#### UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS

GCE Advanced Level

## MARK SCHEME for the November 2005 question paper

# 9701 CHEMISTRY

9701/04

Paper 4 (Structured Questions A2 Core), maximum raw mark 60

This mark scheme is published as an aid to teachers and students, to indicate the requirements of the examination. It shows the basis on which Examiners were initially instructed to award marks. It does not indicate the details of the discussions that took place at an Examiners' meeting before marking began. Any substantial changes to the mark scheme that arose from these discussions will be recorded in the published *Report on the Examination*.

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Page 2	2		Mark Sche	me		Syllabus Paper	
		GCE	A LEVEL - Nov	vember 2005		9701 4	
1 (a)	M <sub>r</sub> (A	gBr) = 108 + 79.9	= 187.9				[1
	mole	$s = 2.5 \times 10^{-12}/187$					
	no. o	f ions = 1.33 x 10	$x^{-14} \times 6 \times 10^{23} = 3$	<b>3.0 x 10<sup>9</sup></b> ions	(correct	ans = [2])	[1
							2
(b)	(i)	<b>A</b> : platinum <b>B</b> : H⁺(aq) <i>or</i> HC <i>i</i> (ac (ignore concentration	q) <i>or</i> H <sub>2</sub> SO <sub>4</sub> (aq) on)	C: voltmeter D: silver (wire)	)		4 x [1
	(ii)	(As [Ag⁺] decrease	s), the potential	will decrease/be	come mor	e negative	[1
	(iii)	K <sub>sp</sub> = [Ag <sup>+</sup> ][Br <sup>-</sup> ]	$= (7.1 \times 10^{-7})^2$	= <b>5.0</b> (41) <b>x 10</b> <sup>-</sup>	<sup>13</sup> mol <sup>2</sup> dm	n <sup>-6</sup>	[1
						uni	<b>ts</b> [1
							7
(c)	(i)	Ag⁺(g) + Br⁻(g) —	→ AgBr(s)				[1
	(ii)	LE	=	$\Delta H_{ m f}$ - (all the r	est)		
			=	-100 – (731 +	285 + 112	- 325)	
			(=	-100 - 731 - 28	35 - 112 +	325)	
			=	<b>-903</b> kJ mol <sup>-1</sup>	(-[1] for e	each error of sign or maths)	[2
	(iii)	LE( <i>AgCl</i> ) should be due to size/radius o	e higher/more ne of C $l$ being less t	gative, than that of Br <sup>-</sup> (	both)		[1
							4
(d)	more	e energy needed, s	ince $r_{Cl} < r_{Br}$ or	ionised electron	nearer to	nucleus	
	or les	ss shielding etc. o	r in terms of I.E.	(C <i>l</i> ) > I.E.(Br)			

P	ade (	3	Mark Scheme	Syllabus	Paper				
-	aye	5	GCE A LEVEL – November 2005	9701	4				
2	(a)	The EMF of a cell made up of the test electrode and a standard hydrogen electrode. ( <i>or</i> the EMF of the electrode compared to the S.H.E.)							
		EMF measured under standard conditions of T, (P) and concentration. ( <i>or</i> at 298K and 1 mol dm <sup>-3</sup> )							
	(b)	The stronge	The stronger the halogen is as an oxidising agent, the more positive is its $E^{e}$ value.						
		Two examp	les of $F_2/F^-$ , $Cl_2/Cl$ ; $Br_2/Br^-$ , $I_2/I^-$ quoted						
		(data:	$F_2/F^- = +2.87V$ $Cl_2/Cl = +1.36V$ $Br_2/Br^- = +1.07V$						
			$I_2/I = +0.54V$ )						
	(c)	(i) or	$\begin{array}{rcl} H_2O_2 \ + \ 2I^{\scriptscriptstyle -} \ + \ 2H^{\scriptscriptstyle +} & \longrightarrow & I_2 \ + \ 2H_2O \\ H_2O_2 \ + \ 2KI \ + \ 2H^{\scriptscriptstyle +} & \longrightarrow & 2K^{\scriptscriptstyle +} \ + \ I_2 \ + \ 2H_2 \end{array}$	0					
				E <sup>e</sup> = 1.77 - 0.54 =	<b>1.23</b> ∨				
		(ii) or	$Cl_2 + SO_2 + 2H_2O \longrightarrow 2Cl^- + SO_4^{2-}$ $Cl_2 + SO_2 + 2H_2O \longrightarrow 2HCl + H_2SO_4$	+ 4H <sup>+</sup>					
				E <sup>e</sup> = 1.36 – 0.17 =	<sup>:</sup> 1.19 ∨				
	(d)	(d) since $E^{e}(I_2/I^{-})$ is +0.54V, tin will be oxidised to $Sn^{4+}$ ( $E^{e}$ for $Sn^{2+}/Sn = -0.14V$ and $E^{e}$ for $Sn^{4}/Sn^{2} = +0.15V$ )							
		Thus: S	$n + 2I_2 \longrightarrow SnI_4$						

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Page 4		Mark Scheme	Syllabus	Paper
3 (a) (	(i) r	nelting point: graph showing (Si (+ Ge): medium) and C: higher than Si/Ge Sn + Pb: lower than Si/Ge		-
	(	conductivity: graph showing (Si (+ Ge): medium) and C: lower (or higher!) than Si/G Sn + Pb: higher than Si/Ge	е	
	[	for your information, the actual figures are shown below]		
(	(ii) (ii) (ii) (ii) (ii) (ii) (ii) (ii)	Sn, Pb (and C(graphite)) have delocalised electrons/metallic Si, Ge (and C(diamond)) have localised electrons/covalent b [for [2] marks carbon has to be mentioned onc	bonds onds æ, and the allotrop	be mentioned
		must in in with the conductivity snownj		
(b) (	b) (i) e.g. CO burns to give $CO_2 [2CO + O_2 \longrightarrow 2CO_2]$ or CO reduces $Fe_2O_3 [3CO + Fe_2O_3 \longrightarrow 3CO_2 + 2Fe]$			
(	(ii) e	e.g. PbO₂ decomposes on heating [2PbO₂ → 2PbO + 0 two valid examples two balanced equati [two valid and balance]	D₂] ons ced equations war	[1] + rrants [3] mark
<b>(c)</b> ι	use:	oottery/china/porcelain etc + property: hardness, high meltin (any one use + one relevant property)	g point, insulator e	etc.
(d) (	(i) a	amphoteric		
(	(ii) e	e.g. SnO + 2HC $l \longrightarrow$ SnC $l_2$ + H <sub>2</sub> O		
	e	e.g. SnO + 2NaOH $\longrightarrow$ Na <sub>2</sub> SnO <sub>2</sub> + H <sub>2</sub> O		

### (Actual figures for (a) (i):)

element	m.pt./°C	conductivity
C(graph)	3652	2 x 10 <sup>3</sup>
C(dia)	3550	1 x 10 <sup>-15</sup>
Si	1410	2 x 10 <sup>-2</sup>
Ge	937	2 x 10 <sup>-2</sup>
Sn	232	9 x 10⁴
Pb	328	5 x 10⁴

Page 5			Mark S	cheme	Syllabus	Paper
		GCE	A LEVEL –	November 2005	9701	4
4 (a)	) HC ( <i>or</i>	$D-C_6H_4-NH_2 + 2AgBr C_6H_7NO)$	+ 20H <sup>-</sup> →	$O = C_6 H_4 = O + H_2 O + N_6 (or C_6 H_4 O_2)$	H <sub>3</sub> + 2Ag + 2Br <sup>-</sup>	[1
(b)	) roc	linol should be <b>less ba</b>	asic than N	H <sub>3</sub>		[1
	be	ecause the lone pair on N is delocalised over/overlaps with the aryl ring				
(c)	<b>) E</b> i	$H_2N-C_6H_4-O^-Na^+$	or	H <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -ONa		[1
	Fi	s HO-C <sub>6</sub> H <sub>4</sub> NH <sub>3</sub> <sup>+</sup> C $l$	or	HO-C <sub>6</sub> H <sub>4</sub> NH <sub>3</sub> Cl		[1
	G	s HO-C <sub>6</sub> H <sub>2</sub> Br <sub>2</sub> -NH <sub>2</sub>	up to	HO-C <sub>6</sub> Br <sub>4</sub> -NH <sub>2</sub> (ignore of	rientation)	[1
(d	) (i)	HNO <sub>3</sub> (aq) or dil HNO	<b>D</b> <sub>3</sub>	(NOT conc., and NOT +	conc. H <sub>2</sub> SO <sub>4</sub> )	[1
	(ii)	reduction				[1
	(iii	<b>)</b> Sn + HC <i>l</i> (aq)				[1
(e)	) (i)	phenol, amide				[1] + [1
	(ii)	$CH_3COCl$ or $(CH_3)$	CO) <sub>2</sub> O			[1
						total: 1

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I	Page	6	Mark Scheme Syllabus			Paper
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5	(a)	(i)	addition (polymerisation)			[1]
		(ii)	condensation (polymerisation)			[1]
						2
	(b)	hyd	ogen bonding			[1]
						1
	(c)	(i)	HO <sub>2</sub> CCH <sub>2</sub> CH <sub>2</sub> CO <sub>2</sub> H			[1]
		(ii)	ester (accep	t "covalent")		[1]
						2
	(d)	(i)	heat with $H_3O^+$ or heat with $OH^-(a_1)$	q)		[1]
		(ii)	H <sub>2</sub> N-CH <sub>2</sub> -CH(OH)-CH <sub>2</sub> -NH <sub>2</sub> o	$H_3N^+-CH_2-CH(OH)-CH_2-N_3N^+$	${\rm IH_3}^+$	[1]
			HO <sub>2</sub> C-CH(OH)-CH(OH)-CO <sub>2</sub> H or	<sup>-</sup> O <sub>2</sub> C-CH(OH)-CH(OH)-CC	$D_2^{-1}$	[1]
			(allow bonus mark if the acid/bashydrolysis)	se forms are consistent with	the reagent us	ed for the [1]
						4 max 3
	(e)	(i)	$NC-CH_2-CO_2^-K^+$			[1]
		(ii)	II: H <sub>2</sub> + Ni <i>or</i> Na in ethanol [allow	w LiA <i>1</i> H <sub>4</sub> ]		[1]
			<b>III:</b> dilute HCl or $H_2SO_4$ or $H^+(aq)$	)		[1]
						3