

Universal gas constant  $R = 8.314 \text{ J K}^{-1} \text{ mol}^{-1}$ 

\* Avagardro's Constant  $N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$ 

Time 3 Hours

\* You are allowed to show the alkyl groups in the contrected form

eg: 
$$H - C - C$$
 group can be shown as  $CH_3CH_2$ 

#### Part - A Structured essays

- \* Answer all questions in this paper.
- \* Provide answers for each question in the space given . This space is sufficient for the answer and we do not expect long answers

Part B & Part C - Essay

- \* Select two questions from each part and answer four questions all together.
- \* Attach part A, Part B, and part C together, keeping Paper A on top of the answer script and hand over the completed answer script to the supervisor.
- \* You are permitted to take Part B of the question paper out of the examination centre.

## For Examiners' use only.

Part	Question No	Marks	
A	2		
11	3		
	4		
	5		
В	6		
	7		
	8		
С	9		
	10		
Total			
Percentage	7		

Final Mark				
In Figures				
In Letters				

#### **Code Numbers**

Examiner	
Checked by 1.	
2.	
Supervision	2.37

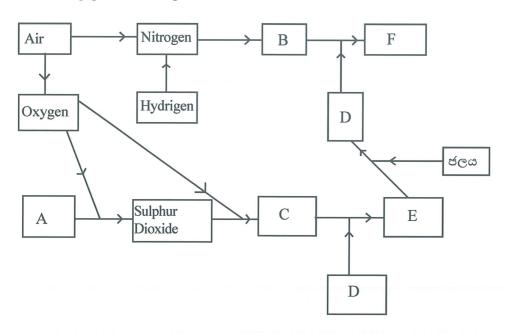
# Part A - Structured Essy type

Answer all four questions on this paper itself ( marks allocared for each question is 10)

1. (	(a) i) Write the election configuration of element X which has the atomic number 24 in the normal way like 1s², 2s²
	ii) Mention the for common oxidation states of the element X
	iii) Write the chemical formula of the oxide which is derived from the highest oxidation state of X
	iv) If the oxide mentioned in (iii) reacts with conc. HBr to give write down the balanced chemica equation for the relevant reaction.
	(b) Answer the following parts (I) to (v) based on the structure of hydroayonic acid given below  HOCN  (i) Drow the most acceptable Lewis Liner structure of this compand
	(ii) Draw the resonance Structures for the above compound and give reasons for their relative stability.
	<ul> <li>(iii) Deduce the shape ,around each atom given below of the structure you mentioned in (b)(I) using VSEPR theory</li> <li>i) Around O ,</li> <li>2) Around C</li> </ul>
	(iv) Identity the orbitals that participate in forming of bonds C-N and O-H hydrocyamic acid.  (1) C

	sabstance	Primary Interaction (As Ionic/polar covalent/Non-	Secondary Interaciton (As Bipolar-Bipalar/Hydrogen bonds/ London forces)
		Polar Covalent)	
i)	Ice (Solid)		·
i)	Silicondioxide(solid)		
įiii	Hydrogen fluoride		
iv)	Magnesium Sulphide (solid)		
(v)	Chlorine (gas)		
(vi)	Mercury (Liquid)		-
(vi)	Mercury (Liquid)  (i) Write the formulae for the lariodic table. And mention their a Formula	Acidic-basic nature  Acidic-basic nature	m to chlorine in the third period the ly. Formula Acidic-basic nature
(vi)	Mercury (Liquid)  (i) Write the formulae for the lariodic table. And mention their a Formula	Acidic-basic nature Acidic-basic nature	ly. Formula Acidic-basic nature

(b) Given below is a flow chart that shows two important industrial processes. Answer the following questions using it.



(i) Give the formulae of the substances denoted by A,B,C,D,E, and F

A	В
C	D
E	F

.....

- (ii) Mention another use of  $N_2(g)$  and  $O_2(g)$ .
- (iii)  $N_2(g)$  සහ  $O_2(g)$  හි වෙනත් පුයෝජනයක් බැගින් සඳහන් කරන්න.

$$N_2(g)$$
..... $O_2(g)$ .....

(iv) Name another important product obtained by the process in (ii) and mention one of its uses.

(v) State desperately the catalysts used in the two industrial processes shown in the above flow chart

Industrial Process	Catalysts	

(c)	of	ICl <sub>n</sub> v	nd chlorine react with each other to give a compound named ${\rm ICl_n}$ when $1{\rm x}10^{\text{-3}}{\rm mo}1$ was reacted with excess amount of KI, all the iodine in that compound was converted to ${\rm Imol~dm^{-3}~Na_2S_2O_3}$ solution was used to react completely with that ${\rm I_2}$
		(i) (	Calculated the value of n.
		(ii) <sup>1</sup>	Write the balanced chemical equation for the reaction between Cl <sub>2</sub> and I <sub>2</sub>
3.	a)	(i).	I Deduce Charlis law using the equation PV = nRT.
	(ii)	pre	hass of 12.0 g of an inert gas was in a vessel of volume V dm³ cooperative t $^{\circ}$ C .The source of that gas was $1.0\times10^{5}$ Nm⁻² . The pressure of the gas increased by 10% when the perature was increased by 10 $^{\circ}$ C
		(	IE. Molar mass of the inert gas is 120g mol <sup>-1</sup> )
			I) Calculate the initial temperature t of the gas
			II) Calculate the volume,V of the vessel
			III) write douwn the assumptions you make for the above calculation.

reach equilibrium. Then it is observed that 0.80 g of X isleft behind in aqueous phase.  (Motar mass of Xis වල මවුලික ස්කන්ධය 125 g mol+)  (I) What is the mass of X, extracted in to ether?  (II) Calculate the coefficient of distribution of X, between ether and water at the relevant temperature.  (III) If the extraction was done by using 4 samples of 25 cm² of ether each, instead of 100 cm² of ether what is the mass of X extracted in to ether?  (III) If the extraction was done by using 4 samples of 25 cm² of ether each, instead of 100 cm² of ether what is the mass of X extracted in to ether?  (III) Answer the following questions using the data given for the reaction  2 CO(g) + O <sub>2</sub> (g)	b)			organic compound,X is in 50.0 cm <sup>3</sup> of aqueous solution. To extract the solution was well stirred with 100.0 cm <sup>3</sup> of ether and allowed to	
(I) What is the mass of X, extracted in to ether?  (II) Calculate the coefficient of distribution of X, between ether and water at the relevant temperature.  (III) If the extraction was done by using 4 samples of 25 cm³ of ether each, instead of 100 cm³ of ether what is the mass of X extracted in to ether?  (C) Answer the following questions using the data given for the reaction  2 CO(g) + O₂(g) : → 2 CO₂(g)  9 හිම්යාව හා සම්බන්ධව දී ඇති දක්ක ඇතුරින් අසා ඇති පුශ්ණ වලට පිළිතුරු සපයන්න.  (B) G₁ CO₂(g) -394.4 kJ mol¹  G₂ CO(g) -137.2 kJ mol¹  A S for the reaction -0.188 kJ mol¹ k¹  1) Show that the above reaction is exothermic at 300 K			ch equilibrium. Then it is	s observed that 0.80 g of X isleft behind in aqueous phase.	
(II) Calculate the coefficient of distribution of X, between ether and water at the relevant temperature.  (III) If the extraction was done by using 4 samples of 25 cm³ of ether each, instead of 100 cm³ of ether what is the mass of X extracted in to ether?  (C) Answer the following questions using the data given for the reaction  2 CO(g) + O <sub>2</sub> (g) → 2 CO gSSSSSSSSSSSSSSSSSSSSSSSSSSSSSSSSSSSS		(I)			
temperature.  (III) If the extraction was done by using 4 samples of 25 cm³ of ether each, instead of 100 cm³ of ether what is the mass of X extracted in to ether?  C) Answer the following questions using the data given for the reaction  2 CO(g) + O₂(g) → 2 CO₂(g)  9 තිබුයාව හා සමබන්ධව දී ඇති දත්ත ඇසුරින් අසා ඇති පුශ්ණ වලට පිළිකුරු සපයන්න.  G₁ CO₂(g) -394.4 kJ mol¹  G₂ CO(g) -137.2 kJ mol¹  Δ S for the reaction -0.188 kJ mol¹ k¹  1) Show that the above reaction is exothermic at 300 K		( )			
temperature.  (III) If the extraction was done by using 4 samples of 25 cm³ of ether each, instead of 100 cm³ of ether what is the mass of X extracted in to ether?  C) Answer the following questions using the data given for the reaction  2 CO(g) + O₂(g) → 2 CO₂(g)  9 තිබුයාව හා සමබන්ධව දී ඇති දත්ත ඇසුරින් අසා ඇති පුශ්ණ වලට පිළිකුරු සපයන්න.  G₁ CO₂(g) -394.4 kJ mol¹  G₂ CO(g) -137.2 kJ mol¹  Δ S for the reaction -0.188 kJ mol¹ k¹  1) Show that the above reaction is exothermic at 300 K					
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of ether what is the mass of X extracted in to ether?  C) Answer the following questions using the data given for the reaction $ 2 CO(g) + O_2(g) \longrightarrow 2 CO_2(g) $ පතිතියාව හා සම්බන්ධව දී ඇති දක්ක ඇසුරින් අසා ඇති පුශ්ණ වලට පිළිතුරු සපයන්න. $ G_f CO_2(g) \qquad -394.4 \text{ kJ mol}^{-1} $ $ G_6 CO(g) \qquad -137.2 \text{ kJ mol}^{-1} $ T) Show that the above reaction is exothermic at 300 K					
2 CO(g) + O₂(g)		(III)			<sub>1</sub> 3
2 CO(g) + O₂(g)					
2 CO(g) + O₂(g)					
$G_{\rm f}  {\rm Co}_2({\rm g}) \qquad \qquad -394.4  {\rm kJ  mol^{-1}}$ $G_{\rm f}  {\rm Co}({\rm g}) \qquad \qquad -137.2  {\rm kJ  mol^{-1}}$ $\Delta  {\rm S}  {\rm for  the  reaction} \qquad \qquad -0.188  {\rm kJ  mol^{-1}  k^{-1}}$ $I)  {\rm Show  that  the  above  reaction  is  exothermic  at  300  K}$	C)	Answ	er the following question	ns using the data given for the reaction	
G <sub>g</sub> CO(g) -137.2 kJ mol <sup>-1</sup> A S for the reaction -0.188 kJ mol <sup>-1</sup> k <sup>-1</sup> Show that the above reaction is exothermic at 300 K		2 C පුතිද්	$\mathrm{CO}(\mathrm{g}) + \mathrm{O}_2(\mathrm{g}) \xrightarrow{\cdot}$ බුයාව හා සම්බන්ධව දී	2 C්ଠ୍2(g) ඇති දත්ත ඇසුරින් අසා ඇති පුශ්ණ වලට පිළිතුරු සපයන්න.	
△S for the reaction -0.188 kJ mol <sup>-1</sup> k <sup>-1</sup> I) Show that the above reaction is exothermic at 300 K			$G_{\rm f}  {\rm CO}_2({\rm g})$	-394.4 kJ mol <sup>-1</sup>	
I) Show that the above reaction is exothermic at 300 K			G CO(g)	-137.2 kJ mol <sup>-1</sup>	
			ΔS for the reaction	-0.188 kJ mol <sup>-1</sup> k <sup>-1</sup>	
		I)	Show that the above re	action is exothermic at 300 K	
II) Deduce whether above reaction occurs spontaneously at 300 K					
		II)	Deduce whether above	e reaction occurs spontaneously at 300 K	
III) Mention the assumptionns you made for the above calculations.		III)	Mention the assumpti	onns you made for the above calculations.	

4)	a.) P, Q and R are the three halogenated hydrocarbons with the common molecular formula of $C_5H_{10}Br_2$ but have three different structural formulae. When they are treated separately with aqueous KOH in alcoholic media, it gives same chemical compound S as the main product. Compound Q shows optical (enantiomeric) isomerism, and compound R reacts with aqueous KOH to give a symmetries ketone as the product. Identify the compounds P,Q and R separately and write their structure in the boxes below
I)	Identify the compounds P,Q and R separately and write their structure in the boxes below.
	P Q R
II)	The above compound was subjected to fractional hydrogenation in the presena of S Pt/Cd catalyst.  When the product T thus obtained is brominatid, you can get the compound Q again  Write a suitable mechanism for the conversion of T ↑ Q
b)	i) Show the method for the following conversion in not more than 6 steps. $CH_3$ - $CO$ - $CH_2CH_3$ ————————————————————————————————————
	Use another compound ,which is not a carbonyl compound and have Sp² hybridized carbon atoans.  ii) Drow the Structure of the main products that 2-butanone gives in the presence of a dilute alkali
	iii) Comment on the isomerism of the structures mentioned in (ii)



# G.C.E. (A/L) Student Support Seminar - 2013

Revision Paper - Chemistry II

Launched under the Supervision of Ministry of Education

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Time 2 hours

- \* Universal gas constant  $R = 8.314 \text{ J K}^{-1} \text{ mol}^{-1}$
- \* Avogadro constant  $N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$

### Part B --- Essay type

Answer **Two** questions only (15 marks for each question)

- (05) (a) (I) I clarify the following terms using suitable balanced equations.
  - (i) Standard Enthalpy of atomization of bromine
  - (ii) Standard Enthalpy of bond dissociation of bromine
  - (iii) Enthhalpy of neutralization of acetic acid
  - (II) State and experiment that can be done in the laboratory to find the enthalpy of neutralization of acetic acid show how the calculation is done mentioning the assumptions make
  - (III) The stable allotropic form of phosphorous  $P_4$  (white phosphorous) borms in excess oxygen and gives a white powder with formula  $P_4O_{10}$  Calculate the enthalpy of sublimation of in symbolic form  $P_4O_{10}$  Therme-Cheis done mentioning the assumptions make

Enthalpy of formation of  $P_{(g)}$   $\Delta H_f P_{(g)} = y kJ mol^{-1}$ 

Enthalpy of formation of  $O_{(g)}$   $\Delta H_f O_{(g)} = x \text{ kJ mol}^{-1}$ 

Enthalpy of combustion of of  $P_{4(s)}$   $\triangle H_c P_{4(s)} = z \text{ kJ mol}^{-1}$ 

Enthalpy for the conversion of 142.0 g of  $P_4O_{10(s)}$  to gaseous atoms = a kJ mol<sup>-1</sup>

(P = 31; O = 16)

- (b) A Certain mass of solid  $NH_4Cl$  was kept in a vessel the volume of which was 0.5 dm³, at the temperature of 400 K and allowed to attain to the equilibrium at that temperature. The barometer reading pf that system in equilibrium was  $8314 \times 10^5 \, N \, m^{-2}$ .
  - (i) If the partial pressure of the gas ammonic at 400 K is 05 atm. What should be the partial pressure of the gas HCL? (1 atm = 1 x  $10^5$  Nm<sup>-2</sup>)
  - (ii) What should be the  $K_{c}$  of the system at that temperature?
  - (iii) Another 10 mal of  $NH_4Cl$  was added to the system, maintaining the same temperature and allowed to attain to the equilibium. calculate the mass of HCL in the system in equlibrium.

$$(H = 1; S = 16)$$

 $_{\mu}O_{10(s)}$ 

- (iv) If 2.0 moles of He gas was added to the system in equilibrium in part (III), how will it affect to the partial pressure of HCL in the system? Explain this qualitatively.
- (v) Suggest a suitable experiment to find the equilibrium constant K<sub>p</sub> of the system in equilibrium formed by dissociating the above NH<sub>4</sub>Cl<sub>(s)</sub> at 500 K
   (Experimenttal details are not necessary)
- (06) (a) (I) What do you mean by the form buffer solution?
  - (II) A buffer solution containing the salt BCL results when excess amount of the base BOH in reacted with HCL. Show that the , pOH value of this solution.

$$pOH = pK_b + log \frac{[BCl_{(aq)}]}{[BOH_{(aq)}]}$$

(III) A volume of 400cm<sup>3</sup> of the week base BOH was reacted with a solution of HCL in the concentration of 0.1 mol dm<sup>-3</sup> .volumes of HCL added and pH values of the solution in each instance is given below.

	Valuve of HCl Valuve	pH of the solution
1.	5.00 cm <sup>3</sup>	10.04
2.	20.00 cm <sup>3</sup>	9.14

A value of Vcm³ HCl, was necessary to neutrolize the base competely.

- (i) Give two expressions for the concentiations of the ase, left behind in the solution without reacting, in the instances that pH is 10.04 and 9.14
- (ii) what is the voluve of above V cm<sup>3</sup>?
- (iii) calculate the dissociation censtant ((K<sub>b</sub>)of the base at the relevent temperature.
- (iv) calculate the dissociation censtant (k<sub>b</sub>) of the base at the relevent temperature (i.e; Consider that ionic product of water is 1 x 10<sup>-14</sup> mol <sup>2</sup>dm-<sup>6</sup> at relevent temperature.)
- (b) (i) Hg<sub>2</sub>I<sub>2(s)</sub> is a solid ionic compound which is slightly soluble in water. Deduce and expression for the Kspof this compound using relevant principles.
  - (ii) There are calions Ag<sup>+</sup>(aq) and Hg<sub>2</sub><sup>2+</sup>(aq) in and aqueous solution. Concentrations of each of there are  $\Im$ 1 mol dm<sup>-3</sup> in relation to each other. Show that which cation precipitates first as as iodide, when a solution of NaI is added gradually in to the above solution

( Ksp of AgI 
$$_{(S)}$$
 = 8.5 x  $10^{-17}$  mol $^2$ dm $^{-6}$  Hg $_2$ I $_2$   $_{(S)}$  of Ksp of = 2.5 x  $10^{-26}$  mol $^3$ dm $^{-9}$  at the relevant temperature)

- (iii) At the Moment that second cation starts to precipitale.
  - (a) What is the precentage cation concentration left behind, which stared to precipate frist?
  - (b) What is the precipitaled cation concentration which started to precipitate frist?

- (7) (a) Give reasons for the followings
  - (i) A pure sample of CCl<sub>4</sub> can not be prepared by the reaction of methene gas with chlorine gas in the presence of difused sunlight.
  - (ii) Acidic properties are higher in mitrophenols than in phenol.
  - (b) (i) State how the following conversion is done minimum number of steps using CH<sub>3</sub>MgBr as the only onlyrganic compound, and using the given inorganic reagents appropriatily.

$$\begin{array}{c} {\rm C_6H_5CH_2Br} \rightarrow {\rm C_6H_5CH_2}\text{- C=N CH_2C_6H_5} \\ {\rm CH_3} \\ {\rm [inorgamic\ recigents:\ H_2O,\ dil.\ H_2SO_4\ NH_3(\emph{I})\ CaC_2(s)$} \\ {\rm HgSO_4(aq),\ LiAlH_4,\ KMnO_4]} \end{array}$$

- (c) How do you deferentiate the following pairs of compounds using minimum number of chemical tests?
  - (i) C<sub>6</sub>H<sub>5</sub>CONHC<sub>6</sub>H<sub>5</sub> &D C<sub>6</sub>H<sub>5</sub>CONHCH<sub>2</sub>C<sub>6</sub>H<sub>5</sub> (ii) CH<sub>3</sub>CH<sub>2</sub>Cl &D (CH<sub>3</sub>)<sub>3</sub>CCl (iii) HCOOCH<sub>3</sub> &D CH<sub>3</sub>COOCH<sub>3</sub>

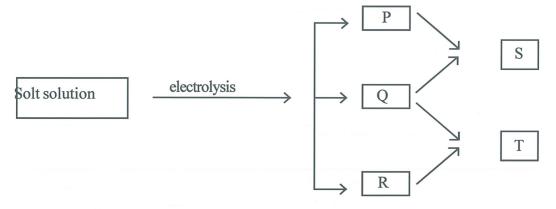
# Part C- Essay Type Answer Two Questions only (15 Marks for eachquestion)

- (8) (a) There is only one unpaired electron in M, which ia a 3d element in the periodic table .M does not react with dil.HCL, but reacts with did. HNO<sub>3</sub> and gives a colorless gas G and a coloured solution L
  - i. Identify M, giving reasons
  - ii. Mention two industrial uses of M.
  - iii. State a chemical test, to identify the gas G.
  - iv. Write the balanced chemical equation for the reaction of M with dil-HNO,
  - v When excess conc.HCl is added to solution L, the result is cloured solution Q. And when excess conc. NH<sub>3</sub> is added to solution L, the result is coloured solution R. Mention the colours of L,Q andR; the shapes of those coloured species and their UPAC name separately.
  - (b) X, Y, Z and aqueous solutions of the salts of three elements that belong to the d -block .Aqueous NaOH solution was added to each solutions above Separately
    - X: A white preciitation was obtained.it dissalved in excess aqueous NaOH solution, as well as in excess aqueous NH<sub>3</sub> solution
    - y: A green precipitation was obtained. it did not dissolve in excess NaOH solution, but dissolved in excess aqueous NH<sub>3</sub> solution.
    - Z: A yellowish brown precipitation was obtained it did not dissolve in excess NaOH solution or in excess NH<sub>3</sub> solution.
    - i Identity X,Y, and Z using the above observations.
    - ii if you are suppled with a solid mixture of the elements X,Y, and Z in part (I) how do you determine the percentage, by weight, of those elements?

(c) 2.39g of a solid mixture, which contains only ammonium ahloride and ammonium sulphate, was dissolved in water and the solution was prepared to a volume of 250.0 cm<sup>3</sup> .25.0 cm<sup>3</sup> of that solution was taken in to a beaker and 50.0 cm<sup>3</sup> of 0.2 mol dm -3 NaOH solution was added. this mixture was boiled to remove all the ammonia and was litrated with 0.3 mol dm -3 HCl solution valume of Hcl used was 20.0 Cm<sup>3</sup>

$$(N=14; H=1; S=32; Cl=35.5; O=16)$$

- i Write the balnce chemical reactions occured here
- ii Calculate the mole presentage of NH<sub>4</sub>Cl in the mixture.
- (9) a Substance P,Q and R, obtained by electrotysius a solution of NaCl (common solt)are used as raw materials for many industrial processes.)



(Here P,Q,R,S and T are not accepted symbols)

- i Write the formulae of the sulstances P, Q, R, S, and T
- ii Mention one use each of P, Q, R, S, and T
- iii Clarify the Clarify the physio-chemical principles applied in the above eletrolytial process in industry.
- iv what are the adverse effects to the environment in this industrial process ?(only four points will be susufficient)
- A solution the volume of which is 1000cm³ was prepared by dissolving sodium oxalate and anhydrous oxalic acid in water. 25.0 cm³ of this solution was litrated with 0.1 mol dm³ NaOH using phenolphthalane as indicator volume of NaOH solution used here was 17.6 cm³ 40 cm³ of 0.2 mol dm³ KMnO₄ solution was used to neatralise another 25.0 cm³ of the inertial solution under acidic condition. Calculate separately the concentrations of sodium oxalate and oxalic acid in the solution.
- Disprin the common pain killer contains acetyl solicilyc acid mixed with CaCO<sub>3</sub> and starch.
- (i) wwhat is the comon name for acetyl salicily acid?

- (ii) what is the chemical reaction that you expect, when a tablet of disprin is put in to water?
- (iii) How do you show, in the laboratary, that a tablet of disprin contains the three constituents mentioned?
- (iv) Mention briefly a test to determine the amount of CaCO<sub>3</sub> quantitatively, in a tablet of disprin.
- 10 (a) I. Consider the following reaction

$$2NO(g) + Br_2(g) \longrightarrow 2NOBr(g) \triangle H = -23kJ \text{ mol}^{-1}$$

This is a second order reaction with respect to [NO] and is a first order reaction with respect to  $[Br_2]$ . Activation enargy of the above reaction is  $+ 5.4 \text{ kJ mol}^{-1}$ .

- i Write the rate equation and mentioned the units of rate rate constant for the above reaction, dipicting all the energy changers
- ii Draw a tablet energy diagram to show the pathway of the above reaction depicting all the energy changers.
- II NO(g) gas reacts with O<sub>2</sub> (g) to give NO<sub>2</sub> (g) according to the following equation.

 $2NO(g) + O_2(g)$   $2NO_2(g)$  results of an experiment done to determine the order of above reaction is given below.

Experiment	Initial concentration (mol dm <sup>-3</sup> ,)		Initial rate of formation
Experiment	[NO]	Ο,	( mol dm <sup>-3</sup> s <sup>1</sup> )
1	1×10 ³	1×10 <sup>-3</sup>	7×10 <sup>-6</sup>
2	1×10-3	2×10 <sup>-3</sup>	14×10 <sup>-6</sup>
3	1×10 <sup>-3</sup>	3×10 <sup>-3</sup>	21×10 <sup>-6</sup>
4	2×10 <sup>-3</sup>	3×10 <sup>-3</sup>	84×10 <sup>-6</sup>
5	3×10 <sup>-3</sup>	3×10 <sup>-3</sup>	189×10 <sup>-6</sup>

- i) Determine the rate in relation to NO and O<sub>3</sub>
- ii) Write the rate equation that shows the steps od determing the above speed.
- iii) It is seen that the steps determining the speed have is abnormal. what could be the reasen for that?
- (b) i) Graphite layer is applied on plastic items before they electroplated with silver. clarify the reason for this
  - ii) what is the other electrode used in the electrolytic cell, in the above electro plating?
  - iii) Draw a labeled diagram of the electrical circuit with the electrolytic cell used for the above electroplating
  - iv) How long should b an electrical current of 0.1 A be flown through the electrolytic cell to deposit 1.089g of silver in the plastic item?

(Ag = 108) Faraday constant 96500 C mol<sup>-1</sup>)

v) Is it suitable to apply copper initially on the plastic item, instead of graphite, in this electroplating? Give your ideas.

- (c) Chlorine is used in water purification.
  - i) Write the balanced equation for the reaction of chlorine with water.
  - ii) Clarify how bacteria are destroyed by this solution
  - iii)  $P^h$  value of the above solution should be controlled carefully to maintain the activity of  $Cl_2$  in it. Clarify the reason for this using the equation in (c) I.
  - iv) Mentioned two other methods other than chlorination ,to purity water