Section A

| Question | Marking guidance | Mark | AO | Comments |
|----------|--|------|--------|--|
| 01.1 | A mixture of liquids is heated to boiling point for a prolonged time | 1 | AO1b | |
| | Vapour is formed which escapes from the liquid mixture, is changed back into liquid and returned to the liquid mixture | 1 | AO1b | |
| | Any ethanal and ethanol that initially evaporates can then be oxidised | 1 | AO2g | |
| 01.2 | $CH_3CH_2OH + H_2O \longrightarrow CH_3COOH + 4H^+ + 4e^-$ | 1 | AO2d | |
| 01.3 | Mixture heated in a suitable flask / container | 1 | AO3 2a | A labelled sketch illustrating these points scores the |
| | With still head containing a thermometer | 1 | AO3 2a | marks |
| | Water cooled condenser connected to the still head and suitable <u>cooled</u> collecting vessel | 1 | AO3 2a | |
| | Collect sample at the boiling point of ethanal | 1 | AO3 2a | |
| | Cooled collection vessel necessary to reduce evaporation of ethanal | 1 | AO3 2a | |
| 01.4 | Hydrogen bonding in ethanol and ethanoic acid or no hydrogen bonding in ethanal | 1 | AO1a | |
| | Intermolecular forces / dipole-dipole are weaker than hydrogen bonding | 1 | AO1a | |

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| 01.5 | Reagent to confirm the presence of ethanal: | | | |
|------|--|---|-------|--|
| | Add Tollens' reagent / ammoniacal silver nitrate / aqueous silver nitrate followed by 1 drop of aqueous sodium hydroxide, then enough aqueous ammonia to dissolve the precipitate formed | 1 | AO1b | |
| | OR | | | |
| | Add Fehling's solution | | | |
| | | | 4.04h | MO and MO approximate diff. M4 is given |
| | Warm | 1 | AO1b | M2 and M3 can only be awarded if M1 is given correctly |
| | | | | |
| | Result with Tollen's reagent: | | | |
| | Silver mirror / black precipitate | 1 | AO1b | |
| | OR | | | |
| | Result with Fehling's solution: | | | |
| | Red precipitate / orange-red precipitate | | | |
| | | | | |
| | Reagent to confirm the absence of ethanoic acid | | | |
| | Add sodium hydrogencarbonate or sodium carbonate | 1 | AO1b | |
| | Result; no effervescence observed; hence no acid present | 1 | AO1b | M5 can only be awarded if M4 is given correctly |
| | OR | | | |
| | Reagent; add ethanol and concentrated sulfuric acid and warm | | | |
| | Result; no sweet smell / no oily drops on the surface of the liquid, | | | |
| | hence no acid present | | | 2 |
| | | | | Z |

| Question | Marking guidance | Mark | AO | Comments |
|----------|--|------|------|--|
| 02.1 | | | | Extended response |
| | Stage 1: Moles of acid at equilibrium | | | |
| | Moles of sodium hydroxide in each titration = $(3.20 \times 2.00 \times 10^{-1}) / 1000 = 6.40 \times 10^{-4}$ | 1 | AO2h | |
| | Sample = 10 cm^3 so moles of acid in 250 cm^3 of equilibrium mixture | | | |
| | $= 25 \times 6.40 \times 10^{-4} = 1.60 \times 10^{-2}$ | 1 | AO2h | M2 can only be scored if = answer to M1 \times 25 |
| | Stage 2: Moles of ester and water formed | | | |
| | Moles of acid reacted = $8.00 \times 10^{-2} - 1.60 \times 10^{-2} = 6.40 \times 10^{-2}$ | | | |
| | = moles ester and water formed | 1 | AO2h | M3 is 8.00 × 10^{-2} – M2 |
| | Stage 3: Moles of ethanol at equilibrium | | | |
| | Moles of ethanol remaining = $1.20 \times 10^{-1} - 6.40 \times 10^{-2} = 5.60 \times 10^{-2}$ | 1 | AO2h | M4 is 1.20 × 10 ⁻¹ – M3 |
| | Stage 4: Calculation of equilibrium constant | | | |
| | $K_c = [CH_3COOCH_2CH_3] [H_2O] / [CH_3COOH] [CH_3CH_2OH]$ | 1 | AO1b | |
| | $= (6.40 \times 10^{-2})^2 / (1.60 \times 10^{-2})(5.60 \times 10^{-2})$ | | | |
| | = 4.5714 = 4.57 | 1 | AO2h | M6 is $M3^2 / M2 \times M4$ Answer must be given to 3 significant figures |

| 02.2 | | | 1 | | | | | |
|------|---|-------------------|----------|-------------|----------|---|--------|--|
| | | Rough | 1 | 2 | 3 | | | |
| | Final burette reading / cm ³ | 4.60 | 8.65 | 12.85 | 16.80 | | | |
| | Initial burette reading / cm ³ | 0.10 | 4.65 | 8.65 | 12.85 | | | |
| | Titre / cm ³ | 4.50 | 4.00 | 4.20 | 3.95 | 1 | AO1b | |
| | | | | | | | | |
| 02.3 | Mean = 4.00 + 3.95 / 2 = 3.98 (| cm ³) | | | | 1 | AO3 1a | Allow 3.975 (cm ³) |
| | Titres 1 and 3 are concordant | | | | | | AO3 1a | Allow titre 2 is not concordant |
| 02.4 | Thymol blue | | | | | | AO1b | |
| 02.5 | Percentage uncertainty: 0.15/3.98 × 100 = 3.77% | | | | | | AO2h | Allow consequential marking on mean titre from 2.3 |
| 02.6 | Use a lower concentration of NaOH | | | | | | AO3 2b | |
| | So that a larger titre is required titre) | (reduces | percenta | ige uncerta | ainty in | 1 | AO3 2b | |

| Question | Marking guidance | Mark | AO | Comments |
|----------|---|------|--------|-----------------|
| 03.1 | Wear plastic gloves: | | | |
| | Essential – to prevent contamination from the hands to the plate | 1 | AO3 1a | |
| | Add developing solvent to a depth of not more than 1 cm ³ : | | | |
| | Essential – if the solvent is too deep it will dissolve the mixture from the plate | 1 | AO3 1a | |
| | Allow the solvent to rise up the plate to the top: | | | |
| | Not essential – the Rf value can be calculated if the solvent front does not reach the top of the plate | 1 | AO3 1a | |
| | Allow the plate to dry in a fume cupboard: | | | |
| | Essential – the solvent is toxic | 1 | AO3 1a | Allow hazardous |
| 03.2 | Spray with developing agent or use UV | 1 | AO1b | |
| | Measure distances from initial pencil line to the spots (x) | 1 | AO2h | |
| | Measure distance from initial pencil line to solvent front line (y) | 1 | AO2h | |
| | R_f value = x / y | 1 | AO1b | |
| 03.3 | Amino acids have different polarities | 1 | AO1b | |
| | Therefore, have different retention on the stationary phase or different solubility in the developing solvent | 1 | AO1b | |

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| Question | | Marking guidance | Mark | AO | Comments |
|----------|--|---|------|--------|--|
| 04.1 | Scheme Ir question. Level 3 5–6 marks Level 2 3–4 marks | tion is marked using levels of response. Refer to the Mark instructions for Examiners for guidance on how to mark this All stages are covered and the explanation of each stage is generally correct and virtually complete. Answer is communicated coherently and shows a logical progression from stage 1 and stage 2 to stage 3. Steps in stage 3 must be complete, ordered and include a comparison. All stages are covered but the explanation of each stage may be incomplete or may contain inaccuracies OR two stages are covered and the explanations are generally correct and virtually complete. Answer is mainly coherent and shows aprogression from Stage 1 and stage 2 to stage 3. | 6 | AO3 1a | Indicative Chemistry content |
| | Level 1 1–2 marks | Two stages are covered but the explanation of each stage may be incomplete or may contain inaccuracies, OR only one stage is covered but the explanation is generally correct and virtually complete. Answer includes some isolated statements, but these are not presented in a logical order or show confused reasoning. | | | The O–H bond in the ethanedioic acid is more polarised / H becomes more δ+ More dissociation into H⁺ ions Ethanedioic acid is stronger than ethanoic acid |
| | Level 0 0 marks | Insufficient correct Chemistry to warrant a mark. | | | |

| 04.2 | | | | Extended response |
|------|---|---|------|---|
| | Moles of NaOH = Moles of HOOCCOO ⁻ formed = 6.00×10^{-2} | 1 | AO2h | |
| | Moles of HOOCCOOH remaining = $1.00 \times 10^{-1} - 6.00 \times 10^{-2}$ | | | |
| | $= 4.00 \times 10^{-2}$ | 1 | AO2h | |
| | $K_{a} = [H^{+}][A^{-}]/[HA]$ | | | |
| | $[H^+] = \mathcal{K}_{a} \times [HA]/[A^-]$ | 1 | AO2h | |
| | $[H^+] = 5.89 \times 10^{-2} \times (4.00 \times 10^{-2}/V) / (6.00 \times 10^{-2}/V) = 3.927 \times 10^{-2}$ | 1 | AO2h | |
| | $pH = -log_{10}(3.927 \times 10^{-2}) = 1.406 = 1.41$ | 1 | AO1b | Answer must be given to this precision |
| 04.3 | $5H_2C_2O_4 + 6H^+ + 2MnO_4^- \longrightarrow 2Mn^{2+} + 10CO_2 + 8H_2O$ | 1 | AO2d | |
| | OR $5C_2O_4^{2-} + 16H^+ + 2MnO_4^- \longrightarrow 2Mn^{2+} + 10CO_2 + 8H_2O$ | | | |
| | Moles of KMnO ₄ = $20.2 \times 2.00 \times 10^{-2}/1000 = 4.04 \times 10^{-4}$ | 1 | AO2h | |
| | Moles of $H_2C_2O_4 = 5/2 \times 4.04 \times 10^{-4} = 1.01 \times 10^{-3}$ | 1 | AO2h | |
| | Concentration = moles/volume (in dm ³) | | | |
| | = $1.01 \times 10^{-3} \times 1000/25 = 4.04 \times 10^{-2} \text{ (mol dm}^{-3}\text{)}$ | 1 | AO2h | If 1:1 ratio or incorrect ratio used, M2 and M4 can be scored |

| Question | Marking guidance | Mark | AO | Comments |
|----------|---|------|--------|--------------------------------|
| 05.1 | [CH₃OCOCOOH]⁺ | 1 | AO3 1a | Allow names |
| | [CH ₃ OCOCOOCH ₃] ⁺ | 1 | AO3 1a | Do not allow molecular formula |
| 05.2 | Positive ions are accelerated by an electric field | 1 | AO1a | |
| | To a constant kinetic energy | 1 | AO1a | |
| | The positive ions with m/z of 104 have the same kinetic energy as those with m/z of 118 and move faster | 1 | AO2e | |
| | Therefore, ions with m/z of 104 arrive at the detector first | 1 | AO2e | |

Section B

In this section, each correct answer is awarded 1 mark.

| Question | Key | AO |
|----------|-----|--------|
| 6 | A | AO3 1b |
| 7 | В | AO2f |
| 8 | D | AO2d |
| 9 | D | AO2d |
| 10 | В | AO2b |
| 11 | А | AO2b |
| 12 | D | AO1b |
| 13 | В | AO2d |
| 14 | С | AO2h |
| 15 | В | AO1b |
| 16 | D | AO2c |
| 17 | D | AO2c |
| 18 | A | AO3 1b |
| 19 | В | AO3 1a |
| 20 | D | AO3 1a |

| Question | Кеу | AO |
|----------|-----|--------|
| 21 | В | AO1a |
| 22 | В | AO1a |
| 23 | D | AO2h |
| 24 | В | AO1a |
| 25 | С | AO1a |
| 26 | С | AO1a |
| 27 | С | AO3 1a |
| 28 | D | AO1a |
| 29 | В | AO3 1a |
| 30 | D | AO3 1a |
| 31 | В | AO1a |
| 32 | С | AO3 1b |
| 33 | D | AO2a |
| 34 | С | AO3 1a |
| 35 | С | AO1a |