
CHEMISTRY

9701/42

Paper 4 A Level Structured Questions

October/November 2017

MARK SCHEME

Maximum Mark: 100

Published

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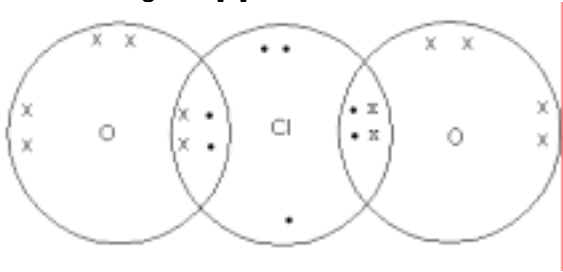
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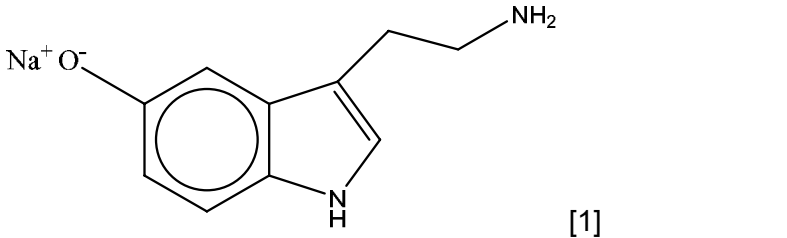
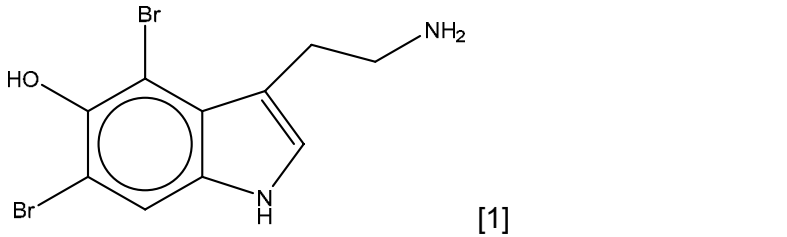
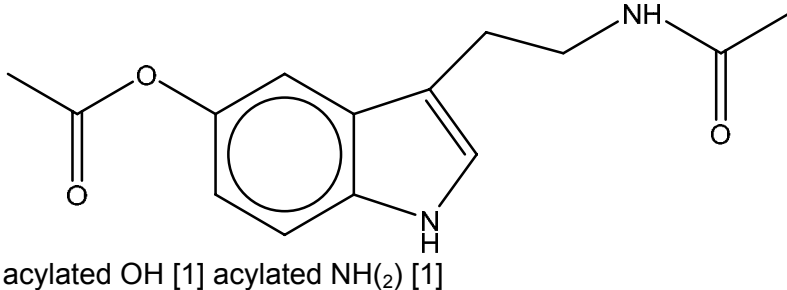
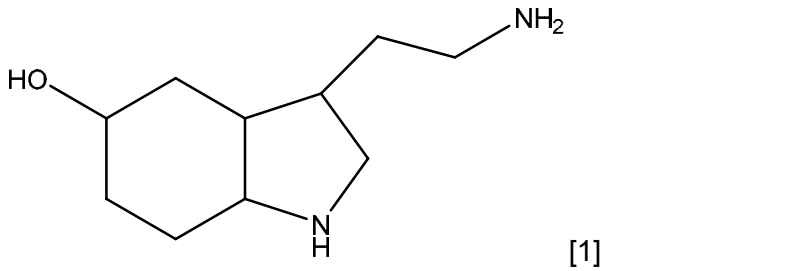
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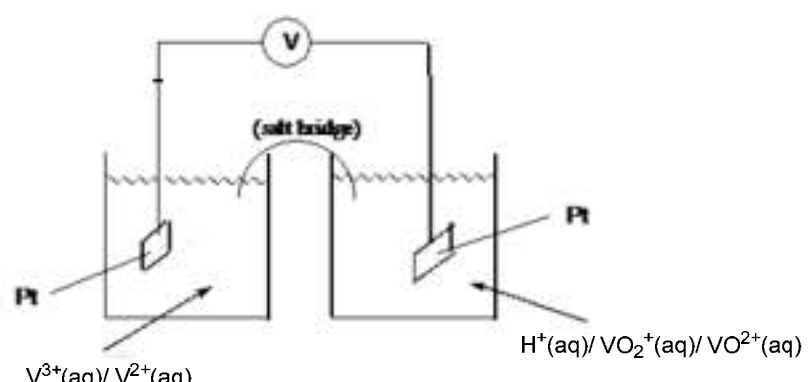
This document consists of **12** printed pages.

Question	Answer	Marks
1(a)	Cl +3 to +4 (and oxidised)	1
	Cl 0 to -1 (and reduced)	1
1(b)	19 electrons total [1] correct diagram [1] 	2
1(c)(i)	the exponent / power to which a concentration is raised in the rate equation	1
1(c)(ii)	$(0.0022 = k(0.01) \times (0.06))$ $k = 3.7$ (3.67)	1
	$\text{mol}^{-1} \text{dm}^3 \text{s}^{-1}$	1
1(c)(iii)	initial rate = 5.50×10^{-3}	1
	$[\text{ClO}_2] = 0.048$	1
1(d)(i)	slowest step (in a multi-step reaction)	1
1(d)(ii)	1 mole of F_2 and 1 mole ClO_2 reacting in the rate-determining step	1
	1st step is rate-determining step and a balanced mechanism consistent with overall equation e.g. $\text{ClO}_2 + \text{F}_2 \rightarrow \text{ClO}_2\text{F}_2$ $\text{ClO}_2 + \text{ClO}_2\text{F}_2 \rightarrow 2\text{ClO}_2\text{F}$ or $\text{ClO}_2 + \text{F}_2 \rightarrow \text{ClO}_2\text{F} + \text{F}$ $\text{ClO}_2 + \text{F} \rightarrow \text{ClO}_2\text{F}$	1
1(e)	k increases (as rate increases)	1

Question	Answer	Marks
2(a)(i)	$\text{Mg}_3\text{N}_2 + 6\text{H}_2\text{O} \rightarrow 3\text{Mg}(\text{OH})_2 + 2\text{NH}_3$	1
2(a)(ii)	moles of $\text{Mg}_3\text{N}_2 = 2.52 / 100.9 = 0.025$ (0.0249)	1
	(moles of $\text{Mg}(\text{OH})_2 = 0.075$ (0.0749)) mass of $\text{Mg}(\text{OH})_2 = (0.075 \times 58.3) = 4.37$ g or 4.4 g	1
2(b)	solubility increases (down the group)	1
	ΔH_{latt} and ΔH_{hyd} both decrease / less exothermic / more endothermic	1
	but ΔH_{latt} decreases more (than ΔH_{hyd} decreases)	1
	ΔH_{sol} becomes more negative / more exothermic / less endothermic	1
2(c)(i)	$K_{\text{sp}} = [\text{Mg}^{2+}][\text{OH}^-]^2$	1
2(c)(ii)	$K_{\text{sp}} = (1.7 \times 10^{-4}) \times (2 \times 1.7 \times 10^{-4})^2 = 2.0 \times 10^{-11}$ (1.97×10^{-11})	1
	$\text{mol}^3 \text{dm}^{-9}$	1
2(d)	cations become bigger / ionic radius increases	1
	polarisation/distortion of anion / hydroxide ion decreases	1

Question	Answer			Marks
3(b)	reagent	structure of product	type of organic reaction	8
	Na	 <p style="text-align: right;">[1]</p>	redox or reduction	
	excess Br ₂ (aq)	 <p style="text-align: right;">[1]</p>	(electrophilic) substitution	
	excess CH ₃ COCl	 <p>acylated OH [1] acylated NH₂ [1]</p>	condensation (or addition + elimination)	
excess H ₂ / Pt catalyst	 <p style="text-align: right;">[1]</p>	reduction or hydrogenation or addition		

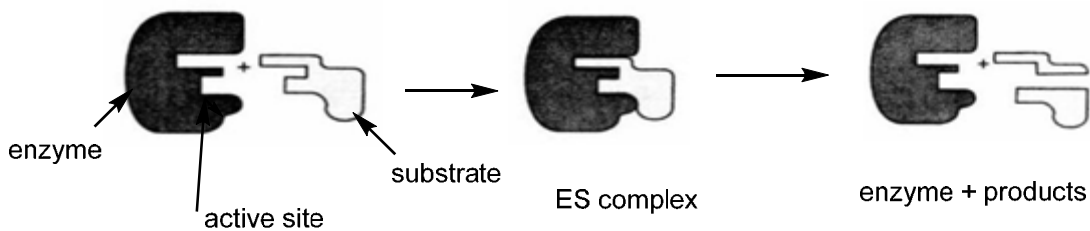
Question	Answer	Marks
3(c)(i)	(spectrum of M) contains a broad peak (for O–H) at 2500–3000 cm^{-1} or (spectrum of M) contains peak (for C=O) at 1640–1750 cm^{-1} or (spectrum of M) lacks (NH ₂ peak) at 3300–3500 cm^{-1}	1
3(c)(ii)	5 or 6 peaks	1
	OH/NH protons exchange with deuterium or $-\text{OH} / -\text{NH} + \text{D}_2\text{O} \rightarrow -\text{OD} / -\text{ND} + \text{DHO}$	1
3(d)	ester and hydrolysed	1

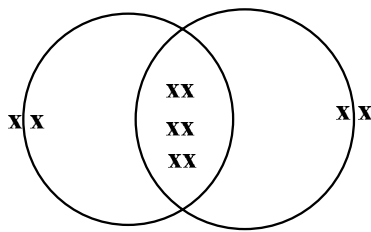
Question	Answer	Marks
4(a)(i)	$E^\ominus_{\text{cell}} = 1.00 - (-0.26) = (+)1.26 \text{ V}$	1
4(a)(ii)	$\text{VO}_2^+ + \text{V}^{2+} + 2\text{H}^+ \rightarrow \text{VO}^{2+} + \text{V}^{3+} + \text{H}_2\text{O}$	1
4(a)(iii)	 <p> $\text{V}^{3+}(\text{aq}) / \text{V}^{2+}(\text{aq})$ </p> <p> $\text{H}^+(\text{aq}) / \text{VO}_2^+(\text{aq}) / \text{VO}^{2+}(\text{aq})$ </p> <p> solutions labelled correctly in one half-cell [1] solutions labelled correctly in both half-cells [1] two graphite or platinum electrodes [1] salt bridge and voltmeter [1] </p>	4

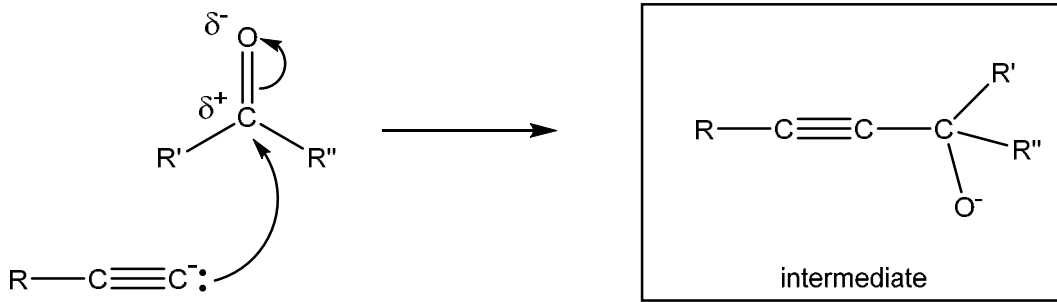
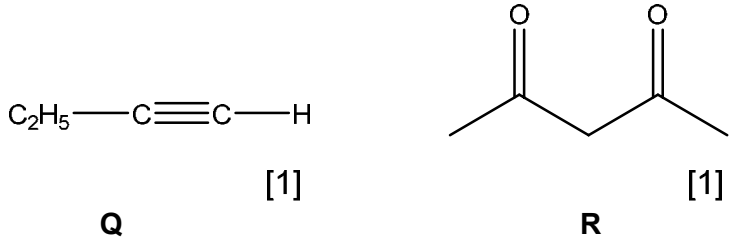
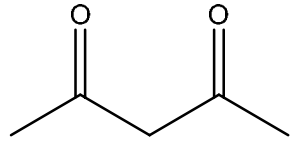
Question	Answer	Marks
4(b)	<ul style="list-style-type: none"> $V^{2+}(aq)$ and $Sn^{4+}(aq)$: yes and $E^{\circ}_{cell} = +0.15 - (-0.26) = +0.41 \text{ V}$ [1] $2V^{2+} + Sn^{4+} \rightarrow 2V^{3+} + Sn^{2+}$ [1] $VO^{2+}(aq)$ and $Fe^{3+}(aq)$ no reaction [1] 	3

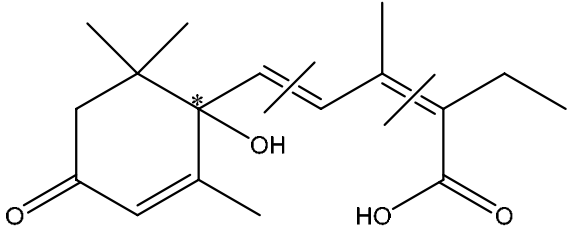
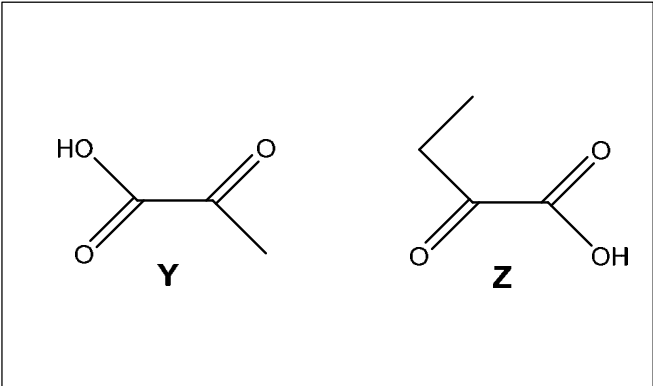
Question	Answer	Marks
5(a)	$(Na^+) 0.095 / 0.181 = 0.525$ and octahedral and co-ordination no. = 6	1
	$(Mg^{2+}) 0.065 / 0.181 = 0.359$ and tetrahedral and co-ordination no. = 4	1
5(b)	enthalpy change = $(-642) - (2 \times -106) = -430$	1
5(c)(i)	$-106 = 147 + 121 + 736 + (-349) + \text{lattice energy}$ lattice energy = -761	3
5(c)(ii)	$MgCl_2$ more exothermic / negative / bigger than $MgCl$ and $NaCl$ more exothermic / negative / bigger than $MgCl$	1
	(reason for $MgCl_2$) higher charge / lower radius of Mg^{2+} cation	1
	(reason for $NaCl$) smaller radius of Na^+ cation	1
5(d)	energy change when 1 mole of atoms / ions each gain an electron or energy change when 1 mole of atoms / ions gain 1 mole of electrons	1
	gaseous	1

Question	Answer	Marks									
6(a)	central metal atom/ion surrounded by (one or more) ligands	1									
6(b)	<table border="1"> <thead> <tr> <th></th> <th>co-ordination number</th> <th>oxidation number</th> </tr> </thead> <tbody> <tr> <td>$[\text{Pt}(\text{NH}_3)_4\text{Cl}_2]^{2+}$</td> <td>6</td> <td>+4</td> </tr> <tr> <td>$[\text{PtCl}_4]^{2-}$</td> <td>4</td> <td>+2</td> </tr> </tbody> </table>		co-ordination number	oxidation number	$[\text{Pt}(\text{NH}_3)_4\text{Cl}_2]^{2+}$	6	+4	$[\text{PtCl}_4]^{2-}$	4	+2	2
	co-ordination number	oxidation number									
$[\text{Pt}(\text{NH}_3)_4\text{Cl}_2]^{2+}$	6	+4									
$[\text{PtCl}_4]^{2-}$	4	+2									
6(c)		2									
6(d)	(HNO ₃ +) AgNO ₃ reagent	1									
	$[\text{Pt}(\text{NH}_3)_4\text{Cl}_2]\text{Br}_2$ with cream ppt. (of AgBr) and $[\text{Pt}(\text{NH}_3)_4\text{Br}_2]\text{Cl}_2$, with white ppt. (of AgCl) observation with both	1									
6(e)	octahedral: both	1									
	square planar: geometric	1									
	tetrahedral: neither	1									

Question	Answer	Marks
6(f)	<p>diagrams</p>  <p>Marks can be awarded from words or diagram. Any three marking points from:</p> <ul style="list-style-type: none"> • substrate shape is complementary to active site • the substrate binds / bonds / fits into the active site • products are released • lower E_A / bonds weakened in substrate 	3

Question	Answer	Marks
7(a)(i)	$\text{CaC}_2 + 2\text{H}_2\text{O} \rightarrow \text{C}_2\text{H}_2 + \text{Ca}(\text{OH})_2$	1
7(a)(ii)		1
7(b)	$\text{C}_n\text{H}_{2n-2}$	1
7(c)(i)	delocalised electrons	1
7(c)(ii)	CH	1
7(c)(iii)	less dense	1

Question	Answer	Marks																				
7(d)(i)	 <p>R—C≡C[−]: R'—C(=O)—R''</p> <p>2 curly arrows [1] dipole [1] intermediate [1]</p>	3																				
7(d)(ii)	nucleophilic addition	1																				
7(d)(iii)	 <p>C₂H₅—C≡C—H </p> <p>Q [1] R [1]</p>	2																				
7(e)	<table border="1" data-bbox="347 1005 1579 1244"> <thead> <tr> <th></th> <th>CH₃CHO</th> <th>HCO₂H</th> <th>CH₃COCH₃</th> <th>HO₂CCO₂H</th> </tr> </thead> <tbody> <tr> <td>hot acidified MnO₄[−](aq)</td> <td>✓</td> <td>✓</td> <td>✗</td> <td>✓</td> </tr> <tr> <td>alkaline I₂(aq)</td> <td>✓</td> <td>✗</td> <td>✓</td> <td>✗</td> </tr> <tr> <td>Tollens' reagent</td> <td>✓</td> <td>✓</td> <td>✗</td> <td>✗</td> </tr> </tbody> </table>		CH ₃ CHO	HCO ₂ H	CH ₃ COCH ₃	HO ₂ CCO ₂ H	hot acidified MnO ₄ [−] (aq)	✓	✓	✗	✓	alkaline I ₂ (aq)	✓	✗	✓	✗	Tollens' reagent	✓	✓	✗	✗	4
	CH ₃ CHO	HCO ₂ H	CH ₃ COCH ₃	HO ₂ CCO ₂ H																		
hot acidified MnO ₄ [−] (aq)	✓	✓	✗	✓																		
alkaline I ₂ (aq)	✓	✗	✓	✗																		
Tollens' reagent	✓	✓	✗	✗																		

Question	Answer	Marks								
8(a)(i)	 <p>circle or asterisk on correct C atom only [1] lines through the two correct bonds only [1]</p>	2								
8(a)(ii)	ketone, (tertiary) alcohol, alkene, carboxylic acid two for each mark	2								
8(a)(iii)	sp carbons = 0 sp ² carbons = 8 sp ³ carbons = 9	1								
8(a)(iv)		2								
8(b)(i)	<table border="1" data-bbox="349 1225 770 1428"> <thead> <tr> <th>compound</th> <th>spot</th> </tr> </thead> <tbody> <tr> <td>J</td> <td>2</td> </tr> <tr> <td>K</td> <td>3</td> </tr> <tr> <td>L</td> <td>1</td> </tr> </tbody> </table>	compound	spot	J	2	K	3	L	1	1
compound	spot									
J	2									
K	3									
L	1									

Question	Answer	Marks
8(b)(ii)	The more polar the compound and stronger attractive forces to the (polar) stationary phase ora: less polar compound and weaker attractive forces to the (polar) stationary phase	1
8(b)(iii)	R_f = retardation factor or retention factor or R_f = distance moved by compound from baseline over distance travelled by solvent front	1