be to no more than $2 / 3 \mathrm{SF}(1)$ | TE on 18(c)(i) <br> Allow 1.4\% / 2 SF <br> Ignore 'data given to 3SF) | 2 |


| Question <br> number | Answer | Additional guidance | Mark |
| :--- | :--- | :--- | :---: |
| $\mathbf{1 9 ( a ) ( \mathbf { i } )}$ | • Amine | Do not award ammine | $\mathbf{1}$ |


| Questio numbe | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 19(a)(ii) | - rewrite the Ideal Gas Equation in terms of mass and molar mass <br> - change the subject of the equation <br> - change volume to $\mathrm{m}^{3}$ and temperature to K <br> native method <br> change equation subject (1) $n=p V / R T$ <br> change volume to $\mathrm{m}^{3}$ and temperature to K (1) $\begin{aligned} & \mathrm{n}=103000 \times 1.57 \times 10^{-4} / 8.31 \times 288=6.7568 \times 10^{-3}(1) \\ & M_{\mathrm{r}}=0.493 / 6.7568 \times 10^{-3}=72.963=73\left(\mathrm{~g} \mathrm{~mol}^{-1}\right)(1) \end{aligned}$ | Example of calculation $P V=\frac{m}{M_{\mathrm{r}}} R T$ $\begin{equation*} M_{\mathrm{r}}=m R T / P V \tag{1} \end{equation*}$ $157 \mathrm{~cm}^{3}=1.57 \times 10^{-4} / 0.000157 \mathrm{~m}^{3}$ <br> Allow conversion to $\mathrm{kPa} \&$ vol to $\mathrm{dm}^{3}$ $15^{\circ} \mathrm{C}=288 \mathrm{~K}$ $\mathrm{M}_{\mathrm{r}}=\frac{0.493 \times 8.31 \times 288}{103000 \times 1.57 \times 10^{-4}}$ $=72.963=73.0 / 73\left(\mathrm{~g} \mathrm{~mol}^{-1}\right)$ <br> Ignore just ' g ' do not award $72.9\left(\mathrm{~g} \mathrm{~mol}^{-1}\right)$ <br> Correct answer with some working scores (4) <br> Correct answer with no working scores zero <br> Ignore premature rounding if final answer correct otherwise allow rounding to at least 3SF Ignore SF except 1 SF | 4 |


| Question <br> number | Answer | Additional guidance | Mark |
| :--- | :--- | :--- | :---: |
| 19(b)(i) | - hydrogen chloride $/ \mathrm{HCl}$ | Allow <br> hydrochloric acid $/ \mathrm{HCl}(\mathrm{aq})$ | $\mathbf{1}$ |


| Question <br> number | Answer | Additional guidance | Mark |
| :--- | :--- | :--- | :---: |
| 19(b)(ii) | $\bullet$ N-substituted amide / -CONHR | Accept <br> amide / acid amide / -CONH $/$ / -CONH- <br> If name and formula are given, both must be <br> correct | $\mathbf{1}$ |


| Question | Answer | Additional guidance |  |  |  |  | Mark |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 19(b)(iii) | - conversion of percentages by mass into moles <br> - evaluation of moles and division by the smallest value to give a ratio <br> - conversion of ratio into an empirical formula | Example of calculation |  |  |  |  | 3 |
|  |  |  | C | H | N | O |  |
|  |  | \% | 62.6 | 11.3 | 12.2 | 13.9 |  |
|  |  | mol | $\begin{aligned} & 62.6 / 12 \\ & (=5.2167) \end{aligned}$ | $\begin{aligned} & 11.3 / 1 \\ & (=11.3) \end{aligned}$ | $\begin{aligned} & 12.2 / 14 \\ & (= \\ & 0.8714) \end{aligned}$ | $\begin{aligned} & 13.9 / 16 \\ & =0.86875 \end{aligned}$ |  |
|  |  | Ratio | 6.00 | 13.01 | 1.00 | 1.00 |  |
|  |  | $\mathrm{C}_{6} \mathrm{H}_{13} \mathrm{NO}$ <br> Correct <br> Ignore | answer with <br> F except 1 | no work <br> FF in mol | g scores <br> calculatio |  |  |



| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| $\begin{aligned} & * 19(c) \\ & \text { cont } \end{aligned}$ | Indicative points <br> - IP1 three peaks indicates three proton environments <br> - IP2 no splitting shows no proton environment is adjacent to another <br> - IP3 chemical shift $=7(\mathrm{ppm})$ indicates $\mathrm{N}-\mathbf{H}$ proton <br> - IP4 nine protons in one environment and no coupling indicates $\left(\mathrm{CH}_{3}\right)_{3} \mathrm{C}$ - <br> - IP5 chemical shift $=2(\mathrm{ppm})$ indicates $\mathrm{CH}_{3} \mathrm{C}=\mathrm{O}$ protons <br> - IP6 structure is | Marks may be awarded for IPs annotated on the NMR or on the structure <br> Allow three types of proton / hydrogen <br> Allow 'peaks have one split' <br> Accept $\mathrm{CH}_{3} \mathrm{C}=\mathrm{O}$ indicated by amide responsible for the peak at 2 (ppm) <br> Allow <br> $\mathrm{CH}_{3} \mathrm{C}=\mathrm{O}$ indicated by use of ethanoyl chloride <br> Allow displayed, structural or skeletal formulae or any combination of these e.g. | 6 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 19(d) |  | Allow displayed, structural or skeletal formulae or any combination of these e.g. <br> Allow TE on the amide in 19(c)providing $\mathbf{X}$ is shown as an amine | 1 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 20(a)(i) | - calculation of the difference between the enthalpies of combustion of cyclohexene and cyclohexa-1,4-diene <br> - subtraction of the calculated difference from the enthalpy of combustion of cyclohexa-1,4-diene to give the enthalpy of combustion of cyclohexa-1,3,5-triene | Example of calculation $\begin{equation*} -3752-(-3584)=-168\left(\mathrm{~kJ} \mathrm{~mol}^{-1}\right) \tag{1} \end{equation*}$ <br> Ignore sign of 168 $-3584-(-168)=-3416\left(\mathrm{~kJ} \mathrm{~mol}^{-1}\right)$ <br> TE on numerical errors in M1 <br> Correct answer with some working scores (2) | 2 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 20(a)(ii) | An explanation that makes reference to <br> - calculation of the difference between the enthalpies of combustion of benzene and cyclohexa-1,3,5-triene <br> - benzene more stable than cyclohexa-1,3,5-triene by $149\left(\mathrm{~kJ} \mathrm{~mol}^{-1}\right)$ <br> - (benzene more stable)because the $\pi$ electrons in benzene are delocalised | $\begin{equation*} -3416-(-3267)=-149\left(\mathrm{~kJ} \mathrm{~mol}^{-1}\right) \tag{1} \end{equation*}$ <br> TE on value calculated in (a)(i) <br> Ignore omission of negative sign <br> Allow enthalpy of combustion of benzene less negative / less exothermic than that of cyclohexa-1,3,5-triene and so benzene is more stable <br> If M1 is scored Allow just 'benzene more stable than cyclohexa-1,3,5-triene' <br> Allow benzene has delocalisation energy / resonance stability /bonds delocalised standalone mark <br> Do not award if the enthalpy of combustion of benzene is more negative than that calculated for cyclohexa-1,3,5-triene | 3 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 20(b) | An answer that makes reference to four of the following: <br> Similarities <br> - both reactions involve electrophilic attack <br> - both reactions form a carbocation <br> Differences <br> - reaction with benzene is substitution (because the stable benzene ring is retained / restored) <br> - reaction with cyclohexene is addition (because $\sigma$ bonds stronger than $\pi$ bonds) <br> - reaction with benzene requires a catalyst (and heat) <br> and <br> whereas reaction with cyclohexene occurs under normal laboratory conditions | This may be shown using a labelled diagram <br> Accept $2 \mathrm{C}-\mathrm{H}$ and $1 \mathrm{C}-\mathrm{C}$ stronger than $\mathrm{C}=\mathrm{C}$ and $\mathrm{Br}-\mathrm{Br}$ <br> Allow cyclohexene reaction does not require catalyst / heat <br> Allow cyclohexene reacts with bromine water <br> Do not award if conditions are incorrect <br> e.g. cyclohexene reaction requires heat | 4 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 20(c)(i) | - 2,4,6-tribromophenol <br> OR | If name and formula are given both must be correct <br> Accept Kekulé structure <br> Allow <br> Correct structure and tribromophenol <br> Ignore punctuation errors in the name such as omission of commas or hyphen and inclusion of spaces | 1 |


| Question <br> number | Answer | Additional guidance | Mark |
| :--- | :--- | :--- | :---: |
| $\mathbf{2 0 ( c ) ( i i )}$ | An explanation that makes reference to <br> - lone pair of electrons on the oxygen <br> - overlap / interact with the $\pi$ electrons of the ring <br> and <br> increasing its electron density / becomes more susceptible to electrophilic <br> attack | (1) | Allow lone pair of electrons on the OH <br> Allow (lone pair) donated to the $\pi$ electrons <br> Ignore ingreasing the reactivity of the ring <br> Ignore reference to phenol being a <br> nucleophile |

## Section C

| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 21(a) | An explanation that makes reference to the following points <br> - the electronic structure of $\mathrm{Fe}(\mathrm{II})$ is $[\mathrm{Ar}] 3 \mathrm{~d}^{6}$ and $\mathrm{Fe}(\mathrm{III})$ is $[\mathrm{Ar}] 3 \mathrm{~d}^{5}$ <br> - $3 \mathrm{~d}^{5}$ is more stable than $3 \mathrm{~d}^{6}$ because the $3 \mathrm{~d}^{6}$ subshell has two paired electrons which results in repulsion / pairing of electrons is (energetically) less favourable | Ar may be given as $1 s^{2} 2 s^{2} 2 p^{6} 3 s^{2} 3 p^{6}$ <br> Allow just Fe (II) is $3 \mathrm{~d}^{6}$ and Fe (III) is $3 \mathrm{~d}^{5}$ <br> Allow half-filled subshell is more stable Allow 1 electron in each orbital is stable If no other mark is scored, just ${ }^{ } \mathrm{Fe}(\mathrm{III})$ has a half-filled 3d subshell scores (1) <br> Allow use of 'orbital' or '(quantum) shell' for subshell | 2 |
| Question number | Answer | Additional guidance | Mark |
| 21(b)(i) | - $\mathrm{Fe}^{2+}(\mathrm{g}) \rightarrow \mathrm{Fe}^{3+}(\mathrm{g})+\mathrm{e}\left({ }^{-}\right)$ | Allow $\mathrm{Fe}^{2+}(\mathrm{g})-\mathrm{e}\left({ }^{-}\right) \rightarrow \mathrm{Fe}^{3+}(\mathrm{g})$ <br> Accept $\mathrm{e}^{(-)}(\mathrm{g})$ | 1 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 21(b)(ii) | An answer that makes reference to the following points <br> M1 <br> - conversion of iron(II) $/ \mathrm{Fe}^{2+}$ to iron(III) $/ \mathrm{Fe}^{3+}$ requires $2958 \mathrm{~kJ} \mathrm{~mol}^{-1}$ OR <br> conversion of iron(II) $/ \mathrm{Fe}^{2+}$ to iron(III) $/ \mathrm{Fe}^{3+}$ requires (large amounts of ) energy <br> OR <br> conversion of iron(II) $/ \mathrm{Fe}^{2+}$ to iron(III) $/ \mathrm{Fe}^{3+}$ is (very) endothermic <br> M2 <br> - this energy is recovered by hydration (which is exothermic) <br> M3 <br> - the hydration of iron(III) $/ \mathrm{Fe}^{3+}$ is more exothermic than iron(II) $/ \mathrm{Fe}^{2+}$ and because the iron(III) has a greater charge | These marks may be awarded if a labelled Hess cycle is used <br> Accept 'charge density' for charge Ignore smaller ionic radius | 3 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 21(c) | An answer that makes reference to the following points <br> - the energy difference between the two sets of (3)d orbitals is different <br> - because the ligands / ions are different <br> - so radiation / light / quanta absorbed from different regions of the visible spectrum | Allow the different energy levels of the (3)d subshell <br> Do not award orbital for orbitals Do not award (quantum) shell for subshell <br> Ignore just 'complexes are different' <br> Accept different radiation / light frequencies or different radiation / light wavelengths transmitted / reflected <br> Do not award radiation / light emitted | 3 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 21(d)(i) | - conversion of $\%$ transmittance into $\log (\%$ transmittance $)$ <br> - use of the graph to obtain a value for the concentration of iron in $\mathrm{mg} \mathrm{dm}^{-3}$ <br> - calculation of the mass of iron in $500 \mathrm{~cm}^{3}$ of iron solution which is also the mass of iron in 20 g of sodium carbonate $\begin{aligned} & \text { OR } \\ & \text { M3 Concentration of } \mathrm{Na}_{2} \mathrm{CO}_{3}=20 \times 1000 / 500 \div 1000 \\ &=40000 \mathrm{~m} \mathrm{~g} \mathrm{dm}^{-3} \end{aligned}$ | Example of calculation $\log (39.8)=1.60$ <br> $[\mathrm{Fe}]=0.44\left(\mathrm{mg} \mathrm{dm}^{-3}\right)($ allow $0.42-0.46)$ <br> $0.5 \times 0.44=0.22 \mathrm{mg}\left(\right.$ in $\left.500 \mathrm{~cm}^{3} / 20 \mathrm{~g}\right)$ <br> Accept $0.21-0.23(\mathrm{mg})$ <br> OR <br> $0.00022 / 2.2 \times 10^{-4} \mathrm{~g}$ (in $500 \mathrm{~cm}^{3} / 20 \mathrm{~g}$ ) <br> $\left(10^{6} \times 2.2 \times 10^{-4}\right) \div 20=11 \mathrm{ppm}$ <br> (So Fe is within the stated specification) <br> TE at each stage <br> Correct answer with some working scores | 4 |


| Question number | Answer | Additional guidance | Mark |
| :---: | :---: | :---: | :---: |
| 21(d)(ii) | - (thioglycolic acid is a) bidentate (ligand) <br> - because there are three ligands per complex ion and the coordination number of $\mathrm{Fe}^{3+}$ is (usually) six <br> OR <br> can form dative bonds using the lone pairs on the SH and the COOH groups | Allow three ligands replace six (monodentate) water ligands <br> Accept S and O atoms Ignore just 'forms two dative covalent bonds' <br> Do not award two dative covalent bonds from COOH | 2 |
| Question number | Answer | Additional guidance | Mark |
| 21(e)(i) | - catalyst and reactants are in the same phase | Accept 'same state' <br> Allow all species are in aqueous solution <br> Allow 'It' is in the same phase / state as the <br> reactants <br> Ignore reference to producta | 1 |


| Question <br> number | Answer | Additional guidance | Mark |  |
| :--- | :--- | :--- | :--- | :---: |
| 21(e)(ii) |  | (1) | $\mathrm{S}_{2} \mathrm{O}_{8}{ }^{2-}+2 \mathrm{Fe}^{2+} \rightarrow 2 \mathrm{SO}_{4}{ }^{2-}+2 \mathrm{Fe}^{3+}$ |  |
|  | $\bullet$ equation for oxidation of iron(II) by peroxodisulfate | (1) | $2 \mathrm{I}^{-}+2 \mathrm{Fe}^{3+} \rightarrow \mathrm{I}_{2}+2 \mathrm{Fe}^{2+}$ |  |
|  |  | equation for oxidation of iodide ions by iron(III) | Allow multiples $/$ equations in any order <br> Ignore state symbols even if incorrect <br> Two unbalanced equations with all species <br> correct scores (1) | $\mathbf{2}$ |


| Question <br> number | Answer | Additional guidance | Mark |
| :--- | :--- | :--- | :---: |
| 21(e)(iii) | uncatalysed reaction involves two negatively charged ions reacting <br> and <br> catalysed steps involve oppositely charged ions reacting (which is <br> kinetically more favourable / lowers the activation energy) | Ignore general definitions of catalysts | $\mathbf{1}$ |


| Question <br> number | Answer | Additional guidance | Mark |
| :--- | :--- | :--- | :---: |
| $\mathbf{2 1 ( f )}$ | - chloride ions are large so steric hindrance is too great for six ligands to |  |  |
| coordinate around the central ion |  |  |  |$\quad$| Allow just 'chloride ions are large' |
| :--- |
| Allow chlorine ligands are (too) large |
| Do not award chlorine atoms |
| Ignore reference to repulsion between |
| negative chloride ions |$\quad$|  |
| :--- |

