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UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS GCE Advanced Level

MARK SCHEME for the May/June 2011 question paper for the guidance of teachers

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.

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1 (a) N≡N triple bond is (very) strongor the N₂ molecule has no polarity

[1]

(b)
$$3Mg(s) \rightarrow 3Mg^{2+}(g)$$
 $\Delta H_1 = 3 \times 148 + 3 \times 2186 = 7002$ $N_2(g) \rightarrow 2N^{3-}(g)$ $\Delta H_2 = 994 + 2 \times 2148 = 5290$

LE =
$$-\Delta H_1 - \Delta H_2 - 461$$
 = **-12,753** (kJ mol⁻¹)

(-[1] for each error) [3]

(c) (i)
$$\text{Li}_3\text{N} + 3\text{H}_2\text{O} \rightarrow \text{NH}_3 + 3\text{LiOH}$$
 (balanced equation) [1]

(ii) advantage: no high pressure/temperature/catalyst needed/standard conditions used [1]

disadvantage: Li is expensive

or Li would need to be recycled/removedor LiOH by-product is corrosive/strongly basic

or this would be a batch, rather than continuous process [1]

(d) (i)
$$\text{Li}_3\text{N}: 100 \times 14/35 = 40\% \text{ N}$$
 [1] urea: $100 \times 28/60 = 47\% \text{ N}$ [1]

(iii)
$$NH_2CONH_2 + H_2O \rightarrow 2NH_3 + CO_2$$

 $or \rightarrow NH_2CO_2H + NH_3$
 $or NH_2CONH_2 + 2H_2O \rightarrow 2NH_3 + H_2CO_3$ [1]

(iv) The LiOH would be strongly alkaline
 or would increase the pH of the soil
 or would 'burn' the crops/reduce plant growth/stunt plants
 or would contaminate the environment

[Total: 12]

[1]

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- 2 (a) (i) One that can go in either direction.
 - (ii) both forward & reverse reactions are going on at the same time, but the concentrations of all species do not change (owtte)

 or rate of forward = rate of backward reaction

 [1]

(b) (i)
$$K_c = [H^{\dagger}][OH^{-}]/[H_2O]$$
 [1]

(ii)
$$K_w = [H^+][OH^-]$$
 [1] rearrangement of equation in (i) gives $K_c[H_2O] = [H^+][OH^-] \& K_w = K_c[H_2O]$ (owtte)
or the $[H_2O]$ is contained within K_w [1]

(iii) K_w will be higher in hot water **because** reaction is endothermic [1]

(c) (i)
$$[OH^-] = 5 \times 10^{-2}$$
; $[H^+] = (1 \times 10^{-14})/5 \times 10^{-2} = 2 \times 10^{-13}$ [1] $pH = -log_{10}[H^+] = 12.7$ (correct ans = [2]) ecf [1]

(ii)
$$[NH_4^+] = [OH^-] (= x)$$
 [1] $x^2 = 1.8 \times 10^{-5} \times 0.05 \implies x (= [OH^-]) = 9.49 \times 10^{-4} \text{ (mol dm}^{-3})$ (correct ans = [2]) [1]

(iii)
$$[H^{+}] = K_w/[OH^{-}] = (1 \times 10^{-14})/9.49 \times 10^{-4} = 1.05 \times 10^{-11} (\text{mol dm}^{-3})$$
 ecf [1]

[Total: 12 max 11]

[1]

(b)
$$PCl_5$$
 fizzes or white/misty fumes or heat evolved [1] $PCl_5 + 4H_2O \rightarrow H_3PO_4 + 5HCl$ or $PCl_5 + 3H_2O \rightarrow HPO_3 + 5HCl$ (allow partial hydrolysis: $PCl_5 + H_2O \rightarrow POCl_3 + 2HCl$) [1]

(c) (i)
$$P = 30.4/31 = 0.98$$
 $Cl = 69.6/35.5 = 1.96$ [1] Thus E.F = $PC l_2$

$$M_r(PCl_2) = 102$$
, so $2 \times PCl_2 = 204 \approx 200$, so M.F. = P_2Cl_4 [1]

(iii) O.N. =
$$(+)2$$

(iv)
$$(HO)_2P-P(OH)_2$$
 or $H(HO)P(=O)-P(=O)(OH)H$ ecf from structure in (ii) [1] Allow HO-P-OH or HO-P=O H

[Total: 10]

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4 (a)
$$N_2 + 2O_2 \rightarrow 2NO_2$$
 (or via NO) or $2NO + O_2 \rightarrow 2NO_2$ [1]

(ii)
$$NO_2 + 2CO \rightarrow \frac{1}{2} N_2 + 2CO_2$$
 or similar
Allow $2NO_2 + CH_4 \rightarrow CO_2 + N_2 + 2H_2O$ [1]

Allow formed from N₂ and O₂ in air during combustion

(ii) NO +
$$\frac{1}{2}$$
 O₂ \rightarrow NO₂ [1]

(iii)
$$K_p = (p_{NO}.p_{SO_3})/(p_{NO_2}.p_{SO_2})$$
 [1] units: dimensionless/none (don't accept just a blank!)

(iv)
$$K_p = 99.8^2/0.2^2 = 2.5 \times 10^5$$
 [1]

[Total: 11]

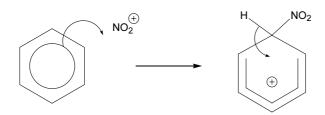
5 (a)

transformation	reagent + conditions
$C_2H_4 \rightarrow C_2H_5Cl$	HC <i>I</i> , no light or catalyst
$C_2H_5OH \rightarrow C_2H_5Cl$	conc $HCl + ZnCl_2$ or $SOCl_2$ or PCl_5 or PCl_3 and heat
$C_2H_6 \rightarrow C_2H_5Cl$	C 1 ₂ + light
$C_2H_4 \rightarrow C_2H_4Cl_2$	C l ₂ , no light or catalyst
CH₃CO₂H → CH₃COC <i>l</i>	SOC l_2 or PC l_5 or PC l_3 and heat
H ₃ C — H ₃ C — Cl	C l ₂ + A lC l ₃
— CH ₂ C1	C 1 ₂ + light or heat

[6]

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(b) (i) production of
$$NO_2^+$$
: $2H_2SO_4 + HNO_3 \rightarrow 2HSO_4^- + H_3O^+ + NO_2^+$ [1] (accept $H_2SO_4 + HNO_3 \rightarrow HSO_4^- + H_2O + NO_2^+$)



curly arrow from ring to NO_2^+ and from C-H bond to ring [1] correct intermediate, including charge in the right place

Note charge area must be more than half ring [1]

(ii)
$$\mathbf{C}$$
 is $C_6H_5CO_2H$ [1]

(iii) step 1: reagent is hot acidified or alkaline
$$KMnO_4$$
 [1] step 2: reagent is $Br_2 + FeBr_3/A_1C_1$ etc. (H_2O or light negates) [1]

(If ${\bf C}$ is given as 3-bromotoluene, then allow the last [2] marks if steps 1 and 2 are reversed.)

[Total: 12]

6 (a) (i) aqueous alkaline iodine
$$or I_2 + OH^-(aq)$$
 allow NaC1O + KI [1]

(ii)
$$CH_3CO$$
- or $CH_3CH(OH)$ - [1]

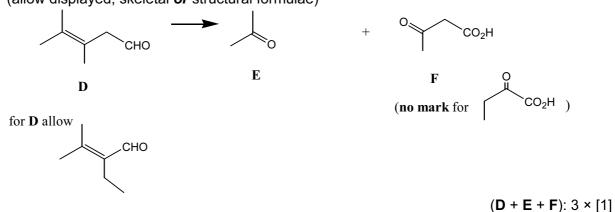
(iv)

compound	result
CH₃OH	×
CH₃CH₂OH	✓
CH₃CHO	✓
CH₃CO₂H	×
СНО	×
СОСН₃	√

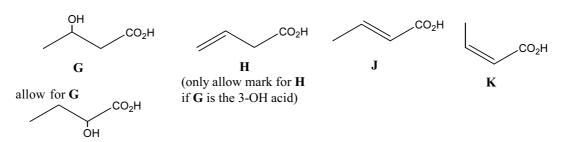
• √ • √ • √ [3]

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(b) (allow displayed, skeletal *or* structural formulae)



(c) (allow displayed, skeletal and structural formulae) Must be consistent with F



(N.B. letters **H**, **J**, **K** can be swapped around)

 $(G + H + J + K): 4 \times [1]$

geometrical or cis-trans isomerism

[1]

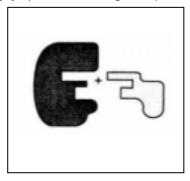
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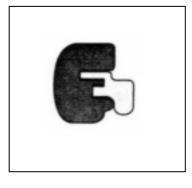
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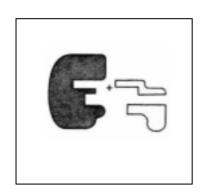
7 (a) The tertiary/3-dimensional structure/shape is held together by hydrogen/ionic/van der Waals bonds [1]

These break (relatively) easily/are weak/break at/above 45 °C [1]

(b) (or similar diagrams)







Enzyme + substrate

Enzyme-substrate complex

Enzyme + products

3 × [1]

(c) a competitive inhibitor combines with the enzyme's active site (so preventing the substrate from binding) [1]

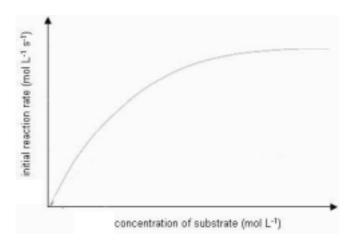
non-competitive inhibitor bonds with the enzyme away from the active site/at an allosteric site [1]

this changes the shape of the active site

[1]

Also allow competitive inhibition can be overcome by increasing [substrate] **or** non-competitive inhibition cannot be removed by increasing [substrate] for the 3rd mark

(d) (i)



Line must be of similar shape to original but level out below original line

[1]

(ii) Inhibitor reduces the number of enzymes with 'working' active sites (owtte)

[1]

[Total: 10]

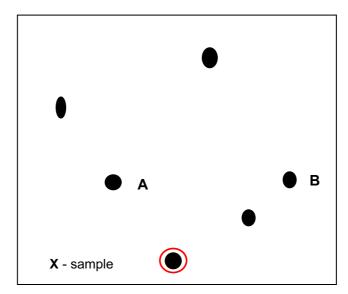
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8 (a) partition – separation due to the different solubilities of compounds in two solvents/phases

[1]

adsorption – separation due to the different attractions between the compounds and thestationary phase, relative to their solubility in the solvent[1]Note, if candidates do not refer to different solubilities and different attractionsmax 1

(b)



Ring: [1]

A + B: [1]

(c) (i) X is bromine – M and (M+2) peaks almost same height [1]

(ii)
$$\frac{M}{M+1} = \frac{100}{1.1} \times \frac{9}{n} = \frac{100}{0.3}$$
 1.1 × n

Hence
$$n = \frac{100 \times 0.3}{1.1 \times 9} = 3.03$$
 $p = 3$

(answer + working) [1]

(If the mass peak is at 122 and the compound contains Br and 3 C atoms then Q = (122 - 79 - 36)) thus Q = 7 ecf from (ii) [1]

(The compound is C_3H_7Br)

(iii) (R is at m/e 43), hence
$$C_3H_7^+$$
 [1]

(d) Any two from
$$H_2$$
, H_2O , CO , C_2H_4 , C_2H_2 , CH_4 $2 \times [1]$

[Total: 10]

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9 (a) (i) One [1]

(ii) Any alkene (or allow a cyclic amide, as in caprolactam) [1]

(b) Any TWO from: addition needs unsaturated/double bonds/alkene

condensation eliminates a small molecule

condensation needs a molecule other than a hydrocarbon

empirical formula of addition polymer is the same as that of its monomer

condensation needs two different functional groups

(**NOT** – "condensation needs two different monomers") 2 × [1]

(ii) O—CH₂

'sticks' to rest of molecule

Note: candidates need only show 'brackets' if more than one repeat unit shown

(iii) Polyesters [1]

(d) Monomers in *Terylene* have to alternate in order to condense out water (owtte) [1]

Alkenes can link in any order (and still form a polyalkene) (or diagram showing this) [1]

[Total: 10]

[1]