MARKSCHEME

May 2001

CHEMISTRY

Higher Level

Paper 2

14 pages

SECTION A

1. (a) (i) Tl **[1]**

   (ii) Mn^{2+} (accept Mn) **[1]**

   (iii) Any two of N, P, As, Sb, Bi **[1]**

   (b) Lines (may be shown on diagram) **[1]**
   Convergence at high frequency / energy / short wavelength (may be shown on a diagram) **[1]**
   Electron transfer between energy levels / each transition / line depends on energy differences / $\Delta E = h\upsilon$ **[1]**   **[3 max]**

   (c) remaining electrons in one shell / Li^+ has one less shell or converse **[1]**
   effective nuclear charge (Li^+) greater than that of the atom **[1]**   **[2 max]**

   (d) *Any two of the following, [1] each*
   • Do not form bonds
   • Full outer shells
   • No attraction for electrons
   • Stable / inert / do not react   **[2 max]**

2. (a)

	Pt	Cl	N	H	
Divide by A_r values					**[1]**
No. of moles	0.3332	0.6666	0.667	2	**[1]**

   Empirical formula $PtCl_2N_2H_6$ **[1]**   **[3 max]**

   (b) Accept any diagram that shows two Cl atoms attached to Pt **and** two NH_3 groups attached to Pt. (Allow ECF from (a)). **[1]**

   (c) Moles of $AgNO_3 = 0.0162 \times 0.102 = 0.00165$ **[1]**
   Moles of $NaCl = 0.0148 \times 0.125 = 0.00185$ **[1]**
   Therefore limiting reactant is $AgNO_3$ / 0.00165 moles AgCl produced **[1]**
   Mass of AgCl produced $= 0.00165 \times 143.32 = 0.236\,g$ (accept values in range 0.236 - 0.237 g) **[1]**   **[4 max]**

   If the wrong limiting reagent is used (NaCl), 0.265 g scores [3].
   (N.B. Error carried forward; other routes are possible for the deduction that $AgNO_3$ is limiting / moles $AgCl$ = moles $AgNO_3$)

3.   (a)   (i)   Energy / enthalpy change for the formation of 1 mol of a compound *[1]*
                 from its elements in their standard states / 101 kPa, 298K *[1]*   *[2 max]*

           (ii) $$3C(s) + 4H_2(g) \rightarrow C_3H_8(g)$$ *[1]*

      (b)   (i)   Reference to (particular) bonds in a range of compounds / average values *[1]*
                 Break or make 1 mole of bonds in gaseous state
$$= [2(C—C) + 8(C—H) + 5(O=O)] - [6(C=O) + 8(O—H)] \; [1]$$
$$= [2(348) + 8(412) + 5(496)] - [6(743) + 8(463)] \; [1] \; *$$
$$= -1690 \; [1] \; (units \; not \; required)$$ *[5 max]*

                 *(N.B. correct extraction of **five** bond enthalpy values (even with wrong coefficients) would score [1])*

           (ii)   Positive *[1]*
                 Increase in disorder / less order / more gas (on RHS) *[1]*   *[2 max]*

4.   (a)   (i)   *Any two of the following, [1] each*
                 Some mention of a **reversible** reaction /⇌ *etc.*
                 Indication that rate of left to right = rate right to left
                 At equilibrium no net change in concentration of reactants and products
                 Closed system   *[2 max]*

           (ii) $K_c = \dfrac{[CH_3CO_2C_2H_5][H_2O]}{[CH_3CO_2H][C_2H_5OH]}$ *[1]*   *[1]*

           (iii) $[C_2H_5OH] = [CH_3CO_2H] = 0.33$ *[1]*

                 $K_c = 4.1 \, (\text{accept } 4.0)$ *[1]*   *[2 max]*

                 (Allow for ECF from (ii))

      (b) K_w increases with temperature rise / $[H^+]$ or $[OH^-]$ rises as temperature rises *[1]*
              Equilibrium shifts to right (as temperature increases) *[1]*   *[3 max]*
              Endothermic reaction *[1]*

      (c)   (i)   No effect *[1]*
                 Some description of role of catalyst *e.g.* increases rate of forward **and** reverse reaction (equally) / increases rate of attainment of equilibrium *[1]*   *[2 max]*

           (ii)   No effect *[1]*
                 Concentration of reactants the same *[1]*   *[2 max]*

SECTION B

5.   (a)   Volume of gas method: **or**   Mass loss method:
        Chemicals in flask / beaker *[1]*   Chemicals in flask / beaker *[1]*
        Method of collection *[1]*   Top pan balance *[1]*
        Airtight *[1]*   Not airtight *[1]*

        Measurement and processing:
        Volume / mass monitoring *[1]*
        Timing *[1]*
        Appropriate processing of data *[1]*

        e.g. convert time to rate / $\text{rate} = \dfrac{1}{\text{time}}$ / plot volume–time graph and measure slope   *[6 max]*

   (b)   Slope decreases / slope becomes less steep / rate decreases *[1]*
         Acid / reactant concentration falls / surface area of magnesium / reactant decreases *[1]*   *[2 max]*

   (c)   (i)   For line B:
               Curve shown less steep than A *[1]*
               Curve becomes horizontal at half the height of A / explain why curve goes to
               half height *[1]*
               Acid particles more widely spaced / less frequent collisions between reactant
               particles *[1]*   *[3 max]*

         (ii) For line C:
               Curve shown steeper than A *[1]*
               Curve becomes horizontal at same height as A *[1]*
               Magnesium surface area increased / more frequent collisions between
               reactant particles *[1]*   *[3 max]*

   (d)   Order = 1 / first order *[1]*
         Rate proportional to concentration / graph is straight line *[1]*   *[2 max]*

   (e)   Rate = $k[\text{H}_2\text{SO}_4]$ *[1]*

         e.g. $k = \dfrac{0.008}{0.9} \bigg/ \dfrac{0.006}{0.675} \bigg/ \approx 8.9 \times 10^{-3}$ *[1]*   (*accept answer* $\pm 2 \times 10^{-3}$)

         s^{-1} *[1]*
         The value of k would be greater at a higher temperature *[1]*   *[4 max]*

(f)   Diagram:
Line for products below line for reactants *[1]*
ΔH marked on vertical distance between reactants line and products line *[1]*
y axis label and *x* axis label or reactant and product labels *[1]*
E_a marked on vertical distance between reactants line and intermediate / transition
state *[1]*
E_{cat} marked on vertical distance between reactants line and catalyst intermediate /
transition state *[1]*   *[5 max]*

Energy/enthalpy

reaction coordinate/progress of reaction / time / reaction path
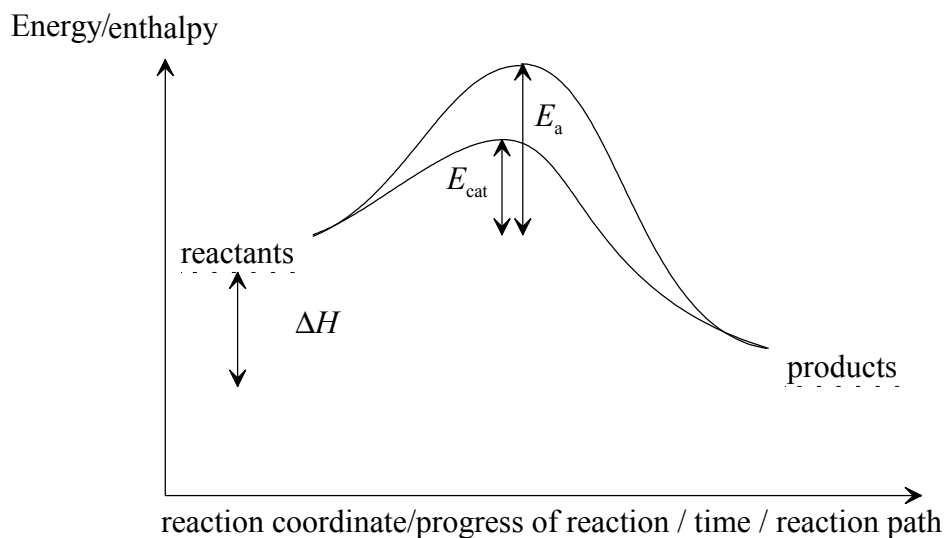
(If reaction is shown as endothermic, award [4 max])

6. (a) **Brønsted–Lowry**: acid is proton (H^+) donor and base is proton acceptor *[1]*

For example, $HNO_3 + H_2O \rightarrow H_3O^+ + NO_3^-$ *[1]* (*accept other valid example*)

Identification of the acid **and** the base on the left hand side *[1]*

Lewis: acid is electron pair acceptor (electrophile) and base is electron pair donor (nucleophile) *[1]*

For example, $BF_3 + NH_3 \rightarrow BF_3NH_3 / Cu^{2+} + 4NH_3 \rightarrow Cu(NH_3)_4^{2+}$ *[1]* (*accept other valid example*)

Identification of **both** the acid and the base on the left hand side *[1]*

Bond formed is (dative) covalent / co-ordinate / indication that one species gives bonding electron pair *[1]*   *[7 max]*

(b) Strong acid completely dissociated, weak acid weakly / partially dissociated *[1]*
example of strong **and** weak acid (names or formulas) *[1]*   *[2 max]*

(c) Molecule and anion equilibrium / $HIn \rightleftharpoons H^+ + In^-$ / indicator is a weak acid or weak base *[1]*
Different colours *[1]*
Addition of H^+ or OH^- / acid or alkali causes shift in equilibrium *[1]*
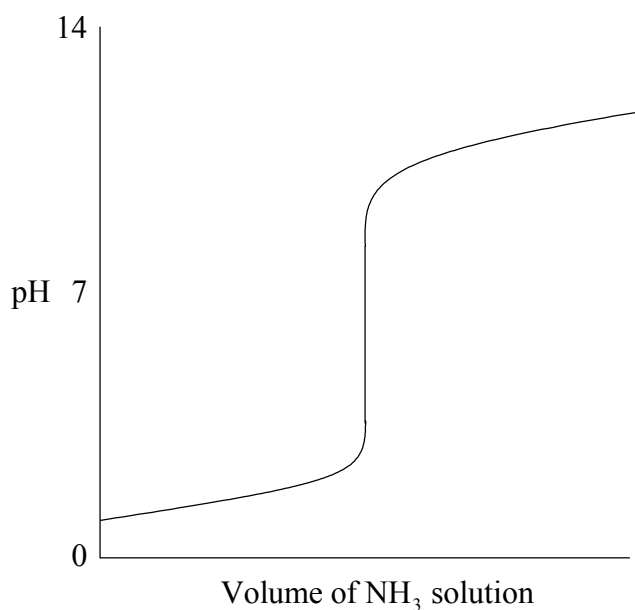$H^+ \rightarrow$ colour A / OH^- causes colour B *[1]*   *[4 max]*

(d) (i)
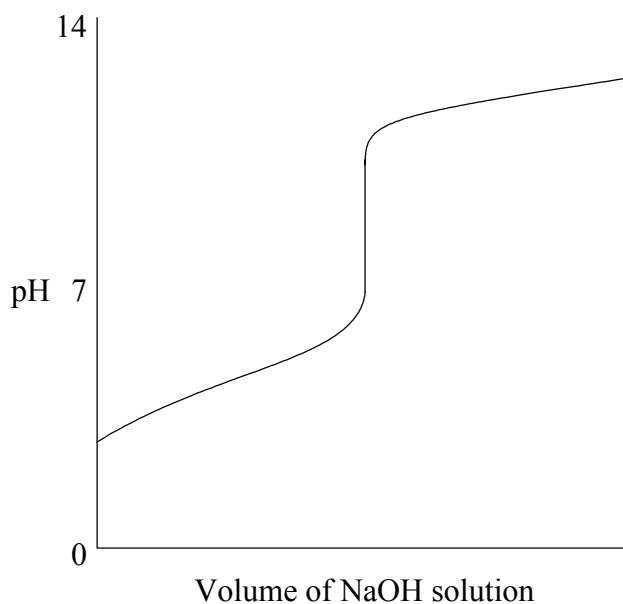
Line starts clearly above $pH = 1$ *[1]*
More of vertical line in alkaline than acidic region *[1]*
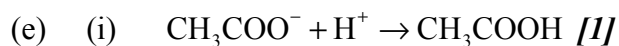Phenolphthalein *[1]*   *[3 max]*

(ii)

Line starts at pH = 1 / same as printed graph *[1]*
More of vertical line in acidic than alkaline region *[1]*
Methyl orange / methyl red / bromophenol blue / bromothymol blue / bromocresol green *[1]*   *[3 max]*

(e)   (i) $CH_3COO^- + H^+ \rightarrow CH_3COOH$ *[1]*
          Added H^+ ions react with conjugate base / CH_3COO^- *[1]*   *[2 max]*

      (ii) $[H^+] = \dfrac{K_a[CH_3COOH]}{[CH_3COO^-]}$ / $pH = pK_a + \log\dfrac{[base]}{[acid]}$ *[1]*
            Correct values of CH_3COOH and CH_3COO^- mol or $mol\,dm^{-3}$ *[1]*
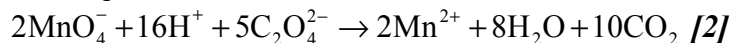   Correct value of $[H^+] = 3.48 \times 10^{-5}$ $(mol\,dm^{-3})$ *[1]*
            pH = 4.4–4.5 (consequential on $[H^+]$ value) *[1]*   *[4 max]*

7. (a) Oxidation number of Mn on left-hand side 7 *[1]*
Oxidation number of C on left-hand side 3 *[1]*
Oxidation number of Mn **and** C on right-hand side 2 and 4 respectively *[1]*

Final equation:
$$2MnO_4^- + 16H^+ + 5C_2O_4^{2-} \rightarrow 2Mn^{2+} + 8H_2O + 10CO_2 \ \textbf{[2]}$$

*(If ratio $MnO_4^- : C_2O_4^{2-}$ is $2:5$ but equation wrong give **[1]**. Do not penalise absence of state symbols)* *[5 max]*

(b) (i) Diagram must show:
Voltmeter and metals *[1]*
Salt bridge correctly labelled *[1]*
$[Ni^{2+}]$ **and** $[Cd^{2+}]$ as $1 \ mol\,dm^{-3}$ *[1]* *[3 max]*

(i) • H_2 gas
• At 1 atm / 101.3 kPa
• 298 K
• $1 \ mol\,dm^{-3} \ H^+$
*(Four correct **[2]**, 2/3 correct **[1]**)*
Pt electrode *[1]* *[3 max]*

(c) (i) $Ni^{2+}(aq) + 2e^- \rightarrow Ni(s)$ *[1]*
$Cd(s) - 2e^- \rightarrow Cd^{2+}(aq) / Cd(s) \rightarrow Cd^{2+}(aq) + 2e^-$ *[1]*
Allow $Cd + Ni^{2+} \rightarrow Cd^2 + Ni$ for *[2]*

Cd is oxidised *[1]*
Oxidising agent is Ni^{2+} *(do not accept Ni)* *[1]* *[4 max]*

(ii) Cd is **A** *[1]*
External electron flow from Cd to Ni *[1]*. *[2 max]*

(iii) Overall voltage is 0.2 volts (accept + or – 0.2 V) *[1]*
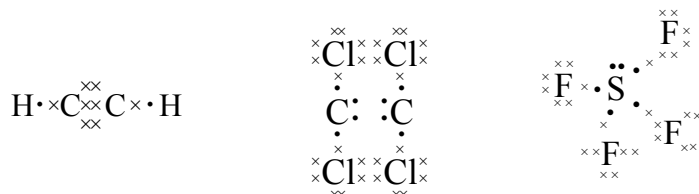ΔG is negative *(consequential on the sign of the potential)* *[1]* *[2 max]*

(d) Silver at cathode / - ve *[1]*
Oxygen at anode / + ve *[1]* *[2 max]*

(e) Number of coulombs $= 20 \times 5 \times 60^2 / 360000$ *[1]*
Number of Faradays $= 3.73$ *[1]*
Moles of $OH^- = 3.73$ *[1]*
Mass of NaOH $= 3.73 \times 40 = 149.2$ g *[1]* *[4 max]*

8. (a) Each structure (**with** non-bonding e⁻ pairs where relevant) *[1]*
(*lines are acceptable instead of pairs of dots/crosses*)

H·×C××C×·H

Cl×× ×Cl×
  C: :C
×Cl× ×Cl×

×F ·S·
F   F
  F

Shapes:
C_2H_2 linear / straight *[1]*
            angle 180° *[1]*

C_2Cl_4 (trigonal) planar *[1]*
            angle 120° *[1]*

SF_4 K-shaped / based on trigonal bipyramid / see saw *[1]*
            angle 90° *[1]*
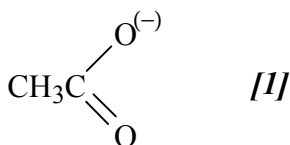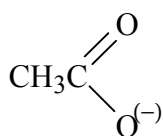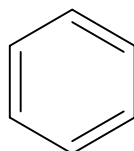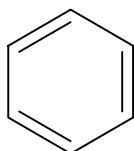   angle 120° *[1]*   *[10 max]*

(b) sp (C_2H_2) *[1]*
     sp² (C_2Cl_4) *[1]*   *[2 max]*

(c) Acceptable resonance structures

CH_3C with =O and O⁻ *[1]* CH_3C with O⁻ and =O *[1]*

(accept delocalized ethanoate ion for *[2]*)

e.g. (CH_3C with O, (−), O scores *[2]*)
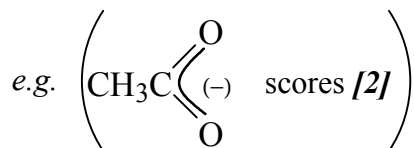
[benzene ring] *[1]*   [benzene ring] *[1]*   *[4 max]*

(accept delocalized benzene for *[2]*)

(d) (i)   C to O in the ion equal *[1]*
            Different bond length in acid *[1]*
            $C = O < C—O$ (*or converse*) *[1]*
            C to O in ethanoate ion intermediate bond length / 1.5 bonds *[1]*   *[4 max]*

     (ii)   Benzene is stable because of delocalization *[1]*
            Addition destroys delocalization *[1]*   *[2 max]*

     (iii) (pK_a values show) ethanoic stronger acid than ethanol *[1]*
            (*Some correct statement about difference in acidity*)

            Reference to relative stability of (both) anions (*they do not have to be shown*) *[1]*

            Anion of acid more stable C = O group e⁻ withdrawing / delocalisation of charge (or resonance) in ethanoate ion / explain instability of ethoxide *[1]*   *[3 max]*