

Mark Scheme Results)

Pearson Edexcel Advanced Level In Chemistry (9CH0) Paper 01 Advanced Inorganic and Physical Chemistry

Edexcel and BTEC Qualifications

Edexcel and BTEC qualifications are awarded by Pearson, the UK's largest awarding body. We provide a wide range of qualifications including academic, vocational, occupational and specific programmes for employers. For further information visit our qualifications websites at www.edexcel.com or www.edexcel.com, you can get in touch with us using the details on our contact us page at www.edexcel.com/contactus.

Pearson: helping people progress, everywhere

Pearson aspires to be the world's leading learning company. Our aim is to help everyone progress in their lives through education. We believe in every kind of learning, for all kinds of people, wherever they are in the world. We've been involved in education for over 150 years, and by working across 70 countries, in 100 languages, we have built an international reputation for our commitment to high standards and raising achievement through innovation in education. Find out more about how we can help you and your students at: www.pearson.com/uk

Summer 2019
Publications Code 9CH0_01_1906_MS
All the material in this publication is copyright
© Pearson Education Ltd 2019

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.
- Mark schemes will indicate within the table where, and which strands of QWC, are being assessed. The strands are as follows:
 - i) ensure that text is legible and that spelling, punctuation and grammar are accurate so that meaning is clear
 - ii) select and use a form and style of writing appropriate to purpose and to complex subject matter
 - iii) organise information clearly and coherently, using specialist vocabulary when appropriate

Using the Mark Scheme

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

The mark scheme gives examiners:

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

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

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

Phrases/words in **bold** indicate that the <u>meaning</u> of the phrase or the actual word is **essential** to the answer.

ecf/TE/cq (error carried forward) means that a wrong answer given in an earlier part of a question is used correctly in answer to a later part of the same question.

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

Quality of Written Communication

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

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

Full marks will be awarded if the candidate has demonstrated the above abilities. Questions where QWC is likely to be particularly important are indicated (QWC) in the mark scheme, but this does not preclude others.

| Question Number | Answer | |
|--------------------|---|-----|
| 1(a) | The only correct answer is A | (1) |
| | B is not correct because oxygen atoms do not have this number of protons | |
| | C is not correct because oxygen atoms do not have this number of protons | |
| | D is not correct because oxygen atoms do not have this number of protons | |

| Question Number | Answer | |
|--------------------|---|-----|
| 1(b) | The only correct answer is D | (1) |
| | A is not correct because there are no peaks for bromine molecules | |
| | B is not correct because the peak for the ⁷⁹ Br- ⁸¹ Br⁺ molecular ion is missing | |
| | \boldsymbol{C} is not correct because the peak for the ⁷⁹ Br- ⁸¹ Br ⁺ molecular ion is not twice the height of the other molecular ion peaks | |

| Question Number | Answer | |
|--------------------|---|--|
| 1(c) | ne only correct answer is C | |
| | A is not correct because this is only the number of electrons in the 3p orbitals of the chlorine atom | |
| | B is not correct because this is only the number of electrons in the 3p orbitals of the chloride ion | |
| | D is not correct because this is the total number of electrons in the chloride ion, not just those in the p orbitals | |

| Question Number | Answer | |
|--------------------|--|--|
| 1(d) | he only correct answer is A | |
| | B is not correct because oxygen has more protons to exert an attractive force to reduce the ionic radius | |
| | C is not correct because sodium has more protons to exert an attractive force to reduce the ionic radius | |
| | D is not correct because aluminium has more protons to exert an attractive force to reduce the ionic radius | |

(Total Question 1 = 4 marks)

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|------------|---|------|
| 2(a)(i) | • equation | Example of equation | (1) |
| | equation | $Cl_2 + 2KBr \rightarrow Br_2 + 2KCl$ | |
| | | Accept ionic equation $Cl_2 + 2Br^- \rightarrow Br_2 + 2Cl^-$ | |
| | | Allow multiples Ignore state symbols even if incorrect | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|--|------|
| 2(a)(ii) | An answer that makes reference to the following points: | | (3) |
| | (M1) the precipitate is a mixture of silver chloride and silver bromide or not all of the bromide ions were oxidised (1) | Allow Some bromide ions are still present/bromide ions were in excess/both chloride and bromide ions are present | |
| | • (M2) silver chloride/AgCl dissolves in dilute ammonia (1) | Do not award references to Cl ⁻ dissolving | |
| | (M3) silver bromide/AgBr does not dissolve in dilute ammonia (1) | Silver bromide only dissolves in concentrated ammonia | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|--|------|
| 2(a)(iii) | An answer that makes reference to the following point: | | (1) |
| | iodine is a weaker oxidising agent than chlorine or iodine cannot oxidise bromide ions or | Accept reverse arguments | |
| | iodine is a stronger reducing agent | Ignore Just references to reactivity/displacement e.g. iodine is less reactive/cannot displace | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---------------------------------------|---|------|
| 2(b)(i) | | Example of equation | (1) |
| | balanced equation | $3Cl_2 + 6OH^- \rightarrow 5Cl^- + ClO_3^- + 3H_2O$ | |
| | | Allow multiples | |
| | | Ignore state symbols even if incorrect | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|--|------|
| 2(b)(ii) | An explanation that makes reference to the following points: | Ignore general definitions of disproportionation | (2) |
| Clip with | | | |
| 2(b)(i) | oxidation number for chlorine changes from 0 to -1 so it is | Accept oxidation numbers and their changes | |
| | reduced (1) | shown with equation | |
| | oxidation number for chlorine changes from 0 to +5 | | |
| | so it is oxidised (1) | Allow 1 out of 2 marks for three correct | |
| | | oxidation numbers of the chlorine | |

(Total Question 2 = 8 marks)

| Question Number | Answer | |
|--------------------|--|-----|
| 3(a)(i) | The only correct answer is D | (1) |
| | A is not correct because the measurement uncertainty is doubled as there are two burette readings | |
| | B is not correct because this gives the largest measurement uncertainty | |
| | C is not correct because the measurement uncertainty is doubled as the pipette is used twice | |

| Question Number | Answer | | Additional Guidance | Mark |
|--------------------|--|-------------------|--|------|
| 3(a)(ii) | | | Example of calculation | (3) |
| | • calculation of Q | (1) | Q=(100 x 4.18 x 6.5=) = 2717 (J) / 2.717 kJ | |
| | calculation of enthalpy change | (1) | ΔH= 2.717 ÷ 0.05= (-)54.340 | |
| | answer to nearest whole number and with | negative sign (1) | $= -54 \text{ (kJ mol}^{-1}\text{)}$ | |
| | | | Allow TE at each stage Correct answer with no working scores (3) | |

| Question Number | Answer | |
|--------------------|---|-----|
| 3(b)(i) | The only correct answer is B | (1) |
| | A is not correct because there is no extrapolation to the largest temperature increase carried out | |
| | C is not correct because the extrapolation is at the wrong time | |
| | D is not correct because the extrapolation extends beyond the time of addition of alkali | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|--|------|
| 3(b)(ii) | An explanation that makes reference to | | (2) |
| | ethanoic acid is a weak(er) acid / only partially ionised/dissociated (1) | Allow hydrochloric acid is a strong(er) acid/fully ionised | |
| | • (some) energy is used to fully/completely ionise the ethanoic acid (1) | Do not award 'more NaOH will react so more energy given off' | |

(Total Question 3 = 7 marks)

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|------------------------------|------|
| 4(a)(i) | An answer that makes reference to the following point: | | (1) |
| | (Identity of gas is) oxygen/O₂ and | Do not award just 'O' | |
| | (test result is that the splint) relights | Allow 'rekindle'/'reignites' | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|---|------|
| 4(a)(ii) | An answer that makes reference to the following point: | If name and formula given then both must be correct | (1) |
| | (Identity of gas is) nitrogen dioxide and | Allow NO ₂ | |
| | (appearance is) brown (gas/fumes) | Do not award NO or N ₂ O ₄ Do not award if liquid referred to Do not award if two gases are given unless one of the gases stated is colourless oxygen | |
| | | Ignore shades of colour | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|------------|--|------|
| 4(a)(iii) | | Example of equation | (1) |
| | • equation | 2NaNO ₃ → 2NaNO ₂ + O ₂ Accept multiples | |
| | | Ignore state symbols even if incorrect | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|--|------|
| 4(a)(iv) | An description that makes reference to following points: | Allow annotated diagrams to illustrate the marking points | (2) |
| | use of a delivery tube to bubble gas into limewater (1) | Example of a diagram that could be given credit solid be limewater Do not award if the apparatus setting would not be feasible such as missing cork/bung or gaps around delivery tube horizontal tube with limewater in bung in the test tube with limewater | |
| | compare the time taken for the limewater to go cloudy (1) | Allow TE from an incorrect/unsuitable method Allow Any length of time, e.g. 10 minutes | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|--|------|
| 4(b) | An explanation that makes reference to | Accept reverse arguments Penalise omission of ion once only | (3) |
| | the magnesium ion/cation is smaller (than the barium ion/cation) (1) | Allow magnesium ion/cation has a higher charge density Do not award reference to just atom | |
| | which polarises the (large) carbonate (ion)/anion (1) | Accept 'which distorts the carbonate electron cloud' Do not award if molecules of carbonate are referred to | |
| | and weakens the carbon-oxygen bond (C-O/C=O) bonds (1) | Allow 'weakens the bonds in the carbonate ion' Allow 'less energy needed to break the C-O bond' Do not award if the bonds referred to are the ionic bonds between the metal ion and the carbonate ion | |

(Total Question 4 = 8 marks)

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|---|------|
| 5(a)(i) | An answer that makes reference to the following point: | | (1) |
| | the two negative ions repel each other | Reference to both charge and repulsion needed | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|---|------|
| 5(a)(ii) | An explanation that makes reference to the following points: | Example of equations | (3) |
| | • equation with oxidation of iron(II) ions (1) | $2Fe^{2+} + S_2O_8^{2-} \rightarrow 2Fe^{3+} + 2SO_4^{2-}$ | |
| | • equation with reduction of iron(III) ions (1) | $2Fe^{3+} + 2I^{-} \rightarrow 2Fe^{2+} + I_{2}$ | |
| | | Allow multiples Ignore state symbols even if incorrect | |
| | | Allow one mark if the two correct equation are given in the wrong order | |
| | (catalysis is possible because) variable oxidation state/iron has more than one oxidation state/number or | Allow reference to iron being oxidised and reduced | |
| | both steps now involve oppositely charged ions (1) | Allow reference to the iron ions being positive and so not repelled | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|--|------|
| 5(b) | | Example of equations | (2) |
| | equation with oxidation of cobalt(II) ions | $2\text{Co}^{2+} + \frac{1}{2}\text{O}_2 + 2\text{H}^+ \rightarrow 2\text{Co}^{3+} + \text{H}_2\text{O}$ | |
| | equation with reduction of cobalt(III) ions | $2\text{Co}^{3+} + \text{SO}_3^{2-} + \text{H}_2\text{O} \rightarrow 2\text{Co}^{2+} + \text{SO}_4^{2-} + 2\text{H}^+$ | |
| | | Allow multiples Ignore state symbols even if incorrect | |

| Question Number | Answer | Mark |
|--------------------|---|------|
| 5(c) | The only correct answer is C | (1) |
| | A is not correct because only Mn^{2+} is an autocatalyst for this reaction B is not correct because only Mn^{2+} is an autocatalyst for this reaction D is not correct because only Mn^{2+} is an autocatalyst for this reaction | |

| Question Number | Answer | Additional Guidance Mark |
|--------------------|--|--|
| 5(d) | Tungsten • (because) adsorption is too strong and so desorption would be too slow Ignore ref | ferences to oxidation or reactivity series (1) |
| | Do not aw | vard 'a b sorption' |

(Total Question 5 = 8 marks)

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|--|------|
| 6(a) | An answer that makes reference to the following points: (M1) (a lower boiling temperature is expected) because water has fewer electrons than hydrogen sulfide (1) | Accept water has 10 electrons but hydrogen sulfide has 18 electrons (per molecule) Ignore reference to Mr/size of atom | (4) |
| | (M2) water has weaker/less London forces (1) (M3) (a higher boiling temperature occurs because) water has hydrogen bonding (1) | Allow van der Waals'/dispersion forces/ instantaneous dipole-induced dipole | |
| | (M4) hydrogen bonding is stronger than London forces and requires more energy to break (and results in a higher boiling temperature) (1) | Accept reverse arguments Ignore references to permanent dipole-dipole forces | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|--|------|
| 6(b) | An explanation that makes reference to the following points: | Accept supporting diagrams which illustrate the following: | (4) |
| | (M1) oxygen is more electronegative than hydrogen and carbon (1) | Accept electronegativity values stated Allow oxygen has a greater force of attraction for the bonded electron pairs than hydrogen or carbon | |
| | (M2) which results in a polar bond with oxygen δ- so carbon and hydrogen δ+ (1) | Penalise the failure to refer to carbon and hydrogen once only | |
| | (M3) carbon dioxide is a symmetrical/linear molecule and so the dipole moments/vectors cancel (1) | Allow 'symmetrical so dipoles/ polar bonds cancel' | |
| | (M4) the lone pairs of electrons of oxygen/ the V-shape of the water molecule mean that the dipole moments/ vectors do not cancel (1) | Allow angular/bent for V-shape | |
| | | Penalise reference to just 'charges' once only in M3 and M4 if dipoles not stated or shown in the answer | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|-----------------------|---|------|
| 6(c) | Two diagrams | Examples of suitable diagrams | (2) |
| | • oxonium ion (1) | $\begin{bmatrix} H & \times & \times \\ & H & \\ & & & \end{bmatrix}^{\dagger}$ | |
| | hydroxide ion (1) | H * Ö × | |
| | | Ignore inner 'shell' of two electrons | |
| | | Ignore missing brackets | |
| | | Ignore covalent circles | |
| | | Penalise all dots or all crosses or use of other | |
| | | symbols for electrons once only | |
| | | Penalise omission of charges once only | |

| Question Number | Answer | | Additional Guidance | Mark |
|--------------------|---|-----|---|------|
| 6(d)(i) | | | Example of calculation: | (2) |
| | • square root of K_w at 310K to get [H ⁺] | (1) | $[H^+] = (\sqrt{2.40} \times 10^{-14}) = 1.549x10^{-7} \text{ (mol dm}^{-3})$ | |
| | • calculation of pH to 2 decimal places | (1) | pH = $(-\log 1.549x10^{-7})$ = (6.809894379) = 6.81 Correct answer with no working scores (2) | |
| | | | Allow TE from incorrect [H ⁺] as long as answer is in the pH range 6.00 – 7.00 inclusive | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|--|------|
| 6(d)(ii) | An answer that makes reference to the following points: • (M1) neutral (1) | Acidic or alkaline scores (0) | (2) |
| | • (M2) because [H ⁺ (aq)] = [OH ⁻ (aq)] /equal amounts of H ⁺ and OH ⁻ ions (1) | Allow both [H ⁺] and [OH ⁻] have increased equally (from 298 K to 310 K) M2 dependent on M1 | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|---|------|
| 6(d)(iii) | An answer that makes reference to the following point: • positive / + sign because K _w increases as the | | (1) |
| | temperature increases | Allow 'positive because' | |
| | | bond breaking requires energy | |
| | | or | |
| | | equilibrium shifts to the right | |
| | | or | |
| | | there is greater/more ionisation/dissociation | |
| | | Ignore 'endothermic' | |

(Total Question 6 = 15 marks)

| Question Number | Ansı | wer | Additional Guidance | Mark |
|--------------------|--|--|---|------|
| *7 | and logically structured answer with linkages and fully sustained reasoning. Marks are awarded for indicative content and for how the answer is structured and shows lines of reasoning. The following table shows how the marks should be awarded | | Guidance on how the mark scheme should be applied: The mark for indicative content should be added to the mark for lines of reasoning. For example, a response with four 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 | (6) |
| | Number of indicative I marking points seen in | Number of marks awarded for indicative marking points 4 3 2 1 0 | structure and some linkages and lines of reasoning). If there were no linkages between the points, then the same indicative marking points would yield an overall score of 3 marks (3 marks for indicative content and zero marks for linkages). | |
| | Answer shows a coherent logical structure with linkages and fully sustained lines of reasoning demonstrated throughout | | If there is any incorrect chemistry, deduct mark(s) from the reasoning. If no reasoning mark(s) awarded do not deduct mark(s). More than one indicative marking point may be made within the same comment or explanation | |

| Ansv | ver is partially structured | 1 | Deduct a reasoning mark if no comparison | |
|--------|--|------------------------------|--|--|
| with | some linkages and lines of | | made | |
| reaso | oning | | | |
| Ansv | ver has no linkages | 0 | Penalise the use of 'atom' instead of ion | |
| betw | een points and is | | once only against any indicative point | |
| unst | ructured | | Ignore incorrect colours | |
| Indica | ative content | | | |
| Simila | rities | | | |
| • | (IP1) the differences in ene | rgy levels determines the | This can be mentioned separately or as a | |
| | colour of the flame test and | d complex ion | comparison | |
| Differ | ences | | | |
| Flame | test | | | |
| • | (IP2) heat (energy) results | in electron promotion | Allow electrons excited by heat | |
| • | (IP3) return of an (excited) state | electron to a lower (energy) | Allow electron is 'de-excited' to a lower (energy) state Do not award if d-d transitions stated | |
| Comp | lex ion | | | |
| Comp | (IP4) d orbital s are split (in | energy by the ligands) | Allow d subshell splitting | |
| | (ii 4) a orbitals are split (iii | chergy by the ligarias) | Do not award singular "d orbital" splitting | |
| | (IP5) light (energy) is need | ed for electron promotion | | |
| • | (IF3) light (energy) is need | ed for electron promotion | Accept "The colour seen is complimentary | |
| | (IP6) the colour not absorb | ed is the colour seen | to that absorbed" | |
| | (ii e) the colour flot absorb | ca is the colour seen | Allow 'colour reflected is the colour seen' | |
| | | | Do not award if colour attributed to 'fall' of | |
| | | | electron to lower energy d orbital | |
| | | | Do not award 'emission of light' | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|---|------|
| 8(a)(i) | An answer that makes reference to the following points: | Allow corrections to be made on the diagram | (2) |
| | • identification and correction of the first error (1) | Error 1 – arrow for enthalpy change of formation should go down/be reversed | |
| | • identification and correction of the second error (1) | Error 2 – the word 'half' should be deleted from the enthalpy change of atomisation of hydrogen | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|---|------|
| 8(a)(ii) | | Example of calculation | (1) |
| | calculation of first electron affinity of hydrogen | 1^{st} EA= -(218+496+107)-56 +804 = -73 (kJ mol ⁻¹) | |
| | | Allow a TE | |
| | | 1 st EA = +39(kJ mol ⁻¹) if the first arrow reversed direction is not identified | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|--|------|
| | Penalise incorrect or missing unit | s in (b)(i) and (b)(ii) once only | |
| 8(b)(i) | • calculation of Δ <i>G</i> (1) | Example of calculation $\Delta G = -56 - (298 \times -76.5)$ 1000 =-33.203 (kJ mol ⁻¹) or $\Delta G = -56000 - (298 \times -76.5)$ = -33203 (J mol ⁻¹) | (2) |
| | • ΔG is negative/ <0 and so reaction is feasible (1) | Ignore SF except 1 Allow ≤0 and so reaction is feasible Standalone mark Allow TE on own ΔG calculated value | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---------------------------|---|------|
| 8(b)(ii) | | Example of calculation | (1) |
| | • calculation of <i>T</i> | $\Delta G = 0$, so $\Delta H = T\Delta S_{(system)}$ or $T = \Delta H/\Delta S_{(system)}$ | |
| | | T = 56/0.0765 = 732 K | |
| | | or | |
| | | T = 56000/76.5 = 732 K | |
| | | or | |
| | | T = 459°C | |
| | | Ignore SF except 1 SF | |
| | | Do not award -732K | |
| | | TE on incorrect values penalised already in (b)(i) | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|---|------|
| 8(c) | | Example of half equation | (1) |
| | reduction half-equation | $12O_2 + H_2O + 2e^- \rightarrow 2OH^-$ | |
| | | Allow multiples | |
| | | Ignore state symbols even if incorrect | |

| Question Number | Answer | Mark |
|--------------------|--|------|
| 8(d) | The only correct answer is B (-804, -711, -2718) | (1) |
| | A is not correct because the lattice energy of magnesium hydride is not exothermic enough | |
| | C is not correct because the lattice energy of potassium hydride should be less exothermic than sodium hydride and also that the lattice energy of magnesium hydride is not exothermic enough | |
| | D is not correct because the lattice energy of potassium hydride should be less exothermic than sodium hydride | |

(Total Question 8 = 8 marks)

| Question Number | Answer | Mark |
|--------------------|---|------|
| | The only correct answer is D (resists changes in pH if small quantities of acid or base are added) | (1) |
| | A is not correct because buffer solutions can be alkaline or acidic | |
| 9(a) | B is not correct because buffers do not always contain equal numbers of moles of the acid and its conjugate base | |
| | C is not correct because a buffer does not prevent any change in pH | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|---|------|
| 9(b) | • (M1) calculation of [H ⁺] (1) | Example of calculation $[H^+] = 10^{-3.9} = 1.2589x 10^{-4} \text{ (mol dm}^{-3}\text{)}$ | (5) |
| | • (M2) rearrangement of K_a and calculation of ethanoate concentration (1) | $[CH_3COO^-] = (1.74 \times 10^{-5} \times 0.800) = 1.2589 \times 10^{-4}$ | |
| | | =0.11057 (mol dm ⁻³) | |
| | • (M3) calculation of the number of moles of ethanoate in the buffer volume (1) | n=(0.11057 x 0.05 =) 5.5285x 10 ⁻³ (mol) | |
| | • (M4) calculation of the mass of sodium ethanoate in the buffer volume (1) | m=(5.5285x 10 ⁻³ x 82=) 0.453339 | |
| | • (M5) answer to 2/3 SF (1) | m=0.45/ 0.453 (g) Award this mark only if there has been some attempt at calculation using an $M_{\rm r}$ | |
| | | TE at each stage | |
| | | If Henderson-Hasselbalch equation used (M1) for calculation of $pK_a = 4.759$ (M2) for rearrangement and calculation of | |
| | | ethanoate concentration Remaining marking points as above | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|--|------|
| 9(c) | An answer that makes reference to the following points: | | (3) |
| | carbon dioxide dissolved in the blood forms carbonic acid (and so this concentration increases) | Can be shown in an equation Do not award 'CO ₂ reacts with H ⁺ to form carbonic acid' | |
| | the equilibrium will shift to the right and produces more H⁺/acid ions (1) | Allow Carbonic acid (partially) dissociates to produce H ⁺ Do not award 'CO ₂ reacts with H ⁺ so equilibrium shifts to the right to produce more H ⁺ ' | |
| | the (high) concentration of hydrogencarbonate ions suppress the ionisation of carbonic acid (to help to control pH) or the (large) reservoir/excess of hydrogencarbonate ions combine with the H⁺ ions (to help to control blood pH) (1) | | |
| | | Ignore general comments about the effects of adding acid and/or alkali to a buffer which do not relate to carbon dioxide | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|--|------|
| 9(d)(i) | | PH 7 25 40 Volume of sodium hydroxide added / cm³ | (4) |
| | • general shape of weak acid-strong base curve (1) | Ignore missing initial rise in pH Vertical part must cover 3-5 pH units between 6 – 11 The curve should reach pH~12-13 by 10cm ³ after vertical section | |
| | • curve starts at pH of 3.0 and ends at pH of 12-13 (1) | The curve must start at zero and continue to 100 cm ³ | |
| | • vertical part of the curve at 40 cm ³ (1) | | |
| | • labelling of area indicating buffer action (1) | Allow buffering area to be labelled anywhere between ~5 and 35 cm ³ inclusive | |

| Question Number | Answer | | Additional Guidance | Mark |
|--------------------|--|--------------------------------------|---|------|
| 9(d)(ii) | An answer that makes reference to | | Standalone marks | (2) |
| ζ(α)(ιι) | determine the pH at the point when neutralised | en half of the acid is (1) | Allow 'pH at half-equivalence point' Allow 'pH at half neutralisation point' | |
| | • $K_a = 10^{-pH}$ / $K_a = 10^{-pK}$ | (1) | Accept description in words such as inverse log of minus pH or value is pK_a and so inverse log of minus value gives K_a Allow $pK_a = -\log K_a$ | |
| | | | Ignore any calculation | |

(Total Question 9 = 15 marks)

| Question Number | Answer | | Additional Guidance | Mark |
|--------------------|--|-----|--|------|
| 10(a)(i) | • (M1) calculation of mole fractions | (1) | Example of calculation $X_{SO_2} = 0.0160 \div 0.8 = 0.02(0)$ $X_{O_2} = 0.0120 \div 0.8 = 0.015$ | (5) |
| | | | $X_{SO_3} = 0.772 \div 0.8 = 0.965$ | |
| | • (M2) calculation of partial pressures | (1) | $P_{SO_2} = 0.02(0) \times 2.40 = 0.048$ | |
| | | | $P_{\rm O_2} = 0.015 \times 2.40 = 0.036$ | |
| | | | $P_{SO_3} = 0.965 \times 2.40 = 2.316$ | |
| | • (M3) expression of K_p | (1) | $K_{p} = \frac{(\boldsymbol{P}_{SO_3})^2}{(\boldsymbol{P}_{SO_2})^2 \times \boldsymbol{P}_{O_2}}$ | |
| | • (M4) calculation of value of K_p | (1) | Do not award square brackets $K_p = \frac{2.316^2}{0.048^2 \times 0.036} = \frac{2.316^2}{0.048^2 \times 0.036}$ | |
| | | | K_p = 64668.4/6.46684 x 10 ⁴ K_p = 65000/6.5 x 10 ⁴ /64700/6.47 x 10 ⁴ Ignore SF except 1 | |
| | • (M5) units | (1) | atm ⁻¹ | |
| | | | Correct final answer without working scores (5) | |

| Answer | Additional Guidance | Mark |
|--|--|--|
| calculation of the number of molecules | Example of calculation $N=(n \times L = 0.0160 \times 6.02 \times 10^{23})$ | (1) |
| | $= 9.632 \times 10^{21}$ | |
| | Ignore SF except 1SF | |
| | | • calculation of the number of molecules |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|--|--|------|
| 10(a)(iii) | An answer that makes reference to the following points: | Standalone marks | (2) |
| | • to ensure that K_p stays the same/ quotient stays the same or only temperature changes the value of K_p (1) | Allow concentration changes have no effect on the value of K_p | |
| | the number of (sulfur dioxide) molecules decreases Either because the equilibrium shifts to the right or because one of the denominators (oxygen) has increased so the other denominator (sulfur dioxide) has to decrease (1) | Allow 'moles' for molecules | |

| Question Number | Answer | Additional Guidance | Mark |
|--------------------|---|---|------|
| 10(b) | An explanation that makes reference to the following points: | 'Equilibrium moves to the right' scores (0) | (2) |
| | • equilibrium position shifts to the left (1) | | |
| | (because) the hydroxide ions combine with/neutralise the H⁺ ions to remove them from the equilibrium (1) | Allow Hydroxide ions react with H ⁺ ions to make water/ hydroxide ions react with H ⁺ ions to reduce their number/concentration | |
| | | Ignore reference to 'more products formed' | |

| Question Number | Answer | Mark |
|--------------------|--|------|
| | The only correct answer is D $K_c = [H_2O]^4$ | (1) |
| | $[H_2]^4$ | |
| 10(c) | A is not correct because the solids should not be included in the expression and the powers of the remaining substituents have been omitted | |
| | B is not correct because the solids should not be included in the expression | |
| | C is not correct because the powers of the substituents have been omitted | |