

Nyakambi, N.

(5)

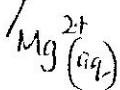
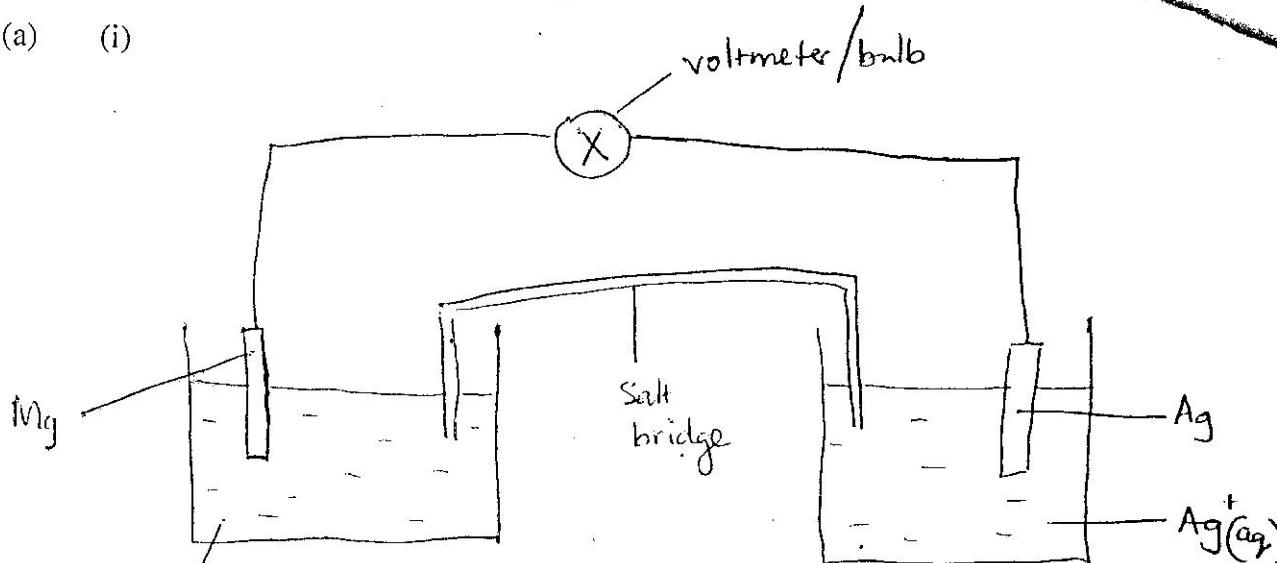
**ZIMBABWE SCHOOL EXAMINATIONS COUNCIL**  
**General Certificate of Education Advanced Level**

**MARKING SCHEME**

**NOVEMBER 2009**

**CHEMISTRY 9189/1**

1 (a) (i)

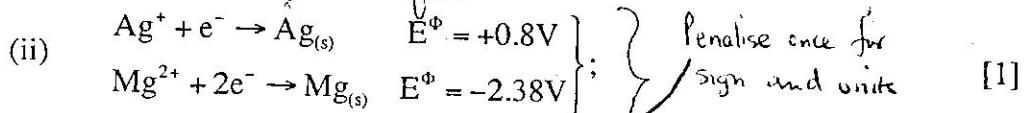


Careful marks for  
eqns.

(diagram only -1)  
labels -1

[2]

Answer



Penalise once for  
Sign and units

[1]

$E^\phi_{cell} = 0.8 - (-2.38) = +3.18 V;$

~~with sign and units~~

[1]

(iii)  $n(Mg) = \frac{0.08}{24} = 3.3 \times 10^{-3}$  moles;

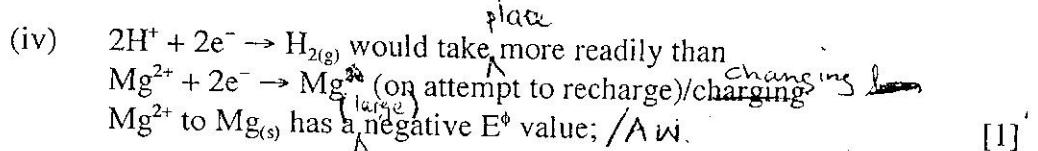
[1]

$n(\text{electrons given off by Mg}) = 2 \times 3.3 \times 10^{-3} = 6.6 \times 10^{-3}$  moles; [1]

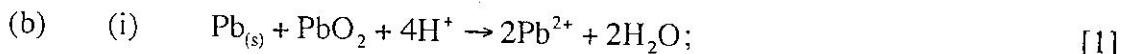
$1 \text{ mole } e^- \rightarrow 96500 \text{ C}$

$6.6 \times 10^{-3} \text{ moles } e^- \rightarrow 6.6 \times 10^{-3} \times 96500 = 636.9 \text{ C};$  [1]

$I = \frac{Q}{t} = \frac{636.9}{5 \times 60} = 2.12 \text{ amp (answer with units);}$  [1]



[1]



[1]

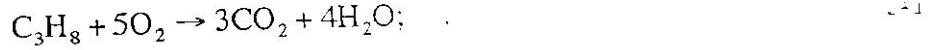
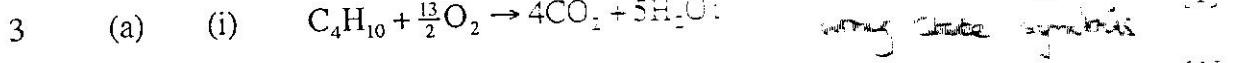
(ii) Limited mileage between recharging; high mass;  
low voltage; (any two)

[2]

/current.

[Total: 12]

- 2 (a) (i) It is when in a reversible reaction the rate of the forward reaction equals the rate of the backward reaction; [1]
- (ii) Temperature - Increase in temperature shifts equilibrium to the endothermic reaction; /AW/vice versa;
- Concentration - Increase in concentration of reactants shift equilibrium to the products side/AW/vice versa;
- Pressure - Increase in pressure shifts equilibrium to the side with fewer moles of gas/vice versa; /AW [2]  
(any two)
- (iii) A catalyst increases the rate of both the forward and the backward reactions; [1]
- (b) (i)  $n(\text{OH}^-) = n(\text{H}^+) = \frac{18}{1000} \times 0.2 = 0.0036 \text{ (moles);}$  [1]
- (ii)  $n(\text{OH}^-) \text{ from } \text{M(OH)}_3 = 0.0036 - n(\text{OH}^-) \text{ from KOH}$   
 $= 0.0036 - \frac{25}{1000} \times 0.1 = 0.0011 \text{ (moles);}$  [1] ~~ecf~~
- $\therefore [\text{OH}^-] \text{ (from } \text{M(OH)}_3) = \frac{0.0011 \times 1000}{25} = 0.044 \text{ (mol dm}^{-3}\text{)}$  [1] ~~ecf~~
- (iii)  $[\text{M}^{3+}] = \frac{1}{3} [\text{OH}^-] \text{ (from } \text{M(OH)}_3) = \frac{1}{3} \times 0.044 = 0.01467 \text{ (mol dm}^{-3}\text{)}$  [1] ~~ecf~~
- (iv) It is the amount of a substance that dissolves in a specific volume of solvent at a specified temperature; /AW. [1]
- Solubility of  $\text{M(OH)}_3$  in  $0.1 \text{ Mol dm}^{-3}$  KOH =  
 $s = 0.01467 \text{ mol dm}^{-3};$  [1] ~~ecf~~  
 (1 mole  $\text{M(OH)}_3$  gives 1 mole  $\text{M}^{3+}$  ions in solution)
- (v)  $K_{\text{sp}} = s(3s + 0.1)^3$  (~~if  $0.1 >> 3s$~~ ) credit.  
 $\therefore K_{\text{sp}} = s \times (0.1)^3 = 0.01467 \times (0.1)^3 = 1.5 \times 10^{-5} \text{ mol}^4 \text{ dm}^{-12};$  [2]
- $\begin{pmatrix} \text{working - 1 mark} \\ \text{answer - 1 mark} \\ \text{with correct units} \end{pmatrix}$
- [Total: 12]



(ii) Let volume of butane be  $x$  and that of propane be  $y$  then:

$$x + y = 1 \text{ dm}^3; \text{ (eqn I)} \quad [1]$$

$$\frac{x}{24} \times 3000 + \frac{y}{24} \times 2280 = 118.08; \text{ (eqn II)} \quad [1]$$

From equ I  $x = 1 - y$ ; substitute for  $x$  in eqn II

$$\left[ \begin{array}{l} \left(1 - y\right) \frac{3000}{24} + \frac{2280}{24} y = 118.08 \\ 125 - 125y + 95y = 118.08 \end{array} \right] \quad \begin{array}{l} \text{Accept} \\ \text{alternative} \\ \text{Working} \\ \text{if} \\ \text{is correct.} \end{array}$$

$$\therefore y = \frac{118.08 - 125}{-30} = 0.2307 \Rightarrow 23.1\%; \quad [1]$$

$$x = 1 - 0.2307 = 0.7693 \Rightarrow 76.9\%; \quad [1]$$

6 max 5

(b) (i) heat absorbed by water =  $\frac{70}{100} \times 118.08 = 82.656 \text{ kJ}$ ; [1]

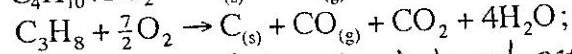
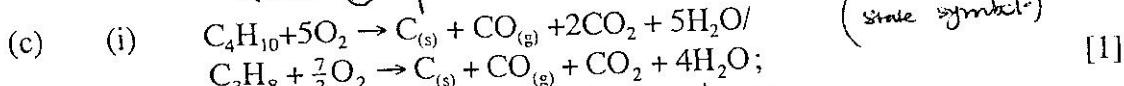
then

$$(82.656 \times 1000 \text{ J} = mc\theta = m \times 4.2 \times (100 - 22)) \quad \begin{array}{l} \text{(accept v from)} \\ \text{m} \end{array} \quad [1]$$

$$m = \frac{82656}{4.2 \times 78} = 252.308 \text{ g}; \quad [1]$$

$$\therefore \text{Volume of water} = 252.3 \text{ cm}^3 / 0.252 \text{ dm}^3; \quad [1]$$

(ii) Assumption - water has a density of  $1 \text{ g/cm}^3$ ; / boiling pt of water at rtp is  $100^\circ\text{C}$ . [1]



accept C and CO for balanced eqn.

(ii) CO - poisonous if inhaled; / Ans. [1]

C - causes smog; / Ans. / gives the atmosphere a dirty appearance. [1]

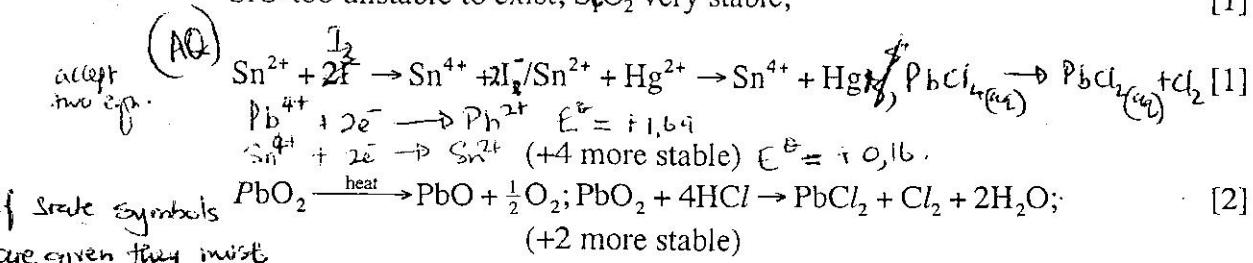
↓  
plants, buildings

[Total: 12]

[13 Max 12]

- 4 (a)  $\text{XO} + \frac{1}{2}\text{O}_2 \rightarrow \text{XO}_2$  (where X is C, Ge or Sn); [1]  
 (+4 more stable)

SiO too unstable to exist,  $\text{SiO}_2$  very stable; [1]



e.g.  $\text{SnO}_{(s)}$  Stability of +2 increases down the group while that of +4 decreases; [1]  
~~(Reject reverse trend)~~

$\text{SnO}_2^{(s)}$  S electrons are less screened by inner electrons compared to p electrons hence tend to behave as inner electrons down the group/inner pair effect; [1]

- (b) (i) Covalent bonding; simple molecular structure; tetrahedral shape; [3] consider separately  
 (ii) Both  $\text{SnI}_4$  and  $\text{I}_2$  are (non polar) covalent substances that dissolve both in organic solvents; [1]

$\text{SnI}_4$  would not form in water as it may be hydrolysed/react in water [1]  
 [Total: 12]

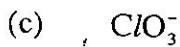
- 5 (a)  $3\text{I}_2 + 6\text{OH}^- \xrightarrow{5\text{T}^-} 5\text{I}^- + \text{IO}_3^- + 3\text{H}_2\text{O}$ ; / [1]  
 $3\text{I}_2 + 6\text{KOH} \rightarrow 5\text{KI} + \text{KIO}_3 + 3\text{H}_2\text{O}$  (reject).

Only  $\text{I}^-$  reacts with con  $\text{H}_2\text{SO}_4$  to reform  $\text{I}_2$ ; [1]  
 $\text{I}^- + \text{H}_2\text{SO}_4 \rightarrow$  accept equation (balanced). [1]  
 $\therefore$  Fraction =  $\frac{5}{6}(0.833)$  [1]

- (b) H - I ; [1]

$\text{HCl} - 431\text{kJmol}^{-1}$   
 $\text{HBr} - 366\text{kJmol}^{-1}$   
 $\text{HI} - 299\text{kJmol}^{-1}$  [1]

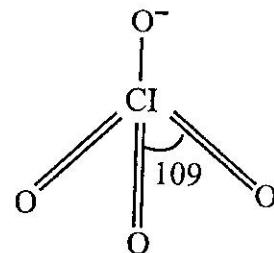
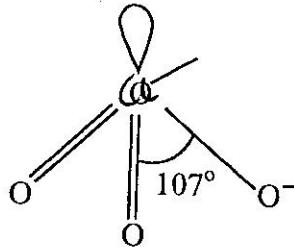
H-X bond energy values decrease down the group [1]  
 $\therefore$  bonds become weaker; / know A/W



one lone pair  
3 bonded pairs  
shape trigonal pyramidal



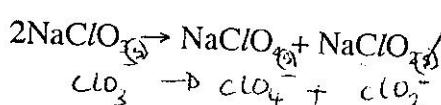
no lone pairs  
4 bonded pairs  
shape tetrahedral

Bond angle  $\approx 107^\circ$ ;

(1 mark for each shape and one mark for each angle)

bond angle  $109^\circ$ ; [4] $109 - 109,5$ 

- (d) Simultaneous oxidation and reduction of the same species; / AW [1]



[1]

[Total: 12]

6

(a)

Treat with hot concentrated KMnO<sub>4</sub>; Then treat product with 2.4 dinitrophenyl hydrazine;  $\text{NaClO}_4$  then  $\text{I}_2$  (aq); then (warm)  $\text{MnO}_4^-$  (orange) accept  $\text{MnO}_4^-$  (alkaline)  $\text{H}^+$  (yellow)

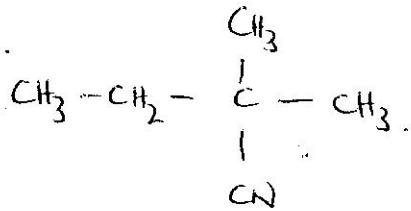
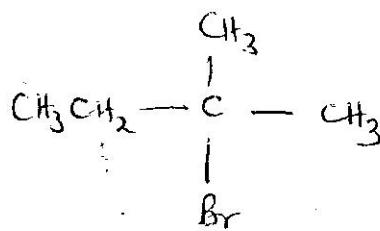
for  $\text{C}_2\text{H}_3$  yet  
only in

A gives an orange-red ppt B does not; accept yellow (reject for triiodomethane test) [3]

(i) II -  $\text{KCN}$  dissolved in ethanol, reflux/heat; reject acid base [2]  
III - heat/reflux in dilute sulphuric/hydrochloric acid; reject heat [2]

(ii) C

D



[2]

- (iii) It would not exhibit optical activity;
- 
- It has no chiral carbon; [2]

b(iii) (c)

Boiling points; / melting pt / volatility for B.  
B has higher boiling point; [1]

More efficient packing of molecules than in branched A; / AW. [1]

accept density (branched A) low

[Total: 12]

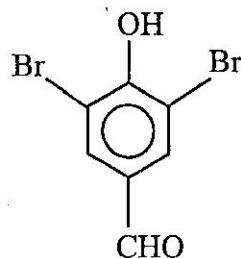
7 (a) (i) E and F – position; E/F and G – functional group; [2]

(ii)  $E < F < G$ ; accept  $G > F > E$  [1]

<sup>molecular</sup>  
E forms intra<sub>A</sub> hydrogen bonding hence only weak Van der Waal's forces hold molecules together; [1]

<sup>molecular</sup>  
Both F and G form inter<sub>A</sub> hydrogen bonds but there is <sup>more</sup> <sup>or</sup> stronger in G than F; [1]

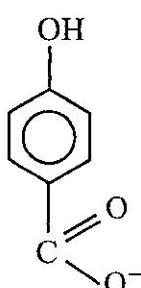
(b) (i) (Bromine decolourised) and white ppt formed;



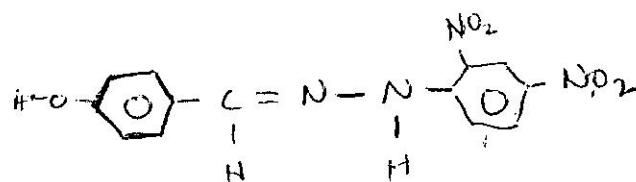
~~reject black~~

(ii) silver mirror;  
black/grey ppt/  
silver ppt;

accept silver lining



(iii) orange/  
orange-red ppt;  
accept yellow



[6]

(c) condensation/nucleophilic substitution; addition-elimination [1]

[Total: 12]

(a)

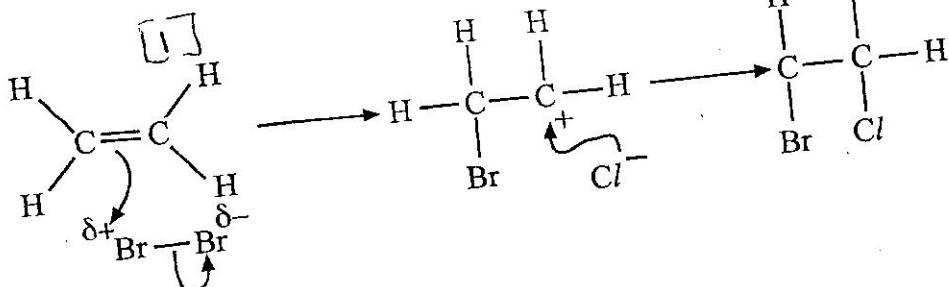


[1]

Alkalis shift above equilibrium to the right releasing  $\text{CN}^-$  to attack carbonyl carbon; [1]

Acids shift above equilibrium to the left hence no  $\text{CN}^-$  to attack carbonyl carbon; [1]

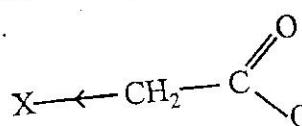
(b)



[3]

/ mechanism described in words

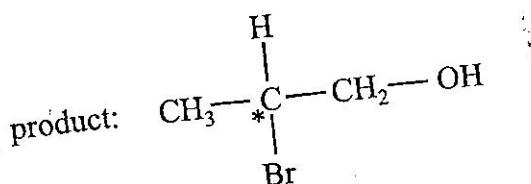
(c) F is more electronegative than I; hence causes a greater negative inductive effect and more stabilisation of the anion: / AW [1]



[1]

more weakening of  $\text{O}-\text{H}$  bond.

(d) An equimolar mixture of D and L forms is formed, effects cancel each other out (racemic mixture); / AW [1]



[1]

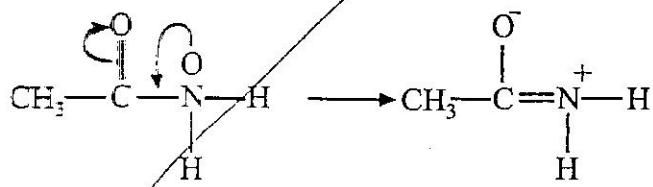
AW

- rotation of light
- equal amounts

- (e)  $\text{CH}_3\text{CH}_2\text{NH}_2$  can accept a proton via the lone pair of electrons on the nitrogen atom;

[1]

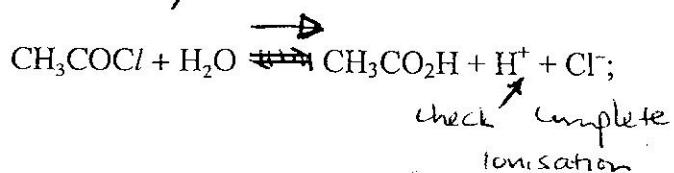
The lone pair on the nitrogen atom of  $\text{CH}_3\text{CONH}_2$  is not available for protonation due to the delocalisation shown below:



- (f) Ethanoic acid partially ionises in solution to give a weakly acidic solution  
 $\text{CH}_3\text{COOH} \rightleftharpoons \text{CH}_3\text{COO}^- + \text{H}^+$

[1]

Ethanoyl chloride is hydrolysed by water to give  $\text{HCl}$  which completely ionises in solution.



[1]

[Total: 12]

~~X~~ Cancelled and not corrected consider.