

**MARK SCHEME for the October/November 2011 question paper  
for the guidance of teachers**

**9701 CHEMISTRY**

**9701/43**

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)  $\text{Cr}^{3+}$ :  $1s^2 2s^2 2p^6 3s^2 3p^6 3d^3$  [1]  
 $\text{Mn}^{2+}$ :  $1s^2 2s^2 2p^6 3s^2 3p^6 3d^5$  [1]  
[2]

- (b) (i) Any two from  
•  $\text{H}^+$  is on the oxidant/L.H. side of each of the  $\frac{1}{2}$ -equations, *or*  $\text{H}^+$  is a reactant  
• (increasing  $[\text{H}^+]$ ) will make  $E^\ominus$  more positive  
• (increasing  $[\text{H}^+]$ ) will drive the reaction over to the R.H./reductant side *or* forward direction  
[1] + [1]

- (ii)  $\text{KMnO}_4$ : Purple/violet to colourless (allow very pale pink) [1]  
 $\text{K}_2\text{Cr}_2\text{O}_7$  Orange to green [1]  
[4]

- (c) (i)  $\text{MnO}_2 + \text{SO}_2 \longrightarrow \text{MnSO}_4$  (*or*  $\text{Mn}^{2+} + \text{SO}_4^{2-}$ ) [1]  
manganese changes/is reduced from +4 to +2 [1]  
sulfur changes/is oxidised from +4 to +6 [1]

- (ii) **No effect**, because  $\text{H}^+$  does not appear in the overall equation *or* its effect on the  $\text{MnO}_2/\text{Mn}^{2+}$  change is cancelled out by its effect on the  $\text{SO}_2/\text{SO}_4^{2-}$  change [1]  
[4]

- (d) (i)  $\text{MnO}_2 + 4\text{H}^+ + \text{Sn}^{2+} \longrightarrow \text{Mn}^{2+} + 2\text{H}_2\text{O} + \text{Sn}^{4+}$  [1]

- (ii)  $n(\text{MnO}_2) = 0.02 \times 18.1/1000 = 3.62 \times 10^{-4} \text{ mol}$  [1]  
 $n(\text{Sn}^{2+}) = 3.62 \times 10^{-4} \times 5/2 = 9.05 \times 10^{-4} \text{ mol}$  [1]  
 $n(\text{Sn}^{2+})$  that reacted with  $\text{MnO}_2 = (20 - 9.05) \times 10^{-4} = 1.095 \times 10^{-3} \text{ mol}$  [1]  
reaction is 1:1, so this is also  $n(\text{MnO}_2)$   
mass of  $\text{MnO}_2 = 1.095 \times 10^{-3} \times (54.9 + 16 + 16) = 0.0952 \text{ g}$  [1]  
 $\Rightarrow$  **95% – 96%**; 2 or more s.f. [1]  
[6]

[Total: 16]

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- 2 (a) (i) A molecule/ion/species with a lone pair (of electrons) or electron pair donor...  
.... that bonds to a metal ion/transition element.... [1]
- (ii) ...by means of a dative/coordinate (covalent) bond [1]  
[2]
- (b) (i) straight line from (0, 0.01) to point at (350, 0.0028) with all points on the line [1]
- (ii) order w.r.t.  $\text{Cr}(\text{CO})_6$  is 1 **and** order w.r.t.  $\text{PR}_3$  is zero [1]  
because (a)  $\text{Cr}(\text{CO})_6$  graph has a constant half-life (which is 700 s)  
or construction lines on graph showing this) [1]  
because (b)  $\text{PR}_3$  graph is a straight line (of constant slope) or line shows a constant rate  
of reaction or no change in rate or shows a linear decrease [1]
- (iii) rate =  $k[\text{Cr}(\text{CO})_6]$  [1]  
 $k = (0.9 - 1.1) \times 10^{-3} \text{ (s}^{-1}\text{)}$  (one or more s.f.) [1]  
*either*  $\text{rate}_0 = 0.01/1020 = 9.8 \times 10^{-6} \text{ mol sec}^{-1}$  when  $[\text{Cr}(\text{CO})_6] = 0.01 \text{ mol dm}^{-3}$   
so  $k = 9.8 \times 10^{-6}/0.01 = 9.8 \times 10^{-4}$   
or  $t_{1/2} \approx 700 \text{ sec}$   
 $k = 0.693/700 = 9.9 \times 10^{-4}$
- (iv) (units of k are)  $\text{sec}^{-1}$  [1]
- (v) N.B. the chosen mechanism must be consistent with the rate equation in (iii). Thus:  
*either* if rate =  $k[\text{Cr}(\text{CO})_6]$   
mechanism **B** is consistent [1]  
because it's the only mechanism that does NOT involve  $\text{PR}_3$  in its slow/rate-determining  
step or only  $\text{Cr}(\text{CO})_6$  is involved in slow step or  $[\text{PR}_3]$  does not affect the rate [1]
- or
- if rate =  $k[\text{Cr}(\text{CO})_6][\text{PR}_3]$ , then  
mechanism **A** or **C** or **D** is consistent [1]  
because both reactants are involved in slow step [1]  
[9]

[Total: 11]

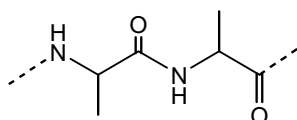
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3 (a) (i) E is  $\text{CH}_3\text{CH}(\text{NH}_2)\text{CN}$  [1]

(ii)  $\text{C}_6\text{H}_5\text{CH}_2\text{CHO}$  [1]  
[2]

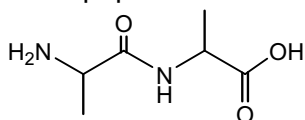
(b) (i) a polymer/polypeptide of amino acids, (joined by peptide bonds)  
(allow 'chain' of amino acids' but not 'sequence': the idea of 'many' has to be conveyed) [1]

(ii)



peptide bond shown in full ( $\text{C}=\text{O}$ ) in an ala-ala fragment in a chain [1]  
two repeat units [1]

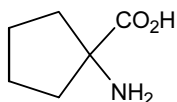
Allow peptide bond shown in full ( $\text{C}=\text{O}$ ) in a dipeptide ala-ala for 1 mark



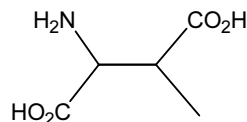
[3]

(c) (i)  $\text{HCl}$  or  $\text{H}_2\text{SO}_4$  or  $\text{NaOH}$  or  $\text{H}^+$  or  $\text{OH}^-$  reagents [1]  
+ heat and  $\text{H}_2\text{O}/\text{aq}$  (allow  $\text{H}_3\text{O}^+$ ).  
If T is quoted,  $80^\circ\text{C} < T < 120^\circ\text{C}$ . NOT warm. conditions [1]

(ii)



and



(if a structural formula, it must have all H atoms) allow protonated or deprotonated versions [1] + [1]

[max 3]

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(d) (i)  $\text{NH}_3^+ - \text{CH}(\text{CH}_3) - \text{CO}_2^-$  [1]

(ii)

compound	zwitterion

[3]

[4]

(e) (i) A buffer is a solution whose pH stays **fairly** constant *or* which maintains **roughly** the same pH *or* which resists/minimises changes in pH [1]  
when **small/moderate** amounts of acid/ $\text{H}^+$  or alkali/ $\text{OH}^-$  are added [1]

(ii)  $\text{NH}_2\text{CH}(\text{CH}_3)\text{CO}_2\text{H} + \text{H}(\text{Cl}) \longrightarrow {}^+\text{NH}_3\text{CH}(\text{CH}_3)\text{CO}_2\text{H} (+ \text{Cl}^-)$  [1]

(iii) blood contain  $\text{HCO}_3^-$  (*or* in an equation) [1]  
which absorbs  $\text{H}^+$  *or* equn  $\text{H}^+ + \text{HCO}_3^- \longrightarrow \text{H}_2\text{CO}_3 (\text{H}_2\text{O} + \text{CO}_2)$  [1]  
*or* absorbs  $\text{OH}^-$  *or* equn  $\text{OH}^- + \text{HCO}_3^- \longrightarrow \text{CO}_3^{2-} + \text{H}_2\text{O}$  [1]

(iv)  $[\text{CH}_3\text{CO}_2\text{Na}] = 0.05$   $[\text{CH}_3\text{CO}_2\text{H}] = 0.075$  [1]  
 $\text{pH} = 4.76 + \log (0.05/0.075) = \mathbf{4.58}$  *or*  $\mathbf{4.6}$  [1]

[7]

[Total: 19]

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4 (a)  $\text{Ca}(\text{NO}_3)_2 \longrightarrow \text{CaO} + 2\text{NO}_2 + \frac{1}{2} \text{O}_2$  [1]  
[1]

(b) (down the group) nitrates become **more stable** or require a higher temperature to decompose [1]  
as size/radius of (cat)ion increases or charge density of ion decreases [1]  
so polarisation/distortion of **anion/nitrate** decreases [1]  
[3]

(c) (i)  $\text{Li}_2\text{CO}_3 \longrightarrow \text{Li}_2\text{O} + \text{CO}_2$  [1]

(ii) radius of Li ion/ $\text{Li}^+$  is less than that of Na ion/ $\text{Na}^+$  (or polarising power of  $\text{M}^+$  is greater) [1]

(iii) Brown/orange fumes/gas would be evolved or glowing splint relights [1]  
Since the nitrate is likely to be thermally unstable or decomposes (just like the carbonate) or the balanced equation:  $2\text{LiNO}_3 \longrightarrow \text{Li}_2\text{O} + 2\text{NO}_2 + \frac{1}{2}\text{O}_2$  [1]  
[4]

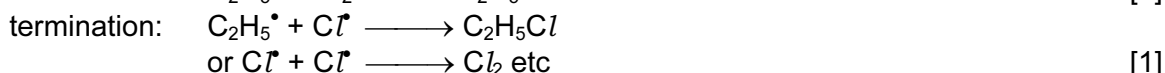
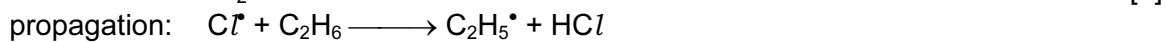
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- 5 (a) Alkanes are non-polar *or* have no dipole *or* C–H bonds are strong *or* C and H have similar electronegativities [1]  
[1]

- (b) (i) (free) radical substitution *or* substitution by homolytic fission [1]

- (ii) initiation:  $\text{Cl}_2 \longrightarrow 2\text{Cl}^\bullet$  [1]



all 3 names [1]

(iii)

structural formula of by-product	formed by
<b><math>\text{CH}_2\text{Cl}-\text{CH}_2\text{Cl}</math> (or isomer)</b>	<b>further substitution</b>
<b><math>\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3</math></b>	<b>(termination of 2 ×) <math>\text{C}_2\text{H}_5^\bullet</math></b>
<b><math>\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{Cl}</math> (or isomer)</b>	<b>substitution of <math>\text{C}_4\text{H}_{10}</math> by-product</b>

[3]

accept in the “formed by” column the formulae of radicals that will produce the compound in the “by-product” column, or the reagents, e.g.  $\text{C}_4\text{H}_9^\bullet + \text{Cl}_2$  *or*  $\text{C}_4\text{H}_9^\bullet + \text{Cl}^\bullet$  *or*  $\text{C}_4\text{H}_{10} + \text{Cl}_2$  (giving  $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{Cl}$ ).

do not allow anything more *Cl*-substituted than **dichlorobutane**.

N.B.  $\text{C}_2\text{H}_5\text{Cl}$  is the **major** product, not a **by**-product, so do not allow  $\text{C}_2\text{H}_5\text{Cl}$ .

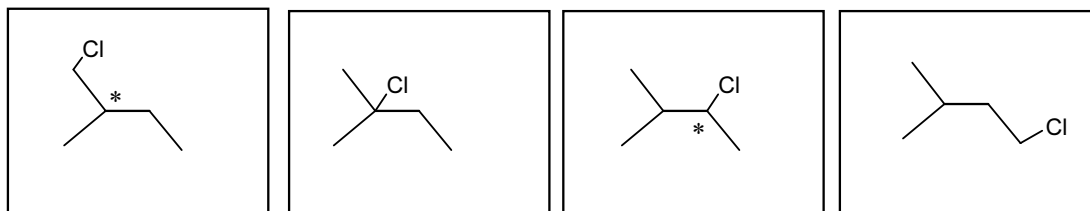
- (iv) J/K = **2.3 : 1** *or* 7:3 *or* 21:9 [2]

(reason: straightforward relative rate suggests 21:1, but there are 9 primary to 1 tertiary, so divide this ratio by 9.  $21/9 = 2.33$ )

allow [1] mark if J/K ratio is given as 21:1;

[10]

(c)



4 isomers  $4 \times [1]$

2 chiral atoms identified correctly, even in incorrect structures

[1] + [1]

**[max 5]**

**[Total: 16]**

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- 6 (a) (i) **K**, because it is the (only) one to contain nitrogen *or* it's an amino acid *or* because it contains CO<sub>2</sub>H *or* NH groups [1]
- (ii) molecule: **J**, polymer: RNA (**not** DNA) [1]  
*or* molecule: **L**, polymer: starch, cellulose, glycogen *or* polysaccharide (**not** carbohydrate) [2]
- (b) (i) Covalent bonding [1]  
(ii) Hydrogen bonding [1]  
(iii) Ionic/electrovalent bonding *or* disulphide/–S–S– bonding *or* van der Waals' forces [1]  
[3]
- (c) (i) Enzymes [1]  
(ii) • change in pH  
• increase in T (NOT decrease; T > 40 °C *or* “too high” are OK)  
• addition of heavy metal ions *or* specific, e.g. Hg<sup>2+</sup>, Ag<sup>+</sup>, Pb<sup>2+</sup> etc.  
any two bullet points [1] + [1]
- change in pH disrupts ionic bonds  
*or* metal ions disrupt ionic bonds  
*or* metal ions disrupt –S–S– bonds  
*or* heating disrupts hydrogen bonds  
any one [1]
- This changes: the 3D structure *or* shape of the enzyme *or* the active site [1]  
[**max 4**]  
[**Total: 9**]



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7 (a)

structural information	analytical technique
three-dimensional arrangement of atoms and bonds in a molecule	X-ray crystallography/diffraction
chemical environment of protons in a molecule	NMR (spectroscopy) <b>only</b>
identity of amino acids present in a polypeptide	Electrophoresis / chromatography / mass spectrometry

[1] + [1] + [1]  
[3]

(b) (i) **paper chromatography;**

The components **partition** between the solvent/moving phase and the water/liquid stationary phase *or* separation relies on different solubilities (of components) in the moving solvent and the stationary water phase. [1]

(ii) **thin-layer chromatography.**

Separation depends on the differential **adsorption** of the components onto the solid particles/phase *or*  $Al_2O_3$  *or*  $SiO_2$ . [1]  
[2]

(c) (i) No. of carbon atoms present =  $\frac{0.2 \times 100}{5.9 \times 1.1} = 3.08$  hence 3 carbons [1]

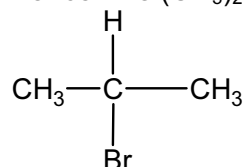
(ii) Bromine [1]

(iii) **One** bromine is present as there is only an M+2 peak / no M+4 peak *or* the M and M+2 peaks are of similar height [1]

(iv) *NMR spectrum shows a single hydrogen split by many adjacent protons and 6 protons in an identical chemical environment. This suggests...*  
two  $-CH_3$  groups and a lone proton attached to the central carbon atom [1]

Empirical formula of **N** is  $C_3H_7Br$  [1]

Hence **N** is  $(CH_3)_2CHBr$  *or*



[1]  
[6]

[Total: 11]

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- 8 (a) (i) Soluble form would be most effective [1]
- (ii) **Q**, since the 'mini-pills'/granules/powder have a larger surface area  
or **P**, because it has no protective casing [1]
- (iii) The gel coat stops it being broken down while passing through the upper part of the digestive system/stomach  
or the gel coat is stable to stomach acid. [1]  
[3]
- (b) The drug is taken quickly/directly to the target  
or more accurate dosing can be achieved [1]
- When the drug is taken by mouth it has to pass through the stomach/intestine wall to get into the bloodstream. or some is digested/lost to the system [1]  
[2]
- (c) (i) condensation (polymerisation) [1]
- (ii) hydrogen bonds or van der Waals' [1]
- (iii) It would change the overall shape of the (drug) molecule  
The 'fit' into the active site would be less effective [1] + [1]
- (iv) Hydrolysis [1]  
[5]
- [Total: 10]