

Time evolution of a shifted initial state:

Let $\phi(x, t)$ be the time evolution of an initial function $\phi(x)$ in the field of the harmonic oscillator. For $x_0 \in \mathbb{R}$ define the shifted function by $\phi_{x_0}(x) = \phi(x - x_0)$. Its time evolution is given by

$$\phi_{x_0}(x, t) = \exp\left(i \frac{x_0^2}{4} \sin 2t\right) e^{-i(x_0 \sin t)x} \phi(x - x_0 \cos t, t) \quad (7.81)$$

We can draw some interesting conclusions from this result: Assume that $\phi = \phi_n$ is an eigenstate of the harmonic oscillator. The eigenstates have a trivial time evolution, they always remain centered at the origin, the expectation values of the position x and momentum p are zero for all times. Now, let's shift the eigenfunction to a new position x_0 , that is, the new initial state is given by

$$\psi(x, 0) = \phi_n(x - x_0). \quad (7.82)$$

Equation (7.81) gives the following result for the time evolution of the translated eigenstate:

$$\psi(x, t) = \exp\left(i \frac{x_0^2}{4} \sin 2t - iE_n t - i(x_0 \sin t)x\right) \phi_n(x - x_0 \cos t). \quad (7.83)$$

In particular, the position probability density is just given by

$$|\psi(x, t)|^2 = |\phi_n(x - x_0 \cos t)|^2. \quad (7.84)$$

Time evolution does not change the shape of $|\phi_n|^2$, it just translates the function to its classical position $x_0 \cos t$.



CD 5.19 shows the motion of the initially translated eigenstates ϕ_1 , ϕ_2 , and their superposition. The evolution of the shifted initial state shows the motion of $\phi_1 + \phi_2$ as in CD 5.5, while the center of the wave packet performs the oscillation $x_0 \cos t$.

7.5. Motion of Gaussian Wave Packets

7.5.1. Coherent states

Here we apply the results of the previous section to the ground state $\phi_0(x) = \pi^{(-1/4)} \exp(-\frac{1}{2}x^2)$. Among the eigenstates it is distinguished also by the property that it is optimal with respect to the uncertainty relation

$$\Delta x \Delta p \geq \frac{1}{2}. \quad (7.85)$$

The ground state satisfies

$$\Delta x \Delta p = \frac{1}{2}. \quad (7.86)$$

For the time evolution of the initially translated ground state we can apply the results obtained in the previous section. We obtain

$$\psi(x, t) = \left(\frac{1}{\pi}\right)^{1/4} \exp\left(i \frac{x_0^2}{4} \sin 2t - i \frac{t}{2}\right) \exp\left(i p_t x - \frac{(x - x_t)^2}{2}\right), \quad (7.87)$$

with $x_t = x_0 \cos t$ and $p_t = -x_0 \sin t$. This is a normalized Gaussian function centered at the average position x_t and with average momentum p_t . Here (x_t, p_t) describes the classical oscillation of a particle with initial position x_0 (in dimensionless units).



CD 5.10 contains movies of a coherent state in position space, momentum space, and in the energy representation. See also Color Plate Fi:HOcoh.

We know already that the function $\psi(x, t)$ is again optimal with respect to the uncertainty relation (see Section 2.8.1). The state $\psi(x, t)$ hence satisfies Eq. (7.86) for all times. The states with minimal uncertainty are called *coherent states*. Their motion is most similar to the oscillation of a particle in classical mechanics. We collect these observations in the following box.

Coherent states of a harmonic oscillator:

The coherent states (states with minimal uncertainty) of a harmonic oscillator are shifted Gaussian functions, initially localized at x_0 , which have the shape of the ground state. The maximum of a coherent state always follows the trajectory of a classical-mechanical particle that starts at x_0 with zero initial momentum. The wavelength of the phase always corresponds to the momentum of the classical particle. During its time evolution a coherent state retains its shape (the shape of the ground state) without spreading.

It is a consequence of Exercise 7.10 that the coherent states minimize the difference between the mean energy and the energy of the classical motion of $\langle x(t) \rangle_\psi$ and $\langle p(t) \rangle_\psi$ because the coherent states have minimal uncertainty. One even has the following result: ψ is a coherent state if and only if

$$\frac{1}{2} \left(\langle p \rangle_{\psi(t)}^2 + \langle x \rangle_{\psi(t)}^2 \right) = \frac{1}{2} \langle p^2 + x^2 \rangle_{\psi(t)}. \quad (7.88)$$

7.5.2. Arbitrary Gaussian function

For the harmonic oscillator potential, the Schrödinger equation with the Gaussian initial function

$$\phi(x) = \left(\frac{a}{\pi}\right)^{1/4} \exp\left(-a \frac{x^2}{2}\right) \quad (7.89)$$

has the solution

$$\psi(x, t) = \left(\frac{a}{\pi}\right)^{1/4} (\cos t + ia \sin t)^{-1/2} \exp\left(-a(t) \frac{x^2}{2}\right), \quad (7.90)$$

where

$$a(t) = \frac{a \cos t + i \sin t}{\cos t + ia \sin t}. \quad (7.91)$$

This result can be obtained using the explicitly known integral kernel of the time evolution. This integral kernel (Mehler's kernel) will be obtained in a later section.

One has to be careful with the definition of the square root

$$(\cos t + ia \sin t)^{-1/2} \quad (7.92)$$

in Eq. (7.90). It is necessary to take that branch of the square root that gives a continuous dependence on t . Hence for $a = 1$ this expression will simply become $\exp(-it/2)$.

By combining Eq. (7.90) with Eq. (7.81) one can easily find an expression for the time evolution of a translated Gaussian function.

7.6. Harmonic Oscillator in Two and More Dimensions

The wave function of a particle in two dimensions depends on a two-dimensional position variable $\mathbf{x} = (x_1, x_2)$. The Hamiltonian operator for the harmonic oscillator in two dimensions can be written as a sum of one-dimensional Hamiltonians

$$\begin{aligned} H &= -\frac{1}{2} \Delta + \frac{\mathbf{x} \cdot \mathbf{x}}{2} \\ &= -\frac{1}{2} \frac{d^2}{dx_1^2} + \frac{x_1^2}{2} - \frac{1}{2} \frac{d^2}{dx_2^2} + \frac{x_2^2}{2} \\ &= H_{x_1} + H_{x_2}, \end{aligned}$$

where H_{x_i} ($i = 1, 2$) acts only on the variable x_i . Hence we can make the same observations as in the case of free particles: The Schrödinger equation in two space dimensions can be solved by a product ansatz

$$\psi(\mathbf{x}, t) = \psi_1(x_1, t) \psi_2(x_2, t), \quad (7.93)$$

where each ψ_i is a solution of the one-dimensional harmonic oscillator equation.

$$\begin{aligned} i \frac{d}{dt} \psi &= i \frac{d\psi_1}{dt} \psi_2 + i\psi_1 \frac{d\psi_2}{dt} \\ &= (H_{x_1} \psi_1) \psi_2 + \psi_1 (H_{x_2} \psi_2) \\ &= (H_{x_1} + H_{x_2}) \psi_1 \psi_2 = H\psi. \end{aligned}$$

If $\psi_{n,m}$ is a product of two eigenfunctions ϕ_n and ϕ_m of the one-dimensional Hamiltonian,

$$\psi_{n,m}(\mathbf{x}) = \phi_n(x_1) \phi_m(x_2), \quad (7.94)$$

then $\psi_{n,m}(\mathbf{x})$ is an eigenfunction of H belonging to the eigenvalue $E_{n,m}$

$$E_{n,m} = E_m + E_n = m + n + 1, \quad H \psi_{n,m} = E_{n,m} \psi_{n,m}. \quad (7.95)$$

Hence it is clear that

$$\psi_{n,m}(\mathbf{x}, t) = \psi_{n,m}(\mathbf{x}) e^{-iE_{n,m}t} = \phi_n(x_1, t) \phi_m(x_2, t) \quad (7.96)$$

is a solution of the time-dependent Schrödinger equation. This solution is a product of solutions of one-dimensional equations for x_1 - and x_2 -coordinates, respectively. Of course, this does not mean that *every* solution of the two-dimensional oscillator is a product of one-dimensional solutions. Nevertheless, any solution can be written as an (infinite) linear combination of products as follows.

As a consequence of the fact that the functions ϕ_n form an orthonormal basis in the Hilbert space $L^2(\mathbb{R})$ it can be shown that the product functions $\psi_{n,m}$ form an orthonormal basis in $L^2(\mathbb{R}^2)$ (this is a property of the tensor product of Hilbert spaces). Hence every initial function $\phi \in L^2(\mathbb{R}^2)$ can be expanded as

$$\phi(\mathbf{x}) = \sum_{n=0}^{\infty} \sum_{m=0}^{\infty} c_{n,m} \psi_{n,m}(\mathbf{x}), \quad (7.97)$$

and the unique solution of the Schrödinger equation with initial condition $\psi(\mathbf{x}, t) = \phi(\mathbf{x})$ is given by

$$\psi(\mathbf{x}, t) = \sum_{n=0}^{\infty} \sum_{m=0}^{\infty} c_{n,m} e^{-i(n+m+1)t} \psi_{n,m}(\mathbf{x}). \quad (7.98)$$

The ground-state energy is $E_0 + E_0 = 1$ and every solution is periodic with period 2π .

EXERCISE 7.12. *Generalize the above considerations to n -dimensions.*

If $\psi(x, t)$ is the time evolution of a one-dimensional harmonic oscillator, then its Fourier transform, the function $\hat{\psi}(k, t)$, is also a solution of the

Schrödinger equation of the harmonic oscillator in one dimension. Hence the wave function in phase space,

$$\Psi(x, k, t) = \psi(x, t) \hat{\psi}(k, t) \quad (7.99)$$

is a solution of the two-dimensional oscillator equation.



CD 5.14–5.18 show solutions of the two-dimensional harmonic oscillator. Among the various Gaussian wave packets, the coherent and squeezed states are of particular interest.

7.7. Theory of the Harmonic Oscillator

7.7.1. Supersymmetry

Define the operators

$$A = \frac{1}{\sqrt{2}} (x + ip) = \frac{1}{\sqrt{2}} \left(\frac{d}{dx} + x \right), \quad (7.100)$$

$$A^\dagger = \frac{1}{\sqrt{2}} (x - ip) = \frac{1}{\sqrt{2}} \left(-\frac{d}{dx} + x \right). \quad (7.101)$$

Here $p = -id/dx$ is the momentum operator, and x denotes the position operator (the operator of multiplication of $\psi(x)$ by x). For suitable (differentiable) wave functions you can see by a partial integration that

$$\langle \phi, A\psi \rangle = \langle A^\dagger \phi, \psi \rangle. \quad (7.102)$$

Therefore, the operators A and A^\dagger are formally adjoint to each other.

Ψ Schwartz space. For the mathematical investigation it is necessary to have a dense domain in the Hilbert space $L^2(\mathbb{R})$, which is invariant under the action of the operators A and A^\dagger . Such a domain is the Schwartz space $\mathcal{S} = \mathcal{S}(\mathbb{R})$. The set \mathcal{S} consists of infinitely differentiable functions. The functions (and all their derivatives) are required to go to zero faster than any inverse power of $|x|$, as $|x|$ tends to infinity. More precisely,

$$\mathcal{S} = \left\{ f \in C^\infty(\mathbb{R}) \mid \text{for all integers } k, l: \sup_{x \in \mathbb{R}} \left| x^k \frac{d^l}{dx^l} f(x) \right| < \infty \right\}. \quad (7.103)$$

Typical examples of functions in \mathcal{S} are Gauss functions $\exp(-ax^2)$, and all the oscillator eigenfunctions. Because all finite linear combinations of functions in \mathcal{S} are again contained in \mathcal{S} , the set \mathcal{S} is a linear subspace of the Hilbert space $L^2(\mathbb{R})$. Theorem 2.2 in Section 2.7 states that the linear subspace spanned by the functions

$$G_q(x) = \left(\frac{1}{\pi} \right)^{1/4} e^{iqx} \exp\left(-\frac{x^2}{2}\right) \quad (7.104)$$

3. Wave function in momentum space
4. Wave function in phase space

CD 5.5. Oscillating state 1+2

1. Wave function in position space
2. Position probability density
3. Wave function in momentum space
4. Wave function in phase space

CD 5.6. Superposition of three eigenstates ($0 + 1 + 2$)

1. Wave function in position space
2. Position probability density
3. Wave function in momentum space

CD 5.7. Linear combination of three eigenstates ($0 + 1 - 2$)

1. Wave function in position space
2. Position probability density

CD 5.8. Superposition of ten eigenstates

1. Sum of first ten eigenstates
2. Boltzmann distribution of energies
3. Random phases

CD 5.9. Eigenfunction expansion

1. Build a Gaussian function (interactive experiment)
2. Energy representation
3. Energy representation of a shifted state
4. Time evolution in energy space

CD 5.10. Coherent state (shifted eigenstate)

1. Wave function in position space
2. Position probability density
3. Wave function in momentum space
4. Energy representation
5. Time period and gauge

CD 5.11. Squeezed state (widespread initial localization)

1. Wave function in position space
2. Position probability density
3. Wave function in momentum space

CD 5.12. Squeezed state (sharper initial localization)

1. Wave function in position space
2. Position probability density
3. Momentum space

CD 5.13. Step function

1. Approximation in position space

2. Position probability density
3. Momentum space

CD 5.14. Coherent state

1. Phase space motion
2. Circular motion in two dimensions
3. Elliptic motion in two dimensions

CD 5.15. Diagonal motion (surface plots)

1. Coherent state in two dimensions
2. Squeezed state in two dimensions (flat)
3. Squeezed state in two dimensions (sharp)

CD 5.16. Particle at rest

1. Centered squeezed state
2. Asymmetric centered state
3. Squeezed state in phase space

CD 5.17. Phase space motion (contour plots)

1. Squeezed state in phase space
2. Closer to the center
3. Very flat Gaussian state
4. Sharp Gaussian peak

CD 5.18. Circular motion (surface plots)

1. Circular motion of Gaussian
2. Another initial condition
3. Yet another initial condition (“the swimmer”)

CD 5.19. Constant shapes (time evolution of shifted eigenstates)

1. Translated first eigenstate
2. Second eigenstate
3. First+second eigenstate

CD 5.20. Gallery of angular momentum eigenstates

6. Special Systems

CD 6.1. Free Fall

1. Dropping a particle
2. Throwing a particle

CD 6.2. Free fall

1. Exact solution in a linear potential

CD 6.3. Free fall in two dimensions

1. Letting a particle drop vertically
2. Vertical throw of a particle