Statistical Mechanics I

In addition to Atkins and the primer, you may (or may not!) find some of the following books useful:

F. C. Andrews, *Equilibrium Statistical Mechanics*, 2nd edition (John Wiley and Sons, Inc 1975)
Peter A. Rock, *Chemical Thermodynamics, Principles and Applications* (The Macmillan company, -Collier Macmillan Canada, Ltd., Toronto, Ontario 1969)
Oscar K. Rice, *Statistical Mechanics, Thermodynamics and Kinetics* (W. H. Freeman and Company 1967)
M. H. Everdell, *Statistical Mechanics and its Chemical applications* (Academic Press, London, 1975)
Robert J. Finkelstein, *Thermodynamics and Statistical Physics, a Short Introduction* (W. H. Freeman and Company, San Fransisco 1959)
Walter Greiner, Ludwig Neise, Horst Stöcker, *Thermodynamics and Statistical Mechanics* (Springer-Verlag New York 1995)
Donald A. McQuarrie, *Statistical Mechanics* (Harper and Row, 1976)

SECTION A

Make notes on the following.

- 1. The Boltzmann distribution and its derivation.
- 2. What is a molecular partition function? What is its physical significance and why is it important in chemistry?
- 3. Show how the molecular partition function can be written as the product of contributions from independent modes of motion of a molecule and derive analytical expressions for the translational, rotational and vibrational partition functions (stating any approximations you make).
- 4. What is a canonical partition function and why is it used? How is it related to the molecular partition function?
- 5. By using standard equations from thermodynamics, show how the canonical partition function can be used to calculate the following macroscopic thermodynamic properties for a system of interacting particles:
 - i) Internal energy, U
 - ii) Entropy, S
 - iii) Helmholtz energy, A
 - iv) Pressure, P
 - v) Enthalpy, H
 - vi) Gibbs free energy, G
 - vii) Heat capacity, C_v

SECTION B

Question 1

- a) i) State the relationship between the translational energy of a particle inside a cubic container, the mass *m* of the particle, the dimension *L* of the container, and the associated quantum numbers.
 - ii) By using this relationship, show that the translational partition function q_{tr} is given by the expression

$$q_{\rm tr} = \frac{V}{\Lambda^3}$$
 where $\Lambda = \frac{h}{(2\pi m k_{\rm B} T)^{1/2}}$ and $V = L^3$

NB: You may need the following integral:
$$\int_0^\infty \exp(-\alpha x^2) dx = \frac{1}{2} \left(\frac{\pi}{\alpha}\right)^{1/2}$$

b) Explain why the expression for q_{tr} is invalid for small values of *L*, and estimate the dimension at which the invalidity will become significant for oxygen molecules at 298 K.

Question 2

a) Using your expression for the vibrational partition function from Section A, show that the contribution to the molar specific heat at constant volume from the vibrational motion of a diatomic molecule is

$$C_{v^{\text{vib}}} = R \left(\frac{q_{\text{vib}}}{T}\right)^2 \frac{\exp(-q_{\text{vib}}/T)}{\left[1 - \exp(-q_{\text{vib}}/T)\right]^2} \quad \text{where} \quad q_{\text{vib}} = \frac{hc\omega_{\text{e}}}{k_{\text{B}}}$$

What are the limiting values of C_v^{vib} in the limits as $T \rightarrow 0$ and at high temperature? Explain the significance of these two results.

- b) The molar specific heat at constant volume of chlorine at 300 K is 25.44 J K⁻¹ mol⁻¹. By trial and error or other means, use this value to estimate the harmonic vibrational wavenumber for chlorine.
- c) In reality, diatomic molecules are anharmonic oscillators. How will this anharmonicity modify the way in which $C_{\nu}^{\nu ib}$ changes as the temperature increases from low values?

Question 3

The two lowest electronic states of a fluorine atom are separated in energy by $\Delta\epsilon$ ~400 cm⁻¹.

- a) Deduce term symbols for these states, specify their respective electronic degeneracies, and explain which of the two has the lower energy.
- b) With the assumption that only these electronic states are thermally accessible, write down an expression for the molecular electronic partition function q_{el} .
- c) Obtain an expression for the electronic contribution $E_{el}(T)$ to the internal energy. Explain physically why, for temperatures $T >> \Delta \varepsilon/k_{B}$, $\Delta E_{el}(T) = E_{el}(T) E_{el}(0)$ tends to the value 1/3 $N\Delta \varepsilon$, where N is the total number of fluorine atoms.

d) Show that the electronic contribution, $C_{el}(T)$, to the specific heat is given by

$$\frac{C_{\rm el}(T)}{Nk_{\rm B}} = 2\left(\frac{T_{\rm e}}{T}\right)^2 \frac{\exp(T_{\rm e}/T)}{\left[1 + 2\exp(T_{\rm e}/T)\right]^2}$$

where $T_e = \Delta \varepsilon / k_B$ is thus defined. Sketch $C_{el}(T)$ as a function of T/T_e , and comment on the limiting behaviour of $C_{el}(T)$ at low and high temperatures.