#### UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS

GCE Advanced Level and GCE Advanced Subsidiary Level

## MARK SCHEME for the May/June 2006 question paper

#### 9701 CHEMISTRY

9701/06

Paper 6

Maximum raw mark 40

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.

All Examiners are instructed that alternative correct answers and unexpected approaches in candidates' scripts must be given marks that fairly reflect the relevant knowledge and skills demonstrated.

Mark schemes must be read in conjunction with the question papers and the Report on the Examination.

The minimum marks in these components needed for various grades were previously published with these mark schemes, but are now instead included in the Report on the Examination for this session.

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Page 1	Mark Scheme	Sylla	er
-	GCE A/AS Level – May/June 2006	9701	

# Biochemistry

1	(a)	(i)	O CH <sub>3</sub> O		age.
			NH <sub>2</sub> – CH <sub>2</sub> – C- NH -CH – C-NH – CH <sub>2</sub> – COOH	[1]	
			Displayed structure	[1]	·
			One peptide linkage shown	[1]	
		(ii)	Condensation	[1]	[4]
	(b)	(i)	Weak intermolecular forces of attraction (1) Van der Waals (1)	[2]	
		(ii)	No attraction/ffinity for water	[1]	
			Non-polar structure	[1]	[4]
	(c)	(i)	Both contain the polyamide structure/-CONH-	[1]	
		(ii)	Bullet proof vests; body armour; ropes; airbags; kayaks; gloves; skis; run-flat tyres; shields for jet engines; helmets; racquets; clothing any <b>one</b>	e [1]	[2]
2	(a)	Dia	gram:	[1]	
2	(a)		gram: nd – van der Waals forces	[1] [1]	
2	(a)	Re			[3]
2	(a) (b)	R e	nd – van der Waals forces	[1] [1]	
2	•	R e	nd – van der Waals forces osphate end - ionic/polar van der Waals interactions/dipole –dipole interactions/temporary dipoles/hydrop	[1] [1] ohobic	
2	•	R e	ond – van der Waals forces osphate end - ionic/polar van der Waals interactions/dipole –dipole interactions/temporary dipoles/hydroginteractions	[1] [1] ohobic [1]	
2	(b)	R e Pho (i)	osphate end - ionic/polar van der Waals interactions/dipole –dipole interactions/temporary dipoles/hydroginteractions with the hydrocarbon part of the bilayer	[1] [1] phobic [1] [1]	;
2	(b)	R e Pho  (ii)  (ii)	osphate end - ionic/polar  van der Waals interactions/dipole –dipole interactions/temporary dipoles/hydrog interactions  with the hydrocarbon part of the bilayer  Disrupt it/distort it/weakens the interactions between the bilayers	[1] [1] phobic [1] [1]	;
2	(b)	RecPho(i) (ii) K+1	and – van der Waals forces  osphate end - ionic/polar  van der Waals interactions/dipole –dipole interactions/temporary dipoles/hydrog interactions  with the hydrocarbon part of the bilayer  Disrupt it/distort it/weakens the interactions between the bilayers  moves into cell, Na <sup>+</sup> moves out of cell	[1] [1] phobic [1] [1] [1]	;

Page 2	Mark Scheme	Sylla	er
	GCE A/AS Level – May/June 2006	9701	

#### **Environmental Chemistry**

- 3 (a)  $Cl_2 + H_2O \rightarrow HOCl + HCl/Cl_2 + H_2O == ClO^- + 2H^+ + Cl^-$ 
  - $\mathsf{HOC}\mathit{l}$  is an oxidising agent which kills bacteria/ $\mathsf{C}\mathit{l}\mathsf{O}^-$  is an oxidising agent which kills bacteria
  - (b) Dissolved chlorine will react with organic pollutants in water [1]
  - (c) (i) Water softener/removes magnesium or calcium ions from water (as insoluble magnesium or calcium phosphate) [1]
    - (ii) Phosphate encourages growth of algae and bacteria to form an algal bloom
       Algal bloom prevents sunlight from reaching plants lower down in water so they stop photosynthesising and die
       [1]
      - Bacteria feeding on dead plants multiply and their respiration uses up all the available dissolved oxygen
    - (iii)  $Al^{3+} + PO_4^{3-} \rightarrow AlPO_4$  [1]
      - Moles of A $lPO_4$  = 4.00/ecf from wrong formula or molar ratio

        Concentration = 0.004 mol dm<sup>-3</sup>/ecf from wrong number of moles

        [1]

        [7]

[1]

[1]

[1]

[1]

[3]

[3]

- 4 (a) (i)  $Si_2AlO_9^{7-}$  [1]
  - (ii) Cations can be adsorbed onto surface of clay/attraction between the negative clay and the cation [1]
    - Plants need certain cations such as potassium and by attraction to clay they cannot be washed out of soil easily
  - (b) (i) Any two from
    Hydrogen ions adsorbed onto surface of clay / attraction between hydrogen ion and negative clay
    - Some cations attached to clay can raise pH because cation is replaced by proton from water [1]
    - Other cations such as aluminium can lower pH when replaced by hydrogen ion from water [1]
    - (ii) If H<sup>+</sup> is low RCO<sub>2</sub>H dissociates to produce H<sup>+</sup>/dissociation equilibrium moves to the right [1]
      - If  $[H^{\dagger}]$  is high RCO<sub>2</sub>H forms RCO<sub>2</sub>H<sub>2</sub><sup> $\dagger$ </sup> [1] [4]
  - (c) Any three from
    - Liming involves adding calcium hydroxide or calcium carbonate to soil [1]
    - $OH^- + H^+ \rightarrow H_2O/CO_3^{2-} + H^+ \rightarrow HCO_3^-/more$  complex equations involving clay [1]
    - Can reduce nitrogen content of the soil/ $NH_4^+ + OH^- \rightarrow NH_3 + H_2O$  [1]
    - Efficiency of liming reduced by acid surges caused by melting of ice containing acidic water

		- A:	
Page 3	Mark Scheme	Sylla	er
	GCE A/AS Level – May/June 2006	9701	

#### Phase Equilibria

5 (a) (i) V.P. of A = vapour pressure of A on own x mol fr. of A

OR  $P_A = P_A x_A$ 

(ii) 
$$0.3 \times 48 = 14.4$$
  
 $0.7 \times 36 = \frac{25.2}{39.6 \text{ k Pa}}$ 

[1] [1]

(iii) Raoult's law obeyed

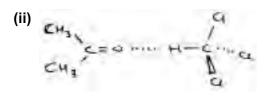
[1]

components are similar/ideal mixture/components not interact

[1] [5]

**(b) (i)** Molecules attract each other OR dipoles align Stronger intermolecular forces than components

[1]



OR Interact in 1:1 ratio

[1] [2]

(c) pure propanone

[1]

since this has lowest b.p. OR highest VP OR is most volatile

(allow discussion of b.p./composition curves)

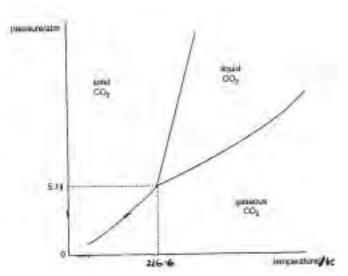
[1]

Then azeotrope or 0.50 composition

[1] [3]

Page 4		Mark Scheme	Sylla er
(a) preservation	GCE	A/AS Level – May/June 2006	9701 dhacambhidge.com





	both axes solid/liquid slope areas T.P. (1) values or label shape		[1] [1] [1] [1]	[5]
(b)	(i)	line drawn at 298K or indicated	[1]	
		value of 60 atm indicated	[1]	
		[Explanation without ref to diagram only scores [1]]		
	(ii)	CO <sub>2</sub> expands from over 60 atm to 1 atm cools	[1]	
		to below triplet point, explains solid	[1]	[4]
(c)	The wat	solid/liquid line has a positive slope for $\text{CO}_2$ rather than the negative slope of er	[1]	[1]

Page 5	Mark Scheme	Sylla	ęr
	GCE A/AS Level – May/June 2006	9701	

## **Spectroscopy**

7	(a)	Two	o absorptions	[1]	Toda
		Asy	mmetric bend (or diagram)	[1]	
		Asy	mmetric stretch (or diagram)	[1]	[3]
	(b)	(i)	1710 cm <sup>-1</sup> – C=O	[1]	
			2260 cm <sup>-1</sup> – C≡N	[1]	
			3200 cm <sup>-1</sup> – O–H	[1]	
		(ii)	NC-CH <sub>2</sub> -CH <sub>2</sub> -CO <sub>2</sub> H	[1]	[4]
	(c)	Nm	r	[1]	
		+ in	dication of absorptions (CH <sub>2</sub> $\sim$ 1.3 $\delta$ , -O-H $\sim$ 4.5 $\delta$ )	[2 x 1]	
		OR	Mass spectrometry	[1]	
			vo examples of likely fragmentations e.g. M-28 (loss of CN) and 7 (loss of –OH)	[2 x 1]	[3]
8	(a)	Stru	ucture II	[1]	
		Αp	eak at 3450 cm <sup>-1</sup> is characteristic of -OH would be seen for structure <b>II</b>	[1]	[2]
	(b)	(i)	Triplet-quartet is characteristic of a CH <sub>3</sub> next to CH <sub>2</sub> group	[1]	
			Standard 1,3,3,1 and 1,2,1 diagrams	[1]	
		(ii)	Singlet (1) at $\delta$ 2.0 – 3.8 (1)	[2]	
		(iii)	Deuterium oxide will exchange protons with -OH group in structure II	[1]	
			Since deuterium does not absorb in this part of the spectrum the –OH peak would disappear	[1]	[6]
	(c)	Stru	ucture <b>II</b> will show (M-17) <sup>+</sup> - loss of OH		
		Stru	ucture I will show (M-31) <sup>+</sup> loss of CH <sub>3</sub> O		

any **two [2]** 

Structure II will show  $(M-43)^{+}$  loss of  $C_3H_7$ 

Page 6	Mark Scheme	Sylla	er
	GCE A/AS Level – May/June 2006	9701	

### **Transitions Elements**

			Transitions Lientents	1	SA. 1
9	(a)	(i)	somewhere between 4% and 20% chromium	[1]	'do
		(ii)	Cr forms its oxide/Cr <sub>2</sub> O <sub>3</sub> on the steel's surface	[1]	
			which is impermeable to oxygen/hard	[1]	[3]
	(b)	(i)	Cr = 33.6/52 = 0.646 O = 20.6/16 = 1.288 Cl = 45.8/35.5 = 1.290	[1]	
			thus <b>A</b> is $CrO_2Cl_2$	[1]	
			O.N. of chromium = +6	[1]	
		(ii)	orange solution contains Cr <sub>2</sub> O <sub>7</sub> <sup>2-</sup>	[1]	
			$2CrO_2Cl_2 + 3H_2O \longrightarrow Cr_2O_7^{2-} + 6H^+ + 4Cl^-$	[1]	
			white ppte is $AgCl \ or \ Ag^+ + Cl^- \longrightarrow AgCl(s)$	[1]	
			yellow solution contains CrO <sub>4</sub> <sup>2-</sup>	[1]	
			$Cr_2O_7^{2-} + 2OH^- \longrightarrow 2CrO_4^{2-} + H_2O$	[1] [8 max 7]	[7]
10	(a)	col	our dues to absorption of visible light	[1]	
		d-o	rbitals are split into two sets at different energies	[1]	
		pho	oton is absorbed when an electron is promoted to higher orbital	[1]	[3]
	(b)	(i)	[Fe(SCN] <sup>2+</sup> is formed - this is red	[1]	
			F <sup>-</sup> is a stronger ligand than SCN <sup>-</sup> or ligand exchange occurs	[1]	
			[FeF <sub>6</sub> ] <sup>3-</sup> is colourless <i>or</i> energy gap between d-orbitals is large	[1]	
		(ii)	reduction occurs	[1]	
			to VO <sup>2+</sup> (which is blue)	[1]	
			$SO_2 + 4H^+ + 2VO_3^- \longrightarrow SO_4^{2-} + 2VO^{2+} + 2H_2O$	[1]	
			(further reduction to) V <sup>3+</sup> (which is green)	[1]	

[1] [8 max 7]

[7]

 $Sn^{2+} + 4H^+ + 2VO^{2+} \longrightarrow Sn^{4+} + 2V^{3+} + 2H_2O$