

Two Particles in an Infinite Well Potential

Julian Avila, Laura Herrera, Sebastian Rodríguez

Universidad Distrital Francisco José de Caldas.

Abstract

Keywords:

1 Problem Setting

We consider two indistinguishable particles, each confined to a one-dimensional infinite potential well of length L . The single-particle Hilbert space is $\mathcal{H}_i \cong L^2([0, L])$. The total Hilbert space for the non-relativistic system is the tensor product of the individual spaces:

$$\mathcal{H} = \mathcal{H}_1 \otimes \mathcal{H}_2. \quad (1)$$

The configuration space is $(x_1, x_2) \in [0, L] \times [0, L]$. The potential imposes Dirichlet boundary conditions, requiring the wavefunction $\Psi(x_1, x_2)$ to vanish at the boundaries.

The system's dynamics are governed by the Hamiltonian \hat{H} , which we decompose into kinetic, external potential, and interaction terms:

$$\hat{H} = \hat{T} + \hat{V}_{\text{ext}} + \hat{V}_{\text{int}}. \quad (2)$$

The total kinetic energy $\hat{T} = \hat{T}_1 + \hat{T}_2$ is the sum of the single-particle operators,

$$\hat{T}_1 = \frac{\hat{p}_1^2}{2m} \otimes \hat{I}, \quad (3)$$

$$\hat{T}_2 = \hat{I} \otimes \frac{\hat{p}_2^2}{2m}, \quad (4)$$

where \hat{I} is the identity operator on the single-particle space.

The external potential \hat{V}_{ext} is zero within the well and infinite otherwise, a constraint already enforced by the boundary conditions. The particles interact via a contact

potential \hat{V}_{int} , which is proportional to a Dirac delta function:

$$\hat{V}_{\text{int}}(x_1, x_2) = g \delta(x_1 - x_2). \quad (5)$$

Here, g represents the coupling strength of the interaction.

In the position representation, the Hamiltonian operator acts on the wavefunction as

$$\hat{H} = -\frac{\hbar^2}{2m} (\partial_1^2 + \partial_2^2) + g \delta(x_1 - x_2). \quad (6)$$

The configuration space is the square $[0, L] \times [0, L]$, and the interaction \hat{V}_{int} is active only along the diagonal $x_1 = x_2$.

2 Symmetry and Indistinguishability

The indistinguishability of the particles implies a fundamental symmetry. We introduce the particle exchange operator \hat{P}_{12} , whose action on the two-particle wavefunction is defined as

$$\hat{P}_{12}\Psi(x_1, x_2) = \Psi(x_2, x_1). \quad (7)$$

This operator commutes with the Hamiltonian, $[\hat{P}_{12}, \hat{H}] = 0$, as both the kinetic term and the interaction term are symmetric under the exchange $x_1 \leftrightarrow x_2$. This commutation is a crucial property: it ensures that the exchange symmetry of a state is conserved over time. Consequently, eigenstates of \hat{H} can be chosen as simultaneous eigenstates of \hat{P}_{12} with eigenvalues $p_{12} = \pm 1$.

- **Bosons (Symmetric):** $p_{12} = +1$. $\Psi_S(x_1, x_2) = \Psi_S(x_2, x_1)$.
- **Fermions (Antisymmetric):** $p_{12} = -1$. $\Psi_A(x_1, x_2) = -\Psi_A(x_2, x_1)$.

The Spin-Statistics Theorem connects this symmetry to the particle's intrinsic spin. In this work, we restrict our analysis to the fermionic case, requiring the total wavefunction to be antisymmetric under particle exchange.

The state vector must therefore belong to the antisymmetric subspace $\mathcal{H}_A \subset \mathcal{H}$. For a state constructed from two distinct single-particle orbitals, $|\phi_a\rangle$ and $|\phi_b\rangle$, the normalized antisymmetric state is

$$|\Psi_A\rangle = \frac{1}{\sqrt{2}} (|\phi_a\rangle \otimes |\phi_b\rangle - |\phi_b\rangle \otimes |\phi_a\rangle). \quad (8)$$

In this notation, the first ket in each product refers to particle 1 and the second to particle 2.

Projecting eq. (8) into the position basis ($\Psi_A(x_1, x_2) = \langle x_1, x_2 | \Psi_A \rangle$) yields the Slater determinant for the wavefunction:

$$\Psi_A(x_1, x_2) = \frac{1}{\sqrt{2}} (\phi_a(x_1)\phi_b(x_2) - \phi_b(x_1)\phi_a(x_2)). \quad (9)$$

Note that if $|\phi_a\rangle = |\phi_b\rangle$, the state vanishes, in accordance with the Pauli Exclusion Principle.

3 Position Representation of the Schrödinger Equation

The dynamics are governed by the time-dependent Schrödinger equation (TDSE). In the position representation, using the Hamiltonian from eq. (6), this reads:

$$i\hbar\partial_t\Psi(x_1, x_2, t) = \left(-\frac{\hbar^2}{2m}(\partial_{x_1}^2 + \partial_{x_2}^2) + g\delta(x_1 - x_2)\right)\Psi(x_1, x_2, t). \quad (10)$$

A central consequence of the fermionic symmetry, discussed in section 2, is the antisymmetry of the wavefunction: $\Psi(x_1, x_2, t) = -\Psi(x_2, x_1, t)$. This requirement has a profound effect on the interaction term. If we evaluate the wavefunction along the diagonal $x_1 = x_2 = x$, the antisymmetry implies

$$\Psi(x, x, t) = -\Psi(x, x, t) \implies \Psi(x, x, t) = 0. \quad (11)$$

The wavefunction must be identically zero for any configuration where the two particles are at the same position.

Because the delta-function potential $\hat{V}_{\text{int}} = g\delta(x_1 - x_2)$ has support only on this diagonal (where the wavefunction vanishes), the interaction term has no effect on the system. We can confirm this by examining the expected value of the interaction potential:

$$\begin{aligned} \langle \hat{V}_{\text{int}} \rangle &= \langle \Psi | g\delta(x_1 - x_2) | \Psi \rangle \\ &= g \int_0^L \int_0^L \Psi^*(x_1, x_2) \Psi(x_1, x_2) \delta(x_1 - x_2) dx_2 dx_1 \\ &= g \int_0^L \Psi^*(x_1, x_1) \Psi(x_1, x_1) dx_1 \\ &= g \int_0^L |0|^2 dx_1 = 0. \end{aligned} \quad (12)$$

Therefore, for fermions, the problem simplifies remarkably. The system behaves as two non-interacting identical particles in an infinite well, and the governing equation reduces to

$$i\hbar\partial_t\Psi(x_1, x_2, t) = -\frac{\hbar^2}{2m}(\partial_{x_1}^2 + \partial_{x_2}^2)\Psi(x_1, x_2, t). \quad (13)$$

We seek stationary-state solutions by applying the separation of variables, positing an ansatz of the form

$$\Psi(x_1, x_2, t) = \psi(x_1, x_2)\varphi(t). \quad (14)$$

Substituting this into eq. (13) and dividing by $\Psi(x_1, x_2, t)$ separates the spatial and temporal components:

$$\frac{1}{\psi(x_1, x_2)} \left[-\frac{\hbar^2}{2m} (\partial_{x_1}^2 + \partial_{x_2}^2) \right] \psi(x_1, x_2) = i\hbar \frac{1}{\varphi(t)} D_t \varphi. \quad (15)$$

The left side depends only on position and the right side only on time, so both must equal a separation constant, which we identify as the total energy E . This yields two independent equations: the Time-Independent Schrödinger Equation (TISE)

$$\left[-\frac{\hbar^2}{2m} (\partial_{x_1}^2 + \partial_{x_2}^2) \right] \psi(x_1, x_2) = E\psi(x_1, x_2), \quad (16)$$

and the temporal equation

$$i\hbar D_t \varphi = E\phi(t). \quad (17)$$

3.1 Time Component

The solution to the temporal equation, eq. (17), is straightforward,

$$\varphi(t) = e^{-iEt/\hbar}, \quad (18)$$

where we have set the initial phase $\phi(0) = 1$. The full time-dependent solution for a stationary state is thus $\Psi(x_1, x_2, t) = \psi(x_1, x_2)e^{-iEt/\hbar}$. The time evolution manifests only as a global phase, which is unobservable.

3.2 Spatial Component and Energy

We now solve the spatial TISE, eq. (16). Since the Hamiltonian is a sum of non-interacting single-particle Hamiltonians ($\hat{H} = \hat{H}_1 + \hat{H}_2$), we can construct the solution from the single-particle energy eigenfunctions.

The normalized stationary states for a single particle in the well are

$$\phi_n(x) = \sqrt{\frac{2}{L}} \sin\left(\frac{n\pi x}{L}\right), \quad n = 1, 2, 3, \dots \quad (19)$$

These are eigenfunctions of the single-particle Hamiltonian, $\hat{H}_i \phi_n(x_i) = E_n \phi_n(x_i)$, with corresponding energy eigenvalues

$$E_n = \frac{n^2 \pi^2 \hbar^2}{2mL^2}. \quad (20)$$

As required for fermions, we construct the two-particle spatial wavefunction as the normalized Slater determinant

$$\psi(n_1, n_2; x_1, x_2) = \frac{1}{\sqrt{2}} [\phi_{n_1}(x_1)\phi_{n_2}(x_2) - \phi_{n_1}(x_2)\phi_{n_2}(x_1)] \quad (21)$$

which is valid for $n_1 \neq n_2$, in accordance with the Pauli Exclusion Principle.

We verify this is an eigenfunction of the total spatial Hamiltonian $\hat{H} = \hat{H}_1 + \hat{H}_2$:

$$\begin{aligned}\hat{H}\psi &= (\hat{H}_1 + \hat{H}_2) \frac{1}{\sqrt{2}} [\phi_{n_1}(x_1)\phi_{n_2}(x_2) - \phi_{n_1}(x_2)\phi_{n_2}(x_1)] \\ &= (E_{n_1} + E_{n_2})\psi_{n_1, n_2}.\end{aligned}\tag{22}$$

By comparing this with the TISE, $\hat{H}\psi = E\psi$, we identify the total energy of the system as the sum of the single-particle energies:

$$E_{n_1, n_2} = E_{n_1} + E_{n_2} = \frac{\pi^2 \hbar^2}{2mL^2}(n_1^2 + n_2^2).\tag{23}$$

The prefactor $1/\sqrt{2}$ in eq. (21) ensures the state is normalized ($\langle\psi|\psi\rangle = 1$) due to the orthonormality of the single-particle orbitals ϕ_n .