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## Chem 30324, Spring 2018, Homework 6

## **Due March 7, 2018**

#### Quantum mechanics of vibrating NO.

The diatomic nitric oxide (NO) is an unusual and important molecule. It has an odd number of electrons, which is a rarity for stable molecule. It acts as a signaling molecule in the body, helping to regulate blood pressure, is a primary pollutant from combustion, and is a key constituent of smog. It exists in several isotopic forms, but the most common,  $^{14}$  N=  $^{16}$ O, has a bond length of 1.15077 Å and harmonic vibrational frequency of 1904 cm $^{-1}$ .

1. The ground vibrational wavefunction of N=O can be written

$$\Psi_{v=0}(x) = \left(\frac{1}{\alpha\sqrt{\pi}}\right)^{1/2} e^{-x^2/2\alpha^2}, \quad x = R - R_{eq}, \quad \alpha = \left(\frac{\hbar^2}{\mu k}\right)^{1/4}$$

where  $x=R-R_{eq}$ . Calculate  $\langle x \rangle$  and  $\langle x^2 \rangle$  for NO in the  $\Psi_{v=0}(x)$  state (you might want to use  $\alpha$  as a length unit).

- 2. Calculate the average potential energy,  $\langle V(x) \rangle$ , in the ground state, in units of hv. Hint: This is trivial to calculate given the answer to question 1!
- 3. Using conservation of energy and your answer to question 2, calculate the average kinetic energy,  $\langle T(x) \rangle$ , in the ground state, in units of  $h\nu$ . Comment on the relationship between the kinetic and potential energies. This is a general result for all  $\nu$ , and is a consequence of the virial theorem for the harmonic potential.

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4. Calculate the classical minimum and maximum values of the  $^{14}$ N= $^{16}$ O bond length for a molecule in the ground vibrational state. Hint: Calculate the classical limits on x, the value of x at which the kinetic energy is 0 and thus the total energy equals the potential energy.

5. Calculate the probability for a quantum mechanical  $^{14}N=^{16}O$  molecule to have a bond length outside the classical limits. This is an example of quantum mechanical tunneling.

## Statistical mechanics of vibrating NO

6. Using your knowledge of the harmonic oscillator and the Boltzmann distribution, complete the table below for the first four harmonic vibrational states of  $^{14}$  N= $^{16}$  O.

Quantum number	Energy (kJ/mol)	Relative population at 400 K	Relative population at 410 K
v = 0			
v = 1			
v = 2			
v = 3			

7. Use the table to estimate the average vibrational energy of a mole of  $^{14}$ N= $^{16}$ O at 400 and 410 K.

8. Use your answer to Question 7 to estimate the vibrational heat capacity ( dE/dT) of a mole of  $^{14}\rm N=^{16}O$  in this temperature range. How does your answer compare to the classical estimate, R=8.314 J/mol K?

9. Predict the harmonic vibrational frequency of the heavier cousin of  $^{14}$  N= $^{16}$  O,  $^{15}$  N= $^{18}$  O, in cm $^{-1}$ . Assume the force constant is independent of isotope. Do you think these two isotopes could be distinguished using infrared spectroscopy?

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# NO goes for a spin

 $^{14}N=^{16}O$  has an equilibrium bond length of 1.15077 Å

- 10. Calculate the moment of inertia of  $^{14}N = ^{16}O$ , in amu  $\text{\AA}^2$ , and the rotational energy constant, B, in kJ mol $^{-1}$  and in cm $^{-1}$ .
- 11. Imagine that the NO molecule is adsorbed flat on a surface upon which it is free to rotate. Plot out the energies of the four lowest-energy rotational quantum states, in units of B, being sure to include appropriate quantum numbers and degeneracies. Also indicate the total rotational angular moment of each state, in units of  $\hbar$ .
- 12. Derive a selection rule for light-induced excitation of the plane-spinning NO molecule. *Hint:* Treat the NO as a 2-D rotor. Find the conditions on  $\Delta m_l$  that make the transition dipole moment integral  $\langle \psi_{m_l} | x | \psi_{m'_l} \rangle$  non-zero. Recall that x can be written r cos  $\Phi$  in polar coordinates.
- 13. Now imagine the NO molecule is free to rotate in three-dimensional space. As in Question 2 above, plot out the energies of the four lowest-energy rotational quantum states, in units of B, being sure to include appropriate quantum numbers and degeneracies. Also indicate the total rotational angular moment of each state, in units of  $\hbar$ .
- 14. Use the vector model to sketch the total angular momentum vectors consistent with  $l=1\,.$