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CHEMISTRY

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Paper 4 A Level Structured Questions MARK SCHEME Maximum Mark: 100

Published

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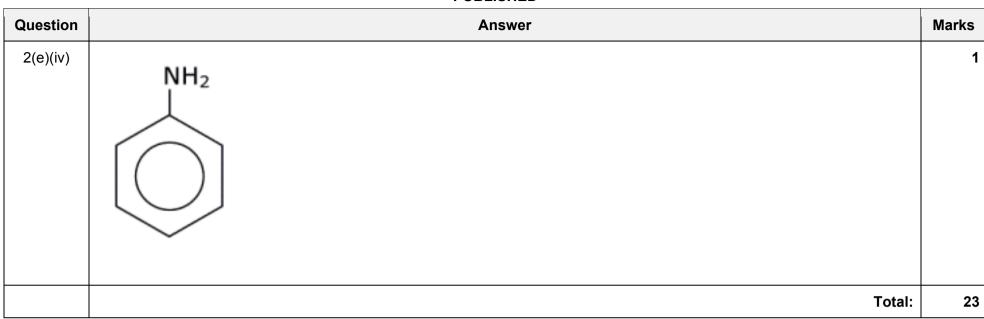
Question	Answer	Marks
1(a)	solubility increases down the group	1
	ΔH_{latt} and ΔH_{hyd} both decrease or ΔH_{latt} and ΔH_{hyd} both become less exothermic / more endothermic	1
	ΔH_{latt} decreases / changes more (than ΔH_{hyd} as OH ⁻ being smaller than M ²⁺)	1
	ΔH_{sol} becomes more exothermic / more negative / less endothermic / less positive	1
1(b)(i)	$\Delta H_{r1} - (538 + 2x230 + 394) = -(1216 + 286)$	1
	$\Delta H_{\rm r1} - 1392 = -1502$	
	$\Delta H_{r1} = -110$	1
1(b)(ii)	let $\Delta H_{f}(HCO_{3}^{-}(aq)) = y$	1
	2y - 538 = -1216 - 394 - 286 - 26	
	y = -692	1
1(b)(iii)	$\Delta H_{r3} - 538 - 2(230 + 394) = -538 - 2(692)$	1
	$\Delta H_{r3} = -136$	
1(b)(iv)	ΔH_{r3} will be identical to ΔH_{r4} , / unchanged	1
	as the reaction is the same, or:	1
	$2OH^{-}(aq) + 2CO_{2}(g) \longrightarrow 2HCO_{3}^{-}(aq)$ or	
	metal ions stay in solution/metal ions are unchanged / are spectators	

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Question	Answer	Marks
1(c)	more gaseous moles are being consumed (in reaction 3) or more CO ₂ moles are being consumed (in reaction 3)	1
	ΔS is therefore expected to be more negative/less positive for reaction 3.	1
	Total:	13

Question	Answer	Marks
2(a)(i)	$ \begin{array}{c} \begin{array}{c} \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} \\ \end{array} \\ \end{array} \\ \begin{array}{c} \end{array} \\ \end{array} $	1 + 1
	16 electrons on each diagram	1
2(a)(ii)	HNC = 115–125° AND NCO = 180°	1
2(a)(iii)	cyanic acid, because it's a stronger / higher bond enthalpy / triple / C≡N / more electrons involved bond	1
2(b)(i)	$[H^+] = \sqrt{([HNCO]K_a)} = \sqrt{(0.1 \times 1.2 \times 10^{-4})} \text{ or } 3.46 \times 10^{-3}$	1
	$pH = log [H^+] = 2.5 (2.46)$	1
2(b)(ii)	$Na_2CO_3 + 2(NH_2)_2CO \longrightarrow 2NaNCO + CO_2 + 2NH_3 + H_2O$	1
2(c)(i)	$(n(OH^{-}) \text{ at start} = (2 \times 0.1 \times 30) / 1000 = 6 \times 10^{-3} \text{ mol})$ $(n(OH^{-}) \text{ reacted} = (0.1 \times 20) / 1000 = 2 \times 10^{-3} \text{ mol})$ $n(OH^{-}) \text{ remaining} = (6-2) \times 10^{-3} = 4 \times 10^{-3} \text{ mol}, (in 50 \text{ cm}^{3})$	1
	so $[OH^{-}]_{end} = (4 \times 10^{-3} \times 1000) / 50 = 0.08 \text{ mol dm}^{-3}$	1

Question	Answer	Marks
2(c)(ii)	$[H^+] = K_w / [OH^-] = (1 \times 10^{-14}) / 0.08 = 1.25 \times 10^{-13} \text{ mol dm}^{-3}$	1
	so pH = $-\log(1.25 \times 10^{-13})$ = 12.9	1
2(c)(iii)	curve starts at 2.46 / 2.5	1
	vertical portion (end point) at vol added = 10.0 cm^3	1
	finishes at pH = 12.9	1
2(d)(i)	monodentate: (a species that) forms one dative / coordinate bond	1
	ligand: a species that uses a lone pair of electrons to form a dative / coordinate bond to a metal atom / metal ion	1
2(d)(ii)	[Ag(NCO) ₂] ⁻ or [Ag(OCN) ₂] ⁻ correct formula	1
	correct charge	1
2(e)(i)	$n(BaCO_3) = 1.66 / 197.3 = 8.4(1) \times 10^{-3} mol$	1
2(e)(ii)	$n(RNCO) = 8.41 \times 10^{-3} \text{ mol, so } M_r = 1 / (8.41 \times 10^{-3}) = 119$	1
2(e)(iii)	molecular formula = C ₇ H ₅ NO	1

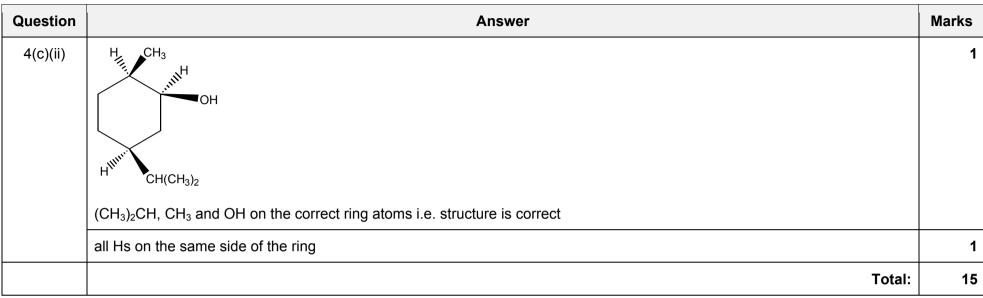


Question	Answer	Marks
3(a)(i)	+3 or Co ³⁺	1
3(a)(ii)	oxidation	1
	ligand displacement / replacement / exchange / substitution	1

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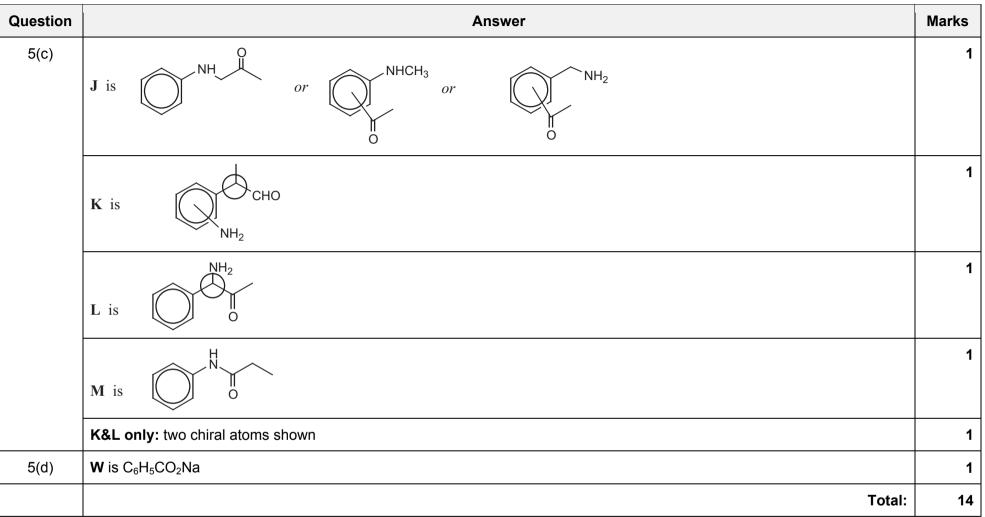
Question	Answer	Ma	larks		
3(a)(iii)	$\begin{bmatrix} I \\ H_{3}N/III, I \\ H_{3}N/III, I \\ H_{3}N/III, I \\ CI \\ CI \end{bmatrix}^{+} or \begin{bmatrix} I \\ H_{3}N/III, I \\ CI \\ H_{3}N/III, I \\ CI \\ H_{3}N/III, I \\ CI \\ H_{3}N/III, I \\ III \\ IIII \\ III \\ IIII \\ III \\ III \\ III \\ III \\ III \\ IIII \\ IIII \\ IIII \\ III \\ IIII \\ III \\ III \\ IIII \\ III \\ IIII \\ IIIII \\ IIII \\ IIIII \\ IIII \\ IIII \\ IIII \\ IIII \\ IIIII \\ IIII \\ IIII \\ IIII \\ IIII \\ IIIIII$		1+1		
			1		
	geometrical or cis-trans		1		
3(b)(i)	The number of bonds / atoms bonded to an atom / ion / species / metal				
3(b)(ii)	C 6 $[Cr(CN)_6]$ –		6		
	D – $[Ni(NH_2CH_2CH_2NH_2)_3]$ 2+/+2				
	E 4 [PtC4] –				
	F 3 – 3–/–3				
3(c)(i)	$K_{\text{stab}(1)} = [\text{FeSCN}^{2+}]/([\text{Fe}^{3+}][\text{SCN}^{-}])$ mol ⁻¹ dm ³		3		
	$K_{\text{stab}(2)} = [\text{FeC}l_4^-]/([\text{Fe}^{3+}][\text{C}l]^4) \text{mol}^{-4} \text{dm}^{12}$				
3(c)(ii)	$K_{eq(3)} = K_{stab(1)} / K_{stab(2)}$		1		
3(c)(iii)	$K_{eq(3)} = 1750$		1		
	mol ³ dm ⁻⁹		1		
		Total:	19		

Question	Answer	Marks
4(a)(i)	optical, because it contains a / one chiral C-atom or chiral C-atoms or chiral atom / centre or C* indicated or C with 4 different groups	1
4(a)(ii)	$C_{10}H_{14}O + 3H_2 \longrightarrow C_{10}H_{20}O$ correct formulae	1
	balancing	1
4(b)(i)	electrophilic substitution	1
4(b)(ii)	step 3 reduction	1
	step 5 substitution / hydrolysis	1
4(b)(iii)	step 1 $(CH_3)_2CHCl + AlCl_3 / AlBr_3 / FeCl_3 / FeBr_3$	1+1
	step 2 $HNO_3 + H_2SO_4$ conc (T < 55 °C)	1
	step 3 Sn + HC1	1
	step 4 HNO_2 (or $NaNO_2 + HCl$) (at T < 10 °C)	1
	the two temperatures for steps 2 and 4	1
4(c)(i)	H_2 + Pt or H_2 + Ni + heat or pressure	1



Question	Answer					
5(a)		J	К	L	Μ	
		amine methyl ketone	aromatic amine aldehyde	amine methyl ketone	amide	
	J and L correct					1 + 1
	K correct					
	M correct					1
5(b)(i)	hydrolysis					1
5(b)(ii)	P is C ₆ H ₅ NH ₂					
	Q is CH ₃ CH ₂ CO ₂ N	Na				1

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Question	Answer	Marks
6(a)	Any of the three methods possible. Any 4 of the 5 points for each method available for maximum 4 marks. Method 1 1 Ensure both solutions (A and B) at 40 °C before mixing 2 mix known volumes of A and B and start the clock 3 at known time take out a sample / X and add it to ice-cold solvent 4 titrate against HC <i>I</i> 5 repeat at time at known time intervals Method 2 1 Ensure both solutions (A and B) at 40 °C before mixing 2 mix known volumes of A and B and start the clock 3 at known time pour into ice-cold solvent or pour ice-cold solvent in 4 titrate against HC <i>I</i> 5 repeat with different concentrations of either A or B, or repeat using different times Method 3 1 Ensure both solutions (A and B) at 40 °C before mixing 2 mix known volumes of A and B and start the clock 3 at known time intervals Method 3 1 Ensure both solutions (A and B) at 40 °C before mixing 2 mix known volumes of A and B and start the clock and add pH meter 3 at a known time 4 record the pH 5 repeat pH readings at known time intervals	4
6(b)(i)	from 1 and 3: when $[RCl]$ is trebled, so is rate, so order w.r.t. $[RCl] = 1$	1
	from 1 and 2: when both concentrations are doubled, rate doubles so [OH ⁻] has no effect on rate, so order w.r.t.[OH ⁻] = 0	1
6(b)(ii)	rate = $k[RCl]$ AND units: sec ⁻¹ 1/s	1
6(b)(iii)	relative rate = 2.0	1

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Question	Answer	Marks
6(c)(i)	$C_{6}H_{5} \xrightarrow{C_{1}O^{-}} C_{6}H_{5} \xrightarrow{C_{6}H_{5}} C_{6}H_{5} \xrightarrow{C_{6}H_{5}} C_{6}H_{5} \xrightarrow{OH} C_{1}H_{1}H_{1}H_{1}H_{1}H_{1}H_{1}H_{1}H$	1
	intermediate cation	1
	OH ⁻ with lone pair and curly arrow	1
6(c)(ii)	 Beginning with candidate's mechanism in (c)(i): If S_N1: racemate / mixture of / two optical isomers will be formed, because: the intermediate is planar / has a plane of symmetry / OH⁻ can approach from top or bottom or from any direction If S_N2: one optical isomer because attack always from fixed direction / from same side / the "configuration" always inverts / there is an asymmetric transition state 	1

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Question	on Answer							Marks
6(d)(i)		δ value	number of H atoms	group	splitting	result with D ₂ O		
		1.4	3	CH₃ / methyl	doublet	peak remains		
		2.7	1	OH / hydroxyl / alcohol	singlet	peak disappears		
		4.0	1	СН	quartet	peak remains		
	the three groups are in their correct places wrt the δ values no. of H atoms for each peak agrees with group column						1	
	splitting patterns doublet, singlet and quartet are assigned to correct groups						1	
	peak identified	as OH di	isappears with D ₂ O, no	o other peak disappears				1
							Total:	16