Mark Scheme - OA4 Organic Synthesis and Analysis

1	(a) Chromophore			[1]	
	(b)	(i)	Melting temperature lower than literature value / melting occurs over a		
			temperature range	[1]	
		(ii)	Identify percentage or amount of impurities (1)		
			Identify the number of compounds present or number of impurities	(1) [2]	
	(c)	(i)	Acidified potassium dichromate (1)		
			Heat and distil (1) do not accept 'reflux'	[2]	
		(ii)	M_r of phenylmethanol = 108.08 M_r of benzenecarbaldehyde = 10 100% conversion would be 10.0 ÷ 108.08 × 106.06 = 9.815g (1)	06.06 (1)	
			86% yield = $9.815 \times 86 \div 100 = 8.44g$ (1)	[3]	
		(iii)	Two resonances in the range 5.8-7.0 ppm (1)		
			These are doublets (1)		
			One singlet at around 11.0 ppm (1)		
			All resonances have the same area (1)	[4]	

Total [13]

- 2 (a) CH₃CH(CH₃)CH₂Cl (1) AlCl₃ / FeCl₃ (1) Room temperature / in the dark (1) [3]
 - (b) (i) 2,4-DNP (1) Orange precipitate (1) [2]
 - (ii) Tollen's reagent (1) Silver mirror with C, no reaction with B (1) [2]
 - (c) Optical isomerism is where a molecule and its mirror image are different / nonsuperimposable (1)

Compound C has a chiral centre / 4 different groups attached to one carbon atom
(1)

The two isomers rotate the plane of polarised light in opposite directions (1) [4]

QWC: organisation of information clearly and coherently; use of specialist vocabulary where appropriate (1) [1]

- (d) Dilute acid (1) heat (1) hydrolysis (1) [3]
- (e) Acidified potassium dichromate (VI) (1) / heat (1)

One step reactions are generally better as they have a better yield / there is waste in each stage (1)

Two step process may be cheaper / use more sustainable reagents / may give a better yield in this case / produce less harmful waste materials / potassium dichromate may react with other parts of the molecule as well / may be easier to separate product (1)

Do not credit same idea twice e.g. if 'better yield' gains first mark, a different point is required to gain second mark [4]

QWC: selection of a form and style of writing appropriate to purpose and to complexity of subject matter [1]

Total [20]

3 (a) (i) Sodium / potassium cyanide

(iii) Sulfuric / hydrochloric acid [1]

(v) eg

(vi) LiAlH₄ / H₂ / sodium, ethanol [1]

(vii) The nitrogen atoms act as electron pair donors / proton acceptors [1]

(b) (i) Molecular formula is C₄H₄O₂ [1]

- (iv) Two of the (remaining) protons are in equivalent environments (and one is not) / there are CH and CH₂ present [1]
- (v) Possibilities

Total [12]

[1]

- 4 (a) (i) 48.5 / 49 % [1]
 - (ii) Find a use for the calcium sulfate [1]
 - (b) Total volume of aqueous sodium hydroxide needed = $\frac{26.40 \times 250}{25.00}$ = 264.0 cm³ (1)

from the graph this is equivalent to 0.011 mole of the acid (1)

$$\therefore M_r \text{ of the acid} = \underbrace{\frac{\text{mass}}{\text{no. of moles}}} = \underbrace{\frac{2.31}{0.011}} = 210 \quad (1)$$

$$C_6H_8O_7$$
. $n H_2O = 210$
 \uparrow
 $192 \therefore n = 18$ (1)

since M_r of water is 18 n = 1 (1) [5]

- (c) The two 'ends' of the double bond have different groups bonded to the carbon atoms (of the double bond) / they have different structural formulae, so cannot be stereo / geometric isomers [1]
- (d) eg sodium ethanoate / ethanoic acid (1) methane (1) [2]
- (e) $C_5H_6O_5 \rightarrow CH_3COCH_3 + 2CO_2$ [1]

(f)

[1]

- (g) (Fractional) distillation / (preparative) gas chromatography / HPLC
- (h) (i) eg An optically active isomer that will rotate the plane of polarised light
 / an isomer with a chiral centre [1]
 - (ii) An equimolar mixture of both enantiomers (that has no apparent effect on the plane of polarised light) [1]

Total [15]

5 (a) (i) A compound that can rotate the plane of polarised light. [1]

(iv) Reflux / heat with H₂O/H⁺ [1]

 (v) It contains an equal amount of the two enantiomers / it is a racemic mixture (1)

The rotating effect of one form exactly cancels out the effect of the other (1) [2]

(ii) H₃C—C—COOH

(c) (i) 2-aminopropanoic acid [1]

(ii) Nitrous acid / nitric(III) acid / HNO₂ [1]

(iii) It exists as a zwitterion (1)
strong electrostatic attractions / ionic bonds between different

zwitterions (1) [2]

Total [12]

[1]

- 6 (a) (i) $CH_3CH_2CH_2CH_3 + Cl_2 \rightarrow CH_3CH_2CH_2CH_2CH_2Cl_1 + HCl_1$ [1]
 - (ii) CH₃CH₂CHCH₃ [1]
 - (b) (Anhydrous) aluminium chloride / iron(III) chloride allow AlCl3 / FeCl3 [1]
 - (c) (i) orange / red precipitate [1]

(ii) O_{CC}CH₃

(1) —COCH3 groups in any positions

It must contain a C=O group but it is not an aldehyde as it does not react with Tollens' reagent (1) [2]

- (d) (i) (Alkaline) potassium manganate(VII) (solution) allow KMnO₄ / MnO₄ [1]
 - (ii) Dilute acid allow HC1/H⁺ [1]
 - (iii) Lithium tetrahydridoaluminate(III)/lithium aluminium hydride allow LiAlH4 [1]

(iv)

(e) Only the infrared spectrum of benzoic acid would have a peak at 1650–1750 cm⁻¹ (1)

This is due to the carbonyl group present in the benzoic acid (1) [2]

Total [12]

(b) Moles of calcium carbide = 500/64.1 = 7.80 (1)

Moles of ethyne = 7.80

Volume of ethyne = $7.80 \times 24.0 = 187 \text{ (dm}^3\text{)}$ (1) [2]

(c) If the process is endothermic left to right then it needs to absorb energy

- hence the high temperature / endothermic reactions need a high temperature [1]

Curly arrows (1), full (1) and partial charges (1) [3]

(f) Any two for (1) each
energy costs / cost of catalyst / problems of separation of products /
time taken / availability of starting materials / percentage yield /
atom economy / relative health and safety [2]

(g)
$$C_6H_5 - C \equiv C - CH_2 - CH_3$$
 (1) C_1H_1 (1) [2]

(ii) I sulfuric acid / H₂SO₄ / phosphoric acid / H₃PO₄ / Al₂O₃ [1]

II 3-hydroxypropanoic acid does not show a C = C absorption at 1620-1670 cm⁻¹ but this is present in propenoic acid [1]

III The CH₃—C / CH₃CH(OH) group is absent [1]

Total [16]

8 (a) (i) 2 mol of ethanol gives 1 mol of ethoxyethane (1)

Moles of ethanol =
$$\frac{69}{46}$$
 = 1.5

- :. Moles of ethoxyethaneif theoretical yield = 0.75
- :. Moles of ethoxyethaneif 45% yield = $0.75 \times 0.45 = 0.34$ (1)

Mass of ethoxyethane = $0.34 \times 74 = 25g$ (1) allow error carried forward

[3]

(ii) Ethene / C2H4

[1]

[2]

(iii) H H

$$\begin{vmatrix} \xi_{\perp} \\ \xi_{-} \\ -C - C - Br \end{vmatrix}$$
 product

- for correct curly arrows (1) for correct δ⁺ and δ⁻
- (iv) They need to have an N-H/O-H/F-H bond/a highly electronegative atom bonded to hydrogen [1]
- (b) (i) For example

Accept any polybrominated species

Do not accept a monobrominated species

- (ii) Bromine decolorised / orange to colourless / white solid
- [1]

- (c) Reagent Iron(III) chloride solution / FeCl₃ (1)
 - Observation Purple coloration / solution (1)

- [2]
- (d) (i) C₁₀H₁₂O₁ [1]

(ii)
$$H \xrightarrow{H} H$$
 $CH_0 - C - C - C$ $CH_0 - C - C - C$ $CH_0 - C$ $CH_$

(e) Displayed formula, for example

Functional group carboxylic acid (1) [2]

Total [15]

9	(a)	(i)	A	[1]
		(ii)	D	[1]
		(iii)	С	[1]
		(iv)	С	[1]
	(b)	(i)	Nucleophilic substitution	[1]
		(ii)	The C–Cl bond in chlorobenzene is stronger than in 1-chlorobutane (1) due to delocalization of electron density from the ring with the bond (1)	
			OR	
			Delocalised electrons in chlorobenzene (1) repel lone pair of electrons on nucleophile / ammonia (1)	[2]
		(iii)	C ₄ H ₉ NH ₂ + CH ₃ COCI	[1]
		(iv)	I Tin and concentrated hydrochloric acid (1)	
			Add sodium hydroxide (after cooling) (1)	
			Steam distillation to separate the product (1)	[3]
			II C₅H₅NN⁺CI⁻	[1]
			III Azo dye / azo compound	[1]

Total [13]

- 10 (a) (i) (Fractional) distillation / (preparative) gas chromatography / HPLC / TLC column chromatography / solvent extraction [1]
 - (ii) the fragmentation pattern would be different / valid examples given [1]
 - (iii) I

II Heated electrically / by a naked flame with a water bath (1) Add compound G to the ethanol until the hot ethanol will (just) not dissolve any more solute (1)

Filter hot (1)

Allow to cool (1)

Filter (1)

Dry in air / window sill / < 60 °C in an oven (1)

Maximum 4 out of 5 total if second marking point not given Note 5 marks maximum here

QWC Information organised clearly and coherently, using specialist vocabulary where appropriate [1]

(iv) I The amine is reacted with so dium nitrite / HCl(aq) or nitrous acid (1) at a temperature of < 10 °C (1) [2]

II

[5]

(b) (i) Nucleophilic addition (1)

Accept a mechanism that shows HCN polarisation and nucleophilic addition as a concerted process

polarisation / charges shown (1) curly arrows on first structure (1) regeneration of ⁻C≡N or capture of H⁺ and curly arrow (1) [4]

(ii) Chromophores (1)
The colour will be black (1) as the compound absorbs blue / other colours (1)

[3]

Total [20]

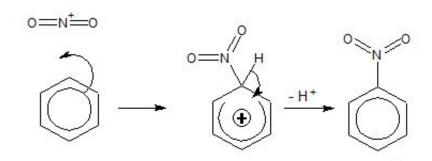
11 (a) (i) Both molecules have lone pairs on nitrogen (1)

The lone pairs can form (coordinate) bonds with H+ ions (1) [2]

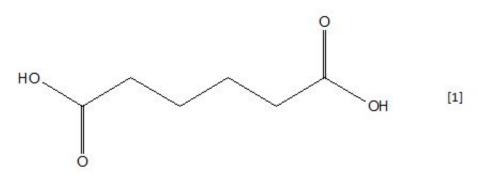
- (ii) Lone pair on N in phenylamine is delocalised over benzene ring (1) therefore less able to accept H⁺ (1)
- (iii) I Arrow in first step (1)

 Cation structure in second step (1)

 Arrow in second step (1)



- II (fractional) distillation / steam distillation
- III Sn and conc. HCl (1) followed by NaOH (1) [2]
- (b) (i)



(ii) Addition polymerisation makes one product only /
 Condensation produces one product plus a small molecule like water (1)

Addition polymerisation uses one starting material / Condensation polymerisation has two different starting materials (1)

Addition polymerisation involves monomer with one functional group /
Condensation polymerisation involves monomer with two functional groups
(1)

(max 2) [2]

[3]

[1]

(iii) Alanine has strong (electrostatic) forces between the zwitterions (1)

Butanoic acid has hydrogen bonding between molecules /
electrostatic forces in alanine are stronger than forces in butanoic acid

(1)

[2]

(iv) Soda lime (1)
$$CH_3CH_2NH_2$$
 (1) [2]

Total [20]