Quantum harmonic oscillator (in 1D)

A harmonic oscillator is a system with a quadratic potential:

$$V(x) = \frac{Kx^2}{2}$$
 (k is sometimes referred to as
"spring" or "force" constant)

Such a simple potential has great importance in physics and quantum mechanics in particular. The reason for this is that this potential can serve as the first approximation (which suffices in many practical applications) to a more realistic interaction, provided that the amplitude of the motion is small enough. Let us consider a particle trapped hear the minimum of a smooth potential. A smooth potential can be approximated as a Taylor series

$$V(x-x_{o}) = V(x_{o}) + V'(x_{o})(x-x_{o}) + \frac{1}{2!}V''(x_{o})(x-x_{o})^{2} + ...$$

The first term in this expansion is just a constant. It results in a trivial shift of the total energy, The second term, $V'(x_0)$, vanishes as we are at a point of minimum. This leaves only terms proportional to $(x-x_0)^2$, $(x_0-x_0)^3$, and so on If $x-x_0$ is small then we can ignore the higher powers of $x-x_0$.

Now the point of the minimum, Xo, Cay be shifted with a simple transformation $X-X_0 \rightarrow X'$ Therefore we will assume for simplicity $X_0=0$. Then $V(X) \approx \frac{V'(0)X^2}{2} = \frac{kX^2}{2} = \frac{m\omega^2 X^2}{2} \qquad \omega = \sqrt{\frac{k}{m}}$

The time idependent Schrödinger equation
for 1D hormonic oscillator takes the form
$$-\frac{\pi^2}{2m}\frac{d^2\psi}{dx^2} + \frac{1}{2}m\omega^2x^2\psi = E\psi$$

It is convenient to introduce the natural scale
for X in this problem. We set
 $\xi \equiv dx = \sqrt{\frac{m\omega}{\pi}} \times$ where $\frac{1}{d}$ is the natural
length scale
Let us also define new "energy" expressed in
units of $\frac{1}{2}tw$:
 $E = \frac{2E}{tw}$
The Schrödinger equation then reads
 $-\frac{d^2\psi}{d\xi^2} + \xi^2\psi = E\psi$ or $\frac{d^2\psi}{dx^2} = (\xi^2 - E)\psi$
In order to simplify things a little we can
unke a substitution
 $\psi(\xi) = h(\xi) = \frac{\xi^2}{2}$
This substitution is unofivated by the fact that
at large ξ the Schrödinger equation above becomes
 $\psi^{\parallel} = \xi^2\psi$ and the solution of that is $\psi z he^{\frac{\xi}{2}} + Be^{\frac{\xi}{2}}$
 $function at $\xi \to \infty$
After the substitution $\psi = he^{\frac{\xi^2}{2}}$ we obtain i
 $\frac{d^2h(\xi)}{d\xi^2} - 2\xi \frac{dh(\xi)}{d\xi} + (E-1)h = 0$ — the Hermite
equation$

The solutions to Hermite's equation can be oftained by employing different techniques. The one we will use below is called the power Series method. In this method we represent h(s) as a power series: $h(\xi) = a_0 + a_1 \xi + a_2 \xi^2 + \dots = \sum_{i=n}^{2} a_i \xi^i$ Its derivatives are $h'(s) = a_1 + 2a_2s + 3a_3s^2 + \dots = \sum_{j=1}^{\infty} ja_j s^{j-1} = \sum_{j=0}^{\infty} ja_j s^{j-1}$ $h''(s) = 2a_2 + 6a_3s + 12a_4s^2 + ... = \sum_{j=2}^{\infty} (j-1)ja_js^{j-2}$ $= \sum_{j=0}^{\infty} (j+1)(j+2) a_{j+2} \xi^{j}$ Plugging these into the equation for h(s) yields $\sum_{j=0}^{\infty} \left[(j+1)(j+2)a_{j+2} - 2ja_j + (\epsilon-1)a_j \right] \xi^j = 0$ Now, this relation must be true for any value of §. Therefore we must require that each coefficient before S vanish $(j+1)(j+2)a_{j+2} - 2ja_j + (\epsilon-1)a_j = 0$ more specifically, $\xi^{\circ}: 2\alpha_{2} + (\varepsilon - 1)\alpha_{\circ} = 0$ $\xi': 2.3a_3 + (\epsilon - 1 - 2)a_1 = 0$ $\xi^{2} = 3.4a_{4} + (\epsilon^{-1-4})a_{2} = 0$

and so on

We can see that $a_{j+2} = \frac{(2j+1-\epsilon)}{(j+1)(j+2)} a_j \leftarrow recursion relation$ The recursion formula relates all even coefficients to an and all odd coefficients to a. The even and odd coefficients are independent of each other As the potential is symmetric (even) in x, i.e. V(-x) = V(x), the eigenstates must be either even or odd functions. Indeed, this can be proven as follows. Suppose 4(x) is a solution of the stationary SE with energy E: $-\frac{\hbar^2}{2m}\frac{\partial^2\Psi(x)}{\partial x^2} + V(x)\Psi(x) = E\Psi(x)$ $-\frac{t^2}{2m}\frac{\partial^2\psi(-x)}{\partial x^2} + V(X)\psi(-x) = E\psi(-x) \qquad [we used V(-x) = V(x)]$ which we can also write as So if $\psi(x)$ is a solution then $\psi(-x)$ is also a solution. The solution of the SE corresponding to a particular energy E is unique up to an arbitrary phase factor. This implies that $\Psi(x) = C \Psi(-x)$ where C is a constant $\Psi(-x) = C \Psi(x) = C \cdot C \Psi(-x) \implies C^2 = 1 \implies C = \pm 1$ Bu + Hence 4(x) must be either even or odd if V(x)=V(-x) Let us now go fack to our harmonic oscillator. For even eigenstates $a_1 = 0$, $a_0 = 1$ (or any other) nonzero value; since the wave function has to be normalized we do not need to worry about the actual value of as now)

For odd eigenstates we choose $a_0 = 0$, $a_1 = 1$. If j is large we have $\frac{a_{j+2}}{a_j} = \frac{2}{j}$ $\frac{a_{j+2}}{a_j} = \frac{2}{j}$ Now recall that $e^{s^2} = 1 + s^2 + \frac{s^4}{2!} + \frac{s^6}{3!} + \dots$ The coefficient of § in the expansion of est is $b_j = \frac{1}{\binom{j}{2}!} \quad \text{and} \quad \frac{b_{j+2}}{b_j} = \frac{\binom{j}{2}!}{\binom{j}{2}+1!} = \frac{1}{j+1} \xrightarrow{>} \frac{2}{j} \quad \text{when} \quad j \to \infty$ Therefore h(z) goes as e^{z^2} if the power series is infinite. This is a disaster because they $\Psi(\xi) = e^{-\frac{\xi^2}{2}}h(\xi) = e^{+\frac{\xi^2}{2}} \rightarrow \infty$ when $\xi \rightarrow \infty$ For hormalizable states this not really an option. Hence, the only way to keep the wave function hormalizable is to require that the series is finite (e.g. h(z) is a polynomial). The series will termihate if E - 1 - 2h = 0This condition gives us the quantization of energy. Only those values of the energy are allowed, which Satisfy the condition $E = \frac{t_{10}}{2} \epsilon = t_{10} \left(N + \frac{1}{2} \right) \quad \text{where} \quad h = 0, 1, 2, \dots \infty$ These are the energy levels for a harmonic oscillator Determining the corresponding wave functions is straightforward. For the allowed E's the recursion formula is $a_{j+2} = -\frac{2(n-j)}{(j+1)(j+2)}a_j$ When n=0 there is only one term:

$$h(\xi) = \alpha_0 \implies \psi_0(\xi) = \alpha_0 e^{-\frac{\xi^2}{2}}$$

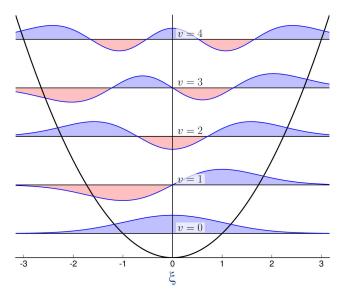
When h=1 we take a = 0 (symmetry consideration) and

$$\begin{split} h_1(\xi) &= a_1 \xi \implies \psi_1(\xi) = a_1 \xi e^{-\frac{\xi^2}{2}} \\ \text{When } h = 2 \quad \text{af } j = 0 \quad \text{we get } a_2 = -2a_0, \quad \text{so} \\ h_2(\xi) &= a_0(1 - 2\xi^2) \implies \psi_2(\xi) = a_0(1 - 2\xi^2) e^{-\frac{\xi^2}{2}} \\ \text{By continuing this way we can determine } \psi_n \\ \text{for any } h \quad \text{In general,} \\ \psi_n(\xi) &= \left(\frac{m\omega}{\pi t}\right)^{N_4} \frac{1}{\sqrt{2^4 n!}} \quad H_n(\xi) e^{-\frac{\xi^2}{2}} \end{split}$$

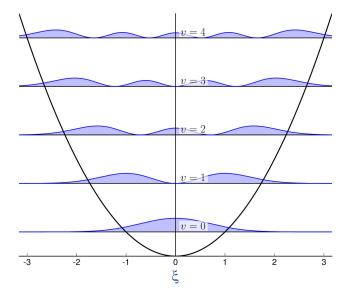
Polynomials H_h(S) are called the Hermite polynomials. They can be generated using the Rodrigues formula:

$$H_{h}(\xi) = (-1)^{h} e^{\xi^{2}} \left(\frac{d}{d\xi}\right)^{h} e^{-\xi^{2}}$$

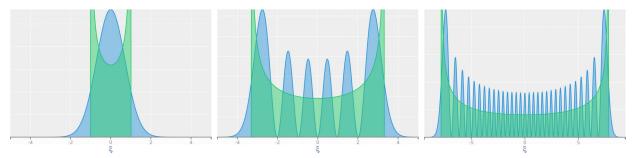
$$H_{0}(\xi) = 1 \qquad H_{1}(\xi) = 2\xi \qquad H_{2}(\xi) = e^{\xi^{2}} \frac{d^{2}}{d\xi^{2}} e^{\xi^{2}} = -2 + 4\xi^{2}$$



Harmonic oscillator wavefunctions, $\psi_n(\xi)$ for n = 0, 1, 2, 3, 4



Harmonic oscillator probability distributions, $|\psi_n(\xi)|^2$ for n = 0, 1, 2, 3, 4



Comparison of quantum (blue) and classical (green) probability distributions for n = 0, 5, 30