

**USEFUL MATHEMATICS**

You can cancel out any 0 and factor out all constants. The trig function are:

$$\sin(x) = O/H$$

$$\cos(x) = A/H$$

$$\tan(x) = O/A$$

For all real  $x$ ,  $x^2 \geq 0$ , so we invent  $i$  such that  $i^2 = -1$ . The Gaussian is

$$P(x) = \frac{1}{\sqrt{2\pi}\Delta} e^{-(x-x_0)^2/2\Delta^2}$$

Integrating in polar coordinates we find

$$\int_{-\infty}^{\infty} dx e^{-ax^2} = \sqrt{\frac{\pi}{a}}$$

Differentiating under the integral sign

$$\int_{-\infty}^{\infty} dx x^{2n} e^{-ax^2} = \left(\frac{-d}{da}\right)^n \sqrt{\frac{\pi}{a}}$$

Integrating directly we find

$$\int_{-\infty}^{\infty} dx e^{-ax} = \frac{1}{a}$$

Integrating by parts is very useful

$$\int_a^b fg'dx = \int_a^b fdg = - \int_a^b gdf + fg|_a^b$$

For example, it gives us the  $\Gamma$  function

$$\Gamma(n+1) = \int_{-\infty}^{\infty} dx x^n e^{-ax} = \frac{n!}{a^{n+1}}$$

Decomposing into even/odd functions

$$f(x) = \frac{f(x) + f(-x)}{2} + \frac{f(x) - f(-x)}{2}$$

We then have  $\int_{-a}^a dx \text{ odd}(x) = 0$  and

$$\int_{-a}^a dx \text{ even}(x) = 2 \int_0^a dx \text{ even}(x)$$

Completing the square can also help

$$3x^2 - 12x = 3((x-2)^2 - 4)$$

The chain rule is also essential

$$\frac{d}{dx}(x \sin(x^2)) = \sin(x^2) + 2x^2 \cos(x^2)$$

The  $u$ -Substitution helps as well

$$\int_1^4 dx 6xe^{-x^2} = \int_1^{16} du 3e^{-u}$$

We can do a  $u$ -substitution like

$$P_u(u)du = P_x(x)dx$$

$$n(u)du = n(E)d(E)$$

**PHYSICAL CONSTANTS**

$$q_e = 1.602 \cdot 10^{-19} \text{ C}$$

$$m_e = 9.109 \cdot 10^{-31} \text{ kg} = 511 \text{ keV}$$

$$k_B = 8.617 \cdot 10^{-5} \text{ eV/K}$$

$$\hbar = 10.55 \cdot 10^{-34} \text{ J s}$$

**PHYSICS 1 KNOWLEDGE**

The units of your answer should be sensible. Energy, momentum, and angular momentum are conserved. We have

$$\mathbf{F} = m\mathbf{a}$$

The work is

$$W = \int \mathbf{F} \cdot d\mathbf{l}$$

The kinetic energy is

$$K = \frac{1}{2}m|\mathbf{v}|^2 = \frac{|\mathbf{p}|^2}{2m}$$

and the Hamiltonian is

$$H = K + U$$

For a spring-mass system

$$\mathbf{F} = k\mathbf{x} \implies \omega = \sqrt{k/m} = 2\pi f$$

For relativistic motion we have

$$K = \sqrt{p^2c^2 + m_0^2c^4} - m_0c^2$$

$$pc = \sqrt{E^2 - m_0^2c^4} = \sqrt{K^2 + 2Km_0c^2}$$

The Lorentz force law is

$$\mathbf{F} = q(\mathbf{E} + \mathbf{v} \times \mathbf{B})$$

In a capacitor we have

$$|\mathbf{E}| = V/d$$

For a wave we have

$$f = c/\lambda$$

Waves can be added, leading to beats

$$\psi_{\text{total}}(x, t) = \psi_1(x, y) + \psi_2(x, t)$$

**PROBABILITY DISTRIBUTIONS**

The expectation value of  $A$  is

$$\langle A \rangle = \int_{-\infty}^{\infty} dx P(x)A(x)$$

In radial coordinates this is

$$\langle A \rangle = \int_0^{\infty} dr P(r)A(r)$$

The probability is normalized if

$$\int_{\mathcal{V}} P(V) dV = 1$$

Physically,  $\psi(x)$  meaningless, but  $P(x) = \psi^*(x)\psi(x) = |\psi(x)|^2$  is meaningful. In radial coordinates we have  $dP(r)/dr = |\psi|^2 A$ , or  $P(r) = r^2 |R(r)|^2$ . From this we insist that  $\psi$  be finite and

$$\int_{-\infty}^{\infty} |\psi(x)|^2 dx = 1$$

The "uncertainty" in a quantity is

$$\Delta q \equiv \sqrt{\langle q^2 \rangle - \langle q \rangle^2}$$

Uncertainty (Cauchy-Schwarz)

$$\Delta p \Delta x \geq \hbar/2$$

$$\Delta E \Delta t \geq \hbar/2$$

**OPERATORS**

We have the operators

$$\hat{x} = x = i\hbar \frac{\partial}{\partial p}$$

where  $\hat{x}$  and  $\hat{p}$  are canonical conjugates

$$\hat{p} = p = -i\hbar \frac{\partial}{\partial x}$$

The Hamiltonian is

$$\hat{H} = \hat{K} + \hat{U}$$

Which is composed of kinetic energy

$$\hat{K} = \frac{\hat{p}^2}{2m} = -\frac{\hbar^2}{2m} \frac{\partial^2}{\partial x^2}$$

and potential energy

$$\hat{U} = U(x)$$

The energy generates time translation

$$\hat{E} = i\hbar \frac{\partial}{\partial t}$$

**SCHRÖDINGER EQUATION**

The Schrödinger equation is

$$\hat{H}\psi = \hat{E}\psi$$

With  $\hat{E} = i\hbar \partial_t$  we have

$$\hat{H}\psi = i\hbar \frac{\partial \psi}{\partial t}$$

For  $E_n$  an eigenvalue of  $\hat{H}$  we have the time-independent Schrödinger equation (solutions have  $\psi_n(t) = \psi_n(0)e^{-iE_n t/\hbar}$ )

$$-\frac{\hbar^2}{2m} \frac{\partial^2 \psi_n(x)}{\partial x^2} + U(x)\psi_n(x) = E_n \psi_n(x)$$

Solving we have ( $\alpha = \sqrt{2m(U-E_n)/\hbar}$ )

$$\psi_n(x) = Ae^{-\alpha x} + Be^{\alpha x}$$

In terms of trig functions ( $k = i\alpha$ )

$$\psi_n(x) = C \cos(kx) + D \sin(kx)$$

This solution holds in regions where  $U$  is constant. The boundaries conditions are that  $\psi$  and its first derivative are continuous (except when  $U$  is infinite). We can superimpose solutions

$$|\psi_1 + \psi_2|^2 = (\psi_1 + \psi_2)(\psi_1 + \psi_2)^*$$

which results in time-dependent beats.

**INFINITE SQUARE WELL**

We consider the potential

$$U(x) = \begin{cases} 0 & 0 \leq x \leq L \\ \infty & \text{otherwise} \end{cases}$$

For which the energy eigenvalues are

$$E_n = n^2 \frac{\pi^2 \hbar^2}{2mL^2}$$

Which correspond to eigenfunctions

$$\psi_n(x) = \sqrt{2/L} \sin(n\pi x/L)$$

### HARMONIC OSCILLATOR

We consider the potential

$$U(x) = \frac{1}{2}kx^2 = \frac{1}{2}m\omega^2x^2$$

For which the energy eigenvalues are

$$E_n = (n + \frac{1}{2})\hbar\omega$$

with eigenfunctions ( $a = \sqrt{\hbar/m\omega}$ )

$$\psi_n(x) = \frac{1}{\sqrt{2^n n! \sqrt{\pi a^2}}} e^{-x^2/2a^2} H_n\left(\frac{x}{a}\right)$$

Where the Hermite polynomials are

$$H_0(x) = 1$$

$$H_1(x) = 2x$$

$$H_2(x) = 4x^2 - 2$$

$$H_n(x) = (-1)^n e^{x^2} \frac{d^n}{dx^n} (e^{-x^2})$$

And the ground state wavefunction is

$$\psi_0(x) = \frac{1}{\sqrt{\sqrt{\pi a^2}}} e^{-x^2/2a^2}$$

### 3D INFINITE SQUARE WELL

In 3D,  $\partial_{xx}^2 \mapsto \nabla^2$  so that the time-independent Schrödinger equation is

$$-\frac{\hbar^2}{2m} \nabla^2 \psi_n(\mathbf{r}) + U(\mathbf{r})\psi_n(\mathbf{r}) = E_n \psi_n(\mathbf{r})$$

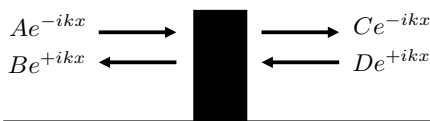
For a right parallelepiped we have

$$E_{ijk} = \frac{\pi^2 \hbar^2}{2m} \left[ \left(\frac{n_i}{L_1}\right)^2 + \left(\frac{n_j}{L_2}\right)^2 + \left(\frac{n_k}{L_3}\right)^2 \right]$$

The linear momentum of the states is

$$|p_i| = n_i \frac{\pi \hbar}{L_i}$$

### QUANTUM TUNNELING



Reflection and transmission coefficients

$$\mathcal{R} \equiv \frac{|B|^2}{|A|^2}, \quad \mathcal{T} \equiv \frac{|C|^2}{|A|^2}$$

Where there is no accumulation so

$$\mathcal{R} + \mathcal{T} = 1$$

We can approximate the transmission

$$\mathcal{T}(E) \approx e^{-\frac{2\sqrt{2m}}{\hbar} \int dx \sqrt{U(x)-E}}$$

Where we define the penetration depth

$$\delta = \hbar / \sqrt{2m(U-E)}$$

The energy levels of a finite well are

$$E_n \approx n^2 \frac{\pi^2 \hbar^2}{2m(L+2\delta)^2}$$

### QUANTUM NUMBERS

Quantum numbers are for counting to index eigenstates (usually nodes of the wavefunction). In hydrogenic atoms (and electronic orbitals in atoms)

- Principle,  $n = 1, 2, 3, \dots$
- Angular momentum,  $l = 0, \dots, n-1$   
 $\circ l=0 \leftrightarrow s, 1 \leftrightarrow p, 2 \leftrightarrow d, 3 \leftrightarrow f$
- Magnetic,  $m_l = -l, \dots, l-1, l$
- Spin,  $s = -S, -S+1, \dots, S$

For electrons and nucleons  $S = 1/2$ .

Ex. The  $4F_{\frac{5}{2}} e^-$  is  $n = 4, l = 3, j = \frac{5}{2}$ .

Pauli exclusion principle: No two fermions may occupy an orbital with the same  $n, l, m_l, m_s$  numbers.

### ANGULAR MOMENTUM

Where the total angular momentum is

$$j = l + s$$

Length of angular momentum vector

$$|\mathbf{J}| = \sqrt{j(j+1)}\hbar$$

The total orbital angular momentum

$$|\mathbf{L}| = \sqrt{l(l+1)}\hbar$$

The angular momentum along  $\hat{z}$  is

$$L_z = m_l \hbar$$

Which leads to “space quantization”

$$\cos(\theta) = \frac{L_z}{|\mathbf{L}|}$$

The spherical wave functions can be decomposed in the complete basis

$$\psi_{nlm_l}(r, \theta, \phi) = R_{nl}(r) Y_l^{m_l}(\theta, \phi)$$

Hund’s rules for orbital occupation

1. Maximize the total spin
2. Maximize  $\mathbf{J}$
3. Maximize  $\mathbf{L}$

### WAVE-PARTICLE DUALITY

Light has momentum

$$p_\gamma = \frac{h}{\lambda}$$

and energy

$$E_\gamma = hf = \frac{hc}{\lambda} = \frac{1240 \text{ eV nm}}{\lambda}$$

Which motivates wave-particle duality and the de Broglie wavelength

$$\lambda = \frac{h}{p} = \frac{hc}{pc}$$

### OPTICAL EVIDENCE FOR QUANTIZATION

Wein Law approximation

$$\lambda_{\max} = \frac{0.002898}{T}$$

Plank’s spectral radiance formula

$$u(f, T) = \frac{8\pi h f^3}{c^3} \left( e^{\frac{hf}{k_B T}} - 1 \right)^{-1}$$

Energy quantization from a cavity

$$E = nhf = n\hbar\omega$$

Photoelectric effect with workfunction

$$K_{\max} = hf - \phi_{\text{WF}}$$

### ELECTRONIC EVIDENCE FOR QUANTIZATION

Bragg law for constructive interference

$$n\lambda = 2d \sin(\theta)$$

Compton scattering relation

$$\Delta\lambda = \frac{h}{m_e c} (1 - \cos(\theta))$$

Thompson’s cathode ray experiment

$$\frac{e}{m_e} = \theta \left( \frac{V}{d} \right) \left( \frac{1}{B^2 l} \right)$$

Millikan oil drop experiment

$$ne = \frac{mg}{E} \left( \frac{v_{\text{ter}} + v_{\text{up}}}{v_{\text{ter}}} \right)$$

### HYDROGEN

Hydrogen has the Coulomb potential and energy levels ( $k = 1/4\pi\epsilon_0$ )

$$E_n = -\frac{ke^2 Z^2}{2a_0 n^2} = -\frac{13.6 Z^2}{n^2} \text{ eV}$$

which have negative energies and are bound states. Real eigenfunctions can always be found for bound states.

Stationary states of hydrogenic atoms

$$\psi(r, \theta, \phi, t) = R_{nl}(r) Y_l^{m_l}(\theta, \phi) e^{-i\omega t}$$

where  $R_{nl}$ , and  $Y_l^{m_l}$  are tabulated.

The “radius” of wavefunctions is

$$r_n = \frac{a_0 n^2}{Z}$$

where  $a_0$  is the Bohr radius

$$a_0 = \frac{\hbar^2}{m_e k e^2} = 0.0529 \text{ nm}$$

### QUANTUM COMBINATORICS

Quantum effects onset at high density

$$\left(\frac{N}{L^{\text{dim}}}\right) \left(\frac{\hbar/2}{\sqrt{mk_B T}}\right)^{\text{dim}} \approx 1$$

Then we sum rather than integrate

$$E_{\text{tot}} = \sum_i n_i E_i$$

Where the number in in state  $i$  is

$$n_i(E_i) = g(E_i) f(E_i)$$

$g_i$  is the density of states and  $f_i$  is the distribution function. In continuum

$$n(E)dE = g(E)f_{\text{dist}}(E)dE$$

### DISTRIBUTION FUNCTIONS

Classical (distinguishable), ex. gasses

$$f_{\text{MB}}(E_i, T) = \left(e^{\frac{E_i}{k_B T}}\right)^{-1}$$

Bosons ( $S = \text{integer}$ , indistinguishable), ex. photons, phonons, gluons

$$f_{\text{BE}}(E_i, T) = \left(e^{\frac{E_i}{k_B T}} - 1\right)^{-1}$$

Fermions ( $S = \text{integer} + \frac{1}{2}$ , indistinguishable), ex. electrons, neutrinos

$$f_{\text{FD}}(E_i, T) = \left(e^{\frac{E_i}{k_B T}} + 1\right)^{-1}$$

### MAXWELL-BOLTZMANN GAS

The speed distribution of a gas is

$$n(v)dv = \frac{4\pi N}{V} \left(\frac{m}{2\pi k_B T}\right)^{\frac{3}{2}} v^2 e^{-\frac{mv^2}{2k_B T}} dv$$

We can then find the mean using

$$\langle v \rangle = \frac{\int_0^\infty dv v n(v)}{\int_0^\infty dv n(v)}$$

or the root-mean-square (RMS) using

$$\sqrt{v^2} = \sqrt{\frac{\int_0^\infty dv v^2 n(v)}{\int_0^\infty dv n(v)}}$$

or the extremal points using

$$\frac{d(n(v))}{dv} = 0$$

### BLACKBODY RADIATION

The spectra radiance is given by

$$u(E)dE = E n(E)dE = \frac{g_\gamma(E)EdE}{e^{\frac{E}{k_B T}} - 1}$$

The total number with frequency is

$$N(f)df = \frac{8\pi f^2 df}{c^3} = \frac{8\pi e^2 dE}{(hc)^3} = g_\gamma dE$$

So we have density of states

$$g_\gamma(E) = \frac{8\pi E^2}{(hc)^3}$$

### EINSTEIN HEAT CAPACITY

The specific heat capacity is defined by

$$C \equiv \frac{dU}{dT}$$

Where for classical materials have

$$U = 3N_A k_B T = 3RT$$

Which gives  $C = 3R$ , corresponding to one  $R$  per degree of freedom. If instead we model the energy in a solid as being sequestered solely in phonons then

$$\langle E \rangle = \frac{\hbar\omega}{e^{\frac{\hbar\omega}{k_B T}} - 1}$$

Or for a macroscopic sample of solid

$$U = 3N_A \langle E \rangle$$

Which gives us the specific heat

$$C = 3R \left(\frac{\hbar\omega}{k_B T}\right)^2 \frac{e^{\hbar\omega/k_B T}}{(e^{\hbar\omega/k_B T} - 1)^2}$$

Where the Einstein temperature is

$$T_E = \frac{\hbar\omega}{k_B}$$

### FREE ELECTRON GAS

For a free-electron gas we have

$$E = \frac{|\mathbf{p}|^2}{2m_e} = \frac{\hbar^2 |\mathbf{k}|^2}{2m_e}$$

Which means that

$$d|\mathbf{k}| = \frac{1}{2} \left(\frac{2m_e}{\hbar^2}\right)^{\frac{1}{2}} E^{-\frac{1}{2}} dE$$

So the density of states is then

$$g(E) = \frac{8\sqrt{2}\pi m_e^{\frac{3}{2}}}{h^3} E^{\frac{1}{2}}$$

Which means the number at  $E$  is

$$n(E)dE = \frac{8\sqrt{2}\pi m_e^{\frac{3}{2}}}{h^3} \frac{E^{\frac{1}{2}} dE}{e^{(E-\mu)/k_B T} + 1}$$

So the chemical potential  $T = 0$  is

$$E_F = \mu(0) = \frac{\hbar^2}{2m_e} \left(\frac{3N}{8\pi V}\right)^{\frac{2}{3}}$$

The Fermi velocity is given by

$$\frac{1}{2} m_e v_F^2 = E_F$$

And we have the Fermi temperature

$$T_F \equiv \frac{E_F}{k_B}$$

And Fermi wavenumber

$$|\mathbf{k}_F| = \sqrt{2m_e E_F}/\hbar$$

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