

# Chem237: Lecture 15

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## 1 Classical Normal Mode Analysis

If we had a problem of a mass attached to a spring, we can immediately find the equations of motion from the harmonic oscillator model. However, we can always imagine a situation where our system consists of having multiple masses connected to each other by springs. For example, a water molecule  $\text{H}_2\text{O}$  can be thought of as 3 atoms connected to each other by 2 springs. In this system of coupled oscillators, the equations of motion that correspond to the molecular vibrations can be solved as if it were a coupled harmonic oscillator problem. Solving this problem using cartesian coordinates can be incredibly tedious and physically impossible if we keep adding more atoms to the system.

This can be made much simpler by performing a transformation from the cartesian basis to the normal mode basis. In the normal mode basis, we can imagine all of our atoms oscillating sinusoidally each with a frequency and phase relation that corresponds to a particular normal mode. Chemists use normal mode analysis to determine whether a normal mode is IR and/or Raman active, which is useful when creating vibrational spectra.

In this lecture, we will show the mathematical basis for normal mode analysis. But before we introduce how it can be applied to a quantum system such as atoms in a molecule, we will start with the classical analog.

### 1.1 Harmonic Approximation

We begin by writing the Hamiltonian for our system of interest with  $N$  vibrational degrees of freedom,

$$H = \sum_{i=1}^N \frac{p_i^2}{2m} + V(r_1, \dots, r_N) \quad (1)$$

where our coordinate vector is  $\mathbf{r} = \begin{bmatrix} r_1 \\ \vdots \\ r_N \end{bmatrix}$ . We can invoke the **Harmonic approximation**, which says that the potential  $V(\mathbf{r})$  can be expressed as a Taylor expansion about a minimum  $r_o$ , which we will take to be 0,

$$V(r) = V(r_o) + \sum_j \left( \frac{\partial V}{\partial r_j} \right) r_j + \frac{1}{2} \sum_{ij} r_i K_{ij} r_j + \dots \quad (2)$$

where  $\mathbf{K}$  is called the **Hessian** and is defined as

$$K_{ij} = \frac{\partial^2 V}{\partial r_i \partial r_j}. \quad (3)$$

Since our potential is expanded about a minimum, we can say that  $\left. \frac{\partial V}{\partial r_j} \right|_{r_j=r_o} = 0$ . If we truncate at the quadratic term and assume that  $V(r_o) \approx 0$ , then we obtain an approximate potential

$$V(r) \approx \frac{1}{2} \sum_{ij} r_i K_{ij} r_j. \quad (4)$$

The Hamiltonian can then be written in terms of matrices as

$$\mathbf{H} \approx \frac{1}{2} \mathbf{p}^T \mathbf{M}^{-1} \mathbf{p} + \frac{1}{2} \mathbf{r}^T \mathbf{K} \mathbf{r} \quad (5)$$

By doing this, we have effectively turned our system into one where each degree of freedom can be approximated as a harmonic oscillator. The equations of motion for each degree of freedom can then be solved by using Hamilton's equations:

$$\begin{aligned}\dot{r}_i &= \frac{\partial H}{\partial p_i} \\ \dot{p}_i &= -\frac{\partial H}{\partial r_i}\end{aligned}\tag{6}$$

However, this can be difficult when we have a system of coupled oscillators.

## 1.2 Normal Mode Coordinates

One way to get around this problem is by transforming our original coordinates into a set of normal mode coordinates. This decouples all of our degrees of freedom and turns our system into one of  $N$  independent harmonic oscillators, each with a normal mode frequency  $\omega_i$ . If our mass matrix  $\mathbf{M}$  is not diagonal, we could perform an eigenvalue decomposition and use the corresponding eigenvectors to find a diagonal matrix for  $\mathbf{M}$ , which would just be a matrix of its eigenvalues. Instead, we will introduce mass scaled coordinates to get rid of our mass matrix from the Hamiltonian. These coordinates can be defined as:

$$\begin{aligned}r'_i &= \sqrt{m_i} r_i & \iff & \mathbf{r}' = \mathbf{M}^{1/2} \mathbf{r} \\ p'_i &= \frac{1}{\sqrt{m_i}} p_i & \iff & \mathbf{p}' = \mathbf{M}^{-1/2} \mathbf{p} \\ K'_{ij} &= \frac{1}{\sqrt{m_i m_j}} K_{ij} & \iff & \mathbf{K}' = \mathbf{M}^{-1/2} \mathbf{K} \mathbf{M}^{1/2}\end{aligned}\tag{7}$$

So  $\mathbf{H}$  becomes,

$$\mathbf{H} = \frac{1}{2} \mathbf{p}'^T \mathbf{p}' + \frac{1}{2} \mathbf{r}'^T \mathbf{K}' \mathbf{r}'\tag{8}$$

where  $\mathbf{K}'$  is now our mass-scaled Hessian.

We can simplify the Hamiltonian even further by turning the mass-scaled Hessian into its diagonal form using an eigendecomposition,

$$\mathbf{K}' = \mathbf{U} \mathbf{\Lambda} \mathbf{U}^\dagger = \sum_{i=1}^N \lambda_i U_i U_i^\dagger\tag{9}$$

where  $\lambda_i$  is the  $i$ th eigenvalue of the Hessian and  $\mathbf{U}$  is a matrix of the eigenvectors. We will also make use of the **resolution of the identity**,

$$\mathbf{U} \mathbf{U}^\dagger = \mathbf{I} \iff \mathbf{I} = \sum_i U_i U_i^\dagger\tag{10}$$

which is a neat trick that is used very often in quantum mechanics. We can insert the identity matrix anywhere in our equation because it is like multiplying by 1.

$$\mathbf{H} = \frac{1}{2} \mathbf{p}'^T \mathbf{p}' + \frac{1}{2} \mathbf{r}'^T \mathbf{K}' \mathbf{r}' \longrightarrow \mathbf{H} = \frac{1}{2} \mathbf{p}'^T \mathbf{U} \mathbf{U}^\dagger \mathbf{p}' + \frac{1}{2} \mathbf{r}'^T \mathbf{U} \mathbf{\Lambda} \mathbf{U}^\dagger \mathbf{r}'\tag{11}$$

Then we will define the normal mode coordinates and momenta as

$$\tilde{\mathbf{r}} = \mathbf{U}^\dagger \mathbf{r}' \quad ; \quad \tilde{\mathbf{p}} = \mathbf{U}^\dagger \mathbf{p}'\tag{12}$$

which gives us the simplified Hamiltonian,

$$\begin{aligned}\mathbf{H} &= \frac{1}{2} \tilde{\mathbf{p}}^T \tilde{\mathbf{p}} + \frac{1}{2} \tilde{\mathbf{r}}^T \mathbf{\Lambda} \tilde{\mathbf{r}} \\ &= \sum_{i=1}^N \left( \frac{1}{2} \tilde{p}_i^2 + \frac{\lambda_i}{2} \tilde{r}_i^2 \right)\end{aligned}\tag{13}$$

where each normal mode oscillates with a single frequency  $\omega_i = \sqrt{\lambda_i}$ .

The analysis described in this lecture is simply transforming from one coordinate system to another so that whenever we are given  $r_1, r_2, \dots$ , for any system, we can immediately use equation (12) to find the coordinates in the normal mode basis. All we need to find is the matrix  $\mathbf{U}$  that does this.