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B.Sc. RNLK-/CC-1(T)/22

2022

Chemistry [First Semester] Paper - CC-1(T) Full Marks : 40 Time : 2 hours The figures in the right-hand margin indicate marks. Candidates are required to give their answers in their own words as far as practicable. Illustrate the answers wherever necessary.

Group - A

Answer any five questions : 5×2=10 (a) A gas expands (i) isothermally from volume v₁ to v₂ (ii) Adiabatically from volume v₃ to v₄ in a leak proof container. Identify the type of system in each case (close / open/ isolated). (b) For a reversible evalue process A System is less than

(b) For a reversible cyclic process Δ System is less than Δ S surroudings justify or criticise.

(Turn Over)

- (2)
- (c) For an ideal gas obeying the Maxwellian distribution of molecular speed in three-dimension. Find the

maximum value of $\left(\frac{1}{n}\right)\frac{dN}{dc}$ for a gas of molar mass 4.0 gm mol⁻¹ kept at 127°C.

- (d) Calculate a_{b} for a gas for which T_{B} =500 K.
- (e) Half-life a reaction is doubled when the initial concentration of a reactant. What is the order of the reaction with respect to the reactant?
- (f) Calculate the percentage increase of rate comtant for a 5% increase in temperature. given E=20KJ Mol⁻¹ and T=300K.
- (g) Define turn over number.
- (h) Catalyst is like a tunnel in chemical reaction -Explain.

Group - B

Answer any four from the following : $5 \times 4 = 20$

2. (a) (i) Derive an expression of average time required to travel unit distance for gas molecule. 3

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(Continued)

- (3)
- (ii) A gas obeying the P(V-b)=RT has the compressibility factor Z=1.0018 at 27°C and 1 atm pressure. Assuming the gas molecule to be spherical in shape. Calculate its molecular weight.
- (b) (i) Find the unit of $\frac{dlnk}{dT}$ for a nth order reaction. 2
 - (ii) Explain energy of activation and entropy of activation and comment on.
- (c) (i) For a reaction is logarithm of concentration vs. time plot is a straight line with (-ve) slope. What is the order of the reaction.
 - (ii) The slope and intercept of the plot $\frac{1}{R^0} vs \frac{1}{[s]_0}$ are $3.5 \times 10^2_{5}$ and $5 \times 10^4 \text{mol}^{-1}$ L respectively where R° and $[s]_0$ are the initial rate and initial rate ain initial substract concentration of entynce catalysed reaction obeying Michelis Menten kinetics. Estimate k_M and turn-over number when the initial concentration of enzyme. $[E]_0 = 2.5 \times 10^{-9}$ mol L⁻¹.

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(Turn Over)

- (4)
- (d) (i) A carnot engine working between 0°C and 100°C taken up 840J from the HTR. Calculate the w and heat rejected. 2

(ii) Show that
$$\left(\frac{\partial s}{\partial T}\right)_V = \frac{Cv}{T}$$
 for a gas. 3

- (e) (i) For the reaction H₂O(g)→H₂(g) + O₂(g) : ΔH° =242 KJ mol⁻¹ at 290K. Find ΔH° at 310K. Assume that over the temperature range c_p values are effectively constant are given by Cp(H₂O, g) =35.5, Cp(H₂,g)=28.8 and Cp(O₂,g)=29.1 all are in units of JK⁻¹mol⁻¹.
 - (ii) Isothermal free expansion of an ideal gas must be adiabatic – justify or criticize.
- (f) Two moles of an ideal gas at 27°C are enclosed in a leak proof cylinder filled with a movable friction len piston and thermally insulated system or not. The pressure on the piston is release very slowly to effect a quasi-static expansion to double its volume. Calculate
 - (i) the final temperature.

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(Continued)

(5)

(ii) the change in enthalpy of the gas and

(iii) the work done by the gas. Given [Cv=1.5R]

5

Group -C

Answer one of the following :

1×10=10

- 3. (a) (i) Derive the expression for most probable velocity from Maxwell-Boltzman distribution law. 3
 - (ii) Dra a carnot cycle with an ideal gas the working substance in U-S plane.
 - (iii) Derive the expression for adiabatic revessible work of an ideal gas. What will be its magnitude when $\gamma \rightarrow 0$? 4
 - (b) (i) Discuss the important kinetic features of the Lindemann theory of unimolecular reaction. 4
 - (ii) Hydrolysis of ethelacetate by a solution of strong acid in an example of specific acid (H⁺) catalysed reaction.- Explain.

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(Turn Over)

- (iii) The effective rate constant of a first order reaction following Lindemann mechanism has the following values :
 - k=2.5×10⁻³ Sec⁻¹ when [A] = 5×10²(M) and k = 4.2×10^{-3} Sec⁻¹ when [A]= 9.8×10^{-2} (M) Find the rate comtant for the activation step (K₂) and the rates of K₋₂/K₁ 3

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