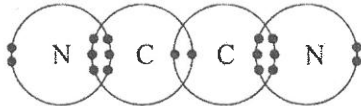


## SECTION B

### Part I

#### Marks

1. (a) (i) The mass number of a  $^{13}\text{C}$  atom is greater than that of a  $^{12}\text{C}$  atom. /  
The mass number of a  $^{12}\text{C}$  atom is 12 while the mass number of a  $^{13}\text{C}$  atom is 13. /  
A  $^{13}\text{C}$  atom has one more neutron than a  $^{12}\text{C}$  atom. /  
A  $^{12}\text{C}$  atom has 6 neutrons while a  $^{13}\text{C}$  atom has 7 neutrons. 1
- (ii) Let  $x$  be the percentage abundance of  $^{12}\text{C}$ . 2  
 $12x + 13(1 - x) = 12.011$   
 $x = 0.989 = 98.9\%$
- (b) Similarity: 1  
 Both diamond and graphite have a giant covalent structure.
- Difference: 1  
 In diamond, each carbon atom forms 4 covalent bonds with other carbon atoms.  
 In graphite, each carbon atom forms 3 covalent bonds with other carbon atoms.
- As there are no delocalised electrons in diamond, it is a poor conductor of electricity. 1  
 As there are delocalised electrons in graphite, it is a good conductor of electricity.
- (c) 1
- 
2. (a) in the presence of air / oxygen and water 1
- (b) phenolphthalein 1
- (c) • Iron loses electrons / is oxidised to form iron(II) ions /  $\text{Fe}^{2+}(\text{aq})$  ions. 1  
 •  $\text{Fe}^{2+}(\text{aq})$  ions react with potassium hexacyanoferrate(III) to form a blue colour. 1
- (d)  $\text{Y} > \text{iron} > \text{X}$  1  
 No blue colour is observed / No rusting occurs in set-up 2, which implies that Y loses electrons / is oxidised more readily than iron / Y gives sacrificial protection for iron.  
 Blue colour is observed / Rusting occurs in set-up 1, which implies that iron loses electrons / is oxidised more readily than X. 1

	Marks
3. (a) voltmeter / multimeter	1
(b) To complete the circuit / balance the charges in the two half cells / allow the flow of ions between the two half cells.	1
(c) (i) $\text{AgNO}_3(\text{aq}) / \text{Ag}^+(\text{aq})$ reacts with $\text{KCl}(\text{aq}) / \text{Cl}^-(\text{aq})$ to form water-insoluble $\text{AgCl}(\text{s})$ , which will block the ion flow in the salt bridge.	1
(ii) saturated / concentrated potassium nitrate solution / $\text{KNO}_3(\text{aq})$	1
(d) (i) Mn	1
Under using the same metal-metal ion in half-cell A, the voltage of chemical cell 3 is the greatest positive value / the highest.	1
(ii) $+1.15 - (-0.46) = +1.61$	1
(iii) $\text{Ag}^+(\text{aq}) + \text{e}^- \rightarrow \text{Ag}(\text{s})$	1
4. (a) (i) No. of moles of H = $3.09 / 1.0 = 3.09$ No. of moles of N = $14.42 / 14.0 = 1.03$ No. of moles of S = $33.06 / 32.1 = 1.03$ No. of moles of O = $49.43 / 16.0 = 3.09$  Mole ratio of H : N : S : O = 3 : 1 : 1 : 3 The empirical formula of sulphamic acid is $\text{H}_3\text{NSO}_3$ .	2
(ii) Let the molecular formula be $(\text{H}_3\text{NSO}_3)_n$ . $(3 \times 1.0 + 14.0 + 32.1 + 3 \times 16.0) \times n = 97.1$ $n = 1$ The molecular formula of sulphamic acid is $\text{H}_3\text{NSO}_3$ .	1
(b) (i) No. of moles of NaOH that reacted with the $\text{H}_2\text{SO}_4(\text{aq})$ $= 2 \times 0.202 \times 24.80 / 1000 = 0.010$	1
(ii) No. of moles of NaOH that reacted with the sulphamic acid $= 0.150 \times 100.0 / 1000 - 0.010 = 0.005$	1
(iii) No. of moles of the sulphamic acid that reacted with $\text{NaOH}(\text{aq})$ $= 0.486 / 97.1 = 0.005$  Mole ratio of $\text{H}_3\text{NSO}_3 : \text{NaOH} = 0.005 : 0.005 = 1 : 1$  The basicity of sulphamic acid is 1. / Sulphamic acid is a monobasic acid.	1
(c) $\text{CaCO}_3(\text{s}) + 2\text{H}^+(\text{aq}) \rightarrow \text{Ca}^{2+}(\text{aq}) + \text{CO}_2(\text{g}) + \text{H}_2\text{O}(\text{l})$	1

5. (a) Fractional distillation 1  
Different hydrocarbons evaporate and condense as different fractions at different temperatures. 1
- (b) (i) cracking 1
- (ii) Prevent sucking back of water as sudden cooling may break the hot boiling tube. 1
- (iii) The gas mixture contains alkenes which react with  $\text{Br}_2(\text{aq})$  to give colourless products. 1
- (iv) • Kerosene vaporises quickly without being cracked. 1  
• The broken unglazed porcelain is not hot enough and so cannot function as a catalyst for cracking of kerosene. 1
6. (a) (i)  $\text{NH}_3(\text{aq})$  and  $\text{FeSO}_4(\text{aq})$  1
- (ii) iron(II) hydroxide 1
- (b) (i)  $\text{Mg}(\text{NO}_3)_2(\text{aq})$  and  $\text{Na}_2\text{CO}_3(\text{aq})$  1
- (ii)  $\text{Mg}^{2+}(\text{aq}) + \text{CO}_3^{2-}(\text{aq}) \rightarrow \text{MgCO}_3(\text{s})$  1
- (c)  $\text{H}_2\text{SO}_4(\text{aq})$  and  $\text{Na}_2\text{CO}_3(\text{aq})$  1
7. (a) (i) 1
- 
- (ii) • C-H bonds are polar because carbon and hydrogen have different electronegativities. 1  
• C-Cl bonds are polar because carbon and chlorine have different electronegativities. 1
- (iii) Dichloromethane is a polar molecule, because the polarities of C-H bonds and C-Cl bonds do not cancel out each other. 1
- (b) • There are weak van der Waals' forces between dichloromethane molecules. 1  
• There are strong hydrogen bonds between ethanol molecules. 1

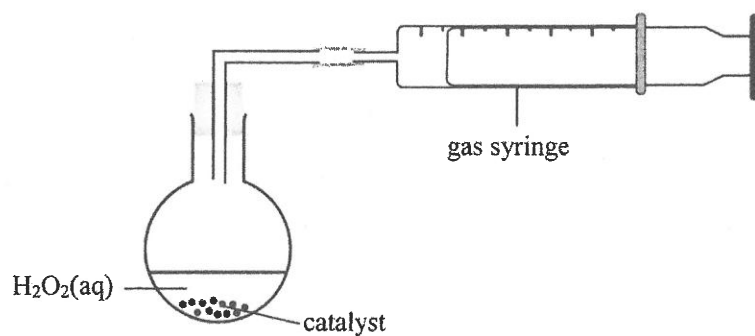
	<u>Marks</u>
8. (a) Energy released from the reaction = $1.0 \times (50.0 + 50.0) \times 4.20 \times 6.8 \div 1000 = 2.856 \text{ kJ}$ Enthalpy change of neutralisation = $-2.856 \div (1.0 \times 50 \div 1000) = -57.12 \text{ kJ mol}^{-1}$	2
(b) All acids and alkalis used in experiments 1 to 4 completely dissociate in water.	1
(c) Enthalpy change for the reaction = $-50.10 - (-57.12)$ = $+7.02 \text{ kJ mol}^{-1}$	2
9. Chemical knowledge	4
<ul style="list-style-type: none"> <li>• Silver can be obtained by heating silver oxide alone / directly.</li> <li>• Lead can be obtained by heating a mixture of lead(II) oxide and carbon / charcoal / coke.</li> <li>• Aluminium can be obtained by electrolysis of aluminium oxide in molten state.</li> <li>• The more reactive a metal is / The higher the position of a metal in the metal reactivity series is, the more difficult the extraction method of the metal is.</li> </ul>	
Communication mark	1

## Part II

Marks

10. (a)

1



(b) A, because it takes the shortest time for the reaction to finish / the initial rate of the reaction is the highest.

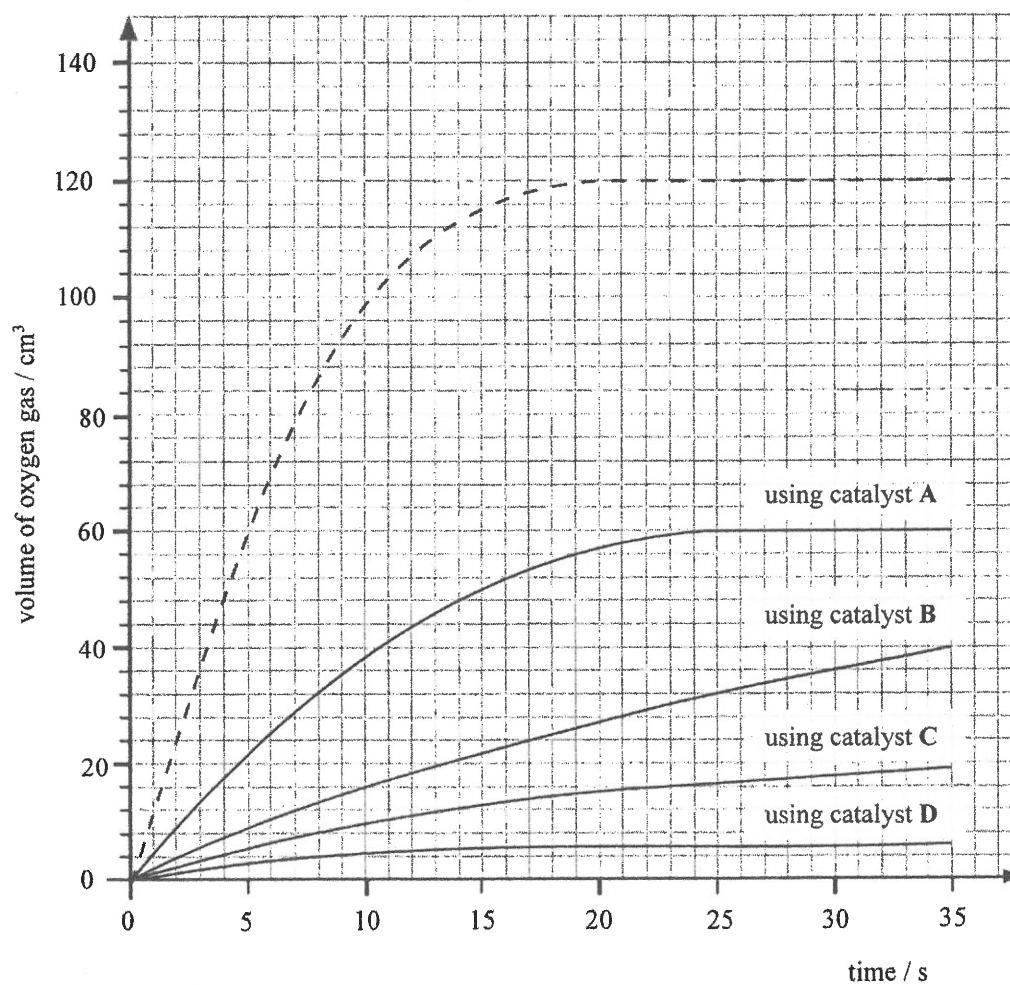
1

(c)  $60 \text{ cm}^3$

1

(d)

2



11. (a) 1.  $\text{LiAlH}_4$ , dry ether 2.  $\text{H}^+(\text{aq})$   
OR  
 $\text{NaBH}_4$

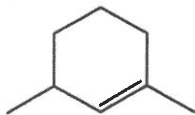
1

- (b) dehydration

1

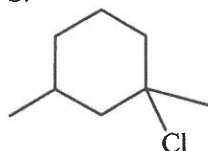
- (c) R:

1



- (d) S:

1



- (e)

Q	R

1

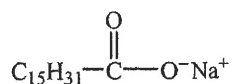
2

12. (a) propane-1,2,3-triol

1

- (b)

1



- (c) (i) There are no two distinct liquid layers after the mixture has rested for a long time. / One persistent milky layer is formed.

1

- (ii) • The ionic head ( $\text{COO}^-$ ) of product **B** dissolves in water / is hydrophilic, while the hydrocarbon tail ( $\text{C}_{15}\text{H}_{31}$ ) dissolves in oil / is hydrophobic.  
• By shaking, the oil breaks up into oil droplets. The repulsion between the negative charges on the oil droplets prevents the oil droplets from joining together again.

1

1

13. (a) •  $\text{Al}_2\text{O}_3$  is a basic oxide and reacts with  $\text{HCl}(\text{aq})$  to form  $\text{AlCl}_3$  and  $\text{H}_2\text{O}$ . /  $\text{Al}_2\text{O}_3(\text{s}) + 6\text{HCl}(\text{aq}) \rightarrow 2\text{AlCl}_3(\text{aq}) + 3\text{H}_2\text{O}(\text{l})$  1
- $\text{Al}_2\text{O}_3$  is an acidic oxide and reacts with  $\text{NaOH}(\text{aq})$  to form  $\text{NaAl}(\text{OH})_4$  /  $\text{NaAlO}_2$  and  $\text{H}_2\text{O}$ . /  $\text{Al}_2\text{O}_3(\text{s}) + 2\text{NaOH}(\text{aq}) + 3\text{H}_2\text{O}(\text{l}) \rightarrow 2\text{NaAl}(\text{OH})_4(\text{aq})$  1
- Aluminium oxide is both an acidic oxide and a basic oxide. 1
- (b) Both  $\text{S}_8$  and  $\text{Cl}_2$  have a simple molecular structure and there are weak van der Waals' forces between their respective molecules. 1
- The molecular size of  $\text{S}_8$  is larger than that of  $\text{Cl}_2$  and the van der Waals' forces between  $\text{S}_8$  molecules are stronger than those between  $\text{Cl}_2$  molecules. 1

## 14. Chemical knowledge

5

Observable change in both cases:

- The equilibrium mixture changes from pink to blue / becomes more intense in blue.

Addition of  $\text{NaCl}(\text{s})$ :

- When the concentration of  $\text{Cl}^-(\text{aq})$  ions increases, the equilibrium position shifts to the right.
- The equilibrium constant /  $K_c$  remains unchanged.

Increase in the temperature:

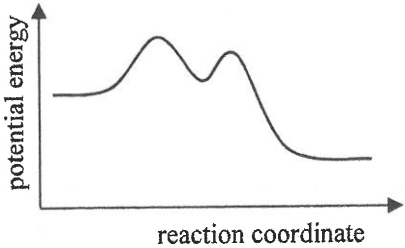
- As the forward reaction is endothermic, an increase in temperature favours the endothermic side of reaction and the equilibrium position shifts to the right.
- The equilibrium constant /  $K_c$  increases.

Communication mark

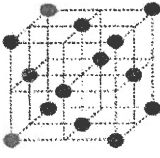
1

Paper 2

Marks

1. (a) (i) (1) Haber process 1
- (2) It is a fertiliser. / It can increase the crop yield. 1
- (ii) (1) concentrated sodium chloride solution / brine 1
- (2) No toxic mercury will be produced by membrane electrolytic cells but toxic mercury may leak out from flowing mercury cells. / The operation of membrane electrolytic cells requires less energy than flowing mercury cells. 1
- (iii) Yeast provides enzyme / catalyst. 1
- (b) (i) 2
- 
- (ii)  $\frac{4 \times 60.0}{2 \times 58.0 + 5 \times 32.0} \times 100\% = 87.0\%$  1
- (iii)  $\text{CH}_3\text{OH} + \text{CO} \rightarrow \text{CH}_3\text{COOH}$  1
- (iv) HI is regenerated at the end of the reaction. 1
- (v) (1) Toxic  $\text{CH}_3\text{OH}$  / CO and corrosive HI are used in Method 2 while these substances are not used in Method 1. 1
- (2) The atom economy is 100% in Method 2 but not in Method 1. / Method 2 has higher atom economy than Method 1. 1
- (c) (i) Use a gas syringe to follow the change in the volume of the gaseous mixture at regular time intervals. / Use a pressure sensor connected to a data logger to follow the change in the pressure of the gaseous mixture at regular time intervals. 1
- (ii) (1)  $a = 1$  1
- (2)  $\log \frac{k_2}{k_1} = \frac{Ea}{2.3R} \left( \frac{1}{T_1} - \frac{1}{T_2} \right)$  2
- $$\log \left( \frac{k}{4.29 \times 10^{-4}} \right) = \frac{108500}{2.3(8.31)} \left( \frac{1}{318} - \frac{1}{338} \right)$$
- $$k = 4.88 \times 10^{-3} \text{ s}^{-1}$$
- (iii) (1) B 1
- (2) • An increase in temperature can increase the average kinetic energy of the molecules. 1
- There are more molecules having kinetic energy equal to or greater than the activation energy. 1
- This will increase the effective collision frequency / the number of effective collisions per unit time. 1

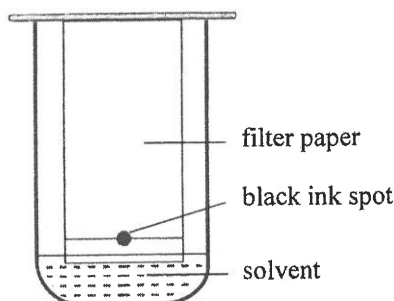


2. (a) (i) • When a rubber band is stretched, the polymer chains move only slightly without breaking the disulphur cross-linkages. 1  
 • When the stretching force is removed, the polymer chains are still held together by the cross-linkages, and they return to their original position. 1
- (ii) (1)  $\text{NHCOCH}_3$  / acetylamino group 1  
 (2) There are hydrogen bonds between the polymer chains of chitin. 1
- (iii) smectic phase 1
- (b) (i) ester group /  $\text{—}\overset{\text{O}}{\parallel}\text{C—O—}$  1
- (ii)  $\begin{array}{c} \text{CH}_3 \\ | \\ \text{HO—C—COOH} \\ | \\ \text{H} \end{array}$  1
- (iii) PLA is formed by joining up the monomer molecules repeatedly with the elimination of small molecules. 1
- (iv) Corn starch is a renewable resource. 1
- (v) PLA is non-toxic / biocompatible. 1
- (vi) Not suitable, because PLA is a thermoplastic. 1
- (vii) Because ethyl ethanoate molecules and PLA molecules are polar, while hexane molecules are non-polar. / The intermolecular attractions in ethyl ethanoate and PLA are similar while that in hexane is different. 1
- (c) (i) 8 1
- (ii) (1)  1
- (2) A-B-C type packing 1
- (iii) The deduction is incorrect. As the temperature increases, the iron metal expands on heating, so the density of iron at 1450 °C is lower. 1
- (iv) (1) Kevlar 1  
 • It has a higher tensile strength than iron. 1  
 • It has a lower density than iron. 1
- (2) • There are more hydrogen bonds between the polymer chains of Kevlar than those of nylon-6,6. 1  
 • Kevlar has many benzene rings which restrict the movement between the polymer chains, but nylon-6,6 does not have. 1

3. (a) (i) • Add 2,4-dinitrophenylhydrazine. 1  
 • Compound **A** forms a yellow / orange / red precipitate while compound **B** does not. 1  
 OR  
 • Add  $\text{Na}_2\text{CO}_3(\text{aq})$  /  $\text{NaHCO}_3(\text{aq})$ . / Add deionised water and then add  $\text{Mg}(\text{s})$ .  
 • Compound **B** gives out a colourless gas while compound **A** does not.

(ii)

2



(iii) Any one of the following:

1

- These solvents are immiscible with water.
- These solvents do not react with the organic compounds.
- These solvents can dissolve more organic compounds than water can.
- These solvents have low boiling points / can evaporate easily.

(b) (i) To ensure that all  $\text{PbCO}_3(\text{s})$  and  $\text{CuCO}_3(\text{s})$  have completely reacted with dilute  $\text{HNO}_3(\text{aq})$ . 1

(ii) A deep blue solution is formed. 1

(iii) (1) • Washing is to remove water soluble impurities and drying is to remove water. 1  
 • To obtain a more accurate mass of  $\text{Pb}(\text{OH})_2(\text{s})$ . 1

(2) lead(II) hydroxide /  $\text{Pb}(\text{OH})_2$  1

(3) Percentage by mass of  $\text{PbCO}_3$  in the solid sample 2

$$= \frac{1.47}{(207.2 + 16.0 \times 2 + 1.0 \times 2)} \times (207.2 + 12.0 + 16.0 \times 3) \div 2.03 \times 100\%$$

$$= 80.22\%$$

(c) (i) The infra-red spectrum of the indoor air sample shows a strong absorption peak in the region from  $1680 \text{ cm}^{-1}$  to  $1800 \text{ cm}^{-1}$  corresponding to the carbonyl group /  $\text{C}=\text{O}$  bond of formaldehyde. 1

(ii)  $\text{HCO}^+$  1

(iii) (1) colorimeter 1

(2) The absorbance is directly proportional to the concentration of formaldehyde. 1

(3) From the graph, the concentration of formaldehyde in solution **S** is  $0.25 \text{ mg dm}^{-3}$ . 2  
 Mass of formaldehyde =  $0.25 \times 0.03 = 0.0075 \text{ mg}$

(4) Concentration of formaldehyde in the indoor air sample 1  
 $= 0.0075 / 0.1 = 0.075 \text{ mg m}^{-3} < 0.1 \text{ mg m}^{-3}$   
 Therefore, the quality of the air inside the room is good.

(5) Instrumental analytical methods are sensitive enough to measure very low levels of formaldehyde. 1