

Fys2160 – 2013 – Oblig 1

Micro- and Macrostates in the Einstein Crystals

In this project we will address the micro- and macro-states of an Einstein crystal. You will learn how to represent and count microstates in a simple model of a crystal consisting of a set of independent oscillators, you will learn how to find the probability of a macrostate for two Einstein crystals in thermal contact, and to find the time evolution of the Einstein crystal using a Monte Carlo simulation technique.

A simple model for a crystal that still captures surprisingly many of the important features of the statistical physics of a crystal is the Einstein crystal. A real crystal consists of a set of atoms in a periodic configurations interacting through interatomic interactions that include both short range and longer ranged forces. As a result, individual atoms will oscillate around an equilibrium position while interacting mostly with its nearest neighbors. As a simplified model for this system we consider each atom, i , to behave like an independent harmonic oscillator with a potential energy U_i :

$$U_i(\vec{r}_i) = \frac{1}{2}k_x(x_i - x_{i,eq})^2 + \frac{1}{2}k_y(y_i - y_{i,eq})^2 + \frac{1}{2}k_z(z_i - z_{i,eq})^2, \quad (1)$$

From quantum mechanics, we know that the energy of a harmonic oscillator i is

$$\epsilon_i = n_i \Delta \epsilon, \quad (2)$$

where n_i is an integer describing the state of oscillator i . We can therefore describe the state of a crystal with N independent (meaning non-interacting) oscillators by the states n_i for $i = 1, \dots, N$. The total energy of the crystal in this simplified model is then:

$$U = \sum_{i=1}^N \epsilon n_i. \quad (3)$$

For simplicity we will measure energy in units of ϵ :

$$q = \frac{U}{\epsilon} = \sum_{i=1}^N n_i, \quad (4)$$

For a system with a given total energy, the sum of all the n_i is constant, but we can still change how the energy is distributed in the system. We can think of the energy a given number of energy units that we are free to distribute between the oscillators. Any distribution is allowed as long as we do not change the total energy.

We describe a microstate of this system by the numbers n_i for each oscillator:

$$\{n_1, n_2, \dots, n_N\} \quad (5)$$

For example, for a system with $N = 4$ and $q = 4$, a possible microstate is $\{1, 0, 2, 1\}$, that is $n_1 = 1$, $n_2 = 0$, $n_3 = 2$, and $n_4 = 1$.

We will now find the various microstates of this system:

- (a) For a system with $N = 2$ oscillators and $q = 3$, list all the possible microstates.
- (b) For a system with $N = 3$ oscillators and $q = 3$, list all the possible microstates.
- (c) For a system with $N = 4$ oscillators and $q = 3$, list all the possible microstates.

The general formula for the number of microstates for N oscillators with q units of energy is:

$$\Omega(N, q) = \binom{q + N - 1}{q} = \frac{(q + N - 1)!}{q!(N - 1)!}. \quad (6)$$

- (d) Check that the results you found above are consistent with this formula.

We can now list and count the number of microstates for an Einstein crystal, and we are ready to address what happens if two Einstein crystals come in contact. First, we start by looking at a system consisting of two isolated Einstein crystals, system A with N_A oscillators and energy q_A and system B with N_B oscillators and energy q_B . Each system is surrounded by an insulating, rigid and impermeable outer wall so that its energy, volume and number of oscillators (particles) is constant. The total system consists of system A and system B, so that $N = N_A + N_B$ and $q = q_A + q_B$. However, the systems are initially isolated – meaning that they are independent systems with constant energy, volume and number of particles.

- (e) For a system consisting of subsystem A with $N_A = 2$ and $q_A = 5$ and subsystem B with $N_B = 2$ and $q_B = 1$ list all possible microstates of the system.

The two systems are put in thermal contact, so that they can exchange energy, but the number of particles and the volume of each subsystem does not change. The total energy $q = q_A + q_B = 6$ is constant, but the energy can now be freely distributed between the two systems. Let us now count the number of possible microstates for each possible value of q_A and q_B .

- (f) For $N_A = 2$, $N_B = 2$, and $q = 6$ what are the possible values of q_A and q_B ? We call a state with a given q_A (and therefore also a given $q_B = q - q_A$) a macrostate for the system.
- (g) For each possible macrostate q_A find the number of compatible microstates.
- (h) Compare the total number of microstates available to the system before and after the systems came in thermal contact. Comment on the result. What aspects of this result do you think is general?
- (i) If all microstates have the same probability, what are the probability of each of the macrostates?
- (j) What is the probability of the initial macrostate before the two systems came in contact?
- (k) What is the probability of finding all the energy in system A?
- (l) What is the probability of finding exactly half the energy in system A?

We will now address larger systems numerically. You therefore need to write a script/program to find the number of macrostates and the probability of the macrostates. We start from the system we had above, but you will need to write a general program you can use for any value of N_A , N_B , q , and q_A .

- (m) For $N_A = N_B = 2$ and $q = 6$ write a program to find the number of microstates for each macrostate q_A and the probability $P(q_A) = \Omega(q_A)/\Omega_{TOT}$ for each macrostate. Compare with your results from above. Plot the probability $P(q_A)$ as a function of q_A .

We will now address a larger system with $N_A = 50$, $N_B = 50$ and $q = q_A + q_B = 100$.

- (n) Plot the probability $P(q_A)$ as a function of q_A for all possible values of q_A .

- (o) What is the most probable macrostate? What is the probability of the system being in the most probable macrostate compared to all other macrostates in the system? Comment on the result.
- (p) We start from a system with $q_A = 0$ and $q_B = 100$ before the systems come in thermal contact. What is the probability of being in this state after the system has reached equilibrium?

Explore if you dare..

We will now address how the system reaches thermal equilibrium. This is not an essential part of the project, but it may give you additional insight into irreversible processes and the second law of thermodynamics. Proceed if you dare...

The microstate of the system is given by the energy of each oscillator given as n_i for each of the oscillators. We start by studying a single system with N oscillators and energy q . You can generate the initial state by placing each energy unit in a random oscillator. (Some oscillators may receive more than one energy units and some may receive none). At each timestep we attempt a transfer of energy from one oscillator to another oscillator using the following algorithm. Select an oscillator n_1 at random. If the oscillator has zero energy, do nothing, if the oscillator has a positive energy, select another oscillator n_2 at random and transfer one unit of energy from n_1 to n_2 . Repeat the process for as many “timesteps” as you want.

- (q) Write a program to generate the initial microstate and the “time” development of the microstate. Plot the initial state and the state after 100 flips.
- (r) Divide your system into two parts, each part having $N/2$ oscillators. We call the first $N/2$ oscillators system A corresponding to oscillators $i = 1, N/2$, and the second $N/2$ oscillators system B, corresponding to oscillators $i = N/2 + 1, N$. Plot q_A/N_A and q_B/N_B as a function of time. Comment on the result.
- (s) Initialize the system with all the energy in the system A only. Plot the average energy per oscillator (q_A/N_A and q_B/N_B) as a function of time and comment on the result.
- (t) Starting from an equilibrium configuration (either by starting from a random configuration or by running a non-random configuration for a long time before starting measurements) plot the probability of the macrostates. Comment on the results.

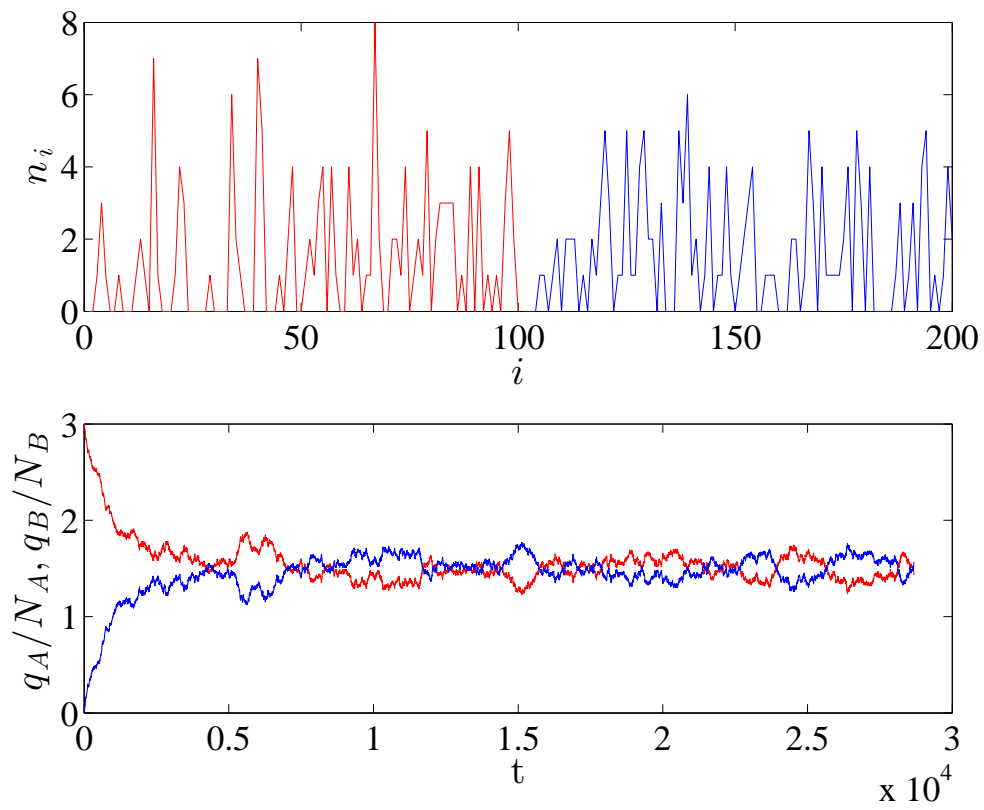


Figure 0.1: Example state (top) and time development of the average energy per oscillator for a system divided into two parts A and B.

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