

CAMBRIDGE INTERNATIONAL EXAMINATIONS

GCE Advanced Level

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MARK SCHEME for the May/June 2014 series

9701 CHEMISTRY

9701/42

Paper 4 (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.

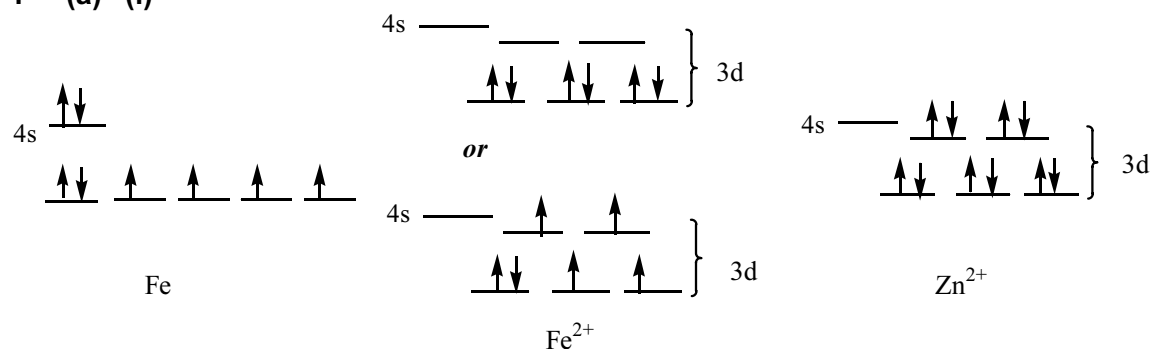
Mark schemes should be read in conjunction with the question paper and the Principal Examiner Report for Teachers.

Cambridge will not enter into discussions about these mark schemes.

Cambridge is publishing the mark schemes for the May/June 2014 series for most IGCSE, GCE Advanced Level and Advanced Subsidiary Level components and some Ordinary Level components.

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1 (a) (i)



[2]

(ii) (colour due to absorbance of visible light)
due to **electron** promoted (from lower) to upper **orbital/energy level**

[1]

in Zn²⁺ there's no space in higher orbital for the electron to go *or* completely filled **d**-orbitals/shell

[1]

4

(b) (i) yellow is due to [CuCl₄]²⁻

[1]

reaction is ligand displacement/exchange

[1]

(ii) (solution goes blue) due to [Cu(H₂O)₆]²⁺

[1]

blue **ppt. or (s)**

[1]

of Cu(OH)₂ *or* [Cu(H₂O)₄(OH)₂] etc.

[1]

purple *or* deep/dark blue **solution or (aq)**

[1]

due to [Cu(NH₃)₄]²⁺ *or* [Cu(NH₃)₄(H₂O)₂]²⁺

[1]

7

(c) (i) 2KI + K₂S₂O₈ → 2K₂SO₄ + I₂ *or*
ionic: 2I⁻ + S₂O₈²⁻ → 2SO₄²⁻ + I₂

[1]

(ii) Fe²⁺ is a **homogeneous catalyst**

[1]

(iii) equations: 2Fe²⁺ + S₂O₈²⁻ → 2Fe³⁺ + 2SO₄²⁻
2Fe³⁺ + 2I⁻ → 2Fe²⁺ + I₂

or verbal equivalent, e.g. reactants are both negative ions, so repel each other *or* Fe²⁺ can be oxidised by S₂O₈²⁻ **and** Fe³⁺ can be reduced by I⁻

[1]

3

[Total: 14]

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- 2 (a) **A:** voltmeter *or* V *or* potentiometer [1]
- B:** platinum *or* Pt [1]
- C:** 1 mol dm⁻³ **and** H⁺ *or* HCl (*or* 0.5 M H₂SO₄) [1]
- D:** lead (metal) *or* Pb [1]
- 4**
- (b) (i) a ✓ in the box next to **-0.17 V** [1]
a comment that the [Pb²⁺] has decreased **plus** a description of the outcome, e.g. as [Pb²⁺] decreases (from 1 mol dm⁻³), Pb²⁺(aq) + 2e⁻ ⇌ Pb(s) goes over to the left hand side, *or* as [Pb²⁺] decreases, Pb²⁺ is less likely to be reduced [1]
- (ii) (K_{sp} =) [Pb²⁺][Cl⁻]² [1]
- (iii) if [PbCl₂] = 3.5 × 10⁻², [Pb²⁺] = 3.5 × 10⁻² and [Cl⁻] = 7.0 × 10⁻²
so K_{sp} = (3.5 × 10⁻²) × (7.0 × 10⁻²)² = **1.715 (1.7) × 10⁻⁴ mol³ dm⁻⁹** (≥2sf) [1] + [1]
- 5**
- (c) (i) the (M²⁺/M) E^o for the two elements are very similar *or* are -0.13 and -0.14 V [1]
E^o (Sn⁴⁺/Sn²⁺) = 0.15 V **and** E^o (Pb⁴⁺/Pb²⁺) = 1.69 V [1]
so Sn²⁺ is quite easily oxidised (to Sn⁴⁺) **or** is a stronger reductant **or** Pb²⁺ is not easily oxidised (to Pb⁴⁺) **or** Pb⁴⁺ is a stronger oxidant **or** Pb⁴⁺ is easily reduced [1]
- (ii) e.g. PbCl₂ + Zn → Pb + ZnCl₂ (*or* ionic) [1]
(other acceptable reductants: Fe, Mg, Ca but **not** Na or K)
- Sn²⁺ + Br₂ → Sn⁴⁺ + 2Br⁻ [1]
(other acceptable oxidants: VO²⁺, Cr₂O₇²⁻, Ag⁺, Cl₂, Br₂, F₂, Fe³⁺, MnO₄⁻)
- 5**
- (d) (i) Pb²⁺(g) + 2Cl⁻(g) → PbCl₂(s) [1]
- (ii) ΔH_f = ΔH_{at} + E(Cl - Cl) + 1st IE + 2nd IE + 2 × E_A(Cl) + LE
-359 = 195 + 242 + 716 + 1450 - 2 × 349 + LE
LE = 2 × 349 - 359 - 195 - 242 - 716 - 1450
LE = **-2264** (kJ mol⁻¹) [3]
- (iii) LE(PbCl₂) > LE(PbBr₂) *or* more exothermic *or* stronger lattice [1]
because Cl⁻/chloride anion has smaller radius/size than Br⁻/bromide [1]

6

[Total: 20]

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- 3 (a) (i) B and D [1] + [1]
- (ii) D [1]
- 3**
- (b) heat with dilute $\text{H}^+(\text{aq})$ or $\text{H}_2\text{SO}_4(\text{aq})$ [1]
- 1**
- (c) (i) K_a larger than that for ethanol because
the ethanoate ion/ CH_3CO_2^- is stabilised by charge delocalisation
or
the O–H bond is weakened due to its proximity to C=O/carbonyl group *or*
the second electronegative/oxygen atom [1]
- K_a smaller than that for chloroethanoic acid because
electron-withdrawing/electronegative chlorine (atom) makes the anion more
stable *or* O–H bond weaker *or* H more easily lost [1]
- (ii) $[\text{H}^+] = \sqrt{([\text{CH}_3\text{CO}_2\text{H}] \times K_a)} = \sqrt{(0.1 \times 1.75 \times 10^{-5})} = \mathbf{1.32(3) \times 10^{-3}}$ (mol dm^{-3}) [1]
- $\text{pH} = -\log_{10}[\text{H}^+] = \mathbf{2.88}$ (2.9) [1]
- 4**
- (d) (i) $n(\text{NaOH})$ at start = $0.1 \times 20/1000 = 2.0 \times 10^{-3}$ mol
 $n(\text{NaOH})$ at finish = $\mathbf{1.0 \times 10^{-3}}$ mol [1]
- (ii) this is in 30 cm^3 of solution,
so $[\text{NaOH}]$ at finish = $1.0 \times 10^{-3}/0.030 = \mathbf{3.3(3) \times 10^{-2}}$ mol dm^{-3} (≥ 2 s.f.) ecf
from (i) [1]
- (iii) $[\text{H}^+] = K_w/[\text{OH}^-] = 1 \times 10^{-14}/3.33 \times 10^{-2} = 3.0 \times 10^{-13}$ mol dm^{-3}
 $\text{pH} = -\log_{10}[\text{H}^+] = \mathbf{12.5(2)}$ [1]
- or* $\text{pOH} = -\log_{10}(3.33 \times 10^{-2}) = 1.48$
 $\text{pH} = \text{p}K_w - \text{pOH} = 14 - 1.48 = \mathbf{12.5(2)}$ [1]
- (iv) pH/vol curve: start at pH 2.88 (2.9) ecf [1]
- vertical (over at least 2 pH units) portion at $V = 10 \text{ cm}^3$ [1]
- levels off at pH 12.5 ± 0.3 ecf [1]
- (v) indicator is thymolphthalein [1]
- 7**
- [Total: 15]**

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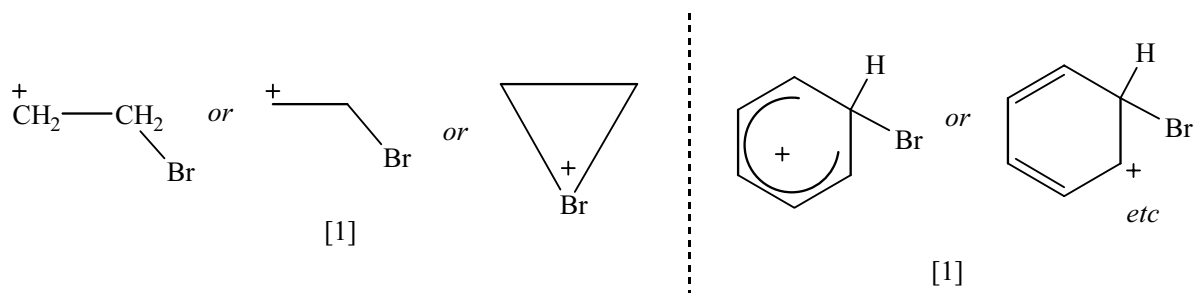
4. (a) (i) addition
AND
(ii) substitution [1]

1



1

- (c) (i) The two intermediate cations:



- (ii) The ring (of π electrons) in benzene is a stable configuration or is unchanged after the reaction. [1]

3

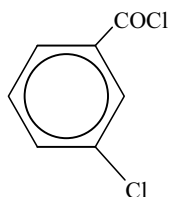
- (d) E is benzoic acid [1]

reaction 1: heat with KMnO_4 (+ OH^- or H^+) [1]

reaction 2: heat with $\text{Cl}_2 + \text{AlCl}_3$ or FeCl_3 [1]

3

- (e) G is [1]



reaction 3: SOCl_2 or PCl_5 [1]

reaction 4: LiAlH_4 [1]

3

[Total: 11]

Page 6	Mark Scheme	Syllabus	Paper
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5. (a) (i) Na reacts with –OH or hydroxyl/ alcohol groups [1]

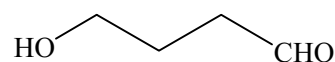
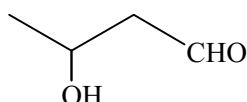
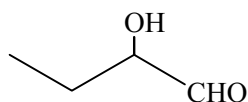
(ii) Fehling's solution reacts with –CHO or aldehyde groups [1]

2

(b) alkene or C=C or carbon double bond or phenol or phenylamine [1]

1

(c) $\text{CH}_3\text{CH}_2\text{CH}(\text{OH})\text{CHO}$ $\text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{CHO}$ $\text{HOCH}_2\text{CH}_2\text{CH}_2\text{CHO}$



[1] + [1] + [1]

3

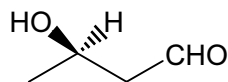
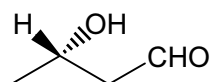
(d) (i) the $\text{CH}_3\text{CH}(\text{OH})$ group or the CH_3CO group or methyl secondary alcohol or methyl ketone [1]

(ii) $\text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{CHO}$ [1]

2

(e) (i) optical isomerism [1]

(ii)



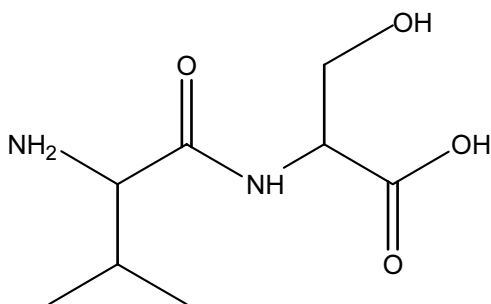
[1]

2

[Total: 10]

Section B

6. (a) (i)



Peptide bond correct [1]
 Rest of structure correct (skeletal, displayed or structural formula, or a mix)

(ii) Condensation *or* nucleophilic substitution *or* addition-elimination [1]

(iii) Water/H₂O [1]

4

(b)

DNA	RNA
Contains deoxyribose	Contains ribose
Contains thymine/T	Contains uracil/U
Double strand/chain/helix <i>or</i> two strands	Single strand/chain

[3]

3

(c) (i) (met) - leu - thr - pro - glu [1]

(ii) Mutations *or* addition/insertion/deletion/substitution/replacement (of a base) [1]

(iii) Changing A (*or* the 14th base) into U [1]

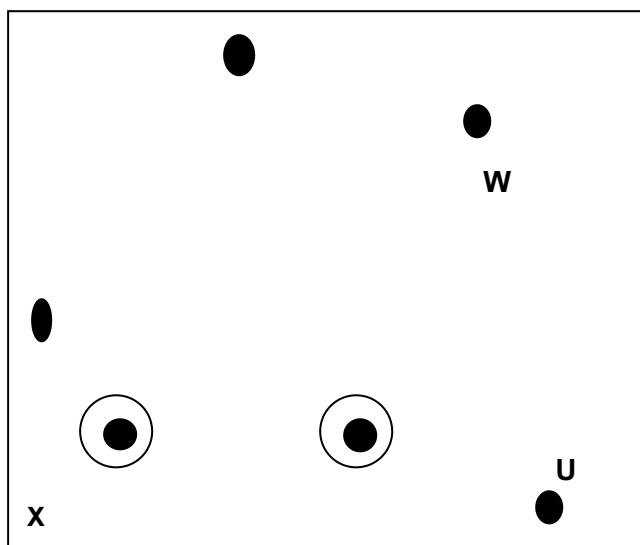
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[Total: 10]

Page 8	Mark Scheme	Syllabus	Paper
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- 7 (a) (i) (Electrophoresis): the size / shape / M_r of the amino acid *or* its charge [1]
- (ii) (Paper chromatography): the partition of the amino acid between, *or* the relative solubility of the compound in, the 2 phases *or* solvent/water and stationary phase/filter paper. [1]
- 2**
- (b) Use ninhydrin as a locating agent [1]
- 1**
- (c) The R_f value *or* retardation/retention factor *or* the distance travelled by the acid compared to that travelled by a standard sample of the amino acid [1]
- 1**
- (d) **R** – glutamic acid; **S** – glycine; **T** – lysine $3 \times [1]$
- 3**

(e)



$3 \times [1]$

3

[Total: 10]

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8. (a) (i) Any addition polymer (e.g. polyethene, polypropene, polystyrene, PVC, PTFE, PVA, *Teflon*) [1]

(ii) Any condensation polymer (e.g. polyamide, polyester, nylon, *Terylene*, PET, PLA, *Kevlar*, *Nomex*) [1]

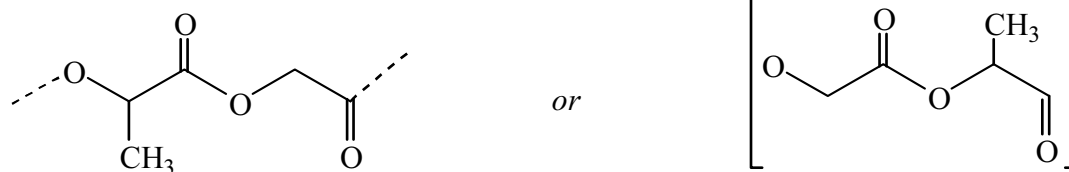
2

(b) Hydrolysis or nucleophilic substitution [1]

Ester and amide/peptide or $-\text{CO}_2-$ and $-\text{CONH}-$ [1]

2

(c)



Correct ester linkage [1]

CH_3 side chain on only one monomer unit [1]

2

(d) Plant materials do not generally contain unsaturated hydrocarbons/alkenes/
 $\text{C}=\text{C}$ [1]

1

(e) (i) Y van der Waals' forces [1]

Z hydrogen bonding [1]

(ii) Z, because it can form hydrogen bonds with water or it contains polar CO
and NH groups [1]

3

[Total: 10]