
Return by tue 9.11. by 2 pm.

1. *Simple model for quantum dots* (See the lecture notes given by Kimmo) Show that the eigenenergies of a two-dimensional harmonic potential are $\varepsilon_l = \hbar\omega(l+1)$, and degeneracies $d_l = l+1$ with $l = 0, 1, 2, \dots$ (Hint: Show that the Hamiltonian separates into a sum of two one-dimensional Hamiltonians and use the energies of one-dimensional harmonic oscillator.)

In the capacitive model of a quantum dot the addition energy is

$$\Delta_2(N) = \varepsilon_{N+1} - \varepsilon_N + e^2/C,$$

where ε_N is the energy of the single-electron level occupied by the N th electron and e^2/C is the capacitive charging energy. Use the simple model of a quantum dot where the non-interacting electrons occupy the levels of the two-dimensional harmonic oscillator, and plot the addition energies $\Delta_2(N)$ as a function of electron number N , for $N=1 \dots 13$. For simplicity use units where $\hbar\omega = e = 1$ and $C = N$, thus assume that the capacitance increases with the electron number. Compare your plot with the experimental one (see the lecture notes). What are the qualitative differences and similarities? **(2 points)**

2. Calculate the atomic polarizability, and hence the dielectric constant ϵ , of liquid Argon, for which the atomic number density is $2.128 \times 10^{28} \text{ m}^{-3}$ and the atomic radius is $1.18 \times 10^{-10} \text{ m}$. Use the so called Clausius-Mossotti relation

$$\frac{\epsilon - 1}{\epsilon + 2} = \frac{n\alpha}{3\epsilon_0}.$$

(The experimental dielectric constant is 1.538) **(2 points)**

3. *Frequency-dependent atomic polarizability.* Allow the local field acting on an ion be frequency-dependent, $\mathbf{E}_{loc} = \mathbf{E}_0 \exp(-i\omega t)$. The simplest classical theory of atomic polarizability treats the ion as an electronic shell of charge Ze and mass Zm tied to a rigid ion core by a harmonic spring, of spring constant $K = Zm\omega_0^2$. Argue that the equation of motion for the electronic shell is

$$Zm\ddot{\mathbf{r}} = -K\mathbf{r} - Ze\mathbf{E}_{loc},$$

where \mathbf{r} is the displacement from equilibrium. By assuming that $\mathbf{r} = \mathbf{r}_0 \exp(-i\omega t)$ show that the frequency-dependent atomic polarizability is

$$\alpha(\omega) = \frac{Ze^2}{m(\omega_0^2 - \omega^2)}.$$

(2 points)