Department of Physics University of Maryland College Park, MD 20742-4111

Physics 603

HOMEWORK ASSIGNMENT #8

Spring 2014

Due date for problems on Thursday, April 24 [deadline on April 29].

- 1. a) PB 8.1:
 - b) PB 8.2
- 2. (5 pt) Using results derived in class, show for Fermi gases that, to leading order in T, S = $(\pi^2/3)k_B^2Tg(\epsilon_F)$
- 3. a) Derive the equivalent of eq. 8.1.9 for arbitrary D and s.
- b) Similarly, find z^{-1} ($\partial z/\partial T$)_P and show that it reduces to the first equation in 8.3 for D=3, s=2.
- c) PB 8.10 b) [Cf. 7.14 for Bose gases.] We did 8.10 a) in class.
- 4, 5. Qual problems (see next pages)

Consider a free-electron gas, with N electrons and dispersion relation $\epsilon = (\hbar |\mathbf{k}|)^2/2m$ in d dimensions, at temperature T=0.

(a) Show that the density of electronic states $G(\epsilon)$ satisfies

$$G(\epsilon) \propto \epsilon^{\alpha}$$
 (1)

and find the value of α as a function of d=1,2,3. (Hint: the sum of these 3 values of α is 0.)

Recall:

$$G(\epsilon) \propto \int \frac{dS_{\epsilon}}{|\nabla_k \epsilon|}$$
 (2)

where the integration in k-space is over a surface (sphere, circle, etc.) of constant energy ϵ . [4 points]

(b) i. What is the [physical] meaning of $G(\epsilon) d\epsilon$? [1 point] ii. The proportionality (1) can be written as an equation in terms of a dimensionless constant A. Using dimensional arguments, show that

$$G(\epsilon) = \frac{AN\epsilon^{\alpha}}{\epsilon_F^{\alpha'}},\tag{3}$$

where ϵ_F is the Fermi energy, and specify the relationship between α' and α . [3 points] iii. Find the [d-dependent] value of A. [2 points]

For graphene (a single planar sheet of graphite), the electronic dispersion relation can be written as $\epsilon = \hbar v_s |\mathbf{k}|$ for small ϵ . (The points in k-space at which $\epsilon = 0$ are hence called Dirac points, and k is measured from them; you do not need this parenthetical information to do this problem.)

- (c) i. Show that the density of states $G(\epsilon)$ is still proportional to $\epsilon^{\alpha}/\epsilon_F^{\alpha'}$, and find the new value of α . [3 points] ii. Does the relationship between α' and α change from part (b)? If yes, how? If not, why not? [1 point]
- (d) i. The low-temperature specific heat of a metal is known to be a power-law of T. Give a quick argument to show what this power is. (It may be helpful to draw a sketch of the change in the Fermi-Dirac distribution when T increases slightly from 0; use of the Sommerfeld expansion is not expected!) [2 points] ii. For graphene with small doping, so that ε_F and G(ε_F) are non-zero, how does the temperature dependence of the low temperature specific heat compare with that found in part (d)-i? [2 points]
- (e) In many cases a small gap (of size E_g) opens around $\epsilon = 0$. If ϵ_F lies inside this gap, write down the leading form of the temperature dependence of the low-T ($T \ll E_g/k_B$) specific heat. In the limit that $T \to 0$, what is the ratio of this specific heat to the specific heat in part (d)-ii? [2 points]

January 2011

Problem I.3

First consider a classical ideal gas at temperature T consisting of N molecules and initially confined in a volume V_i . Then the gas is allowed to expand to a final volume V_f in two different ways:

- (a) Free expansion. The gas is thermally insulated from its environment and experiences free irreversible expansion into a vacuum. Calculate the entropy change of the gas $\Delta S_{\rm irr}^{\rm gas} = S_f S_i$ by comparing the number of accessible states before and after the expansion. (8 points)
- (b) Isothermal expansion. The gas is in thermal contact with a reservoir of temperature T and experiences a slow reversible quasistatic expansion, e.g. produced by a slow motion of a piston that limits the gas volume. Calculate the work W done on the gas in this process, the change $\Delta U = U_f U_i$ of the internal energy of the gas, and the heat Q transferred to the gas from the environment. Calculate the entropy change of the gas $\Delta S_{\text{rev}}^{\text{gas}} = S_f S_i$ in this reversible process by using the formula $\Delta S = Q/T$. Compare your answers for $\Delta S_{\text{irr}}^{\text{gas}}$ and $\Delta S_{\text{rev}}^{\text{gas}}$. Are the two results the same or different? Explain why. (8 points).
- (c) What are the entropy changes in the environment for these two cases: $\Delta S_{\rm irr}^{\rm env}$ and $\Delta S_{\rm rev}^{\rm env}$? What are the total entropy changes in the gas and the environment for these two cases: $\Delta S_{\rm irr}^{\rm tot} = \Delta S_{\rm irr}^{\rm gas} + \Delta S_{\rm irr}^{\rm env}$ and $\Delta S_{\rm rev}^{\rm tot} = \Delta S_{\rm rev}^{\rm gas} + \Delta S_{\rm rev}^{\rm env}$? Are $\Delta S_{\rm irr}^{\rm tot}$ and $\Delta S_{\rm rev}^{\rm tot}$ the same or different? Explain why. (5 points)
- (d) Now consider a non-interacting degenerate Fermi gas made of N spin-1/2 fermions each of mass m and initially confined in a volume V_i at zero temperature T = 0. Calculate the Fermi momentum p_F , the Fermi energy E_F , and the energy per particle U/N. Express U/N in terms of E_F . (7 pts)
- (e) This Fermi gas is thermally insulated from its environment and experiences free irreversible expansion into a vacuum to a final volume V_f . Assume that V_f is sufficiently large so that the Fermi gas becomes non-degenerate, i.e. classical. Calculate the final temperature T_f of the gas after the expansion. Express T_f in terms of E_F obtained above. (6 pts)
- (f) Estimate by what factor V_f/V_i the initially degenerate Fermi gas needs to expand in order to become classical in the final state. (6 pts)

 Hint: A gas behaves classically when the inter-particle separation $d = \sqrt[3]{N/V}$ is much greater than the thermal de Broglie length $\lambda = h/\sqrt{2\pi mkT}$, where h and k are the Planck and the Boltzmann constants.