## Mark Scheme (Results)

## October 2023

Pearson Edexcel International Advanced Level In Chemistry (WCH15)
Paper 01 Unit 5: 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.

## Quality of Written Communication

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

- write legibly, with accurate use of spelling, grammar and punctuation in order to make the meaning clear
- select and use a form and style of writing appropriate to purpose and to complex subject matter
- organise information clearly and coherently, using specialist
vocabulary when appropriate. Full marks will be awarded if the candidate has demonstrated the above abilities.
Questions where QWC is likely to be particularly important are indicated (QWC) in the mark scheme, but this does not preclude others.

| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1}$ | The only correct answer is $\mathbf{D}$ (zinc) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because cobalt forms a stable $\mathrm{Co}^{2+}$ ion with incompletely-filled d-orbitals |  |
|  | $\boldsymbol{B} \quad$ is incorrect because copper forms a stable $\mathrm{Cu}^{2+}$ ion with incompletely-filled d-orbitals |  |
| $\boldsymbol{C} \quad$ is incorrect because nickel forms a stable $\mathrm{Ni}^{2+}$ ion with incompletely-filled d-orbitals |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{2}$ | The only correct answer is $\mathbf{D}\left(\mathrm{VO}_{3}{ }^{-}\right.$and $\left.\mathrm{VO}_{2}{ }^{+}\right)$ | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because chromium has oxidation numbers +6 and +3 respectively |  |
|  | $\boldsymbol{B} \quad$ is incorrect because copper has oxidation numbers +1 and +2 respectively |  |
| $\boldsymbol{C} \quad$ is incorrect because manganese has oxidation numbers +3 and +4 respectively |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{3}$ | The only correct answer is $\mathbf{D}$ (6) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because although there are two different ligands, there are 6 atoms bonded to the central ion |  |
|  | $\boldsymbol{B} \quad$ is incorrect because the charge on Cr is 3+ but there are 6 atoms bonded to the central ion |  |
| $\boldsymbol{C} \quad$ is incorrect because although there are 4 ligands, there are 6 atoms bonded to the central ion |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{4}$ | The only correct answer is $\mathbf{C}\left(\mathrm{Ni}^{2+}\right)$ | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because $\mathrm{Cu}^{2+}$ gives a blue precipitate with aqueous sodium hydroxide and with aqueous ammonia |  |
|  | $\boldsymbol{B} \quad$ is incorrect because the precipitate formed with $F e^{2+}$ and aqueous ammonia is insoluble in excess ammonia |  |
| $\boldsymbol{D} \quad$ is incorrect because $V^{2+}$ is a purple solution |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{5}$ | The only correct answer is $\mathbf{B}\left(\left[\mathrm{Zn}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}+2 \mathrm{NH}_{3} \rightarrow \mathrm{Zn}(\mathrm{OH})_{2}\left(\mathrm{H}_{2} \mathrm{O}\right)_{4}+2 \mathrm{NH}_{4}^{+}\right)$ | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because the precipitate should not have a positive charge and the charges do not balance |  |
| $\boldsymbol{C} \quad$ is incorrect because $\left[\mathrm{Zn}\left(\mathrm{NH}_{3}\right)_{4}\left(\mathrm{H}_{2} \mathrm{O}\right)_{2}\right]^{2+}$ is formed when the precipitate dissolves in excess aqueous ammonia |  |  |
| $\boldsymbol{D} \quad$ is incorrect because $\mathrm{Zn}\left(\mathrm{NH}_{3}\right)_{4}\left(\mathrm{H}_{2} \mathrm{O}\right)_{2}$ should have a 2+ charge and the equation is not balanced |  |  |$\quad$.


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{6}$ | The only correct answer is $\mathbf{B}\left(\mathrm{Mn}^{2+}\right.$ acts as a catalyst; concentration of reactants decreases $)$ | $\mathbf{( 1 )}$ |
|  | $\boldsymbol{A} \quad$ is incorrect because the kinetic energies of the particles do not change |  |
| $\boldsymbol{C} \quad$ is incorrect because $\mathrm{MnO}_{4}{ }^{-}$is not a catalyst and the kinetic energies of the particles do not change |  |  |
| $\boldsymbol{D} \quad$ is incorrect because $\mathrm{MnO}_{4}{ }^{-}$is not a catalyst |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{7}$ | The only correct answer is $\mathbf{C}\left(\Delta S_{\text {total }}\right.$ and $\left.\ln K\right)$ | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because $E_{- \text {cell }}^{o}$ is not directly proportional to $\Delta_{\mathrm{r}} H$ |  |
| $\boldsymbol{B} \quad$ is incorrect because $E_{- \text {cell }}^{o}$ is not directly proportional to $\Delta_{\mathrm{r}} H$ or to $\ln R T$ |  |  |
|  | $\boldsymbol{D} \quad$ is incorrect because $E_{- \text {cell }}^{o}$ is not directly proportional to $\ln R T$ |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{8}$ | The only correct answer is A (standard reduction potential; most negative to most positive) | (1) |
|  | $\boldsymbol{B} \quad$ is incorrect because the electrochemical series has the most negative standard electrode potential first |  |
| $\boldsymbol{C} \quad$ is incorrect because standard cell potentials are determined from two standard electrode potentials |  |  |
| D $\quad$ is incorrect because standard cell potentials are determined from two standard electrode potentials and the |  |  |
| electrochemical series has the most negative standard electrode potential first |  |  |$\quad$.


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{9}$ | The only correct answer is $\mathbf{A}\left(\mathrm{H}_{2}+2 \mathrm{OH}^{-} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}+2 \mathrm{e}^{-}\right)$ | (1) |
|  | $\boldsymbol{B} \quad$ is incorrect because $H^{+}$ions cannot be produced in an alkaline solution |  |
|  | $\boldsymbol{C} \quad$ is incorrect because $H^{+}$ions cannot be produced in an alkaline solution |  |
| $\boldsymbol{D} \quad$ is incorrect because $H^{+}$ions cannot be produced in an alkaline solution |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 0}$ | The only correct answer is B (negative; positive) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because $E_{-c e l l}^{o}=E_{r h s}-E_{\text {lhs }}$ so $0.17-(-0.40)=+0.57$ Vor $0.40-(-0.17)=+0.57 \mathrm{~V}$ |  |
|  | $\boldsymbol{C} \quad$ is incorrect because $E_{-c e l l}^{o}=E_{r h s}-E_{\text {lhs }}$ so $0.17-(-0.40)=+0.57$ V or $0.40-(-0.17)=+0.57 \mathrm{~V}$ |  |
|  | $\boldsymbol{D} \quad$ is incorrect because $E_{-c e l l}^{o}=E_{r h s}-E_{\text {lhs }} \operatorname{so} 0.17-(-0.40)=+0.57$ Vor $0.40-(-0.17)=+0.57 \mathrm{~V}$ |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 1}$ | The only correct answer is C (magnesium) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because $1.635 \div 65.4=0.025 \mathrm{~mol}$ of zinc produced which gives a relative atomic mass of 24.3 for $G$ |  |
|  | $\boldsymbol{B} \quad$ is incorrect because $1.635 \div 65.4=0.025 \mathrm{~mol}$ of zinc produced which gives a relative atomic mass of 24.3 for $G$ |  |
| $\boldsymbol{D} \quad$ is incorrect because $1.635 \div 65.4=0.025 \mathrm{~mol}$ of zinc produced which gives a relative atomic mass of 24.3 for $G$ |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 2}$ | The only correct answer is $\mathbf{D}$ (phenylamine) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because the lone pair of electrons on $N$ in ammonia is not delocalised so can be donated more easily |  |
|  | $\boldsymbol{B} \quad$ is incorrect because the lone pair of electrons on $N$ in butylamine is not delocalised so can be donated more easily |  |
|  | $\boldsymbol{C} \quad$ is incorrect because the lone pair of electrons on $N$ in ethylamine is not delocalised so can be donated more easily |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 3}$ | The only correct answer is B $\left(\mathrm{H}_{2} \mathrm{NCH}\left(\mathrm{CH}_{3}\right) \mathrm{COO}^{-}\right)$ | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because this is the structure of the uncharged molecule |  |
| $\boldsymbol{C} \quad$ is incorrect because this structure would exist at pH less than 6.0 |  |  |
| $\boldsymbol{D} \quad$ is incorrect because this is the structure of the zwitterion |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 4}$ | The only correct answer is $\mathbf{A}\left(\mathrm{CH}_{2}=\mathrm{CHCOOH}\right)$ | (1) |
|  | $\boldsymbol{B} \quad$ is incorrect because phenol does not react with ethanol |  |
|  | $\boldsymbol{C} \quad$ is incorrect because 2-propen-1-ol does not react with sodium hydroxide or ethanol |  |
| $\boldsymbol{D} \quad$ is incorrect because ethanoic acid does not react with hydrogen in the presence of a nickel catalyst |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 5 ( a )}$ | The only correct answer is A (further substitution by a nitro group occurs) | (1) |
|  | $\boldsymbol{B} \quad$ is incorrect because nitrobenzene does not decompose at $80^{\circ} \mathrm{C}$ |  |
|  | $\boldsymbol{C} \quad$ is incorrect because fuming sulfuric acid is needed for the substitution of $\mathrm{SO}_{3} \mathrm{H}$ |  |
| $\boldsymbol{D} \quad$ is incorrect because nitric acid does not decompose at $80^{\circ} \mathrm{C}$ |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 5 ( b )}$ | The only correct answer is C (Sn and concentrated $\mathrm{HCl}(\mathrm{aq})$ are added first, then $\mathrm{NaOH}(\mathrm{aq})$ is added at the end) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because the acid and alkali would react to form a salt if they are added together |  |
| $\boldsymbol{B} \quad$ is incorrect because the acid and alkali would react to form a salt if they are added together |  |  |
| $\boldsymbol{D} \quad$ is incorrect because dilute hydrochloric acid would not react quickly enough with the tin |  |  |


| Question number | Answer | Mark |
| :---: | :---: | :---: |
| 15(c) | The only correct answer is B <br> $\boldsymbol{A}$ is incorrect because the chlorine is not bonded covalently to the nitrogen <br> C is incorrect because the chlorine is not bonded covalently to the nitrogen <br> D is incorrect because the charge should be on the nitrogen on the right not the nitrogen on the left | (1) |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 5 ( d )}$ | The only correct answer is B (alkaline) | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because a phenoxide ion is needed for the reaction and that is produced in alkaline solution |  |
|  | $\boldsymbol{C} \quad$ is incorrect because a phenoxide ion is needed for the reaction and that is produced in alkaline solution |  |
| $\boldsymbol{D} \quad$ is incorrect because a phenoxide ion is needed for the reaction and that is produced in alkaline solution |  |  |


| Question <br> number | Answer | Mark |
| :--- | :--- | :---: |
| $\mathbf{1 6}$ | The only correct answer is $\mathbf{D}\left(8\left(\mathrm{~cm}^{3}\right)\right)$ | (1) |
|  | $\boldsymbol{A} \quad$ is incorrect because $2 \mathrm{~cm}^{3}$ of methane reacts with $4 \mathrm{~cm}^{3}$ of oxygen |  |
|  | $\boldsymbol{B} \quad$ is incorrect because $4 \mathrm{~cm}^{3}$ of methane would react with $4 \mathrm{~cm}^{3}$ of oxygen if they reacted in a 1:1 mole ratio |  |
|  | $\boldsymbol{C} \quad$ is incorrect because $6 \mathrm{~cm}^{3}$ would be the volume of argon if methane reacted with oxygen in a 1:1 mole ratio |  |


| Question number | Answer | Mark |
| :---: | :---: | :---: |
| 17 | The only correct answer is $\mathbf{A}$ ( $x$ is 30 and $y$ is 40 ) <br> B is incorrect because water is a liquid at room temperature <br> C is incorrect because $10 \mathrm{~cm}^{3}$ of but-1-ene reacts with $60 \mathrm{~cm}^{3}$ of oxygen to form $40 \mathrm{~cm}^{3}$ of carbon dioxide so there is an initial decrease of $30 \mathrm{~cm}^{3}$ <br> D is incorrect because $10 \mathrm{~cm}^{3}$ of but-1-ene reacts with $60 \mathrm{~cm}^{3}$ of oxygen to form $40 \mathrm{~cm}^{3}$ of carbon dioxide so there is an initial decrease of $30 \mathrm{~cm}^{3}$ and water is a liquid at room temperature | (1) |



| Question <br> Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 18(b)(i) | An explanation that makes reference to the following points: <br> (concentrated hydrochloric acid) <br> - increases the concentration of $\mathrm{H}^{+}$ions in the first equilibrium (and displaces it to the right) <br> so increases the value of $E / E>1.33$ (V) <br> (concentrated hydrochloric acid) <br> - increases the concentration of chloride ions in the second equilibrium (and displaces it to the left) so decreases the value of $E / E<1.36$ (V) <br> - the difference between 1.33 and 1.36 is (very) small and so using concentrated hydrochloric acid, $E_{\text {cell }}$ will be positive (so the reaction occurs) | Ignore any references to $E_{a} /$ energy <br> Allow just 'when $\left[\mathrm{H}^{+}\right]$increases the first equilibrium shifts to the right' <br> Allow because the coefficient for $\mathrm{H}^{+}$is 14 , the position of equilibrium is very sensitive to the concentration of $\mathrm{H}^{+}$ <br> Allow just 'when $\left[\mathrm{Cl}^{-}\right]$increases the second equilibrium shifts to the left' <br> There must be some indication of the equilibrium referred to but can simply be $\mathrm{Cl}_{2}: 2 \mathrm{Cl}^{-}$ <br> Allow answer in terms of first $E^{\text {o }}$ increasing (above 1.36 (V)) or second $E^{\mathrm{o}}$ decreasing (below $1.33(\mathrm{~V})$ ) so $E_{\text {cell }}$ will be positive for M3 <br> Allow chlorine escapes and displaces second equilibrium to the left and decreases $E^{\circ}$ decreasing below $1.33(\mathrm{~V})$ so $E_{\text {cell }}$ will be positive <br> Ignore references to anode/cathode | (3) |


| Question Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 18(b)(ii) | - left hand side of cell diagram <br> - central vertical lines and right hand side of cell diagram | Example of cell diagram: <br> $\underset{\mathrm{Pt}(\mathrm{s})}{ }\left\|2 \mathrm{Cl}^{-}(\mathrm{aq})\right\| \mathrm{Cl}_{2}(\mathrm{~g}) \\|\left[\mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}(\mathrm{aq})+14 \mathrm{H}^{+}(\mathrm{aq})\right],\left[2 \mathrm{Cr}^{3+}(\mathrm{aq})+7 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})\right] \mid \mathrm{Pt}(\mathrm{s})$ <br> Allow comma between $\mathrm{Cl}^{-}$and $\mathrm{Cl}_{2}$ <br> Do not award missing molar ratio but penalise once only <br> Allow dotted / dashed vertical lines in the cell junction of the cell diagram <br> Allow comma between dichromate ion and proton <br> Allow vertical line between protons and chromium(III) ions <br> Ignore missing / incorrect state symbols <br> Ignore omission of water <br> Ignore missing brackets/use of rounded brackets <br> Penalise inclusion of electrons once only <br> If no other mark is awarded, allow (1) for whole cell diagram written in reverse <br> If no other mark is awarded, allow (1) for electrodes on correct sides but $2 \mathrm{Cl}^{-}$ and $\mathrm{Cl}_{2}$ in reverse order and / or $2 \mathrm{Cr}^{3+}$ and $\mathrm{Cr}_{2} \mathrm{O}_{7}{ }^{2-}+14 \mathrm{H}^{+}$in reverse order <br> Award (1) if $\mathrm{Pt}(\mathrm{s})$ missing both sides but all otherwise correct | (2) |


| Question Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 18(c) | - calculation of $\mathrm{mol}_{\mathrm{MnO}}^{4}-{ }^{-}$and $\mathrm{X}_{2} \mathrm{O}_{5}$ <br> - deduction of mol ratio <br> - final oxidation number of X | Example of calculation: <br> $\mathrm{mol} \mathrm{MnO}_{4}^{-}=\frac{50.0 \times 0.02}{1000}=0.001 / 1.00 \times 10^{-3}$ <br> and <br> $\mathrm{mol} \mathrm{X}_{2} \mathrm{O}_{5}=\frac{25.0 \times 0.1}{1000}=0.0025 / 2.5 \times 10^{-3}$ <br> or <br> $\mathrm{mol} \mathrm{X}=\frac{25.0 \times 0.1 \times 2}{1000}=0.0050 / 5 \times 10^{-3}$ <br> mol ratio $\mathrm{X}: \mathrm{MnO}_{4}^{-}$is $5: 1$ <br> Allow calculation of moles of electrons per Mn and per X giving $5 \times 10^{-3}: 5 \times 10^{-3}$ <br> (there are 5 electrons in the $\mathrm{MnO}_{4}^{-}$half-equation so X 's oxidation number decreased by 1 to <br> (+) 4 <br> Allow $\mathrm{X}^{+4}$ <br> Allow TE of oxidation number (+) 3 from $5: 2$ ratio or from $5 \times 10^{-3} \div 2.5 \times 10^{-3}=2 \text { so }+5-2=(+) 3$ <br> Award (3) for oxidation number (+) 4 provided some working such as number of moles for M1 | (3) |

(Total for Question 18 = 13 marks)


| Question <br> Number | Answer | Additional Guidance | Mark |
| :--- | :--- | :--- | :---: |
| 19(b)(i) | An answer that makes reference to the following point: | (1) |  |
| reaction between two negative ions is slow due to <br> repulsion | Allow negative species for negative ions <br> Allow just 'the negative ions repel' <br> Ignore references to unlikelihood of three negative ions <br> colliding <br> Do not award negative molecules |  |  |


| Question Number | Answer |  | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: | :---: |
| 19(b)(ii) | - ionic equation involving iron(II) <br> - ionic equation involving iron(III) | (1) <br> (1) | $\begin{aligned} & \frac{\text { Examples of ionic equations }}{2 \mathrm{Fe}^{2+}}+2 \mathrm{~S}_{2} \mathrm{O}_{8}{ }^{-2} \xrightarrow{2 \mathrm{Fe}^{3+}}+2 \mathrm{SO}_{4}^{2-} \\ & 2 \mathrm{Fe}^{3+}+2 \mathrm{I}^{-} \rightarrow 2 \mathrm{Fe}^{2+}+\mathrm{I}_{2} \end{aligned}$ <br> Award (1) for balanced equations given in reverse order <br> Allow (1) for two unbalanced equations with all species paired correctly <br> Ignore state symbols even if incorrect | (2) |


| Question Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 19(c)(i) | An explanation that makes reference to the following points: <br> - because it forms one dative (covalent) / co-ordinate bond (to $\mathrm{Fe}^{2+}$ ) <br> - using a lone pair (of electrons) on oxygen | Allow 'a dative/co-ordinate bond' <br> Allow oxygen donates a pair of electrons Ignore water uses a lone pair of electrons | (2) |


| Question <br> Number | Answer | Additional Guidance | Mark |  |
| :--- | :--- | :---: | :--- | :---: |
| 19(c)(ii) | An explanation that makes reference to the following <br> points: <br> - octahedral because there are six pairs of electrons | (1) | Allow this shown on a diagram <br> Allow octahedral because there are 6 coordinate <br> bonds/coordination number is 6 <br> Ignore just octahedral because there are 6 ligands <br> Do not award if bond angle other than $90^{\circ} /$ <br> $90^{\circ}$ and $180^{\circ}$ stated <br> Allow repel/arrange/shape to maximum separation <br> Do not allow repulsion between atoms or water <br> molecules or ligands |  |


| Question Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 19(d) | An explanation that makes reference to the following points: <br> - carbon monoxide replaces / takes the place of the oxygen molecule / ligand <br> - (and it may be toxic) because it binds strongly to the $\mathrm{Fe}^{2+}$ ion | Accept ligand substitution / exchange reaction between oxygen and carbon monoxide COMMENT <br> The question refers to oxygen being carried around and so there needs to be explicit reference and not just implied that to it being replaced/substituted or its place being taken <br> Allow carbon monoxide forms a stronger bond to $\mathrm{Fe}^{2+}$ (than oxygen) <br> Allow carbon monoxide binds (almost) irreversible / permanently to $\mathrm{Fe}^{2+}$ <br> Allow carbon monoxide forms a more stable complex ion with $\mathrm{Fe}^{2+}$ / the complex formed has a larger equilibrium constant <br> Allow prevents / reduces the amount of oxygen being carried to the cells / organs / around the body / blood - scores M2 not M1 <br> Allow just carbon monoxide binds more strongly to haemoglobin/globin | (2) |


| Question <br> Number | Answer |  | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: | :---: |
| 19(e) | An explanation that makes reference to the following points: <br> - there are more particles / moles on the right (of the equation <br> or <br> there is an increase from 3 particles / moles / species on the left of the equation to 5 on the right <br> - so $\Delta S_{\text {system }}$ increases / is positive (and the reaction is thermodynamically feasible) | (1) <br> (1) | Allow species for particles <br> Do not award reference to molecules / atoms /compounds <br> Do not award incorrect numbers <br> Allow $\Delta S_{\text {total }}$ is positive / increasing (and the reaction is thermodynamically feasible) <br> Allow there is an increase in entropy (and the reaction is thermodynamically feasible) <br> Ignore references to increases in disorder | (2) |



|  | Alternative method <br> - (M1) mass of $\mathrm{Cr}_{2} \mathrm{O}_{7}{ }^{2-}$ in $22.55 \mathrm{~cm}^{3}$ <br> - (M2) calculation of $\mathrm{mol} \mathrm{Cr}_{2} \mathrm{O}_{7}{ }^{2-}$ in $22.55 \mathrm{~cm}^{3}$ <br> - (M3)calculation of $\mathrm{mol} \mathrm{Fe}^{2+}$ in $25.0 \mathrm{~cm}^{3}$ <br> - (M4) calculation of mass of $\mathrm{Fe}^{2+}$ in 25.0 $\mathrm{cm}^{3}$ <br> and calculation of total mass of $\left(\mathrm{Fe}^{2+}+\mathrm{Fe}^{3+}\right)$ <br> - (M5) calculation of percentage of $\mathrm{Fe}^{3+}$ | Example of calculation <br> $\operatorname{Mass}\left(\mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}\right)=\frac{2.56 \times 22.55}{1000}=0.057728(\mathrm{~g})$ <br> $\operatorname{Mol}\left(\mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}\right)=\frac{0.057728}{216}=0.00026726 / 2.6726 \times 10^{-4}(\mathrm{~mol})$ <br> $\mathrm{Mol} \mathrm{Fe}{ }^{2+}=0.00026726 \times 6=0.0016036 / 1.6036 \times 10^{-3}(\mathrm{~mol})$ <br> Mass $\mathrm{Fe}^{2+}=0.0016036 \times 55.8=0.089481(\mathrm{~g})$ <br> and <br> Mass $\left(\mathrm{Fe}^{2+}+\mathrm{Fe}^{3+}\right)=\frac{6.28 \times 25.0}{1000}=0.157(\mathrm{~g})$ <br> Mass $\mathrm{Fe}^{3+}=0.157-0.089481=0.067519(\mathrm{~g})$ <br> and <br> $\%$ of $\mathrm{Fe}^{3+}=\frac{0.067519 \times 100}{0.157}=43.0 / 43(\%)$ |
| :---: | :---: | :---: |


| Question Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 20(a)(i) | - equation for the formation of the electrophile <br> - curly arrow within the circle/hexagon to anywhere towards or on $\mathrm{Br}^{+}$ <br> - intermediate structure including charge with horseshoe covering at least 3 carbon atoms and facing the tetrahedral carbon atom and some part of the positive charge must be within the horseshoe <br> - curly arrow from $\mathrm{C}-\mathrm{H}$ bond to anywhere in the hexagon, reforming the delocalised structure | See examples of mechanism on next page <br> $\mathrm{FeBr}_{3}+\mathrm{Br}_{2} \rightarrow \mathrm{Br}^{+}+\mathrm{FeBr}_{4}^{-} /$ <br> $\mathrm{Br}-\mathrm{Br}+\mathrm{FeBr}_{3} \rightarrow \mathrm{Br}^{8+}-\mathrm{Br}^{8-}-{ }_{-}^{--} \mathrm{FeBr}_{3}$ <br> Allow this shown as part of the first step e.g. <br> Allow partial charges on $\mathrm{Br}^{8+}-\mathrm{Br}^{8-}$ <br> Do not award curly arrow starting on or outside the hexagon Do not award missing $+\delta^{+}$on electrophile <br> Do not award curly arrow to a lone pair of electrons on $\mathrm{Br}^{+}$ <br> Do not award dotted bonds to H and Br unless they are part of a 3D structure <br> Ignore missing $\mathrm{H}^{+}$/ involvement of $\mathrm{FeBr}_{4}^{-}$in removal of $\mathrm{H}^{+}$ <br> Ignore reformation of the catalyst even if incorrect | (4) |

Examples of mechanism


Or



| Question <br> Number | Acceptable Answers | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 20(a)(ii)* | This question assesses a 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. <br> The following table shows how the marks should be awarded for structure and lines of reasoning. <br> Comment: <br> Look for the indicative marking points first, then consider the mark for structure of answer and sustained line of reasoning | Guidance on how the mark scheme should be applied: The mark for indicative content should be added to the mark for lines of reasoning. For example, an answer with five indicative marking points 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). <br> If there are no linkages between points, the same five indicative marking points would yield an overall score of 3 marks ( 3 marks for indicative content and no marks for linkages). <br> In general it would be expected that 5 or 6 indicative points would get 2 reasoning marks, and 3 or 4 indicative points would get 1 mark for reasoning, and 0,1 or 2 indicative points would score zero marks for reasoning. <br> General points to note <br> If there is any incorrect chemistry, deduct mark(s) from the reasoning. If no reasoning mark(s) awarded do not deduct mark(s). <br> Accept structures for names throughout <br> If name and formula given both must be correct <br> Deduct a reasoning mark if there is no comparison given for IP1 to IP3 <br> Do not penalise unbalanced / incomplete equations Deduct (mark) from reasoning if any products given are incorrect | (6) |

## Indicative content

- IP1 - Similarity

All are attacked by / react with electrophiles

- IP2 - Types of reaction

Cyclohexene undergoes addition reactions but benzene and/or phenol undergo substitution reactions

## - IP3 - Conditions

Cyclohexene and/or phenol react with (aqueous) bromine / without a catalyst and benzene needs
(a Friedel-Crafts catalyst / iron / iron(III) bromide)

## - IP4 - Benzene

Benzene has delocalised electrons and is (kinetically) stable so the reaction has a high activation energy

## - IP5-Cyclohexene

Cyclohexene has localised electron density in one $\pi$ bond (which increases the electron density and makes it more susceptible to electrophilic attack)

- IP6 - Phenol

Phenol has a lone pair of electrons on the oxygen which is delocalised (within the ring)
and
makes it more susceptible to electrophilic attack

All three need to be mentioned for this IP - evidence for phenol reacting with an electrophile may be seen in IP6

Accept benzene forms monobromo product /
bromobenzene, cyclohexene forms dibromo product / 1,2 dibromocyclohexane and phenol forms tribromo product / 2, 4, 6-tribromophenol
Allow HBr is produced with benzene and phenol but cyclohexene only forms one product

Allow react under normal laboratory conditions / room temperature and pressure
Allow reference to $\mathrm{AlBr}_{3} / \mathrm{AlCl}_{3}$
This IP can be awarded if benzene equation has catalyst and other equation(s) do not
Ignore references to specific temperatures
Allow delocalised ( $\pi$ ) electron ring in benzene is (very) stable
Allow delocalisation of electrons in $\pi$ bonds which decreases the electron density (of the ring) and makes it less susceptible to electrophilic attack

If neither IP4 or IP5 awarded then allow (1) for benzene has delocalised electrons but cyclohexene does not

Allow the lone pair (of electrons) on the oxygen/OH in phenol
and increases the electron density of the (benzene) ring/overlaps with the delocalised ring

| Question Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 20(b) | An answer that makes reference to the following points: <br> - (M1) reagent for step 1 - magnesium and (dry) ether (reacting with bromobenzene) <br> - (M2) first intermediate - phenyl magnesium bromide <br> - (M3) reagent for step 2 - phenyl magnesium bromide with carbon dioxide / $\mathrm{CO}_{2}$ and (followed by hydrolysis with) dilute acid / $\mathrm{H}^{+}$ or methanal and dilute acid $/ \mathrm{H}^{+}$then oxidation <br> - (M4) second intermediate - benzoic acid <br> - (M5) reagent for step 3 - phosphorus(V) chloride / $\mathrm{PCl}_{5}$ <br> - (M6) third intermediate - benzoyl chloride <br> $\bullet(\mathbf{M} 7)$ reagent for step $4-$ ammonia $/ \mathrm{NH}_{3}$ added to an acyl chloride | Allow displayed / structural / skeletal formulae or any combination of these <br> Ignore any references to heat/ incorrect inorganic products <br> Examples of structures of intermediates: <br> Do not award <br> Allow (1) for M3 for the acid hydrolysis of benzonitrile <br> Allow thionyl chloride/ $/ \mathrm{SOCl}_{2}$ <br> Do not award dilute ammonia or ammonia added to benzoic acid <br> M4 to M7 from scheme above can be awarded from benzoic acid however produced | (7) |


| Question Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 20(c)(i) | - repeat unit | Accept skeletal/displayed/structural formulae or combination thereof provided it is correct <br> Example of repeat unit: <br> Accept switching of monomer positions, e.g. <br> Allow amide link to be drawn as $\mathrm{CONH} /-\mathrm{NH}-\mathrm{CO}-$ Allow 'cis' orientation of amide link <br> Ignore bond lengths and bond angles Ignore brackets around repeat unit and $n$ Ignore byproducts such as HCl <br> Do not award additional incomplete repeat units <br> Do not award hydrogen drawn with two single bonds, e.g. $-\mathrm{N}-\mathrm{H}-\mathrm{CO}$ <br> Do not award missing continuation/extension bonds | (1) |


| Question Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 20(c)(ii) | An answer that makes reference to the following point: <br> - because there is hydrogen bonding (and London forces between the chains) in a polyamide <br> - (and this is) stronger than the London forces between the chains in polyalkenes (so more energy is needed to separate the polyamide molecules) <br> or <br> the London forces between the chains in polyalkenes are weaker (than hydrogen bonding so more energy is needed to separate the polyamide molecules) | Reference to breaking of covalent bonds scores (0) Ignore references to (permanent) dipole forces <br> Allow 'it' for the polyamide since it is the subject of the question, so "it has hydrogen bonding" scores M1 <br> Do not award if hydrogen bonding to water stated Do not award if hydrogen bonding shown by $\mathrm{CH}_{2}$ Do not award if ionic bonding or ions referred to <br> Accept dispersion forces / attractions between temporary and induced dipoles for London forces Allow van der Waals' forces for London forces <br> Allow London forces in polyalkenes are easier to overcome (than hydrogen bonding) <br> Note that M2 is awarded for a comparison of the weakness of London forces to the strength of hydrogen bonding. Hence M2 is dependent on M1 or near-miss | (2) |

## Section C

| Question <br> Number | Answer | Additional Guidance |
| :--- | :--- | :--- | :---: |
| 21(a) | 2-hydroxybenzoic acid | Accept 2-hydroxybenzenecarboxylic acid |
|  |  | Allow minor misspellings such as <br> 2-hydroxylbenzenoic acid |
| Ignore missing hyphen or comma instead of |  |  |
| hyphen |  |  |


| $\begin{array}{l}\text { Question } \\ \text { Number }\end{array}$ | Answer | Additional Guidance |
| :--- | :--- | :--- |
| 21(b)(i) | • carboxylic acid and ester and benzene / arene | $\begin{array}{l}\text { (1) } \\ \\ \end{array}$ | \(\left.\begin{array}{l}Allow just 'carboxyl' for carboxylic acid names given in any order <br>

Allow just 'carboxylic' <br>
Allow phenyl for benzene/arene <br>
Allow aromatic ring for benzene/arene <br>

Ignore formulae of groups\end{array}\right]\) Do not award phenol | Do not award carbonyl |
| :--- |



| Question <br> Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 21(b)(iii) | - calculation of amount of salicylic acid <br> - calculation of theoretical mass of acetyl salicylic acid <br> - calculation of actual mass of acetyl salicylic acid | Example of calculation: mol salicylic acid used $=\frac{2.00}{138}=0.014493(\mathrm{~mol})$ <br> theoretical mass of acetyl salicylic acid $=0.014493 \times 180$ $=2.6087(\mathrm{~g})$ <br> TE on M1 <br> actual mass of acetyl salicylic acid $=\frac{2.6087 \times 74.8}{100}=1.9513(\mathrm{~g})$ <br> TE on M2 provided answer $\leq 5.00(\mathrm{~g})$ <br> OR <br> mass salicylic acid converted $=2.00 \times 0.748=1.496(\mathrm{~g})(1)$ <br> mol salicylic acid converted $=\frac{1.496}{138}=0.01084(\mathrm{~mol})(1)$ <br> mass acetyl salicylic acid formed $=0.01084 \times 180$ $=1.9513(\mathrm{~g})$ <br> Ignore SF except 1 SF <br> Correct answer scores without working scores (3) | (3) |


| Question <br> Number | Answer | Additional Guidance | (1) |
| :--- | :--- | :--- | :---: | :---: |
| 21(c)(i) | • completed equation | Example of equation: |  |


| Question Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 21(c)(ii) | An explanation that makes reference to the following points: <br> - acetylsalicylic acid will dissociate less in acidic solution or acetylsalicylic acid dissociate more in alkaline solution <br> - because the additional $\mathrm{H}^{+} / \mathrm{H}_{3} \mathrm{O}^{+}$ions in the acid will shift the equilibrium position to the left <br> - and $\mathrm{OH}^{-}$/ hydroxide ions in the alkali will react with the $\mathrm{H}^{+}$ions and shift the equilibrium position to the right | Penalise reference to change in $K_{\mathrm{a}}$ once only <br> Allow reference to the stomach for 'acidic solution' <br> Allow reference to small intestine for 'alkaline' If both stated then both must be correct <br> Allow the backward reaction is favoured by the additional/higher $\mathrm{H}^{+} / \mathrm{H}_{3} \mathrm{O}^{+}$ions in the acid <br> Accept $\mathrm{H}^{+}+\mathrm{OH}^{-} \rightarrow \mathrm{H}_{2} \mathrm{O}$ <br> and this shifts the equilibrium position to the right <br> Allow $-\mathrm{COOH}+\mathrm{OH}^{-} \rightarrow-\mathrm{COO}^{-}+\mathrm{H}_{2} \mathrm{O}$ <br> Or carboxylic acid group reacts with /neutralises the $\mathrm{OH}^{-}$ <br> and this shifts the equilibrium position to the right/ favours the forward reaction | (3) |


| Question <br> Number | Answer | Additional Guidance | Mark |
| :--- | :--- | :--- | :---: |
| 21(d) | • methanol $/ \mathrm{CH}_{3} \mathrm{OH}$ | Allow displayed formula / combination of structural and displayed formula <br> If name and formula are given then both must be correct <br> Allow methyl alcohol <br> Ignore reference to acid catalyst/ $\mathrm{H}_{2} \mathrm{SO}_{4} / \mathrm{HCl} /$ heat | (1) |
|  |  | Do not award methanal <br> Do not award $\mathrm{CH}_{4} \mathrm{O}$ |  |


| Question Number | Answer |  | Additional Guidance |  |  |  |  | Mark |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 21(e) | - chemical shift ranges for OH and $\mathrm{CH}_{3}$ in acetylsalicylic acid <br> - chemical shift ranges for OH and $\mathrm{CH}_{3}$ in methyl salicylate | (1) <br> (1) | Example of table: |  |  |  |  | (2) |
|  |  |  |  | Acetyls | cylic acid | Methyl | alicylate |  |
|  |  |  | $\begin{array}{\|l} \hline \text { Type of } \\ \text { proton } \end{array}$ | OH | $\mathrm{CH}_{3}$ | OH | $\mathrm{CH}_{3}$ |  |
|  |  |  | $\begin{aligned} & \hline \text { Chemical } \\ & \text { shift / } \\ & \text { ppm } \\ & \hline \end{aligned}$ | $\begin{gathered} 10.0- \\ 12.0 \end{gathered}$ | 1.6-2.8 | 3.8-7.6 | $2.8-4.3$ |  |
|  |  |  | Allow ranges in reverse order e.g. $12.0-10.0$ <br> Allow any range within these ranges $11.8-10.2$ <br> If no other mark is awarded, allow (1) for any two correct ranges <br> If no other mark awarded, allow (1) for any three single values within the correct ranges or two single values with one acceptable range |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |


| Question Number | Answer | Additional Guidance | Mark |
| :---: | :---: | :---: | :---: |
| 21(f) | - (M1) calculation of mol NaOH added at start <br> - (M2) calculation of mol HCl used in titration <br> - (M3) calculation of mol NaOH remaining in $250 \mathrm{~cm}^{3}$ <br> - (M4) calculation of mol acetylsalicylic acid reacted <br> - (M5) calculation of acetylsalicylic acid mass <br> - (M6) calculation of percentage of acetylsalicylic acid and deduction of Brand of tablet | Example of calculation: <br> $\mathrm{mol} \mathrm{NaOH}=\frac{25.0 \times 1.00}{1000}=0.025 / 2.5 \times 10^{-2}(\mathrm{~mol})$ <br> $\mathrm{mol} \mathrm{HCl}=\frac{16.95 \times 0.100}{1000}=0.001695 / 1.695 \times 10^{-3}(\mathrm{~mol})$ <br> ( mol NaOH remaining in $25.0 \mathrm{~cm}^{3}=0.001695 / 1.695 \times 10^{-3}(\mathrm{~mol})$ ) mol NaOH remaining in $250 \mathrm{~cm}^{3}=0.01695 / 1.695 \times 10^{-2}(\mathrm{~mol})$ <br> $\mathrm{mol} \mathrm{NaOH}=0.025-0.01695=0.00805 / 8.05 \times 10^{-3}(\mathrm{~mol})$ mol acetylsalicylic acid $=\frac{0.00805}{2}=0.004025$ <br> mass acetylsalicylic acid $=0.004025 \times 180=0.7245(\mathrm{~g})$ <br> percentage of acetylsalicylic acid $=\frac{0.7245 \times 100}{0.760}$ $=95.329(\%)$ <br> and Brand B <br> Allow TE at each stage <br> Brand / percentage with no working scores (0) <br> Ignore SF except 1 SF in the final mass calculated Ignore incorrect intermediate units <br> Do not credit a division of moles by 2 if carried out before the subtraction | (6) |

