# SET 8 CHEMISTRY PAPER 2 MARKING SCHEME

### **Question 1**

a)  $ZnCO_{3(s)} \ + \ 2HCl_{(aq)} \ \rightarrow ZnCl_{2(aq)} \ + \ Co_{2(g)} \ + \ H_2O_{(l)} \ ^{\sqrt{1}mk}$ i. To remove traces of hydrogen chloride gas  $\sqrt[n]{1mk}$ ii. Some  $Co_2$  gas may dissolve in the water  $\sqrt{1}mk$ iii.  $2NaOH_{(aq)} + H_2O_{(l)} \rightarrow Na_2CO_{3(aq)} + H_2O(l)^{\sqrt{1}mk}$ iv.  $Na_2CO_{3(aq)} + H_2O_{(l)} + CO_{2(g)} \rightarrow 2NaHCO_{3(s)} \sqrt{1mk}$ √1mk Heat the sodium hydrogen carbonate. v.  $2NaHCO_{3(s)} \quad \underline{Heat} \quad Na_2CO_{3(s)} + H_2O_{(l)} + Co_{2(g)} \sqrt{1mk}$ vi. b)  $=\frac{30 \times 1}{1000} \sqrt{\frac{1}{2} \text{ mk}}$  (moles of NaOH that reacted) i.  $= 0.03^{\sqrt{\frac{1}{2}} \text{mk}}$ Initial moles of HCl =  $\frac{50 \times 1}{1000} \sqrt{\frac{1}{2} \text{ mk}}$ ii.  $= 0.05^{\sqrt{\frac{1}{2}mk}}$  $NaOH_{(aq)} + HCl_{(aq)} \rightarrow NaCl_{(aq)} + H_2O_{(l)}$ Acid : base = 1 : 1 Therefore, . : Moles of HCl that reacted with NaOH 0.03 = $\begin{array}{c} 0.05-0.03 \ ^{\sqrt{1/2}\ mk} \\ 0.02 \ ^{\sqrt{1/2}\ mk} \end{array}$ . : Moles of HCl that reacted with XCO<sub>3</sub> = =  $XCO_{3(s)} + 2HCl_{(aq)} \rightarrow 2XCl_{(aq)} + CO_{2(g)} + H_2O_{(l)}$ iii. Acid : carbonate = 2 : 1 Therefore,  $\frac{1}{2} \ge 0.02^{\sqrt{1}/2} = 0.02^{1}$ Moles of XCO<sub>3</sub> that reacted with HCl =  $0.01^{\sqrt{1/2}} \text{ mk}$ = 0.01 moles contain 1g 1 mole has ?  $=\frac{1 x 1}{0.01} g^{\sqrt{1/2} mk}$  $= 100g^{\sqrt{1/2} mk}$  $x + 12 + 48 = 100^{\sqrt{\frac{1}{2} \text{mk}}}$ iv. x + 60 = 100 $x = 40^{\sqrt{1/2} mk}$ 

### **Question 2**

Alkali metals  $\sqrt[1]{1mk}$ i.

ii.

- B is more reactive than D.  $\sqrt{1}$  the outermost energy level electron in D is more firmly held i. than in S  $\sqrt{1}$ mk
- ii. J is more reactive than K.  $\sqrt[1]{2} \frac{mk}{mk}$  the nuclear electron attraction is higher in J than in K.  $\sqrt[1]{2} \frac{mk}{mk}$ E has a larger atomic radius than F.  $\sqrt[1]{1mk}$  nuclear charge increases across the period.  $\sqrt[1]{1mk}$
- iii.
- Before G  $\sqrt[]{1mk}$ iv.
- The melting point increases  $\sqrt[1]{1}$  across the period. Due to increase in the strength of the metallic v. bonds formed as the number of valency electrons increases.  $\sqrt[1/2]{mk}$
- $EK_3^{\sqrt{1}mk}$ vi.
- Ionic / electrovalent bond  $\sqrt[1]{1mk}$  it is formed through transfer of electrons from metal to a non metal. vii. √1mk
- Used in light bulbs.  $\sqrt[1]{1mk}$ viii.

2,8,8,8  $^{\sqrt{1/2}}$  mk ix. С =

2.8 <sup>√ 1/2</sup> mk G =

## **Ouestion 3**

a)

I = polymerization  $\sqrt{1}$  mk i.

II – Thermal Cracking  $\sqrt{1}$ mk

- A = 1,2 dibromopropane  $\sqrt{1}$ mk ii.
- $B = Ethyne \sqrt{1}$
- Asbestos <sup>√1mk</sup> iii.

iv. 
$$\begin{pmatrix} H & CH_3 \\ | & | \\ C & - & C \\ | & | \\ H & H \end{pmatrix}_n^{\sqrt{1}mk}$$

b)

As a fuel  $\sqrt{1}$  mk i.

As ink solvent  $\sqrt[1]{1mk}$ ii.

c)

i. 
$$C_{3}H_{6(g)} \rightarrow CH_{4(g)} + C_{2}H_{2(g)}^{\sqrt{1}mk}$$

 $C_3H_{6(g)} + Br_{2(g)} \rightarrow C_3H_6Br_{2(g)} \sqrt{1}mk$ ii.

## **Ouestion 4**

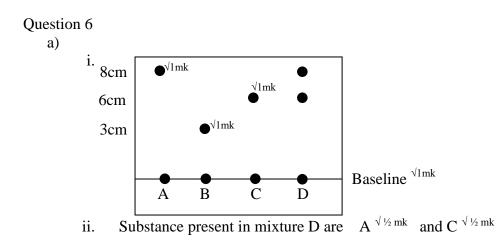
i.

- √1mk a) A – Potassium Nitrate / Sodium Nitrate √1mk
- b) Gentle warming / Moderate temperature
- c) Yellow  $\sqrt[1]{1mk}$  It contains dissolved nitrogen (iv) oxide  $\sqrt[1]{1mk}$
- d) To condense nitric (V) acid fumes  $\sqrt[1]{1mk}$
- e)
- √1mk Nitrogen (ii) oxide
- √1mk ii. Nitrogen (iv) oxide
- Nitrogen (ii) Oxide is oxidized by oxygen  $\sqrt[1]{1mk}$ iii.
- $3 \text{ Cu (NO_3)}_{2(aq)} + 2 \text{NO}_{(g)} + 4 \text{H}_2 \text{O}_{(l)}^{\sqrt{1}\text{mk}}$  $3Cu_{(s)} + 8HNO_{3(ad)}$ iv.  $\rightarrow$

- f)

b) 
$$2H_2O_{2(1)}$$
 MnO<sub>2</sub>  $2H_2O_{(1)} + O_{2(g)}$   $\sqrt{1}mk$ 

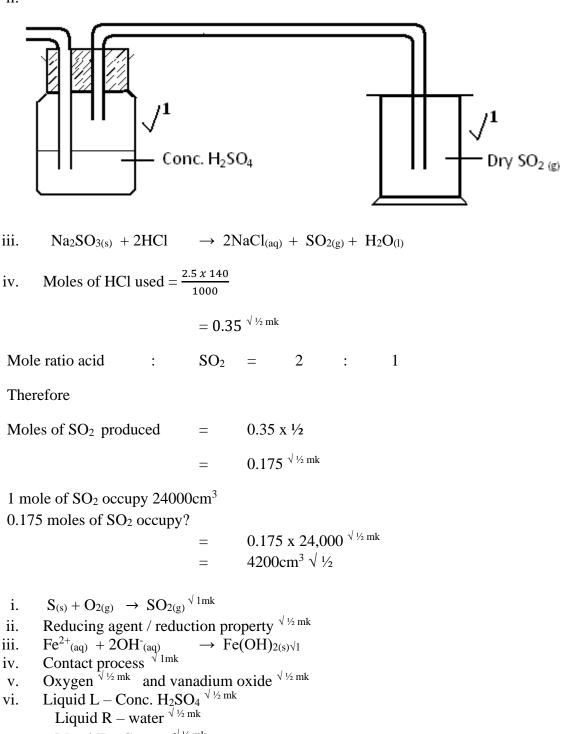
- i. To ensure that all the oxygen has been used up.  $\sqrt{1}$
- ii. For maximum contact between copper and oxygen so that reaction occurs completely.  $\sqrt[1]{lmk}$
- iii. The brown copper metal turned into black copper (ii) oxide.  $\sqrt{1}$  mk
- iv.  $2Cu_{(s)} + O_{2(g)} \rightarrow 2CuO_{(s)}$   $\sqrt{1}mk$
- v. Volume of oxygen used =  $110 87.5 \ cm^{3} \sqrt{\frac{1}{2} \text{ mk}}$ % of oxygen used =  $\frac{22.5}{110} \ x \ 100 \sqrt{\frac{1}{2} \text{ mk}}$ =  $20.5 \sqrt{\frac{1}{2} \text{ mk}}$
- vi. Making oxyacetylene flame used in welding .  $\sqrt[1]{lmk}$



- b) Add water to the mixture, stir to dissolve  $\sqrt[]{\frac{1}{2} \text{ mk}}$  calcium chloride as residue.  $\sqrt[]{}$  Evaporate the filtrate to dryness  $\sqrt[]{\frac{1}{2} \text{ mk}}$  to obtain solid calcium chloride.  $\sqrt[]{\frac{1}{2} \text{ mk}}$
- c)
- I. Fractional distillation  $\sqrt{1}$  mk
- II. Since the two liquids are immiscible, pour both liquids in a separating funnel  $\sqrt[4]{2} \text{ mk}}$  and allow them to settle. The denser liquid will settle at the bottom and the less dense  $\sqrt[4]{2} \text{ mk}}$  will form a second layer on top.  $\sqrt[4]{2} \text{ mk}}$  open tap and run out the liquid in the bottom layer leaving the liquid in the second layer in the funnel.  $\sqrt[4]{2} \text{ mk}}$

## i. Sodium Sulphite $\sqrt{1}$ mk

ii.



- Metal Z Copper  $\sqrt{\frac{1}{2}}$  mk
- vii. If liquid R (water) is used in step V it would react with  $SO_{3(g)}$  so exothermically that the acid vaporizes giving acid mist  $\sqrt[1]{2} \text{ mk}$ . The mist is not easily condensed and therefore results to serious explosions while use of liquid L.(Conc. H<sub>2</sub>SO<sub>4</sub>) in step vi would not cause dilution of oleum  $\sqrt[1]{1/2} \text{ mk}$ .

b)