## Name

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ME 501

Exam \#2
November 15, 2007
Prof. Lucht
ME 255

## 1. POINT DISTRIBUTION

## Problem \#1

Problem \#2
Problem \#3
Problem \#4
Problem \#5

30 points
30 points
30 points
30 points
30 points

## 2. EXAM INSTRUCTIONS

- You must do four (4) out of the five (5) problems and indicate clearly which problem you decided not to have graded. Please write "DO NOT GRADE" in the space next to the problem that you decide not to do.
- This exam is closed book and closed notes.
- Several equation sheets are attached.
- When working the problems, list all assumptions, and begin with the basic equations.
- If you do not have time to complete evaluation of integrals or of terms numerically, remember that the significant credit on each problem will be given for setting up the problem correctly and/or obtaining the correct analytical solution.

1. ( $\mathbf{3 0}$ points) The time-independent Schrödinger wave equation for the harmonic oscillator is

$$
-\left(\hbar^{2} / 2 \mu\right) \frac{d^{2} \psi}{d x^{2}}+V(x) \psi(x)=\varepsilon \psi(x)
$$

where $V(x)=k x^{2} / 2$ and $x=r-r_{e}$. The internuclear spacing is $r$, and the equilibrium internuclear spacing is $r_{e}$. The normalized eigenfunction and energy eigenvalue for the ground state $(\mathrm{v}=0)$ are given by

$$
\begin{aligned}
\psi_{0}(x) & =\beta \exp \left(-a x^{2}\right), \quad a=\frac{\pi \mu v}{\hbar}, \quad \beta=\left(\frac{2 a}{\pi}\right)^{1 / 4} \\
\varepsilon_{0} & =\left(v+\frac{1}{2}\right) h v=\frac{1}{2} h v, \quad k=(2 \pi v)^{2} \mu
\end{aligned}
$$

Calculate the expectation values in Joules for the potential energy $\langle V(x)\rangle$ and for the kinetic energy $\left\langle p_{x}^{2} / 2 \mu\right\rangle$, for $N_{2}$ in its ground vibrational state. For nitrogen, the mass of each nucleus is 14 amu and the vibrational frequency $(v / c)=2330 \mathrm{~cm}^{-1}$. Hint: you may find it useful to express the spring constant $k$ in terms of the parameter $a$. Also note:

$$
\begin{aligned}
& \int_{0}^{\infty} \exp \left(-b x^{2}\right) d x=\frac{1}{2} \sqrt{\frac{\pi}{b}} \quad \text { for } b>0 \\
& \int_{0}^{\infty} x^{2 n} \exp \left(-b x^{2}\right) d x=\frac{1 \times 3 \times 5 \times \ldots \ldots \times(2 n-1)}{2^{n+1} b^{n}} \sqrt{\frac{\pi}{b}} \quad \text { for }(n=1,2, \ldots \ldots, b>0) \\
& \int_{0}^{\infty} x^{2 n+1} \exp \left(-b x^{2}\right) d x=\frac{n!}{2 b^{n+1}} \quad \text { for }(n=0,1,2, \ldots \ldots, b>0)
\end{aligned}
$$

Answers: $\langle V(x)\rangle=\left\langle\frac{p_{x}^{2}}{2 \mu}\right\rangle=1.16 \times 10^{-20} J$

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2. ( 30 points) The system shown below has energy levels of $\left(\varepsilon_{j} / k_{B}\right)=0 \mathrm{~K}, 5 \mathrm{~K}, 10 \mathrm{~K}, 150 \mathrm{~K}$, and 200 K . The degeneracy of each of three lower levels is $10^{8}$ and the two upper levels is $10^{10}$. The system has only these energy levels; there are no translational, rotational, or vibrational modes. The number of particles in the assembly is $10^{5}$.
(a) Calculate the energy of the assembly at temperatures of 5 , 25 , and 50 K , assuming corrected Maxwell-Boltzmann statistics. Why is CMB statistics a good assumption?
(b) Estimate the heat capacity of the system at 15 K and at 37.5 K by considering the results of part (a). Comment on the results.
(c) Calculate the assembly entropy S (in $\mathrm{J} / \mathrm{K}$ ) at temperatures of 5,25 , and 50 K . Assume corrected-Maxwell-Boltzmann statistics. Comment on the results.

|  | $j$ | $\varepsilon_{j} / k_{B}(\mathrm{~K})$ | $g_{j}$ |
| :---: | :---: | :---: | :---: |
|  | 4 | 200 | $10^{10}$ |
|  | 3 | 150 | $10^{10}$ |
|  |  |  |  |
|  | 2 | 10 | $10^{8}$ |
|  | 1 | 5 | $10^{8}$ |
|  | 0 | 0 | $10^{8}$ |

Answers (a) $E(50 \mathrm{~K})=1.64 \times 10^{-16} \mathrm{~J} \quad$ (b) $S(50 \mathrm{~K})=1.73 \times 10^{-17} \mathrm{~J} / \mathrm{K}$

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3. (30 points)
(a) Given the translational energy distribution for a Maxwell-Boltzmann gas

$$
f\left(\varepsilon_{t r}\right) d \varepsilon_{t r}=\frac{2\left(\varepsilon_{t r}\right)^{1 / 2}}{\pi^{1 / 2}\left(k_{B} T\right)^{3 / 2}} \exp \left(-\varepsilon_{t r} / k_{B} T\right) d \varepsilon_{t r}
$$

show that the speed distribution is given by

$$
f(V) d V=\sqrt{\frac{2}{\pi}}\left(\frac{m}{k_{B} T}\right)^{3 / 2} V^{2} \exp \left(-m V^{2} / 2 k_{B} T\right) d V
$$

(b) Calculate the value of the mean reciprocal speed $\left\langle\frac{1}{V}\right\rangle$ for an assembly of nitrogen $\left(\mathrm{N}_{2}\right)$ molecules at a temperature of 1000 K and a pressure of 2 bars. For $\mathrm{N}_{2}$, the mass of each nucleus is 14 amu . You will need to use the definite integral formulae from Problem \#1.

Answer: (b) $\left\langle\frac{1}{V}\right\rangle=1.46 \times 10^{-3} \frac{\mathrm{~s}}{\mathrm{~m}}$
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4. ( 30 points) A rigid pressure vessel with a volume of $1 \mathrm{~m}^{3}$ contains 0.1 kmols of diatomic oxygen $\left(\mathrm{O}_{2}\right)$ and 0.1 kmols of diatomic nitrogen $\left(\mathrm{N}_{2}\right)$ at 300 K . The gas is then heated to 3000 K . At 3000 K , find the equilibrium ratio $\frac{N_{N} N_{N O}}{N_{O} N_{N_{2}}}$ for the reaction. Consider the reaction

$$
O+N_{2} \Leftrightarrow N O+N
$$

Assume that the molecules are rigid rotators and harmonic oscillators, and use the following data:

| Species | $\theta_{\text {rot }}(\mathrm{K})$ | $\theta_{\text {vib }}(\mathrm{K})$ | $\left(\mathrm{D}_{0} / \mathrm{hc}\right)\left(\mathrm{cm}^{-1}\right)$ |
| :---: | :--- | :---: | :--- |
| $\mathrm{N}_{2}$ | 1.998 | 2358 | 78,700 |
| NO | 1.704 | 1904 | 52,300 |

For the calculation of the electronic partition functions, consider the following electronic levels:
O: Level 0 [ $\left.g_{0}=5, \varepsilon_{0} / \mathrm{hc}=0 \mathrm{~cm}^{-1}\right]$, Level $1\left[\mathrm{~g}_{1}=3,\left(\varepsilon_{1} / \mathrm{hc}\right)=158 \mathrm{~cm}^{-1}\right]$,
Level $2\left[\mathrm{~g}_{2}=1,\left(\varepsilon_{2} / \mathrm{hc}\right)=226 \mathrm{~cm}^{-1}\right]$
$\mathrm{N}_{2}$ : Level 0 [ $\left.\mathrm{g}_{0}=1,\left(\varepsilon_{0} / \mathrm{hc}\right)=0 \mathrm{~cm}^{-1}\right]$
NO: Level 0 [ $\left.g_{0}=2,\left(\varepsilon_{0} / \mathrm{hc}\right)=0 \mathrm{~cm}^{-1}\right]$, Level $1\left[\mathrm{~g}_{1}=2,\left(\varepsilon_{1} / \mathrm{hc}\right)=121 \mathrm{~cm}^{-1}\right]$
N: Level 0 [ $\left.g_{0}=2,\left(\varepsilon_{0} / \mathrm{hc}\right)=0 \mathrm{~cm}^{-1}\right]$

Answer: $\frac{N_{N} N_{N O}}{N_{O} N_{N_{2}}}=7.00 \times 10^{-6}$
5. ( 30 points) The temperature of a $1.0 \mathrm{~cm}^{3}$ copper metal crystal is raised from $T_{1}=400 \mathrm{~K}$ to $T_{2}=500 \mathrm{~K}$. Calculate the internal energy change of the crystal $\left(U_{2}-U_{1}\right)$ due to lattice vibrations $\left(U_{2}-U_{1}\right)_{\text {vib }}$ and due to the free electron gas $\left(U_{2}-U_{1}\right)_{\text {elec }}$. Assume that the temperature is high enough that the Einstein model is accurate for determining the internal energy due to lattice vibrations. The Einstein temperature of the copper crystal is $\theta_{E}=265 \mathrm{~K}$. The density of the copper is $9000 \mathrm{~kg} / \mathrm{m}^{3}$, the atomic weight of the metal is copper is 63.5 $\mathrm{kg} / \mathrm{kmol}$, and there is one free electron per lattice site.

Answers: $\left(U_{2}-U_{1}\right)_{\text {vib }}=347 \mathrm{~J} \quad\left(U_{2}-U_{1}\right)_{\text {elec }}=3.20 \mathrm{~J}$

