

**UNIVERSITY COLLEGE LONDON**

University of London

**EXAMINATION FOR INTERNAL STUDENTS**

For The Following Qualifications:–

*B.Sc.*    *M.Sci.*

**Physics 2B28: Statistical Thermodynamics and Condensed Matter Physics**

**COURSE CODE            :    PHYS2B28**

**UNIT VALUE             :    0.50**

**DATE                     :    11-MAY-05**

**TIME                     :    14.30**

**TIME ALLOWED         :    2 Hours 30 Minutes**

**Answer ALL questions from section A and THREE questions from section B**

The numbers in square brackets in the right hand margin indicate the provisional allocation of marks per sub-section of a question.

$$\text{Boltzmann's constant } k = 1.38 \times 10^{-23} \text{ J K}^{-1}$$

$$\text{Planck's constant } h = 6.627 \times 10^{-34} \text{ J s}$$

$$\text{Electronic charge } e = 1.6 \times 10^{-19} \text{ J}$$

$$\text{Avogadro's number } N_A = 6 \times 10^{23}$$

$$\text{Mass of electron } m_e = 9.1 \times 10^{-31} \text{ kg}$$

$$\beta = (kT)^{-1}$$

**SECTION A**

1. (a) What is meant by the terms *macrostate*, *microstate* and *statistical weight* in statistical thermodynamics? [2]  
  
(b) Write down Boltzmann's expression for the entropy of an isolated system, explaining the meaning of each symbol in the expression. [1]  
  
(c) An isolated system of 6 magnetic dipoles is in a macrostate with 4 dipoles 'up' and 2 dipoles 'down'. Calculate the statistical weight of this macrostate. [2]
  
2. (a) Write down the Kelvin *and* Clausius statements of the second law of thermodynamics [2]  
  
(b) State the third law of thermodynamics, and discuss the possibility of reducing the temperature of a system to absolute zero. [4]
  
3. (a) State the *Boltzmann distribution* of a system in equilibrium at a temperature  $T$ , explaining the meaning of each term. [3]  
  
(b) Show that the average energy of a system at equilibrium at a temperature  $T$  can be expressed as  $-\left(\partial \ln Z / \partial \beta\right)$  where the symbols have their usual meaning. [4]

4. (a) What are the possible spin values of identical particles which obey Fermi-Dirac statistics (i.e. fermions)? Give 2 examples of such particles. [2]

(b) The *Fermi-Dirac occupation function* is given by

$$n(E) = \{ \exp [(E-E_F)/kT] + 1 \}^{-1}$$

How are the *Fermi energy*  $E_F$  and *Fermi temperature*  $T_F$  defined? [2]

Draw  $n(E)$  as a function of  $E$ , at  $T = 0$ , and at a temperature  $T \ll T_F$ , marking  $E_F$  clearly on your diagram. [3]

5. Two *identical* particles can occupy three single-particle states with energies  $\epsilon_1 = 0$ ,  $\epsilon_2 = \epsilon$  and  $\epsilon_3 = 2\epsilon$ .

(a) If the two particles are *fermions*, draw all of the possible arrangements in which they can occupy the single-particle states. State the total energy of the two particles in each case, and hence write down the partition function. [4]

(b) Draw the additional ways in which two *bosons* can occupy the single particle states, giving their total energies. What is the partition function in this case? [3]

6. Explain briefly what is meant by the *free-electron model* of a metal. [2]

Show that the density of states of a free electron gas, as a function of wave vector  $k$ , is given by  $f(k) dk = V k^2 dk / (2\pi^2)$ . [5]

How is  $f(k)$  modified when the effect of electron spin is included? [1]

## SECTION B

7. (a) A particular atom can exist in three energy states:  $E_1 = 1.1 \times 10^{-22}$  J,  $E_2 = 1.9 \times 10^{-22}$  J and  $E_3 = 3.2 \times 10^{-22}$  J, with degeneracies  $g_1 = 1$ ,  $g_2 = 3$  and  $g_3 = 5$ , respectively. If the atom is in equilibrium at a temperature of 10 K, calculate:

(i) the value of the partition function  $Z$  [2]

(ii) the individual probabilities that the atom has energy  $E_i = E_1, E_2$  and  $E_3$  [3]

(iii) the mean energy per atom [2]

(b) Now use the expression  $S = -k \sum p_i \ln p_i$  to calculate the entropy per atom [5]

(c) The energy levels of a simple spin  $1/2$  magnetic system in a magnetic field  $B$  are  $E_1 = -\Delta$  and  $E_2 = +\Delta$ , where  $\Delta = \mu_B B$ . Using the expression in part (b) show that the entropy of the system is

$$S = 2 \Delta \{T [1 + \exp(2 \Delta/kT)]\}^{-1} + k \ln [1 + \exp(-2 \Delta/kT)] \quad [6]$$

(d) By considering this formula, deduce the entropy of the system in both the low temperature and high temperature limits. [2]

8. (a) State Clausius' entropy formulation of the Second Law of Thermodynamics? What condition does the entropy of an isolated system satisfy when it is in thermodynamic equilibrium? [2]

(b) An isolated system is partitioned into two sub-systems, 1 and 2. By considering the change in entropy as heat flows from sub-system 1 to sub-system 2, derive the condition that must be satisfied for sub-systems 1 and 2 to be in thermal equilibrium. Show that this leads to the definition of temperature:

$$1/T = (\partial S / \partial E)_{N,V} \quad [4]$$

(c) If the two sub-systems discussed in part(b) are *not* in thermal equilibrium, show that heat flows from the sub-system at higher temperature to the sub-system at lower temperature. [4]

(d) A linear chain of  $N+1$  spins forms a one-dimensional paramagnetic system. Each spin interacts with its neighbours in such a way that the total domain wall energy of the system is  $E = n\epsilon$ , where  $n$  is the number of domain walls separating regions of down

spins from up spins, and  $\epsilon$  is the energy of one domain wall. These domain walls are indicated by vertical lines in the diagram below:



How many ways can the  $n$  domain walls be arranged in the chain of  $N+1$  spins? [2]  
 Calculate the entropy  $S(E)$ , and hence show that the energy  $E$  at a temperature  $T$  is [8]

$$E = N\epsilon \{ [\exp(\epsilon/kT)] + 1 \}^{-1}$$

[note Stirling's formula for large  $N$ :  $\ln(N!) \sim N \ln(N) - N$ ]

9. (a) State the conditions when an ideal gas of identical quantum particles may be described as a *classical gas*, or as being in the *classical regime*. [2]

(b) Considering only the translational motion, show that in the classical regime the partition function of a single gas atom of mass  $m$  in a volume  $V$  is given by [6]

$$Z(1, V, T) = V \left( \frac{2\pi mkT}{h^2} \right)^{3/2}$$

You may assume that the density of momentum states is given by

$$f(p) dp = (V/h^3) 4\pi p^2 dp \text{ and that } \int_0^\infty x^2 \exp(-ax^2) dx = (1/4a) (\pi/a)^{1/2}$$

(c) Show that the validity of the classical regime can be expressed as [4]

$$\frac{N}{V} \left( \frac{h^2}{2\pi mkT} \right)^{3/2} \ll 1$$

(d) The mass of a  $^4\text{He}$  atom is  $6.67 \times 10^{-27}$  kg. Consider whether the classical regime is valid for (i) liquid helium at 4.2 K, for which  $N/V \sim 2 \times 10^{28} \text{ m}^{-3}$  and (ii) gaseous helium at 273 K, for which  $N/V \sim 1 \times 10^{26} \text{ m}^{-3}$  [4]

(e) Define the validity of the classical regime in terms of the *de Broglie wavelength* of the particles. Use this definition to confirm your conclusion in (d) for the validity of the classical regime for gaseous helium at 273K. [4]

- 10 (a) Consider an ideal Fermi-Dirac gas of  $N$  electrons of mass  $m$  in a volume  $V$ . Using the expression for the density of states as a function of wave vector  $k$ ,  $f(k) dk = V k^2 dk / (2\pi^2)$ , express the density of states as a function of energy, and hence show that the Fermi energy  $\epsilon_F$  is given by : [8]

$$\epsilon_F = \frac{h^2}{2m_e} \left( \frac{3N}{8\pi V} \right)^{2/3}$$

- (b) Show that the internal energy of a Fermi-Dirac gas at  $T=0$  is given  $(3/5) N\epsilon_F$ . [3]

- (c) Gold (Au) is a monovalent metal, *i.e.* it has one conduction electron per atom. The atomic mass of Au is 196.97 and its density is  $19300 \text{ kg m}^{-3}$ .

Calculate the Fermi energy (in eV), the Fermi temperature, *and* the Fermi velocity for the conduction electrons in gold: [6]

Comment on the ratio of the Fermi velocity to the velocity of light [1]

Calculate the internal energy due to the conduction electrons in 1 gram of Au at  $T=0$ . [2]

- 11.(a) Define the Gibbs function  $G$  of a system. Write down the differential form ( $dG$ ). [2]

- (b) Deduce the condition for two phases of a homogeneous one-component system to be in equilibrium at constant pressure and temperature. [3]

- (c) Derive the Clausius-Clapeyron equation

$$\frac{dp}{dT} = \frac{L}{T\Delta V}$$

relating to equilibrium between phases subject to a first-order phase transition, where  $L$  is the latent heat and  $\Delta V$  is the volume change in passing from one phase to the other. [8]

- (d) Sketch the  $p, T$  phase diagram applicable to most simple substances. Label the solid, liquid and vapour phases, and mark clearly the triple and critical points. How does the phase diagram for  $\text{H}_2\text{O}$  differ from this diagram? [4]

- (e) When lead is melted at atmospheric pressure, the melting point is  $327.5^\circ\text{C}$ , the density decreases from  $11010$  to  $10650 \text{ kg m}^{-3}$ , and the latent heat is  $24.5 \text{ kJ kg}^{-1}$ . What is the melting point of lead at a pressure of 100 atmospheres? Assume atmospheric pressure is  $10^5 \text{ Pa}$ . [3]