# Drexel Physics 2005 Modern Qual Solutions

2014 entering class

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## 1 Problem 1

We need expected value of the particle energy in the well to be less than  $E=h\nu$ . The energy levels of the finite square well are XXX. We then compute the partition function and take the derivative to find  $\langle E \rangle$  as a function of temperature.

### 2 Problem 2

With  $S=kT\ln\Omega$  and  $\frac{\partial S}{\partial E}=\frac{1}{T},$  we need to add:

$$E = kT^2 \ln d\Omega = (Joules) = (calories)$$
 (2.1)

to increase the number of accessible states by 1000.

# 3 Problem 3

Electron tunneling in transistors.

Angular momentum quantization as measured in the Stern-Gerlach experiment. When a beam of spin  $\frac{1}{2}$  particles passes through a magnetic field they are split into two distinct beams instead of smeared over an angular region.

#### Problem 4 4

To compute the normalization constant we need to solve:

$$C^2 \int_V R^2 = 1 \tag{4.1}$$

$$\int (e^{-r/a})^2 r^2 dr = \frac{1}{C^2}$$
 (4.2)

$$\int r^2 e^{-2r/a} dr = \frac{1}{C^2}$$
 (4.3)

$$\frac{2}{(2/a)^3} = \frac{1}{C^2} \tag{4.4}$$

$$\int r^{2} e^{-2r/a} dr = \frac{1}{C^{2}}$$

$$\frac{2}{(2/a)^{3}} = \frac{1}{C^{2}}$$

$$C = 2\left(\frac{Z}{a_{0}}\right)^{3/2}$$
(4.3)
(4.4)

#### Problem 5 **5**

Free expansion of a gas is one irreversible process. The theory of microscopic reversability only holds for systems that remain at equilibrium.

#### 6 Problem 6

The cloud of gas also has gravitational potential energy and electronic potential energy, so the total system energy may decrease even if the kinetic energy increases.

# 7 Problem 7

### 8 Problem A1

We see that  $S_x = \frac{S_+ + S_-}{2}$ . We find the eigenvalues in the usual manner:

$$S_x = \frac{\hbar}{2} \begin{bmatrix} 0 & \sqrt{2} & 0\\ \sqrt{2} & 0 & \sqrt{2}\\ 0 & \sqrt{2} & 0 \end{bmatrix}$$
 (8.1)

$$\begin{vmatrix} -\lambda & \sqrt{2}/2\hbar & 0\\ \sqrt{2}/2\hbar & -\lambda & \sqrt{2}/2\hbar\\ 0 & \sqrt{2}/2\hbar & -\lambda \end{vmatrix} = 0 \tag{8.2}$$

$$-\lambda(\lambda^2 - \hbar^2/2) - \hbar\sqrt{2}/2(-\lambda\hbar\sqrt{2}/2) = 0$$
 (8.3)

$$-\lambda^3 + \lambda \hbar^2 = 0 \tag{8.4}$$

$$\lambda = \{-\hbar, \hbar\} \tag{8.5}$$

We now solve  $S_x \mathbf{e} = \lambda \mathbf{e}$  for the eigenvectors.

$$e_1 = e_3$$
 (8.6)

$$e_2 = \frac{1}{\lambda} \sqrt{2\hbar} e_1 \tag{8.7}$$

$$\lambda = -\hbar : \mathbf{e}_{-\hbar} = \{1, -\sqrt{2}, 1\} \frac{1}{2}$$
 (8.8)

$$\lambda = \hbar : \mathbf{e}_{\hbar} = \{1, \sqrt{2}, 1\} \frac{1}{2}$$
 (8.9)

We now calculate the probability of measuring  $S_x = \hbar$ . (The question says S + x = 1, but 1 is not an eigenvalue of  $S_x$ .

$$P(S_x = \hbar) = \langle e_{\hbar} | u \rangle^2 = \left( \frac{1}{2\sqrt{2c}} (1 * 1 + 4 * \sqrt{2} - 3 * 1) \right)^2$$
 (8.10)

$$P(S_x = \hbar) = \frac{1}{2c}(9 - 4\sqrt{2}) \tag{8.11}$$

After the measurement the system has a definite value of  $S_x = \hbar$  and is in state  $e_{\hbar}$ . The eigenvector corresponding to  $S_z = \hbar$  is (1,0,0), so we calculate the probability the same way.

$$P(S_z = \hbar) = \langle \mathbf{e_1} | \mathbf{e_{\hbar}} \rangle^2 = \left(1 * \frac{1}{2} + 0 + 0\right)^2$$
 (8.12)

$$P(S_z = \hbar) = \frac{1}{4}$$
 (8.13)

# 9 Problem A2

a) We open up the brackets and find  $\Psi(x,0) = 2x^3 - 10x_1^22x$ .

$$\int_0^3 (C\Psi(x,0))^2 dx = 1 \tag{9.1}$$

$$\frac{1}{C^2} = \left(\int_0^3 2x^3 - 10x^2 + 12x dx\right)^2 \tag{9.2}$$

$$\frac{1}{C^2} = (-\frac{1}{2})^2 = \frac{1}{4} \tag{9.3}$$

$$C = 2 (9.4)$$

$$\Psi(x,0) = 2(2x^3 - 10x^2 + 12x) \tag{9.5}$$

b) The wavefunction has a single 0 in the range at x=2, so it most closely resembles the standard wavefunction  $\sin \frac{2\pi}{3}x$ . c) The wavefunctions can be derived from the Schrödinger equation:

$$-\frac{\hbar^2}{2m}\frac{d^2\psi}{dx^2} = e\psi \tag{9.6}$$

$$\frac{d^2\psi}{dx^2} = -\frac{2mE}{\hbar^2}\psi\tag{9.7}$$

With  $k \equiv \frac{\sqrt{2mE}}{\hbar}$ , the solutions are of the form  $A\cos kx + B\sin kx$ . The infinite square well requires  $\psi(0) = 0$ , so A = 0. This square well also requires  $\psi(3) = 0$ , so k takes on discrete values determined by:

$$\sin\frac{n\pi x}{3} = 0\tag{9.8}$$

$$k = \frac{n\pi}{3}, \ n = 1, 2, 3...$$
 (9.9)

$$\frac{\sqrt{2mE}}{\hbar} = \frac{n\pi}{3} \tag{9.10}$$

$$E = \frac{n^2 \pi^2 \hbar^2}{18m}, \ n = 1, 2, 3... \tag{9.11}$$

So we can estimate the expectation value of the energy for n=2 as  $\frac{2\pi^2\hbar^2}{9m}$ .

### 10 Problem A3

The energies of states are  $E_0 = \frac{1}{2}\hbar\omega$  and  $E_2 = \frac{5}{2}\hbar\omega$  so the time-evolving wavefunction is:

$$|\Psi(t)\rangle = C_0 e^{i\frac{1}{2}\omega t} |0\rangle + C_2 e^{i\frac{5}{2}\omega t} |2\rangle$$
 (10.1)

b) We can write the momentum and position operators in terms of the creation and annihilation operators, which come from "factoring" the Hamiltonian.

$$H = \frac{p^2}{2m} + \frac{1}{2}kx^2 \tag{10.2}$$

$$a = \left(\frac{m\omega}{2\hbar}\right)^{1/2} \hat{X} + i \left(\frac{1}{2m\omega\hbar}\right)^{1/2} \hat{P}$$
 (10.3)

$$a^{\dagger} = -\left(\frac{m\omega}{2\hbar}\right)^{1/2}\hat{X} + i\left(\frac{1}{2m\omega\hbar}\right)^{1/2}\hat{P} \tag{10.4}$$

$$\hat{P} = -\frac{i}{2}\sqrt{2m\omega\hbar}(a+a^{\dagger}) \tag{10.5}$$

$$\hat{X} = \sqrt{\frac{\hbar}{2m\omega}} (a + a^{\dagger}) \tag{10.6}$$

We now apply  $\hat{P}$  to the wavefunction.

$$\langle \hat{P} \rangle = \langle \Psi(t) | \hat{P} | \Psi(t) \rangle$$
 (10.7)

$$\langle \hat{P} \rangle = \left( -\frac{1}{2} m \omega \hbar \right) \Psi(t) \left( (a + a^{\dagger}) \left( C_0 e^{i \frac{1}{2} \omega t} \left| 0 \right\rangle + C_2 e^{i \frac{5}{2} \omega t} \left| 2 \right\rangle \right) \right)$$
 (10.8)

$$\langle \hat{P} \rangle = \left( -\frac{1}{2} m \omega \hbar \right) \Psi^*(t) \left( 0 + C_2 e^{i \frac{5}{2} \omega t} (-\hbar \omega) \left| 1 \right\rangle + C_0 e^{i \frac{1}{2} \omega t} (\hbar \omega) \left| 1 \right\rangle + C_2 e^{i \frac{5}{2} \omega t} (\hbar \omega) \left| 3 \right\rangle \right)$$

$$(10.9)$$

$$\langle \hat{P} \rangle = 0 \tag{10.10}$$

In the last step we used the orthogonality of the harmonic oscillator eigenfunctions, since  $\Psi(t)$  only includes states with energy levels 0 and 2. c) The expectation value of  $\hat{X}$  is:

$$\langle \hat{X} \rangle = \langle \Psi(t) | \hat{X} | \Psi(t) \rangle \tag{10.11}$$

(10.12)

Since  $\hat{X} == \sqrt{\frac{\hbar}{2m\omega}}(a+a^{\dagger})$ , we again see the combination of creation and annihilation operators. Once again the orthogonality of the harmonic oscillator eigenfunctions and the fact that the initial wavefunction contains only states 0 and 2 cause the expectation value to be 0. The expectation of both  $\hat{P}$  and  $\hat{X}$  will be 0 whenever the initial state contains only odd levels or only even levels, corresponding to combinations of only odd/even wavefunctions.

### 11 Problem B1

a)

i) Each distinguishable particle may be in one of 4 states, so there are 16

total states. 4 states have energy 0, 8 states have energy  $\epsilon$ , and 4 states have energy  $2\epsilon$ .

$$Z = \sum_{1}^{8} e^{-\beta \epsilon} + \sum_{1}^{8} e^{-2\beta \epsilon} + \sum_{1}^{8}$$
 (11.1)

$$Z = 4(1 + 2e^{-\beta\epsilon} + 1e^{-2\beta\epsilon}) \tag{11.2}$$

ii) For Bosons there are three spin combinations and three energy combinations for a total of 9 states. Three of the states have energy 0, three have energy  $\epsilon$  and three have energy  $2\epsilon$ .

$$Z = 3(1 + e^{-\beta\epsilon} + e^{-2\beta\epsilon}) \tag{11.3}$$

iii) For Fermions the Pauli exclusion principle will eliminate the states with both particles having the same spin. There are now 3 states, with one for each possible value of the energy.

$$Z = 1 + e^{-\beta\epsilon} + e^{-2\beta\epsilon} \tag{11.4}$$

b) As  $T\to\infty$ ,  $\beta=\frac{1}{kT}\to 0$  and  $e^{-\beta\epsilon}\to 1$ . For the classical particles, each state will have probability  $\frac{1}{16}$ , so a quarter of the systems will have E=0. For the bosons, each state will have probability  $\frac{1}{9}$ , so one third of the systems will have E=0.

For the fermions each state will have probability  $\frac{1}{3}$ , so one third of the systems will have E = 0.

c) We can't use  $\langle E \rangle = \frac{\partial \ln Z}{\partial (-\beta)}$  since we have a small number of states (?? I'm not actually sure if this is correct, I may have just screwed up the calculation). Instead, we calculate  $\langle E \rangle$  directly.

For the classical particles:

$$\langle E \rangle = \frac{\sum E_s e^{-\beta E_s}}{Z} \tag{11.5}$$

$$= \frac{4 \cdot 0 + 8\epsilon e^{-\beta\epsilon} + 4 \cdot 2\epsilon e^{-2\beta\epsilon}}{4(1 + 2e^{-\beta\epsilon} + e^{-2\beta\epsilon})}$$
(11.6)

$$= 2\epsilon \frac{e^{-\beta\epsilon} + e^{-2\beta\epsilon}}{1 + 2e^{-\beta\epsilon} + e^{-2\beta\epsilon}}$$
(11.7)

For the bosons:

$$\langle E \rangle = \frac{\sum E_s e^{-\beta E_s}}{Z} \tag{11.8}$$

$$= \frac{3 \cdot 0 + 3\epsilon e^{-\beta\epsilon} + 3 \cdot 2\epsilon e^{-2\beta\epsilon}}{3(1 + e^{-\beta\epsilon} + e^{-2\beta\epsilon})}$$
(11.9)

$$= \epsilon \frac{e^{-\beta\epsilon} + 2e^{-2\beta\epsilon}}{1 + e^{-\beta\epsilon} + e^{-2\beta\epsilon}}$$
 (11.10)

For the fermions:

$$\langle E \rangle = \frac{\sum E_s e^{-\beta E_s}}{Z} \tag{11.11}$$

$$= \frac{0 + \epsilon e^{-\beta \epsilon} + 2\epsilon e^{-2\beta \epsilon}}{1 + e^{-\beta \epsilon} + e^{-2\beta \epsilon}}$$

$$= \epsilon \frac{e^{-\beta \epsilon} + 2e^{-2\beta \epsilon}}{1 + e^{-\beta \epsilon} + e^{-2\beta \epsilon}}$$

$$(11.12)$$

$$= \epsilon \frac{e^{-\beta \epsilon} + 2e^{-2\beta \epsilon}}{1 + e^{-\beta \epsilon} + e^{-2\beta \epsilon}} \tag{11.13}$$

d) In the high temperature limit,  $\langle E \rangle = \epsilon$  for all types of particles.

#### 12Problem B2

a) The heat capacity  $C_v = (\frac{\partial U}{\partial T})_V$ . If the heat capacity is constant then  $U \sim T$ . Then A = U - TS = CT - TS = T(C - S) with C some constant.

b) For a monatomic ideal gas the Hamiltonian is  $H = \frac{p^2}{2m}$ . The partition function of one atom is:

$$Z_1 = \frac{1}{h^3} \int d^3x \ d^3p \ e^{-\beta \frac{p^2}{2m}}$$
 (12.1)

$$Z_1 = \frac{V}{h^3} \int d^3p \ e^{-\beta \frac{p^2}{2m}} \tag{12.2}$$

Each momentum component integral  $\int_0^\infty e^{-\beta \frac{p^2}{2m}} dp = \sqrt{2m\pi kT}$ . With three independent components, the total partition function of one atom is:

$$Z_1 = \frac{V}{h^3} (2m\pi kT)^{3/2} \tag{12.3}$$

(12.4)

The system partition function (indistinguishable particles) is:

$$Z_N = \frac{1}{N!} (\frac{V}{h^3})^N (2m\pi kT)^{3N/2}$$
 (12.5)

The Helmholtz free energy A is  $A = -kT \ln Z_N$ .

$$A = -NkT(\ln\frac{V}{N!} + \frac{3}{2}\ln\frac{2m\pi kT}{h^2})$$
(12.6)

c) The energy levels of the harmonic oscillator are  $(n+\frac{1}{2})\hbar\omega$ . The partition function for one atom is:

$$\sum_{s=0}^{\infty} e^{-\beta(s+\frac{1}{2})\hbar\omega} = e^{-\beta\frac{1}{2}\hbar\omega} (1 + e^{-\beta\hbar\omega} + e^{-2\beta\hbar\omega} + \dots)$$
 (12.7)

$$=e^{-\beta\frac{1}{2}\hbar\omega}\left(\frac{1}{1-e^{\beta\hbar\omega}}\right) \tag{12.8}$$

The partition function for N atoms is  $Z_1^N = \left(e^{-\beta \frac{1}{2}\hbar\omega}(\frac{1}{1-e^{\beta\hbar\omega}})\right)^N$ . The Helmholtz free energy is:

$$A = -kT \ln Z_N \tag{12.9}$$

$$A = -NkT(-\beta \frac{1}{2}\hbar\omega + \beta\hbar\omega)$$

$$A = -\frac{N}{2}\hbar\omega$$
(12.10)
$$(12.11)$$

$$A = -\frac{N}{2}\hbar\omega\tag{12.11}$$