

Further Quantum Mechanics

University of Cambridge Part II Natural Sciences Tripos

Yue Wu

*Yusuf Hamied Department of Chemistry
Lensfield Road,
Cambridge, CB2 1EW*

yw628@cam.ac.uk

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Nothing in these lecture notes is original. They are largely based on the notes by Dr. John Morgan, who lectured this course in 2025. Moreover, they are nowhere near accurate representations of what was actually lectured, and in particular, all errors are almost surely mine.

Preface

This course focuses on quantum mechanics, and it is slightly more advanced than what you have learned in Part IB Chemistry A: *Introduction to Quantum Mechanics*. It will mostly focus on perturbation theory, including both the time independent and the time dependent cases, and it also covers topics that are removed from the A4 *Theoretical Techniques* course this year, namely normal modes. This course will avoid the rigorous mathematical formulation of quantum mechanics, and especially, it will not introduce concepts like projective Hilbert space or functional analysis. If you want a more mathematical approach to quantum mechanics, you can find my notes on Mathematical Tripos Part II: *Principles of Quantum Mechanics*.

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1 Foundational Principles

We will start from a revision of the foundational principles of quantum mechanics that should be familiar from part IB Chemistry A.

1.1 Wavefunctions and Operators

In quantum mechanics, all physical information about a system is embodied in its wavefunction, denoted Ψ . The wavefunction is complex-valued, and we will use the *position representation* of the wavefunction, so it is a function of the spatial coordinates. In the Born's interpretation, the probability density of finding a particle at \mathbf{r} is

$$P(\mathbf{r}) \propto |\Psi(\mathbf{r})|^2 . \quad (1.1)$$

Wavefunctions should be single-valued and (at least) twice differentiable. Under such interpretation, we would often choose to normalise the wavefunction such that

$$\int d\tau \Psi^* \Psi = 1 , \quad (1.2)$$

where $d\tau$ is a shorthand notation for integrating over all spatial coordinates. This integral should converge for a proper wavefunction, so that it can be normalised.

A quantum mechanical system is defined by its Hamiltonian, \hat{H} , the total energy operator. The Hamiltonian usually includes the kinetic and potential energies of the particles. A quantum mechanical system evolves according to the time-dependent Schrödinger equation

$$i\hbar \frac{\partial}{\partial t} \Psi = \hat{H} \Psi . \quad (1.3)$$

Often, the Hamiltonian operator is independent of time. For such time-independent system, the wavefunction satisfies the time-independent Schrödinger equation (which we often directly refer to as the Schrödinger equation)¹

$$\hat{H}\psi = E\psi , \quad (1.5)$$

where the constant E is the energy of the system² and $\psi(\mathbf{r})$ is now a wavefunction independent of time. This is a (partial) differential equation, so it is only analytically solvable in a few limited cases, most of which you have seen already. Approximations are generally needed to solve for more complex systems, and one of the most important approximation techniques, the perturbation theory, is exactly the main theme of this course.

In general, the Schrödinger equation will have multiple (usually a countably infinite number of) solutions, which can be indexed by a quantum number n such that

$$\hat{H}\psi_n = E_n\psi_n . \quad (1.6)$$

The ψ_n are different states available to the system, and the state with the lowest energy E_n is known as the ground state. We usually arrange the states in sequence so that the ground state is labelled ψ_0 (or ψ_1 if you find it more convenient to start numbering from $n = 1$). We say two or more states are degenerate if they have the same energy. Any linear combination of degenerate wavefunctions is also a solution to the Schrödinger equation with the same energy.

¹This is because if \hat{H} is independent of t , then a special class of solutions of the time-dependent Schrödinger equation exists

$$\Psi(\mathbf{r}, t) = \psi(\mathbf{r})e^{-iEt/\hbar} , \quad (1.4)$$

where $\psi(\mathbf{x})$ is independent of time and satisfies the time-independent Schrödinger equation.

²This is essentially the conservation of energy — if a system has time-translational symmetry, then the energy of the system is conserved. This is an example of the Noether's theorem.

The Schrödinger equation is a specific example of the more general eigenvalue equation. In quantum mechanics, all physical observables A has a corresponding operator \hat{A} , if the wavefunction satisfies

$$\hat{A}\psi = a\psi, \quad (1.7)$$

then the value of A will always be measured to be a . If this equation is not satisfied, then measured values of A will be drawn from a probability distribution. The expectation and the uncertainty of A is

$$\langle A \rangle = \frac{\int d\tau \psi^* \hat{A} \psi}{\int d\tau \psi^* \psi} \quad (1.8)$$

$$\Delta A = \sqrt{\langle A^2 \rangle - \langle A \rangle^2}. \quad (1.9)$$

To avoid writing integrations over and over again, we introduce the Dirac bra-ket notation, in which a bra is $\langle \psi | \equiv \psi^*$ and a ket is $|\psi\rangle \equiv \psi$, and a pair of a bra and a ket forms a bra-ket (or just bracket), which implies integrations³

$$\langle \psi | \psi \rangle := \int d\tau \psi^* \psi. \quad (1.10)$$

An operator can be placed in middle of a bra-ket, so

$$\langle \psi | \hat{A} | \psi \rangle := \int d\tau \psi^* \hat{A} \psi. \quad (1.11)$$

To know what the quantum operator \hat{A} corresponding to a physical observable A is, we write the classical expression of A in terms of position and momentum, and we replace the position and momentum with their quantum operators⁴

$$\hat{x} \longrightarrow x \quad (1.12)$$

$$\hat{p}_x \longrightarrow -i\hbar \frac{\partial}{\partial x}. \quad (1.13)$$

In general, a wavefunction cannot simultaneously be an eigenfunction of two (or more) operators. However, this will happen if and only if the commutator of the two operators, defined as

$$[\hat{A}, \hat{B}] := \hat{A}\hat{B} - \hat{B}\hat{A} \quad (1.14)$$

is zero. If this is the case, then we say the two operators commute. If the condition is not met, then the commutator places a lower bound on the product of the uncertainties

$$\Delta A \Delta B \geq \frac{1}{2} |\langle [A, B] \rangle|. \quad (1.15)$$

1.2 Hermitian Operators

Not all operators can correspond to physical observables. First of all, it has to be linear, so that

$$\hat{A}(a|\phi\rangle + b|\psi\rangle) = a\hat{A}|\phi\rangle + b\hat{A}|\psi\rangle, \quad (1.16)$$

³Actually more formally, $|\psi\rangle$ what truly fundamental, and is the quantum state of a system living in an abstract vector space called a (projective) Hilbert space \mathcal{H} . The wavefunction $\psi(\mathbf{x})$ is just the position-space representation of the vector $|\psi\rangle$. If a quantum state is a vector in the Hilbert space, $|\psi\rangle \in \mathcal{H}$, then an operator is an operator in \mathcal{H} , $\hat{A} : \mathcal{H} \rightarrow \mathcal{H}$, and a bra will be a dual vector living in the dual space of \mathcal{H} , $\langle \psi | \in \mathcal{H}^*$. What allows us to do this is the Riesz representation theorem, which shows for any $|\psi\rangle \in \mathcal{H}$, there is always a corresponding $\langle \psi | \in \mathcal{H}^*$ and *vice versa*.

⁴This is the position-space representations of these operators, which we can act on wavefunctions that are written as a function of spatial coordinates. These operators have different representations in different basis.

where $a, b \in \mathbb{C}$ are constants. Moreover, operators that correspond to physical observable must be Hermitian. For any linear operator \hat{A} , we define its adjoint \hat{A}^\dagger to be the operator such that

$$\langle \phi | \hat{A} | \psi \rangle = \langle \hat{A}^\dagger \phi | \psi \rangle . \quad (1.17)$$

A operator is Hermitian if it is self-adjoint,⁵ meaning $\hat{A} = \hat{A}^\dagger$, so that

$$\langle \phi | \hat{A} | \psi \rangle = \langle \hat{A} \phi | \psi \rangle = \langle \psi | \hat{A} | \phi \rangle^* . \quad (1.18)$$

It is not difficult to show that taking the adjoint has the following properties:

$$(a\hat{A})^\dagger = a^* \hat{A}^\dagger \quad (1.19)$$

$$(\hat{A} + \hat{B})^\dagger = \hat{A}^\dagger + \hat{B}^\dagger \quad (1.20)$$

$$(\hat{A}\hat{B})^\dagger = \hat{B}^\dagger \hat{A}^\dagger \quad (1.21)$$

$$(\hat{A} | \psi \rangle)^\dagger = \langle \psi | \hat{A}^\dagger . \quad (1.22)$$

A Hermitian operator has the following nice properties:

- (i) The eigenvalues are all real.
- (ii) Eigenfunctions with different eigenvalue are orthogonal. Even if some eigenfunctions are degenerate, orthogonal eigenfunctions can always be constructed.
- (iii) The eigenfunction form a complete basis set.

Two functions are orthogonal means that their inner product, defined as $\langle \phi | \psi \rangle$, is zero. Moreover, the completeness of the basis set means that any wavefunction ψ satisfying the boundary conditions can be expressed as a linear combination of the eigenfunctions,

$$\psi = \sum_i c_i \phi_i . \quad (1.23)$$

We can always make this basis set, which we denote as $\{\phi_i\}$, orthonormal by normalising the orthogonal eigenfunctions, so that $\langle \phi_i | \phi_j \rangle = \delta_{ij}$. Then taking the inner product of the above eigenfunction expansion with ϕ_j , we have

$$\begin{aligned} \langle \phi_j | \psi \rangle &= \left\langle \phi_j \left| \sum_i c_i \phi_i \right. \right\rangle \\ &= \sum_i c_i \langle \phi_j | \phi_i \rangle \\ &= \sum_i c_i \delta_{ij} = c_j , \end{aligned} \quad (1.24)$$

and so

$$|\psi\rangle = \sum_i \langle \phi_i | \psi \rangle |\phi_i\rangle . \quad (1.25)$$

From now on, we will denote the eigenstates just by its quantum number in the bra-ket notation: $|i\rangle \equiv |\phi_i\rangle$, so that for example, we have

$$|\psi\rangle = \sum_i \langle i | \psi \rangle |i\rangle . \quad (1.26)$$

⁵There is actually some extremely subtle differences between self-adjoint and Hermitian operators in functional analysis, but they would only bother pure mathematicians.

1.3 Unitary Operators

Apart from position space representations, there are many other basis we can represent the wavefunctions, such as the momentum space, or the basis spanned by eigenstates. An important class of linear operators is the *unitary operators* that we use here for transformations of basis.⁶ To define unitary operators, we first need to define the identity operator. We define the identity operator to be an operator \hat{I} such that

$$\hat{I} |\psi\rangle = |\psi\rangle \quad (1.27)$$

for any $|\psi\rangle$. An useful way of constructing the identity operator is the *resolution of identity*. For any complete orthonormal basis set $\{|\phi_i\rangle\}$, we have

$$\hat{I} = \sum_i |\phi_i\rangle \langle \phi_i| . \quad (1.28)$$

This is because if we act this operator on any $|\psi\rangle$, we get its eigenfunction expansion is that basis

$$\hat{I} |\psi\rangle = \sum_i |\phi_i\rangle \langle \phi_i | \psi \rangle = \sum_i \langle \phi_i | \psi \rangle |\phi_i\rangle = |\psi\rangle , \quad (1.29)$$

where we moved $\langle \phi_i | \psi \rangle$ to the front to make the expression more obvious — we are allowed to do this because $\langle \phi_i | \psi \rangle$ is just a number.

Having defined the identity operator, we can define the inverse of an operator \hat{A} , denoted \hat{A}^{-1} , to be the operator such that

$$\hat{A} \hat{A}^{-1} = \hat{A}^{-1} \hat{A} = \hat{I} . \quad (1.30)$$

A unitary operator \hat{U} is one for which the adjoint is equal to the inverse,

$$\hat{U}^{-1} = \hat{U}^\dagger , \quad (1.31)$$

so that

$$\hat{U} \hat{U}^\dagger = \hat{U}^\dagger \hat{U} = \hat{I} . \quad (1.32)$$

As we claimed before, unitary operators are used to define transformations between different representations. Therefore, if we have an operator \hat{A} , we would like to know what its representation has transformed into after the transformation. Suppose the operator \hat{A} acts on a general wavefunction ψ to get

$$\hat{A} \psi = \phi , \quad (1.33)$$

and after the transformation, we have

$$\tilde{\psi} = \hat{U} \psi , \quad \tilde{\phi} = \hat{U} \phi . \quad (1.34)$$

Then we would like to find \tilde{A} , which is the representation of \hat{A} after the transformation, such that

$$\tilde{A} \tilde{\psi} = \tilde{\phi} . \quad (1.35)$$

We have

$$\tilde{A} \hat{U} \psi = \hat{U} \phi = \hat{U} \hat{A} \psi \implies \tilde{A} \hat{U} = \hat{U} \hat{A} . \quad (1.36)$$

Therefore,

$$\tilde{A} = \hat{U} \hat{A} \hat{U}^\dagger \quad \text{or} \quad \hat{A} = \hat{U}^\dagger \tilde{A} \hat{U} . \quad (1.37)$$

By taking the adjoint of the above equation, we can see that a Hermitian operator after unitary transformation is still an Hermitian operator.

⁶Technically, the “unitary operator” we introduce here is not quite the same as the unitary operators you will see in standard Quantum Mechanics textbook. Here, the “unitary operators” are used for basis transformations, just like unitary matrices for basis transformations in vector spaces. However, in other texts, “unitary operators” usually refer to the ones that actually changes the quantum state itself, such as evolving a quantum state over time.

Moreover, a transformed eigenfunction of an operator is still an eigenfunction of the transformed operator with the same eigenvalue. This means that if

$$\hat{A}\psi_n = a_n\psi_n, \quad (1.38)$$

then since we can insert an identity operator anywhere,

$$\hat{A}\hat{U}^\dagger\hat{U}\psi_n = a_n\hat{U}^\dagger\hat{U}\psi_n, \quad (1.39)$$

and so

$$\hat{U}\hat{A}\hat{U}^\dagger\hat{U}\psi_n = a_n\hat{U}\psi_n, \quad (1.40)$$

which implies

$$\tilde{A}\tilde{\psi}_n = a_n\tilde{\psi}_n. \quad (1.41)$$

We can also show that the expectation values are the same:

$$\begin{aligned} \int d\tau \psi^* \hat{A} \psi &= \int d\tau \psi^* \hat{U}^\dagger \hat{U} \hat{A} \hat{U}^\dagger \hat{U} \psi \\ &= \int d\tau (\hat{U} \psi)^* \hat{U} \hat{A} \hat{U}^\dagger \hat{U} \psi \\ &= \int d\tilde{\tau} \tilde{\psi}^* \tilde{A} \tilde{\psi}. \end{aligned} \quad (1.42)$$

If we let $\hat{A} = \hat{I}$, then

$$\int d\tilde{\tau} \tilde{\psi}^* \tilde{\psi} = \int d\tau \psi^* \psi, \quad (1.43)$$

so a normalised state remains normalised after transformation.⁷

1.4 The Momentum Space Representation

Consider the transformation, in one-dimensional space, defined by the unitary operator

$$\phi(p) = \hat{U}\psi(x) = \frac{1}{\sqrt{2\pi}} \int_{-\infty}^{\infty} dx e^{-ipx} \psi(x), \quad (1.44)$$

which you may recognise as the *Fourier transform*. This is a unitary transformation because its inverse transform is given by

$$\psi(x) = \hat{U}^\dagger \phi(p) = \frac{1}{\sqrt{2\pi}} \int_{-\infty}^{\infty} dp e^{ipx} \phi(p), \quad (1.45)$$

which is exactly the adjoint of the forward transformation. This transforms our wavefunction $\phi(x)$ from the position basis to the momentum basis.

Before we proceed, we first introduce a useful result.

Proposition 1.1.

$$\frac{1}{2\pi} \int_{-\infty}^{\infty} dx e^{i(p-p')x} = \delta(p-p'), \quad (1.46)$$

where $\delta(x)$ is the *Dirac delta function*.⁸

⁷We are deliberately not using the bra-ket notation here because technically, ψ and $\tilde{\psi}$ corresponds to the same quantum state $|\psi\rangle$, just being represented in different basis. For example, if ψ_n is represented in the position (x) basis and $\tilde{\psi}_n$ is represented in the momentum (p) basis, then $\psi_n(x) = \langle x|n\rangle$ and $\tilde{\psi}_n(p) = \langle p|n\rangle$, where $|x\rangle$ and $|p\rangle$ are the position and momentum eigenstates.

⁸Dirac delta function $\delta(x)$ is a weird ‘function’ defined such that $\delta(x) \neq 0 \ \forall x \neq 0$ but for any $\epsilon > 0$, we have

$$\int_{-\epsilon}^{\epsilon} dx \delta(x) = 1. \quad (1.47)$$

You can think of this as a normalised Gaussian in the limit of its variance $\rightarrow 0$.

Dirac delta has a nice property.

$$\int_{-\infty}^{\infty} dx f(x) \delta(x - x') = f(x'). \quad (1.48)$$

Now it's time to investigate the momentum space representation. Specifically, we would like to find out how the position and momentum operators look like in the momentum representation. We do this by rewriting the position-space expectation value of momentum into the momentum space, and we find

$$\begin{aligned} \langle p \rangle &= \int_{-\infty}^{\infty} dx \psi^*(x) \left(-i \frac{\partial}{\partial x} \right) \psi(x) \\ &= \frac{1}{2\pi} \int_{-\infty}^{\infty} dx \left[\int_{-\infty}^{\infty} dp \phi(p)^* e^{-ipx} \right] \left(-i \frac{\partial}{\partial x} \right) \left[\int_{-\infty}^{\infty} dp' \phi(p') e^{+ip'x} \right] \\ &= \frac{1}{2\pi} \int_{-\infty}^{\infty} dx \int_{-\infty}^{\infty} dp \int_{-\infty}^{\infty} dp' p' \phi(p)^* \phi(p') e^{i(p-p')x} \\ &= \int_{-\infty}^{\infty} dp \int_{-\infty}^{\infty} dp' p' \phi(p)^* \phi(p') \delta(p' - p) \\ &= \int_{-\infty}^{\infty} dp \phi(p)^* p \phi(p). \end{aligned} \quad (1.49)$$

This is exactly the momentum-space average of the operator p . We see that in the momentum representation, the momentum operator is unsurprisingly p itself — and in fact that is why we call this representation the momentum representation.

To work out the momentum representation of the position operator, we do the same trick again.

$$\begin{aligned} \langle x \rangle &= \int_{-\infty}^{\infty} dx \psi(x)^* x \psi(x) \\ &= \int_{-\infty}^{\infty} dx \left[\int_{-\infty}^{\infty} dp' \phi(p')^* e^{-ip'x} \right] x \left[\int_{-\infty}^{\infty} dp \phi(p) e^{ipx} \right]. \end{aligned} \quad (1.50)$$

Since

$$-i \frac{\partial}{\partial p} e^{ipx} = x e^{ipx}, \quad (1.51)$$

we can rewrite it as

$$\langle x \rangle = \frac{1}{2\pi} \int_{-\infty}^{\infty} dx \int_{-\infty}^{\infty} dp' \int_{-\infty}^{\infty} dp \phi(p')^* \phi(p) \left(-i \frac{\partial}{\partial p} e^{i(p-p')x} \right). \quad (1.52)$$

We now use integration by part, and we get

$$\begin{aligned} \langle x \rangle &= \frac{1}{2\pi} \int_{-\infty}^{\infty} dx \int_{-\infty}^{\infty} dp' \int_{-\infty}^{\infty} dp \phi(p')^* e^{i(p-p')x} \left(i \frac{\partial}{\partial p} \phi(p) \right) \\ &= \int_{-\infty}^{\infty} dp' \int_{-\infty}^{\infty} dp \delta(p - p') \phi(p')^* \left(i \frac{\partial}{\partial p} \phi(p) \right) \\ &= \int_{-\infty}^{\infty} dp \phi(p)^* \left(i \frac{\partial}{\partial p} \right) \phi(p). \end{aligned} \quad (1.53)$$

This is the expectation integral of the operator $i\partial/\partial p$.

Proposition 1.2. The momentum representation of the position and momentum operators are

$$\tilde{x} = \hat{U} \hat{x} \hat{U}^\dagger = i \frac{\partial}{\partial p} \quad (1.54)$$

$$\tilde{p} = \hat{U} \hat{p} \hat{U}^\dagger = p. \quad (1.55)$$

1.4.1 The State Vector

You can see that the even though we transformed a wavefunction from the position representation to the momentum representation, it is still fundamentally the same thing — both representations describe the same underlying quantum state. Although we may define various unitary transformations and trying to represent a quantum state in different representations, there must be a invariant underlying object. Just as we may choose different basis to describe a vector, and the vector may have different coordinates in different basis, it is still the same vector. We can do the same thing in quantum mechanics.

The true underlying quantum state is called a *state vector*, denoted $|\psi\rangle$, and it lives in a special vector space called a *Hilbert space*. The position representation of a wavefunction is just writing out the coefficients of a state vector in the position basis $\{|x\rangle\}$, and so $\psi(x) = \langle x|\psi\rangle$. Similarly the momentum representation is $\tilde{\psi}(p) = \langle p|\psi\rangle$.

Let's think again what we are doing when transforming the position wavefunction $\psi(x) = \langle x|\psi\rangle$ to the momentum wavefunction $\tilde{\psi}(p) = \langle p|\psi\rangle$. What we are doing is actually inserting an identity operator⁹

$$\hat{I} = \int dx |x\rangle \langle x| \quad (1.56)$$

into the momentum wavefunction:

$$\tilde{\psi}(p) = \langle p|\psi\rangle = \langle p| \int dx |x\rangle \langle x|\psi\rangle = \int dx \langle p|x\rangle \langle x|\psi\rangle = \int dx \langle p|x\rangle \psi(x). \quad (1.57)$$

All we need to do is to find $\langle p|x\rangle$. We can transform it into a more familiar object, $\langle p|x\rangle = \langle x|p\rangle^*$ — it is the Hermitian conjugate of the position representation of momentum eigenstate wavefunctions. We can find it as the eigenfunction of $-\mathrm{i}\hbar/dx$. You should have done this in Part IB Chemistry A: *Introduction to Quantum Mechanics*, $\langle x|p\rangle = e^{\mathrm{i}px}/\sqrt{2\pi}$. Therefore, we have

$$\tilde{\psi}(p) = \frac{1}{\sqrt{2\pi}} \int dx e^{-\mathrm{i}px} \psi(x). \quad (1.58)$$

We can use any other basis we want to represent a quantum state, but it should better be orthonormal and it should be complete, meaning that the basis should span the whole Hilbert space so that we can represent any quantum state we want in such a basis — the eigenstates of a Hermitian operator seems perfect for this purpose.¹⁰ Then a quantum states is defined by the countably infinite number of coefficients of each basis vector. This leads to the *vectors and matrices representations*.

1.5 Vectors and Matrices Representations

If we have a quantum state $|\psi\rangle$ and a complete, orthogonal basis set $\{|\phi_i\rangle\}$, then we can expand and write

$$|\psi\rangle = \sum_i c_i |\phi_i\rangle, \quad (1.59)$$

where as we claimed before, if the basis functions $|\phi_i\rangle$ are orthonormal, then the coefficients c_i are given by

$$c_i = \langle \phi_i|\psi\rangle. \quad (1.60)$$

This expansion be rewritten in matrix notation as

$$|\psi\rangle = (|\phi_1\rangle \quad |\phi_2\rangle \quad \cdots) \begin{pmatrix} c_1 \\ c_2 \\ \vdots \end{pmatrix} = \boldsymbol{\phi}^T \mathbf{c}, \quad (1.61)$$

⁹We are using an integral instead of a sum since we have a continuum of position eigenstates.

¹⁰In fact the position and the momentum representations are one of the worst representations. They are examples of continuum bases, and their eigenstates like $\langle x|x'\rangle = \delta(x-x')$ are not normalisable.

where ϕ is the column vector of the basis set and \mathbf{c} is the column vector of coefficients.

Now, let's do a basis transformation. We define a new basis set $|\theta_j\rangle = \sum_i |\phi_i\rangle U_{ij}$ for some set of coefficients $U_{ij} = \langle \phi_i | \theta_j \rangle$, which can be written in the matrix form as

$$(|\theta_1\rangle \quad |\theta_2\rangle \quad \dots) = (|\phi_1\rangle \quad |\phi_2\rangle \quad \dots) \begin{pmatrix} U_{11} & U_{12} & \dots \\ U_{21} & U_{22} & \dots \\ \vdots & \vdots & \ddots \end{pmatrix}, \quad (1.62)$$

or

$$\boldsymbol{\theta}^T = \boldsymbol{\phi}^T \mathbf{U}. \quad (1.63)$$

If we want to keep the new basis orthonormal, then we must have

$$\begin{aligned} \delta_{ij} &= \langle \theta_i | \theta_j \rangle \\ &= \sum_{k,l} U_{ki}^* U_{lj} \langle \phi_k | \phi_l \rangle \\ &= \sum_k U_{ki}^* U_{kj}, \end{aligned} \quad (1.64)$$

which implies

$$\mathbf{U}^\dagger \mathbf{U} = \mathbf{I}, \quad (1.65)$$

and so \mathbf{U} is a unitary matrix. The transformation between orthonormal bases is given by a unitary matrix.

We hope to represent the same $|\psi\rangle$ in this new transformed basis $\{|\theta_i\rangle\}$

$$|\psi\rangle = \boldsymbol{\theta}^T \mathbf{d}. \quad (1.66)$$

Then clearly, we should have

$$|\psi\rangle = \boldsymbol{\theta}^T \mathbf{d} = \boldsymbol{\phi}^T \mathbf{U} \mathbf{d} = \boldsymbol{\phi}^T \mathbf{c}, \quad (1.67)$$

so

$$\mathbf{c} = \mathbf{U} \mathbf{d} \quad \text{or} \quad \mathbf{d} = \mathbf{U}^\dagger \mathbf{c}. \quad (1.68)$$

The components transform in the opposite way as the basis.

Just as for the position and momentum transformation, by inserting the resolution of identity we can write

$$c_i = \langle \phi | \psi \rangle = \langle \phi_i | \sum_j |\theta_j\rangle \langle \theta_j | \psi \rangle = \sum_j \langle \phi_i | \theta_j \rangle d_j, \quad (1.69)$$

and so

$$U_{ij} = \langle \phi_i | \theta_j \rangle. \quad (1.70)$$

The same thing goes for operators. If a Hermitian operator \hat{A} is represented in the $|\phi_i\rangle$ basis with matrix elements $A_{ij} = \langle \phi_i | \hat{A} | \phi_j \rangle$, then in the new basis $|\theta_i\rangle$, the matrix element A' will be given by

$$\begin{aligned} A'_{ij} &= \langle \theta_i | \hat{A} | \theta_j \rangle \\ &= \sum_{k,l} U_{ki}^* U_{lj} \langle \phi_k | \hat{A} | \phi_l \rangle \\ &= \sum_{k,l} U_{ki}^* A_{kl} U_{lj}, \end{aligned} \quad (1.71)$$

and so

$$\mathbf{A}' = \mathbf{U}^\dagger \mathbf{A} \mathbf{U}. \quad (1.72)$$

1.5.1 Symmetry Orbitals

The above transformation is exactly what we are doing when we transform from the atomic orbital basis (denoted $|\phi_i\rangle$) to symmetry orbitals (denoted θ_i) in Hückel theory. If we have figured out the symmetry orbitals to be $|\theta_j\rangle = \sum_i |\phi_i\rangle U_{ij}$ and the Hamiltonian in the atomic orbital basis to be $H_{ij} = \langle \phi_i | \hat{H} | \phi_j \rangle$, then the Hamiltonian in the symmetry orbital basis set is

$$H' = U^\dagger H U. \quad (1.73)$$

In practise we don't usually do this by matrix multiplication because we already know that a lot of the matrix elements will be zero, so we would rather work out H'_{ij} element by element. The goal of Hückel theory is usually to work out the Hückel molecular orbitals as well as their energies, which are the eigenvectors and eigenvalues of H . By doing a unitary basis transformation, the eigenvalues of H' should be the same as H , and the eigenvectors should be the same ones but represented in the symmetry orbital basis. The point is that the block diagonal form of H' makes them easier to find.

1.6 Uncertainty Principle

We have asserted that the uncertainties of two observables are related to the expectation value of their commutator. It is time to see where the uncertainty principle comes from.

In statistics, the variance of a random variable A is given by

$$\sigma_A^2 = \langle A^2 \rangle - \langle A \rangle^2 = \langle (A - \langle A \rangle)^2 \rangle. \quad (1.74)$$

This can be translated into the quantum mechanical expression of the uncertainty of a physical observable

$$\Delta A^2 = \left\langle \psi \left| \left(\hat{A} - \langle \hat{A} \rangle \right)^2 \right| \psi \right\rangle. \quad (1.75)$$

If \hat{A} is a Hermitian operator, then we can write this as

$$\Delta A^2 = \left\langle \left(\hat{A} - \langle \hat{A} \rangle \right) \psi \left| \left(\hat{A} - \langle \hat{A} \rangle \right) \psi \right\rangle. \quad (1.76)$$

If we denote $\psi_A = \left(\hat{A} - \langle \hat{A} \rangle \right) \psi$, then ΔA^2 is exactly the norm of ψ_A :

$$\Delta A^2 = \langle \psi_A | \psi_A \rangle. \quad (1.77)$$

Suppose we have another observable with operator \hat{B} , then if $\psi_B = \left(\hat{B} - \langle \hat{B} \rangle \right) \psi$, we have

$$\Delta B^2 = \langle \psi_B | \psi_B \rangle \quad (1.78)$$

by exactly the same argument. The product of the two variances is

$$\Delta A^2 \Delta B^2 = \langle \psi_A | \psi_A \rangle \langle \psi_B | \psi_B \rangle. \quad (1.79)$$

Next, we will use Cauchy–Schwarz inequality.

Lemma 1.3 (Cauchy–Schwarz inequality). Let V be a vector space with an inner product \cdot , then for all $\mathbf{u}, \mathbf{v} \in V$,

$$|\mathbf{u} \cdot \mathbf{v}|^2 \leq (\mathbf{u} \cdot \mathbf{u})(\mathbf{v} \cdot \mathbf{v}). \quad (1.80)$$

Therefore, we have

$$\Delta A^2 \Delta B^2 \geq |\langle \psi_A | \psi_B \rangle|^2. \quad (1.81)$$

For any complex number $z \in \mathbb{C}$, we have

$$|z|^2 = \text{Re}(z)^2 + \text{Im}(z)^2 = \left(\frac{z + z^*}{2} \right)^2 + \left(\frac{z - z^*}{2i} \right)^2, \quad (1.82)$$

and since $\langle \psi_A | \psi_B \rangle = \langle \psi_B | \psi_A \rangle^*$, we have

$$\Delta A^2 \Delta B^2 \geq \left(\frac{\langle \psi_A | \psi_B \rangle + \langle \psi_B | \psi_A \rangle}{2} \right)^2 + \left(\frac{\langle \psi_A | \psi_B \rangle - \langle \psi_B | \psi_A \rangle}{2i} \right)^2. \quad (1.83)$$

To simplify things up, we need to work out $\langle \psi_A | \psi_B \rangle$ and $\langle \psi_B | \psi_A \rangle$.

$$\begin{aligned} \langle \psi_A | \psi_B \rangle &= \left\langle \left(\hat{A} - \langle \hat{A} \rangle \right) \psi \middle| \left(\hat{B} - \langle \hat{B} \rangle \right) \psi \right\rangle \\ &= \left\langle \psi \middle| \left(\hat{A} - \langle \hat{A} \rangle \right) \left(\hat{B} - \langle \hat{B} \rangle \right) \middle| \psi \right\rangle \\ &= \left\langle \psi \middle| \hat{A} \hat{B} - \hat{A} \langle \hat{B} \rangle - \hat{B} \langle \hat{A} \rangle + \langle \hat{A} \rangle \langle \hat{B} \rangle \middle| \psi \right\rangle \\ &= \langle \hat{A} \hat{B} \rangle - \langle \hat{A} \rangle \langle \hat{B} \rangle. \end{aligned} \quad (1.84)$$

Similarly

$$\langle \psi_B | \psi_A \rangle = \langle \hat{B} \hat{A} \rangle - \langle \hat{A} \rangle \langle \hat{B} \rangle. \quad (1.85)$$

Defining the *anti-commutator*

$$\{\hat{A}, \hat{B}\} := \hat{A} \hat{B} + \hat{B} \hat{A}, \quad (1.86)$$

we have

$$\langle \psi_A | \psi_B \rangle + \langle \psi_B | \psi_A \rangle = \langle \{\hat{A}, \hat{B}\} \rangle - 2 \langle \hat{A} \rangle \langle \hat{B} \rangle, \quad (1.87)$$

$$\langle \psi_A | \psi_B \rangle - \langle \psi_B | \psi_A \rangle = \langle [\hat{A}, \hat{B}] \rangle. \quad (1.88)$$

This gives what is known as the Robertson–Schrödinger uncertainty principle.

Theorem 1.4 (Robertson–Schrödinger uncertainty principle). For two observables A and B ,

$$\Delta A \Delta B \geq \sqrt{\left(\frac{1}{2} \langle \{\hat{A}, \hat{B}\} \rangle - \langle \hat{A} \rangle \langle \hat{B} \rangle \right)^2 + \left(\frac{1}{2i} \langle [\hat{A}, \hat{B}] \rangle \right)^2}. \quad (1.89)$$

This version of the uncertainty principle is very general, but it simplifies in the special case of uncorrelated operators. The first parenthesised quantity in (1.89) is

$$\text{cov}(\hat{A}, \hat{B}) := \frac{1}{2} \langle \hat{A} \hat{B} + \hat{B} \hat{A} \rangle - \langle \hat{A} \rangle \langle \hat{B} \rangle, \quad (1.90)$$

which is known as the *covariance* of the operators. The corresponding quantity in statistics is

$$\text{cov}(X, Y) := \langle XY \rangle - \langle X \rangle \langle Y \rangle, \quad (1.91)$$

but in quantum mechanics we have to be a little bit more careful because in general $\langle \hat{A} \hat{B} \rangle \neq \langle \hat{B} \hat{A} \rangle$. The covariance is a measure of the correlation between operators, and is 0 if the operators are uncorrelated. If this is the case, then (1.89) reduces to the more familiar Robertson uncertainty principle.

Theorem 1.5 (Robertson Uncertainty Principle). If A and B are two uncorrelated physical observables, then

$$\Delta A \Delta B \geq \frac{1}{2} \left| \langle [\hat{A}, \hat{B}] \rangle \right|. \quad (1.92)$$

Note that if the two operators are correlated, then the Robertson–Schrödinger lower bound is larger than the Robertson lower bound, in other words the Robertson–Schrödinger provides a stronger condition.

2 Normal Modes

In Part IB Chemistry A, we discussed the use of symmetry and group theory to make qualitative conclusions about normal modes.

Example. For example, consider the C – H stretching modes in benzene, with point group D_{6h} .

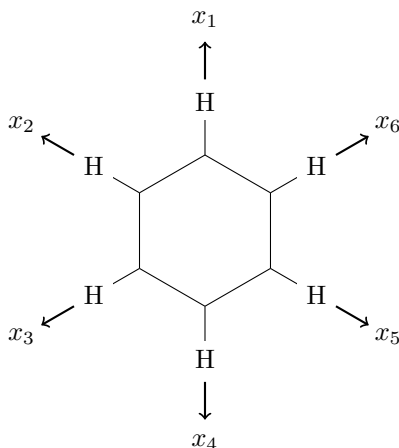


Figure 1: The basis set for representing the C – H stretches in benzene.

Reducing the representation formed by these basis vectors, we get

$$\Gamma^{C-H} = A_{1g} \oplus E_{2g} \oplus B_{1u} \oplus E_{1u}. \quad (2.1)$$

Since there is only one combination for each irreducible representation, it should be easy to determine the symmetry adapted linear combinations of displacements along the C – H bonds that transform according to the symmetries. Based on the Cartesian functions listed in the character table, intuition, or the projection operators, we can determine the normal modes to be

$$Q_{A_{1g}} = \frac{1}{\sqrt{6}}(x_1 + x_2 + x_3 + x_4 + x_5 + x_6) \quad (2.2)$$

$$Q_{E_{2g},1} = \frac{1}{2}(x_2 - x_3 + x_5 - x_6) \quad (2.3)$$

$$Q_{E_{2g},2} = \frac{1}{\sqrt{12}}(2x_1 - x_2 - x_3 + 2x_4 - x_5 - x_6) \quad (2.4)$$

$$Q_{E_{1u},1} = \frac{1}{\sqrt{12}}(2x_1 + x_2 - x_3 - 2x_4 - x_5 + x_6) \quad (2.5)$$

$$Q_{E_{1u},2} = \frac{1}{2}(x_2 + x_3 - x_5 - x_6) \quad (2.6)$$

$$Q_{B_{1u}} = \frac{1}{\sqrt{6}}(x_1 - x_2 + x_3 - x_4 + x_5 - x_6). \quad (2.7)$$

This method is nice and easy, especially for simple cases like this with restricted basis sets. But it does leave unanswered questions. What if we have more than one symmetry adapted combination transforming as a particular IR? These symmetry adapted combinations of displacement vectors are like symmetry orbitals, and some process analogous to combining symmetry orbitals to form molecular orbitals should be indicated. Most of the combinations we have here are only approximations (albeit useful ones) to normal modes, as we can see a lot of them would move the centre of mass. Also, it is impossible that only H atoms are moving during the vibrations — we also need the motion of C atoms as well as displacements in other directions. In such a complete basis set, how to correctly combine

the symmetry adapted displacement vectors to form the true normal modes would be a problem. Additionally, some indication of the frequencies would be useful.

To do this, we have to treat the mechanical problem of molecular vibration seriously.

2.1 Diatomic Vibration

We begin by looking at the vibration of a diatomic molecule, and hopefully this can give us some insight on treating the general problem. Each atom in a diatomic can move in three independent directions (x , y and z), so there are in total six possible independent motions. We know that three of them are translations and rotations, leaving us with only one vibrational mode. It turns out that only the motion along the internuclear axis is useful for vibration. To label the positions of the atoms relative to their equilibrium positions, we define two basis vectors \mathbf{x}_1 and \mathbf{x}_2 . Then the configuration of the atom can be labelled by a vector

$$\mathbf{v} = x_1\mathbf{x}_1 + x_2\mathbf{x}_2, \quad (2.8)$$

where x_1 and x_2 are the components, meaning that atom 1 has displaced by length x_1 along the direction \mathbf{x}_1 , and atom 2 has displaced by x_2 along \mathbf{x}_2 . We can put the components into a single column vector $\mathbf{x} = \begin{pmatrix} x_1 \\ x_2 \end{pmatrix}$.

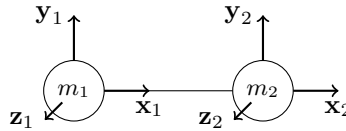


Figure 2: A basis set representing the motion of a diatomic molecule.

The simplest model for the internuclear potential is the harmonic oscillator, with the potential given by

$$V_{\text{HO}} = \frac{1}{2}k(x_1 - x_2)^2, \quad (2.9)$$

where k is the force constant. We can write the Hamiltonian as¹¹

$$\hat{H}(\mathbf{x}) = -\frac{\hbar^2}{2m_1}\frac{\partial^2}{\partial x_1^2} - \frac{\hbar^2}{2m_2}\frac{\partial^2}{\partial x_2^2} + \frac{1}{2}k(x_1 - x_2)^2, \quad (2.10)$$

where the first two terms are the kinetic energies of the two particles with masses of m_1 and m_2 respectively. Now this is a non-separable Hamiltonian — we have cross terms x_1x_2 in the potential energy, so that we can not write the Hamiltonian as $\hat{H}(x_1) + \hat{H}(x_2)$. We would not like to solve this kind of partial differential equations directly.

To make progress, we want to try some coordinate transformation that would make the cross terms vanishes. The transformation

$$\begin{aligned} w_1 &= \frac{1}{\sqrt{2}}(x_1 - x_2) \\ w_2 &= \frac{1}{\sqrt{2}}(x_1 + x_2) \end{aligned} \quad (2.11)$$

of the components seems promising, since it eliminates the cross terms in the potential energy by making it into a single term kw_1^2 . Moreover, these two coordinates have solid physical meaning — w_1 is the compression/extension along the internuclear axis, which is exactly the vibrational motion

¹¹We treat this problem using Hamiltonian. If you do Part IB Mathematics, you will solve this type of problems using Lagrangian — you will find that these two methods are closely related.

we are trying to model, and w_2 describes the translation along the internuclear axis, one of the other motion of the molecules. This is described by the unitary transformation matrix

$$\mathbf{U} = \frac{1}{\sqrt{2}} \begin{pmatrix} 1 & -1 \\ 1 & 1 \end{pmatrix} \quad (2.12)$$

generated by putting the components of the transformed components in terms of the original components in rows,¹² so that the components transforms as desired:

$$\begin{pmatrix} w_1 \\ w_2 \end{pmatrix} = \mathbf{U} \begin{pmatrix} x_1 \\ x_2 \end{pmatrix} = \frac{1}{\sqrt{2}} \begin{pmatrix} 1 & -1 \\ 1 & 1 \end{pmatrix} \begin{pmatrix} x_1 \\ x_2 \end{pmatrix}, \quad (2.13)$$

or

$$\mathbf{w} = \mathbf{U}\mathbf{x}. \quad (2.14)$$

We can be satisfied that no physical properties will be altered by this unitary transformation.

However, we will see that the problem of this basis transformation is that although the potential terms is nicely simplified, the kinetic term is screwed up. From chain rule, we have

$$\begin{aligned} \frac{\partial}{\partial x_1} &= \frac{\partial w_1}{\partial x_1} + \frac{\partial}{\partial w_1} + \frac{\partial w_2}{\partial x_1} \frac{\partial}{\partial w_2} \\ &= \frac{1}{\sqrt{2}} \left(\frac{\partial}{\partial w_1} - \frac{\partial}{\partial w_2} \right), \end{aligned} \quad (2.15)$$

$$\begin{aligned} \frac{\partial}{\partial x_2} &= \frac{\partial w_1}{\partial x_2} + \frac{\partial}{\partial w_1} + \frac{\partial w_2}{\partial x_2} \frac{\partial}{\partial w_2} \\ &= \frac{1}{\sqrt{2}} \left(\frac{\partial}{\partial w_1} + \frac{\partial}{\partial w_2} \right). \end{aligned} \quad (2.16)$$

Therefore in this transformed basis, the Hamiltonian is

$$\hat{H}(\mathbf{w}) = -\frac{\hbar^2}{4m_1} \left(\frac{\partial}{\partial w_1} - \frac{\partial}{\partial w_2} \right)^2 - \frac{\hbar^2}{4m_2} \left(\frac{\partial}{\partial w_1} + \frac{\partial}{\partial w_2} \right)^2 + kw_1^2. \quad (2.17)$$

The kinetic part contains a mixed partial derivative

$$\frac{\hbar^2}{2m_1} \left(\frac{1}{m_1} - \frac{1}{m_2} \right) \frac{\partial^2}{\partial w_1 \partial w_2}, \quad (2.18)$$

which is non-vanishing for $m_1 \neq m_2$.

It seems that by doing a unitary transformation, we can only separate one of the kinetic and the potential part. Whenever we separate one of the terms, the other will necessarily be screwed up.

However, the above trial did provide us some inspiration. The mixed partial derivative term do vanish if the masses of the two particles are equal. What if we do some scaling to the coordinates based on the masses? In turns out that for this particular question, the useful scaling is

$$Q_1 = \sqrt{\frac{m_1 m_2}{m_1 + m_2}} (x_1 - x_2) \quad (2.19)$$

$$Q_2 = \frac{1}{\sqrt{m_1 + m_2}} (m_1 x_1 + m_2 x_2) \quad (2.20)$$

We will explain how we obtained this later, but now let's try this out. Again, the potential is straightforward to rewrite in the \mathbf{Q} basis, as

$$V_{\text{HO}} = \frac{k(m_1 + m_2)}{2m_1 m_2} Q_1^2. \quad (2.21)$$

¹²Then the basis vector would transform as \mathbf{U}^\dagger . Remember that the basis vectors and the components transform in the inverse way.

For the kinetic energies, we again need the chain rule, and we obtain

$$\frac{\partial}{\partial x_1} = \frac{1}{\sqrt{m_1 + m_2}} \left(\sqrt{m_1 m_2} \frac{\partial}{\partial Q_1} + m_1 \frac{\partial}{\partial Q_2} \right), \quad (2.22)$$

$$\frac{\partial}{\partial x_2} = \frac{1}{\sqrt{m_1 + m_2}} \left(-\sqrt{m_1 m_2} \frac{\partial}{\partial Q_1} + m_2 \frac{\partial}{\partial Q_2} \right). \quad (2.23)$$

If we substitute this into the expression of the Hamiltonian, we will see that all the mixed derivatives magically cancels out and we get

$$\hat{H}(Q) = -\frac{\hbar^2}{2} \frac{\partial^2}{\partial Q_1^2} + \frac{k(m_1 + m_2)}{2m_1 m_2} Q_1^2 - \frac{\hbar^2}{2} \frac{\partial^2}{\partial Q_2^2}. \quad (2.24)$$

This can be nicely separated into

$$\hat{H}(\mathbf{Q}) = \hat{H}(Q_1) + \hat{H}(Q_2), \quad (2.25)$$

where

$$\hat{H}(Q_1) = -\frac{\hbar^2}{2} \frac{\partial^2}{\partial Q_1^2} + \frac{k(m_1 + m_2)}{2m_1 m_2} Q_1^2, \quad (2.26)$$

$$\hat{H}(Q_2) = -\frac{\hbar^2}{2} \frac{\partial^2}{\partial Q_2^2}. \quad (2.27)$$

If we compare (2.26) with the Hamiltonian of a canonical harmonic oscillator, we see that this is an harmonic oscillator with a mass 1 and a modified force constant of

$$k' = \frac{k(m_1 + m_2)}{m_1 m_2} =: \frac{k}{\mu}, \quad (2.28)$$

where we defined the familiar reduced mass $\mu := m_1 m_2 / (m_1 + m_2)$. This gives an angular frequency

$$\omega = \sqrt{\frac{k'}{1}} = \sqrt{\frac{k}{\mu}}. \quad (2.29)$$

This is a familiar result. The vibration of a diatomic molecule is the same as the vibration of a single molecule with reduced mass of the system. The second equation (2.27) has only a kinetic term. This is the translation of the whole molecule along the internuclear axis — it can also be thought of as a harmonic oscillator with zero force constant, and so a zero frequency.

2.1.1 Mass-Weighted Coordinates

Now let's consider what is going on. The transformation we proposed can be expressed by the matrix

$$\mathbf{A} = \frac{1}{\sqrt{m_1 + m_2}} \begin{pmatrix} \sqrt{m_1 m_2} & -\sqrt{m_1 m_2} \\ m_1 & m_2 \end{pmatrix}. \quad (2.30)$$

You may verify that this is not a unitary matrix — the easiest way to see this is by checking its determinant. Therefore, physical observable do not have to be preserved by this transformation. For example, we can see that all masses has reduced to 1, and the actual masses has somehow been taken into the modified force constant. However, the observable we really care about, the (angular) frequency of the oscillator ω , is related to the ratio of these, and this is unchanged by our transformation.

To see what it really going on in our transformation, we can split our transformation into two phases. The first phase is to mass-weight the coordinates, defining $q_i = \sqrt{m_i} x_i$, which is a diagonal transform, but is not unitary. This is represented by the diagonal matrix

$$\mathbf{B} = \begin{pmatrix} \sqrt{m_1} & 0 \\ 0 & \sqrt{m_2} \end{pmatrix}. \quad (2.31)$$

The Hamiltonian in this mass-weighted basis set is

$$\hat{H}(\mathbf{q}) = -\frac{\hbar^2}{2} \left(\frac{\partial^2}{\partial q_1^2} + \frac{\partial^2}{\partial q_2^2} \right) + \frac{k}{2} \left(\frac{q_1}{\sqrt{m_1}} - \frac{q_2}{\sqrt{m_2}} \right)^2. \quad (2.32)$$

This transformation does not remove the cross term in the potential, but it has reduced all the kinetic terms to a nice and simple form where all the masses are unity. This is exactly the case that we wanted before — if the masses are equal, then when you do the unitary transformation to simplify the potential term, all the cross terms in the kinetic term after the transformation will automatically cancel out! We are then ready to do our second transformation defined by

$$Q_1 = \frac{1}{\sqrt{m_1 + m_2}} (\sqrt{m_2} q_1 - \sqrt{m_1} q_2), \quad (2.33)$$

$$Q_2 = \frac{1}{\sqrt{m_1 + m_2}} (\sqrt{m_1} q_1 + \sqrt{m_2} q_2), \quad (2.34)$$

which can be represented by the unitary matrix

$$\mathbf{C} = \frac{1}{\sqrt{m_1 + m_2}} \begin{pmatrix} \sqrt{m_2} & -\sqrt{m_1} \\ \sqrt{m_1} & \sqrt{m_2} \end{pmatrix}. \quad (2.35)$$

The combination of these two transformations is the one we claimed before:

$$\mathbf{A} = \mathbf{C}\mathbf{B}. \quad (2.36)$$

2.2 Quadratic Form

Before investigating more complicated cases, you may find it extremely useful to formulate the above process in matrices. Hence, we may rewrite the Hamiltonian in the following matrix form

$$\begin{aligned} \hat{H}(\mathbf{x}) &= \frac{\hbar^2}{2} \begin{pmatrix} \frac{\partial}{\partial x_1} & \frac{\partial}{\partial x_2} \end{pmatrix} \begin{pmatrix} \frac{1}{m_1} & 0 \\ 0 & \frac{1}{m_2} \end{pmatrix} \begin{pmatrix} \frac{\partial}{\partial x_1} \\ \frac{\partial}{\partial x_2} \end{pmatrix} + \frac{1}{2} \begin{pmatrix} x_1 & x_2 \end{pmatrix} \begin{pmatrix} k & -k \\ -k & k \end{pmatrix} \begin{pmatrix} x_1 \\ x_2 \end{pmatrix} \\ &=: -\frac{\hbar^2}{2} \dot{\mathbf{x}}^T \mathbf{T}_x \dot{\mathbf{x}} + \frac{1}{2} \mathbf{x}^T \mathbf{H}_x \mathbf{x}. \end{aligned} \quad (2.37)$$

The subscripts x here stands for the x basis, and \mathbf{H} is confusingly the canonical notation for the Hessian matrix which unfortunately happens to share its notation with the Hamiltonian.

Now we can do the transformations. The first step is to mass-scale the coordinates by

$$\mathbf{q} = \mathbf{B}\mathbf{x}, \quad (2.38)$$

where

$$\mathbf{B} = \begin{pmatrix} \sqrt{m_1} & 0 \\ 0 & \sqrt{m_2} \end{pmatrix}. \quad (2.39)$$

Note that by chain rule, the derivatives transform in the opposite way as the coordinates, so

$$\dot{\mathbf{q}} = \mathbf{B}^{-1} \dot{\mathbf{x}}, \quad (2.40)$$

and so

$$\dot{\mathbf{x}}^T \mathbf{T}_x \dot{\mathbf{x}} = \dot{\mathbf{q}}^T \mathbf{B}^T \mathbf{T}_x \mathbf{B} \dot{\mathbf{q}} =: \dot{\mathbf{q}}^T \mathbf{T}_q \dot{\mathbf{q}}. \quad (2.41)$$

This transformation is designed to reduce the \mathbf{T} matrix to the identity matrix in this basis:

$$\mathbf{T}_q := \mathbf{B}^T \mathbf{T}_x \mathbf{B} = \mathbf{I}, \quad (2.42)$$

and so

$$\dot{\mathbf{x}}^T \mathbf{T}_x \dot{\mathbf{x}} = \dot{\mathbf{q}}^T \mathbf{I} \dot{\mathbf{q}} = \dot{\mathbf{q}}^T \dot{\mathbf{q}}. \quad (2.43)$$

Similarly for the Hessian matrix,

$$\mathbf{x}^T \mathbf{H}_x \mathbf{x} = \mathbf{q}^T (\mathbf{B}^{-1})^T \mathbf{H}_x \mathbf{B}^{-1} \mathbf{q} =: \mathbf{q}^T \mathbf{K} \mathbf{q}, \quad (2.44)$$

where the Hessian matrix in this mass-scaled coordinate is called the *dynamical matrix* and is given the special notation \mathbf{K} . We can calculate this as

$$\mathbf{K} := (\mathbf{B}^{-1})^T \mathbf{H}_x \mathbf{B}^{-1} = \begin{pmatrix} \frac{k}{m_1} & -\frac{k}{\sqrt{m_1 m_2}} \\ -\frac{k}{\sqrt{m_1 m_2}} & \frac{k}{m_2} \end{pmatrix}. \quad (2.45)$$

Similarly, we then do the unitary transformation that diagonalises \mathbf{K} as well, and we finally get

$$\mathbf{T}_Q = \mathbf{I}, \quad \mathbf{H}_Q = \begin{pmatrix} 0 & 0 \\ 0 & \frac{k}{\mu} \end{pmatrix} \quad (2.46)$$

in the \mathbf{Q} basis. Since both matrices are diagonal, we are then able to separate the Hamiltonian.

2.3 The General Case

Now we are ready to generalise the above method for all molecules. Consider a molecule consisting of M atoms. The position of each atom is described by a displacement vector $\mathbf{x}_i = (x_i, y_i, z_i)$ ($1 \leq i \leq N$) from its equilibrium position. We can compactly string them together as a $3N$ -dimensional vector

$$\mathbf{x} = (x_1, y_1, z_1, \dots, x_N, y_N, z_N). \quad (2.47)$$

A general Hamiltonian can be written as

$$\hat{H} = -\frac{\hbar^2}{2} \sum_i \frac{1}{m_i} \frac{\partial^2}{\partial x_i^2} + \frac{1}{2} \sum_i \sum_{j>i}^{3N} k_{ij} (x_i - x_j)^2. \quad (2.48)$$

Note that we sum over $i > j$ in the potential term to avoid double counting, and we allow a potential term to rise between any two coordinates, although in practise we can set most of them to be zero. We can write this into the matrix form introduced above

$$\hat{H} = -\frac{\hbar^2}{2} \dot{\mathbf{x}}^T \mathbf{T}_x \dot{\mathbf{x}} + \frac{1}{2} \mathbf{x}^T \mathbf{H}_x \mathbf{x}, \quad (2.49)$$

where \mathbf{T} is a diagonal matrix whose elements are the reciprocal masses

$$T_{x,ii} = \frac{1}{m_i}, \quad (2.50)$$

and the Hessian matrix elements are

$$H_{x,ij} = \frac{\partial^2 V}{\partial x_i \partial x_j} = \begin{cases} \sum_n k_{in} & \text{if } i = j \\ -k_{ij} & \text{if } i \neq j \end{cases}. \quad (2.51)$$

We first scale the Hamiltonian into the mass-weighted basis so that

$$\hat{H}(\mathbf{q}) = -\frac{\hbar^2}{2} \dot{\mathbf{q}}^T \dot{\mathbf{q}} + \frac{1}{2} \mathbf{q}^T \mathbf{K} \mathbf{q}. \quad (2.52)$$

The matrix \mathbf{T}_q in the mass-weighted basis is the identity matrix so we have omitted it, and the dynamical matrix is simply

$$K_{ij} = \frac{1}{\sqrt{m_i m_j}} H_{x,ij}. \quad (2.53)$$

We now need to transform the basis such that the \mathbf{K} is made diagonal. We can always do this because \mathbf{K} is a real symmetric matrix, which is always diagonalisable by a orthogonal matrix (a real unitary matrix). This transformation is guaranteed to keep \mathbf{T} matrix still diagonalised because the identity matrix is always the identity matrix in any basis. This is done by the orthogonal matrix \mathbf{C} such that

$$\mathbf{H}_Q = \mathbf{C}^\dagger \mathbf{K} \mathbf{C}, \quad (2.54)$$

where each column of \mathbf{C} is the normalised eigenvector of \mathbf{K} , and \mathbf{H}_Q is a diagonal matrix whose elements are eigenvalues $\{\lambda_i\}$ of \mathbf{K} in corresponding order. Finally in this basis, we have

$$\begin{aligned} \hat{H}(\mathbf{Q}) &= -\frac{\hbar^2}{2} \dot{\mathbf{Q}}^\mathbf{T} \dot{\mathbf{Q}} + \frac{1}{2} \mathbf{Q}^\mathbf{T} \mathbf{H}_Q \mathbf{Q} \\ &= \sum_i^{3N} \left(-\frac{\hbar^2}{2} \frac{\partial^2}{\partial Q_i^2} + \frac{1}{2} \lambda_i Q_i^2 \right) \end{aligned} \quad (2.55)$$

separated into $3N$ one-dimensional harmonic oscillator with modified force constants $\{\lambda_i\}$. The angular frequencies of the modes are therefore

$$\omega_i = \sqrt{\lambda_i}. \quad (2.56)$$

The eigenvectors of \mathbf{K} are the normal modes in terms of the \mathbf{q} coordinates, which can be transformed into the \mathbf{x} coordinates. Note that although the normal modes are orthogonal in the \mathbf{q} coordinates, they are not necessarily so in the \mathbf{x} coordinates since the mass scaling is not unitary. However, we may still often find them to be orthogonal, as they must be so if they transform according to different irreducible representations.

Note that since we used all $3N$ displacement vectors, there will be 6 (or 5) normal modes corresponding to translations or rotations. If we use a restricted basis set, there may be less of them. They can be easily identified as modes with frequencies zero.

2.4 Example: AAA Linear Molecule

As an simple example, let's look at the AAA linear molecule of three identical atoms connected by two bonds of equal strengths. An example of this would be the azide anion. For simplicity, we only consider the motions along the internuclear axis.

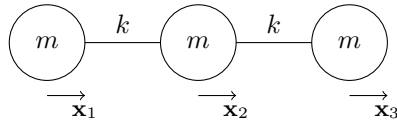


Figure 3: An AAA molecule.

The potential can be modelled as

$$V = \frac{k}{2}(x_1 - x_2)^2 + \frac{k}{2}(x_2 - x_3)^2. \quad (2.57)$$

The Hessian matrix can be found by either taking the partial derivative or by expanding the above function.

$$\mathbf{H}_x = \begin{pmatrix} k & -k & 0 \\ -k & 2k & -k \\ 0 & -k & k \end{pmatrix}. \quad (2.58)$$

The dynamical matrix is

$$\mathbf{K} = \begin{pmatrix} \frac{k}{m} & -\frac{k}{m} & 0 \\ -\frac{k}{m} & \frac{2k}{m} & -\frac{k}{m} \\ 0 & -\frac{k}{m} & \frac{k}{m} \end{pmatrix}. \quad (2.59)$$

We then simply need to diagonalise this matrix. The eigenvalues and the corresponding normal mode frequencies are

$$\lambda_1 = 0 \qquad \lambda_2 = \frac{k}{m} \qquad \lambda_3 = \frac{3k}{m} \quad (2.60)$$

$$\omega_1 = 0 \qquad \omega_2 = \sqrt{\frac{k}{m}} \qquad \omega_3 = \sqrt{\frac{3k}{m}}. \quad (2.61)$$

We can also find the normalised eigenvectors, which give the normal mode coordinates in the \mathbf{q} basis (which is also the normal coordinates in the \mathbf{x} basis since all atoms have equal masses)

$$Q_1 = \frac{1}{\sqrt{3}} \begin{pmatrix} 1 \\ 1 \\ 1 \end{pmatrix} \cdot \mathbf{q} \qquad Q_2 = \frac{1}{\sqrt{2}} \begin{pmatrix} -1 \\ 0 \\ 1 \end{pmatrix} \cdot \mathbf{q} \qquad Q_3 = \frac{1}{\sqrt{6}} \begin{pmatrix} 1 \\ -2 \\ 1 \end{pmatrix} \cdot \mathbf{q}. \quad (2.62)$$

In mode 1, all atoms move in the same direction with the same magnitude. This is the translation. Mode 2 is the symmetric stretch, in which both bonds are simultaneously stretched/compressed, and mode 3 is the antisymmetric stretch, in which when one bond is stretched, the other is compressed.



Figure 4: Normal modes of an AAA molecule.

2.5 The Use of Symmetry

The dynamical matrix in the above example is 3×3 , which we can easily diagonalise directly. For larger molecules, we will encounter the same issue that we had when constructing MOs: any bigger matrix will be too tedious to diagonalise directly by hand. The solution is the same too: we can use an intermediate symmetry-adapted basis set to transform the dynamical matrix to a block diagonal form.

We will summarise the process of constructing normal modes by symmetry as well as molecular orbitals in the table below.

2.6 Example: ABBBA Linear Molecule

Consider the motions along the bond for a centrosymmetric ABBBA linear molecule, for example acetylene. The masses of the two types of atoms are now different, and we also allow the strength of the A – B bond to be different from the B – B bond.

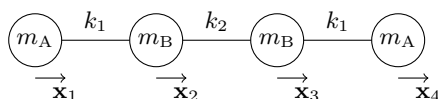


Figure 5: A centrosymmetric ABBBA linear molecule.

MOs	Normal modes
1. Write down the Hamiltonian matrix in the AO basis.	1. Write down the dynamical matrix in the mass-weighted basis.
2. Determine the IRs spanned by the AO basis set.	2. Determine IRs spanned by the mass-weighted basis set.
3. Construct symmetry adapted combinations of the AOs.	3. Construct symmetry adapted combinations of the mass-weighted coordinates.
4. Transform the Hamiltonian matrix into the symmetry-adapted basis set.	4. Transform the dynamical matrix into the symmetry-adapted basis set.
5. The eigenvalues are the orbital energies and the eigenvectors are the MO coefficients in the symmetry-adapted basis set.	5. The eigenvalues are the frequencies squared and the eigenvectors are the normal mode coordinates in the symmetry-adapted basis set.
6. The eigenvectors can be transformed back to the AO basis set.	6. The eigenvectors can be transformed back to the coordinate displacement basis set.

If we proceed without using symmetry, then we have a 4×4 dynamical matrix to diagonalise

$$\mathbf{K} = \begin{pmatrix} \frac{k_1}{m_A} & -\frac{k_1}{\sqrt{m_A m_B}} & 0 & 0 \\ -\frac{k_1}{\sqrt{m_A m_B}} & \frac{k_1+k_2}{m_B} & -\frac{k_2}{m_B} & 0 \\ 0 & -\frac{k_2}{m_B} & \frac{k_1+k_2}{m_B} & -\frac{k_1}{\sqrt{m_A m_B}} \\ 0 & 0 & -\frac{k_1}{\sqrt{m_A m_B}} & \frac{k_1}{m_A} \end{pmatrix}. \quad (2.63)$$

To simplify things up, we will create a $\tilde{\mathbf{q}}$ basis from symmetry adapted combinations of \mathbf{q} . The point group of the molecule is $D_{\infty h}$, and we can split the basis into two two-dimensional sets $\{\mathbf{q}_1, \mathbf{q}_4\}$ and $\{\mathbf{q}_2, \mathbf{q}_3\}$. The irreducible representations and the symmetry adapted combinations of basis vectors should be easy to find¹³

$$\tilde{\mathbf{q}}_1 = \frac{1}{\sqrt{2}}(\mathbf{q}_1 - \mathbf{q}_4) \quad \Sigma_g^+ \quad (2.64)$$

$$\tilde{\mathbf{q}}_2 = \frac{1}{\sqrt{2}}(\mathbf{q}_2 - \mathbf{q}_3) \quad \Sigma_g^+ \quad (2.65)$$

$$\tilde{\mathbf{q}}_3 = \frac{1}{\sqrt{2}}(\mathbf{q}_1 + \mathbf{q}_4) \quad \Sigma_u^+ \quad (2.66)$$

$$\tilde{\mathbf{q}}_4 = \frac{1}{\sqrt{2}}(\mathbf{q}_2 + \mathbf{q}_3) \quad \Sigma_u^+. \quad (2.67)$$

This is the basis transformation represented by the unitary matrix

$$\tilde{\mathbf{C}} = \frac{1}{\sqrt{2}} \begin{pmatrix} 1 & 0 & 1 & 0 \\ 0 & 1 & 0 & 1 \\ 0 & -1 & 0 & 1 \\ -1 & 0 & 1 & 0 \end{pmatrix} \quad (2.68)$$

so that the components transforms as

$$\tilde{\mathbf{q}} = \tilde{\mathbf{C}}^\dagger \mathbf{q}. \quad (2.69)$$

We must have

$$\mathbf{q}^\dagger \mathbf{K} \mathbf{q} = \tilde{\mathbf{q}}^\dagger \tilde{\mathbf{K}} \tilde{\mathbf{q}} = \mathbf{q}^\dagger \tilde{\mathbf{C}} \tilde{\mathbf{K}} \tilde{\mathbf{C}}^\dagger \mathbf{q}, \quad (2.70)$$

¹³Recall when constructing a representation, we transform the basis, not the components.

and so¹⁴

$$\begin{aligned}\tilde{K} &= \tilde{C}K\tilde{C} \\ &= \begin{pmatrix} \frac{k_1}{m_A} & -\frac{k_1}{\sqrt{m_A m_B}} & 0 & 0 \\ -\frac{k_1}{\sqrt{m_A m_B}} & \frac{k_1+2k_2}{m_B} & 0 & 0 \\ 0 & 0 & \frac{k_1}{m_A} & -\frac{k_1}{\sqrt{m_A m_B}} \\ 0 & 0 & -\frac{k_1}{\sqrt{m_A m_B}} & \frac{k_1}{m_A} \end{pmatrix}.\end{aligned}\quad (2.71)$$

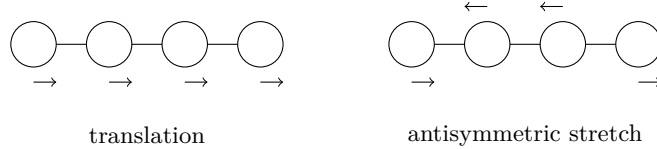
This is block diagonal as expected. For simplicity, we will only work out the Σ_u^+ (bottom right) block. The eigenvalues are

$$\lambda_3 = 0 \qquad \lambda_4 = \frac{k_1(m_A + m_B)}{m_A m_B}, \quad (2.72)$$

and the eigenvectors in the various basis sets are

$$\begin{aligned}Q_3 &= \frac{1}{\sqrt{m_A + m_B}} \begin{pmatrix} \sqrt{m_A} \\ \sqrt{m_B} \end{pmatrix} \cdot \tilde{\mathbf{q}}_{\Sigma_u^+} \\ &= \frac{1}{\sqrt{2(m_A + m_B)}} \begin{pmatrix} \sqrt{m_A} \\ \sqrt{m_B} \\ \sqrt{m_B} \\ \sqrt{m_A} \end{pmatrix} \cdot \mathbf{q} \\ &= \frac{1}{2} \begin{pmatrix} 1 \\ 1 \\ 1 \\ 1 \end{pmatrix} \cdot \mathbf{x} \\ Q_4 &= \frac{1}{\sqrt{m_A + m_B}} \begin{pmatrix} \sqrt{m_B} \\ -\sqrt{m_A} \end{pmatrix} \cdot \tilde{\mathbf{q}}_{\Sigma_u^+} \\ &= \frac{1}{\sqrt{2(m_A + m_B)}} \begin{pmatrix} \sqrt{m_B} \\ -\sqrt{m_A} \\ -\sqrt{m_A} \\ \sqrt{m_B} \end{pmatrix} \cdot \mathbf{q} \\ &= \frac{\sqrt{m_A m_B}}{\sqrt{2(m_A^2 + m_B^2)}} \begin{pmatrix} \sqrt{\frac{m_B}{m_A}} \\ -\sqrt{\frac{m_A}{m_B}} \\ -\sqrt{\frac{m_A}{m_B}} \\ \sqrt{\frac{m_B}{m_A}} \end{pmatrix} \cdot \mathbf{x}.\end{aligned}\quad (2.73)$$

Mode three is the uniform translation, and mode four is the antisymmetric stretch in which when one A – B bond is stretched, the other is compressed.



2.7 Beyond Harmonic Oscillators

Real molecules are not harmonic oscillators in general. We will have some complicated potential as a function of the nuclear coordinates, which defines the nuclear *potential energy surface* (PES). We can perform a Taylor expansion around an equilibrium position to get

$$V(\mathbf{R}) = V(\mathbf{R}_{\text{eq}}) + \sum_{i=1}^{3N} x_i \frac{\partial V}{\partial x_i} \Big|_{\mathbf{R}_{\text{eq}}} + \frac{1}{2} \sum_{i=1}^{3N} \sum_{j=1}^{3N} x_i x_j \frac{\partial^2 V}{\partial x_i \partial x_j} \Big|_{\mathbf{R}_{\text{eq}}} + O(x_i x_j x_k). \quad (2.74)$$

$V(\mathbf{R}_{\text{eq}})$ is the energy at the bottom of the potential well, which we can set to 0, and the first derivatives the the equilibrium positions should also be 0. For sufficiently small oscillations, the cubic terms and above are not significant, so we can truncate out expression at the quadratic term, which is equivalent to making a harmonic approximation. We are left with

$$V(\mathbf{R}) \approx \frac{1}{2} \sum_{i,j=1}^{3N} x_i x_j \frac{\partial^2 V}{\partial x_i \partial x_j} \Big|_{\mathbf{R}_{\text{eq}}} = \mathbf{x}^T \mathbf{H}_x \mathbf{x} \quad (2.75)$$

¹⁴An easier thing to do is to evaluate the matrix component by component, using $\tilde{K}_{ij} = \tilde{\mathbf{q}}_i^\dagger \mathbf{K} \tilde{\mathbf{q}}_j$.

with Hessian matrix elements

$$H_{x,ij} = \left. \frac{\partial^2 V}{\partial x_i \partial x_j} \right|_{\mathbf{R}_{\text{eq}}} . \quad (2.76)$$

For harmonic potentials, this is equivalent to finding the quadratic coefficients of V . If we want to include the effect of cubic terms and above, we can do this by perturbation theory, which is the topic of the rest of this course.

2.8 Transition States

The eigenvalues of the Hessian matrix represents the second derivatives of the potential energies along the principal directions. Recall that a positive second derivative corresponds to a minimum. Ignoring the six directions that do not corresponds to an internal change of the molecule, we can classify a stationary point on the potential energy surface according to the number of negative Hessian eigenvalues. If there are no negative Hessian eigenvalues, then this is a minimum on the PES. If all the eigenvalues are negative, the point is a maximum.

The Murrell–Laidler definition of a *transition state* is an index-1 saddle point, that is a stationary point with only one negative Hessian value. Therefore a transition state is a maximum along one direction and a minimum in all other orthogonal directions. The normal mode corresponding to the negative Hessian eigenvalue is the reaction coordinate at the transition state. Since the vibrational frequency is the square root of the eigenvalue, the frequency in such a direction is imaginary.¹⁵

¹⁵This statement should make you question the derivation of transition state theory presented in the Part IB MELT course. A rigorous treatment of reaction kinetics will be left to Part III.

3 Non-Degenerate Perturbation Theory

3.1 Non-Degenerate Perturbation Theory

Often the Schrödinger equation of a Hamiltonian \hat{H} will be too complicated to be solved directly:

$$\hat{H}\psi_n = E_n\psi_n, \quad (3.1)$$

where $n \in \mathbb{N}$ labelling the quantum states. Then we need to start to make approximations. The starting point of the perturbation theory is to find a simpler Hamiltonian, which we call $\hat{H}^{(0)}$, that somewhat resembles H but whose eigenfunctions $\psi_n^{(0)}$ and eigenvalues $E_n^{(0)}$ are known:

$$\hat{H}^{(0)}\psi_n^{(0)} = E_n^{(0)}\psi_n^{(0)}. \quad (3.2)$$

At this state we will assume that the states are non-degenerate.

The next thing we do to consider the whole one-parameter family of Hamiltonian

$$\hat{H}(\lambda) = \hat{H}^{(0)} + \lambda\hat{H}^{(1)} \quad (3.3)$$

such that

$$\hat{H}(\lambda^*) = \hat{H} \quad (3.4)$$

at some specific value of $\lambda = \lambda^*$. $\hat{H}^{(1)}$ is the *perturbation* to the reference Hamiltonian. Usually we would just let

$$\hat{H}^{(1)} = \hat{H} - \hat{H}^{(0)} \quad (3.5)$$

so that $\hat{H}(\lambda)$ becomes the Hamiltonian \hat{H} of interest at $\lambda = 1$. However, sometimes the perturbation to the reference Hamiltonian has a natural perturbation in it. For example, if we consider applying a small electric field $\mathcal{E}\hat{\mathbf{z}}$ to a molecule with dipole $\boldsymbol{\mu}$, then the strength of the electric field \mathcal{E} will be a natural perturbation parameter.

Then the fundamental assumption in perturbation theory is to assume that both the energies and the quantum states are analytic near $\lambda = 0$, so that if we gradually turn on the perturbation from the reference state where $\lambda = 0$ to some small non-zero λ , the energies and the wavefunctions will also change smoothly given by the power series

$$\psi_n(\lambda) = \psi_n^{(0)} + \lambda\psi_n^{(1)} + \lambda^2\psi_n^{(2)} + \dots \quad (3.6)$$

$$E_n(\lambda) = E_n^{(0)} + \lambda E_n^{(1)} + \lambda^2 E_n^{(2)} + \dots \quad (3.7)$$

In hope that these series do converge to the true values, we can then evaluate these power series expansions (usually the first few correction terms only) at the desired value of λ to get a approximation of the true E_n and ψ_n .

We will employ the conventional *intermediate normalisation*, where the unperturbed wavefunction will be normalised

$$\langle \psi_n^{(0)} | \psi_n^{(0)} \rangle \equiv \langle n | n \rangle = 1. \quad (3.8)$$

This choice means that the perturbed wavefunctions $\psi_n(\lambda)$ will not be normalised. However, since $\psi_n^{(1)}, \psi_n^{(2)}, \dots$ are corrections to $\psi_n^{(0)}$, we may require that

$$\langle n | \psi_n^{(i)} \rangle = 0 \quad (3.9)$$

for all $i > 0$, from which it follows that

$$\langle n | \psi_n \rangle = 1. \quad (3.10)$$

We can claim this because if $\psi_n^{(i)}$ has some non-zero $\psi_n^{(0)}$ component for $i > 0$, we can absorb it into $\psi_n^{(0)}$ and renormalise ψ_n to make the coefficient of $\psi_n^{(0)}$ to be 1 again.



Figure 6: The perturbation theory is appropriate in the first case, in which the reference Hamiltonian is “close enough” to the true Hamiltonian such that the perturbation only leads to small changes in the energies and wavefunctions. The perturbation theory may not work in the second case where the changes in energy levels is comparable with the energy level separations. The reference Hamiltonian is too far away from the true Hamiltonian. Either we will then need too many terms in the series expansion to get a satisfactory result, or the series just may not converge. We will discuss this convergence issue in appendix section A.

3.1.1 The Perturbation Equations

To work out the coefficients $E_n^{(i)}$ and $\psi_n^{(i)}$ of the power series, we simply need to plug the expansion ansatz (3.7) and (3.6) into the Schrödinger equation (3.1) to get

$$(\hat{H}^{(0)} + \lambda \hat{H}^{(1)} - E_n^{(0)} - \lambda E_n^{(1)} - \lambda^2 E_n^{(2)} + \dots)(\psi_n^{(0)} + \lambda \psi_n^{(1)} + \lambda^2 \psi_n^{(2)} + \dots) = 0 \quad (3.11)$$

For this equation to be satisfied for all λ sufficiently small such that the series converges, the coefficients of all powers of λ must be zero. Gathering the terms in $\lambda^0, \lambda^1, \lambda^2, \dots$, we get a series of equations

$$(\hat{H}^{(0)} - E_n^{(0)}) \left| \psi_n^{(0)} \right\rangle = 0 \quad (3.12)$$

$$(\hat{H}^{(0)} - E_n^{(0)}) \left| \psi_n^{(1)} \right\rangle + (\hat{H}^{(1)} - E_n^{(1)}) \left| \psi_n^{(0)} \right\rangle = 0 \quad (3.13)$$

$$(\hat{H}^{(0)} - E_n^{(0)}) \left| \psi_n^{(2)} \right\rangle + (\hat{H}^{(1)} - E_n^{(1)}) \left| \psi_n^{(1)} \right\rangle - E_n^{(2)} \left| \psi_n^{(0)} \right\rangle = 0 \quad (3.14)$$

\vdots

The zeroth-order equation (3.12) is exactly the unperturbed Schrödinger equation (3.2), which we should have already solved. The next equation of interest is the first-order equation (3.13). If we contract it with $\langle \psi_n^{(0)} | \equiv \langle n |$, we get

$$\langle n | H^{(0)} - E_n^{(0)} | \psi_n^{(1)} \rangle + \langle n | \hat{H}^{(1)} - E_n^{(1)} | n \rangle = 0. \quad (3.15)$$

Since $\hat{H}^{(0)}$ is Hermitian, the first term vanishes:

$$\langle n | H^{(0)} - E_n^{(0)} | \psi_n^{(1)} \rangle = \langle \psi_n^{(1)} | H^{(0)} - E_n^{(0)} | n \rangle^* = 0, \quad (3.16)$$

and we are left with

$$E_n^{(1)} = \langle n | \hat{H}^{(1)} | n \rangle. \quad (3.17)$$

This is the key result in perturbation theory. The first-order correction in energy is simply the expectation value of the perturbation for the unperturbed state. This also implies that

$$E_n^{(0)} + \lambda E_n^{(1)} = \langle n | \hat{H}^{(0)} + \lambda \hat{H}^{(1)} | n \rangle = \langle n | \hat{H} | n \rangle. \quad (3.18)$$

The energy corrected to first order is the expectation value of the complete Hamiltonian with the unperturbed wavefunction.

A similar procedure yields the second-order energy. If we contract the second-order perturbation equation (3.14) with $\langle n|$, we get

$$\langle n|\hat{H}^{(0)} - E_n^{(0)}|\psi_n^{(2)}\rangle + \langle n|\hat{H}^{(1)} - E_n^{(1)}|\psi_n^{(1)}\rangle - \langle n|E_n^{(2)}|n\rangle = 0. \quad (3.19)$$

The first term again vanishes since $\hat{H}^{(0)}$ is Hermitian. By the intermediate normalisation, we also have $E_n^{(1)}\langle n|\psi_n^{(1)}\rangle$ in the second term vanishes and the third term $\langle n|E_n^{(2)}|n\rangle = E_n^{(2)}$. Therefore, the second-order energy is

$$E_n^{(2)} = \langle n|\hat{H}^{(1)}|\psi_n^{(1)}\rangle. \quad (3.20)$$

To use this expression we will need to obtain the first-order wavefunction $\psi_n^{(1)}$. This can be obtained by solving the complicated first-order equation (3.13), which we don't expect to have a simple closed form solution in general unless in very simple cases. We will introduce alternative methods to work this out later.

3.1.2 Higher-order Energies and Wigner's $2n + 1$ Theorem

By expanding higher order terms of (3.11), one can show that the n^{th} -order correction in energy is given by

$$E_n^{(i)} = \langle n|\hat{H}^{(1)}|\psi_n^{(i-1)}\rangle, \quad (3.21)$$

so it seems that we need to know the wavefunction to the order $i - 1$ to calculate the energy to order i . In fact, we can do better than this.

Theorem 3.1 (Wigner's $2n + 1$ rule). The $2n + 1^{\text{th}}$ -order correction of energy can be calculated from the first n^{th} -order perturbation of wavefunctions.

This is more easily seen in a different formulation of the perturbation theory called the *Wigner-Brillouin perturbation theory*, but it is a far more complicated topic. A brief proof of this is in appendix section B, and here we will only show explicitly how to construct $E_n^{(3)}$ from $\psi_n^{(1)}$.

If we contract the first-order perturbation equation (3.13) with the second-order wavefunction $\langle \psi_n^{(2)}|$, we get a overall third-order expression

$$\langle \psi_n^{(2)}|\hat{H}^{(0)} - E_n^{(0)}|\psi_n^{(1)}\rangle + \langle \psi_n^{(2)}|\hat{H}^{(1)} - E_n^{(1)}|n\rangle = 0. \quad (3.22)$$

By (3.21) and the intermediate normalisation, we identify the second term as $E_n^{(3)}$, and so

$$E_n^{(3)} = -\langle \psi_n^{(2)}|\hat{H}^{(0)} - E_n^{(0)}|\psi_n^{(1)}\rangle. \quad (3.23)$$

We can get another third-order expression by contracting the second-order equation (3.14) with the first-order wavefunction $\langle \psi_n^{(1)}|$

$$\langle \psi_n^{(1)}|\hat{H}^{(0)} - E_n^{(0)}|\psi_n^{(2)}\rangle + \langle \psi_n^{(1)}|\hat{H}^{(1)} - E_n^{(1)}|\psi_n^{(1)}\rangle = 0. \quad (3.24)$$

We identify the first term in the above expression as the negative third-order energy by (3.23), and so

$$E_n^{(3)} = \langle \psi_n^{(1)}|\hat{H}^{(1)} - E_n^{(1)}|\psi_n^{(1)}\rangle, \quad (3.25)$$

which contains only the first-order wavefunction as claimed.

We can obtain higher order expression using the same trick as above, but they will be much more complicated to derive and are rarely needed. For a problem to be properly treated using perturbation theory, the higher order corrections should be small and so the series should converge quickly.

3.2 The Hellmann–Feynman Theorem

We have formulated the perturbation problem as the eigenvalue problem of a family of one-parameter Hamiltonians

$$\hat{H}(\lambda)\psi(\lambda) = E(\lambda)\psi(\lambda). \quad (3.26)$$

How the energy $E(\lambda)$ changes against λ actually tells us something more about the system.

Theorem 3.2 (Hellmann–Feynman theorem). Let $E(\lambda)$ and $\psi(\lambda)$ be the eigenvalues and eigenstates of the of one-parameter family of Hamiltonians

$$\hat{H}(\lambda)\psi(\lambda) = E(\lambda)\psi(\lambda), \quad (3.27)$$

then

$$\frac{dE(\lambda)}{d\lambda} = \left\langle \psi(\lambda) \left| \frac{d\hat{H}(\lambda)}{d\lambda} \right| \psi(\lambda) \right\rangle. \quad (3.28)$$

Proof. Let $\psi(\lambda)$ be normalised. Then by the product rule,

$$\begin{aligned} \frac{dE(\lambda)}{d\lambda} &= \frac{d}{d\lambda} \left\langle \psi(\lambda) \left| \hat{H}(\lambda) \right| \psi(\lambda) \right\rangle \\ &= \left\langle \frac{d\psi(\lambda)}{d\lambda} \left| \hat{H}(\lambda) \right| \psi(\lambda) \right\rangle + \left\langle \psi(\lambda) \left| \frac{d\hat{H}(\lambda)}{d\lambda} \right| \psi(\lambda) \right\rangle + \left\langle \psi(\lambda) \left| \hat{H}(\lambda) \right| \frac{d\psi(\lambda)}{d\lambda} \right\rangle \\ &= \left\langle \psi(\lambda) \left| \frac{d\hat{H}(\lambda)}{d\lambda} \right| \psi(\lambda) \right\rangle + E(\lambda) \left[\left\langle \frac{d\psi(\lambda)}{d\lambda} \left| \psi(\lambda) \right\rangle + \left\langle \psi(\lambda) \left| \frac{d\psi(\lambda)}{d\lambda} \right\rangle \right] \right. \\ &= \left\langle \psi(\lambda) \left| \frac{d\hat{H}(\lambda)}{d\lambda} \right| \psi(\lambda) \right\rangle + E(\lambda) \frac{d}{d\lambda} \|\psi(\lambda)\| \\ &= \left\langle \psi(\lambda) \left| \frac{d\hat{H}(\lambda)}{d\lambda} \right| \psi(\lambda) \right\rangle. \end{aligned} \quad (3.29)$$

□

Let's make a connection with the perturbation theory. We can expand $E(\lambda)$ in a Taylor series

$$E(\lambda) = E(0) + \left. \frac{dE}{d\lambda} \right|_{\lambda=0} \lambda + \frac{1}{2} \left. \frac{d^2E}{d\lambda^2} \right|_{\lambda=0} \lambda^2 + \dots \quad (3.30)$$

Then we may identify the n^{th} derivatives of $E(\lambda)$ at $\lambda = 0$ as the n^{th} -order correction in energy

$$E^{(1)} = \left. \frac{dE}{d\lambda} \right|_{\lambda=0} \quad (3.31)$$

$$E^{(2)} = \left. \frac{1}{2} \frac{d^2E}{d\lambda^2} \right|_{\lambda=0} \quad (3.32)$$

⋮

Now, for example, consider a molecule/atom in an electric field of strength \mathcal{E} parallel to the z axis, for which the Hamiltonian is

$$\hat{H}(\mathcal{E}) = \hat{H}^{(0)} - \mathcal{E}\hat{\mu}_z, \quad (3.33)$$

where $\hat{H}^{(0)}$ is the molecular Hamiltonian without external fields and $\hat{\mu}_z = z$ is the z component of the dipole moment operator. We have

$$\frac{d\hat{H}(\mathcal{E})}{d\mathcal{E}} = -\hat{\mu}_z. \quad (3.34)$$

Using the Hellmann–Feynman theorem, we can write

$$E^{(1)} = \left. \frac{dE}{d\mathcal{E}} \right|_{\mathcal{E}=0} = - \langle \psi(\mathcal{E}=0) | \hat{\mu}_z | \psi(\mathcal{E}=0) \rangle . \quad (3.35)$$

This is the expectation value of the dipole moment in the unperturbed state, which is, by definition, the permanent dipole moment,

$$E^{(1)} = -\mu_z^0 . \quad (3.36)$$

Looking at the second-order contribution, we have

$$E^{(2)} = \left. \frac{1}{2} \frac{d^2 E}{d\mathcal{E}^2} \right|_{\mathcal{E}=0} = \left. \frac{1}{2} \frac{d}{d\mathcal{E}} (-\mu_z) \right|_{\mathcal{E}=0} . \quad (3.37)$$

The definition of the polarisability is exactly

$$\alpha_{ij} := \left. \frac{d\mu_i}{d\mathcal{E}_j} \right|_{\mathcal{E}_j=0} , \quad (3.38)$$

and so

$$E^{(2)} = -\frac{1}{2} \alpha_{zz} . \quad (3.39)$$

Example. Quadratic Stark effect and the polarisability of hydrogen atom.

Consider the ground electronic state of the hydrogen atom, where the electronic Hamiltonian, ground state wavefunction and energy are

$$\hat{H}^{(0)} = -\frac{1}{2} \nabla^2 - \frac{1}{r} \quad (3.40)$$

$$\psi_0^{(0)} = \frac{1}{\sqrt{\pi}} e^{-r} \quad (3.41)$$

$$E_0^{(0)} = -\frac{1}{2} . \quad (3.42)$$

We again consider the perturbation of a external electric field of strength \mathcal{E} in the z direction, giving a perturbing Hamiltonian

$$\lambda = \mathcal{E} \quad \hat{H}^{(1)} = z . \quad (3.43)$$

The first-order energy is

$$E_0^{(1)} = \langle 0 | z | 0 \rangle = 0 , \quad (3.44)$$

which can easily be evaluated by symmetry. This shows that a ground state hydrogen atom has no permanent dipole moment. Using the first-order perturbation equation (3.13), one can obtain the differential equation

$$\left(-\frac{1}{2} \nabla^2 - \frac{1}{r} + \frac{1}{2} \right) \psi_0^{(1)} + \frac{1}{\sqrt{\pi}} z e^{-r} = 0 . \quad (3.45)$$

This is one of the extremely rare cases where the first-order wavefunction can be solved analytically in a closed form,¹⁶ which gives

$$\psi_0^{(1)} = -\frac{1}{\sqrt{\pi}} \left(1 + \frac{r}{2} \right) z e^{-r} . \quad (3.46)$$

One can then obtain the exact second-order energy

$$E_0^{(2)} = \langle 0 | z | \psi_0^{(1)} \rangle = -\frac{9}{4} . \quad (3.47)$$

¹⁶Even in this case, this equation is not easy to solve at all! You need to change to parabolic coordinates and do fancy stuffs.

Hence, the electronic energy of hydrogen in an electric field of strength \mathcal{E} is given by

$$E = -\frac{1}{2} - \frac{9}{4}\mathcal{E}^2 + O(\mathcal{E}^4), \quad (3.48)$$

where one can actually argue that only even power terms are permitted by symmetry. This tells us that the ground state hydrogen atom has a polarisability of $\alpha = 9/2$ exactly.

This is a particularly important example, which we will use as a benchmark for the methods introduced later since we know the answer exactly.

3.3 Matrix Formulation

Let's say we have an unperturbed wavefunction expressed in some orthonormal basis $\{\phi_i\}$

$$|\psi_n^{(0)}\rangle = \sum_i c_{ni} \phi_i = \boldsymbol{\phi}^T \mathbf{c}_n. \quad (3.49)$$

Then if $|\psi_n^{(0)}\rangle$ and hence \mathbf{c}_n is normalised, we can express the first-order energy as

$$\begin{aligned} E_n^{(1)} &= \left\langle \sum_i c_{ni} \phi_i \left| \hat{H}^{(1)} \right| \sum_j c_{nj} \phi_j \right\rangle \\ &= \sum_{i,j} c_{ni}^* c_{nj} \left\langle \phi_i \left| \hat{H}^{(1)} \right| \phi_j \right\rangle \\ &= \mathbf{c}_n^\dagger \mathbf{H}^{(1)} \mathbf{c}_n, \end{aligned} \quad (3.50)$$

where $\mathbf{H}^{(1)}$ is the matrix representation of $\hat{H}^{(1)}$ in the $\{\phi_i\}$ basis

$$H_{ij}^{(1)} = \left\langle \phi_i \left| \hat{H}^{(1)} \right| \phi_j \right\rangle. \quad (3.51)$$

3.3.1 Perturbations to Normal Modes

The Hamiltonian of a (unperturbed) molecular vibration is

$$\hat{H}^{(0)}(\mathbf{q}) = -\frac{\hbar^2}{2} \dot{\mathbf{q}}^T \dot{\mathbf{q}} + \frac{1}{2} \mathbf{q}^T \mathbf{K}^{(0)} \mathbf{q}. \quad (3.52)$$

in the mass-weighted basis. If we perturb the system by changing some of the bond strengths or the masses, then this will lead to a change in the dynamical matrix

$$\mathbf{K} = \mathbf{K}^{(0)} + \mathbf{K}^{(1)}. \quad (3.53)$$

If the first-order change in the eigenvalue of \mathbf{H} (energy) is given by $\mathbf{c}^\dagger \mathbf{H}^{(1)} \mathbf{c}$, where \mathbf{c} is the eigenvectors of $\mathbf{H}^{(0)}$ (eigenstates), then we can equally show that the first-order change in the eigenvalues of \mathbf{K} (frequency squared) is given by $\mathbf{Q}^\dagger \mathbf{K}^{(1)} \mathbf{Q}$, where \mathbf{Q} is the eigenvectors of $\mathbf{K}^{(0)}$ (normal modes).¹⁷ Therefore, we can write

$$(\omega_n^2)^{(1)} = \lambda_n^{(1)} = \mathbf{Q}_n^\dagger \mathbf{K}^{(1)} \mathbf{Q}_n. \quad (3.55)$$

¹⁷Can be seen by replacing all the H to K and ψ to Q in the above derivation of perturbation theory. Or equally, you can see

$$\mathbf{H}^{(1)}(\mathbf{q}) = \frac{1}{2} \mathbf{q}^T \mathbf{K}^{(1)} \mathbf{q}, \quad (3.54)$$

so the matrix representation of $\mathbf{H}^{(1)}$ in the $\{\mathbf{q}_i\}$ basis is $\frac{1}{2} \mathbf{K}^{(1)}$. Since $\mathbf{H}^{(0)}$ and $\mathbf{K}^{(0)}$ share the same set of eigenvectors (normal modes), the first-order change in the eigenvalues of $\mathbf{K}^{(0)}$ will be a twice the change in the eigenvalues of $\mathbf{H}^{(0)}$. Hence, if $E_n^{(1)} = \mathbf{c}^\dagger \mathbf{H}^{(1)} \mathbf{c}$, we naturally have $(\omega_n^2)^{(1)} = \lambda_n^{(1)} = \mathbf{Q}_n^\dagger \mathbf{K}^{(1)} \mathbf{Q}_n$.

Example. AAB linear triatomic.

We will consider a AAB triatomic linear molecule as a perturbation of the AAA linear molecule which we have already solved. We will let atom B have a different mass than atom A, and let the bond strength of A – B and A – A be different. Of course this still leads to a 3×3 dynamical matrix, which we can directly diagonalise. However, the algebra will be very tedious, so we will try to treat it using perturbation theory.

The dynamical matrices of interest are

$$\mathbf{K} = \begin{pmatrix} \frac{k_1}{m_A} & -\frac{k_1}{m_A} & 0 \\ -\frac{k_1}{m_A} & \frac{k_1+k_2}{m_A} & -\frac{k_2}{m_A m_B} \\ 0 & -\frac{k_2}{\sqrt{m_A m_B}} & \frac{k_2}{m_B} \end{pmatrix} \quad (3.56)$$

$$\mathbf{K}^{(0)} = \begin{pmatrix} \frac{k_1}{m_A} & -\frac{k_1}{m_A} & 0 \\ -\frac{k_1}{m_A} & \frac{2k_1}{m_A} & -\frac{k_1}{m_A} \\ 0 & -\frac{k_1}{m_A} & \frac{k_1}{m_A} \end{pmatrix} \quad (3.57)$$

$$\begin{aligned} \mathbf{K}^{(1)} &= \mathbf{K} - \mathbf{K}^{(0)} \\ &= \begin{pmatrix} 0 & 0 & 0 \\ 0 & \frac{k_2-k_1}{m_A} & \frac{k_1}{m_A} - \frac{k_2}{m_A m_B} \\ 0 & \frac{k_1}{m_A} - \frac{k_2}{m_A m_B} & \frac{k_2}{m_B} - \frac{k_1}{m_A} \end{pmatrix}. \end{aligned} \quad (3.58)$$

We also know that for the reference system, the three modes are

$$\lambda_1^{(0)} = 0 \quad \mathbf{Q}_1 = \frac{1}{\sqrt{3}} \begin{pmatrix} 1 \\ 1 \\ 1 \end{pmatrix} \quad (3.59)$$

$$\lambda_2^{(0)} = \frac{k_1}{m_A} \quad \mathbf{Q}_2 = \frac{1}{\sqrt{2}} \begin{pmatrix} -1 \\ 0 \\ 1 \end{pmatrix} \quad (3.60)$$

$$\lambda_3^{(0)} = \frac{3k_1}{m_A} \quad \mathbf{Q}_3 = \frac{1}{\sqrt{6}} \begin{pmatrix} 1 \\ -2 \\ 1 \end{pmatrix}. \quad (3.61)$$

Therefore, the first-order eigenvalues are

$$\lambda_1^{(1)} = \mathbf{Q}_1^\dagger \mathbf{K}^{(1)} \mathbf{Q}_1 = \frac{k_2}{3m_A m_B} (\sqrt{m_A} - \sqrt{m_B})^2 \quad (3.62)$$

$$\lambda_2^{(1)} = \mathbf{Q}_2^\dagger \mathbf{K}^{(1)} \mathbf{Q}_2 = \frac{1}{2} \left(\frac{k_2}{m_B} - \frac{k_1}{m_A} \right) \quad (3.63)$$

$$\lambda_3^{(1)} = \mathbf{Q}_3^\dagger \mathbf{K}^{(1)} \mathbf{Q}_3 = \frac{1}{6m_A} \left(\frac{k_2}{m_B} (2\sqrt{m_B} + \sqrt{m_A})^2 - 9k_1 \right). \quad (3.64)$$

The frequencies correct to first order are

$$\omega_i = \sqrt{\lambda_i^{(0)} + \lambda_i^{(1)}}. \quad (3.65)$$

3.4 Rayleigh–Schrödinger Perturbation Theory

Since the eigenstates of a Hermitian operator forms an complete orthonormal set, we can try to expand the first-order wavefunction $|\psi_n^{(1)}\rangle$ in the basis of the unperturbed wavefunction $\{|k\rangle\}$. As we have shown above, $|\psi_n^{(1)}\rangle$ should have no component of $|n\rangle$, so we can write

$$|\psi_n^{(1)}\rangle = \sum_{k \neq n} c_k |k\rangle. \quad (3.66)$$

If we substitute this into the first-order equation (3.13), we get

$$\sum_{k \neq n} c_k (E_k^{(0)} - E_n^{(0)}) |k\rangle + (\hat{H}^{(1)} - E_n^{(1)}) |n\rangle = 0. \quad (3.67)$$

To find the coefficient c_j of a particular $|j\rangle$, we contract the above equation with $\langle j|$ and get

$$\sum_{k \neq n} \langle j | E_k^{(0)} - E_n^{(0)} | k \rangle + \langle j | \hat{H}^{(1)} - E_n^{(1)} | n \rangle = 0, \quad (3.68)$$

and then by orthogonality, we have¹⁸

$$c_j = -\frac{\langle j | \hat{H}^{(1)} | n \rangle}{E_j^{(0)} - E_n^{(0)}} = -\frac{H_{jn}^{(1)}}{E_j^{(0)} - E_n^{(0)}}. \quad (3.69)$$

If we substitute this back (3.66), we get an expansion of the first-order wavefunction

$$|\psi_n^{(1)}\rangle = -\sum_{j \neq n} \frac{H_{jn}^{(1)}}{E_j^{(0)} - E_n^{(0)}} |j\rangle. \quad (3.70)$$

We can also substitute this into (3.20) to get an expansion of the second-order energy

$$\begin{aligned} E_n^{(2)} &= \langle n | \hat{H}^{(2)} | \psi_n^{(1)} \rangle \\ &= -\sum_{j \neq n} \frac{H_{nj}^{(1)} H_{jn}^{(1)}}{E_j^{(0)} - E_n^{(0)}} \\ &= -\sum_{j \neq n} \frac{|H_{jn}^{(1)}|^2}{E_j^{(0)} - E_n^{(0)}}, \end{aligned} \quad (3.71)$$

where notice that $H^{(1)}$ is Hermitian. This is the key result of Rayleigh–Schrödinger perturbation theory. Higher-order results may be obtained in analogous ways, but they are rarely needed.¹⁹

Example. Polarisability of hydrogen Revisited.

Last time using the exact result of the first-order wavefunction, we have worked out that the polarisability of hydrogen is $\alpha = 4.5$ exactly. Now let's see what Rayleigh–Schrödinger gives us.

Since we have identified $\alpha = -2E_0^{(2)}$ from Hellmann–Feynman theorem, the ground state polarisability of hydrogen is given by

$$\alpha = 2 \sum_{k \neq 0} \frac{|\langle 0 | z | k \rangle|^2}{E_k^{(0)} - E_0^{(0)}}. \quad (3.72)$$

The ground state $|0\rangle$ is totally symmetric, so for the integral $\langle 0 | z | k \rangle$ to be non-zero, k should transform the same as the irreducible representation as the Cartesian function z — these are exactly the

¹⁸This expression goes well with our assumption that $|n\rangle$ is non-degenerate. If $|n\rangle$ is degenerate with some $|j\rangle$, then this expression will diverge. We will comment a bit more on this when discussing the degenerate perturbation theory.

¹⁹In fact due to some converging issue that we will discuss later, stopping at the first or the second-order correction is usually a better choice.

$2p_z, 3p_z, \dots$ orbitals.²⁰ The first of these is

$$\psi_{2p_z}^{(0)} = \frac{1}{\sqrt{32\pi}} z e^{-r/2} \quad (3.74)$$

with energy $-\frac{1}{8}$. This gives a contribution 2.96 to α . Clearly this is a poor estimate of α since we have only calculated the first term. You will evaluate the $3p_z$ contribution in the exercises. As you include more and more contributions, the result should converge, albeit very slowly since the denominator of (3.72) does not get any larger than $\frac{1}{2}$, so we are only relying on the numerator for convergence.

However, if you really calculate this series, summing up over all np_z orbitals, you will find that it actually converges to a wrong value! This is because there are also continuum (unbounded) states above the bounded energy levels. We also need to include these into our summation. This is very tricky to do, even numerically on a computer, and it is included in section C together with an alternative method by A. Dalgarno and J.T. Lewis that can evaluate the exact polarisability of hydrogen in a clever way.

A much easier thing to do is to give an upper bound for this series. Since $E_k^{(0)} - E_0^{(0)} > E_{n=2}^{(0)} - E_0^{(0)} = \frac{3}{8}$ for all (bounded and unbounded) states k ,²¹ we have

$$\alpha < \frac{16}{3} \sum_{k \neq 0} |\langle 0|z|k \rangle|^2. \quad (3.75)$$

If we write

$$\sum_{k \neq 0} \langle 0|z|k \rangle \langle k|z|0 \rangle, \quad (3.76)$$

we see something very similar to the resolution of identity in the middle of this expression, except it excluded $|0\rangle$ from all states $|0\rangle$. This is actually not a problem, since $\langle 0|z|0\rangle$ is exactly the first-order energy, which we have identified to be zero. Therefore,

$$\sum_{k \neq 0} \langle 0|z|k \rangle \langle k|z|0 \rangle = \sum_k \langle 0|z|k \rangle \langle k|z|0 \rangle = \langle 0|z^2|0 \rangle. \quad (3.77)$$

We have reduced this infinite series into a single term! Due to the spherical symmetry of $|0\rangle$, this can also be easily evaluated as

$$\langle 0|z^2|0 \rangle = a_0^2, \quad (3.78)$$

where the Bohr radius $a_0 = 1$ in atomic units. Hence, we have an upper bound

$$\alpha < \frac{16}{3}. \quad (3.79)$$

The exact value $\alpha = 4.5$ does sit in this range.

Despite its practical difficulty, Rayleigh–Schrödinger perturbation theory provides some useful physical insight. The expression (3.72) shows that an atom will have a large polarisability if it possess low-lying excited states with an appropriate symmetry.

²⁰Since the wavefunction $|k\rangle \equiv |n, \ell, m\rangle = R_{n,\ell}(r)Y_{\ell,m}(\theta, \varphi)$ is a product of totally symmetric radial wavefunction and a spherical harmonic $|\ell, m\rangle = Y_{\ell,m}(\theta, \varphi)$, we need $Y_{\ell,m}(\theta, \varphi)$ to transform as the Cartesian function z . These are the $|1, 0\rangle = Y_{1,0}(\theta, \varphi)$ spherical harmonics, which combines with the radial wavefunctions $R_{n,1}(r)$, $n \geq \ell = 1$ to give np_z orbitals, $n \geq 1$.

More generally, we have

$$\langle n', \ell', m' | z | n \ell m \rangle = 0 \quad \text{unless} \quad \begin{cases} \ell' = \ell \pm 1 \\ m' = m. \end{cases} \quad (3.73)$$

This comes from the Wigner–Eckart theorem with $T_{q=0}^{(1)}$ and parity consideration. See my notes on Mathematical Tripos Part II: *Principles of Quantum Mechanics* for more detail.

²¹We cannot give a lower bound of α using this method because the energy of unbounded states can be arbitrarily high. A lower bound can be given using variation perturbation theory, which will be introduced in section 3.6.

3.5 Ladder Operators

A *ladder operator* in quantum mechanics is an operator that acts on an eigenstate of another operator to produce an eigenfunction with a different value. Since the changes to the quantum number are usually $+1$ (raising operator) or -1 (lowering operator), they got their names by imaging stepping up or down onw ‘rung’ on the ‘ladder’ of eigenstates.

If we have an Hermitian operator \hat{B} , and another operator \hat{a} which satisfy the commutating relationship

$$[\hat{B}, \hat{a}] = c\hat{a} \quad (3.80)$$

for some real constant c , then \hat{a} is a ladder operator²² for \hat{B} . To show this, consider acting \hat{a} on the eigenstate of \hat{B} , $|b\rangle$ with eigenvalue b . We have

$$\begin{aligned} \hat{B}\hat{a}|b\rangle &= \hat{a}\hat{B}|b\rangle + [\hat{B}, \hat{a}]|b\rangle \\ &= b\hat{a}|b\rangle + c\hat{a}|b\rangle \\ &= (b+c)\hat{a}|b\rangle, \end{aligned} \quad (3.81)$$

and so $\hat{a}|b\rangle$ is an eigenstate of \hat{B} with eigenvalue $b+c$. Depending on the sign of c , this state is either raised or lowered.

By taking the adjoint of (3.80), we have

$$[\hat{B}, \hat{a}^\dagger] = -c\hat{a}^\dagger. \quad (3.82)$$

Hence, if \hat{a} is a raising (lowering) operator, then its Hermitian conjugate \hat{a}^\dagger will be a lowering (raising) operator, changing the magnitude of the eigenvalue by the same amount.

3.5.1 Angular Momentum Ladder Operators

By the commutation relationship of angular momentum operator

$$[\hat{J}_i, \hat{J}_j] = \sum_k \epsilon_{ijk} \hat{J}_k \quad [\hat{\mathbf{J}}^2, \hat{J}_i] = 0 \quad (3.83)$$

for $i, j, k \in \{x, y, z\}$, one can show that the operators

$$\hat{J}_+ = \hat{J}_x + i\hat{J}_y \quad (3.84)$$

$$\hat{J}_- = \hat{J}_x - i\hat{J}_y \quad (3.85)$$

satisfies the commutation relationships

$$[\hat{J}_z, \hat{J}_\pm] = \pm\hat{J}_\pm \quad (3.86)$$

$$[\hat{\mathbf{J}}^2, \hat{J}_\pm] = 0, \quad (3.87)$$

and so they are ladder operators for \hat{J}_z . We have

$$\hat{J}_z\hat{J}_\pm|J, M\rangle = (M \pm 1)\hat{J}_\pm|J, M\rangle, \quad (3.88)$$

$$\hat{\mathbf{J}}^2\hat{J}_\pm|J, M\rangle = J(J+1)\hat{J}_\pm|J, M\rangle, \quad (3.89)$$

where $|J, M\rangle = Y_{JM}$ is the angular momentum eigenstates. Therefore,

$$\hat{J}_\pm|J, M\rangle \propto |J, M \pm 1\rangle. \quad (3.90)$$

²²The notation *a* here stands for *annihilation*, as \hat{a} and \hat{a}^\dagger are more formally known as the *annihilation* and *creation* operator. The create/destroy thw quantum of vibration (phonon).

3.5.2 Harmonic Oscillator Ladder Operator

The Hamiltonian of a standard harmonic oscillator takes the form

$$\hat{H} = -\frac{1}{2} \frac{d^2}{dq^2} + \frac{1}{2} q^2. \quad (3.91)$$

We will first show how to use the ladder operators to infer the eigenvalues of harmonic oscillator without needing to solve any differential equations. The ladder operators of the Harmonic oscillator are given by

$$\hat{a} = \sqrt{\frac{1}{2}} \left(q + \frac{d}{dq} \right) \quad (3.92)$$

$$\hat{a}^\dagger = \sqrt{\frac{1}{2}} \left(q - \frac{d}{dq} \right). \quad (3.93)$$

It is easy to see that \hat{a} lowers the eigenvalue of an eigenstate by 1, and \hat{a}^\dagger raise the eigenvalue by 1. Additionally, we find that the ladder operators nicely “factor out” the Hamiltonian

$$\hat{a}\hat{a}^\dagger = \hat{H} + \frac{1}{2} \quad \hat{a}^\dagger\hat{a} = \hat{H} - \frac{1}{2}. \quad (3.94)$$

We define the *numbering operator* $\hat{N} = \hat{a}^\dagger\hat{a} = \hat{H} - \frac{1}{2}$, with $\hat{N}|n\rangle = n|n\rangle$.²³ Then $|n\rangle$ is also an eigenstate of \hat{H} , with $\hat{H}|n\rangle = (n + \frac{1}{2})|n\rangle$. However, since

$$n = \langle n|\hat{N}|n\rangle = \langle n|\hat{a}^\dagger\hat{a}|n\rangle = \|\hat{a}|n\rangle\|^2 \geq 0, \quad (3.95)$$

we find that we cannot lower the eigenstate $|n\rangle$ infinitely. We must have $n \geq 0$, so the lowering process must somehow terminate at some point to avoid negative values of n . This will be our ground state, and is given by $\hat{a}|n\rangle = 0$. Therefore, we must have $n = 0$ to be the ground state. The excited states can be obtained by repeatedly acting \hat{a}^\dagger on $|0\rangle$, so the eigenvalues of \hat{H} are given by $k + \frac{1}{2}$, where $k \in \mathbb{N}_0$.

Using this method, we can also easily find the eigenfunctions of \hat{H} . If we directly solve the Schrödinger equation, then this is a second-order inhomogeneous differential equation, which we can't easily solve. The standard method would then be to solve this using an infinite series. However, we have claimed that the ground state satisfies

$$\hat{a}\psi(q) = \sqrt{\frac{1}{2}} \left(q\psi + \frac{d\psi}{dq} \right) = 0. \quad (3.96)$$

This is now a first-order, linear, homogeneous equation, which we know the general solution. The ground state wavefunction is therefore

$$\psi(q) = \frac{1}{\sqrt{2\pi}} e^{-\frac{1}{2}q^2}. \quad (3.97)$$

The wavefunction of the excited state can be obtained by acting \hat{a}^\dagger repeatedly.

We know that $\hat{a}|n\rangle \propto |n-1\rangle$ and $\hat{a}^\dagger|n\rangle \propto |n+1\rangle$, but it might not be normalised. Suppose $\hat{a}|n\rangle = c_n|n-1\rangle$, where c_n is real to keep all eigenfunctions real. Then consider

$$\langle n|\hat{a}^\dagger\hat{a}|n\rangle = |c_n|^2 \langle n-1|n-1\rangle = |c_n|^2. \quad (3.98)$$

We also have

$$\langle n|\hat{a}^\dagger\hat{a}|n\rangle = \langle n|\hat{N}|n\rangle = n \langle n|n\rangle = n. \quad (3.99)$$

Equating the above two results give $c_n = \sqrt{n}$. We can use the same method to work out the coefficient for the raising operator.

$$\hat{a}|n\rangle = \sqrt{n}|n-1\rangle \quad \hat{a}^\dagger|n\rangle = \sqrt{n+1}|n+1\rangle. \quad (3.100)$$

²³It is named so because it counts the number of quantum of vibrations (phonon) in a vibrational eigenstate.

Example. Harmonic oscillator with cubic perturbation potential.

Now consider adding a cubic perturbation $\hat{H}^{(1)} = \lambda q^3$ to the harmonic oscillator Hamiltonian. We anticipate to evaluate integrals like $\langle j | q^3 | n \rangle$, which seems algebraically tedious. However, this can be much easily done using ladder operators.

From the definition of ladder operators, one can write

$$q = \frac{1}{\sqrt{2}}(\hat{a} + \hat{a}^\dagger). \quad (3.101)$$

so if we act it on some eigenstate of \hat{H} , we will get

$$q |n\rangle = \sqrt{\frac{n+1}{2}} |n+1\rangle + \sqrt{\frac{n}{2}} |n-1\rangle. \quad (3.102)$$

Now, let's consider what $q^3 |n\rangle$ will give us. By taking the cube of (3.101), we see that q^3 constitutes of 8 different combination raising and lowering operators. One of the combinations is triple raising operator $\hat{a}^\dagger \hat{a}^\dagger \hat{a}^\dagger$ which will raise $|n\rangle$ by three quanta, and another is triple lowering operator $\hat{a} \hat{a} \hat{a}$. There will be three combinations that raise $|n\rangle$ by one quantum ($\hat{a}^\dagger \hat{a}^\dagger \hat{a}$, $\hat{a}^\dagger \hat{a} \hat{a}^\dagger$, $\hat{a} \hat{a}^\dagger \hat{a}^\dagger$), and three that lowers $|n\rangle$ by one quantum ($\hat{a}^\dagger \hat{a} \hat{a}$, $\hat{a} \hat{a}^\dagger \hat{a}$, $\hat{a} \hat{a} \hat{a}^\dagger$). Therefore, when acting on $|n\rangle$, we expect to see a mixture of four states: $|n \pm 3\rangle$ and $|n \pm 1\rangle$.

Let's first consider the first-order variation in energy, given by

$$E_n^{(1)} = \langle n | q^3 | n \rangle. \quad (3.103)$$

Since $q^3 |n\rangle$ is only a combination of $|n \pm 3\rangle$ and $|n \pm 1\rangle$, it will be zero by orthogonality.

Next, we will consider the second-order energy

$$E_n^{(2)} = -\lambda^2 \sum_{j \neq n} \frac{|\langle j | q^3 | n \rangle|^2}{j - n}. \quad (3.104)$$

For simplicity, we will consider the ground state $|n\rangle = |0\rangle$. Since we cannot lower $|0\rangle$ anymore, all the single lowering and triple contributions vanishes. Of the remaining four terms, we find that

$$\hat{a}^\dagger \hat{a}^\dagger \hat{a} |0\rangle = 0 \quad (3.105)$$

also vanishes since it firsts lowers $|0\rangle$. The other three terms are then

$$\hat{a}^\dagger \hat{a} \hat{a}^\dagger |0\rangle = \sqrt{1}\sqrt{1}\sqrt{1} |1\rangle = |1\rangle \quad (3.106)$$

$$\hat{a} \hat{a}^\dagger \hat{a}^\dagger |0\rangle = \sqrt{1}\sqrt{2}\sqrt{2} |1\rangle = 2 |1\rangle \quad (3.107)$$

$$\hat{a}^\dagger \hat{a}^\dagger \hat{a}^\dagger |0\rangle = \sqrt{1}\sqrt{2}\sqrt{3} |3\rangle = \sqrt{6} |3\rangle, \quad (3.108)$$

and so

$$\langle 1 | q^3 | 0 \rangle = \frac{1}{\sqrt{8}}(2+1) \langle 1 | 1 \rangle = \frac{3}{2\sqrt{2}} \quad (3.109)$$

$$\langle 3 | q^3 | 0 \rangle = \frac{1}{\sqrt{8}}\sqrt{6} \langle 3 | 3 \rangle = \frac{\sqrt{3}}{2}. \quad (3.110)$$

Therefore, the second-order energy is

$$E_0^{(2)} = -\lambda^2 \left(\frac{|\langle 1 | q^3 | 0 \rangle|^2}{1-0} + \frac{|\langle 3 | q^3 | 0 \rangle|^2}{3-0} \right) = -\frac{11}{8} \lambda^2. \quad (3.111)$$

This is an exact result. Although the Rayleigh–Schrödinger sum runs over an infinite series of excited states, only two of them have non-zero contributions. In contrast to the polarisability example, the series converges really fast here.

Using a similar approach, one can find the exact first-order wavefunction of the ground state to be

$$\psi_0^{(1)} = -\lambda \left(q + \frac{1}{3}q^3 \right) \psi_0^{(0)}. \quad (3.112)$$

3.6 Variational Perturbation Theory

You should be familiar with the variational principles.

Lemma 3.3 (Variational principles). Let \hat{H} be a Hamiltonian with ground state wavefunction ψ_0 with energy E_0 . Let $\tilde{\psi}$ be any trial wavefunction, then

$$\tilde{E} = \frac{\langle \tilde{\psi} | \hat{H} | \tilde{\psi} \rangle}{\langle \tilde{\psi} | \tilde{\psi} \rangle} \geq E_0, \quad (3.113)$$

where the equality holds if and only if $\tilde{\psi} = \psi_0$.

Now consider the ground state energy corrected to the first order,

$$E_0^{(0)} + \lambda E_0^{(1)} = \langle 0 | \hat{H}^{(0)} | 0 \rangle + \langle 0 | \lambda \hat{H}^{(1)} | 0 \rangle = \langle 0 | \hat{H} | 0 \rangle. \quad (3.114)$$

This can be seen as using the unperturbed ground state $|0\rangle$ as a trial function for the perturbed Hamiltonian, so we must have

$$E_0^{(0)} + \lambda E_0^{(1)} \geq E_0. \quad (3.115)$$

The ground state energy corrected to first order is greater than the true ground state energy.

However, if we consider the second-order energy

$$E_0^{(2)} = \langle \psi_0^{(1)} | \hat{H}^{(1)} | 0 \rangle, \quad (3.116)$$

we see that it is not a expectation value. Therefore, the variational principles do not apply, and the energy corrected to the second order $E_0^{(0)} + \lambda E_0^{(1)} + \lambda^2 E_0^{(2)}$ may be lower than the true ground state energy E_0 .

3.6.1 An Upper Bound to the Second Order Energy

Although the variational principles do not apply, we can get an upper bound for $E_0^{(2)}$ using some trial first-order wavefunction. This will be helpful if we can't solve the first-order wavefunction exactly, nor do we want to expand it as a series, but we have some intuitive guess of what its form might be.

Let φ be a trial function that approximates the true (unknown) ground-state first-order wavefunction $\psi_0^{(1)}$. Let $\eta = \varphi - \psi_0^{(1)}$. As usual, we require the first-order wavefunctions to be orthogonal to the zeroth-order wavefunction, and we also impose this property on the trial function

$$\langle \varphi | 0 \rangle = \langle \eta | 0 \rangle = 0. \quad (3.117)$$

Now consider the quantity

$$X^{(2)} = \langle \varphi | \hat{H}^{(0)} - E_0^{(0)} | \varphi \rangle + \langle \varphi | \hat{H}^{(1)} | 0 \rangle + \langle 0 | \hat{H}^{(1)} | \varphi \rangle. \quad (3.118)$$

Substituting $\varphi = \psi_0^{(1)} + \eta$ and expanding, we have

$$\begin{aligned} X^{(2)} = & \langle \eta | \hat{H}^{(0)} - E_0^{(0)} | \eta \rangle + \langle \psi_0^{(1)} | \hat{H}^{(0)} - E_0^{(0)} | \eta \rangle \\ & + \langle \eta | \hat{H}^{(0)} - E_0^{(0)} | \psi_0^{(1)} \rangle + \langle \psi_0^{(1)} | \hat{H}^{(0)} - E_0^{(0)} | \psi_0^{(1)} \rangle \\ & + \langle \psi_0^{(1)} | \hat{H}^{(1)} | 0 \rangle + \langle \eta | \hat{H}^{(1)} | 0 \rangle + \langle 0 | \hat{H}^{(1)} | \psi_0^{(1)} \rangle + \langle 0 | \hat{H}^{(1)} | \eta \rangle . \end{aligned} \quad (3.119)$$

We can use the first-order perturbation equation (3.13) to replace $(\hat{H}^{(0)} - E_0^{(0)}) |\psi_0^{(1)}\rangle$ by $-(\hat{H}^{(1)} - E_0^{(1)}) |0\rangle$, and then a lot of terms cancels out or vanishes by orthogonality, and we are left with

$$X^{(2)} = \langle \eta | \hat{H}^{(0)} - E_0^{(0)} | \eta \rangle + \langle 0 | \hat{H}^{(1)} | \psi_0^{(1)} \rangle . \quad (3.120)$$

The first term is positive by the variational principle, and the second term is the second-order energy. Hence we can conclude that

$$X^{(2)} \geq E_0^{(2)} , \quad (3.121)$$

so $X^{(2)}$ is an upper bound to the second-order energy of the ground state. As usual, we can include some parameters in φ , and minimise $X^{(2)}$ to get the tightest upper bound.

Example. Polarisability of hydrogen revisited.

Let's consider again the polarisability of hydrogen. Since we are applying an electric field along the z direction, we might reasonably choose a trial function

$$\varphi = Az\psi_0^{(0)} \quad (3.122)$$

for the first-order wavefunction, where A is some constant. Note that $\langle \varphi | 0 \rangle = 0$ as we imposed before. If we substitute this ansatz into (3.118), we get

$$X^{(2)} = A^2 \langle z\psi_0^{(0)} | \hat{H}^{(0)} - E_0^{(0)} | z\psi_0^{(0)} \rangle + 2A \langle 0 | z^2 | 0 \rangle . \quad (3.123)$$

This is evaluated to be

$$X^{(2)} = \frac{1}{2}A^2 + 2A . \quad (3.124)$$

$X^{(2)}$ is minimised when $A = -2$, taking the value $X^{(2)} = -2$, which is our upper bound to the second-order energy. Recalling that the polarisability is $\alpha = -2E^{(2)}$, we conclude that $\alpha = 4$ is an lower bound to the ground state polarisability of the hydrogen atom.

Comparing with the exact value $\alpha = 4.5$, we see that this is a really good one-term estimation, significantly better than the first term estimation in the Rayleigh–Schrödinger.

3.7 Multiple Perturbations

Sometimes we have two or more perturbations happening at the same time. It is helpful to treat such cases according to the relative sizes of the perturbations.

3.7.1 Perturbations of Unequal Magnitudes

If one effect is substantially stronger than the other, it makes sense to regard the weaker one as a second-order perturbation to the Hamiltonian, and we can write

$$\hat{H} = \hat{H}^{(0)} + \lambda \hat{H}^{(1)} + \lambda^2 \hat{H}^{(2)} . \quad (3.125)$$

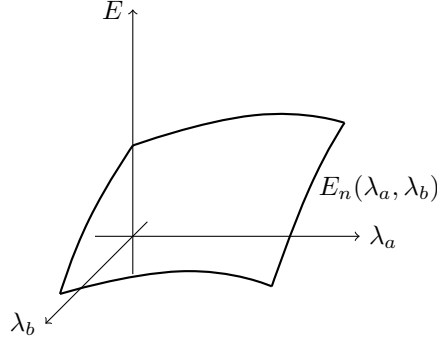


Figure 7: The energy of state n as a function of the parameters λ_a , λ_b when imposing two perturbations of comparable strengths.

We again substitute the same series expansions of ψ_n and E_n to the perturbed Schrödinger equation. The first-order equation remains unchanged, so the first-order energy and wavefunction are the same. However, the second-order equation now has an extra term and becomes

$$(\hat{H}^{(0)} - E_n^{(0)}) \left| \psi_n^{(2)} \right\rangle + (\hat{H}^{(1)} - E_n^{(1)}) \left| \psi_n^{(1)} \right\rangle + (\hat{H}^{(2)} - E_n^{(2)}) |n\rangle = 0. \quad (3.126)$$

Contracting with $\langle n|$ gives the expression of the second-order energy

$$E_n^{(2)} = \langle n | \hat{H}^{(1)} | \psi_n^{(1)} \rangle + \langle n | \hat{H}^{(2)} | n \rangle, \quad (3.127)$$

which now has an extra term compared with the single perturbation case.

3.7.2 Perturbations of Comparable Strengths

When two perturbations of similar strengths are present, it is more appropriate to treat them as two first-order perturbations, each with its own prefactor

$$\hat{H} = \hat{H}^{(0)} + \lambda_a \hat{H}_a^{(1)} + \lambda_b \hat{H}_b^{(1)}. \quad (3.128)$$

This is a two dimensional family of Hamiltonians, and so the energies and wavefunctions are given by two dimensional series

$$\begin{aligned} E_n &= \sum_{j,k} \lambda_a^j \lambda_b^k E_n^{(j,k)} \\ &= E_n^{(0,0)} + \underbrace{\lambda_a E_n^{(1,0)} + \lambda_b E_n^{(0,1)}}_{\text{first order}} + \underbrace{\lambda_a^2 E_n^{(2,0)} + \lambda_a \lambda_b E_n^{(1,1)} + \lambda_b^2 E_n^{(0,2)}}_{\text{second order}} + \dots \end{aligned} \quad (3.129)$$

$$\begin{aligned} |\psi_n\rangle &= \sum_{j,k} \lambda_a^j \lambda_b^k \left| \psi_n^{(j,k)} \right\rangle \\ &= \left| \psi_n^{(0,0)} \right\rangle + \underbrace{\lambda_a \left| \psi_n^{(1,0)} \right\rangle + \lambda_b \left| \psi_n^{(0,1)} \right\rangle}_{\text{first order}} + \underbrace{\lambda_a^2 \left| \psi_n^{(2,0)} \right\rangle + \lambda_a \lambda_b \left| \psi_n^{(1,1)} \right\rangle + \lambda_b^2 \left| \psi_n^{(0,2)} \right\rangle}_{\text{second order}} + \dots \end{aligned} \quad (3.130)$$

We can then substitute these series ansatz into the Schrödinger equation of the doubly perturbed Hamiltonian. Since λ_a and λ_b are independent variables, we need to gather and equate terms in each combination of them separately. The terms in λ_a and λ_b leads to two first-order equations

$$(\hat{H}_a^{(1)} - E_n^{(1,0)}) \left| \psi_n^{(0,0)} \right\rangle + (\hat{H}^{(0)} - E_n^{(0,0)}) \psi_n^{(1,0)} = 0 \quad (3.131)$$

$$(\hat{H}_b^{(1)} - E_n^{(0,1)}) \left| \psi_n^{(0,0)} \right\rangle + (\hat{H}^{(0)} - E_n^{(0,0)}) \psi_n^{(0,1)} = 0, \quad (3.132)$$

and contracting with $\langle \psi_n^{(0,0)} | \equiv \langle n |$ yields two first-order energy expressions analogous to (3.17)

$$E_n^{(1,0)} = \langle n | \hat{H}_a^{(1)} | n \rangle \quad (3.133)$$

$$E_n^{(0,1)} = \langle n | \hat{H}_b^{(1)} | n \rangle. \quad (3.134)$$

The second-equations in λ_a^2 and λ_b^2 are analogous to (3.14). However, we also have a second-order equation in $\lambda_a \lambda_b$ containing cross terms

$$(\hat{H}^{(0)} - E_n^{(0,0)}) |\psi_n^{(1,1)}\rangle + (\hat{H}_a^{(1)} - E_n^{(1,0)}) |\psi_n^{(1,0)}\rangle + (\hat{H}_b^{(1)} - E_n^{(0,1)}) |\psi_n^{(1,0)}\rangle - E_n^{(1,1)} |n\rangle = 0. \quad (3.135)$$

Contracting with $\langle n |$ gives

$$E_n^{(1,1)} = \langle n | \hat{H}_a^{(1)} | \psi_n^{(0,1)} \rangle + \langle n | \hat{H}_b^{(1)} | \psi_n^{(1,0)} \rangle. \quad (3.136)$$

3.7.3 Interchange Theorem

This expression contains both $|\psi_n^{(1,0)}\rangle$ and $|\psi_n^{(0,1)}\rangle$ — we can do slightly better than that. If we take inner products of (3.131) and (3.132) with $|\psi_n^{(0,1)}\rangle$ and $|\psi_n^{(1,0)}\rangle$, respectively, and we get two mixed second-order equations. From these two equations, we can easily get

$$\langle n | \hat{H}_a^{(1)} | \psi_n^{(0,1)} \rangle = \langle \psi_n^{(1,0)} | \hat{H}_b^{(1)} | n \rangle. \quad (3.137)$$

This allows us to rewrite the mixed second-order energy as

$$E_n^{(1,1)} = \langle n | \hat{H}_a^{(1,0)} | \psi_n^{(0,1)} \rangle + \langle \psi_n^{(0,1)} | \hat{H}_a^{(1)} | n \rangle. \quad (3.138)$$

This is known as the *interchange theorem*. Analogous results hold for mixed higher-order energy expressions as well, although they are much more complicated.

Example. Molecular polarisability.

In previous example, we have examined the polarisability of a hydrogen atom. Due to its spherical symmetry, the first-order energy vanishes, and a single number is sufficient to assign its (isotropic) polarisability. However, a molecule may possess a permanent dipole moment and its polarisability may be anisotropic. We must therefore consider the orientations of the molecules and the electric field.

Suppose a molecule consists of a collection of N point charges (including nuclei and electron) e_i located at positions $\{\mathbf{r}_i\}$. In the presence of a uniform electric field \mathcal{E} , there is an extra energy, which we see as a perturbation, given by

$$\hat{H}^{(1)} = - \sum_{i=1}^N e_i \mathcal{E} \cdot \mathbf{r}_i = -\mathcal{E} \cdot \hat{\boldsymbol{\mu}}, \quad (3.139)$$

where we defined the *dipole moment operator* to be

$$\hat{\boldsymbol{\mu}} = \sum_i e_i \mathbf{r}_i. \quad (3.140)$$

In component form,

$$\hat{H}^{(1)} = - \sum_{\beta} \mathcal{E}_{\beta} \hat{\mu}_{\beta}, \quad (3.141)$$

where $\beta \in \{x, y, z\}$. This is three independent perturbation of comparable magnitudes.

By Hellmann–Feynman theorem, we can write

$$E_0 = E_0^{(0)} - \sum_{\beta} \mu_{\beta}^0 \mathcal{E}_{\beta} - \frac{1}{2} \sum_{\beta, \gamma} \alpha_{\beta\gamma} \mathcal{E}_{\beta} \mathcal{E}_{\gamma} - \dots \quad (3.142)$$

for $\beta, \gamma \in \{x, y, z\}$, where $\boldsymbol{\mu}^0$ is the permanent dipole moment vector and α is the polarisability tensor. The first-order energy is

$$E_0^{(1)} = \langle 0 | \hat{H}^{(1)} | 0 \rangle = - \sum_{\beta} \mathcal{E}_{\beta} \langle 0 | \hat{\mu}_{\beta} | 0 \rangle, \quad (3.143)$$

and we indeed have $\boldsymbol{\mu}^0 = \langle 0 | \hat{\boldsymbol{\mu}} | 0 \rangle$. For the second-order energy, we apply have the Rayleigh–Schrödinger result

$$\begin{aligned} E_0^{(2)} &= - \sum_{j \neq 0} \frac{|\langle j | -\boldsymbol{\mathcal{E}} \cdot \hat{\boldsymbol{\mu}} | 0 \rangle|^2}{E_j^{(0)} - E_0^{(0)}} \\ &= - \sum_{\beta, \gamma} \mathcal{E}_{\beta} \mathcal{E}_{\gamma} \sum_{j \neq 0} \frac{\langle j | \hat{\mu}_{\beta} | 0 \rangle \langle 0 | \hat{\mu}_{\gamma} | j \rangle}{E_j^{(0)} - E_0^{(0)}}. \end{aligned} \quad (3.144)$$

We can identify the latter sum as $\frac{1}{2} \alpha_{\beta\gamma}$, but the polarisability is usually put into a symmetric form, although any antisymmetric part will cancel in the total second-order energy in (3.142). Therefore, we will choose the components of the polarisability to be in the symmetrised form

$$\alpha_{\beta\gamma} = \sum_{j \neq 0} \frac{\langle j | \hat{\mu}_{\beta} | 0 \rangle \langle 0 | \hat{\mu}_{\gamma} | j \rangle + \langle j | \hat{\mu}_{\gamma} | 0 \rangle \langle 0 | \hat{\mu}_{\beta} | j \rangle}{E_j^{(0)} - E_0^{(0)}}. \quad (3.145)$$

4 Degenerate Perturbation Theory

Degeneracy presents issues in the above formulation of perturbation theory. First, it is obvious to see that if any state is degenerate to the unperturbed state, the denominator in the Rayleigh–Schrödinger sum (3.70) will be zero. This suggests that an arbitrarily small perturbation will cause an extremely dramatic shift in the degenerate states of the system.

This is not a unfamiliar effect. If we put a small ball in a bowl, then it will stay at the distinct, non-degenerate ground state at the bottom. If we tilt the bowl slightly, then the ball will also move slightly in the direction we tilt it to find the new minimum. However, if we put a ball on a flat table, where any point on it is degenerate, tilting the table however slightly will make the ball run away to infinity in the direction we tilt it. Perturbation lifts degeneracy.

Suppose we impose a perturbation that lifts the degeneracy, thereby leading to unique perturbed states. If we gradually decrease the strength λ of the perturbation to zero, we expect that the system will return to a particular linear combination of the degenerate unperturbed state. This combination should resemble the perturbed states for small λ . The key to degenerate perturbation theory is thus to identify those correct zeroth-order wavefunctions stable to perturbation.

4.1 Secular Equations for the Unperturbed Wavefunctions

Suppose there is a M -fold degenerate state n with energy $E_n^{(0)}$. We denote the i^{th} component of the unperturbed state by $|\psi_{ni}^{(0)}\rangle$, $i \in \{1, \dots, M\}$, where these degenerate wavefunctions are mutually orthogonal $\langle \psi_{ni}^{(0)} | \psi_{nj}^{(0)} \rangle = \delta_{ij}$. Any linear combination of these functions has unperturbed energy $E_n^{(0)}$. We write the zeroth-order wavefunctions that perturbed states returns to in the limit $\lambda \rightarrow 0$ as $|\Phi_{nj}^{(0)}\rangle$. They are just linear combinations of the basis $\{|\psi_{ni}^{(0)}\rangle\}$, so we may write

$$|\Phi_{nj}^{(0)}\rangle = \sum_{i=1}^M |\psi_{ni}^{(0)}\rangle c_{ij} \quad (4.1)$$

The perturbation expansions of the states are therefore

$$|\psi_{nj}(\lambda)\rangle = |\Phi_{nj}^{(0)}\rangle + \lambda |\psi_{nj}^{(1)}\rangle + \lambda^2 |\psi_{nj}^{(2)}\rangle + \dots, \quad (4.2)$$

giving the first-order equations

$$(\hat{H}^{(0)} - E_n^{(0)}) |\psi_{nj}^{(1)}\rangle + (\hat{H}^{(1)} - E_{nj}^{(1)}) \sum_{i=1}^M |\psi_{ni}^{(0)}\rangle c_{ij}^{(0)} = 0. \quad (4.3)$$

Contracting with some $\langle \psi_{nk}^{(0)} |$ gives

$$\sum_i \langle \psi_{nk}^{(0)} | \hat{H}^{(1)} - E_{nj}^{(1)} | \psi_{ni}^{(0)} \rangle c_{ij} = 0. \quad (4.4)$$

This is a set of $M \times M$ simultaneous equations for $j, k \in \{1, \dots, M\}$, which we may put into the matrix form as

$$(\mathbf{H}^{(1)} - \mathbf{I} E_{nj}^{(1)}) \mathbf{c}_j = 0 \quad (4.5)$$

for $j \in \{1, \dots, M\}$, where $H_{ki}^{(1)} = \langle \psi_{nk}^{(0)} | \hat{H}^{(1)} | \psi_{ni}^{(0)} \rangle$ is the matrix element of $\mathbf{H}^{(1)}$ in the degenerate basis $\{|\psi_{ni}^{(0)}\rangle\}$, and $\mathbf{c}_j = (c_{1j}, \dots, c_{Mj})$. The non-trivial solutions of c_{ij} are given by the secular equation

$$|\mathbf{H}^{(1)} - \mathbf{I} E_{nj}^{(1)}| = 0. \quad (4.6)$$

Therefore, the appropriate zeroth-order wavefunctions are given by the eigenvectors of $H^{(1)}$, with the first-order energies given by the eigenvalues. We would also add the additional constraint

$$\langle \Phi_{nj}^{(0)} | \Phi_{nj}^{(0)} \rangle = \sum_i |c_{ij}|^2 = 1 \quad (4.7)$$

so that the zeroth-order wavefunctions are normalised.

Since $\{|\Phi_{ni}^{(0)}\rangle\}$ diagonalises the perturbation Hamiltonian $H^{(1)}$ in the basis spanned by themselves

$$\langle \Phi_{nk}^{(0)} | \hat{H}^{(1)} | \Phi_{nj}^{(0)} \rangle = \delta_{kj} E_{nj}^{(1)}, \quad (4.8)$$

we claim that the second-order energy for any one member of the degenerate set therefore involves a sum that excluded all members of the degenerate set

$$E_{nj}^{(2)} = - \sum_{i \neq n} \frac{|\langle i | \hat{H}^{(1)} | \Phi_{nj}^{(0)} \rangle|^2}{E_i^{(0)} - E_n^{(0)}}. \quad (4.9)$$

Unfortunately, the expression of the first-order wavefunction is not simple, and it is explained in the appendix section D together with the proof of this second-order energy expression.

Example. Linear Stark effect in the hydrogen atom.

We have examined the response of the 1s orbital energy of an hydrogen atom to a small and uniform electric field. The energy shift is proportional to the square of the electric field in the leading order, so it is known as the *quadratic Stark effect*. We treated it using non-degenerate perturbation theory because the 1s state of hydrogen is non-degenerate.

To analyse the shift of the degenerate 2p energy levels, we must instead use the degenerate perturbation theory. For the perturbation Hamiltonian $\hat{H}^{(1)} = z$, the matrix elements are evaluated as

$$H^{(1)} = \begin{pmatrix} 0 & 0 & 0 & 3 \\ 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 \\ 3 & 0 & 0 & 0 \end{pmatrix}, \quad (4.10)$$

where the only non-zero matrix elements are $\langle 2s | \hat{H}^{(1)} | 2p_z \rangle = \langle 2p_z | \hat{H}^{(1)} | 2s \rangle = 3$. Therefore, the $2p_x$ and $2p_y$ orbitals are unaffected by the electric field to the linear order. The other two perturbed orbitals are $|\Phi_{\pm}^{(0)}\rangle = (|2s\rangle \pm |2p_z\rangle)/\sqrt{2}$, with first-order energies $E_{\pm}^{(1)} = \pm 3$.

The electric fields therefore partially lifts the degeneracy of the hydrogen atom's $n = 2$ states with energies $E_{\pm} = -\frac{1}{8} \pm 3\mathcal{E}$, and leaving two orbitals unaffected to the first order. This is known as the *linear Stark effect*. In a many-electron atom, the 2s and $2p_z$ orbitals are no longer degenerate, and so the Stark effect is again quadratic.

4.2 Nearly Degenerate Perturbation Theory

If we have the state of interest non-degenerate, but very close in energy to some other states, we cannot use non-degenerate perturbation theory because although the individual terms in the Rayleigh–Schrödinger sum do not blow up, the perturbation series may fail to converge even for very small λ as the perturbation gets easily larger than the energy level separation. We need nearly degenerate perturbation theory.

Since we have

$$(\hat{H}^{(0)} - E_n^{(0)}) |\psi_{ni}^{(0)}\rangle = 0, \quad (4.11)$$

we may multiply this expression by c_{ij} and add to the secular equation (4.4) multiplied by λ to get

$$\sum_i \left\langle \psi_{nk}^{(0)} \left| \hat{H}^{(0)} + \lambda \hat{H}^{(1)} - E_n^{(0)} - \lambda E_{nj}^{(1)} \right| \psi_{ni}^{(0)} \right\rangle c_{ij} = 0 \quad (4.12)$$

for degenerate states. Since $\hat{H} = \hat{H}^{(0)} + \lambda \hat{H}^{(1)}$ is the total Hamiltonian, and $E_{nj} = E_n^{(0)} + \lambda E_{nj}^{(1)}$ is the total energy corrected to the first order. For a set of exactly degenerate states, solving the secular equations²⁴

$$(\mathbf{H} - \mathbf{I}E_{nj})\mathbf{c}_j = \mathbf{0} \quad (4.14)$$

using the total Hamiltonian is entirely equivalent to solving for the secular equation for just the perturbing Hamiltonian in the degenerate perturbation theory.

The advantage of this form of secular equation is that it can also be applied when a set of states lie close in energy but are not rigorously degenerate. We can think of this as redefining the $\hat{H}^{(0)}$ and $\hat{H}^{(1)}$ in a way such that the small differences in the energy of the unperturbed states is absorbed into the perturbation, so that the unperturbed states are now somehow degenerate.

Example. Fermi resonance in CO₂.

Under harmonic approximation (2.75), the molecular vibrational potential can be approximated as a sum of quadratic potential, which can be decoupled into a sum of independent Hamiltonians of harmonic potentials (2.55). We will show later that the transition intensity between the initial state $|\psi_i\rangle$ and the final state $|\psi_f\rangle$ is proportional to the squared modulus of the transition dipole moment²⁵

$$\mathbf{R}_{f \leftarrow i} = \langle \psi_f | \hat{\boldsymbol{\mu}} | \psi_i \rangle. \quad (4.15)$$

One can write the dipole moment operator into a sum of the normal coordinates Q_k , which can in turn be written as a sum of the ladder operators \hat{a}_k and \hat{a}_k^\dagger .²⁶ This will show that only transitions between states in which the quantum number of only a single mode changes by one can happen, i.e. $\Delta n_k = \pm 1$.

However, anharmonicity brings interesting complications. Since raising and lowering operators no longer holds for anharmonic states, overtones and combination lines are now allowed. Moreover, the fact that the Hamiltonian itself changes leading to the coupling of normal modes can bring interesting features to the IR and Raman spectra. The harmonic states $|n_1, n_2, \dots\rangle$ which originally diagonalises the harmonic Hamiltonian $\langle n'_1, n'_2, \dots | \hat{H}_{\text{harmonic}} | n_1, n_2, \dots \rangle = \sum_i \omega_i (n_i + \frac{1}{2}) \delta_{n_1 n'_1} \delta_{n_2 n'_2} \dots$ no longer does so. This effect can be particularly strong if the two states with the same symmetry happens to be close in energy. We will investigate this effect using nearly degenerate perturbation theory.

The classic example is CO₂. We will take the harmonic approximation as the reference Hamiltonian. In this case, the molecule has 4 normal modes: a symmetric stretch (Q_1) of symmetry Σ_g^+ , a doubly-degenerate bend (Q_{2x}, Q_{2y}) of symmetry Π and an antisymmetric stretch (Q_3) of symmetry Σ_u^- . We label a vibrational state by the three quantum numbers $|n_1, n_2, n_3\rangle$, where $n_2 = n_{2x} + n_{2y}$. Denoting $Q_2^2 = Q_{2x}^2 + Q_{2y}^2$, the unperturbed Hamiltonian and energy are

$$\hat{H}^{(0)} = -\frac{1}{2} \left(\frac{\partial^2}{\partial Q_1^2} + \frac{\partial^2}{\partial Q_{2x}^2} + \frac{\partial^2}{\partial Q_{2y}^2} + \frac{\partial^2}{\partial Q_3^2} \right) + \frac{1}{2} (\omega_1^2 Q_1^2 + \omega_2^2 Q_2^2 + \omega_3^2 Q_3^2), \quad (4.16)$$

$$E_{n_1, n_2, n_3}^{(0)} = \left(n_1 + \frac{1}{2} \right) \omega_1 + \left(n_2 + \frac{1}{2} \right) \omega_2 + \left(n_3 + \frac{1}{2} \right) \omega_3. \quad (4.17)$$

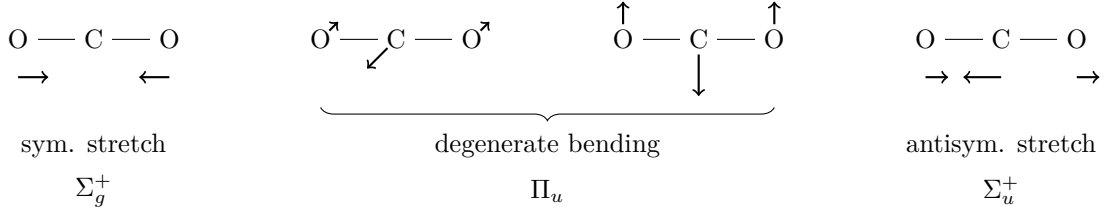
²⁴This is essentially the Schrödinger's equation

$$\mathbf{H}\mathbf{c}_j = E_{nj}\mathbf{c}_j \quad (4.13)$$

in the confined basis set of the nearly degenerate orbitals using the full Hamiltonian, so we should be convinced that this equation is appropriate no matter the states is degenerate or not.

²⁵See time-dependent perturbation theory later.

²⁶One can first integrate out the common electronic wavefunction to work out $\hat{\boldsymbol{\mu}}(\mathbf{Q})$ as a function of nuclear coordinates only, then expand it around the equilibrium nuclear geometry. See *B8: Symmetry* for detail.



The symmetric stretch and bend frequencies are $\omega_1 = 1354.07 \text{ cm}^{-1}$ and $\omega_2 = 672.95 \text{ cm}^{-1}$ respectively. Relative to the ground state, the doubly-excited bend $|020\rangle$ at energy $2\omega_2 = 1345.90 \text{ cm}^{-1}$, lying just $\Delta = 8.17 \text{ cm}^{-1}$ below the $|100\rangle$ state. Moreover, the symmetry of the $|020\rangle$ states are given by $\Pi \otimes \Pi = \Sigma_g^+ + [\Sigma_u^+] + \Pi_g$, so the component of $|020\rangle$ with zero angular momentum,

$$|02^0 0\rangle = \frac{1}{\sqrt{2}}(|02_x 0\rangle + |02_y 0\rangle) \quad (4.18)$$

has the correct symmetry to interact with $|100\rangle$ if there exists appropriate coupling perturbation.

We will add cubic potentials as perturbation. Since the Hamiltonian must be totally symmetric, not all cubic potentials $Q_i Q_j Q_k$ are permitted to present. In turns out that the only allowed terms in the cubic potential are

$$\hat{H}^{(1)} = aQ_1^3 + bQ_1Q_2^2 + cQ_1Q_3^2. \quad (4.19)$$

Using the nearly degenerate perturbation theory, we can see that this perturbing Hamiltonian indeed couples these two states. The total Hamiltonian matrix in the basis of the two states are given by

$$\mathbf{H} = \begin{pmatrix} \omega_1 & W \\ W & 2\omega_2 \end{pmatrix}, \quad (4.20)$$

where W is the *Fermi coupling constant* given by

$$W = \langle 100 | \hat{H}^{(0)} + \hat{H}^{(1)} | 02^0 0 \rangle = \langle 100 | bQ_1Q_2^2 | 02^0 0 \rangle. \quad (4.21)$$

This coupling is due to cubic term the $Q_1Q_2^2$. We can again use ladder operators

$$Q_i = \frac{1}{\sqrt{2\omega_i}}(\hat{a}_i + \hat{a}_i^\dagger) \quad (4.22)$$

to work this out. We write

$$\langle 100 | bQ_1Q_2^2 | 02^0 0 \rangle = b \langle 1 | Q_1 | 0 \rangle \langle 0 | Q_2^2 | 2 \rangle \langle 0 | 0 \rangle. \quad (4.23)$$

The first term is easy to evaluate

$$\langle 1 | Q_1 | 0 \rangle = \frac{1}{\sqrt{2\omega_1}} \langle 1 | \hat{a} + \hat{a}^\dagger | 0 \rangle = \frac{1}{2\omega_1}, \quad (4.24)$$

but the second degenerate mode is a bit more subtle. We need to expand it into its components

$$\begin{aligned} \langle 0 | Q_2^2 | 2^0 \rangle &= \frac{1}{\sqrt{2}} (\langle 0_x, 0_y | Q_{2x}^2 | 2_x, 0_y \rangle + \langle 0_x, 0_y | Q_{2y}^2 | 0_x, 2_y \rangle) \\ &= \sqrt{2} \langle 0 | Q_{2x}^2 | 2_x \rangle \end{aligned} \quad (4.25)$$

as the two components are equivalent. Then

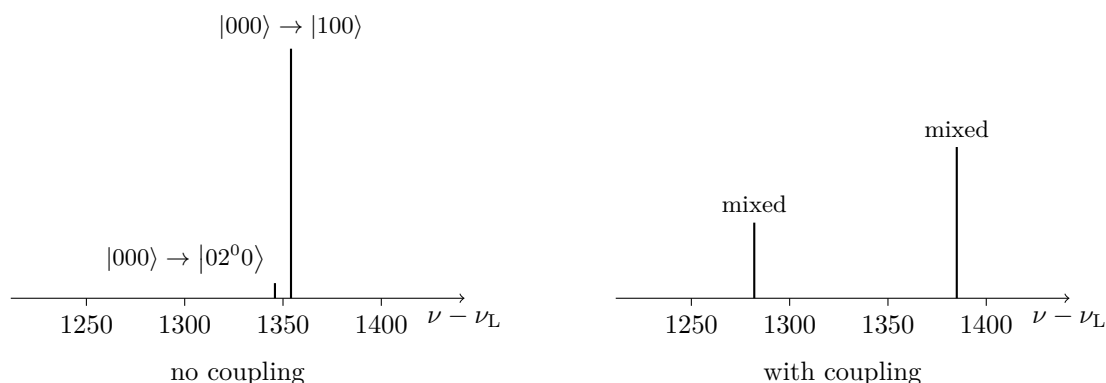
$$\begin{aligned} \sqrt{2} \langle 0 | Q_{2x}^2 | 2_x \rangle &= \frac{1}{\sqrt{2\omega_2}} \langle 0 | \hat{a}^2 | 2 \rangle \\ &= \frac{1}{\omega_2}. \end{aligned} \quad (4.26)$$

Combining the results gives

$$W = \frac{b}{\sqrt{2\omega_1\omega_2}}. \quad (4.27)$$

Solving the secular equation, we find that the perturbed states are separated by $\sqrt{4W^2 + \Delta^2}$, which is clearly bigger than the original separation Δ . This effect is known as the *Fermi resonance*. The two lines are observed to be 102.78 cm^{-1} apart in the Raman spectrum, implying $|W| = 51.2 \text{ cm}^{-1}$ — a rather large coupling.

The mixing of the vibrational states also means that the $|000\rangle \rightarrow |100\rangle$ and $|000\rangle \rightarrow |02^0_0\rangle$ transitions are no longer pure. Consequently, the latter transition, which would normally be a weak overtone, borrows intensity from the strong fundamental symmetric stretch. The pair of mixed levels are known as a *Fermi dyad*.



Example. Nearly free electron gas.

You should have met the concept of nearly free electron gas model for solids in Part IB Chemistry A. The idea of degenerate perturbation theory gives us some more insight on it.

For our electron gas to be ‘nearly free’, let’s consider the situation where a particle moves in a region having a weakly periodic potential. By weakly periodic, we mean that the free energy of the particle is much greater than the periodic potential present: the particle floats well above a corrugated base. For example, let’s assume that the underlying potential has the form

$$V(x) = 2V \cos\left(\frac{2\pi x}{a}\right) = V(e^{iGx} + e^{-iGx}), \quad (4.28)$$

where $G = 2\pi/a$ characterises the periodicity of the lattice in the form of a wavevector. Suppose the system has total length $L = Na$, so it comprises N cells of size a . For a real periodic potential in a periodic crystal lattice, we would replace this by a Fourier series, but let’s now focus on this simple case.

The fact that the periodic potential is weak makes it perfect for perturbation theory. We take the unperturbed Hamiltonian to be the free electron gas

$$\hat{H}^{(0)} = -\frac{\hbar^2}{2m} \frac{\partial^2}{\partial x^2}, \quad (4.29)$$

and the unperturbed wavefunction is

$$\psi_k^{(0)} = \frac{1}{\sqrt{L}} e^{ikx} \quad (4.30)$$

with unperturbed energy

$$E_k^{(0)} = \frac{\hbar^2 k^2}{2m}. \quad (4.31)$$

We impose the periodic boundary condition: $\psi(x) = \psi(x + L)$ so k is discretised

$$k = \frac{2\pi n}{L}, \quad n \in \mathbb{Z}. \quad (4.32)$$

The perturbing Hamiltonian is our periodic potential

$$\hat{H}^{(1)} = 2V \cos\left(\frac{2\pi x}{a}\right). \quad (4.33)$$

To proceed, we would like to find the matrix elements of the perturbation

$$\begin{aligned} \langle k | \hat{H}^{(1)} | k' \rangle &= \frac{2V}{L} \int_0^L dx e^{i(k' - k)x} \cos\left(\frac{2\pi x}{a}\right) \\ &= \frac{V}{L} \int_0^L dx e^{i(k' - k + G)x} + e^{i(k' - k - G)x} \\ &= V[\delta_{k - k', -G} + \delta_{k - k', +G}]. \end{aligned} \quad (4.34)$$

Therefore, the only terms that survive are those for which

$$k' - k = \pm G = \pm \frac{2\pi}{a}, \quad (4.35)$$

or

$$n' - n = \pm n. \quad (4.36)$$

State $|k\rangle$ will only interact with states $|k + G\rangle$ and $|k - G\rangle$.

In general, the states $|k\rangle$ and $|k'\rangle$ are non-degenerate, and the new energies can be calculated by non-degenerate perturbation theory

$$\begin{aligned} E_k &= E_k^{(0)} + \langle k | \hat{H}^{(1)} | k \rangle + \sum_{k' \neq k} \frac{|\langle k' | \hat{H}^{(1)} | k \rangle|^2}{E_k^{(0)} - E_{k'}^{(0)}} \\ &= E_k^{(0)} + \frac{|V|^2}{E_k^{(0)} - E_{k+G}^{(0)}} + \frac{|V|^2}{E_k^{(0)} - E_{k-G}^{(0)}}. \end{aligned} \quad (4.37)$$

However, the above method will break down in the neighbourhood of $k \approx \pm \frac{\pi}{a}$, since by then, $E_k^{(0)} \approx E_{k+G}^{(0)}$. We would then need to use nearly degenerate perturbation theory. For example, if we are interested in a state with $k \approx +\frac{\pi}{a}$, which would interact with a nearly-degenerate state at $k' = k - \frac{2\pi}{a} \approx -\frac{\pi}{a}$, then the appropriate Hamiltonian matrix to diagonalise is

$$H = \begin{pmatrix} E_k & V \\ V & E_{k'} \end{pmatrix}, \quad (4.38)$$

giving first-order energies

$$E = \frac{E_k + E_{k'}}{2} \pm \frac{1}{2} \sqrt{(E_k - E_{k'})^2 + 4V}. \quad (4.39)$$

The two levels are now separated by a gap larger than their original separation:

$$\sqrt{(E_k - E_{k'})^2 + 4V} > (E_k - E_{k'}). \quad (4.40)$$

If we consider $k = +\frac{\pi}{a}$ and $k' = -\frac{\pi}{a}$, so the two interacting states are exactly degenerate, then the perturbation will result in a gap of $2V$. A band gap of $2V$ is opened at $k = \pm \frac{\pi}{a}$.

For a real periodic potential that can be written in a Fourier series

$$V(x) = \sum_{m=1}^{\infty} V_m \cos\left(\frac{m\pi x}{V}\right), \quad (4.41)$$

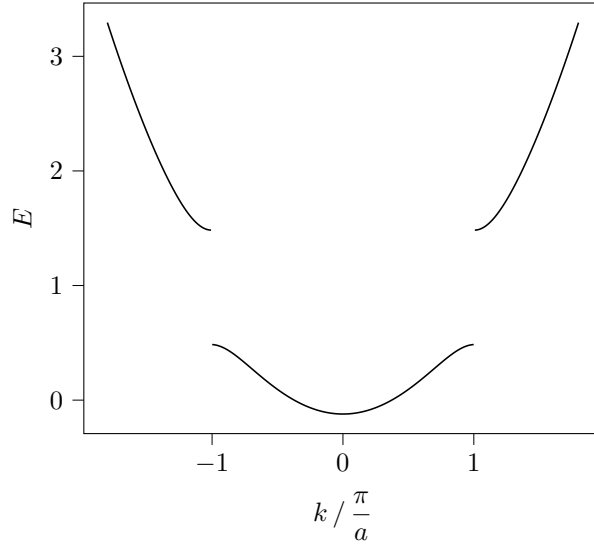


Figure 8: The dispersion curve of a nearly free electron gas model calculated using nearly degenerate perturbation theory, with V set to be 0.5 in arbitrary unit. A band gap of 1 is opened at $k = \pm \frac{\pi}{a}$.

the term $\cos(m\pi x/a) = e^{+imGx} + e^{-imGx}$ will lead to the interaction between $|k\rangle$ and $|k \pm mG\rangle$, and hence band gaps will open at $\pm \frac{m\pi}{a}$. For a most general periodic potential, $V_m \neq 0$ for all m , so band gaps are open at all positive integer multiples of $\pm \frac{\pi}{a}$, leading to the band structure you met in IB Chemistry A.

5 Time-Dependent Systems

5.1 Time-Dependent Schrödinger Equation

For a time-dependent system, we have to turn to consider the time-dependent Schrödinger equation

$$\hat{H}\Psi(\mathbf{x}, t) = i\hbar \frac{\partial \Psi(\mathbf{x}, t)}{\partial t}. \quad (5.1)$$

The wavefunction is now a function of time as well as the spatial coordinates. We will first focus on the special case that the Hamiltonian is independent on time. This is then a straightforward first-order differential equation in time, with solutions given by

$$\Psi(\mathbf{x}, t) = e^{-i\hat{H}t/\hbar} \Psi(\mathbf{x}, 0). \quad (5.2)$$

$\Psi(\mathbf{x}, 0)$, the wavefunction at fixed time $t = 0$, solves the time-independent Schrödinger equation

$$\hat{H}\Psi(\mathbf{x}, 0) = E_k \Psi(\mathbf{x}, 0), \quad (5.3)$$

where it is common to denote the time-independent wavefunction as $\psi_k(\mathbf{x}) \equiv \Psi(\mathbf{x}, 0)$. $e^{-i\hat{H}t/\hbar}$ is the *time-evolution operator*, which is unitary. The exponential of the operator is defined via the Maclaurin series

$$\begin{aligned} e^{-i\hat{H}t/\hbar} |\psi_k\rangle &= 1 - i\frac{t}{\hbar} \hat{H} |\psi_k\rangle - \frac{t^2}{2\hbar^2} \hat{H}^2 |\psi_k\rangle + i\frac{t^3}{6\hbar^3} \hat{H}^3 |\psi_k\rangle + \dots \\ &= 1 - i\frac{t}{\hbar} E_k |\psi_k\rangle - \frac{t^2}{2\hbar^2} E_k^2 |\psi_k\rangle + i\frac{t^3}{6\hbar^3} E_k^3 |\psi_k\rangle \\ &= e^{-iE_k t/\hbar} |\psi_k\rangle. \end{aligned} \quad (5.4)$$

Therefore, for a time-independent Hamiltonian, there is a special class of solutions that can be written as

$$\Psi(\mathbf{x}, t) = e^{-iE_k t/\hbar} \psi_k(\mathbf{x}). \quad (5.5)$$

These are known as *stationary states* because it is just a time-independent state multiplied by a phase factor rotating with time. None of the physical observables will change with time, as one might expect by the action of a unitary operator.

From now on, we will adopt atomic units, so all \hbar 's are gone from our expressions.

5.2 Ehrenfest Theorem

The Ehrenfest theorem concerns the time evolution of expectation values of physical observable. We begin with

$$\begin{aligned} \frac{d}{dt} \langle A \rangle &= \frac{d}{dt} \langle \Psi | \hat{A} | \Psi \rangle \\ &= \left\langle \frac{\partial \Psi}{\partial t} | \hat{A} | \Psi \right\rangle + \left\langle \Psi | \frac{\partial \hat{A}}{\partial t} | \Psi \right\rangle + \left\langle \Psi | \hat{A} | \frac{\partial \Psi}{\partial t} \right\rangle. \end{aligned} \quad (5.6)$$

We can use the time-dependent Schrödinger equation to substitute for the time derivatives of the wavefunction

$$\begin{aligned}
\frac{d}{dt} \langle A \rangle &= \langle -i\hat{H}\Psi | \hat{A} | \Psi \rangle + \left\langle \Psi \left| \frac{\partial \hat{A}}{\partial t} \right| \Psi \right\rangle + \langle \Psi | \hat{A} | -i\hat{H}\Psi \rangle \\
&= i \langle \Psi | \hat{H}\hat{A} | \Psi \rangle + \left\langle \frac{\partial \hat{A}}{\partial t} \right\rangle - i \langle \Psi | \hat{A}\hat{H} | \Psi \rangle \\
&= -i \langle [\hat{A}, \hat{H}] \rangle + \left\langle \frac{\partial \hat{A}}{\partial t} \right\rangle .
\end{aligned} \tag{5.7}$$

This result is general, but the Ehrenfest theorem relates to the specific case of position and momentum.

Theorem 5.1 (Ehrenfest theorem).

$$m \frac{d}{dt} \langle x \rangle = \langle p \rangle \tag{5.8}$$

$$\frac{d}{dt} \langle p \rangle = - \left\langle \frac{dV(x)}{dx} \right\rangle . \tag{5.9}$$

They are the quantum equivalents of the familiar classical expressions

$$p = mv \tag{5.10}$$

$$F = ma , \tag{5.11}$$

but they are not quite the same. To match up with the classical expression exactly, we would need

$$\frac{d}{dt} \langle p \rangle \stackrel{?}{=} - \frac{dV(\langle x \rangle)}{dx} , \tag{5.12}$$

but this would imply

$$\left\langle \frac{dV(x)}{dx} \right\rangle \stackrel{?}{=} \frac{dV(\langle x \rangle)}{dx} , \tag{5.13}$$

which is not in general true. For example, if we set $V(x) = x^3$, then this equation implies $\langle x^2 \rangle \stackrel{?}{=} \langle x \rangle^2$. The difference in them is related to Δx , so if we are approaching the classical limit, $\Delta x \rightarrow 0$, then the Ehrenfest equations would approach the Newton's equations. This is an example of the correspondence principle.

It is possible to go the other way around and derive the time-dependent Schrödinger equation from the Ehrenfest theorem, so the Ehrenfest theorem is just as fundamental as the Schrödinger equation and could be used as the starting point for quantum mechanics.

5.3 Time-Dependent Perturbation Theory

We will consider the time-dependent Hamiltonian of the general form

$$\hat{H}(\mathbf{x}, t) = \hat{H}^{(0)}(\mathbf{x}) + \lambda \hat{H}^{(1)}(\mathbf{x}, t) , \tag{5.14}$$

where the unperturbed Hamiltonian is time-independent and we know its eigenstates $\psi_k(\mathbf{x})$ and eigenvalues E_k exactly. The time dependence appears only in the perturbation $\hat{H}^{(1)}$, which is ‘small’. In the absence of the perturbation, there are stationary solutions

$$|\Psi_k(t)\rangle = e^{-iE_k t} |\psi_k\rangle , \tag{5.15}$$

where

$$\hat{H}^{(0)} |\psi_k\rangle = E_k |\psi_k\rangle . \tag{5.16}$$

We have dropped the superscript (0) on ψ_k and E_k because the stationary state is necessarily unperturbed.

As in the Rayleigh–Schrödinger perturbation theory, we now expand the perturbed wavefunction in terms of the unperturbed state. In the method developed by Dirac, known as the variation of constants,²⁷ the coefficient depend on time

$$|\Psi(t)\rangle = \sum_k a_k(t) |\Psi_k(t)\rangle = \sum_k a_k(t) e^{-iE_k t} |\psi_k\rangle. \quad (5.17)$$

We must be able to do this because $\{|\psi_k\rangle\}$ is a complete orthonormal basis, so we can expand $|\Psi(t)\rangle$ at each time.

There is no subscript to label the state, since the state is changing in time. Moreover, the squared modulus of the coefficient is the probability of observing the system in the stationary state k with energy E_k at time t .²⁸ Substituting into the time-dependent Schrödinger equation, we get

$$\begin{aligned} \sum_k a_k \left(E_k + \lambda \hat{H}^{(1)} \right) |\Psi_k\rangle &= i \sum_k \left(\frac{da_k}{dt} - i a_k E_k \right) |\Psi_k\rangle \\ \lambda \sum_k \hat{H}^{(1)} a_k |\Psi_k\rangle &= i \sum_k \frac{da_k}{dt} |\Psi_k\rangle. \end{aligned} \quad (5.18)$$

To work out the contribution of a particular $|\Psi_j\rangle$, we contract it with $\langle\Psi_j| = e^{+iE_j t} \langle\psi_j|$ and get

$$\lambda \sum_k a_k e^{-i(E_k - E_j)t} \langle j | \hat{H}^{(1)} | k \rangle = i \sum_k \frac{da_k}{dt} e^{-i(E_k - E_j)t} \langle j | k \rangle, \quad (5.19)$$

and by orthogonality of unperturbed wavefunction, we get

$$\frac{da_j}{dt} = -i\lambda \sum_k \hat{H}_{jk}^{(1)}(t) a_k(t) e^{i\omega_{jk}t}, \quad (5.20)$$

where $\omega_{jk} = E_j - E_k$. This is a coupled system of differential equation, telling us how the weights of various states evolve with time. The rate of change of one weight depends on the current values of all the weights.

To break the interdependency of the coefficients, we must make approximations. We expand the coefficients $a_k(t)$ in a power series of λ

$$a_k(t) = a_k^{(0)}(0) + \lambda a_k^{(1)}(t) + \lambda^2 a_k^{(2)}(t) + \dots \quad (5.21)$$

Note that the unperturbed system is time-independent, so $a_k^{(0)}$ does not depend on time. We can then solve for each power in λ individually. Note the presence of λ on the right hand side of (5.20) means that the dependence of first-order terms is given purely by the time-independent zeroth-order terms. Therefore we have successfully decoupled the time dependence of the coefficients. We have

$$\frac{da_j^{(0)}}{dt} = 0 \quad (5.22)$$

$$\frac{da_j^{(1)}}{dt} = -i \sum_k a_k^{(0)}(0) \hat{H}_{jk}^{(1)}(t) e^{i\omega_{jk}t} \quad (5.23)$$

⋮

²⁷This is formally known as the *interaction picture*, or the *Dirac picture*. We will not introduce it in the main text, because we can work out the first-order time-dependent perturbation theory without using this formal framework. However, this is very useful if we want to go to higher orders to explain things like Raman scattering, so it is introduced in the appendix, together with the other two pictures: the Schrödinger picture and the Heisenberg picture.

²⁸Technically the stationary state Ψ_k is the energy eigenstate of the system only if perturbing Hamiltonian is not present, so we should remove it before measuring the energy if we are interested in the question ‘which state of Ψ_k has our system evolved into’.

The zeroth-order equation reassures that the a_j is unchanging because $\hat{H}^{(0)}$ is time-independent. The first-order equation is our main interest, telling us how the state is evolving to the leading order.

If we now suppose that at time $t = 0$, the system is known to be in one of the stationary state $|\psi_n\rangle$, such that $a_k^{(0)} = \delta_{nk}$, we can remove the sum and get

$$\frac{da_j}{dt} = -iH_{jn}^{(1)}(t)e^{i\omega_{jn}t}, \quad (5.24)$$

where we dropped the superscript because we are only interested in the first-order contribution, so this is the rate of change of our total a_k . We may integrate with respect to time and get

$$a_j(t) = -i \int_0^t dt' H_{jn}^{(1)}(t') e^{i\omega_{jn}t'}. \quad (5.25)$$

Heuristically, this amounts to considering only direct transitions from initial state $|n\rangle$ to the state $|j\rangle$ of interest. Excitations that proceed via intermediate states are neglected. We therefore derived a first-order result, where the perturbation is only allowed to act once. A second-order expression can be obtained by substituting (5.25) back to (5.20), but we will do it using the interaction picture in the appendix section E.

Example. A slowly-switched-on perturbation.

Let's first investigate the response of a system to a perturbation that is gradually switched on and approaching a constant value. A possible form of this can be

$$\hat{H}^{(1)}(\mathbf{x}, t) = (1 - e^{-kt})\hat{V}(\mathbf{x}) \quad (5.26)$$

for some time-independent perturbation $\hat{V}(\mathbf{x})$. Suppose we are initially in a state $|n\rangle$, then the coefficient of state $|j\rangle$ is

$$\begin{aligned} a_j(t) &= -iV_{jn} \int_0^t dt' (1 - e^{-kt'}) e^{i\omega_{jn}t'} \\ &= -iV_{jn} \left(\frac{e^{i\omega_{jn}t} - 1}{i\omega_{jn}} - \frac{e^{(i\omega_{jn}-k)t} - 1}{i\omega_{jn} - k} \right), \end{aligned} \quad (5.27)$$

where $V_{jn} = \langle j | \hat{V} | n \rangle$. After some non-trivial amount of algebra, one may show that the probability of transition from state $|n\rangle$ to $|j\rangle$, $|a_j(t)|^2$, is given by

$$|a_j(t)|^2 = \frac{V_{jn}^2}{\omega_{jn}^2} \frac{[\omega_{jn}(1 - e^{-kt}) - k \sin \omega_{jn}t]^2 + k^2(1 - \cos \omega_{jn}t)^2}{k^2 + \omega_{jn}^2}. \quad (5.28)$$

This is not a tidy expression, and we will have a brief analysis of this later. We will now focus on the long-time behaviour limit, $kt \gg 1$, and suppose that the switching is very slow in comparison with the relevant frequency offset, $k \ll \omega_{jn}$. We then have

$$a_j(t) = -iV_{jn} \frac{e^{i\omega_{jn}t}}{i\omega_{jn}} = -\frac{V_{jn}}{E_j - E_n} e^{i\omega t}. \quad (5.29)$$

Evidently, this result cannot be applied to the initial state $|n\rangle$, but we argue that since the perturbation is small, the transition is negligible and we will have $a_n(t) = a_n(0) = 1$ to the leading order. The complete wavefunction is

$$\begin{aligned} \Psi(t) &= \Psi_n(t) + \sum_{j \neq n} a_j(t) \Psi_j(t) \\ &= |n\rangle e^{-iE_n t} - \sum_{j \neq n} \frac{V_{jn}}{E_j - E_n} e^{i\omega_{jn}t} |j\rangle e^{-iE_j t} \\ &= \left(|n\rangle - \sum_{j \neq n} \frac{V_{jn}}{E_j - E_n} |j\rangle \right) e^{-iE_n t}. \end{aligned} \quad (5.30)$$

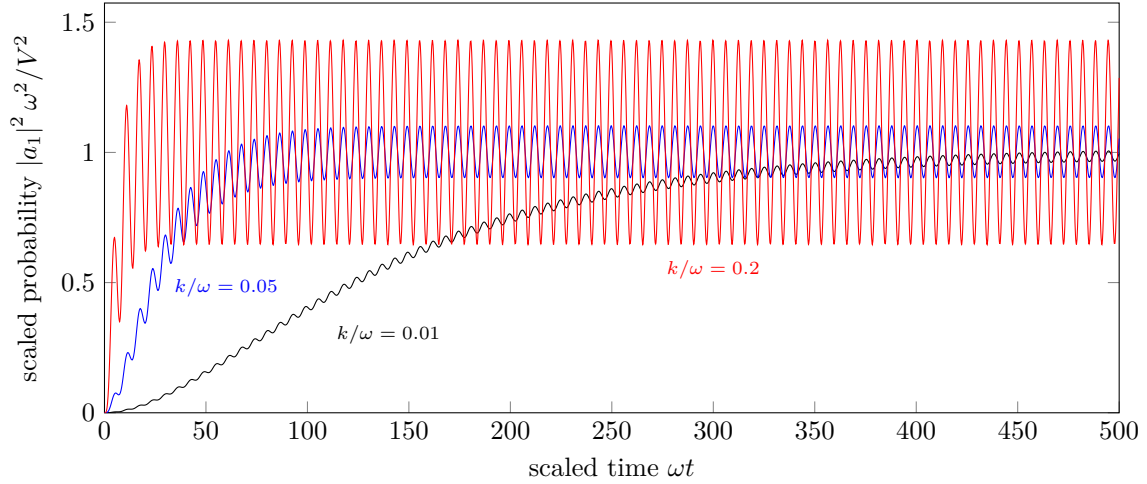


Figure 9: Scaled probability of transition to excited state in a two-level system when a perturbation is switched on at different rates.

We observe that this is exactly the Rayleigh–Schrödinger sum of the first-order wavefunction multiplied by a time-dependent phase factor of constant modulus. Therefore, in the long-time limit,

$$\Psi(t) = (\psi_n^{(0)} + \psi_n^{(1)})e^{-iE_n t}, \quad (5.31)$$

which is just the first-order time-independent perturbation result multiplied with its phase factor.

This justifies the application of time-independent perturbation theory to the study of external perturbations to a system, provided that the perturbation is switched on sufficiently slowly. The system remains in the state that can be properly labelled as n described by the time-independent perturbation theory. We say that such a system is *adiabatic*, standing for slowly varying. This is the *adiabatic theorem* due to Max Born and Vladimir Fock.

Theorem 5.2 (Adiabatic theorem). A physical system remains in its instantaneous eigenstate if a given perturbation is acting on it slowly enough and if there is a gap between the eigenvalue and the rest of the Hamiltonian’s spectrum.

This is closely linked to the adiabatic (Born–Oppenheimer) approximation, in which since the motion of the nuclei is varying slowly with respect to electrons, the electrons can be thought of as in the electronic eigenstates of the instantaneous nuclei geometry at each instant.

If k is not sufficiently small compared to ω then the switching cannot be regarded as adiabatically slow, and we must use the more complex form of the transition probability (5.28). We consider a two-level system with $E_0 = 0$ and $E_1 = \omega$, initially residing in the ground state, and a perturbation $V_{01} = V$ is turned on with rate k . The transition probability is plotted in scaled time ωt and is normalised by the adiabatic long-time limit V^2/ω^2 .

We see that switching at a finite rate induces oscillations in the occupation of the upper state, and the magnitude of the oscillation increases with the switching rate. The oscillations continue even after their mean has settled down to its final value. If you are careful enough, you can see that the mean does not coincide with the adiabatic limit.²⁹

²⁹You may replace all the oscillatory terms (\sin and \cos) with 0 in (5.28), and get

$$\frac{k^2 + \omega^2(1 - e^{-kt})^2}{k^2 + \omega^2}, \quad (5.32)$$

which does tends to 1 as $t \rightarrow \infty$ regardless of k . However, oscillatory terms do shift the mean value because it is squared.

Example. Fermi's golden rule.

We now study the interaction of oscillating electromagnetic radiation with atoms or molecules. We will investigate the stimulated absorption and stimulated emission, which should be familiar from *A3: High Resolution Molecular Spectroscopy*.

The most general form of the perturbing Hamiltonian should be

$$\hat{H}^{(1)}(t) = \hat{V}e^{i\omega t} + \hat{V}^\dagger e^{-i\omega t} \quad (5.33)$$

for some time-independent operator \hat{V} . We let \hat{V} to be Hermitian so that

$$\hat{H}^{(1)} = 2\hat{V} \cos \omega t. \quad (5.34)$$

We insert this into (5.25) and we will get

$$a_j(t) = -V_{jn} \left(\frac{e^{i(\omega_{jn}+\omega)t} - 1}{\omega_{jn} + \omega} + \frac{e^{i(\omega_{jn}-\omega)t} - 1}{\omega_{jn} - \omega} \right). \quad (5.35)$$

We can see that if ω_{jn} is positive ($|j\rangle$ above $|n\rangle$) and ω is close to ω_{jn} , then the second fraction will dominate. This corresponds to a *spontaneous absorption* process, in which $|a_{jn}|^2$ described the probability of the molecule moving from $|n\rangle$ to a higher energy state $|j\rangle$. If ω_{jn} is negative and is close to $-\omega$, then the first term will dominate, and this corresponds to *spontaneous emission*.

We will focus on absorption and ignore the first term. The probability of finding the system in $|j\rangle$ after time t is

$$P_j(t) = |a_j(t)|^2 = \frac{4|V_{jn}|^2}{(\omega_{jn} - \omega)^2} \sin^2 \frac{(\omega_{jn} - \omega)t}{2}. \quad (5.36)$$

In practise, the radiation is never monochromatic. We will describe this by the frequency density of states $\rho(\omega)$ so that $\rho(\omega)d\omega$ is the number of photons with frequency between ω and $\omega + d\omega$. The probability of observing the system in $|j\rangle$ after time t is therefore the integral of (5.36) weighted by the density $\rho(\omega)$:

$$\begin{aligned} P_j(t) &= \int d\omega \frac{4|V_{jn}|^2}{(\omega_{jn} - \omega)^2} \sin^2 \frac{(\omega_{jn} - \omega)t}{2} \rho(\omega) \\ &= \int d\omega \frac{\sin^2 \frac{1}{2}\Omega t}{\Omega^2 t} 4|V_{jn}|^2 \rho(\omega) t, \end{aligned} \quad (5.37)$$

where $\Omega = \omega_{jn} - \omega$. We see an interesting function: $\sin^2 \frac{1}{2}\Omega t / \Omega^2 t$. Viewing it as a function of Ω , and t is a parameter, we see that it is sharply peaked at $\Omega = 0$, showing that only photons with frequency ω close to ω_{jn} contributes significantly to the transition. Moreover, as t increases, the function gets more and more sharply peaked, and in fact, in the $t \rightarrow \infty$ limit, it tends to $\frac{1}{2}\pi\delta(\Omega)$.

Therefore, in the $t \rightarrow \infty$ limit, we can write

$$\begin{aligned} P_j(t) &= \int d\omega 2\pi\delta(\omega_{jn} - \omega) |V_{jn}|^2 t \rho(\omega) \\ &= 2\pi |V_{jn}|^2 \rho(\omega_{jn}) t. \end{aligned} \quad (5.38)$$

This is linear in time. The transition rate from $|n\rangle$ to $|j\rangle$ is

$$R_{nj} = \frac{dP_j(t)}{dt} = 2\pi |V_{jn}|^2 \rho(\omega_{jn}), \quad (5.39)$$

or if we are not using atomic units,

$$R_{nj} = \frac{2\pi}{\hbar^2} |V_{jn}|^2 \rho(\omega_{jn}). \quad (5.40)$$

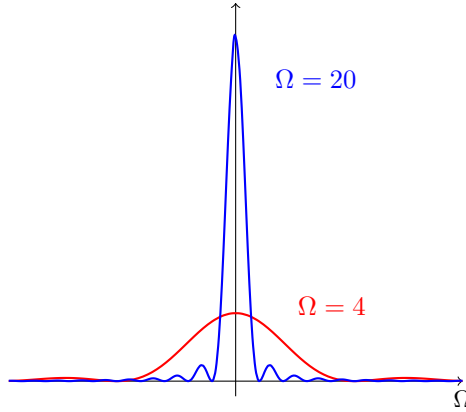


Figure 10: A plot of $\sin^2 \frac{1}{2} \Omega t / \Omega^2 t$ as a function of Ω . It gets more and more sharply peaked as $t \rightarrow \infty$.

This is the *Fermi's golden rule*.

To move further, we must consider the nature of the \hat{V} operator. For our purpose, it should describe the interaction between the molecule and the oscillating electric field. The ‘oscillating’ part is already dealt with by the \cos term in (5.34), so we need \hat{V} to be a time-independent operator describing the effect of a electric field of some polarisation (direction) and magnitude, represented by a vector \mathcal{E} . We will make some further assumptions:

- The field is continuous. Rigorously, the electromagnetic wave should be quantised into packages of photons, and this is necessary if we want to investigate effects like spontaneous emission. However, this is the realm of quantum field theory, and is far beyond the scope of this course.³⁰
- We assume the amplitude of the field is constant over the molecule. This is known as the *dipole approximation*, and it holds as long as the wavelength of the photon is significantly larger than the size of the molecule, and so it is fine up to UV-Vis spectroscopy. It will be problematic if we are interested in ionisations by X-rays.
- We assume that the field is isotropic, so the polarisation can have any random orientation with respect to the molecule.

With in this approximation, we can write the perturbing Hamiltonian as

$$\hat{H}^{(1)} = -\mathcal{E} \cdot \hat{\mu} \cos(\omega t), \quad (5.41)$$

and so

$$\hat{V} = -\frac{1}{2} \mathcal{E} \cdot \hat{\mu} = -\frac{1}{2} \mathcal{E} \hat{\mu} \cos \theta, \quad (5.42)$$

where θ is the direction between the polarising plane of the electromagnetic wave and the dipole moment of the molecule. Note the factor of $\frac{1}{2}$ comes from (5.34). Averaging over the direction θ , we get

$$R_{nj} = \frac{\pi}{2\hbar^2} |\langle j | \mathcal{E} \cdot \hat{\mu} | n \rangle|^2 \rho(\omega_{jn}) \langle \cos^2 \theta \rangle, \quad (5.43)$$

where

$$\langle \cos^2 \theta \rangle = \frac{\int_0^{2\pi} d\phi \int_0^\pi d\theta \sin \theta \cos^2 \theta}{\int_0^{2\pi} d\phi \int_0^\pi d\theta \sin \theta} = \frac{1}{3}, \quad (5.44)$$

³⁰It is fluctuations in the zero-point value of the electromagnetic field that allow the atom to spontaneously emit a photon and decay; heuristically, the fluctuating value of the quantum electromagnetic field — even in the vacuum — ‘tickles’ the excited atom prompting the decay. Treating the electromagnetic field classically is appropriate if the radiation comes from a high intensity laser, where the energy density of the field is so high that the stimulated emissions and absorptions dominate. By contrast, emission of light from the atoms in a humble candle is an inherently quantum phenomenon, occurring purely by spontaneous emission once the flame is lit — candlelight shines even in ambient darkness.

and so

$$R_{nj} = \frac{\pi}{6\hbar^2} \mathcal{E}^2 |\langle j|\hat{\mu}|n\rangle|^2 \rho(\omega_{jn}). \quad (5.45)$$

The transition dipole moment can only be non-zero if its integrand, $\psi_j^* \hat{\mu} \psi_n$ is totally symmetric under the operations of the molecule's point group. This requirement restricts the combinations of states between which transitions may be observed, and is the origin of spectroscopic selection rules.

We have given the transition rate for an individual particle, but we can easily adjust to giving the rate of change of the number of particles in the initial state for a collection of non-interacting particles by multiplying the concentration of particles,

$$\frac{dn_n}{dt} = -\frac{\pi}{6\hbar^2} \mathcal{E}^2 |\langle j|\hat{\mu}|n\rangle|^2 \rho(\omega_{jn}) n_n. \quad (5.46)$$

This is not quite the same as the expression we saw in A3, this is because in there instead used the energy density of photons, $\rho'(\nu)$, such that $\int_{\nu_1}^{\nu_2} d\nu \rho'(\nu)$ is the energy per unit volume of photons in the frequency range ν_1 to ν_2 . Here, we defined $\rho(\omega)$ to be such that $\int_{\omega_1}^{\omega_2} d\omega \rho(\omega)$ is the number of particles in the frequency range ω_1 to ω_2 . Therefore, we need the converting factor

$$\rho(\omega) = \frac{V}{2\pi\hbar\omega} \rho'(\nu). \quad (5.47)$$

Also, from Maxwell's equations, we may show that the electric field per photon for a collection of photons in a given volume V is

$$\mathcal{E} = \sqrt{\frac{2\hbar\omega}{\epsilon_0 V}}. \quad (5.48)$$

Combining all of these together, we get the rate of stimulated absorption

$$\frac{dn_n}{dt} = -B_{nj} \rho'(\nu_{nj}) n_n, \quad (5.49)$$

where the *Einstein B coefficient* is

$$B_{nj} = \frac{2\pi}{4\pi\epsilon_0 3\hbar^2} |\langle j|\hat{\mu}|n\rangle|^2 = \frac{8\pi^3}{4\pi\epsilon_0 3\hbar^2} |\langle j|\hat{\mu}|n\rangle|^2 \quad (5.50)$$

as claimed in A3: *High Resolution Molecular Spectroscopy*. The exactly same equation also holds for stimulated emissions.

Example. Dynamic polarisability.

We see that near-resonant radiations will induce transition. Let us turn to the response of molecules to oscillating electric fields whose frequency ω is far from any transition frequency. To ensure that the molecule is not shocked by the field, we switch it on slowly, with

$$\hat{H}^{(1)} = \hat{V}(e^{i\omega t} + e^{-i\omega t})(1 - e^{-kt}) = 2\hat{V}(1 - e^{-kt}) \cos \omega t, \quad (5.51)$$

where $k \ll \omega$. For simplicity, we will assume z polarised radiation, so that $\hat{V} = -\frac{1}{2}\hat{\mu}_z \mathcal{E}_z$. Substituting into (5.25), performing the time integral, and taking the long time limit $t \gg k^{-1}$, we obtain

$$a_j(t) = \frac{1}{2} \mathcal{E}_z \langle j|\hat{\mu}_z|n\rangle \left(\frac{e^{i(\omega_{jn}+\omega)t}}{\omega_{jn}+\omega} + \frac{e^{i(\omega_{jn}-\omega)t}}{\omega_{jn}-\omega} \right). \quad (5.52)$$

Since our radiation frequency is not close to the energy gaps ω_{jn} , we don't expect any one of the two terms dominate, and we have to consider both of them.

$$\Psi(t) = \Psi_n(t) + \sum_{j \neq n} a_j(t) \Psi_j(t). \quad (5.53)$$

We are interested in how the electron distribution, and hence the electric dipole of the molecule fluctuates in response to the imposed oscillating field, so we would like to evaluate the time-dependent response of the molecular dipole moment.

$$\begin{aligned}
\mu_z(t) &= \langle \Psi | \hat{\mu}_z | \Psi \rangle \\
&= \left\langle \Psi_n + \sum_{j \neq n} a_j \Psi_j \left| \hat{\mu}_z \right| \Psi_n + \sum_{j \neq n} a_j \Psi_j \right\rangle \\
&= \langle \Psi_n | \hat{\mu}_z | \Psi_n \rangle + \sum_{j \neq n} [a_j \langle \Psi_n | \hat{\mu}_z | \Psi_j \rangle + a_j^* \langle \Psi_j | \hat{\mu}_z | \Psi_n \rangle] + O(\mathcal{E}_z^2) \\
&= \langle n | \hat{\mu}_z | n \rangle + \sum_{j \neq n} [a_j \langle n | \hat{\mu}_z | j \rangle e^{-i\omega_{jn}t} + a_j^* \langle j | \hat{\mu}_z | n \rangle e^{i\omega_{jn}t}] + O(\mathcal{E}_z^2), \tag{5.54}
\end{aligned}$$

where we have kept only the first-order terms in the field strength by discarding contributions that are quadratic in the coefficients a_j .³¹ Substituting the expression of $a_j(t)$, we arrive at

$$\mu_z(t) = \mu_{z,n}^{(0)} + \frac{1}{2} \mathcal{E}_z (e^{i\omega t} + e^{-i\omega t}) \sum_{j \neq n} |\langle j | \hat{\mu}_z | n \rangle|^2 \left(\frac{1}{\omega_{jn} + \omega} + \frac{1}{\omega_{jn} - \omega} \right). \tag{5.55}$$

We see that the dipole oscillates harmonically about its permanent value $\mu_{z,n}^{(0)}$ in the absence of a field with the same frequency ω as the driven frequency. Since, in general,

$$\mu = -\frac{\partial E}{\partial \mathcal{E}} = \mu^{(0)} + \alpha \mathcal{E} + \dots, \tag{5.56}$$

we may interpret the sum in (5.55) as a *frequency-independent* or *dynamic polarisability*,

$$\alpha_{zz}(\omega) := 2 \sum_{j \neq n} \frac{\omega_{jn} |\langle j | \hat{\mu}_z | n \rangle|^2}{\omega_{jn}^2 - \omega^2}, \tag{5.57}$$

so that

$$\mu_z(t) = \mu_z^{(0)} + \alpha_{zz}(\omega) \mathcal{E}_z \cos \omega t. \tag{5.58}$$

The subscript zz emphasises that it measures the dipole induced in the z direction by a field along z direction — the polarisability is a tensor.

Note the the dynamic polarisability (5.57) reduces to the static polarisability (3.145) as $\omega \rightarrow 0$. Also the polarisability diverges if $\omega \rightarrow \omega_{jn}$ since then a electric dipole will not induce a dipole but instead cause a transition.

³¹The term proportional to \mathcal{E}_z^2 should also have contributions from the second-order perturbation coefficient $a_j^{(2)}$. We don't have it, so including $|a_j^{(1)}|^2$ here is meaningless.

6 Intermolecular Interactions

6.1 The Dispersion Energy

Weak attractive forces exist even between uncharged non-polar particles, and are significant enough to result in weakly bound dimers of noble gas atoms. Although atoms are spherical on average, and therefore have no permanent dipole moment, fluctuations in the electron density constitute instantaneous dipole moments. We will show in this section that the average interaction between fluctuations on different atoms lowers the energy, and we will derive an expression for the strength of the resulting *dispersion force*.

For a two-atoms system, A and B, we let the zeroth-order Hamiltonian be the sum of the Hamiltonian of the isolated atoms, and contains no terms of interaction between them:

$$\hat{H}^{(0)} = \hat{H}_A^{(0)} + \hat{H}_B^{(0)}. \quad (6.1)$$

The zeroth-order wavefunctions are therefore the products of the individual wavefunctions, $\psi_{mn}^{(0)} = \psi_{A,m}^{(0)} \psi_{B,n}^{(0)}$, and we will denote this by $|mn\rangle$.

We now add the interaction between the atoms as a perturbation. The largest contribution comes from the energy of the instantaneous dipole moments. From classical electrostatic, and replacing the classical quantities by their quantum mechanical operators, the energy of two point dipole $\boldsymbol{\mu}_A$ and $\boldsymbol{\mu}_B$ at separation \mathbf{R} is³²

$$\hat{H}^{(1)} = \frac{1}{R^3} \left[\hat{\boldsymbol{\mu}}_A \cdot \hat{\boldsymbol{\mu}}_B - \frac{3}{R^2} (\hat{\boldsymbol{\mu}}_A \cdot \mathbf{R})(\hat{\boldsymbol{\mu}}_B \cdot \mathbf{R}) \right]. \quad (6.5)$$

Without loss of generality, we can take \mathbf{R} to be parallel to z axis, giving the simpler form

$$\hat{H}^{(1)} = \frac{1}{R^3} (\hat{\mu}_{Ax} \hat{\mu}_{Bx} + \hat{\mu}_{Ay} \hat{\mu}_{By} - 2\hat{\mu}_{Az} \hat{\mu}_{Bz}). \quad (6.6)$$

The first-order energy $E_0^{(1)} = \langle 00 | \hat{H}^{(1)} | 00 \rangle$ only contain terms

$$\langle 00 | \hat{\mu}_{Ai} \hat{\mu}_{Bi} | 00 \rangle = \langle 0_A | \hat{\mu}_{Ai} | 0_A \rangle \langle 0_B | \hat{\mu}_{Bi} | 0_B \rangle, \quad (6.7)$$

³²Let's put dipole $\boldsymbol{\mu}_A$ at origin and a dipole $\boldsymbol{\mu}_B = q\mathbf{d}$ at \mathbf{R} , with $\|\mathbf{R}\| \gg \|\mathbf{d}\|$. The potential generated by $\boldsymbol{\mu}_B$ at origin is

$$\begin{aligned} \phi &= \frac{1}{4\pi\epsilon_0} \left(\frac{q}{R} - \frac{q}{\|\mathbf{R} + \mathbf{d}\|} \right) \\ &= \frac{q}{4\pi\epsilon_0} \left[\frac{1}{R} - \left(\frac{1}{R} + \mathbf{d} \cdot \nabla \frac{1}{R} + \frac{1}{2} (\mathbf{d} \cdot \nabla)^2 \frac{1}{R} + \dots \right) \right] \\ &= \frac{q}{4\pi\epsilon_0} \frac{\mathbf{d} \cdot \mathbf{R}}{R^3} \\ &= \frac{\boldsymbol{\mu}_B \cdot \mathbf{R}}{4\pi\epsilon_0 R^3}. \end{aligned} \quad (6.2)$$

Then the electric field by $\boldsymbol{\mu}_B$ at origin is

$$\begin{aligned} \mathbf{E} &= -\nabla\phi = -\frac{1}{4\pi\epsilon_0} \mathbf{e}_i \frac{\partial}{\partial x_i} \frac{\mu_{Bj} x_j}{(x_k x_k)^{3/2}} \\ &= -\frac{1}{4\pi\epsilon_0} \mathbf{e}_i \left[\frac{\mu_{Bj} \delta_{ij} (x_k x_k)^{3/2} - \mu_{Bj} x_j \cdot \frac{3}{2} (x_l x_l)^{1/2} \cdot 2\delta_{ik} x_k}{(x_m x_m)^3} \right] \\ &= \frac{1}{4\pi\epsilon_0} \mathbf{e}_i \frac{3(\mathbf{R} \cdot \boldsymbol{\mu}_B) R R_i - \mu_{Bj} R^3}{R^6} \\ &= \frac{\mu_B}{4\pi\epsilon_0 R^3} [3(\hat{\mathbf{R}} \cdot \hat{\boldsymbol{\mu}}_B) \hat{\mathbf{R}} - \hat{\boldsymbol{\mu}}_B] \end{aligned} \quad (6.3)$$

using summation convention. Note a vector with a hat here means unit vector. Then finally, the interaction energy of the two dipole is

$$E = -\mathbf{E} \cdot \boldsymbol{\mu}_A = \frac{1}{4\pi\epsilon_0 R^3} \left[\boldsymbol{\mu}_A \cdot \boldsymbol{\mu}_B - \frac{3}{R^2} (\mathbf{R} \cdot \boldsymbol{\mu}_A)(\mathbf{R} \cdot \boldsymbol{\mu}_B) \right]. \quad (6.4)$$

where $i \in \{x, y, z\}$. All of these are zero since an atom has no permanent dipole moment in any direction, so $E_0^{(1)} = 0$.

The second-order energy is

$$E_0^{(2)} = - \sum_{m \neq 0} \sum_{n \neq 0} \frac{|\langle 00 | \hat{H}^{(1)} | mn \rangle|^2}{E_{A,m}^{(0)} + E_{B,n}^{(0)} - E_{A,0}^{(0)} - E_{B,0}^{(0)}}. \quad (6.8)$$

Since $\hat{H}^{(1)}$ contains three terms, the square in the numerator contains nine. However, all the cross terms like

$$\langle 00 | \hat{\mu}_{Ai} \hat{\mu}_{Bi} | mn \rangle \langle mn | \hat{\mu}_{Aj} \hat{\mu}_{Bj} | 00 \rangle = \langle 0_A | \hat{\mu}_{Ai} | m_A \rangle \langle 0_B | \hat{\mu}_{Bi} | n_B \rangle \langle m_A | \hat{\mu}_{Aj} | 0_A \rangle \langle n_B | \hat{\mu}_{Bj} | 0_B \rangle \quad (6.9)$$

vanishes, where $i \neq j \in \{x, y, z\}$. This is because we can reverse the direction of, say, axis i on atom A, and this will change the sign of the expression. However, energy should be totally symmetric, so it can only be zero. The only non-vanishing terms are

$$\langle 00 | \hat{\mu}_{Ai} \hat{\mu}_{Bi} | mn \rangle \langle mn | \hat{\mu}_{Ai} \hat{\mu}_{Bi} | 00 \rangle = \langle 0_A | \hat{\mu}_{Ai} | m_A \rangle \langle 0_B | \hat{\mu}_{Bi} | n_B \rangle \langle m_A | \hat{\mu}_{Ai} | 0_A \rangle \langle n_B | \hat{\mu}_{Bi} | 0_B \rangle, \quad (6.10)$$

where $i \in \{x, y, z\}$. Moreover, by spherical symmetry, this is the same for x , y and z , and we have 6 such terms in total (1 from x , 1 from y and $2^2 = 4$ from z). This gives us the total expression

$$E_0^{(2)} = - \frac{6}{R^6} \sum_{m,n \neq 0} \frac{|\langle 0_A | \hat{\mu}_{Az} | m_A \rangle|^2 |\langle 0_B | \hat{\mu}_{Bz} | m_B \rangle|^2}{(E_{A,m}^{(0)} - E_{A,0}^{(0)}) + (E_{B,n}^{(0)} - E_{B,0}^{(0)})}. \quad (6.11)$$

Since both the numerator and denominator are positive, the dispersion force is attractive and scales as R^{-6} .

Equation (6.11) resembles the polarisability, and we might expect that the strength of dispersion energy would be related to the polarisability of the atom as they need to respond to the instantaneous field generated by each other. However, the connection is not obvious to work out explicitly. We need the seemingly arbitrary identity

$$\frac{2}{\pi} \int_0^\infty d\omega \frac{xy}{(x^2 + \omega^2)(y^2 + \omega^2)} = \frac{1}{x + y}. \quad (6.12)$$

Putting $x = \omega_{m0} = E_{A,m}^{(0)} - E_{A,0}^{(0)}$ and $y = \omega_{n0} = E_{B,n}^{(0)} - E_{B,0}^{(0)}$, the equation becomes

$$E_0^{(2)} = - \frac{12}{\pi R^6} \sum_{m,n \neq 0} \int_0^\infty d\omega \frac{\omega_{m0} |\langle 0_A | \hat{\mu}_{Az} | m_A \rangle|^2 \omega_{n0} |\langle 0_B | \hat{\mu}_{Bz} | m_B \rangle|^2}{(\omega_{m0}^2 + \omega^2)(\omega_{n0}^2 + \omega^2)} \quad (6.13)$$

$$= - \frac{3}{\pi R^6} \sum_{m,n \neq 0} \int_0^\infty d\omega \alpha_{A,zz}(i\omega) \alpha_{B,zz}(i\omega), \quad (6.14)$$

where

$$\alpha_{zz}(i\omega) = 2 \sum_{n \neq 0} \frac{\omega_{n0} |\langle 0 | \hat{\mu}_z | n \rangle|^2}{\omega_{n0}^2 + \omega^2} \quad (6.15)$$

is the dynamic polarisability (5.57) with imaginary frequency.

The advantage of (6.14) is that it is formally exact and of practical use, since the polarisabilities can be calculated both experimentally and computationally. The dispersion is often written $-C_6/R^6$, where the coefficient C_6 are often treated as an empirical quantity. Equation (6.14), however, allows us to compute it analytically.

6.2 The Induction Energy

The situation would be much more complex for molecules. If the molecule possess permanent dipole, then the first-order energy

$$E_0^{(1)} = \langle 00 | \hat{H}^{(1)} | 00 \rangle = \frac{1}{R^3} (\mu_{Ax}^{(0)} \mu_{Bx}^{(0)} + \mu_{Ay}^{(0)} \mu_{By}^{(0)} - 2\mu_{Az}^{(0)} \mu_{Bz}^{(0)}) \quad (6.16)$$

is no longer zero. But since the interaction is now orientation-dependent, we need to integrate over all possible relative orientations weighted by the corresponding Boltzmann factor since not all orientations are equally probable and obtain a weighted average of interaction. This is too complicated to deal with, so we will not do it here. We will quote the result:

$$\langle E_0^{(1)} \rangle = \frac{\mu_A^2 \mu_B^2}{k_B T R^3}. \quad (6.17)$$

This is simply the electrostatic interaction of two point dipoles, and scales as R^{-3} .

In the second-order energy, we should only exclude $|m, n\rangle = |0, 0\rangle$ from the Rayleigh–Schrödinger sum, but in the dispersion force of atom, (6.8), we have excluded all states with either $m = 0$ or $n = 0$ since they evaluate to zero. The dispersion energy can be thought as an interaction between simultaneous excited states of two atoms. However, for polar molecules, terms of form $\langle 00 | \hat{H}^{(1)} | m0 \rangle$ and $\langle 00 | \hat{H}^{(1)} | 0n \rangle$, where only one of the two species is excited, may be non-zero. It is therefore conventional to separate the second-order energy into three contributions,

$$E_0^{(2)} = E_{\text{disp}} + E_{A,\text{ind}} + E_{B,\text{ind}}, \quad (6.18)$$

where

$$E_{\text{disp}} = - \sum_{m \neq 0} \sum_{n \neq 0} \frac{|\langle 00 | \hat{H}^{(1)} | mn \rangle|^2}{E_{A,m}^{(0)} + E_{B,n}^{(0)} - E_{A,0}^{(0)} - E_{B,0}^{(0)}}, \quad (6.19)$$

$$E_{A,\text{ind}} = - \sum_{m \neq 0} \frac{|\langle 00 | \hat{H}^{(1)} | m0 \rangle|^2}{E_{A,m}^{(0)} - E_{A,0}^{(0)}}, \quad (6.20)$$

$$E_{B,\text{ind}} = - \sum_{n \neq 0} \frac{|\langle 00 | \hat{H}^{(1)} | 0n \rangle|^2}{E_{B,n}^{(0)} - E_{B,0}^{(0)}}. \quad (6.21)$$

The dispersion term (6.19) is again the interaction between the instantaneous dipole of A and the instantaneous dipole of B, but we now have two extra contributions. The induction terms (6.20) and (6.21) are the interactions between permanent dipole of A with the instantaneous dipole of B, and the permanent dipole of B with the instantaneous dipole of A, respectively.

7 Relativistic Effects

Everything we discussed so far ignores *relativity*, which refers to the breakdown of classical mechanics when particles have relative speeds approaching the speed of light in vacuum. Special relativity is the simpler theory of relativity, which deals with situations in which gravity is negligible. This is almost always the case as far as chemistry is concerned since electromagnetic forces are orders of magnitudes larger than gravity. The incorporation of special relativity into quantum mechanics was first achieved by Dirac in 1928. General relativity is much more complicated and the unification of quantum mechanics and general relativity is still an important open question in theoretical physics.

Although chemists often ignore relativity, as you have previously seen in the IA inorganic and materials chemistry course, relativistic effects play a significant effect for heavy atoms. However, any sufficient precise prediction for any atom, including hydrogen, must include relativistic effect to be accurate.

The *Dirac equation* for a free electron is

$$\left(\beta m_e c^2 + c \sum_{n=1}^3 \alpha_n \hat{p}_n \right) \psi(\mathbf{x}, t) = i\hbar \frac{\partial \psi(\mathbf{x}, t)}{\partial t}. \quad (7.1)$$

This looks reminiscent to the time-dependent Schrödinger equation, but with a few important differences. The wavefunction ψ is no longer a scalar function, but a 4-component vector, which corresponds to a spin-up and spin-down electron, and a spin-up and spin-down positron. The operator in brackets on the left replaces the non-relativistic Hamiltonian. α_n and β are 4×4 matrices that obey certain symmetry and commutation properties. m_e is the rest mass of the electron, c is the speed of light and \hat{p}_n is the n^{th} component of the momentum operator. To include the potential due to hydrogen nucleus, we must add in the potential

$$\left(\beta m_e c^2 + c \sum_{n=1}^3 \alpha_n \hat{p}_n + \frac{e^2}{4\pi\epsilon_0 r} \right) \psi(\mathbf{x}, t) = i\hbar \frac{\partial \psi(\mathbf{x}, t)}{\partial t}. \quad (7.2)$$

The Dirac equation for free spin- $\frac{1}{2}$ massive particle is often recast in a more compact form as

$$\left(i\hbar \gamma^\mu \hat{\partial}_\mu - mc \right) \psi = \mathbf{0}, \quad (7.3)$$

where γ^μ is a set of 4×4 matrices formed from α_n and β . $\hat{\partial}_\mu$ is the component of the 4-gradient (essentially ∇ with a fourth dimension of time), and Einstein summation applies to sum over the four components μ .

As gravity is completely negligible in chemical systems, exact solutions to the Dirac equation offer a huge improvement over the non-relativistic Schrödinger equation. However, we already know that the exact solutions to the Schrödinger equation are rare for chemical systems, the Dirac equation is only even harder to deal with.

“The underlying physical laws necessary for the mathematical theory of a large part of physics and the whole of chemistry are thus completely known, and the difficulty is only that the exact application of these laws leads to equations much too complicated to be soluble.”

— Paul Dirac (1929), *The quantum mechanics of many-electron systems*.

Since Dirac’s claim, significant progress has been made in the field of numerical solutions, particularly with the advent of fast digital computation, and it is no longer the case that Dirac’s equation is too complicated to be (numerically) solvable. It is, however, still complicated to be soluble using pen and paper.

Beyond the Dirac equation is quantum field theory, in which properties of the underlying fields, such as the electromagnetic field, are also quantised. Moreover, elementary particles like electrons are also thought of as ‘ripples’ on an ‘electron field’ spanning throughout the whole universe. This is a much deeper subject, but it can be chemically significant — in particular, spontaneous emission can only be explained by QFT.

Instead, to investigate the relativistic effect, we will use perturbation theory to add the terms that arise from the non-relativistic limit of Dirac’s equation into Schrödinger equation as small perturbations. Our main example would be hydrogen spectrum, the observations of which were also historically a significant driver in the development of relativistic quantum mechanics.

7.1 Hydrogen Fine Structure

The *Lyman- α line* in the hydrogen spectrum is the $n = 1 \rightarrow 2$ transition observed at around 121.57 nm. Using the familiar exact solution to the Schrödinger equation, we can predict that the line should be observed at

$$h\nu = E_2 - E_1 = R_H \left(1 - \frac{1}{4}\right) = \frac{3}{4}R_H, \quad (7.4)$$

where R_H is the Rydberg constant.

There is actually a subtlety concerning this constant that worth mentioning. The collection of constant that emerges from the normal Schrödinger equation for the Rydberg constant is

$$R_\infty = \frac{m_e e^4}{8\epsilon_0^2 h^2} = 2.17987 \times 10^{-18} \text{ J}. \quad (7.5)$$

Using this value, we will predict the Lyman- α line should appear at 121.50 nm, noticeably different from the experimental value. The error is arising from the Born–Oppenheimer approximation: we have treated the nucleus as immobile, or equivalently having infinite mass. The hydrogen nucleus is indeed massive compared with the electron, but it is only heavier by a factor of ~ 1836 . The error of this approximation is easily observable in accurate spectroscopic measurement. Instead, we should replace the electron mass with the reduced mass of the electron-proton system

$$\mu = \frac{m_e m_p}{m_e + m_p}, \quad (7.6)$$

which gives the true Rydberg constant

$$R_H = \frac{\mu e^4}{8\epsilon_0^2 h^2} = 2.17869 \times 10^{-18} \text{ J}. \quad (7.7)$$

Using this value, we predict the Lyman- α line at 121.57 nm, in agreement with experiment. Incidentally, this also implies that the line for deuterium should be at a slightly shorter wavelength than hydrogen due to a slightly larger effective mass, appearing at 121.54 nm. This is actually how the existence of deuterium was proved in 1931, surprisingly slightly before the discovery of neutron in 1932.

Even in the late nineteenth century, the precision of UV spectrometers exceeds the one-hundredth of a nm we have just quoted. In 1887 Michelson and Morley³³ observed that the Lyman- α for hydrogen is split into a doublet, with lines at 121.5668 nm and 121.5674 nm. This splitting, which is also observed for other lines, is called the *fine structure* of the spectrum. It is a result of relativistic effects.

³³They are more well-known for disproving the existence of aether.

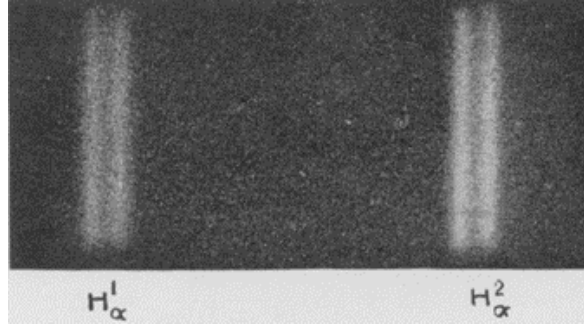


Figure 11: H_α ($n = 2 \rightarrow n = 3$) lines of hydrogen and deuterium. Each are shown to be a doublet. Figure adapted from G.N. Lewis and F.H. Spedding, Phys. Rev. 43, 964 (1933).

7.1.1 Relativistic Kinetic Energy

The first relativistic contribution is the easiest to understand. It arises from the relativistic correction to the kinetic energy. Classically, the non-relativistic kinetic energy is

$$T = \frac{\mathbf{p}^2}{2m}, \quad (7.8)$$

where $\mathbf{p} = m\mathbf{v}$ is the classical non-relativistic momentum. However, in special relativity, the momentum is given by

$$T = (\gamma - 1)mc^2 = \left(\frac{1}{\sqrt{1 - \frac{v^2}{c^2}}} - 1 \right) mc^2. \quad (7.9)$$

Written in terms of non-relativistic momentum, and expand for small $v \ll c$, we get

$$T = mc^2 \left(\frac{1}{\sqrt{1 - \frac{\mathbf{p}^2}{m^2 c^2}}} - 1 \right) \quad (7.10)$$

$$= \frac{\mathbf{p}^2}{2m} - \frac{\mathbf{p}^4}{8m^3 c^2} + \dots \quad (7.11)$$

The first term is just the non-relativistic kinetic energy, and the second term is the relativistic correction to the leading order. Therefore, for a hydrogen atom, we use perturbative Hamiltonian³⁴

$$\hat{H}_T^{(1)} = -\frac{\hat{\mathbf{p}}^4}{8\mu^3 c^2}. \quad (7.12)$$

This term is the first of our three relativistic corrections, and we use a subscript T to stand for the kinetic contribution to the relativistic correction. The first-order perturbation to the energy is³⁵

$$E_{T,n\ell m}^{(1)} = \langle n, \ell, m | \hat{H}_T^{(1)} | n, \ell, m \rangle = -\frac{1}{8\mu^3 c^2} \langle n, \ell, m | \hat{\mathbf{p}}^4 | n, \ell, m \rangle. \quad (7.13)$$

We can obtain a general expression of this by some algebraic manipulations. The Schrödinger equation of the reference Hamiltonian gives

$$\left(\frac{\hat{\mathbf{p}}^2}{2\mu} - \frac{1}{r} \right) |n, \ell, m\rangle = E_n^{(0)} |n, \ell, m\rangle \quad (7.14)$$

$$\hat{\mathbf{p}}^2 |n, \ell, m\rangle = 2\mu \left(\frac{1}{r} + E_n^{(0)} \right) |n, \ell, m\rangle \quad (7.15)$$

³⁴The official handout is again being rather loose here and directly set $\mu = m_e = 1$ in atomic unit.

³⁵One might be worrying about the degeneracies. Note that our system is rotationally invariant, so in particular $[L_z, \hat{H}_T^{(1)}] = 0$ and $[\mathbf{L}^2, \hat{H}_T^{(1)}] = 0$. Therefore $\langle n, \ell', m' | \hat{H}_T^{(1)} | n, \ell, m \rangle = 0$ unless both $\ell' = \ell$ and $m' = m$. Thus our perturbation does not mix degenerate states of the gross structure, so non-degenerate perturbation theory is sufficient.

in atomic units, where $E_n^{(0)} = -\frac{\mu}{2n^2}$ is the non-relativistic energy of state $|n, \ell, m\rangle$ of hydrogen atom.³⁶ Therefore,

$$\begin{aligned} E_{T,n\ell m}^{(1)} &= -\frac{1}{8\mu^3 c^2} \langle n, \ell, m | \hat{\mathbf{p}}^2 \hat{\mathbf{p}}^2 | n, \ell, m \rangle \\ &= -\frac{1}{2\mu c^2} \left(\frac{\mu^2}{4n^4} - \frac{\mu}{n^2} \left\langle \frac{1}{r} \right\rangle + \left\langle \frac{1}{r^2} \right\rangle \right). \end{aligned} \quad (7.16)$$

For hydrogen orbital, it can be shown that³⁷

$$\left\langle \frac{1}{r} \right\rangle = \frac{\mu}{n^2}, \quad (7.17)$$

$$\left\langle \frac{1}{r^2} \right\rangle = \frac{\mu^2}{(\ell + \frac{1}{2}) n^3}. \quad (7.18)$$

Combining all these, we get

$$E_{T,n\ell m}^{(1)} = -\frac{\mu}{8n^4 c^2} \left(\frac{4n}{\ell + \frac{1}{2}} - 3 \right). \quad (7.19)$$

7.1.2 Spin-Orbit Coupling

Due to its spin, an electron has a magnetic dipole moment. Since electron is moving around the nucleus, from the perspective of the electron, the nucleus is also moving around the electron, so the electron is effectively experiencing a magnetic field. The magnetic dipole moment of the electron can interact with the magnetic dipole, and the Hamiltonian of this perturbation is³⁸

$$\hat{H}_{\text{SO}}^{(1)} = \frac{1}{2\mu^2 c^2} \frac{\hat{\mathbf{l}} \cdot \hat{\mathbf{s}}}{r^3}, \quad (7.24)$$

and so the first-order change in the energy is

$$E_{\text{SO},n\ell m}^{(1)} = \frac{1}{2\mu^2 c^2} \langle \hat{\mathbf{l}} \cdot \hat{\mathbf{s}} \rangle \left\langle \frac{1}{r^3} \right\rangle. \quad (7.25)$$

³⁶You might be more familiar with the energy $E_n^{(0)} = -\frac{1}{2n^2}$, but this is under Born–Oppenheimer approximation.

³⁷The first result is obvious from the virial theorem, which states that for a potential $V(r) = ar^k$, we have $2\langle T \rangle = k\langle V \rangle$, and so for the Coulomb potential, $\langle \frac{1}{r} \rangle = -\langle V \rangle = -2E_n^{(0)}$. The second result can also be evaluated using some tricks of defining an effective potential. See my notes on Principles of Quantum Mechanics.

³⁸The magnetic dipole moment of the electron is given by

$$\mathbf{m} = -\frac{g_s}{2\mu} \mathbf{s}, \quad (7.20)$$

where g_s is the electron g factor. It is exactly 2 in Dirac's theory, but in quantum field theory (and in reality), it takes the value $g_s \approx 2.002319\dots$ due to a effect called anomalous magnetic dipole moment. When placed in a magnetic field, this magnetic dipole will have a energy $U = -\mathbf{m} \cdot \mathbf{B}$. If the electron has velocity $\mathbf{v} = \mathbf{p}/\mu\gamma$, then classically the Lorentz transformation of the electric field generated by the nucleus gives the magnetic field experienced by the electron

$$\mathbf{B} = \frac{\gamma}{c^2} \mathbf{v} \times \mathbf{E} = \frac{1}{\mu c^2} \mathbf{p} \times \left(-\frac{\hat{\mathbf{r}}}{r^2} \right) = \frac{1}{\mu c^2 r^3} \mathbf{l}. \quad (7.21)$$

This contributes

$$\frac{g_s}{2\mu^2 c^2} \frac{\mathbf{l} \cdot \mathbf{s}}{r^3} \quad (7.22)$$

to the energy, and is known as the Lamor interaction. There is another complicated effect called Thomas precession, which reduces the energy by $\mathbf{l} \cdot \mathbf{s} / 2\mu^2 c^2 r^3$. Therefore, the total interaction energy is

$$U = \frac{(g_s - 1)}{2\mu^2 c^2 r^3} \mathbf{l} \cdot \mathbf{s} \approx \frac{1}{2\mu^2 c^2 r^3} \mathbf{l} \cdot \mathbf{s} \quad (7.23)$$

using the approximate value $g_s = 2$ (ignoring QFT effects).

We have³⁹

$$\left\langle \frac{1}{r^3} \right\rangle = \frac{\mu^3}{n^3 \ell(\ell + \frac{1}{2})(\ell + 1)}. \quad (7.26)$$

Next, to evaluate the spin-orbit expectation, we introduce the total angular momentum operator $\hat{\mathbf{j}} = \hat{\mathbf{l}} + \hat{\mathbf{s}}$. The reference wavefunctions are eigenfunctions of $\hat{\mathbf{l}}^2$ and $\hat{\mathbf{s}}^2$ with eigenvalues $\ell(\ell + 1)$ and $s(s + 1) = \frac{3}{4}$ respectively, and are also eigenfunctions of $\hat{\mathbf{j}}^2$ with eigenvalues $j(j + 1)$. j is the quantum number for the magnitude of the total angular momentum with possible values given by the Clebsch–Gordan series

$$j \in \{\ell + s, \ell + s - 1, \dots, |\ell - s|\}. \quad (7.27)$$

Since $s = \frac{1}{2}$, the series reduces to

$$j = \begin{cases} \ell \pm \frac{1}{2} & \ell \neq 0 \\ \frac{1}{2} & \ell = 0 \end{cases}. \quad (7.28)$$

Since

$$\langle \hat{\mathbf{j}}^2 \rangle = \langle (\hat{\mathbf{l}} + \hat{\mathbf{s}})^2 \rangle = \langle \hat{\mathbf{l}}^2 \rangle + \langle \hat{\mathbf{s}}^2 \rangle + 2 \langle \hat{\mathbf{l}} \cdot \hat{\mathbf{s}} \rangle, \quad (7.29)$$

we have

$$\begin{aligned} \langle \hat{\mathbf{l}} \cdot \hat{\mathbf{s}} \rangle &= \frac{1}{2} (\langle \hat{\mathbf{j}}^2 \rangle - \langle \hat{\mathbf{l}}^2 \rangle - \langle \hat{\mathbf{s}}^2 \rangle) \\ &= \frac{1}{2} \left(j(j + 1) - \ell(\ell + 1) - \frac{3}{4} \right). \end{aligned} \quad (7.30)$$

Putting these together, we have the spin-orbit coupling energy

$$E_{\text{SO}, n\ell m; j}^{(1)} = \frac{\mu}{4c^2} \frac{j(j + 1) - \ell(\ell + 1) - \frac{3}{4}}{n^3 \ell(\ell + \frac{1}{2})(\ell + 1)}. \quad (7.31)$$

7.1.3 The Darwin Term

The third relativistic term is the Darwin term⁴⁰ and is somewhat more mysterious in origin. It has various interpretations, including a small rapid oscillation of the electron (*zitterbewegung*), a small mixing of with the positron component of the wavefunction, or the effect of spontaneously appearing and annihilating electron-positron pairs. Regardless of the origin, it is a perturbation to the potential energy and has the form

$$\hat{H}_{\text{D}}^{(1)} = \frac{\pi}{2\mu^2 c^2} \delta(\mathbf{r}), \quad (7.32)$$

where $\delta(\mathbf{r})$ is the Dirac delta. Since it only picks out the value of a function at origin in a integral, only orbitals that are non-zero at origin will be affect, which is only the s orbitals. For $\ell \neq 0$, the first-order energy is zero. For s orbitals, the values of their wavefunctions at origin are

$$\psi_{n00}^{(0)}(\mathbf{0}) = \sqrt{\frac{\mu^3}{n^3 \pi}}, \quad (7.33)$$

giving a Darwin contribution

$$E_{\text{D}, n\ell m} = \frac{\pi}{2\mu^2 c^2} \left| \psi_{n00}^{(0)}(\mathbf{0}) \right|^2 \delta_{\ell 0} = \frac{\mu}{2c^2 n^3} \delta_{\ell 0}. \quad (7.34)$$

³⁹This can also be calculated using effective potential. See Principles of Quantum Mechanics.

⁴⁰This is due to Charles Galton Darwin (1887-1962), not the more famous Charles Robert Darwin (1809-1882).

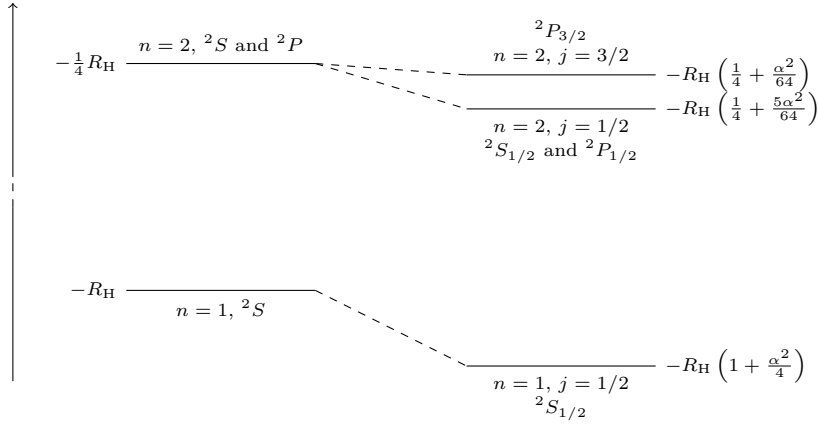


Figure 12: The Schrödinger and Dirac pictures of the hydrogen energy levels. Note that this is not to scale.

7.1.4 Total Effect

If we combine the three first-order relativistic corrections for hydrogen, we the total expression

$$\begin{aligned}
 E_{nj}^{(1)} &= E_{\text{T}}^{(1)} + E_{\text{SO}}^{(1)} + E_{\text{D}}^{(1)} \\
 &= -\frac{\mu}{2c^2 n^4} \left(\frac{n}{j + \frac{1}{2}} - \frac{3}{4} \right) \\
 &= \frac{E_n^{(0)}}{c^2 n^2} \left(\frac{n}{j + \frac{1}{2}} - \frac{3}{4} \right). \tag{7.35}
 \end{aligned}$$

Note that the ℓ -dependence has magically disappeared, and the energy is now only dependent on n and j . The first order energy here is always negative, so the relativistic correction is a decrease in energy.

As with much of spectroscopy, measurements and fitting were carried out before the theory has existed to explain the observations. For historical reason, (7.35) is often written as

$$E_{nj}^{(1)} = \frac{E_n^{(0)} \alpha^2}{n^2} \left(\frac{n}{j + \frac{1}{2}} - \frac{3}{4} \right), \tag{7.36}$$

where α is the *fine-structure constant*. In atomic unit, it is

$$\alpha = \frac{1}{c} \approx \frac{1}{137}. \tag{7.37}$$

Since α is dimensionless, it is the same number in real units

$$\alpha = \frac{e^2}{4\pi\epsilon_0 \hbar c} \approx \frac{1}{137}. \tag{7.38}$$

We can now calculate the magnitudes of the relativistic corrections to the hydrogen energy levels $n = 1$ and $n = 2$ and predict the observed fine-structure for the Lyman- α line. Using the expressions derived above, we predict the two Lyman- α lines are spaced by 0.54 pm in wavelength. The experimental value is 0.60 pm. We have accounted for a large proportion of this shift.

There are a lot of sources of remaining errors. The exact solution of Dirac's equation for a hydrogenic atom gives

$$E_{nj} = mc^2 \left[1 + \left(\frac{Z\alpha}{n - j - \frac{1}{2} + \sqrt{(j + \frac{1}{2})^2 - (Z\alpha)^2}} \right)^2 \right]^{-1/2}. \tag{7.39}$$

Expanding in $1/(Z\alpha)$ gives

$$E_{n,j} = mc^2 \left(1 - (Z\alpha)^2 \frac{1}{2n^2} + (Z\alpha)^4 \left(\frac{3}{4n} - \frac{2}{2j+1} \right) \frac{1}{2n^3} + \dots \right). \quad (7.40)$$

The first term is the rest mass energy of the electron. The second term is the usual non-relativistic hydrogen binding energy, while the third term is the fine-structure corrections that we have laboriously computed above. The higher order terms contribute a large part of our error.

The remaining error comes from QFT effects. In particular, there is a very small difference between the S and P levels with $n = 2$ and $j = \frac{1}{2}$, while we (as well as Dirac equation) predicted them to be degenerate. This is known as the Lamb shift, and it was only accurately measured in 1947, winning a Nobel prize in 1955. The Lamb shift cannot be understood using the kind of single-particle quantum mechanics that we're discussing in this course. It is caused by quantum fluctuations of the electromagnetic field and needs the full machinery of quantum field theory, specifically *quantum electrodynamics*, or QED for short. Historically the experimental discovery of the Lamb shift was one of the prime motivations that led people to develop the framework of quantum field theory.

7.1.5 Hyperfine Structure

Both the fine structure corrections and the QED corrections above treat the nucleus of the atom as a point-like object. This means that, although the corrections are complicated, the problem always has rotational symmetry. In reality, however, the nucleus has structure. This structure affects the atomic energy levels, giving rise to what is called *hyperfine structure*. The most important effects come from the fact that the nucleus also carries a magnetic dipole moment. We denote the spin angular momentum of the nucleus as \mathbf{I} , then the hyperfine structure will depend on the total angular momentum of the whole atom, including the nucleus, $\mathbf{F} = \mathbf{I} + \mathbf{J}$. It is of the order $\alpha^2 m_e/M$, where M is the mass of the nucleus, so it is further depressed by a factor of electron-nucleus mass ratio than the fine structure.

Since the hydrogen nucleus (a proton) has $I = \frac{1}{2}$, each fine-structure level of some J value will be further split into two levels, $F = J + \frac{1}{2}$ and $F = J - \frac{1}{2}$. For $n = 1$, the $F = 1$ and $F = 0$ states are only separated by 5.88×10^{-6} eV, corresponding to a electromagnetic wavelength of 21 cm.

Although this effect is small, the 21 cm line provides the most powerful way of tracing diffuse gas in interstellar and intergalactic space. This wavelength is much longer than the size of typical specks of dust, so 21 cm radiation can propagate with little absorption right through clouds of dust and gas that do absorb visible light. It was radiation this radiation that first revealed the large scale structure of our galaxy. The line is intrinsically very narrow, so the temperature and radial (with the Earth as origin) velocity of the hydrogen that emitted the radiation can be accurately measured from the Doppler shift and broadening of the observed spectral line.

7.2 Larger Atoms

7.2.1 Hydrogenic Atoms

If we repeated our analysis with a hydrogenic atom of nuclear charge Z , we would find that the leading relativistic correction is

$$E_{nj}^{(1)} = \frac{E_n^{(0)} Z^2 \alpha^2}{n^2} \left(\frac{n}{j + \frac{1}{2}} - \frac{3}{4} \right) \quad (7.41)$$

from which we see that the magnitude of the relativistic correction increases as the fourth power of Z (since there is a Z^2 within $E_n^{(0)}$). Or we could say that the energy corrected to first-order is

$$E_{nj} = E_n^{(0)} \left(1 + \frac{Z^2 \alpha^2}{n^2} \left(\frac{n}{j + \frac{1}{2}} - \frac{3}{4} \right) \right). \quad (7.42)$$

We can derive the approximated form of this dependence using an approximate argument stated in Part IA IMC. Using a completely classical model, the orbiting speed of electron around the nucleus is $\approx Zc/137 \approx Zc\alpha$. The effective relativistic mass of the electron is therefore

$$m_e^* = \gamma m_e = \frac{m_e}{\sqrt{1 - Z^2 \alpha^2}}. \quad (7.43)$$

Replacing this with m_e in the definition of the Rydberg constant gives⁴¹

$$R_H^* \approx \frac{R_H}{\sqrt{1 - Z^2 \alpha^2}}. \quad (7.44)$$

Taking the first order expansion gives

$$E_n = E_n^{(0)} \left(1 + \frac{Z^2 \alpha^2}{2} \right), \quad (7.45)$$

which gives the correct dependence on Z and α , although it naturally misses the subtleties regarding n and j .

7.2.2 Terms and Levels for Real Atoms

Real atoms include interactions between electrons and a full exact relativistic (or even non-relativistic) treatment become analytically impossible. We will have a brief qualitative analysis of what is happening.

We start by specifying the configuration for electrons in an atom. The configuration places electrons into hydrogenic orbitals from the orbital approximation. Due to coulombic interactions between electrons, they exchange angular momentum, and so it is no longer possible to specify an ℓ and s value for each electron. What we can do is to specify the total quantum number L and S for the total angular momentum over all electrons. Each such combination is called a *term*. The electron-electron repulsion means different terms within the same configuration generally have different energies. Then relativistic effects, or spin-orbit coupling in particular, allow exchange between L and S , making them no longer well-defined as well. We can only specify the total angular momentum J . Individual states with a particular J are called *levels*. As long as the splitting from spin-orbit coupling is small compared to the energy difference between terms, we can treat this effect as a small perturbation to the terms and we can still specify the L and S from which each level originates.⁴²

This is illustrated for the ground state carbon atom in figure 13. We first couple individual l and s using Clebsch–Gordan series to give the L and S values. This will give all the terms. Then, we again use the Clebsch–Gordan series to couple L and S to give the possible J values for each term, giving the levels.

7.2.3 The Clebsch–Gordan Series

The easiest way to derive the Clebsch–Gordan series is to consider the direct products of the irreducible representations of the full rotation group. This is explained in *B8: Symmetry*.

⁴¹Our argument here is too crude to even concern the difference between R_H and R_∞ , so we simply take $R_H \approx R_\infty \propto m_e$.

⁴²Actually J and I also couples to give hyperfine structures as described above. However, this is too weak and we usually ignore them.

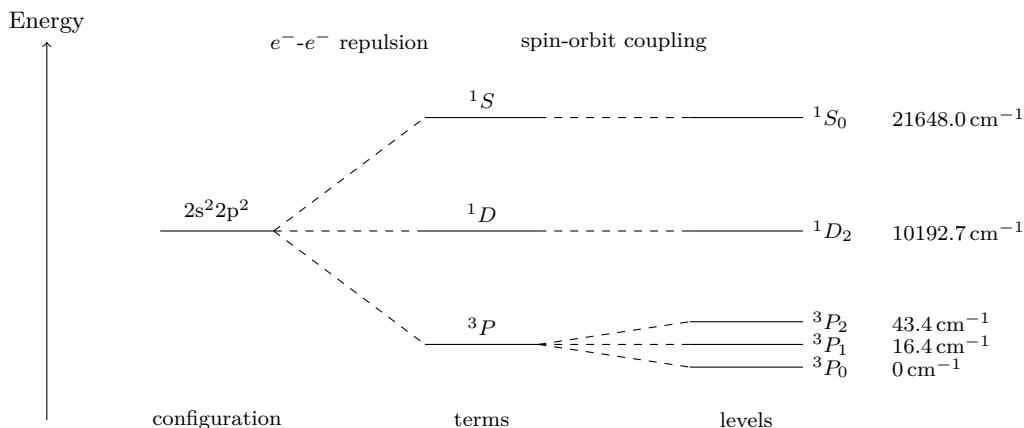


Figure 13: Terms and levels of the carbon ground state configuration.

Here, we will work it out in a more explicit way. Suppose there are two angular momenta with quantum numbers L and S , and call the result of their coupling J . These quantum numbers correspond to the magnitudes of vectors, so cannot be directly added, because in general $\|\mathbf{a}\| + \|\mathbf{b}\| \neq \|\mathbf{a} + \mathbf{b}\|$. What can be added are their z components, M_L and M_S . Following the usual rules of the rigid rotor, M_L takes values from L to $-L$ in integer steps, so there are $2L + 1$ such values, and the equivalent for M_S . Therefore there must be $(2L + 1)(2S + 1)$ values of M_J . The largest possible value of M_J is from the addition of the maximum values of M_S and M_L , so $L + S$. There must be a $J = L + S$ level to which this M_J state belongs. This J level has other states, in fact $2(L + S) + 1$ in total, so there are $4LS$ states remaining unaccounted for. Now consider $M_J = L + S - 1$: there are two such states, as we could have $M_L = L - 1$ and $M_S = S$, or we could have $M_L = L$ and $M_S = S - 1$. One of those states is accounted for by the $J = L + S$ level, leaving the other to belong to a $J = L + S - 1$ level, the next term in the Clebsch-Gordan series. $2(L + S - 1) + 1$ states belong to this level. If we iterate this procedure, keeping track of the number of states accounted for, we run out of states after including the $J = |L - S|$ level.

One question remains concerning the allowed terms in figure 13: why do we not have 1P , 3D and 1S terms? The answer lies with the Pauli exclusion principle: when we consider the allowed values of individual m_l and m_s , we cannot have two electrons with the same m_l and m_s values. There are two approaches to deciding which combinations of L and S are allowed. The simplest is to keep track of which L and S come from the symmetrised and antisymmetrised product, and only match a symmetrised L with an antisymmetrised S , and *vice versa*, which is what we did in *B8: Symmetry*. An alternative and more laborious procedure is to consider every possible combination of m_s and m_l and the M_L and M_S they result in. Then infer which combinations of L and S give rise to the set of M_L and M_S states you have generated.

7.2.4 Hund's Rules

Hund's rules allows us to determine the ground state term and level.

Theorem 7.1 (Hund's rules).

1. the ground term is the one with the highest S ;
2. if two or more terms have the same maximum S , the ground state is the one with the highest L ;
3. if the shell is less than half-full, the ground state level is the one with the lowest J , and if the shell is more than half-full, the ground state level is the one with the highest J .

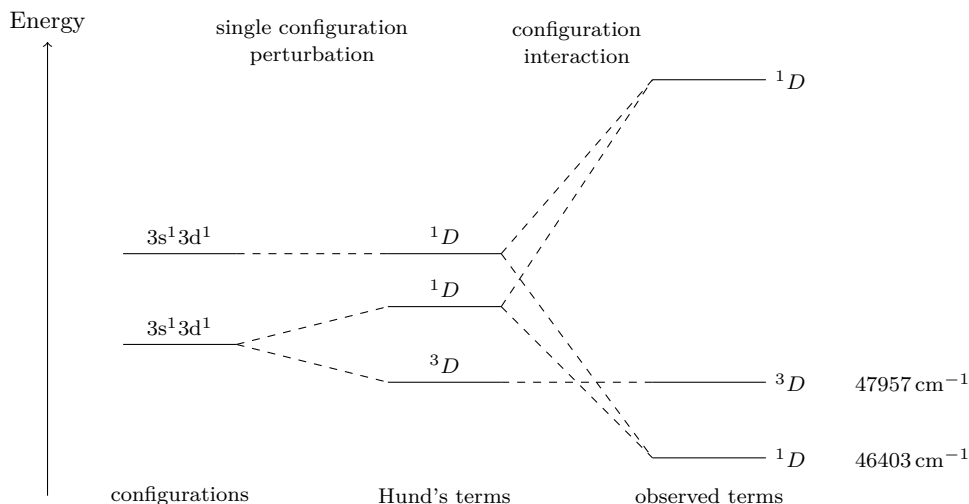


Figure 14: The ordering of the excited Mg violets the Hund's rules due to configuration interaction.

The justification for Hund's first rule lies with Fermi holes and has been described previously. Hund's second rule can be justified by the classical analogy of orbiting electrons. Maximising L means that the m_l of individual electrons are, as far as possible, aligned, so they are 'orbiting in the same direction'. Two electrons orbiting in the same direction will 'approach each other' less frequently than those orbiting in opposite directions, so the average repulsion is reduced.

For Hund's third rule, consider that maximising J would mean maximising M_L and M_S , so aligning L and S as far as possible. Remembering that spin-orbit coupling is a magnetic interaction, this arrangement corresponds to aligning two magnets, which is higher energy than opposing them. Therefore the lower J value is lower energy. If the shell is half full, then the ground state term must have $L = 0$, so there is only one level and no decision to make. If the shell is more than half full, consider instead a shell that is less than half full of holes. We can apply the exact same reasoning to the holes as for the electrons, but now maximising the energy of the holes means minimising the energy of the electrons, so the rules reverses.

As you can tell, our above arguments are rather qualitative. This is because there is no rigorous proof of Hund's rules as they are not rigorously true and exceptions exist. They tend to work well for predicting the ground state level of the ground state configuration, but should not be trusted for ordering beyond that. The classic example of an exception is for the excited configuration of Mg $3s^1 3d^1$. For this configuration, we find terms 3D and 1D and Hund's first rule predicts that the 3D term is lower in energy. The opposite is observed: the 1D is lower in energy. The origin of the problem is that configurations are not a real thing (they arise from the orbital approximation) and when we apply electron-electron repulsion, the perturbation can cause a mixing of the configurations, an example of nearly-degenerate perturbation theory. Here, the $3p^2$ configuration lies not much higher in energy and gives rise to a 1D term, but not a 3D term. The 1D terms from the two configurations mix and the resultant terms contain some contribution from both configurations. As ever with two-state mixing, the lower energy term is lowered and the higher energy term is raised. Mixing like this always happens, but the only question that matters qualitatively is whether it is strong enough to change the order of the terms.

7.2.5 jj Coupling

At the start of this section we said 'as long as the splitting from spin-orbit coupling is small compared to the energy difference between terms...'. This regime is called '*LS coupling*' or '*Russell-Saunders coupling*'. The assumption is that L and S are reasonably well-defined and the spin-orbit coupling is

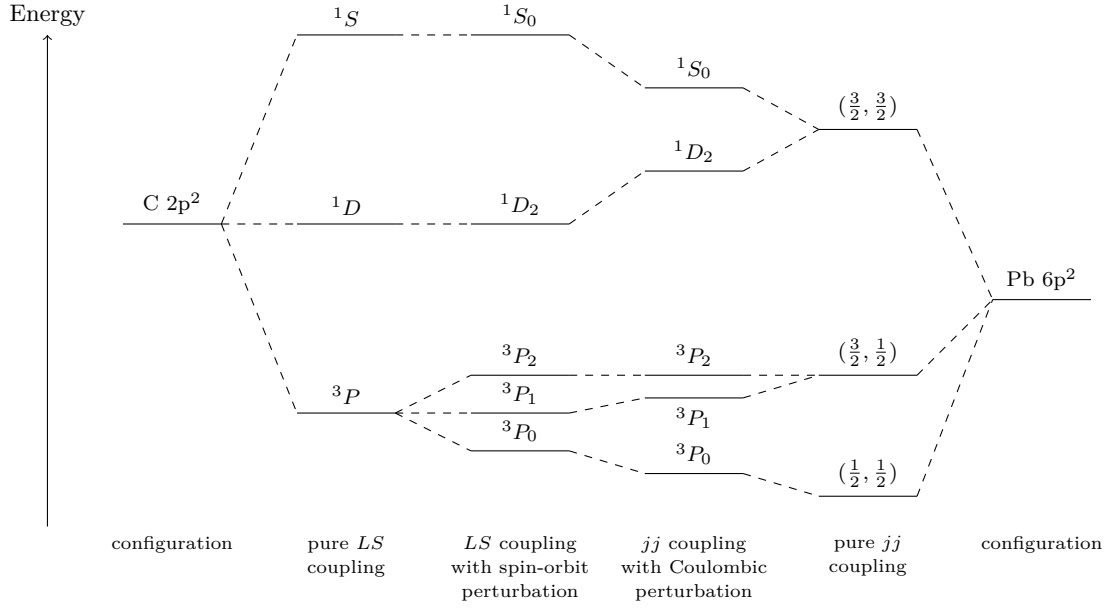


Figure 15: A qualitative correlation diagram showing the levels in the np^2 ground state configurations of C and Pb.

a small perturbation. If we had included the nuclear charge in (7.31), we would have

$$E_{\text{SO},n\ell m;j}^{(1)} = \frac{Z^4}{4c^2} \frac{j(j+1) - \ell(\ell+1) - \frac{3}{4}}{n^3 \ell(\ell + \frac{1}{2})(\ell + 1)}. \quad (7.46)$$

The factor Z^4 shows that the magnitude of the spin-orbit coupling increases rapidly with Z , and when we arrive at around Lanthanoids, the electron-electron repulsions and spin-orbit coupling are about the same magnitude, making the idea of treating spin-orbit coupling as a small perturbation no longer appropriate. As we further increase Z , the spin-orbit coupling would be so large that the Coulombic splitting between terms to be small compared to the spin-orbit splitting between levels. By then, we should use a different scheme, called jj coupling. There is no hard cutoff between LS coupling and jj coupling, as in between both effects are similar in magnitude. Beyond about gold we are safely in the jj coupling regime.

To apply jj coupling, first we generate an individual j for each electron, using the Clebsch-Gordan series to couple the l and s of each electron. Then the individual j are then coupled using the Clebsch-Gordan series to generate the overall J possibilities.

Example. Consider the ground state configuration of Pb, $6p^2$ as an example. As with Russell-Saunders coupling, all full shells have zero angular momentum so can be ignored. Each individual electron has $l = 1$ and $s = \frac{1}{2}$, so for each electron the possibilities for j are $\frac{1}{2}$ and $\frac{3}{2}$. Then we have several ways of combining to get J values. The Clebsch-Gordan series suggests that for $(\frac{1}{2}, \frac{1}{2})$ then $J = 0, 1$; for $(\frac{3}{2}, \frac{1}{2})$ then $J = 1, 2$; and for $(\frac{3}{2}, \frac{3}{2})$ then $J = 0, 1, 2, 3$. Evaluating the degeneracies of each of these levels $(2J + 1)$ and summing gives 30 states, and yet we know that there should be 15 states: 6 ways to place the first electron, 5 ways to place the second, and the order doesn't matter. This is because we have omitted the Pauli principle, which is relevant because both electrons are in the same shell.

The statement of the Pauli principle in jj coupling is that electrons in the same shell with the same j values must have different m_j values. As with Russell-Saunders coupling, we can list all the possible combinations of m_j values, generate the list of M_J values by addition, and then infer the J levels that give rise to them. Doing so removes the $J = 1$ level from the $(\frac{1}{2}, \frac{1}{2})$ set leaving only $J = 0$. It also removes the $J = 3$ and $J = 1$ levels from the $(\frac{3}{2}, \frac{3}{2})$ set, leaving the $J = 0$ and $J = 2$

levels. There is no effect on the $(\frac{3}{2}, \frac{1}{2})$ levels as the electrons have different values of j . Overall then, we have $J = 0, 0, 1, 2, 2$, for the expected total of 15 states.

Hund's third rule still applies and we can suggest that sets with smaller j values will be lower in energy. In the absence of Coulombic effects, the different J values from the same (j_1, j_2) set would have the same energies, but treating the Coulombic effect as a small perturbation, the levels will split. Quite which way is not easy to see, as we need to apply Hund's first and second rules but we have not worked out S and L . We make the observation that there must be a continuous and smooth change from the carbon levels to the lead levels as Z increases, with J staying unchanged, so we can correlate the levels between the two situations. States with the same J are also subject to an avoided-crossing rule, so we can unambiguously decide which level ends up where. The correlation diagram is shown in figure 15. The diagram implies that even when we are in the jj coupling regime, levels can be labelled with their Russell–Saunders term symbols. Although on the jj side, both $J = 2$ levels, for example, will have a significant contribution from both 3P and 1D .

Appendices

A Convergence of Perturbation Series

We began our study of perturbation theory by assuming that the states and energy eigenvalues of the full Hamiltonian depended analytically on a dimensionless parameter λ controlling the perturbation. Even when the individual coefficients of powers of λ are finite, this is often not the case because the infinite perturbative series itself may fail to converge, or may converge only for some range of λ . The issue is that the coefficients of λ may grow too rapidly. Heuristically, the condition for convergence is thus that the typical energy splitting $\langle m | H^{(1)} | n \rangle$ induced by the perturbation should be much smaller than the initial energy difference $E_m - E_n$. However, a detailed criterion is often hard to come by since the higher terms in the perturbation expansion involve complicated sums (or integrals) over many different intermediate states.

To illustrate this in a simple context, let's consider in turn the following three perturbations of a 1D harmonic oscillator potential

$$\hat{H} = \frac{\hat{p}^2}{2m} + \frac{1}{2}m\omega^2 x^2 + \begin{cases} -\lambda m\omega^2 x_0 x \\ +\frac{1}{2}\lambda m\omega^2 x^2 \\ +\lambda \epsilon x^4, \end{cases} \quad (\text{A.1})$$

where x_0 and ϵ are constants. Of course, the first two can be solved exactly — we'd never really use perturbation theory to study them.

In the first case, we have

$$\hat{H} = \frac{\hat{p}^2}{2m} + \frac{1}{2}m\omega^2 (x - \lambda x_0)^2 - \frac{\lambda^2}{2}m\omega^2 x_0^2, \quad (\text{A.2})$$

from which we easily see that the exact energies are

$$E_n(\lambda) = \left(n + \frac{1}{2}\right) \hbar\omega - \frac{\lambda^2}{2}m\omega^2 x_0^2 \quad (\text{A.3})$$

with corresponding position space wavefunction $\langle x | n_\lambda \rangle = \langle x - \lambda x_0 | n \rangle$ just a translation of the usual harmonic oscillator wavefunction $\langle x | n \rangle$. If we instead tackled this problem using perturbation theory, we'd find

$$\begin{aligned} E_n(\lambda) &= E_n - \lambda m\omega^2 x_0 \langle n | x | n \rangle + \lambda^2 m^2 \omega^4 x_0^2 \sum_{k \neq n} \frac{|\langle k | x | n \rangle|^2}{(n - k) \hbar\omega} + O(\lambda^3) \\ &= \left(n + \frac{1}{2}\right) \hbar\omega - \frac{\lambda^2}{2}m\omega^2 x_0^2 + O(\lambda^3). \end{aligned} \quad (\text{A.4})$$

To obtain this result, we note that the first-order term vanishes (e.g. by symmetry), while since x is a linear combination of creation and annihilation operators \hat{a}^\dagger and \hat{a} , only the $j = n + 1$ and $j = n - 1$ terms can contribute to the sum in the second-order term. Going further, we'd find that there are no higher corrections in λ — the $O(\lambda^3)$ terms are in fact zero — though this is not easy to see directly. Thus, in this case, the perturbative result converges to the exact result, and the radius of convergence is infinite. This reflects the fact that the perturbation $-\lambda m\omega^2 x_0 x$ didn't really change the character of the original Hamiltonian. No matter how large λ is, for large enough x the perturbation remains negligible.

Turning to the second case, it's again immediate that the exact energy levels are

$$E_n(\lambda) = \left(n + \frac{1}{2}\right) \hbar\omega(\lambda), \quad (\text{A.5})$$

where $\omega(\lambda) = \omega\sqrt{1+\lambda}$ is the modified frequency. Viewing as a complex function, it has a branch cut starting at $\lambda = -1$, so the energy is only analytic in the disc $|\lambda| < 1$. Again using perturbation theory, we find

$$\begin{aligned} E_n(\lambda) &= E_n + \frac{\lambda}{2} m\omega^2 \langle n|x^2|n \rangle + O(\lambda^3) \\ &= \left(n + \frac{1}{2}\right) \hbar\omega \left(1 + \frac{\lambda}{2} - \frac{\lambda^2}{8} + O(\lambda^3)\right), \end{aligned} \quad (\text{A.6})$$

agreeing to this order with the Taylor expansion of the exact answer. Continuing further, we'd find that this Taylor series does indeed converge provided $|\lambda| < 1$, and that it then converges to the exact answer. The physical reason why the perturbation series diverges when $|\lambda| \geq 1$ is simply that if $\lambda = -1$, the 'perturbation' has completely cancelled the original harmonic oscillator potential, so we are no longer studying a system that can be treated as a harmonic oscillator in the first instance. Once $\lambda < -1$ the harmonic oscillator potential is turned upside down, and we do not expect our system to possess any stable bound states.

Finally, consider the case

$$\hat{H} = \hat{H}^{(0)} + \lambda \epsilon x^4. \quad (\text{A.7})$$

I do not know whether this model has been solved exactly, but it can be treated perturbatively. After a fair amount of non-trivial calculation⁴³ one obtains the series

$$E_0(\lambda) = \hbar\omega + \sum_{n=1}^{\infty} (\lambda\epsilon)^n a_n \quad (\text{A.8})$$

for the ground state energy including the quartic interaction, where the coefficients behave as

$$a_n = \frac{(-1)^{n+1} \sqrt{6}}{\pi^{3/2}} 3^n \Gamma\left(n + \frac{1}{2}\right) \left(1 - \frac{95}{72} \frac{1}{n} + O(n^{-2})\right). \quad (\text{A.9})$$

On account of the Γ -function, these grow faster than factorially with n , so the series (A.8) has radius of convergence $\lambda = 0$! Once again, this is easy to see from the form of the perturbed Hamiltonian: even though we may only care about $\lambda > 0$, our assumption that the perturbation expansion is analytic in λ at $\lambda = 0$ means that, if it converges, it will do so for a disc $\lambda \in D \subset \mathbb{C}$. For any $\lambda \in \mathbb{R}_{<0}$, the Hamiltonian of the quartic oscillator is unbounded below, so there cannot be any stable bound states that are analytic in λ at $\lambda = 0$.

Let me comment that even when perturbative series do not converge, they may still provide very useful information as an asymptotic series. Briefly, we say a series $S_N(\lambda) = \sum_{n=0}^N a_n \lambda^n$ is *asymptotic* to an exact function $S(\lambda)$ as $\lambda \rightarrow 0^+$ (written $S_N(\lambda) \sim S(\lambda)$ as $\lambda \rightarrow 0^+$) if

$$\lim_{\lambda \rightarrow 0^+} \frac{1}{\lambda^N} \left| S(\lambda) - \sum_{n=0}^N a_n \lambda^n \right| = 0. \quad (\text{A.10})$$

In other words, if we just include a fixed number N of terms in our series, then for small enough $\lambda \geq 0$ these first N terms differ from the exact answer by less than $\epsilon \lambda^N$ for any $\epsilon > 0$ (so the difference is $o(\lambda^N)$). However, if we instead try to fix λ and improve our accuracy by including more and more terms in the series, then an asymptotic series will eventually diverge. Most of the perturbative series one meets in the quantum world (including most Feynman diagram expansions in the fancy quantum field theory) are only asymptotic series. Just as in our toy examples above, the radius of convergence of such series is often associated with interesting physics.

⁴³You can find the details in Bender, C. and Wu, T.T., *Anharmonic Oscillator II: A Study of Perturbation Theory in Large Order*, Phys. Rev. D7, 1620-1636 (1973)

B Wigner's $2n + 1$ Theorem

Let's first reformulate our problem a little bit. We know that the energy of a normalised state ψ is given by the expectation value of the Hamiltonian $E = \langle \psi | \hat{H} | \psi \rangle$. This allows to consider the energy as a functional

$$E[\psi, \lambda] = \langle \psi | \hat{H} | \psi \rangle = \langle \psi | \hat{H}^{(0)} + \lambda \hat{H}^{(1)} | \psi \rangle. \quad (\text{B.1})$$

For example, in the unperturbed situation where $\lambda = 0$, the ground state energy is given by the minimum of the energy functional

$$E_{\min} = E[\psi_0^{(0)}, 0], \quad (\text{B.2})$$

where $\psi_0^{(0)}$ is the ground state wavefunction of the unperturbed Hamiltonian. By the Rayleigh–Ritz variational principles, all eigenstates of the Hamiltonian \hat{H} are the stationary values of the energy functional. Therefore, to find the k^{th} eigenstate of \hat{H} for any λ , we can write $\psi_k = \psi_k^{(0)} + \Delta\psi_k$, and find the $\Delta\psi_k$ which satisfies

$$\frac{\partial E[\psi_k^{(0)} + \Delta\psi_k, \lambda]}{\partial(\Delta\psi_k)} = 0, \quad (\text{B.3})$$

and then the eigenvalue (energy) will given by $E_k(\lambda) = E[\psi_k^{(0)} + \Delta\psi_k, \lambda]$. From now on, we will drop the subscript k to refer to any eigenstate. If the perturbation λ is small, then we can expand the eigenstates and the eigenvalues as a Taylor series (as what we did in the normal perturbation theory)

$$\Delta\psi = \sum_{n=1}^{\infty} \frac{1}{n!} \frac{\partial^n \psi}{\partial \lambda^n} \Big|_{\lambda=0} \equiv \sum_{n=1}^{\infty} \psi^{(n)} \lambda^n \quad (\text{B.4})$$

$$E = \sum_{n=0}^{\infty} \frac{1}{n!} \frac{\partial^n E}{\partial \lambda^n} \Big|_{\lambda=0} \equiv \sum_{n=0}^{\infty} E^{(n)} \lambda^n. \quad (\text{B.5})$$

Then the what we want to claim is that we only need to know $\psi^{(i)}$ for $i = 1, \dots, n$ to calculate $E^{(2n+1)}$. Before proving this, we need some preparations.

First, let's expand the functional $E[\psi^{(0)} + \Delta\psi, \lambda]$ in both $\Delta\psi$ and λ . We get the double series

$$E = E[\psi^{(0)} + \Delta\psi, \lambda] = \sum_{k=0}^{\infty} \sum_{p=0}^{\infty} \frac{1}{k!} \frac{1}{p!} \frac{\delta^{k+p} E[\psi^{(0)} + \Delta\psi, \lambda]}{\delta(\Delta\psi)^k \delta \lambda^p} \Big|_{\Delta\psi=0, \lambda=0} (\Delta\psi)^k \lambda^p. \quad (\text{B.6})$$

Then for $\psi = \psi^{(0)} + \Delta\psi$ to be an eigenstate, we need to set $\partial E[\psi^{(0)} + \Delta\psi, \lambda] / \partial(\Delta\psi)$ to zero, which is evaluated as

$$\frac{\partial E[\psi^{(0)} + \Delta\psi, \lambda]}{\partial(\Delta\psi)} = \sum_{k=1}^{\infty} \sum_{p=0}^{\infty} \frac{1}{(k-1)!} \frac{1}{p!} \frac{\delta^{k+p} E[\psi^{(0)} + \Delta\psi, \lambda]}{\delta(\Delta\psi)^k \delta \lambda^p} \Big|_{\Delta\psi=0, \lambda=0} (\Delta\psi)^{k-1} \lambda^p. \quad (\text{B.7})$$

For simplicity, we denote

$$f^{(p)} := \sum_{k=1}^{\infty} \frac{1}{(k-1)!} \frac{1}{p!} \frac{\delta^{k+p} E[\psi^{(0)} + \Delta\psi, \lambda]}{\delta(\Delta\psi)^k \delta \lambda^p} \Big|_{\Delta\psi=0, \lambda=0} (\Delta\psi)^{k-1}, \quad (\text{B.8})$$

and so the above expression reduces to

$$\frac{\partial E[\psi^{(0)} + \Delta\psi, \lambda]}{\partial(\Delta\psi)} = \sum_{p=0}^{\infty} f^{(p)} \lambda^p = 0. \quad (\text{B.9})$$

For this to hold for all λ , we must have $f^{(p)} = 0$ for all p .

Next, let's have a closer look at our energy expansion

$$E = \sum_{n=0}^{\infty} E^{(n)} \lambda^n = \sum_{k=0}^{\infty} \sum_{p=0}^{\infty} c_{kp} (\Delta\psi)^k \lambda^p \quad (\text{B.10})$$

for some complicated coefficients c_{kp} given in (B.6). Here we can use the series expansion (B.4) for $\Delta\psi$

$$\Delta\psi = \sum_{n=1}^{\infty} \psi^{(n)} \lambda^n \quad (\text{B.11})$$

so that

$$E = \sum_{n=0}^{\infty} E^{(n)} \lambda^n = \sum_{k=0}^{\infty} \sum_{p=0}^{\infty} c_{kp} \left(\sum_{n=1}^{\infty} \psi^{(n)} \lambda^n \right)^k \lambda^p \quad (\text{B.12})$$

Now consider the terms contributing to $E^{(2n+1)} \lambda^{2n+1}$ in the above expansion. We claim that the terms must be linear in $\psi^{(G)} \lambda^G$ for all $G > n$. This is because if there is any quadratic (or above) term, then

$$E^{(2n+1)} \lambda^{2n+1} \sim [\psi^{(G)} \lambda^G]^2 \sum_{p=0}^{\infty} \lambda^p = \psi^{(2G)} \underbrace{\lambda^{2G} \sum_{p=0}^{\infty} \lambda^p}_{\lambda^{2n+2} \text{ or higher}}. \quad (\text{B.13})$$

The power of λ is at least $2n+2$, so it cannot contribute to $E^{(2n+1)} \lambda^{2n+1}$. Then, we can write $(\Delta\psi)^k$ in the following way:

$$\begin{aligned} (\Delta\psi)^k &= \left[\sum_{n=1}^k \psi^{(n)} \lambda^n \right]^k = [\psi^{(1)} \lambda + \psi^{(2)} \lambda^2 + \dots + \psi^{(G)} \lambda^G + \dots] [\psi^{(1)} \lambda + \psi^{(2)} \lambda^2 + \dots] \dots \\ &= \sum_{a=k}^{nk} P^{(a)}(\psi^{(1)}, \psi^{(2)}, \dots, \psi^{(n)}) \lambda^a + k \psi^{(G)} \lambda^G (\Delta\psi)^{k-1} + [\text{higher order terms.}] \end{aligned} \quad (\text{B.14})$$

The first term is a polynomial P of $\psi^{(1)}$ up until $\psi^{(n)}$, coming from picking a lower-than- $\psi^{(n)} \lambda^n$ term from each of the square brackets in the first line. The second term pulls out a single $\psi^{(G)} \lambda^G$ from one of the brackets, and there should be k terms like that. Technically this term is at least linear in $\psi^{(G)} \lambda^G$, since $(\Delta\psi)^{k-1}$ will certainly contain $\psi^{(G)} \lambda^G$ in each factor of $\Delta\psi$, but this expansion is good enough to prove what we want. Finally, we are left with terms in which $\psi^{(G)} \lambda^G$ appears twice or more often, and we are not interested in those.

Finally, to prove $2n+1$ theorem, we only need to plug this horrible-looking expansion of $(\Delta\psi)^k$ into our Taylor expansion of E (B.6).

$$\begin{aligned} E &= \sum_{k=0}^{\infty} \sum_{p=0}^{\infty} \frac{1}{k!} \frac{1}{p!} \frac{\delta^{k+p} E[\psi^{(0)} + \Delta\psi, \lambda]}{\delta(\Delta\psi)^k \delta \lambda^p} \Big|_{\Delta\psi=0, \lambda=0} \\ &\quad \left[\sum_{a=k}^{nk} P^{(a)}(\psi^{(1)}, \dots, \psi^{(n)}) \lambda^a + k \psi^{(G)} \lambda^G (\Delta\psi)^{k-1} + [\text{H.O.T.}] \right] \lambda^p, \end{aligned} \quad (\text{B.15})$$

where $G > n$. Now consider terms contributing to $E^{(2n+1)} \lambda^{2n+1}$. They are

$$\begin{aligned} E^{(2n+1)} \lambda^{2n+1} &= \sum_{k=0}^{\infty} \sum_{p=0}^{\infty} \frac{1}{k! p!} \frac{\delta^{k+p} E[\psi^{(0)} + \Delta\psi, \lambda]}{\delta(\Delta\psi)^k \delta \lambda^p} \Big|_{\Delta\psi=0, \lambda=0} \sum_{a=k}^{nk} P^{(a)}(\psi^{(1)}, \dots, \psi^{(n)}) \lambda^{a+p} \Big|_{a+p=2n+1} \\ &\quad + \underbrace{\sum_{p=0}^{\infty} \sum_{k=1}^{\infty} \frac{1}{(k-1)! p!} \frac{\delta^{k+p} E[\psi^{(0)} + \Delta\psi, \lambda]}{\delta(\Delta\psi)^k \delta \lambda^p} \Big|_{\Delta\psi=0, \lambda=0}}_{f^{(p)}} (\Delta\psi)^{k-1} \psi^{(G)} \lambda^{G+1} \Big|_{p+G=2n+1}. \end{aligned} \quad (\text{B.16})$$

Since $f^{(p)} = 0$, the second term vanishes, and we are only left with

$$E^{(2n+1)} \lambda^{2n+1} = \sum_{k=0}^{\infty} \sum_{p=0}^{\infty} \frac{1}{k! p!} \frac{\delta^{k+p} E[\psi^{(0)} + \Delta\psi, \lambda]}{\delta(\Delta\psi)^k \delta \lambda^p} \Big|_{\Delta\psi=0, \lambda=0} \sum_{a=k}^{nk} P^{(a)}(\psi^{(1)}, \dots, \psi^{(n)}) \lambda^{a+p} \Big|_{a+p=2n+1}. \quad (\text{B.17})$$

Only $\psi^{(1)}, \dots, \psi^{(n)}$ are present in this expression. This completes the proof. \square

C More on the Polarisability of Hydrogen

C.1 Rayleigh–Schrödinger Sum Involving Continuum States

The Rayleigh–Schrödinger sum of the polarisability of hydrogen atom is given by

$$\alpha = \sum_{k \neq 0} \frac{|\langle 0|z|k\rangle|^2}{E_k^{(0)} - E_0^{(0)}}. \quad (\text{C.1})$$

Here $|k\rangle$ should include both the bound states and the continuum states. Let's first consider the bound states, which can be labelled by $|n\ell m_\ell\rangle$. We can separate this integral into radial and angular parts,

$$\langle 0|z|n\ell m_\ell\rangle = \langle Y_{00}|\cos\theta|Y_{\ell m_\ell}\rangle \langle R_{1s}|r|R_{n\ell}\rangle, \quad (\text{C.2})$$

where $Y_{\ell m_\ell}$ are the spherical harmonics. Since

$$Y_{00} = \frac{1}{2}\sqrt{\frac{1}{\pi}}, \quad Y_{10} = \frac{1}{2}\sqrt{\frac{3}{\pi}}\cos\theta, \quad (\text{C.3})$$

we can evaluate the angular part to

$$\langle Y_{00}|\cos\theta|Y_{\ell m_\ell}\rangle = \frac{1}{\sqrt{3}} \langle Y_{10}|Y_{\ell m_\ell}\rangle = \frac{1}{\sqrt{3}}\delta_{1\ell}\delta_{0m_\ell} \quad (\text{C.4})$$

by the orthonormality of spherical harmonics. Thus we need the p_z orbitals only. We can also put in the energy expressions $E_0^{(0)} = -\frac{1}{2}$, $E_n^{(0)} = -\frac{1}{2n^2}$, giving

$$\alpha_{\text{bound}} = \frac{4}{3} \sum_{n=2}^{\infty} \frac{|\langle R_{1s}|r|R_{np}\rangle|^2}{1 - n^{-2}}. \quad (\text{C.5})$$

To proceed, we need the radial wavefunctions. In atomic units, it is given by

$$R_{n\ell}(r) = \sqrt{\left(\frac{2}{n}\right)^3 \frac{(n-\ell-1)!}{2n(n+1)!}} e^{-\frac{r}{n}} \left(\frac{2r}{n}\right)^\ell L_{n-\ell-1}^{(2\ell+1)}\left(\frac{2r}{n}\right), \quad (\text{C.6})$$

where $L_m^{(\alpha)}$ is the generalised (associated) Laguerre polynomial of degree m .⁴⁴ We will only need the 1s radial wavefunction

$$R_{1s}(r) = 2e^{-r} \quad (\text{C.8})$$

and the np wavefunction

$$R_{np}(r) = 4\sqrt{\frac{1}{n^7(n^2-1)}} r e^{-r/2} L_{n-2}^{(3)}\left(\frac{2r}{n}\right). \quad (\text{C.9})$$

There is no good reason to assume that we can treat this sum analytically, so we can do this numerically on a computer.

We see that the sum converges to something close to $\alpha_{\text{bound}} \approx 3.66326$. As stated in the main text, the sum converges slowly, and to the wrong value. We also need to include the contributions of the unbounded states with energies above 0. They correspond to ionised free electrons, and the energy becomes continuous rather than discrete, so they are also known as the continuum states.

⁴⁴It is defined as the power series solution of the differential equation

$$xy'' + (\alpha + 1 - x)y' + my = 0, \quad (\text{C.7})$$

where m is a non-negative integer (necessary for the solution to be a finite-term power series), and α can be any real number.

n	Contributions summed up to n
2	2.959621106
5	3.552510445
10	3.634363556
20	3.655782791
50	3.662031178
100	3.662948342
200	3.663180129
500	3.663245412
1000	3.663254768
2000	3.663257109
5000	3.663257765

Table 1: Numerical sum of the hydrogen polarisability including contributions up to np orbital.

A inspection on the Hamiltonian reveals that these continuum state should have the same angular dependence as the bound states, and the only way to get a positive energy from the expression is if $n^2 < 0$, so we would have a imaginary n . These observations suggest that we need to evaluate

$$\alpha_{\text{unbound}} = \frac{4}{3} \int_0^{\infty} dn \frac{|\langle R_{1s}|r|R_{np}\rangle|^2}{1 - n^{-2}}. \quad (\text{C.10})$$

This integral is over purely imaginary n , so a first step would be to change the integration variable to a real variable. The energy E is a good choice, so

$$\alpha_{\text{unbound}} = \frac{2}{3} \int_0^{\infty} dE \frac{|\langle R_{1s}|r|R_{Ep}\rangle|^2}{E + \frac{1}{2}}. \quad (\text{C.11})$$

However, we would still need to adapt the radial wavefunction for complex n , as the generalised polynomials are only defined for integer degree. Generalised Laguerre polynomials can be written as

$$L_a^{(b)}(x) = \frac{(a+b)!}{a!b!} {}_1F_1(-a, b+1, x), \quad (\text{C.12})$$

where ${}_1F_1$ is the confluent hypergeometric function of the first kind. It is defined for non-integer parameters,⁴⁵ and we can also replace the factorials by Gamma function. Therefore, the radial continuum wavefunctions can be written as

$$R_{E\ell}(r) = \sqrt{\frac{2k}{\pi}} e^{\frac{\pi}{2k}} \frac{|\Gamma(\ell+1+\frac{i}{k})|}{(2\ell+1)!} (2kr)^\ell e^{-ikr} {}_1F_1\left(\ell+1+\frac{i}{k}, 2\ell+2, 2ikr\right), \quad (\text{C.14})$$

where $k = \sqrt{2E}$ is the wavenumber. This expression is actually the same form as the bound state expression other than the normalisation factor. The key difference is the e^{-ikr} term. For the unbound states, the wavenumber is real, meaning a complex exponential, so oscillations of the wavefunction. For the bound states, the energy is negative so the wavenumber is complex, giving a real exponential that decays with r .

We are only interested in the $\ell = 1$ case, so the expression simplifies to

$$R_{Ep}(r) = \sqrt{\frac{2k}{r}} e^{\frac{\pi}{2k}} \left| \Gamma\left(2+\frac{i}{k}\right) \right| \frac{kr}{3} e^{-ikr} {}_1F_1\left(2+\frac{i}{k}, 4, 2ikr\right). \quad (\text{C.15})$$

Evaluating this integral numerically gives a continuum contribution $\alpha_{\text{unbound}} \approx 0.83674$.⁴⁶ Combining all of these together, we get $\alpha \approx 3.66326 + 0.83674 \approx 4.5$.

⁴⁵The confluent hypergeometric function (also known as the Kummer's function) is the solution to the power series solution to the differential equation

$$xy'' + (b-x)y' - ay = 0. \quad (\text{C.13})$$

We can see that if a is a non-positive integer, then this reduces to the differential equation defining the generalised Laguerre polynomials.

⁴⁶In practise, there are some numerical issues to worry about. The continuum wavefunctions oscillate forever and

C.2 Alternative Method Finding the Exact Hydrogen Polarisability

There is actually a way finding out the polarisability of hydrogen atom, originally due to A. Dalgarno and J.T. Lewis, not involving solving the first-order wavefunction analytically in the parabolic coordinates, or evaluating the complicated Rayleigh-Schrödinger sum numerically.

Suppose there is an operator \hat{X} such that $\hat{H}^{(1)} = [\hat{X}, \hat{H}^{(0)}]$, then

$$\begin{aligned}\langle m | \hat{H}^{(1)} | n \rangle &= \langle m | \hat{X} \hat{H}^{(0)} | n \rangle - \langle m | \hat{H}^{(0)} \hat{X} | n \rangle \\ &= (E_n^{(0)} - E_m^{(0)}) \langle m | \hat{X} | n \rangle.\end{aligned}\quad (\text{C.17})$$

We can then simplify the Rayleigh-Schrödinger sum of the second-order energy as

$$\begin{aligned}E_n^{(2)} &= - \sum_{m \neq n} \frac{\langle n | \hat{H}^{(1)} | m \rangle (E_n^{(0)} - E_m^{(0)}) \langle m | \hat{X} | n \rangle}{E_m^{(0)} - E_n^{(0)}} \\ &= \sum_{m \neq n} \langle n | \hat{H}^{(1)} | m \rangle \langle m | \hat{X} | n \rangle \\ &= \left\langle n \left| \hat{H}^{(1)} \left(\sum_m |m\rangle \langle m| \right) \hat{X} \right| n \right\rangle - \langle n | \hat{H}^{(1)} | n \rangle \langle n | \hat{X} | n \rangle \\ &= \langle n | \hat{H}^{(1)} \hat{X} | n \rangle - \langle n | \hat{H}^{(1)} | n \rangle \langle n | \hat{X} | n \rangle,\end{aligned}\quad (\text{C.18})$$

where we have cleverly used the resolution of identity to eliminate the sum over all states. Now the task reduces to finding the operator \hat{X} . It turns out that this is not difficult to do. We have

$$[\hat{X}, \hat{H}^{(0)}] |0\rangle = \hat{H}^{(1)} |0\rangle = \mathcal{E} z |0\rangle. \quad (\text{C.19})$$

This is just a differential equation that can be solved by separation of variable, and we have

$$\hat{X} = -\mathcal{E} \left(\frac{1}{2} r + 1 \right) z. \quad (\text{C.20})$$

The expectation values in (C.18) is easily calculated by exploiting spherical symmetry. The second term vanishes since $\langle n | \hat{H}^{(1)} | n \rangle = 0$, so we have

$$\begin{aligned}E_0^{(2)} &= \mathcal{E} \langle 0 | z \hat{X} | 0 \rangle \\ &= -\mathcal{E}^2 \left\langle 0 \left| \left(\frac{1}{2} r + 1 \right) z^2 \right| 0 \right\rangle \\ &= -\frac{1}{3} \mathcal{E}^2 \left\langle 0 \left| \left(\frac{1}{2} r + 1 \right) (x^2 + y^2 + z^2) \right| 0 \right\rangle \\ &= -\frac{1}{3} \mathcal{E}^2 \left[\frac{1}{2} \langle 0 | r^3 | 0 \rangle + \langle 0 | r^2 | 0 \rangle \right].\end{aligned}\quad (\text{C.21})$$

This will give us $E_0^{(2)} = -\frac{9}{4}$ and so $\alpha = \frac{9}{2}$ exactly.

to not converge to 0 as $r \rightarrow \infty$. However, the exponential decay of the 1s wavefunction do ensure a bounded result of $\mu_E = \langle R_{1s} | r | R_{Ep} \rangle$. We cannot evaluate μ_E at $E = 0$ since the wavefunction is undefined there, but it tends to 0 everywhere as $E \rightarrow 0$ from above. The increasing denominator of the energy also ensures the contribution decreases as the energy increases and the energy integral

$$\alpha_{\text{unbound}} = \frac{2}{3} \int_0^\infty dE \frac{\mu_E^2}{E + \frac{1}{2}} \quad (\text{C.16})$$

does converges. A lower limit of $r = 10^{-7} a_0$ and an upper limit of $r = 100 a_0$ would give sufficient precision, with the greater error coming from a lower bound above 0.

D First-Order Wavefunctions in the Degenerate Perturbation Theory

Following the main text, we have split the unperturbed wavefunctions into two subsets: $\{|\psi_i^{(0)}\rangle\}$ that are non-degenerate with the state of interest, and $\{|\Phi_{nj}^{(0)}\rangle\}$ that are degenerate with the state of interest. All of the states in the two sets combined are orthonormal, and we impose the intermediate normalisation such that

$$\langle \Phi_{nj}^{(0)} | \psi_{nj}^{(1)} \rangle = 0. \quad (\text{D.1})$$

We will split the first-order wavefunction into the contribution from the non-degenerate states labelled ND and the degenerate states labelled D

$$|\psi_{nj}^{(1)}\rangle = |\psi_{nj,\text{ND}}^{(1)}\rangle + |\psi_{nj,\text{D}}^{(1)}\rangle = \sum_{k \neq n} c_{k,\text{ND}} |\psi_k^{(0)}\rangle + \sum_{l \neq j} c_{l,\text{D}} |\Phi_{nl}^{(0)}\rangle. \quad (\text{D.2})$$

The first-order equation for a wavefunction in the degenerate set is

$$(\hat{H}^{(0)} - E_n^{(0)}) |\psi_{nj}^{(1)}\rangle + (\hat{H}^{(1)} - E_{nj}^{(1)}) |\Phi_{nj}^{(0)}\rangle = 0. \quad (\text{D.3})$$

If we contract this with $\langle \Phi_{nm}^{(0)} |$ for some m , we get expression of the first-order energy that we find in the main text. If we instead contract with $\langle \psi_i^{(0)} |$, we obtain

$$c_{i,\text{ND}} = \langle \psi_i^{(0)} | \psi_{nj}^{(1)} \rangle = - \frac{\langle \psi_i^{(0)} | \hat{H}^{(1)} | \Phi_{nj}^{(0)} \rangle}{E_i^{(0)} - E_n^{(0)}}, \quad (\text{D.4})$$

and so the non-degenerate contribution to the first-order wavefunction is

$$|\psi_{nj,\text{ND}}^{(1)}\rangle = - \sum_{k \neq n} \frac{H_{kj}^{(1)}}{E_k^{(0)} - E_n^{(0)}} |\psi_k^{(0)}\rangle \quad (\text{D.5})$$

as in the normal Rayleigh–Schrödinger sum.

To work out the degenerate coefficients, we need a bit more efforts. Consider the second-order equation

$$(\hat{H}^{(0)} - E_n^{(0)}) |\psi_{nj}^{(2)}\rangle + (\hat{H}^{(1)} - E_{nj}^{(1)}) |\psi_{nj}^{(1)}\rangle - E_{nj}^{(2)} |\Phi_{nj}^{(0)}\rangle = 0. \quad (\text{D.6})$$

Contracting with $\langle \Phi_{nm}^{(0)} |$ for some m gives

$$\begin{aligned} E_{nj}^{(2)} \delta_{jm} &= \langle \Phi_{nm}^{(0)} | \hat{H}^{(1)} - E_{nj}^{(1)} | \psi_{nj}^{(1)} \rangle \\ &= \sum_{k \neq n} c_{k,\text{ND}} \langle \Phi_{nm}^{(0)} | \hat{H}^{(1)} - E_{nj}^{(1)} | \psi_k^{(0)} \rangle + \sum_{l \neq j} c_{l,\text{D}} \langle \Phi_{nm}^{(0)} | \hat{H}^{(1)} - E_{nj}^{(1)} | \Phi_{nl}^{(0)} \rangle \\ &= \sum_{k \neq n} c_{k,\text{ND}} \langle \Phi_{nm}^{(0)} | \hat{H}^{(1)} | \psi_k^{(0)} \rangle + c_{m,\text{D}} (E_{nm}^{(1)} - E_{nj}^{(1)}) \end{aligned} \quad (\text{D.7})$$

using the fact that $\{|\Phi_{nl}^{(0)}\rangle\}$ diagonalises $\hat{H}^{(1)}$ in its span. If we choose $m = j$, the second term vanishes and we arrive at the second-order energy expression we claimed in the main text

$$E_{nj}^{(2)} = \sum_{k \neq n} c_{k,\text{ND}} \langle \Phi_{nj}^{(0)} | \hat{H}^{(1)} | \psi_k^{(0)} \rangle = - \sum_{k \neq n} \frac{|H_{jk}^{(1)}|^2}{E_k^{(0)} - E_n^{(0)}}. \quad (\text{D.8})$$

If we instead choose $j \neq m$, the term on the left vanishes and we get

$$c_{m,D} = -\frac{1}{E_{nm}^{(1)} - E_{nj}^{(1)}} \sum_{k \neq n} c_{k,ND} \left\langle \Phi_{nm}^{(0)} \left| \hat{H}^{(1)} \right| \psi_k^{(0)} \right\rangle = \frac{1}{E_{nm}^{(1)} - E_{nj}^{(1)}} \sum_{k \neq n} \frac{H_{mk}^{(1)} H_{kj}^{(1)}}{E_k^{(0)} - E_n^{(0)}}. \quad (\text{D.9})$$

This gives us the total expression of the first-order wavefunction

$$\left| \psi_{nj}^{(1)} \right\rangle = -\sum_{k \neq n} \frac{H_{kj}^{(1)}}{E_k^{(0)} - E_n^{(0)}} \left| \psi_k^{(0)} \right\rangle + \sum_{k \neq n} \sum_{l \neq j} \frac{H_{lk}^{(1)} H_{kj}^{(1)}}{(E_k^{(0)} - E_n^{(0)})(E_{nl}^{(1)} - E_{nj}^{(1)})} \left| \Phi_{nl}^{(0)} \right\rangle. \quad (\text{D.10})$$

E The Three Pictures

Our formulation of quantum mechanics so far is known as the *Schrödinger's picture*: a quantum state evolve with time following the Schrödinger equation

$$i\hbar \frac{d}{dt} |\psi\rangle = \hat{H} |\psi\rangle, \quad (\text{E.1})$$

and observables are associated with operators that (usually) do not change with time, such that the expectation value of a physical quantity A is given by

$$\langle A \rangle(t) = \left\langle \psi(t) \left| \hat{A} \right| \psi(t) \right\rangle. \quad (\text{E.2})$$

We shall have a bit more careful consideration on the role of time.

E.1 State Propagation

Any general state can be expanded as a linear combination of the eigenstates of \hat{H}

$$|\psi(t)\rangle = \sum_n c_n |n\rangle. \quad (\text{E.3})$$

We can also expand it in the eigenstates of any other Hermitian operators. Hence, the set of possible outcomes of a measurement and eigenstate of a observable do not change with time. It is the relative weights $c_n(t)$ that changes with time.

For now, let's assume that the Hamiltonian is time independent. Substituting the expansion (E.3) into the Schrödinger equation (E.1), we get

$$\begin{aligned} i\hbar \sum_n \frac{dc_n(t)}{dt} |n\rangle &= \sum_n c_n(t) \hat{H} |n\rangle, \\ i\hbar \frac{dc_n(t)}{dt} &= c_n(t) E_n \end{aligned} \quad (\text{E.4})$$

Solving this gives

$$c_n(t) = c_n(0) e^{-iE_n t/\hbar}. \quad (\text{E.5})$$

Therefore, for a time independent Hamiltonian, the state evolves as

$$|\phi(t)\rangle = \sum_n e^{-iE_n t/\hbar} c_n(0) |n\rangle, \quad (\text{E.6})$$

where it is common to denote $\omega_n := E_n/\hbar$. If only one frequency component is present, the energy is certain and the form of the states does not change with time — it is known as a stationary state.

The function of a operator is defined as

$$f(\hat{X}) |\psi_n\rangle = f(x_n) |\psi_n\rangle , \quad (\text{E.7})$$

where $\hat{X} |\psi_n\rangle = x_n |\psi_n\rangle$, and its action on a general state is defined by eigenstate expansion, so it has the ket-bra form

$$f(\hat{X}) = \sum_n f(x_n) |\psi_n\rangle \langle \psi_n| . \quad (\text{E.8})$$

Therefore, we can write

$$|\psi(t)\rangle = e^{-i\hat{H}t/\hbar} |\psi(0)\rangle . \quad (\text{E.9})$$

This is often denoted as the *time shift operator*

$$\hat{U}(t, t_0) = e^{-i\hat{H}(t-t_0)/\hbar} = \sum_n e^{-i\omega_n(t-t_0)} |n\rangle \langle n| \quad (\text{E.10})$$

so that

$$|\psi(t)\rangle = \hat{U}(t, t_0) |\psi(t_0)\rangle . \quad (\text{E.11})$$

It is easily verified that $\hat{U}(t - t_0)$ is unitary.

E.2 Time Dependent Hamiltonian

Now we allow the Hamiltonian to have time dependence. If we put $|\psi(t')\rangle = \hat{U}(t', t_0) |\psi(t_0)\rangle$ into the Schrödinger equation, we get

$$i\hbar \frac{d}{dt'} \hat{U}(t', t_0) = \hat{H}(t') \hat{U}(t', t_0) . \quad (\text{E.12})$$

We integrate both sides from t_0 to t and we get

$$\hat{U}(t, t_0) = 1 + \frac{-i}{\hbar} \int_{t_0}^t dt' \hat{H}(t') \hat{U}(t', t_0) , \quad (\text{E.13})$$

where we used $\hat{U}(t_0, t_0) = 1$. This can be solved by iteration, where we substitute the expression itself to $\hat{U}(t', t_0)$ on the right hand side repeatedly and get

$$\begin{aligned} \hat{U}(t, t_0) = & 1 + \frac{-i}{\hbar} \int_{t_0}^t dt_1 \hat{H}(t_1) + \left(\frac{-i}{\hbar} \right)^2 \int_{t_0}^t dt_1 \int_{t_0}^{t_1} dt_2 \hat{H}(t_1) \hat{H}(t_2) \\ & + \left(\frac{-i}{\hbar} \right)^2 \int_{t_0}^t dt_1 \int_{t_0}^{t_1} dt_2 \int_{t_0}^{t_2} dt_3 \hat{H}(t_1) \hat{H}(t_2) \hat{H}(t_3) + \dots \end{aligned} \quad (\text{E.14})$$

Since the Hamiltonian at one time does not commute with the Hamiltonian at a different time, the order $\hat{H}(t_1) \hat{H}(t_2) \hat{H}(t_3) \dots$ must be preserved.

Consider the double integral

$$\begin{aligned} \int_{t_0}^t dt_1 \int_{t_0}^{t_1} dt_2 \hat{H}(t_1) \hat{H}(t_2) &= \int_{t_0}^t dt_1 \int_{t_0}^t dt_2 \hat{H}(t_1) \hat{H}(t_2) \Theta(t_1 - t_2) \\ &= \int_{t_0}^t dt_1 \int_{t_0}^t dt_2 \hat{H}(t_2) \hat{H}(t_1) \Theta(t_2 - t_1) \\ &= \frac{1}{2} \int_{t_0}^t dt_1 \int_{t_0}^t dt_2 \overleftarrow{\mathcal{T}} [\hat{H}(t_2) \hat{H}(t_1)] , \end{aligned} \quad (\text{E.15})$$

where Θ is the Heaviside step function and we have introduced the *time ordering operator*

$$\overleftarrow{\mathcal{T}} [\hat{H}(t_2) \hat{H}(t_1)] = \begin{cases} \hat{H}(t_2) \hat{H}(t_1) & \text{if } t_2 \geq t_1 \\ \hat{H}(t_1) \hat{H}(t_2) & \text{if } t_2 < t_1 \end{cases} \quad (\text{E.16})$$

ordering the operator by time. Similarly, one can show that the n^{th} integral in E.14 is given by

$$\overleftarrow{\mathcal{T}} \left[\frac{1}{n!} \int_{t_0}^t dt_1 \int_{t_0}^t dt_2 \dots \int_{t_0}^t dt_n \hat{H}(t_1) \hat{H}(t_2) \dots \hat{H}(t_n) \right]. \quad (\text{E.17})$$

Therefore, for $t \geq t_0$, we can write (E.14) as

$$\begin{aligned} \hat{U}(t, t_0) &= \overleftarrow{\mathcal{T}} \left[\sum_{n=0}^{\infty} \left(\frac{-i}{\hbar} \right)^n \int_{t_0}^t dt_1 \int_{t_0}^t dt_2 \dots \int_{t_0}^t dt_n \hat{H}(t_1) \hat{H}(t_2) \dots \hat{H}(t_n) \right] \\ &= \overleftarrow{\mathcal{T}} \exp \left[\frac{-i}{\hbar} \int_{t_0}^t dt' \hat{H}(t') \right] \end{aligned} \quad (\text{E.18})$$

taking the Taylor expansion of exponential function to define the *time ordering exponential*. One way of thinking this is that time is divided into a number of discrete intervals, and the Hamiltonian is approximately constant during each of those intervals. The eigenvectors and eigenvalues of $\hat{H}(t')$ can then be used to propagate the states over a short period.

Similarly, one can show that when $t < t_0$, the operator propagating a state reverse in time is

$$\hat{U}(t, t_0) \overrightarrow{\mathcal{T}} \exp \left[\frac{-i}{\hbar} \int_{t_0}^t dt' \hat{H}(t') \right], \quad (\text{E.19})$$

where $\overrightarrow{\mathcal{T}}$ is the *anti-time ordering operator* ordering the time in reverse e.g.

$$\overrightarrow{\mathcal{T}} [\hat{H}(t_2) \hat{H}(t_1)] = \begin{cases} \hat{H}(t_1) \hat{H}(t_2) & \text{if } t_2 \geq t_1 \\ \hat{H}(t_2) \hat{H}(t_1) & \text{if } t_2 < t_1. \end{cases} \quad (\text{E.20})$$

One can show that \hat{U} is unitary and $\hat{U}^\dagger(t_2, t_1) = \hat{U}(t_1, t_2)$.

E.3 Heisenberg's Picture

There is another approach to quantum mechanics, which is analogous to the Schrödinger picture. Instead of putting the time dependency on the state, it puts all of time dependency on the operators.

In the Schrödinger's picture, the expectation value of any observable is given by

$$\langle A \rangle(t) = \langle \psi(t) | \hat{A}^S(t) | \psi(t) \rangle. \quad (\text{E.21})$$

We allow the operator to have time dependence (e.g. potential energy for a varying potential) as well as the time evolution of the states. Note that we have added a superscript S to explicitly denote the Schrödinger's picture.

However, we know that $\psi(t) = \hat{U}(t, t_0) |\psi(t_0)\rangle$, and so we may write

$$\langle A \rangle(t) = \langle \psi(t_0) | \hat{U}^\dagger(t, t_0) \hat{A}^S(t) \hat{U}(t, t_0) | \psi(t_0) \rangle. \quad (\text{E.22})$$

If we define the operator in *Heisenberg's picture* to be

$$\hat{A}^H(t) = \hat{U}^\dagger(t, t_0) \hat{A}^S(t) \hat{U}(t, t_0), \quad (\text{E.23})$$

then we may write

$$\hat{A}(t) = \langle \psi(t_0) | \hat{A}^H(t) | \psi(t_0) \rangle. \quad (\text{E.24})$$

As we promised, this gives an alternative view of quantum mechanics. The quantum state itself is not changing — it is always $|\psi(t_0)\rangle$, and it is the operator that is evolving in time.⁴⁷

Note that the two pictures coincide at $t = t_0$ because $\hat{U}(t_0, t_0) = 1$.

Operators in the Heisenberg picture satisfy a differential equation. Consider

$$\frac{d\hat{A}^H}{dt} = \frac{d}{dt}\hat{U}^\dagger \hat{A}^S \hat{U} = \frac{d\hat{U}^\dagger}{dt} \hat{A}^S \hat{U} + \hat{U}^\dagger \frac{d\hat{A}^S}{dt} \hat{U} + \hat{U}^\dagger \hat{A}^S \frac{d\hat{U}}{dt}. \quad (\text{E.25})$$

Since $i\hbar \frac{d}{dt} \hat{U} = \hat{H} \hat{U}$,

$$\begin{aligned} i\hbar \frac{d\hat{A}^H}{dt} &= -\hat{U}^\dagger \hat{H}^\dagger \hat{A}^S \hat{U} + i\hbar \hat{U}^\dagger \frac{d\hat{A}^S}{dt} \hat{U} + \hat{U}^\dagger \hat{A}^S \hat{H} \hat{U} \\ &= -\hat{U}^\dagger \hat{H}^\dagger \hat{U} \hat{U}^\dagger \hat{A}^S \hat{U} + i\hbar \hat{U}^\dagger \frac{d\hat{A}^S}{dt} \hat{U} + \hat{U}^\dagger \hat{A}^S \hat{U} \hat{U}^\dagger \hat{H} \hat{U} \\ &= [\hat{A}^H, \hat{H}^H] + i\hbar \hat{U}^\dagger \frac{d\hat{A}^S}{dt} \hat{U}. \end{aligned} \quad (\text{E.26})$$

If the operator \hat{A}^S is time independent in Schrödinger's picture, then the third term vanishes. If further the Hamiltonian is time independent, \hat{H} and \hat{U} commutes so that $\hat{H}^H(t) = \hat{H}$, giving

$$i\hbar \frac{d\hat{A}^H}{dt} = [\hat{A}^H, \hat{H}]. \quad (\text{E.27})$$

This is the *Heisenberg's equation*. It neatly shows that any operator that commutes with \hat{H} is a constant of motion.

E.4 The Interaction Picture

We now come to a particularly valuable tool: the *interaction picture*. It is called so because it was originally devised to handle scattering, but now it is used widely.

Let's write the Hamiltonian in the form

$$\hat{H}^S(t) = \hat{H}_0^S + \hat{V}^S(t). \quad (\text{E.28})$$

This would occur, for example, in a system where a collection of particle is subjected to a time dependent external force $\hat{V}^S(t)$. \hat{H}_0^S would then be the Hamiltonian of the free system, so it is called the *free Hamiltonian*.

We know that $|\psi^S(t)\rangle = \hat{U}(t, t_0) |\psi^S(t_0)\rangle$. We can unwrap the time-independent phase factor caused by \hat{H}_0 from the fully evolved state to define the state in the Interaction picture to be

$$\begin{aligned} |\psi^I(t)\rangle &= e^{+i\hat{H}_0(t-t_0)/\hbar} |\psi^S(t)\rangle \\ &= e^{+i\hat{H}_0(t-t_0)/\hbar} \hat{U}(t, t_0) |\psi^S(t_0)\rangle. \end{aligned} \quad (\text{E.29})$$

⁴⁷In fact, all this has a precise analogue in classical mechanics. Classically, there are also two ways of thinking about time evolution. On the one hand, we can think of a particle moving in some way through phase space M . If we know it's location $(\mathbf{x}(t), \mathbf{p}(t)) \in M$ for every time t by solving Newton's second law $\frac{d\mathbf{p}}{dt} = F(\mathbf{x}(t))$ from a initial condition $(\mathbf{x}(t_0), \mathbf{p}(t_0))$, we can compute any physical quantity we wish, represented by some function $f : M \rightarrow \mathbb{R}$, by evaluating $f(\mathbf{x}(t), \mathbf{p}(t))$. This also suggests a perspective in which the "state" of our particle is simply a choice of initial conditions. These initial conditions do not themselves evolve, rather, it is the quantities we measure that vary in time. Thus, instead of thinking of a physical quantity f as a map from phase space, we treat it just as a map from time, so $f : [t_0, \infty) \rightarrow \mathbb{R}$. You'll examine these classical pictures further if you study classical Hamiltonian mechanics.

We evolve the state from t_0 to t by the full Hamiltonian, and then evolve back from t to t_0 using the free Hamiltonian. Now the expectation value is

$$\begin{aligned}\langle A \rangle(t) &= \langle \psi^S(t) | \hat{A}^S(t) | \psi^S(t) \rangle \\ &= \langle \psi^I(t) | e^{+i\hat{H}_0(t-t_0)/\hbar} \hat{A}^S(t) e^{-i\hat{H}_0(t-t_0)/\hbar} | \psi^I(t) \rangle \\ &= \langle \psi^I(t) | \hat{A}^I(t) | \psi^I(t) \rangle ,\end{aligned}\tag{E.30}$$

where the operator in the interaction picture is

$$\hat{A}^I(t) = e^{+i\hat{H}_0(t-t_0)/\hbar} \hat{A}^S(t) e^{-i\hat{H}_0(t-t_0)/\hbar} .\tag{E.31}$$

Moreover, we have

$$\begin{aligned}i\hbar \frac{\partial}{\partial t} |\psi^I(t)\rangle &= i\hbar \frac{\partial}{\partial t} e^{+i\hat{H}_0(t-t_0)/\hbar} |\psi^S(t)\rangle \\ &= e^{+i\hat{H}_0(t-t_0)/\hbar} \left[-\hat{H}_0 + i\hbar \frac{\partial}{\partial t} \right] |\psi^S(t)\rangle \\ &= e^{+i\hat{H}_0(t-t_0)/\hbar} \left[-\hat{H}_0 + \hat{H}^S(t) \right] |\psi^S(t)\rangle \\ &= e^{+i\hat{H}_0(t-t_0)/\hbar} \hat{V}^S(t) e^{-i\hat{H}_0(t-t_0)/\hbar} |\psi^I(t)\rangle \\ &= \hat{V}^I(t) |\psi^I(t)\rangle .\end{aligned}\tag{E.32}$$

Therefore, in the interaction picture, the operators evolve according to the time-independent free part of the Hamiltonian, whereas states evolve according to the time-dependent interaction part of the Hamiltonian.

The solution to (E.32) can be written as

$$|\psi^I(t)\rangle = \hat{S}(t, t_0) |\psi^I(t_0)\rangle ,\tag{E.33}$$

where $\hat{S}(t, t_0)$ is the *scattering operator* given by

$$\hat{S}(t, t_0) = \overleftarrow{\mathcal{T}} \exp \left[\frac{-i}{\hbar} \int_{t_0}^t dt' \hat{V}^I(t') \right] ,\tag{E.34}$$

for $t \geq t_0$, while for $t < t_0$,

$$\hat{S}(t, t_0) = \overrightarrow{\mathcal{T}} \exp \left[\frac{-i}{\hbar} \int_{t_0}^t dt' \hat{V}^I(t') \right] .\tag{E.35}$$

F Time Dependent Perturbation Theory in the Interaction Picture

The formulism of the Interaction Picture makes the treatment of time-dependent perturbation theory particularly easy, because the Hamiltonian it considered (E.28) is essentially a time-dependent perturbation theory Hamiltonian

$$\hat{H} = \hat{H}^{(0)} + \hat{H}^{(1)}(t)\tag{F.1}$$

time-independent Hamiltonian $\hat{H}^{(0)} \equiv \hat{H}_0$ with a time-dependent perturbation $\hat{H}^{(1)}(t) \equiv \hat{V}(t)$. All we have to do is to ensure that perturbing Hamiltonian is small everything in our expansions later will converge.

We expand the perturbed state in the interaction picture using the eigenstates of the unperturbed system,

$$|\psi^I(t)\rangle = \sum_n a_n(t) |n\rangle ,\tag{F.2}$$

and substitution into the (E.32) gives

$$i\hbar \frac{da_m(t)}{dt} = \sum_n V_{mn}^I(t) a_n(t), \quad (\text{F.3})$$

where the matrix elements of the perturbation are

$$\begin{aligned} V_{mn}^I(t) &= \langle m | \hat{V}^I(t) | n \rangle \\ &= \langle m | e^{+i\hat{H}_0 t/\hbar} \hat{V}^S(t) e^{-i\hat{H}_0 t/\hbar} | n \rangle \\ &= e^{i\omega_{mn}t} V_{mn}^S(t), \end{aligned} \quad (\text{F.4})$$

where $\omega_{mn} = \omega_m - \omega_n$, and V_{mn}^S is $H_{mn}^{(1)}$ using the notation in the main text. Therefore, we have

$$i\hbar \frac{da_m(t)}{dt} = \sum_n e^{i\omega_{mn}t} V_{mn}^S(t) a_n(t), \quad (\text{F.5})$$

or equivalently in the matrix form

$$i\hbar \frac{d\mathbf{a}}{dt} = \mathbf{W} H^{(1)} \mathbf{W}^\dagger \mathbf{a}, \quad (\text{F.6})$$

where \mathbf{W} is the diagonal matrix of phase factors. Although this equation is exact, it is a coupled differential equation (of possibly infinite-dimensional), so it is rarely used.

What we rather do is to use the scattering operator

$$\begin{aligned} \hat{S}(t) &= \overleftarrow{\mathcal{T}} \exp \left[\frac{-i}{\hbar} \int_0^t dt' \hat{V}^I(t') \right] \\ &= \overleftarrow{\mathcal{T}} \left[\sum_{n=0}^{\infty} \left(-\frac{i}{\hbar} \right)^n \frac{1}{n!} \int_0^t dt_1 \dots \int_0^t dt_n \hat{V}^I(t_1) \dots \hat{V}^I(t_n) \right]. \end{aligned} \quad (\text{F.7})$$

Recall how we interpret this time-ordered exponential. It treats the effect of the perturbation as a sequence of impulses: we evolve according to $\hat{H}^{(0)}$ for some time $t_n \geq 0$ then feel the effect $\hat{V}(t_n)$ of the perturbation just at this time, before evolving according to H_0 for a further time $t_n - t_{n-1} \geq 0$ and feeling the next impulse from the perturbation, and so on. Finally, we integrate over all the possible intermediate times at which the effects of the perturbation were felt, and sum over how many times it acted. This method, closely related to Green's functions for ODEs and PDEs, is also the basis for Feynman diagrams in QFT.

Since the perturbation is ‘small’, the more times that the perturbation has acted on our system, the smaller the weight it should contribute to the time evolution of the system. Hence, to the first order, we can consider only the first term in the expansion of the scattering operator

$$\hat{S} = \hat{I} + \frac{-i}{\hbar} \int_0^t dt_1 \hat{V}^I(t_1). \quad (\text{F.8})$$

Now suppose at $t = 0$ we're in the state

$$|\psi(0)\rangle = \sum_n a_n(0) |n\rangle, \quad (\text{F.9})$$

then at a later time t the interaction picture state will be

$$|\psi^I(t)\rangle = \hat{S}(t) |\psi(0)\rangle \quad (\text{F.10})$$

and can again be described by a superposition $\sum_n a_n(t) |n\rangle$. Contracting with a state $\langle j|$ gives

$$a_j(t) = \langle j | \hat{S}(t) | \psi^I(0) \rangle \quad (\text{F.11})$$

in the interaction picture. Thus, using the scattering operator expanded to the first order (F.8), we have

$$\begin{aligned}
a_j(t) &\approx a_j(0) + \frac{-i}{\hbar} \sum_n a_n(0) \int_0^t dt' \langle j | \hat{V}^I(t) | n \rangle \\
&= a_j(0) + \frac{-i}{\hbar} \sum_n a_n(0) \int_0^t dt' \langle j | e^{+i\hat{H}_0 t'/\hbar} \hat{V}^S(t') e^{-i\hat{H}_0 t'/\hbar} | n \rangle \\
&= \underbrace{a_j(0)}_{a_j^{(0)}(t)} + \underbrace{\frac{-i}{\hbar} \sum_n a_n(0) \int_0^t dt' e^{i(E_j - E_n)t'/\hbar} H_{jn}^{(1)}(t')}_{a_j^{(1)}(t)}. \tag{F.12}
\end{aligned}$$

to the first non-trivial order. The first term is the unperturbed free evolution, and the second term is the first order perturbation we derived in the main text.

This can be trivially generalised to give the second-order time dependent perturbation theory — we just need to expand the scattering operator by one more term.

$$\hat{S} = \hat{I} - \frac{i}{\hbar} \int_0^t dt_1 \hat{V}^I(t_1) - \frac{1}{\hbar^2} \int_0^t dt_1 \int_0^{t_1} dt_2 \hat{V}^I(t_1) \hat{V}^I(t_2). \tag{F.13}$$

This will give an extra second order coefficient in our $a_j(t)$

$$a_j(t) \approx a_j^{(0)}(t) + a_j^{(1)}(t) + a_j^{(2)}(t), \tag{F.14}$$

where

$$\begin{aligned}
a_j^{(2)}(t) &= -\frac{1}{\hbar^2} \sum_n a_n(0) \int_0^t dt_1 \int_0^{t_1} dt_2 \langle j | \hat{V}^I(t_1) \hat{V}^I(t_2) | n \rangle \\
&= -\frac{1}{\hbar^2} \sum_n \sum_k a_n(0) \int_0^t dt_1 \int_0^{t_1} dt_2 \langle j | \hat{V}^I(t_1) | k \rangle \langle k | \hat{V}^I(t_2) | n \rangle \\
&= -\frac{1}{\hbar^2} \sum_n \sum_k a_n(0) \int_0^t dt_1 \int_0^{t_1} dt_2 e^{i\omega_{jk}t_1} e^{i\omega_{kn}t_2} H_{jk}^{(1)}(t_1) H_{kn}^{(1)}(t_2), \tag{F.15}
\end{aligned}$$

where we have inserted an identity operator. If we have $a_n(0) = 1$ for some particular n , and $a_m(0) = 0$ for all other $m \neq n$, i.e. we are initially at $|n\rangle$ at $t = 0$, then the above second order coefficient simplifies to

$$a_j^{(2)}(t) = -\frac{1}{\hbar^2} \sum_k \int_0^t dt'' \int_0^{t''} dt' e^{i\omega_{jk}t''} e^{i\omega_{kn}t'} H_{jk}^{(1)}(t'') H_{kn}^{(1)}(t'). \tag{F.16}$$

Under this framework, generalisation to higher order perturbations are trivial, although the integrals will be harder and harder to evaluate.

F.1 Raman Activity

Now we will use this to investigate Raman scattering. We do have the added complication that there are two interactions, corresponding to the ‘first’ photon and the ‘second’ photon (although thinking of these events happening sequentially is not very ‘quantum’) and the key feature of Raman scattering is that these two photons have different frequencies. Therefore we will write our perturbation matrix elements as

$$\hat{H}_{jk}^{(1)}(t') = V_{kn} \left(e^{i\omega' t'} + e^{-i\omega' t'} \right) \quad \hat{H}_{jk}^{(1)}(t'') = V_{jk} \left(e^{i\omega'' t''} + e^{-i\omega'' t''} \right), \tag{F.17}$$

where ω' is the frequency of the first photon and ω'' is the frequency of the second photon, and as previously⁴⁸

$$V_{kn} = -\frac{1}{2}\mathcal{E}\langle k|\hat{\mu}|n\rangle. \quad (\text{F.18})$$

With these definitions, we can perform the integrations directly, and we will get a large number of terms

$$\begin{aligned} a_j^{(2)}(t) = & -\sum_k V_{jk}V_{kn} \left[\frac{e^{i(\omega_{jn}+\omega'+\omega'')t}-1}{(\omega_{jn}+\omega'+\omega'')(\omega_{kn}+\omega')} - \frac{e^{i(\omega_{jk}+\omega'')t}-1}{(\omega_{jk}+\omega'')(\omega_{kn}+\omega')} \right. \\ & + \frac{e^{i(\omega_{jn}+\omega'-\omega'')t}-1}{(\omega_{jn}+\omega'-\omega'')(\omega_{kn}+\omega')} - \frac{e^{i(\omega_{jk}-\omega'')t}-1}{(\omega_{jk}-\omega'')(\omega_{kn}+\omega')} \\ & + \frac{e^{i(\omega_{jn}-\omega'+\omega'')t}-1}{(\omega_{jn}-\omega'+\omega'')(\omega_{kn}-\omega')} - \frac{e^{i(\omega_{jk}+\omega'')t}-1}{(\omega_{jk}+\omega'')(\omega_{kn}-\omega')} \\ & \left. + \frac{e^{i(\omega_{jn}-\omega'-\omega'')t}-1}{(\omega_{jn}-\omega'-\omega'')(\omega_{kn}-\omega')} - \frac{e^{i(\omega_{jk}-\omega'')t}-1}{(\omega_{jk}-\omega'')(\omega_{kn}-\omega')} \right]. \quad (\text{F.19}) \end{aligned}$$

Various of these terms correspond to interesting second-order effects, some of which are used in spectroscopic experiments, but if we are interested in non-resonant Raman scattering, we require $\omega_{jn} \approx \pm(\omega' - \omega'')$, so that the difference in energy between the initial and excited state is comparable to the difference in energy of the two photons, but there is no state at energy $E_n^{(0)} \pm \omega'$.⁴⁹ The plus corresponds roughly to absorption of the first photon and emission of the second photon. The less intuitive minus corresponds to stimulated emission caused by the first photon and absorption of the second photon.

With the non-resonant Raman condition, of the eight terms present in (F.19), it is the third and fifth that will dominate. Noting that ω_{jn} , ω' and ω'' do not depend on k , so can be taken outside the sums, we have

$$a_j^{(2)} = -\frac{e^{i(\omega_{jn}+\omega'-\omega'')t}-1}{(\omega_{jn}+\omega'-\omega'')(\omega_{kn}+\omega')} \sum_k \frac{V_{jk}V_{kn}}{\omega_{kn}+\omega'} - \frac{e^{i(\omega_{jn}-\omega'+\omega'')t}-1}{(\omega_{jn}-\omega'+\omega'')(\omega_{kn}-\omega')} \sum_l \frac{V_{jl}V_{ln}}{\omega_{ln}-\omega'}. \quad (\text{F.20})$$

Once again, we must take the square modulus of this expression to generate a transition probability. The resulting probability has four terms, but we can note that the cross terms, which can only simultaneously satisfy one of the plus or minus in the non-resonant Raman condition, will be much smaller than the diagonal terms and we will drop them, giving

$$P_j(t) = 4t \frac{\sin^2[\frac{1}{2}(\omega_{jn}+\omega'-\omega'')]t}{(\omega_{jn}+\omega'-\omega'')^2 t} \left| \sum_k \frac{V_{jk}V_{kn}}{\omega_{kn}+\omega'} \right|^2 + 4t \frac{\sin^2[\frac{1}{2}(\omega_{jn}-\omega'+\omega'')]t}{(\omega_{jn}-\omega'+\omega'')^2 t} \left| \sum_l \frac{V_{jl}V_{ln}}{\omega_{ln}+\omega'} \right|^2. \quad (\text{F.21})$$

Deploying the same tricks concerning integrating over a frequency distribution and tending to a δ function as we did with Fermi's golden rule, we obtain

$$R_{nj} = 2\pi\rho(\omega') \left(\left| \sum_k \frac{V_{jk}V_{kn}}{\omega_{kn}+\omega'} \right|^2 + \left| \sum_l \frac{V_{jl}V_{ln}}{\omega_{ln}+\omega'} \right|^2 \right). \quad (\text{F.22})$$

If we again assume that the cross terms being negligible, effectively undoing our previous approxima-

⁴⁸In the actual Raman, the only external photon source is the laser beam, so the second photon is actually emitted by spontaneous emission. However, we can't treat spontaneous emission without quantising the field, so we will model this as a stimulated process here.

⁴⁹The case of resonant Raman scattering, where there does exist a state close in energy to the initial state plus or minus the laser energy, such that $\omega_{kn} \approx \pm\omega'$ and $\omega_{jk} \approx \pm\omega''$, is more complicated and is not considered here.

tion, we can rewrite R_{nj} as

$$\begin{aligned}
R_{nj} &= 2\pi\rho(\omega') \left| \sum_k \frac{V_{jk}V_{kn}}{\omega_{kn} + \omega'} + \frac{V_{jk}V_{kn}}{\omega_{kn} - \omega'} \right|^2 \\
&= 2\pi\rho(\omega') \left| \sum_k \frac{(\omega_{kn} - \omega')V_{jk}V_{kn} + (\omega_{kn} + \omega')V_{jk}V_{kn}}{\omega_{kn}^2 - \omega'^2} \right|^2.
\end{aligned} \tag{F.23}$$

This sum is related to (5.57) for the dynamic polarisability. The difference is that (5.57) refers specifically to the polarisability evaluated for a single quantum state, $\langle n|\hat{\alpha}|n\rangle$, whereas here it is the extension to a transition polarisability $\langle j|\hat{\alpha}|n\rangle$.

$$\langle j|\hat{\alpha}_{\beta\gamma}(\omega')|n\rangle = \sum_k \frac{(\omega_{kn} - \omega') \langle j|\hat{\mu}_\beta|k\rangle \langle k|\hat{\mu}_\gamma|n\rangle + (\omega_{kn} + \omega') \langle j|\hat{\mu}_\gamma|k\rangle \langle k|\hat{\mu}_\beta|n\rangle}{\omega_{kn}^2 - \omega'^2}. \tag{F.24}$$

We can see that this reduces to (5.57) in the case of $|j\rangle = |n\rangle$. Thus, we can write

$$R_{nj} \propto |\langle j|\hat{\alpha}(\omega')|n\rangle|^2. \tag{F.25}$$

Note that while we normally think about the static polarisability for Raman activity, it is in fact the dynamic polarisability at the laser frequency that matters, although the distinction has no consequences if we are only interested in the symmetry argument of whether or not the integral is non-zero.

The inclusion of the intermediate states $|k\rangle$ is what gives the common interpretation of there existing a transitory virtual state in Raman scattering, although we see that interpretation is only loosely correct. It is not that the system moves to one virtual state, but rather that it includes all possible states, although with a greater contribution from those with energies closer to the initial energy plus or minus the laser energy.