CAMBRIDGE INTERNATIONAL EXAMINATIONS

GCE Advanced Level

MARK SCHEME for the October/November 2013 series

9701 CHEMISTRY

9701/41

Paper 4 (A2 Structured Questions), maximum raw mark 100

This mark scheme is published as an aid to teachers and candidates, to indicate the requirements of the examination. It shows the basis on which Examiners were instructed to award marks. It does not indicate the details of the discussions that took place at an Examiners' meeting before marking began, which would have considered the acceptability of alternative answers.

Mark schemes should be read in conjunction with the question paper and the Principal Examiner Report for Teachers.

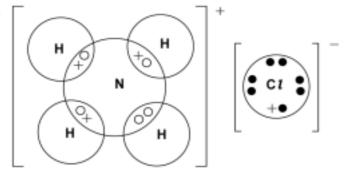
Cambridge will not enter into discussions about these mark schemes.

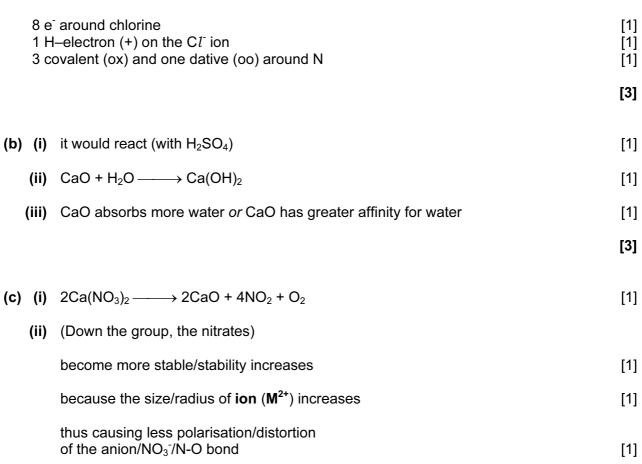
Cambridge is publishing the mark schemes for the October/November 2013 series for most IGCSE, GCE Advanced Level and Advanced Subsidiary Level components and some Ordinary Level components.



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1 (a)





[Total: 10]

[4]

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|--|----|

- 2 (a) (i) Si-Si bonds are weaker (than C-C bonds) [1]
 - (ii) metallic (Sn) is weaker than (giant) covalent (Ge) [1]

[2]

(b) (i)
$$SiCl_4 + 2H_2O \longrightarrow SiO_2 + 4HCl$$

 $or SiCl_4 + 4H_2O \longrightarrow Si(OH)_4 + 4HCl$
 $or SiCl_4 + 3H_2O \longrightarrow H_2SiO_3 + 4HCl$
(partial hydrolysis is *not sufficient* e.g. to $SiCl_3OH + HCl$) [1]

(ii)
$$PbCl_4 \longrightarrow PbCl_2 + Cl_2$$
 [1]

(iii)
$$SnCl_2 + 2FeCl_3 \longrightarrow SnCl_4 + 2FeCl_2$$
 [1]

(iv)
$$SnO_2 + 2NaOH \longrightarrow Na_2SnO_3 + H_2O$$

 $or SnO_2 + 2NaOH + 2H_2O \longrightarrow Na_2Sn(OH)_6$
 or ionic equation $SnO_2 + 2OH^- \longrightarrow SnO_3^{2-} + H_2O$ [1]

[4]

[Total: 6]

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|---------|--|---------------------------|----------|------------|
| (a) (i) | $NH_3 + HZ \longrightarrow NH_4^+ + Z^-$ $CH_3OH + HZ \longrightarrow CH_3OH_2^+ + Z^-$ | | | [1] [1] |
| (ii) | $NH_3 + B^- \longrightarrow NH_2^- + BH$ $CH_3OH + B^- \longrightarrow CH_3O^- + BH$ | | | [1] [1] |
| | | | | [4] |
| (b) (i) | a reaction that can go in either direction | | | [1] |
| (ii) | rate of forward = rate of backward reaction or forward/back reactions occurring but concentrations of all | l species do no | t change | [1] |
| | | | | [2] |
| (c) (i) | a solution that resists changes in pH | | | [1] |
| | when small quantities of acid or base/alkali are added | | | [1] |
| (ii) | in the equilibrium system HZ + $H_2O = Z^- + H_3O^+$ | | | [1] |
| | addition of acid: reaction moves to the left $or H^{+}$ combines with Z^{-} and forms HZ | | | [1] |
| | addition of base: the reaction moves to the right or H ⁺ combines with OH ⁻ and more Z ⁻ formed | | | [1] |
| | | | [5 ma | x 4] |
| (d) (i) | $[H^+] = \sqrt{(0.5 \times 1.34 \times 10^{-5})} = 2.59 \times 10^{-3} \text{ (mol dm}^{-3})$ | | | [1] |
| | pH = 2.59/2.6 (min 1 d.p) | | ecf | [1] |
| (ii) | $CH_3CH_2CO_2H + NaOH \longrightarrow CH_3CH_2CO_2Na + H_2O$ | | | [1] |
| (iii) | n(acid) in 100 cm ³ = $0.5 \times 100/1000 = 0.05$ mol n(acid) remaining = $0.05 - 0.03 = 0.02$ mol [acid remaining] = 0.2 (mol dm ⁻³) | | | [1] |
| | likewise, $n(salt) = 0.03 \text{ mol}$ [salt] + 0.3 (mol dm ⁻³) | | | [1] |
| (iv) | pH = 4.87 + log(0.3/0.2) = 5.04–5.05 | | ecf | [1] |
| | | | | [6] |
| (e) G | s CH ₃ CH ₂ COC <i>l</i> | | | |
| `´ H | s SOC l_2 or PC l_5 s NaC l | | | [2] |
| (or | corresponding Br compounds for \textbf{G},\textbf{H} and $\textbf{J};\text{CH}_3\text{CH}_2\text{COBr},$ | SOBr ₂ , NaBr) | [Total: | |
| | | | | |

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- 4 (a) (the energy change) when 1 mol of bonds [1]
- is broken in the gas phase [1]
 - **(b) (i)** (C-X bond energy) decreases/becomes weaker (from F to I) [1]
 - due to bond becoming longer/not such efficient orbital overlap [1]
 - (ii) (as the bond energy of C-X decreases) the halogenalkanes become more reactive (answer must imply that it is from F to I) [1]
 - (c) The C-Cl bond is weaker than the C-F <u>and</u> C-H bonds or C-Cl bond (E = 340) and C-H (E = 410) [1]
 - so is (easily) broken to form Cl^{\bullet}/Cl radicals/Cl atoms [1] causing the breakdown of O_3 into O_2 [1]
 - (d) C*l*-CH₂CH₂-CO₂H [1] HO-CH₂CH₂-C*l* [1]

- (e) (i) light/UV/hv *or* 300°C [1]
 - (ii) (free) radical substitution [1]

(iii)
$$\Delta H = E(C-H) - E(H-Cl) = 410 - 431 = -21 \text{ kJ mol}^{-1}$$
 [1]

(iv)
$$\Delta H = E(C-H) - E(H-I) = 410 - 299 = +111 \text{ kJ mol}^{-1}$$
 ecf [1]

(v) The reaction with iodine is endothermic $or \Delta H$ is positive or requires energy [1]

(vi)
$$Cl_2 \longrightarrow 2Cl^*$$
 [1]
 $CH_3CH_2^* + Cl_2 \longrightarrow CH_3CH_2Cl + Cl^*$ [1]
 $CH_3CH_2^* + Cl^* \longrightarrow CH_3CH_2Cl$ [1]

[Total: 19]

[8]

[2]

[3]

[3]

[3]

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5 (a) (i) many monomers form a polymer

[1]

- (ii) addition [1]
- (iii) C=C/double/ π bond is broken **and** new C-C single bond<u>s</u> are formed or double bond breaks and forms single bonds with other monomers

[1]

[3]

(b) propenoic acid [1]

[1]

(c) (i) CO₂Na CO₂Na

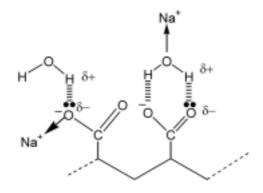
carbon chain **and** CO₂H [1]

at least one sodium salt [1]

(ii) 120° to 109(.5)° [1] due to the change from a trigonal/sp² carbon to a tetrahedral/sp³ carbon [1]

[4]

(d) (i)



Any four:

hydrogen bond labelled

water H-bonded to O through H atom $\delta+/\delta$ - shown on each end of a H-bond

lone pair shown on O⁻ or C=O or H₂O on a correct H-bond

Na⁺ shown as coordinated to a water molecule

[3]

(ii) Solution became paler **and** Cu⁽²⁺⁾ swapped with Na⁽⁺⁾ or darker in colour **and** polymer absorbs water

[1]

[4]

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| (e) (i) | alke | ne(1), amide(1) | | [2] |
| (ii) | NH ₃ | | | [1] |
| (iii) | H ₂ O | | | [1] |
| (iv) | | (aq)/H₃O⁺ and heat/reflux (not warm) H⁻ (aq), heat and acidify | | [1] |
| | 0, 0 | ir (aq), ricat and acidity | | [5] |
| | | | | [Total: 17] |

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Section B

6 (a) (i) **six/6** (gsv, sgv, gvs, vgs, svg, vsg)

[1]

(ii) H_3C CH_3 OH OH

(iii) valine (allow glycine) [1]

(iv) any two of: hydrogen bonds and CO_2H or OH or NH_2 or CONH or CO or NH or CO_2^- ionic bonds and NH_3^+ or CO_2^- van der Waals' and $-CH_3$ or -H

2 × [1]

[6]

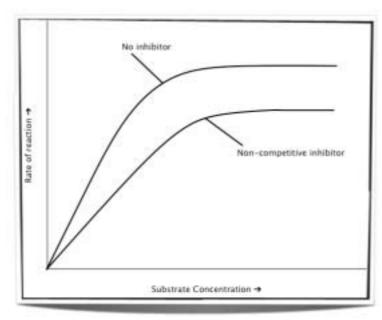
[1]

(b) (i) same shape/structure as substrate

(inhibitor) competes/blocks/binds/bonds to **active site**or substrate cannot bind to **active site**[1]

(ii) binds with enzyme and changes shape/3D structure (of enzyme/active site) [1]





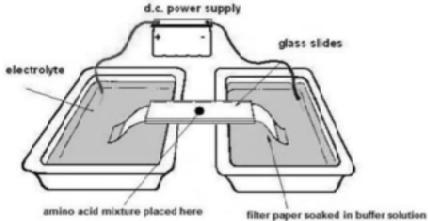
[1]

[4]

[Total: 10]

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7 (a)



power supply (idea of complete circuit) electrolyte/buffer solution gel/filter paper/absorbent paper (amino acid) sample/mixture [centre of plate]

4 × [1]

[4]

(b) any two from:

size/ $M_{\rm r}$ (of the amino acid species) charge (on the amino acid species) temperature

2 × [1]

[2]

(c) Ratio of the <u>concentration</u> of a solute in each of two (immiscible) solvents or equilibrium constant representing the distribution of a solute between two solvents or $PC = [X]_a/[X]_b$ (at a constant temperature)

[1]

[1]

(d) (i) $K_{pc} = [Z \text{ in ether}]/[Z \text{ in H}_2O] - \text{allow reverse ratio}$ 40 = (x/0.05)/((4-x)/0.5)

[1]

= 3.2 g

ecf [1]

(ii) First extraction 40 = (x/0.025)/((4-x)/0.5)x = 2.67 g

ecf [1]

(iii) Second extraction: 1.33g remain in solution Second extraction 40 = (y/0.025)/((1.33-y)/0.5) y = **0.887** g

mass extracted = 2.67 + 0.89 = 3.56/3.6 g

ecf [1]

[4]

[Total: 11]

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- 8 (a) (i) (nitrates are) soluble
 - (ii) $Ba^{(2+)}$ and $Pb^{(2+)}$ [1]
 - SO₄⁽²⁻⁾
 - BaCO₃/PbCO₃/CaSO₄ are insoluble [1]
 - [4]
 - (b) (i) fertilisers/animal manure [1]
 - (ii) washing powder/detergents/fertilisers/animal manure [1]
 - (iii) growth/production of algae/weeds/plants or eutrophication [1]
 - (c) (i) any one of:

$$2SO_2 + O_2 \longrightarrow 2SO_3$$
 and $SO_3 + H_2O \longrightarrow H_2SO_4$

or
$$SO_2 + NO_2 \longrightarrow SO_3 + NO$$
 and $SO_3 + H_2O \longrightarrow H_2SO_4$

or
$$SO_2 + \frac{1}{2}O_2 + H_2O \longrightarrow H_2SO_4$$
 [1]

(ii) roasting sulfide ores/extraction of metals from sulfide ores [1]

[2]

[3]

[1]

[Total: 9]