- (a) Calculate adiabatic lapse rate if the molecular weight of air is 0.029 Kg mol⁻¹ and γ for air is 1.4.
 - (b) Calculate the critical constants of a gas, given a=10⁻⁵, the unit of pressure being 1 atm and b=10⁻³, the unit of volume being Ig molecular volume at NTP.
 - (c) Show diagrammatically how first order phase transition is different from second order phase transition.
 - (d) 50 g of water at 40° C is converted into ice at -10° C at constant atmospheric pressure. If the specific heat of ice at constant pressure is 0.5 cal g-1K-1, calculate the total change in entropy of the system. Latent heat of ice = 80 cal/g.

- (e) Give any two differences between reversible and irreversible processes with one example for each process.
- (f) At what temperature will the average speed of molecules of H₂ gas be four times the average speed of O₂ molecules. (6x3)
- 2. (a) Show that for an adiabatic change in a perfect

gas
$$TP^{\left(\frac{1-\gamma}{\gamma}\right)} = constant.$$
 (7)

(b) Show that for an adiabatic reversible process:

$$\frac{\partial T}{\partial v} = \frac{C_v - C_P}{\alpha v C_v}$$

where C_{ν} , and C_{p} are the specific heat at constant volume and pressure respectively, ν is the specific volume and α is the volume coefficient of expansion. (6)

(c) Prove that adiabatic elasticity of a gas is γ times
 the isothermal elasticity.

- (a) Give Kelvin-Planck and Clausius statements for second law of thermodynamics. Show that both the statements are equivalent to each other. (6)
 - (b) Show that the efficiency of a Carnot engine is dependent only on the temperatures of the source and sink by explaining the various cycles it undergoes.
 (8)
 - (c) A reversible engine converts one-fourth of heat into work. When the temperature of the sink is reduced by 50° C, it converts one half of heat input into work. Calculate the temperatures of the source and sink.

 (4)
 - 4. (a) Starting from the Maxwell's thermodynamical

relation prove that
$$C_P - C_v = \frac{T\alpha^2 v}{\beta_T}$$

where C_v , and C_p are the specific heat at constant volume and pressure respectively, T is the absolute temperature, β_T is the isothermal compressibility, α is the coefficient of volume expansion and ν is the specific volume. (5)

(b) Derive Clausius-Clapeyron's equation

$$dP/dT = L/T(V_2 - V_1)$$

from Maxwell's the thermodynamical relations & also explain the effect of pressure on

- (i) Boiling point of liquids and
- (ii) Melting point of solids (6)
- (c) Discuss in detail the concept of Clausius inequality and hence show that the difference in entropy for an irreversible process is greater than zero. (7)

- 5. (a) What do you mean by magneto-caloric effect?

 Giving a brief description of the experimental procedure, derive the expression for the fall in temperature of the specimen. Under what conditions will the fall in temperature would be more?

 (9)
 - (b) Show that when two phases of a one component system are in equilibrium, then specific Gibb's energies have the same value in both the phases. Hence, derive Ehrenfest's equations for the second order phase transitions. (9)
- (a) Derive an expression for the mean free path and discuss its dependence on temperature and pressure.
 - (b) Derive an expression for the coefficient of thermal conductivity using kinetic theory of gases. (9)

- (c) Calculate the root mean square velocity of neutrons and electrons at 400K, taking the mass of neutron and electron as 1.675×10⁻²⁷ kg and 9.11×10⁻³¹ kg respectively.
- (a) Discuss the results of Andrew's experiments on carbon dioxide. Hence, give a comparison of van der Waals' and Andrew's isotherms. (6)
 - (b) What do you mean by throttling process? Explain by giving a brief discussion of the experiment. Derive an expression for Joule-Thomson coefficient for ideal gases.
 (8)
 - (c) Calculate the drop in temperature when carbon dioxide gas suffers Joule-Thomson expansion at 30°C. The pressures on the two sides of the porous plug are given as 40 atm and 1 atm respectively. The van der Waals' constants of the gas are a =

36.5 x 10^{-2} Nm⁴ mol⁻², b=5.28 x 10^{-5} m³mol⁻¹. (C_p =36.575 JK⁻¹ mol⁻¹ and R = 8.31 JK⁻¹mol⁻¹)

Values of Constants:

Boltzmann's constant, $k = 1.38 \times 10^{-23} \text{ JK}^{-1}$ Universal gas constant, $R = 8.31 \text{ Jmol}^{-1} \text{K}^{-1}$