M1.	(a)	(base) elimination (penalise other words before 'elimination' e.g. nucleophilic)	
	M1 :	curly arrow from lone pair of electrons on oxygen of hydroxide ion (insist on a lone pair of electrons on the oxygen atom and a negative charge, but only credit this mark if the attack is to a correct H atom)	
	M2 : curly arrow from the middle of the C-H bond to the middle of the C-C bond		
		(only credit this mark if the arrow originates from the correct C–H bond and if an attempt has been made at M1)	
		M3 : curly arrow from the <u>middle of the C–Br bond</u> towards/alongside the Br atom	
		(credit M3 independently unless the bond breaking is contradicted by an additional arrow)	
		(penalise curly arrow if the C–Br has a formal positive charge)	
		(credit full marks for an E1 mechanism, with M2 awarded for a correct curly arrow on the correct carbocation)	
		(award a maximum of two marks for either an incorrect haloalkane or an incorrect organic product)	
		(maximum 2 marks for use of 'sticks' for the haloalkane, unless RE from 2(b), when credit can be given)	
(b)	(i)	M1: compounds with the same structural formula	
		M2 : but the bonds/groups/atoms have different spatial arrangements or orientation or configuration/are arranged differently in space/3D	
		(ignore reference to the same molecular formula for M1)	
	(ii)	M1 : correct structural representation for cis-but-2-ene <u>and</u> its name or its identification as the cis isomer	
		M2: correct structural representation for trans-but-2-ene	

(accept representations which are 90° to linear)

(award one mark for two correct structures but either wrong/no names)
(maximum 1 mark for an incorrect alkene)

ini i mark for an incorrect alkene)

(iii) geometric(al) or cis-trans

1

1

(c) nucleophile or electron pair donor (penalise 'base')

1

(d) CH₃CH₂CH₂Br + 2NH₃ → CH₃CH₂CH₂CH₂NH₂ + NH₄Br
 (M1 correct product)
 (M2 balanced equation using 2NH₃ and leading to NH₄Br)
 (penalise M1 for use of C₄H₃NH₂ or for incorrect haloalkane, but allow consequent correct balancing of equation with 2 moles of ammonia)

2

(1-)butylamine

(credit 1-aminobutane and butyl-1-amine) (award QoL mark for correct spelling)

1

[13]

- **M2.** (a) (i) UV light OR sunlight OR T \geq 450°C (1) NOT high T
 - (ii) (free) radical substitution (1)
 - (iii) CCI₄ (1) OR named

3

- (b) (i) $CH_3CI + KCN \rightarrow CH_3CN + KCI$ (1) $CN^ Cl^-$
 - (ii) <u>nucleophilic substitution</u> (1)

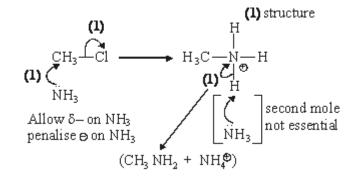
(iii) <u>C-Br bond</u> is <u>weaker</u> (than C-Cl bond)
OR <u>C-Br bond</u> enthalpy is <u>less than</u> C-Cl **(1)** *Ignore electronegativity*

3

(c) CH₃COOH OR ethanoic acid (1)

1

- (d) (i) $\overset{\delta_{+}}{C} \overset{\delta_{-}}{C} \overset{\delta_{-}}{C} = \underbrace{OR \ C-Cl \ is \ polar}_{C}$ (1) OR C atom is electron deficient / δ_{+}
 - (ii) methylamine (1) only
 - (iii) S_N1 scores full marks



[13]

M3.B

[1]

M4.A

[1]

M5. (a) (i) (Free) radical substitution (Both words needed)

1

1

M2 ultra-violet light OR sunlight OR 1000°C ≥ T ≥ 450 °C

(Ignore reference to temperature if included with uv light)

(Penalise "high temperature" for M2)

1

(iii) $^{2\dot{C}H_3} \rightarrow C_2H_6$ (OR CH_3CH_3 as alternative to C_2H_6)

1

(iv) $CH_3Br + Br_2 \rightarrow CH_2Br_2 + HBr$

(b) (i) <u>Electron pair donor</u> OR species with an <u>electron pair</u> able to form a covalent <u>bond</u>.

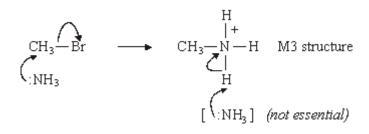
1

(ii) Methylamine

(Credit "aminomethane")

1

(iii)



1

M1 arrow to show breakage of C - Br bond

1

M2 arrow from lone pair on N of NH₃ to form bond with C

1

M4 arrow from bond of N – H to N atom of CH₃NH₃ (Ignore partial charges on haloalkane but penalise if incorrect)

(Accept
$${^{C\!H_3}}^{\stackrel{+}{N}}{^{H_3}}$$
 for M3)

(Full credit for carbocation mechanism; M1 for C – Br bond breakage and M2 for lone pair attack on carbocation)
(Second mole of ammonia not essential to mechanism for

[11]

1

M6.D

[1]

M7.D

[1]

M8. (a) (i) Electrophilic addition (Both words required)

1

(ii) M1 the reaction to form 1-bromopropane goes *via* the primary carbocation OR 1° carbocation

OR via ${^{\text{CH}_3\text{CH}_2}}^{\overset{+}{\text{C}}\text{H}_2}$

M2 primary carbocations are less stable than secondary carbocations

(Credit converse arguments for M1 and M2 i.e. the reaction to form 2-bromopropane goes via the secondary carbocation, M1, and secondary carbocations are more stable than primary carbocations, M2)

(Accept the use of "carbonium ions" as an alternative to carbocation)

1

(b) M1 NaOH OR KOH OR correct name

1

M2 aqueous or solution in water (ignore heat, reflux etc.)

(Penalise M1 for hydroxide ion alone, but mark on and credit M2)

(Credit M2 ONLY for H₂O as reagent and heat / warm / T=50 to 100°C)

(NaOH(aq) scores M1 and M2 provided it is not contradicted) (Penalise M2 if NaOH(aq) followed by concentrated or ethanol)

(Penalise M1 and M2 if followed by acid)

1

(c) Ethanolic OR alcoholic OR CH3CH2OH / CH3OH solvent OR aqueous ethanol/alcohol

OR higher temperature (must be comparative)

(Ignore heat or heat under reflux) (Credit part (c) independently from part (b)) (Penalise "ethanoic")

1

(d) (i) Secondary OR 2°

1

M1 arrow from double bond to H of H - O bond

M2 arrow from bond to oxygen atom to show H - O bond breakage M4 arrow from lone pair of electrons to carbon atom of carbocation

(Penalise M1 if arrow goes to H₂SO₄ or to formal positive charge on H, but ignore partial charges on sulphuric acid unless wrong)

(Credit M2 for H⁺ ion)

(For M4, accept negative charge anywhere on the ion)

4

1

(iii) Catalyst ONLY

(Ignore homogeneous, heterogeneous)

[12]