

Last Time

The Position of a Particle Executing a Random Walk

The Probability of Finding a Particle at a Position after a Random Walk

Treating the Concentration as Time-Dependent Probability Distribution

Relating the Self-Diffusivity to a Random Walk

A Puzzle: Why for a random walk is $\nabla\mu = 0$?

3.21 Spring 2001: Lecture 14

The Successful Jump Frequency as an Activated Process

The treatment of diffusion as a statistical process permitted a physical correspondence between the macroscopic diffusivity and microscopic parameters for, average jump distance $\langle r \rangle$, jump correlation f , and the average frequency at which a jumper makes a finite jump Γ .

In this lecture, the statistical evaluation of microscopic process will be applied to the successful jump frequency Γ . A physical correspondence for Γ that is related to microscopic processes of attempt or natural atomic vibration frequency and the difference in energy between the potential energy of site and the maximum value of the minimum potential energy (the saddle energy) as the atom moves from one equilibrium site to the next.

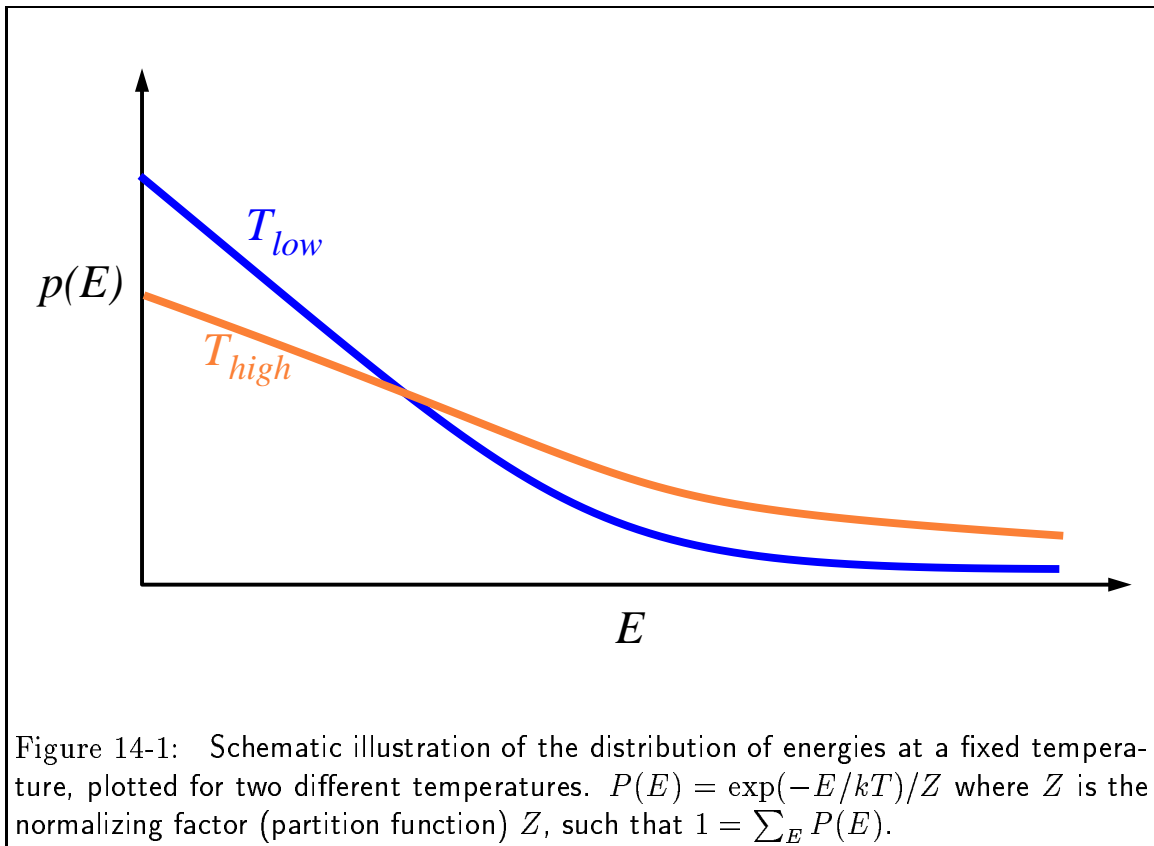
The result that will be obtain, that the frequency of successful hops,

$$\Gamma = \nu e^{-\frac{E_{saddle} - E_{equil.}}{kT}} \quad (14-1)$$

is related to the natural frequency multiplied by a Boltzmann factor has remarkable general application.

Distribution of Energy among Particles

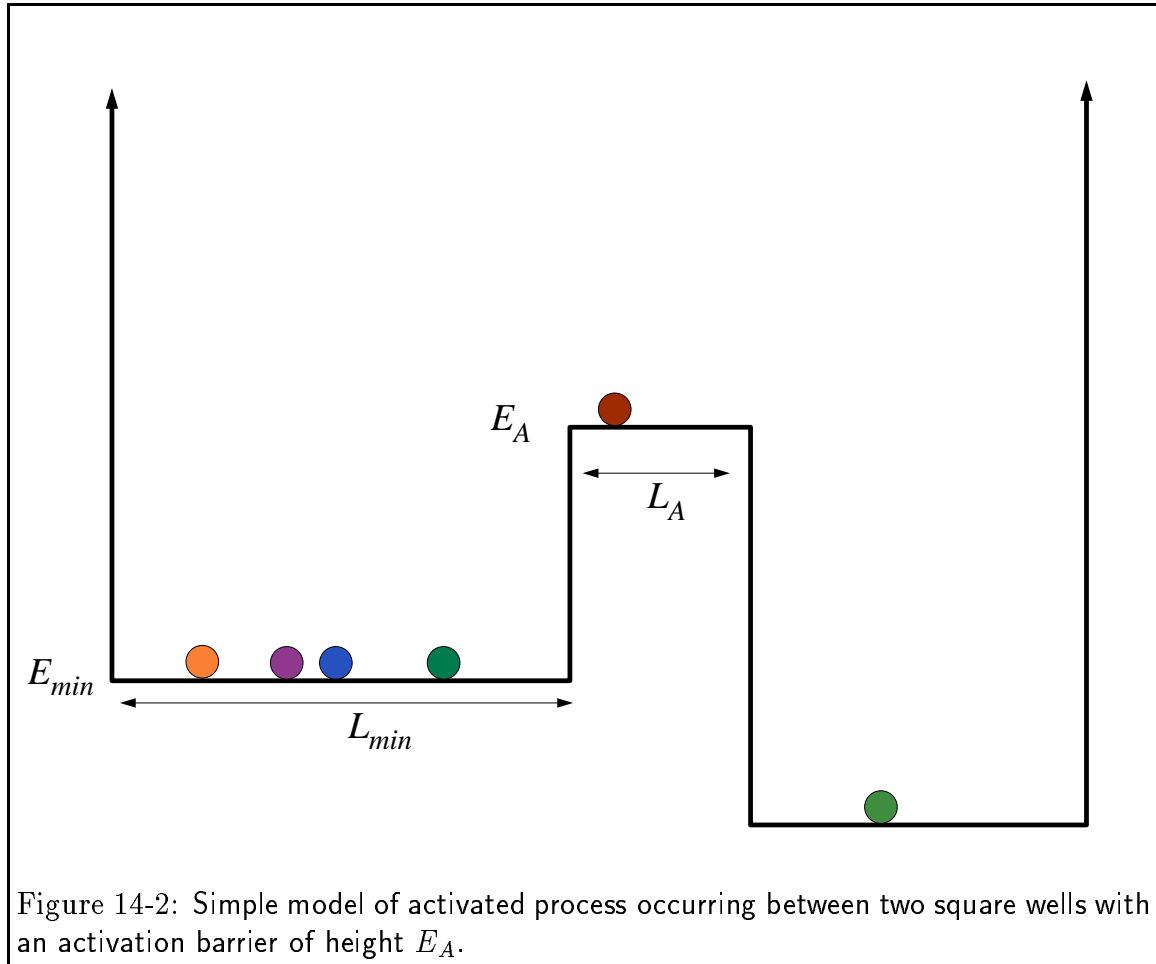
A fundamental result from statistical mechanics is that for an ensemble of atoms at a fixed temperature T , that the energies of the atoms has a characteristic probability distribution:



Below, the rate of successful jumps for simple models of activated processes will be derived. Each derivation will depend on the distribution of energies given above. It will be supposed that a single atom will assume all possible values of energy with probabilities given by the Boltzmann distribution over time (the ergodic assumption). In other words, the distribution is considered to apply to the atoms at a time scale that is rapid compared to the natural frequency of the atoms—no correlation is made for the loss (or gain) of energy as an atom hops from one equilibrium site to the next.

Activation Processes in Square Wells

Consider an ensemble of particles with distributed energies moving about on the following energy landscape:



The characteristic time it takes a particle to cross the activated state is

$$\tau_{cross} \approx \frac{L_A}{v_{rms}} \approx L_A \sqrt{\frac{m}{kT}} \quad (14-2)$$

where $v_{rms} = \sqrt{\langle v^2 \rangle}$ and m is the mass of the particle with characteristic thermal energy kT .

The total rate, R_{cross} , that particles cross the barrier is

$$\begin{aligned}
 R_{cross} &= \frac{\text{(number of particles in activated state)}}{\tau_{cross}} \\
 &= \frac{N_{tot} \text{(probability of being in activated state)}}{\tau_{cross}} \\
 &= N_{tot} \sqrt{\frac{kT}{m}} \frac{1}{L_A} \frac{Z_A}{Z_{min}}
 \end{aligned} \tag{14-3}$$

where Z_A and Z_{min} are the partition functions for the activated and minimum states.

The rate that single particle crosses, Γ , is:

$$\Gamma = \sqrt{\frac{kT}{m}} \frac{1}{L_A} \frac{Z_A}{Z_{min}} \tag{14-4}$$

$$\frac{Z_A}{Z_{min}} = \frac{\int_{L_A} e^{-\frac{E(x)}{kT}} dx}{\int_{L_{min}} e^{-\frac{E(x)}{kT}} dx} = \frac{L_A}{L_{min}} e^{-\frac{E_A - E_{min}}{kT}} \tag{14-5}$$

Therefore,

$$\Gamma = \sqrt{\frac{kT}{m}} \frac{1}{L_{min}} e^{-\frac{E_A - E_{min}}{kT}} \tag{14-6}$$

The term that multiplies the Arrhenius factor (the $1/T$ exponential) is the characteristic time it takes a particle to make an attempt at the activated state.

Activation Processes in Harmonic Wells

Consider the following modification of the above simple case, the minima are treated as harmonic wells:

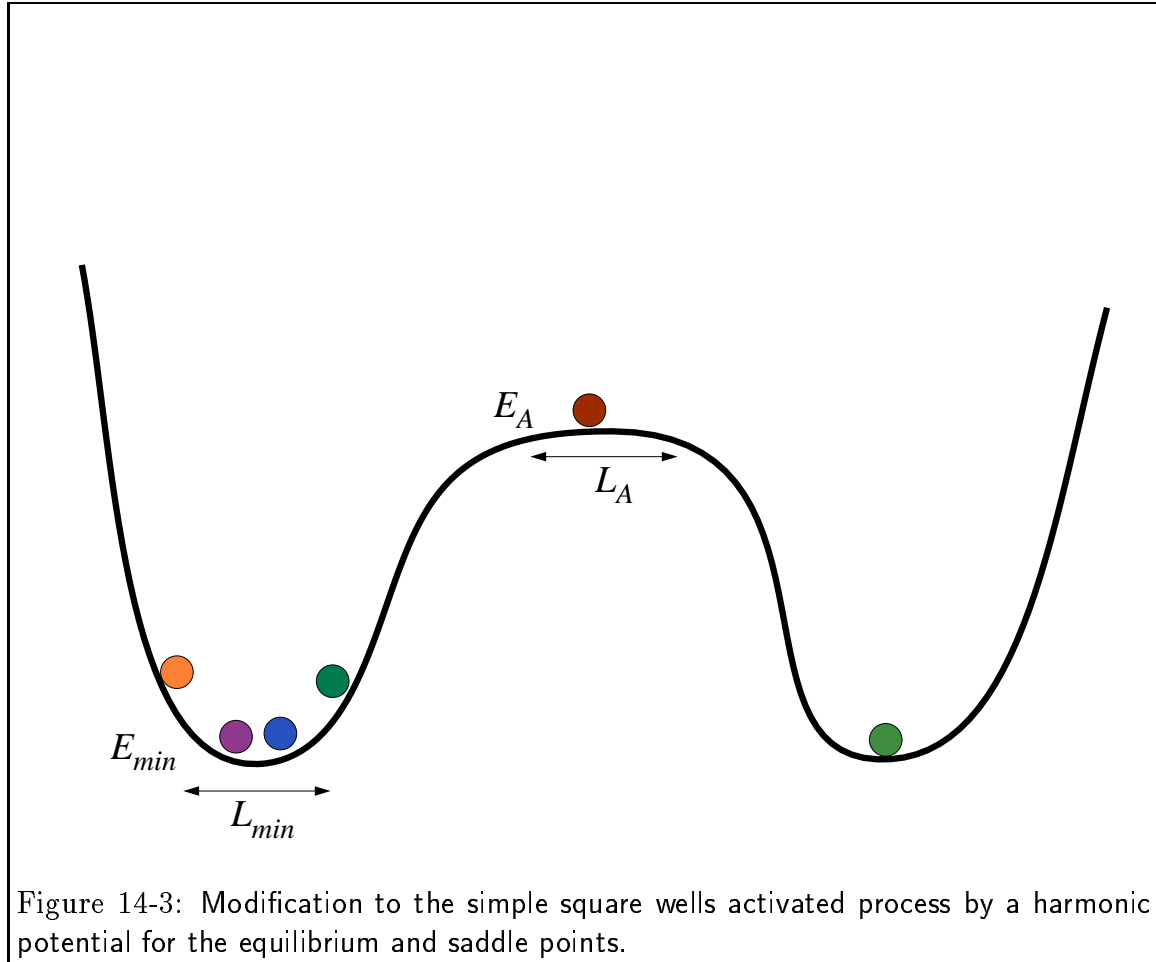


Figure 14-3: Modification to the simple square wells activated process by a harmonic potential for the equilibrium and saddle points.

The minima can be approximated by

$$E(x) = E_{min} + \frac{\beta}{2}(x - x_{min})^2 \quad (14-7)$$

The analysis is similar to the case of the square wells, but for the ratio of the partition functions:

$$\frac{Z_A}{Z_{min}} = \frac{\int_{L_A} e^{-\frac{E(x)}{kT}} dx}{e^{-\frac{E_{min}}{kT}} \int_{-\frac{L_{min}}{2}}^{\frac{L_{min}}{2}} e^{-\frac{\beta(x-x_{min})^2}{kT}} dx} \quad (14-8)$$

Approximating,

$$\int_{-\frac{L_{min}}{2}}^{\frac{L_{min}}{2}} e^{-\frac{\beta(x-x_{min})^2}{kT}} dx \approx \int_{-\infty}^{\infty} e^{-\frac{\beta(x-x_{min})^2}{kT}} dx \quad (14-9)$$

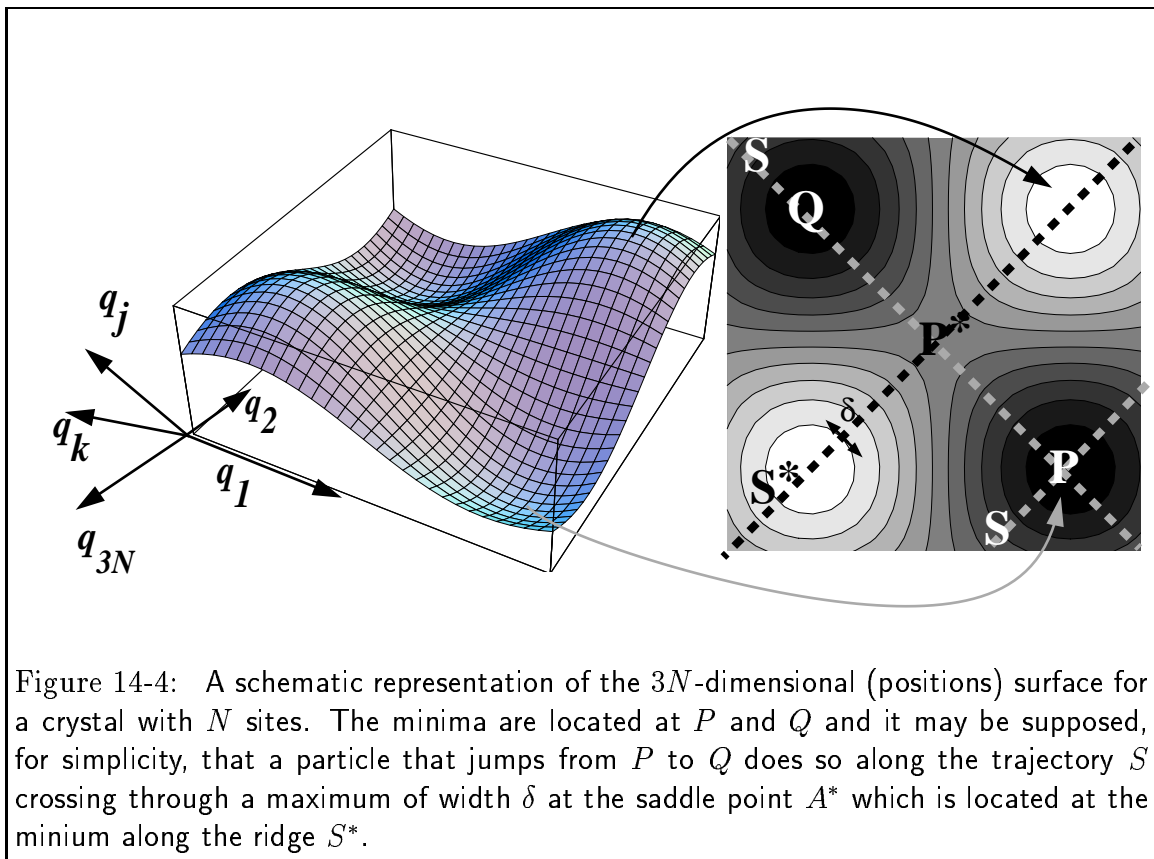
and carrying out the integration,

$$\Gamma = \sqrt{\frac{\beta}{m}} \frac{1}{\sqrt{2\pi}} e^{-\frac{E_A - E_{min}}{kT}} = \omega_o e^{-\frac{E_A - E_{min}}{kT}} \quad (14-10)$$

where ω_o is the characteristic oscillating frequency at the minima of a particle with mass m sitting in a well of curvature β .

Many-Body Theory of Activated Processes at Constant Pressure

In a real system, an atom or a vacancy does not make a successful hop without affecting (or getting effected by) its neighbors—all of the particles are vibrating and saddle point energy is an oscillating target produced by the random vibrations of all the atoms. The energy-surface that an atom, interstitial, or vacancy travels upon is a complicated and changing surface. If there are N spherical particles, then there are $6N$ -degrees of freedom to this surface, but it will be assumed that the momentum variables can be averaged out so that only a $3N$ -dimensional potential surface remains:



The minima, or equilibrium values of momenta and positions, can be approximated by harmonic wells:

$$E(q_i) = E_{min} \sum_{i=1}^{3N} \frac{m_i \omega_i^2}{2} q_i^2 \quad (14-11)$$

This is approximated by a one-dimensional problem by assuming that all states during a hop lie on the surface S in the figure.

Let the first coordinate be in the direction of the crossing (parallel to S), then the average (rms) momentum p_1 in that direction is related to the an average rate of attempts. The result that was derived for the harmonic potential can be re-used in this case:

$$\Gamma = \sqrt{\frac{kT}{m}} \frac{1}{L_A} \left(\begin{array}{l} \text{fraction} \\ \text{particles} \\ \text{activated} \\ \text{state} \end{array} \right) \text{ of in} \quad (14-12)$$

where L_A is recognized to be the width δ in Figure. 14-4.

However, in this case, the particle may have a different volume in the activated state compared to the equilibrium state:²⁷

$$V_{part}^{mig} = V_{part}^A - V_{part}^{eq} \quad (14-13)$$

For the case where the volume may vary, but pressure is constant, the canonical constant pressure partition function must be used:

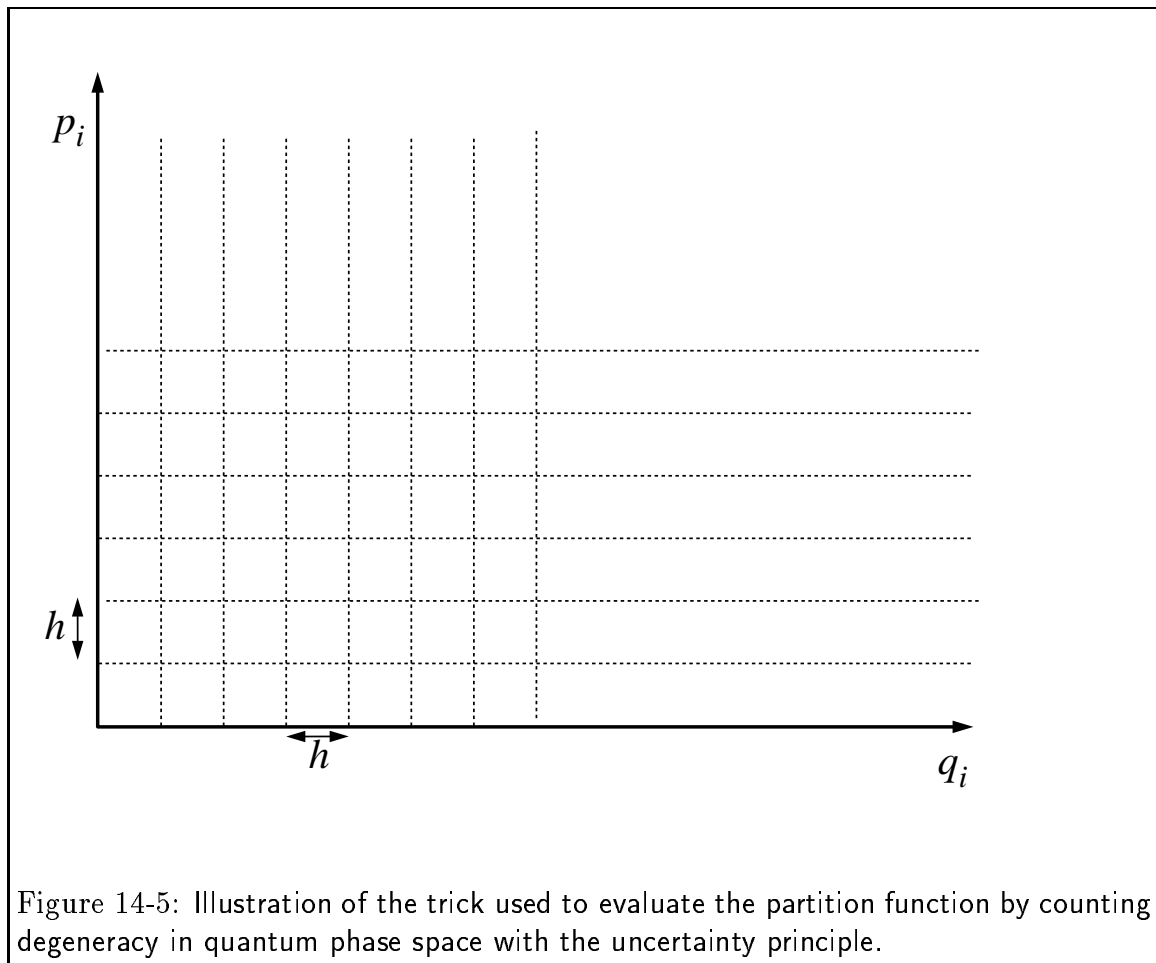
$$Z_P = \sum_{\text{energies, volumes}} e^{-\frac{(E+PV)}{kT}} \quad (14-14)$$

Therefore Γ_{part} picks up an additional factor:

$$\Gamma_{part} = \sqrt{\frac{kT}{m}} \frac{1}{L_A} e^{-\frac{-PV_{part}^{mig}}{kT}} \frac{Z_A}{Z_{min}} \quad (14-15)$$

It remains to evaluate the partition functions by summing over all energies: $Z = \sum_i^{3N} e^{-E(p_i, q_i)/kT}$. The partition function is evaluated by passing to the classical limit by dividing up the quantum phase space into cells of side-length equal to Planck's constant, h :

²⁷This is certainly important for the case for migrating particles that have a large compliance (low stiffness) such as vacancies.



Because of the uncertainty principle:

$$\Delta p_i \Delta q_i \geq h \quad (14-16)$$

Each elementary volume, $(\Delta p \Delta q)_i$, in phase space must be considered to have degeneracy:

$$\frac{\Delta p_i \Delta q_i}{h^{3N}} \quad (14-17)$$

Therefore,

$$Z = \frac{1}{h^{3N}} \sum_i e^{-\frac{E_i}{kT}} \quad (14-18)$$

In the classical limit,

$$Z = \frac{1}{h^{3N}} \int \int \dots \int_{-\infty}^{\infty} \sum_i e^{-\frac{E(p_i, q_i)}{kT}} dp_1 dp_2 \dots dp_{3N} dq_1 dq_2 \dots dq_{3N} \quad (14-19)$$

Using the Harmonic approximations for the minima and carrying out the integration and letting all the masses have the same value:

$$Z_{min} = \left(\frac{2\pi kT}{h} \right)^{3N} \left(\prod_{i=1}^{3N} \frac{1}{\omega_i} \right) e^{-\frac{E_{min}}{kT}} \quad (14-20)$$

Carrying out the same process for the activated state (which has one less degree of freedom) and adding the momentum near the activated state to the integral:

$$Z_A = L_A e^{-\frac{E_A}{kT}} \left(\frac{2\pi kT}{h} \right)^{3N-1} \left(\sqrt{\frac{m}{2\pi kT}} \right) \left(\prod_{i=2}^{3N-1} \frac{1}{\omega_i^A} \right) \quad (14-21)$$

The products over the vibrational modes can be related to the entropies of the states, i.e.,

$$\left(\prod_{i=2}^{3N-1} \omega_i^A \right) = \left(\frac{2\pi kT}{h} \right)^{3N-1} e^{-\frac{S^A}{k}} \quad (14-22)$$

Putting this all back into the expression for the rate of jumps,

$$\Gamma_{part} = \frac{kT}{h} e^{-\frac{(E_{part}^{mig} + P V_{part}^{mig} - T S_{part}^{mig})}{kT}} = \Gamma_{part} = \frac{kT}{h} e^{-\frac{G_{part}^{mig}}{kT}} \quad (14-23)$$
